MICROCOPY RESOLUTION TEST CHART
NATIONAL BUREAU OF STANDARDS
STANDARD REFERENCE MATERIAL 1010a
(ANSI and ISO TEST CHART No. 2)
A method of systematically controlling endoparasites and ectoparasites of warm-blooded animals by orally or percutaneously administering to the animal a parasiticidally effective amount of an acyl urea compound.
Use of Acyl Urea Compounds for Controlling Endoparasites and Ectoparasites of Warm-Blooded Animals

This application is a continuation-in-part of U.S. Patent Application Serial No. 723,588, filed April 15, 1985, which is a continuation-in-part of U.S. Patent Application Serial No. 687,249, filed December 28, 1984.

Brief Summary of the Invention

Technical Field

This invention relates to the use of acyl urea compounds for the systemic control of endoparasites and ectoparasites of warm-blooded animals.

Background of the Invention

Animal parasites, such as mites, lice, ticks, grubs, biting flies, coccidia, flukes, tapeworms, hookworms and other helminths, cause severe economic losses to livestock producers. Infestations may result in reduced feed efficiency, weight loss, decreased production of milk or eggs, and increased incidence of disease or death. The control of animal parasites is therefore one of the most important problems in the field of animal husbandry.

Certain insecticides, such as coumaphos, carlaryl and lindane, are currently available for control of some of these parasites. Insect growth regulators, such as methoprene (juvenile hormone mimic) and diflubenzuron, have been used as feed additives to control fly larvae in livestock and

Systemic control of animal parasites is accomplished by absorbing a parasiticide in the bloodstream or other tissues of the host animal. Parasites which eat or come into contact with the parasiticide-containing blood or tissue are killed, either by ingestion or contact. Only a very few insecticides have been found to be sufficiently nontoxic to be used systemically in animals.

Accordingly, the object of this invention is to provide a method for the use of certain acylurea compounds to systemically control animal parasites. This and other objects will readily become apparent to those skilled in the art in light of the teachings herein set forth.

**Disclosure of the Invention**

This invention relates to a method for controlling parasites of warm-blooded animals which comprises administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein R₁, R₂, R₃, R₄ and Y are as defined hereinafter.

Detailed Description

As indicated above, this invention relates to a method of controlling parasites of warm-blooded animals by administration of certain acyl urea compounds. More particularly, this invention involves a method for controlling endoparasites and ectoparasites of warm-blooded animals by orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
\text{O} & \quad Y \\
R₁ - C - N - C - N - R₄ \\
R₂ & \quad R₃
\end{align*}
\]

wherein:

R₁ and R₄ are independently a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system, a bicyclic aromatic or nonaromatic ring system, a polycyclic aromatic or nonaromatic ring system, and a bridged ring system which may be saturated or unsaturated in
which the permissible substituents (Z) are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio, thiocyanocarbonylalkylthio, hydroxyimino, alkoxycarbonylthio, trialkylsilyloxy, aryl dialkylsilyloxy, triarylsilyloxy, foraminidino, alkylsulfamido, dialkylsulfamido, alkoxysulfonyl, poly haloalkoxy sulfon yl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxy carbonyl and derivative salts, formamidino, alkyl, alkoxy, poly haloalkyl, poly haloalkoxy, alkoxycarbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or poly haloalkenyl; alkylthio, poly haloalkthio, alkylsulfanyl, poly haloalkylsulfanyl, alkylsulfon yl, poly haloalkylsulfon yl, alkylsulfon ylamino, alkylcarbonylamino, poly haloalkylsulfon ylamino, poly haloalkylcarbonylamino, trialkylsilyl, aryl dialkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, poly haloalkenyl, alkenyloxy, alkynyl, alknyloxy, poly haloalkenyloxy, polyfluoroalkyl, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted arylox yiminomethyl, hydrazonomethyl, unsubstituted
or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X_1 \quad \text{or} \quad \begin{array}{c}
\text{Y}_1 \\
\text{Y}_1
\end{array} \]

\[-X = R_5, \quad \text{or} \quad \begin{array}{c}
\text{Y}_1 \\
\text{Y}_1
\end{array} \]

\[-X - R_5, - P - Y_2 R_6, - Y_4 - P - Y_2 R_6 \]

\[-Y_3 R_7 \\
\text{Y}_3 R_7
\]

\[-Y_3 R_7 \\
\text{Y}_3 R_7
\]

wherein \( R_5 \) is a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system, a bicyclic aromatic or nonaromatic ring system, a polycyclic aromatic or nonaromatic ring system, and a bridged ring system which may be saturated or unsaturated in which the permissible substituents (Z) are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxycarbonylalkyl, alkoxycarbonylalkylthio, polyhaloalkenylthio, thiocyanato, propargylthio, hydrazinyl, alkoxyimino, trialkylsilyloxy, arylalkylsilyloxy, triarylsilyloxy, foraminio, alkylsulfamido, dialkylsulfamido, alkoxy sulfonyl, polyhaloalkoxy sulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts.
formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxymethyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonfyl, polyhaloalkylsulfonfyl, alkylcarbonylamino, polyhaloalkylcarbonylamino, polyhaloalkylcarbonylamino, trialkylsilyl, aryldialkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkenyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonyl hydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,
$Y_1$ and $Y_4$ are independently oxygen or sulfur,
$Y_2$ and $Y_3$ are independently oxygen, sulfur,
amino or a covalent bond, $R_6$ and $R_7$ are
independently hydrogen or substituted or
unsubstituted alkyl, polyhaloalkyl, phenyl or benzyl
in which the permissible substituents ($Z$) are the
same or different and are one or more hydrogen,
halogen, alkylcarbonyl, alkylcarbonylalkyl,
alkoxycarbonylalkyl, alkoxyalkylcarbonylalkyl,
polyhaloalkenylthio, thiocyanato, propargylthio,
hydroxyimino, alkoxyimino, trialkylsilyloxy,
aryldialkylsilyloxy, triarylsilyloxy, formamidino,
alkylsulfamido, dialkylsulfamido, alkoxyalkylsulfonyle,
polyhaloalkylsulfonyle, hydroxy, amino,
aminocarbonyl, alkylaminocarbonyl,
dialkylaminocarbonyl, aminothiocarbonyl,
alkylaminothiocarbonyl, dialkylaminothiocarbonyl,
nitro, cyano, hydroxycarbonyl and derivative salts,
formamido, alkyl, alkoxy, polyhaloalkyl,
polyhaloalkoxy, alkoxycarbonyl, substituted amino in
which the permissible substituents are the same or
different and are one or two propargyl, alkoxyalkyl,
aldehydialkyl, alkyl, alkenyl, haloalkenyl or
polyhaloalkenyl; alkylthio, polyhaloalkylthio,
aldehydialkyl, polyhaloalkylsulfanyl, alkylsulfonyle,
polyhaloalkylsulfonyle, alkylsulfonamido,
alkylcarbonylamino, polyhaloalkylsulfonamido,
polyhaloalkylcarbonylamino, trialkylsilyl,
aryldialkylsilyl, triarylsilyl, sulfonic acid and
derivative salts, phosphonic acid and derivative
salts, alkoxyalkylamino, alkylaminocarbonyloxy,
dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl,
akynyl, alkynyl, alkynloxy, polyhaloalkenyl,
polyhaloalkynyl, polyhaloalkynyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

- X, = X, - X = R₅, = X-R₅,

\[
\begin{align*}
Y₁ & \quad Y₁ \\
-X-R₅ & \quad P \quad -Y₂R₆ & \quad -Y₄ \quad P \quad -Y₂R₆ \\
\quad Y₃R₇ & \quad Y₃R₇
\end{align*}
\]

or

\[
\begin{align*}
Y₂R₆ & \quad Y₃R₇
\end{align*}
\]

and

X is a covalent single bond or double bond, a substituted or unsubstituted heteroatom or substituted carbon atom, or a substituted or unsubstituted, branched or straight chain containing two or more carbon atoms or heteroatoms in any combination in which the permissible substituents (Z) are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkythio, poly haloalkenylthio, thiocyanato, propargylthio, hydroxyiminio, alkoxyiminio, trialkylsilyloxy, aryl dialkylsilyloxy, triarylsilyloxy, foramidino, alkylsulfamido, dialkyl sulfamido, alkoxy sulfonyl, polyhaloalkoxysulfonyl, hydroxy, amino,
aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonyl, polyhaloalkylsulfonyl, alkylsulfonylamino, alkylcarbonylamino, polyhaloalkylsulfonylamino, polyhaloalkylcarbonylamino, trialkylsilyl, aryldialkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkenyloxy, polyhaloalken yloxy, polyhaloalkynyl, polyhaloalkyn yloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[ -X = X, \quad -X = R_5, = X-R_5, \]
\[ Y_1 \]
\[ -X - R_5, -P - Y_2 R_6 \]
\[ \frac{Y_1}{Y_3 R_7} \]
\[ -Y_4 + P - Y_2 R_6 \]
\[ \frac{Y_3 R_7}{Y_3 R_7} \]
or

\[ Y_2R_6 \]

\[ Y_3R_7 \]

or \( R_4 \) is hydrogen;

\( Y \) is oxygen or sulfur; and

\( R_2 \) and \( R_3 \) are independently hydrogen, alkyl, substituted alkyl in which the permissible substituents are the same or different and are one or more halogen, alkoxy, alkylthio, or cyano; cycloalkyl, cycloalkenyl, substituted benzyl in which the permissible substituents are the same or different and are one or more halogen, hydroxy, nitro, cyano, alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy; hydroxy, alkoxy, polyhaloalkoxy, alkylthio, polyhaloalkylthio, acyl, alkoxy carbonyl, alkoxy thiocarbonyl, alkylsulfanyl, substituted phenylsulfanyl in which the permissible substituents are the same or different and are one or more halogen, nitro, cyano or polyhaloalkyl; substituted phenylthio in which the permissible substituents are the same or different and are one or more halogen, alkyl, nitro, cyano, polyhaloalkyl, or substituted or unsubstituted alkoxy carbonyl, alkenoxycarbonyl, alkynoxycarbonyl, cycloalkoxycarbonyl, phenoxy carbonyl, hydroxycarbonyl or the alkali metal salt or ammonium salt thereof, alkyl, alkoxy, alkylthio, cycloalkyl, cycloalkoxy, cycloalkylthio, alkenyl, alkenylthio, alkanoyl, alkylsulfanyl, alkynyl, phenyl, phenoxy, phenylthio or amino; 

\[ N-(alkylcarbonyl)-N-alkylaminothio, alkoxy carbonylthio, trialkylsilyl, dialkylarylsilyl, thiarylsilyl, \]
or $R_2$ and $R_3$ are independently

$X, \quad -X = R_5,$

\[
\begin{array}{c}
Y_1 \\
\end{array}
\quad \begin{array}{c}
Y_1 \\
\end{array}
\]

$-X - R_5, \quad -P - Y_2 R_6, \quad -Y_4 - P - Y_2 R_6$

\[
\begin{array}{c}
Y_3 R_7 \\
\end{array}
\quad \begin{array}{c}
Y_3 R_7 \\
\end{array}
\]

or

\[
\begin{array}{c}
Y_2 R_6 \\
\end{array}
\quad \begin{array}{c}
Y_3 R_7 \\
\end{array}
\]

or $R_2$ and $R_3$ may be linked together to form a substituted or unsubstituted, heterocyclic ring system which may be saturated or unsaturated and in which the permissible substituents (Z) are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, poly haloalkenylthio, thiocyanato, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, aryl diaalkylsilyloxy, triarylsilyloxy, formamido, alkylsulfamido, dialkyl sulfamido, alkoxy sulfonyle, poly haloalkoxy sulfonyle, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxy carbonyl and derivative salts, formamido, alkyl, alkoxy, poly haloalkyl, poly haloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or
polyhaloalkeny; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonyl, polyhaloalkylsulfonyl, alkylcarboxylamino, polyhaloalkylcarboxylamino, polyhaloalkylcarboxylamino, trialkylsilyl, arylalkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkenyl, polyhaloalkenyl, alkenyl, alkynyl, alkynylthio, polyhaloalkenyloxy, polyhaloalkenyl, polyhaloalkynyl, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyaminomethyl, unsubstituted or substituted aryloximinomethyl, hydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

- \( -X, = X, -X = R_5, = X-R_5, \)

\[ \begin{align*}
&Y_1 \\
&-X - R_5, -P - Y_2R_6 \quad -Y_4 - P - Y_2R_6 \\
&Y_3R_7 \quad Y_3R_7
\end{align*} \]

or

\[ \begin{align*}
&Y_2R_6 \\
&Y_3R_7
\end{align*} \]

Certain \( Z \) groups which may be substituted or unsubstituted include aryloximinomethyl and arylhydrazonomethyl. Other groups within the definition of \( R_2 \) and \( R_3 \) which may be substituted or unsubstituted include alkoxy carbonyl, alkenoxycarbonyl, alkynoxycarbonyl, cycloalkoxy carbonyl, phenoxy carbonyl, hydroxycarbonyl or the alkali metal salt or ammonium salt thereof, alkyl, alkoxy, alkylthio, cycloalkyl, cycloalkyloxy, cycloalkylthio, alkenyl, alkenylthio, alkanoyl.
alkylsulfonyl, alkynyl, phenyl, phenoxy, phenylthio and amino. Permissible substituents for these groups include independently one or more halogen, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, cyano, nitro, alkylthio, alkylsulfinyl, alkylsulfonyl, polyhaloalkylthio, polyhaloalkylsulfonyl, trialkylsilyl and arylidialkylsilyl in any combination.

The alkyl-containing moieties above may contain from about 1 to about 100 carbon atoms or greater, preferably from about 1 to about 30 carbon atoms, and more preferably from about 1 to about 20 carbon atoms. The polysaccharide moiety may contain up to about 50 carbon atoms.

Monocyclic ring systems encompassed by R₁, R₄ and R₅ in formula 1 may be represented by generalized formula 2 as follows:

![Diagram](image)

wherein B₁ represents a saturated or unsaturated carbon atom and A₁ represents a ring-forming chain of atoms which together with B₁ forms a cyclic system containing from 0 to 3 double bonds or from 0 to 2 triple bonds. A₁ may contain entirely from 2 to 12 carbon atoms, may contain a combination of from 1 to 11 carbon atoms and from 1 to 4 heteroatoms which may be selected independently from N, O, S, P or other heteroatoms, or may contain ring-forming heteroatoms alone.

Ring-forming heteroatoms may in some cases bear oxygen atoms as in aromatic N-oxides and ring systems containing the sulfinyl, sulfonyl, selenoxide and phosphono moieties.
Selected carbon atoms contained in cycles formed by \( B_1 \) and \( A_1 \) containing more than 3 ring-forming atoms may bear carbonyl, thiocarbonyl, substituted or unsubstituted imino groups or substituted or unsubstituted methylidene groups.

The group designated as \( Z \) represents one or more substituents selected independently from among the group of substituents defined for \( Z \) herein. When the cycle formed by \( B_1 \) and \( A_1 \) contains fewer than 4 ring forming members, it should be a saturated carbocycle, i.e. cyclopropyl. When the cycle formed by \( B_1 \) and \( A_1 \) contains fewer than 5 ring-forming members, it should contain no more than 1 heteroatom.

Representative illustrations of \( A_1 \) include the following:

1. A chain of from 2-12 carbon atoms which together with \( B_1 \) contains 0 double bonds;
2. A chain of from 4-12 carbon atoms which together with \( B_1 \) contains 1 double bond;
3. A chain of from 4-12 carbon atoms which together with \( B_1 \) contains 2 double bonds;
4. A chain of from 5-12 carbon atoms which together with \( B_1 \) contains 3 double bonds;
5. A chain containing 1 heteroatom and from 2 to 11 carbon atoms which together with \( B_1 \) contains 0 double bonds;
6. A chain containing 1 heteroatom and from 3-11 carbon atoms which together with \( B_1 \) contains 1 double bond;
7. A chain containing 1 heteroatom and from 3-11 carbon atoms which together with \( B_1 \) contains 2 double bonds;
8. A chain containing 1 heteroatom and from 4-11 carbon atoms which together with \( B_1 \) contains 3 double bonds;
(9) A chain containing 2 heteroatoms and from 2-10 carbon atoms which together with $B_1$ contains 0 double bonds;

(10) A chain containing 2 heteroatoms and from 2-10 carbon atoms which together with $B_1$ contains 1 double bond;

(11) A chain containing 2 heteroatoms and from 2-10 carbon atoms which together with $B_1$ contains 2 double bonds;

(12) A chain containing 2 heteroatoms and from 3-10 carbon atoms which together with $B_1$ contains 3 double bonds;

(13) A chain containing 3 heteroatoms and from 1-9 carbon atoms which together with $B_1$ contains 0 double bonds;

(14) A chain containing 3 heteroatoms and from 1-9 carbon atoms which together with $B_1$ contains 1 double bond;

(15) A chain containing 3 heteroatoms and from 1-9 carbon atoms which together with $B_1$ contains 2 double bonds;

(16) A chain containing 3 heteroatoms and from 2-9 carbon atoms which together with $B_1$ contains 3 double bonds;

(17) A chain containing 4 heteroatoms and from 0-8 carbon atoms which together with $B_1$ contains 0 double bonds;

(18) A chain containing 4 heteroatoms and from 0-8 carbon atoms which together with $B_1$ contains 1 double bond;

(19) A chain containing 4 heteroatoms and from 0-8 carbon atoms which together with $B_1$ contains 2 double bonds; and
(20) A chain containing 4 heteroatoms and from 1-8 carbon atoms which together with $B_1$ contains 3 double bonds.

Illustrative monocyclic ring structures which are encompassed by $R_1$, $R_4$ and $R_5$ in formula 1 include the following:
wherein \( Z \) is as defined herein.

Bicyclic ring systems encompassed by \( R_1 \), \( R_4 \) and \( R_5 \) in formula 1 may be represented by generalized formulae 3 and 4 as follows:

wherein \( B_2 \) and \( B_3 \) may be independently a saturated or unsaturated carbon atom or a saturated nitrogen atom, \( A_2 \) and \( A_3 \) independently represent
the ring-forming chains of atoms described below and
Z represents one or more substituents selected
independently from among the group of substituents
defined for Z herein. Combinations of A_2 and A_3
may be selected from among the arrays of atoms
listed below in items (21) through (98) and may
contain in combination with either B_1 or B_2, but
not both, from 0 to 2 double bonds. A_2 and A_3,
independent of B_2 and B_3, may contain entirely
from 1 to 11 carbon atoms, may contain a combination
of 1 to 3 heteroatoms which may be selected
independently from among N, O, S, P or other
heteroatoms together with from 1 to 10 carbon atoms
or may contain from 1-3 ring-forming heteroatoms
alone.

Ring-forming heteroatoms may in some cases
bear oxygen atoms, as in aromatic N-oxides and ring
systems containing the sulfinyl, sulfonyl,
selenoxide and phosphono groups. Selected carbon
atoms contained in A_2 and A_3 may bear carbonyl,
thiocarbonyl, substituted or unsubstituted imino
groups or substituted or unsubstituted methylidene
groups.

In regard to structures encompassed within
formulae 3 and 4, it is noted as follows:
(a) When B_2 and B_3 are both nitrogen,
the groups A_2 and A_3 should each contain no
fewer than three ring atoms;
(b) When B_2 but not B_3 is nitrogen,
either of A_2 or A_3 should contain at least three
ring atoms and the other at least two ring atoms;
(c) When either of groups A₂ or A₃ contains fewer than three ring atoms, the other should contain at least three ring atoms and the bridgehead atoms should be saturated;

(d) When the group A₂ or A₃ contains a carbon atom bearing a carbonyl, thiocarbonyl, imino or methylidene group, it should together with B₂ and B₃ form a cycle having at least four members;

(e) When an annular double bond is exocyclic to either of the two rings represented in structures 3 and 4, it should be contained in a ring containing at least five members and be exocyclic to a ring containing at least five members; and

(f) When a group A₂ or A₃ is joined to the bridgehead atoms B₁ and B₂ by 2 double bonds, the group A₂ or A₃ is understood to contain one double bond and the bridgehead atoms are considered to be unsaturated.

It is recognized that bycyclic ring systems defined for R₁, R₄ and R₅ may be spirocyclic ring systems and are not limited to the fused bicyclic structures of formulae 3 and 4. Spirocyclic ring systems may be saturated or unsaturated carbocyclic or heterocyclic and may be independently substituted by one or more substituents Z as defined herein.

Illustrative combinations of A₂ and A₃ which are encompassed within generalized formulae 3 and 4 are listed below as follows:

(21) A₂ is a chain containing from 1-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds and A₃ is a chain

SUBSTITUTE CHECK
containing 3-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds;

    (22) A₂ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing 3-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds;

    (23) A₂ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond;

    (24) A₂ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 2 double bonds and A₃ is a chain containing 3-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds;

    (25) A₂ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 2 double bonds and A₃ is a chain containing 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond;

    (26) A₂ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 2 double bonds and A₃ is a chain containing 4-11 carbon atoms which together with B₂ or B₃ contains 2 double bonds;

    (27) A₂ is a chain containing 1 heteroatom and from 0-10 carbon atoms which together with B₂ or B₃ contains 0 double bonds and A₃ is a chain containing 1-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds;
(28) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing 1-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds;

(29) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds;

(30) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(31) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 4-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;

(32) \(A_2\) is a chain containing 1 heteroatom and from 0-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(33) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;
(34) A₂ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing from 4-11 carbon atoms which together with B₂ or B₃ contains 2 double bonds;

(35) A₂ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond;

(36) A₂ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B₂ or B₃ contains 0 double bonds and A₃ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond;

(37) A₂ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds;

(38) A₂ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B₂ or B₃ contains 2 double bonds and A₃ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 0 double bonds;

(39) A₂ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing from 3-11 carbon atoms which together with B₂ or B₃ contains 1 double bond;
(40) \(A_2\) is a chain containing 2 heteroatoms and from 2-9 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(41) \(A_2\) is a chain containing 2 heteroatoms and from 2-9 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;

(42) \(A_2\) is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;

(43) \(A_2\) is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(44) \(A_2\) is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;

(45) \(A_2\) is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 2-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds;
(46) \(A_2\) is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds;

(47) \(A_2\) is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds;

(48) \(A_2\) is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(49) \(A_2\) is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(50) \(A_2\) is a chain containing 3 heteroatoms and from 2-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond;

(51) \(A_2\) is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;
(52) \(A_2\) is a chain containing 3 heteroatoms and from 2-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;

(53) \(A_2\) is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing from 3-11 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds;

(54) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 0 double bonds and \(A_3\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) and \(B_3\) contains 0 double bonds;

(55) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) and \(B_3\) contains 0 double bonds;

(56) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 2 double bonds and \(A_3\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) and \(B_3\) contains 0 double bonds;

(57) \(A_2\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) or \(B_3\) contains 1 double bond and \(A_3\) is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with \(B_2\) and \(B_3\) contains 1 double bond;

D-14312-2
(58) $A_2$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ and $B_3$ contains 1 double bond;

(59) $A_2$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ and $B_3$ contains 2 double bonds;

(60) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ and $B_3$ contains 0 double bonds;

(61) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ and $B_3$ contains 0 double bonds;

(62) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ and $B_3$ contains 0 double bonds;
(63) A_2 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 0 double bonds and A_3 is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B_2 and B_3 contains 1 double bond;

(64) A_2 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 1 double bond and A_3 is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B_2 and B_3 contains 1 double bond;

(65) A_2 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 2 double bonds and A_3 is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B_2 and B_3 contains 1 double bond;

(66) A_2 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 0 double bonds and A_3 is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B_2 and B_3 contains 2 double bonds;

(67) A_2 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 1 double bond and A_3 is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B_2 and B_3 contains 2 double bonds;
(68) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds;

(69) $A_2$ is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(70) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(71) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(72) $A_2$ is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond;
(73) A₂ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 1 double bond;

(74) A₂ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with B₂ or B₃ contains 2 double bonds and A₃ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 1 double bond;

(75) A₂ is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with B₂ or B₃ contains 0 double bonds and A₃ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 2 double bonds;

(76) A₂ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with B₂ or B₃ contains 1 double bond and A₃ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 2 double bonds;

(77) A₂ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with B₂ or B₃ contains 2 double bonds and A₃ is a chain containing 1 heteroatom and from 2-10 carbon atoms which together with B₂ or B₃ contains 2 double bonds;
(78) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(79) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 2 heteroatoms and from 2-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(80) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(81) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond;

(82) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond;
(83) $A_2$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 2-9 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds;

(84) $A_2$ is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(85) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(86) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(87) $A_2$ is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond;
(88) A_2 is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with B_2 or B_3 contains 1 double bond and A_3 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 1 double bond;

(89) A_2 is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with B_2 or B_3 contains 2 double bonds and A_3 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 1 double bond;

(90) A_2 is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with B_2 or B_3 contains 0 double bonds and A_3 is a chain containing 2 heteroatoms and from 1-9 carbon atoms which together with B_2 or B_3 contains 2 double bonds;

(91) A_2 is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with B_2 or B_3 contains 1 double bond and A_3 is a chain containing 2 heteroatoms and from 2-9 carbon atoms which together with B_2 or B_3 contains 2 double bonds;

(92) A_2 is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with B_2 or B_3 contains 2 double bonds and A_3 is a chain containing 2 heteroatoms and from 2-9 carbon atoms which together with B_2 or B_3 contains 2 double bonds;
(93) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds and $A_3$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(94) $A_2$ is a chain containing 3 heteroatoms and from 2-8 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 3 heteroatoms and from 2-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(95) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 3 heteroatoms and from 0-8 carbon atoms which together with $B_2$ or $B_3$ contains 0 double bonds;

(96) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond and $A_3$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond;

(97) $A_2$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 2 double bonds and $A_3$ is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with $B_2$ or $B_3$ contains 1 double bond; and
(98) \( A_2 \) is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with \( B_2 \) or \( B_3 \) contains 2 double bonds and \( A_3 \) is a chain containing 3 heteroatoms and from 1-8 carbon atoms which together with \( B_2 \) or \( B_3 \) contains 2 double bonds.

Illustrative bicyclic ring structures which are encompassed by \( R_1, R_4 \) and \( R_5 \) in formula 1 included the following:
Polycyclic ring systems, i.e., greater than 2 rings, encompassed by $R_1$, $R_4$ and $R_5$ in formula 1 may be represented by generalized formulae 5, 6, 7 and 8 as follows:
wherein B₄, B₅, B₆ and B₇ may be independently a saturated or unsaturated carbon atom or a saturated nitrogen atom, and A₄, A₅, A₆ and A₇ independently represent ring forming chains of atoms which may contain together with one or the other (but not both) of their two associated bridgehead atoms, from 0-2 double bonds. The groups Z represent one or more substituents selected independently from among the group of substituents defined for Z herein.

The ring-forming elements of A₄, A₅, A₆ and A₇ independent of B₄, B₅, B₆, and B₇ may contain entirely from 1-11 carbon atoms, may contain a combination of from 1-10 carbon atoms and from 1-3 heteroatoms which may be selected independently from among N, O, S, P or other heteroatoms, or may contain from 1-3 heteroatoms alone. Ring-forming heteroatoms may in some cases bear oxygen atoms as in aromatic N-oxides and ring systems containing the sulfinyl, sulfonyl, selenoxide and phosphono groups. The group A₆ may at times be defined as a bond. Selected carbon atoms contained in A₄, A₅, A₆ and A₇ may bear one or more carbonyl, thiocarbonyl or substituted or unsubstituted imino groups.
On structure 8 the groups $B_8$, $B_9$ and $B_{10}$ represent independently a saturated or unsaturated carbon atom or a saturated nitrogen atom. The group $B_{11}$ may represent a saturated or unsaturated carbon atom or a nitrogen or phosphorous atom. The groups $A_8$, $A_9$ and $A_{10}$ represent ring-forming chains of atoms which may contain together with 1 of the groups $B_8$, $B_9$, $B_{10}$ and $B_{11}$ from 0-2 double bonds.

The ring-forming elements of groups $A_8$, $A_9$ and $A_{10}$ independent of groups $B_8$, $B_9$, $B_{10}$ and $B_{11}$ may contain entirely from 2-10 carbon atoms, may contain from 1-10 carbon atoms in combination with 1-3 heteroatoms which may be selected independently from among N, O, S, P or other heteroatoms, or may contain from 2-3 heteroatoms alone. Ring-forming heteroatoms may in some cases bear oxygen atoms as in aromatic N-oxides and in ring systems containing the sulfinyl, sulfonyl, selenoxide and phosphono groups. Selected carbon atoms contained in groups $A_8$, $A_9$ and $A_{10}$ may bear one or more carbonyl, thiocarbonyl or substituted or unsubstituted imino groups.

It is recognized that polycyclic ring systems defined for $R_1$, $R_4$ and $R_5$ may be spirocyclic ring systems and are not limited to the fused polycyclic structures of formulae 5, 6, 7 and 8. Spirocyclic ring systems may be saturated or unsaturated, carbocyclic or heterocyclic and may be independently substituted by one or more substituents 2 as defined herein.

Illustrative polycyclic ring structures which are encompassed by $R_1$, $R_4$ and $R_5$ in formula 1 include the following:
Bridged bycyclic structures encompassed by \( R_1, R_4 \) and \( R_5 \) in formula 1 may be represented by generalized formulae 9, 10, and 11 as follows:

\[
\begin{align*}
9 & \quad \begin{array}{c}
\text{A}_{11} \quad \text{A}_{12} \quad \text{A}_{13} \\
\text{B}_{12} \quad \text{Z} \\
\text{Z} \quad \text{B}_{13} \quad \text{Z}
\end{array} \\
10 & \quad \begin{array}{c}
\text{A}_{11} \quad \text{A}_{13} \quad \text{A}_{12} \\
\text{B}_{12} \quad \text{Z} \\
\text{Z} \quad \text{B}_{13} \quad \text{Z}
\end{array} \\
11 & \quad \begin{array}{c}
\text{A}_{12} \quad \text{A}_{11} \quad \text{A}_{13} \\
\text{B}_{12} \quad \text{Z} \\
\text{Z} \quad \text{B}_{13} \quad \text{Z}
\end{array}
\end{align*}
\]

wherein \( B_{12} \) and \( B_{13} \) may be independently a saturated carbon atom optionally substituted by \( Z \) or a nitrogen atom, and the groups \( A_{11}, A_{12} \) and \( A_{13} \) independently represent ring-forming chains of atoms which may contain, independently of \( B_{12} \) and \( B_{13} \), from 0-2 double bonds. The groups \( Z \) represent one or more substituents selected independently from among the groups of substituents defined for \( Z \) herein.

The ring-forming elements of \( A_{11}, A_{12} \) and \( A_{13} \), independent of \( B_{12} \) and \( B_{13} \), may contain entirely from 1-11 carbon atoms, may contain a combination of from 1-10 carbon atoms and from 1-3 heteroatoms which may be selected independently from among \( N, O, S, P \) or other heteroatoms, or may contain from 1-3 heteroatoms alone with the proviso
that when one of the groups $A_{11}$, $A_{12}$ and $A_{13}$ is a single heteroatom, the other two groups should contain two or more ring-forming atoms. A second proviso is that when one or both of the groups $B_{12}$ and $B_{13}$ is nitrogen, the groups $A_{11}$, $A_{12}$ and $A_{13}$ should contain at least two saturated ring-forming atoms.

Ring-forming heteroatoms may in some cases bear oxygen atoms as in the sulfinyl, sulfonyl, selenoxide and phosphono moieties. Selected carbon atoms contained in $A_{11}$, $A_{12}$ and $A_{13}$ may bear one or more carbonyl, thiocarbonyl or substituted or unsubstituted imino groups.

Illustrative bridged bicyclic structures which are encompassed by $R_1$, $R_4$ and $R_5$ in formula 1 include the following:
The substituent X may be an unsubstituted heteroatom such as an oxygen or sulfur, as in carbonyl and thiocarbonyl systems, or may be a substituted heteroatom or carbon atom. X may also be a covalent single or double bond, or an array of atoms serving to join the ring system $R_5$ to the remainder of the acyl urea molecule. The group X may be joined to the ring system $R_5$ by means of a single or double bond. Likewise, X may be attached to the remainder of the acyl urea molecule by means of a double or single bond. X may be a saturated or unsaturated, branched or straight chain of carbon atoms; a branched or straight, saturated or unsaturated chain of atoms consisting of both carbon atoms and heteroatoms; or may be a branched or straight, saturated or unsaturated chain consisting entirely of heteroatoms. Selected heteroatomic components of X may bear oxygen atoms as in the case of groups containing the sulfonyl, sulfinyl, N-oxide and phosphono moieties. Selected heteroatomic components of X may bear one or more substituents Z as defined herein. Selected carbon atoms
participating in $X$ may bear carbonyl, thiocarbonyl, substituted or unsubstituted imino, substituted or unsubstituted alkylidene or one or more substituents $Z$ as defined herein.

Illustrative structures which are encompassed by substituent $X$ include the following:
wherein \( m \) is a value of from 0 to 8, \( n \) is a value of from 0 to 2, and \( R_8, R_9, R_{10} \) and \( R_{11} \) are independently hydrogen or substituted or unsubstituted alkyl, polyhaloalkyl, phenyl or benzyl in which the permissible substituents are as defined for \( Z \) herein.

\( R_2 \) and \( R_3 \) may be linked together to form a heterocyclic saturated or unsaturated ring.
This ring system may be part of a saturated or unsaturated, aromatic or nonaromatic, carbo- or hetero-, bicyclic or polycyclic ring system. Illustrative of suitable heteroatoms include oxygen, sulfur, nitrogen, silicon and phosphorous. The ring system can be optionally substituted with one or more of the substituents Z as defined herein.

Illustrative linking structures which are encompassed by R₂ and R₃ of formula 1 include the following:

\[ \begin{array}{c}
\text{N} & \text{O} \\
\text{N} & \text{O} \\
\text{N} & \text{O} \\
\text{N} & \text{O} \\
\text{N} & \text{O} \\
\end{array} \]

\[ \begin{array}{c}
\text{R}_{12} & \text{R}_{13} \\
\text{R}_{12} & \text{R}_{13} \\
\text{R}_{12} & \text{R}_{13} \\
\text{R}_{12} & \text{R}_{13} \\
\text{R}_{12} & \text{R}_{13} \\
\end{array} \]

wherein \( R_{12}, R_{13}, R_{14} \) and \( R_{15} \) are independently hydrogen, halogen, or substituted or...
unsubstituted alkyl polyhaloalkyl, phenyl or benzyl in which the permissible substituents are as defined for Z herein.

It is recognized that the acyl urea linkage in formula 1 may be converted to different forms including the following:
wherein \( R_{73}, R_{74} \) and \( R_{75} \) are independently hydrogen, alkyl, substituted alkyl in which the permissible substituents are the same or different and are one or more halogen, alkoxy, alkylthio or cyano; cycloalkyl, cycloalkenyl, substituted benzyl in which the permissible substituents are the same or different and are one or more halogen, hydroxy, nitro, cyano, alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy; hydroxy, alkoxy, polyhaloalkoxy, alkylthio, polyhaloalkylthio; acyl, alkoxy carbonyl,
alkoxythiocarbonyl, alkylsulfonyl, substituted phenylsulfonyl, in which the permissible substituents are the same or different and are one or more halogen, nitro, cyano or alkyl; substituted phenylthio in which the permissible substituents are the same or different and are one or more halogen, alkyl, nitro, cyano, polyhaloalkyl, or substituted or unsubstituted alkoxy carbonyl, alkenoxycarbonyl, alkynoxycarbonyl, cycloalkoxycarbonyl, phenoxycarbonyl, hydroxycarbonyl or the alkali metal salt or ammonium salt thereof, alkyl, alkoxy, alkylthio, cycloalkyl, cycloalkyloxy, cycloalkylthio, alkenyl, alkenylthio, alkanoyl, alkylsulfonyl, alkynyl, phenyl, phenoxy, phenylthio or amino; N-(alkylcarbonyl)-N-alkylaminothio, alkoxy carbonylthio, trialkylsilyl, dialkylarylsilyl, triarylsilyl and

\[ Y_1 \]
\[ \text{P} \]
\[ Y_2 \]
\[ R_6 \]
\[ Y_3 \]
\[ R_7 \]

wherein \( Y_1, Y_2, Y_3, R_6 \) and \( R_7 \) are as defined herein; and \( A \) can be \(-\text{SSR}_7\), 1-benzimidazolyl, 1-benztriazolyl and 1-pyrazolyl which can be substituted with one or more substituents \( Z \) defined herein. \( A \) can also be 1-diazolyl or 1-imidazolyl optionally substituted with alkyl. Such modifications and embodiments can be made without departing from the spirit and scope of this invention.
It is readily apparent that formula 1 encompasses a wide variety of urea compounds as more particularly described hereinbelow. For example, U.S. Patent 4,426,385, incorporated herein by reference, discloses and claims bicyclooxyphenyl urea compounds which are encompassed by the above formula 1 and which may be used in the method of this invention. Such bicyclooxyphenyl urea compounds can be prepared by the methods described in U.S. Patent 4,426,385, and are available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the bicyclooxyphenyl urea compounds of U.S. Patent 4,426,385 which may be useful in the method of this invention include the following:

1-[4-(4-chloro-1-naphthoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-3,5-dichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-3-methylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-3-methylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-3-methylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2-methyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2-methyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2-methyl-5-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(4-chloro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[2,3,5-trichloro-4-(4-chloro-1-naphthoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[2,3,5-trichloro-4-(4-chloro-1-naphthoxy)phenyl]-3-(2-fluorobenzoyl)urea;
1-[3,5-dimethyl-4-(4-chloro-1-naphthoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dimethyl-4-(4-chloro-1-naphthoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dimethyl-4-(4-chloro-1-naphthoxy)phenyl]-3-(2-fluorobenzoyl)urea;
1-(3,5-dimethyl-4-(4-chloro-1-naphthoxy)phenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-(2,5-dimethyl-3-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-3-(2,6-difluorobenzoyl)urea;
1-(2,5-dimethyl-3-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-3-(2-fluorobenzoyl)urea;
1-(2,5-dimethyl-3-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-3-(2-chlorobenzoyl)urea;
1-[3-methyl-4-(4-chloro-1-naphthoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(3-methyl-4-(4-chloro-1-naphthoxy)phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-methyl-4-(4-chloro-1-naphthoxy)phenyl)-3-(2-chlorobenzoyl)urea;
1-(3,5-dichloro-4-[4-nitro-1-naphthoxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-cyano-1-naphthoxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-trifluoromethyl-1-naphthoxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3,5-dichloro-4-[2,2-dimethyl-2,3-dihydro-4-trifluoromethyl-7-benzofuranyloxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-nitro-5,6,7,8-tetrahydro-1-naphthoxy]-phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-benzothiényloxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-5-n-pentyl-4-[2-trifluoromethyl-7-trifluoromethylsulfinyl-4-benzodioxalanyloxy]phenyl)-3-(2,6-dichlorobenzoyl)urea;
1-(4-[2-chloro-4-nitro-5-oxo-5,6,7,8-tetrahydro-1-naphthoxy]-3,5-dibromophenyl)-3-(2,6-dichlorobenzoyl)urea;
1-(3,5-difluoro-4-[4-phenylsulfinyl-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dichloro-4-[2,3-dihydro-2,2-dimethyl-7-benzofuranyloxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-chloro-2,3-dihydro-2,2-dimethyl-7-benzofuranyloxy]phenyl)-3-(2-chlorobenzoyl)urea;
1-(4-[4-chloro-1-naphthoxy]-3,5-dichlorophenyl)-3-[[2,4-dichloro-3-pyridiny]l carbonyl]urea;
1-(4-[2,4-dichloro-1-naphthoxy]-3,5-dichlorophenyl)-3-(3-trifluoromethyl-2-pyrazinyl)carbonyl)urea;
1-(4-[1,6-dichloro-2-naphthoxy]-3-chlorophenyl)-3-(2-pyrimidinyl)carbonyl)urea;
1-(4-[4-chloro-1-naphthoxy]-3,5-dichlorophenyl)-3-(5-chloro-4-pyrimidinyl)carbonyl)urea;
1-(4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-[4-chloro-1-naphthoxy]-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-[1-naphthoxy]-3-chlorophenyl)-3-(2,6-dichlorobenzoyl)urea;
1-(4-[2-naphthoxy]-phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-methoxy-1-naphthoxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-4-[4-chloro-1-naphthoxy]-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-[4-chloro-1-naphthoxy]-2,5-dimethylphenyl)-3-(2-chlorobenzoyl)urea;
1-(3-chloro-4-[4-chloro-1-naphthoxy]-2,5-dimethylphenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(4-[4-chloro-5,6,7,8-tetrahydro-1-naphthoxy]-3,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-[4-chloro-5,6,7,8-tetrahydro-1-naphthoxy]-5-methylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-[4-chloro-5,6,7,8-tetrahydro-1-naphthoxy]-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-(3,5-dichloro-4-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-4-[4-chloro-5,6,7,8-tetrahydro-1-naphthoxy]-5-methylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-5-methylphenyl)-3-(2-chloro-6-fluoro-
benzoyl)urea;
1-(3,5-dimethyl-4-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]phenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-5-methyl-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-methyl-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dichloro-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(2-methyl-4-[4-chloro-1-naphthoxy]-5-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-trifluoromethyl-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dimethyl-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,5-difluorobenzoyl)urea;
1-(2,3,5-trichloro-4-[4-chloro-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(2-methyl-3-chloro-4-[4-chloro-1-naphthoxy]-5-difluoromethyl-phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(2-methyl-3-chloro-4-[4-chloro-1-naphthoxy]-5-carboxyphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-[4N-ethyl-N-phenylsulfamido-5,6,7,8-tetrahydro-1-naphthoxy]phenyl)-1-(2-methoxy-6-
methylbenzoyl)urea;
1-(3,5-difluoro-4-[4-phenylsulfanyl-1-naphthoxy]phenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-dichloromethyl-4-[4-phenylsulfonyl-1-naphthoxy]phenyl)-3-(2-trifluoromethylbenzoyl)urea;
1-(3,5-dichloro-4-[4-chloro-1-naphthoxy]phenyl)-N-methyl-N'-methyl-3-(2,6-dichlorobenzoyl)urea;
1-(4-[4-N-(4-chlorophenyl)sulfamido-1-naphthoxy]-3-iodo-5-methylphenyl)-3-(2-trichloromethoxybenzoyl)urea;
1-(3,5-dichloro-4-[4-chloro-1-naphthoxy]phenyl-3-(2-chloro-6-nitrobenzoyl)urea;
1-(4-[7-2,4-dichlorophenylsulfonyl-3-isopropoxy-5-benzofuranyloxy]phenyl)-3-(2-ethoxybenzoyl)urea;
1-(3-methoxy-4-[4-chloro-1-naphthoxy]phenyl-3-(2,6-difluorobenzoyl)urea;
1-(3-perfluoroisopropoxy-4-[2,2,6-trimethyl-4-benzodioxalanyl oxy]phenyl)-3-(2-iodobenzoyl)urea;
1-(3-chloro-4-[2-chloro-4-ethylsulfonyl-1-naphthoxy]phenyl)-3-(2-chlorobenzoyl)urea;
1-(3-chlorodifluoromethyl-4-[5,8-dichloro-4-trichloromethylsulfonyl-1-naphthoxy]phenyl)-3-(2-chlorobenzoyl)urea;
1-(3,5-dichloro-4-[4-methylsulfonyl-5,6,7,8-tetrahydro-1-naphthoxy]phenyl)-3-(2,6-dichlorobenzoyl)urea; and
1-(4-[4-N-(4-nitrophenyl)-N-isopropyl-sulfamido-5,6,7,8-tetrahydro-1-naphthoxy]phenyl)-3-(2-chloro-6-methylbenzoyl)urea.

U.S. Patent Application Serial No. 495,331, filed May 20, 1983, incorporated herein by reference, discloses and claims 1-(alkylphenoxyaryl)-3-benzoyl urea compounds which may be used in the method of this invention. Such 1-(alkylphenoxyaryl)-3-benzoyl urea compounds can be prepared by the methods described in the application, and are available from Union Carbide.
Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the 1-(alkylphenoxyaryl)-3-benzoyl urea compounds of U.S. Patent Application Serial No. 495,331 which may be useful in the method of this invention include the following:

1-[2,5-dimethyl-3-chloro-4-(2,5-dimethyl-4-chlorophenoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[2,5-dimethyl-3-chloro-4-(2,5-dimethyl-4-chlorophenoxy)phenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,3-dimethyl-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,3-dimethyl-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(2-methyl-3,4-dichlorophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-3,4-dichlorophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-methylphenox)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methylphenox)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-methylphenox)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-methylphenox)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2-methylphenox)-3,6-dimethyl-5-chlorophenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(2-methylphenox)-2,3,5,6-tetramethylphenyl]-3-(2,6-dimethylbenzoyl)urea;
1-[4-(2-methylphenox)-2,3,5,6-tetramethylphenyl]-3-(2,6-methylbenzoyl)thiourea;
1-[4-(2-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-methylphenoxy)-2,3,5-trimethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-methylphenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
4-(2-methyl-4-bromophenoxyc)-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-bromophenoxy)-3,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2-bromo-4-methylphenoxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-methylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2-bromo-4-methylphenoxy)-3,5-dimethylphenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(2-methyl-4-t-buty1phenoxy)3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-t-buty1phenoxy)-2,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-methyl-4-t-buty1phenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-t-buty1phenoxy)-3,5-dichlorophenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(2-methyl-4-t-buty1phenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-t-buty1phenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2-methyl-4-t-buty1phenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-methyl-4-t-buty1phenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-methyl-4-chlorophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-chlorophenoxy)-3,5-dichlorophenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(2-methyl-4-chlorophenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-chlorophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-dimethoxybenzoyl)urea;
1-[4-(4-nonylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(4-nonylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(4-nonylphenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(4-nonylphenoxy)-3,5-dimethylphenyl]-3-(2-methylbenzoyl)urea;
1-[4-(2-chloro-4-methylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-chloro-4-methylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2-chloro-4-methylphenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-chloro-4-methylphenoxy)2,3,5,6-tetramethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3,4,5-trimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3,5,6-trimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(3,4,5-trimethylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(3,4,5-trimethylphenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,3,5-trimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-dimethoxybenzoyl)urea;
1-[4-(2,3,5-trimethylphenoxy)-2,3,5,6-tetramethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,3,5-trimethylphenoxy)-3,3-propylphenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(2,3,5-trimethylphenoxy)-3-trifluoromethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-trifluoromethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-trifluoromethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2-methyl-4-trifluoromethylphenoxy)-3,5-dichlorophenyl]-3-(2,6-dimethoxybenzoyl)urea;
1-[4-(2-methyl-4-trifluoromethylphenoxy)-2,3,5-trimethylphenyl]-3-(2-methylbenzoyl)thiourea;
1-[4-(2,4-dimethylphenoxy)2,3,6-trimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3,5-dimethylphenoxy)-2,6-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-methoxyphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-dimethylaminophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-methylthiophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-methylsulfonylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-methoxyphenoxy)-3-chloro-6-methylphenyl]-3-(2-methylbenzoyl)urea;
1-[4-(2-methyl-4-methoxyphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(3,4-dichlorobenzoyl)urea;
1-[4-(2,3-dimethyl-4-bromophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-chlorophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-ethoxycarbonylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-trifluoromethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-trifluoromethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-trifluoromethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-trifluoromethoxyphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-trifluoromethoxyphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3,5-dimethyl-4-chlorophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4,5-trimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4,5-trimethylphenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-bromophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-cyanophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dimethyl-4-cyanophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-methyl-4-cyanophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-methyl-4-cyanophenoxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dimethylphenoxy)-2,5-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dimethylphenoxy)-2,5-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-{4-(2,4-dimethylphenoxy)-2,6-dimethyl-5-chlorophenyl}-3-(2-chlorobenzoyl)urea;
1-{4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chlorophenyl}-3-(2,6-difluorobenzoyl)urea;
1-{4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chlorophenyl}-3-(2-fluorobenzoyl)urea;
1-{4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chlorophenyl}-3-(2-chlorobenzoyl)urea;
1-{4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chlorophenyl}-3-(2,6-dichlorobenzoyl)urea;
1-{4-(2,5-dichloro-4-methylphenoxy)-3,6-dimethyl-5-chlorophenyl}-3-(2,6-difluorobenzoyl)urea;
1-{4-(2,5-dichloro-4-methylphenoxy)-3,6-dimethyl-5-chlorophenyl}-3-(2-chlorobenzoyl)urea;
1-{4-(2-methyl-4,5-dichlorophenoxy)-3,6-dimethyl-5-chlorophenyl}-3-(2,6-difluorobenzoyl)urea;
1-{4-(2-methyl-4,5-dichlorophenoxy)-3,6-dimethyl-5-chlorophenyl}-3-(2-chlorobenzoyl)urea; and
1-{4-(2,5-dimethyl-4-chlorophenoxy)-3-methylphenyl}-3-(2,6-difluorobenzoyl)urea.

U.S. Patent Application Serial No. (D-13541-2), filed March 29, 1985, incorporated herein by reference, discloses and claims 1-(phenoxyphenyl)-3-benzoyl urea compounds which may be used in the method of this invention. Such 1-(phenoxyphenyl)-3-benzoyl urea compounds can be prepared by the methods described in the application, and are available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the 1-(phenoxyphenyl)-3-benzoyl urea compounds of U.S. Patent Application Serial No. (D-13541-2) which may
be useful in the method of this invention include
the following:
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-3-chloro-
phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-3-chloro-
phenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-3-chloro-
phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-3-chloro-
phenyl]-3-(2-bromo-6-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-3-chloro-
phenyl]-3-(2-fluoro-6-bromobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-
phenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,4-dibromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)thiourea;  
1-[4-(2,4-dibromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-dichlorobenzoyl)thiourea;  
1-[4-(2,4-dibromophenoxy)2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;  
1-[4-(2,4-dibromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea,  
1-[4-(2,4-dibromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;  
1-[4-(2,4-dibromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chloro-6-bromobenzoyl)urea;  
1-[4-(2-fluoro-4-chlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;  
1-[4-(2,4-difluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;  
1-[4-(2,4-difluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;  
1-[4-(2,4-difluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;  
1-[4-(2-chloro-4-fluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;  
1-[4-(2-chloro-4-fluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;  
1-[4-(2-chloro-4-fluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;  
1-[4-(2-fluoro-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;  
1-[4-(2-fluoro-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;  
1-[4-(2-fluoro-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
MICROCOPY RESOLUTION TEST CHART
NATIONAL BUREAU OF STANDARDS
STANDARD REFERENCE MATERIAL 1010a
(ANSI and ISO TEST CHART No. 2)
1-[4-(2-bromo-4-fluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4-fluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2,4,5-trichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4,5-trichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4,5-trichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,4,5-trichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dichloro-4-bromophenoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dichloro-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dichloro-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(2,5-dichloro-4-bromophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,5-dimethyl-7-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2-difluorobenzoyl)urea;
1-[4-(2,5-dichloro-4-fluorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,3,5-trichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(2-bromo-5-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-5-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2-bromo-5-chlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4,5-trichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4,5-trichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4,5-trichlorophenoxy)-2,3,5-trichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-3,5-dimethyl-2-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trimethylphenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,5-trimethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,5-trimethylphenyl]-3-(2-fluorobenzoyl)urea; and
1-[4-(2,4-dibromophenoxy)-2,3,5-trimethylphenyl]-3-(2,6-difluorobenzoyl)urea.

U.S. Patent Application Serial No. (D-13541-3), filed March 29, 1985, incorporated herein by reference, discloses and claims 1-(phenoxyphenyl)-3-benzoyl urea compounds which may be used in the method of this invention. Such 1-(phenoxyphenyl)-3-benzoyl urea compounds can be prepared by the methods described in the application, and are available from Union Carbide Agricultural Products Company, Inc., Research
Triangle Park, North Carolina. Illustrative of the 1-(phenoxyphenyl)-3-benzoyl urea compounds of U.S. Patent Application Serial No. (D-13541-3) which may be useful in the method of this invention include the following:

1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-bromo-6-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-fluoro-6-bromobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-bromobenzoyl)thiourea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-bromo-6-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-halo-6-fluorobenzoyl)urea.
1-[4-(2,4-dibromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2,4-dibromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-dichlorobenzoyl)thiourea;
1-[4-(2,4-dibromophenoxy)2,6-dimethyl-3-chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chloro-6-bromobenzoyl)urea;
1-[4-(2-fluoro-4-chlorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-difluorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-difluorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-difluorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2,4-difluorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(2-chloro-4-fluorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(2-chloro-4-fluorophenoxy)2,6-dimethyl-3-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-chloro-4-fluorophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-fluoro-4-bromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-fluoro-4-bromophenoxy)-2,6-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[(4-(2-fluoro-4-bromophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-bromo-4-fluorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-bromo-4-fluorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2,5-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2,5-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(2,5-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)3-(2-fluorobenzoyl)thiourea;
1-[(4-(2,5-dichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2-fluorobenzoyl)urea;
1-[(4-(2,4,5-trichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2-fluorobenzoyl)urea;
1-[(4-(2,4,5-trichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2,4,5-trichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2-chlorobenzoyl)thiourea;
1-[(4-(2,4,5-trichlorophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2,5-dichloro-4-bromophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(2,5-dichloro-4-bromophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2,4,5-dichloro-4-bromophenoxy)-2,6-dimethyl-3-chlorophenyl)-3-(2,6-difluorobenzoyl)thiourea;
1-[(4-(2,5-dichloro-4,5-dichlorophenoxy)-2,6-dimethyl-7-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(2-bromo-4,5-dichlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2-difluorobenzoyl)urea;
1-[4-(2,5-dichloro-4-fluorophenoxy)-2,6-dimethyl-3-
chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trichlorophenyl]-3-
(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trichlorophenyl]-3-
(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trichlorophenyl]-3-
(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trichlorophenyl]-3-
(2,6-dichlorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2-difluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,5-dibromo-4-chlorophenoxy)-2,3,6-trichloro-
phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-(4-(2-bromo-4-chlorophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-dichlorobenzoyl)urea;
1-(4-(2,4-dibromophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2,4-dibromophenoxy)-2,3,6-trichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-(4-(2,4-dibromophenoxy)-2,3,6-trichlorophenyl)-3-(2-fluorobenzoyl)urea;
1-(4-(2,5-dichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2-fluorobenzoyl)urea;
1-(4-(2,5-dichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2,5-dichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2,5-dichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(4-(2-bromo-5-chlorophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2-bromo-5-chlorophenoxy)-2,3,6-trichlorophenyl)-3-(2-fluorobenzoyl)urea;
1-(4-(2-bromo-5-chlorophenoxy)-2,3,6-trichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-(4-(2,4,5-trichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2,4,5-trichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2,4,5-trichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-(4-(2,4,5-trichlorophenoxy)-2,3,6-trichlorophenyl)-3-(2-fluorobenzoyl)urea;
1-(4-(2,4,5-dichlorophenoxy)-3,6-dimethyl-2-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-(2,4,5-dichlorophenoxy)-3,6-dimethyl-2-chlorophenyl)-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-3,6-dimethyl-2-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-3,6-dimethyl-2-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-3,6-dimethyl-2-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-3,6-dimethyl-2-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dibromophenoxy)-3,6-dimethyl-2-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trimethylphenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(2,4-dichlorophenoxy)-2,3,6-trimethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,6-trimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,6-trimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(2-bromo-4-chlorophenoxy)-2,3,6-trimethylphenyl]-3-(2-fluorobenzoyl)urea; and
1-[4-(2,4-dibromophenoxy)-2,3,6-trimethylphenyl]-3-(2,6-difluorobenzoyl)urea.

U.S. Patent Application Serial No. 712,195, filed March 15, 1985, incorporated herein by reference, discloses and claims alpha-cyanobenzyl phenyl benzoyl urea compounds which may be used in the method of this invention. Such alpha-cyanobenzyl phenyl benzoyl urea compounds can be prepared by the methods described in the
application, and are available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the alpha-cyanobenzyl phenyl benzoyl urea compounds of U.S. Patent Application Serial No. (D-14858) which may be useful in the method of this invention include the following:

1-[2,5-dichloro-4-(alpha-cyano-4-fluorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2-methyl-5-chloro-4-(alpha-cyano-4-chlorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dichloro-4-(alpha-cyano-4-chlorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(alpha-cyano-4-methylbenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dichloro-4-(alpha-cyano-2,4-dimethylbenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dichloro-4-(alpha-cyano-4-fluorobenzyl)phenyl]-3-(2-fluorobenzoyl)urea;
1-[2-methyl-5-chloro-4-(alpha-cyano-alpha-methyl-4-chlorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dichloro-4-(alpha-cyano-4-methylbenzyl)phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-2,5-dimethyl-4-(alpha-cyano-4-chlorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,3,5-trichloro-4-(alpha-cyano-4-methylbenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dichloro-4-(alpha-cyano-alpha-propyl-4-chlorobenzyl)phenyl]-3-(2-chlorobenzoyl)urea;
1-[2,5-dichloro-4-(alpha-cyano-4-chlorobenzyl)phenyl]-3-(2,6-dimethoxybenzoyl)urea;
1-[2-methyl-5-chloro-4-(alpha-cyano-4-chlorobenzyl)phenyl]-3-(2,6-trifluoromethylbenzoyl)urea;
1-[(2,5-dichloro-4-(alpha-cyano-4-trifluoromethoxybenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[(2,5-dichloro-4-(alpha-cyano-2-bromo-4-chloro benzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[(2-methyl-5-chloro-4-(alpha-cyano-4-bromobenzyl) phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[(2,3-dichloro-4-(alpha-cyano-4-chlorobenzyl). phenyl]-3-(2-trifluoromethoxybenzoyl)urea;
1-[(2-chloro-5-trifluoromethyl)-4-(alpha-cyano-4- cyanobenzyl)phenyl]-3-(2,6-difluorobenzoyl) urea;
1-[(2-chloro-5-ethoxycarbonyl-4-(alpha-cyano-2,4- dichlorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea;

U.S. Patent Application Serial No. 428,994, filed September 30, 1982, incorporated herein by reference. discloses and claims bicyclooxyheterocycl cycl cycl aryl urea compounds which may be used in the method of this invention. Such bicyclooxyheterocycl cycl aryl urea compounds can be prepared by the methods described in the application, and are available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the bicyclooxyheterocycl cycl aryl urea compounds of U.S. Patent Application Serial No. 428,994 which may be useful in the method of this invention include the following:

1-[(2-[(4-chloro-1-naphthoxy]-5-pyridyl)-3-(2,6-difluoro benzoyl)urea;
1-[(2-[(4-chloro-1-naphthoxy]-5-pyridyl)-3-(2-chloro-6- fluorobenzoyl)urea;
1-(3-chloro-2-[4-chloro-5,6,7,8-tetrahydro-1-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-2-[4-chloro-5,6,7,8-tetrahydro-1-naphthoxy]-5-pyridyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-2-[4-chloro-1-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-2-[4-chloro-1-naphthoxy]-5-pyridyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(2-[4-chloro-1-naphthoxy]-3-methyl-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(2-[4-chloro-1-naphthoxy]-3-methyl-5-pyridyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-2-[1,6-dibromo-2-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-2-[1,6-dibromo-2-naphthoxy]-5-pyridyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(3-chloro-2-[4-methoxy-1-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-methoxy-2-[4-methoxy-1-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-bromo-2-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-bromo-2-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-5-pyridyl)-3-(2-chlorobenzoyl)urea;
1-(3-chloro-2-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-5-pyridyl)-3-(2-chlorobenzoyl)urea;
1-(3-chloro-2-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-5-pyridyl)-3-(2,6-difluorobenzoyl)urea;
1-(2-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-3-methyl-5-pyridyl)-3-(2-chloro-6-fluorobenzoyl)urea; and
1-(2-[4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy]-3-methyl-5-pyridyl)-3-(2-chlorobenzoyl)urea.
U.S. Patent Application Serial No. 454,847, filed December 30, 1982, incorporated herein by reference, discloses and claims 1-(pyridyloxyaryl)-3-benzoyl urea compounds which may be used in the method of this invention. Such 1-(pyridyloxyaryl)-3-benzoyl urea compounds can be prepared by the methods described in the application, and are available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the 1-(pyridyloxyaryl)-3-benzoyl urea compounds of U.S. Patent Application Serial No. 454,847 which may be useful in the method of this invention include the following:

1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2-chloro-6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-trifluoromethylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,5-dibromo-phenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,5-dibromo-phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,5-dibromo-phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,5-dibromo-phenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-methyl-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-methyl-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-methyl-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-methyl-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-methylbenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-trifluoromethylbenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3-methylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridylthio)phenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridylthio)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridylthio)phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridylthio)phenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(3-chloro-5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(4-chlorobenzoyl)urea;
1-[4-(5-cyano-2-pyridyloxy)-3,6-dimethyl-5-chlorophenyl]-3-(4-chlorobenzoyl)thiourea;
1-[4-(5-cyano-2-pyridyloxy)-3,5-dibromophenyl]-3-(4-chlorobenzoyl)thiourea;
1-[4-(3-methyl-5-nitro-2-pyridyloxy)-3,5-dibromophenyl]-3-(2,6-difluorobenzoyl)thiourea,
1-[4-(3-methyl-5-nitro-pyridyloxy)-3,5-dibromophenyl]-3-(4-chlorobenzoyl)thiourea;
1-[4-(3-cyano-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-cyano-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-cyano-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-cyano-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(3-cyano-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(4-chlorobenzoyl)thiourea;
1-[4-(3-cyano-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-dimethylbenzoyl)urea;
1-[4-(3-cyano-4,6-dimethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(3-cyano-4,6-dimethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(3-cyano-4,6-dimethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-cyano-4,6-dimethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-cyano-4,6-dimethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dimethylphenyl]-3-(4-chlorobenzoyl)thiourea;
1-[4-(5-cyano-6-chloro-2-pyridyloxy)-3,5-dimethylphenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(3-chloro-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(3-chloro-5-methyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-benzoyl thiourea;
1-[4-(3-chloro-5-methyl-2-pyridyloxy)-3-methylphenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-methyl-2-pyridyloxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[(4-(3,5-dimethyl-2-pyridylthio)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(3,5-dimethyl-2-pyridylthio)-3,5-dichlorophenyl)-(3-chlorobenzoyl)thiourea;
1-[(4-(3,5-dimethyl-2-pyridylsulfonyl)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(3-methyl-5-chloro-2-pyridylthio)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(3-methyl-5-chloro-2-pyridylthio)-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)thiourea;
1-[(4-(3-methyl-5-chloro-2-pyridylsulfonyl)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(3-methyl-5-chloro-2-pyridylthio)-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(3-methyl-5-chloro-2-pyridylthio)-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)thiourea;
1-[(4-(3-methyl-5-chloro-2-pyridylthio)-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)thiourea;
1-[(4-(3-methyl-5-chloro-2-pyridylsulfuryl)-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(3-methyl-5-chloro-2-pyridylsulfonyl)-3,5-dichlorophenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(3,6-dimethyl-2-pyridylthio)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(4,6-dimethyl-2-pyridyloxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(4,6-dimethyl-2-pyridyloxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(3-chloro-5-methyl-2-pyridylthio)-2-methyl-5-chlorophenyl)-3-(2-chlorobenzoyl)urea; and
1-[(4-(3-chloro-5-methyl-2-pyridylthio)-2-methyl-5-chlorophenyl)-3-(2-chlorobenzoyl)thiourea.

U.S. Patent Application Serial No. 454,849, filed December 30, 1982, incorporated herein by reference, discloses and claims phenoxyphenyl and phenoxypyridyl urea compounds which may be used in the method of this invention. Such phenoxyphenyl and phenoxypyridyl urea compounds can be prepared by the methods described in the application, and are
available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the phenoxyphenyl and phenoxypyridyl urea compounds of U.S. Patent Application Serial No. 454,849 which may be useful in the method of this invention include the following:

1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2-chloro-6-fluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2-trifluoromethylbenzoyl)-3-(3,6-dimethyl-5-chloro-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2,6-dichlorobenzoyl)-3-(3,5-dimethyl-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2,6-difluorobenzoyl)-3-(3,5-dimethyl-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2-chloro-6-fluorobenzoyl)-3-(3,5-dimethyl-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2-chlorobenzoyl)-3-(3,5-dimethyl-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-\[(4-(4-chlorophenylsulfonyl)phenoxy)phenyl\]urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)thiourea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)thiourea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)thiourea;
1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)thiourea;
1-(2-chloro-6-fluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)thiourea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)thiourea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,5-dimethyl-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,5-dimethyl-4-[4-(4-chlorophenylsulfenyl)phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3,5-dimethyl-4-[4-(4-chlorophenylsulfinyl)phenoxylphenyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(3,5-dimethyl-4-[4-(4-chlorophenylsulfinyl)phenoxylphenyl)urea;
1-(2-chlorobenzoyl)-3-(3,5-dimethyl-4-[4-(4-chlorophenylsulfinyl)phenoxylphenyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2,6-dichlorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(2-[4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2-chlorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-bromo-5-pyridyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-chloro-5-pyridyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-chloro-5-pyridyl)urea;
1-(2-chlorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-chloro-5-pyridyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-chloro-5-pyridyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-chloro-5-pyridyl)urea;
1-(2-chlorobenzoyl)-3-(2-[4-(4-chlorophenylsulfonyl)phenoxylphenyl)-3-chloro-5-pyridyl)urea;
1-(2-chlorobenzoyl)-3-(2-[4-(4-chlorophenylsulfinyl)phenoxy]-3-chloro-5-pyridyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-[4-(4-chlorophenylsulfinyl)phenoxy]-3-chloro-5-pyridyl)thioure; 
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-dimethylbenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-dimethylbenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-trifluoromethylphenylsulfonyl)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-chloro-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-chloro-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[3-chloro-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2-chloro-6-fluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-chloro-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-chloro-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-dimethoxybenzoyl)-3-(3,6-dimethyl-5-chloro-4-[3-chloro-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-methyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-methyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-methyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)urea;
1-(2,6-dimethoxybenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-methyl-4-(2,4-dimethylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-methyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)thiourea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[3-chloro-4-(2-t-butyl-4-methylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-chloro-4-(2-t-butyl-4-methylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[3-chloro-4-(2-t-butyl-4-methylphenoxy)-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[3-isopropyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)urea;
1-(2,6-dichlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[3-isopropyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-isopropyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)urea;
1-(2-chlorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[2-isopropyl-4-(2,4-dichlorophenoxy)-phenoxy]phenyl)thiourea;
1-(2-dichlorobenzoyl)-3-(3-chloro-4-[4-(4-chloro-4-t-butylthiophenylsulfonyl)-phenoxylphenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,6-dimethyl-5-chloro-4-[4-(4-trifluoromethylphenoxy)phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3,5-dichloro-4-[4-(4-chloro-benzoyl)phenoxylphenyl)urea;
1-benzoyl-3-(3,5-dichloro-4-[4-(4-chlorophenylsulfonyl)phenoxy]phenyl)thiourea;
1-(2,6-difluorobenzoyl)-3-(3,5-dichloro-4-[2-trifluoromethyl-4-(4-chlorophenylsulfonyl)phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(2-methyl-5-chloro-4-[4-(4-chloro-phenylsulfonyl)phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(2,5-dimethyl-4-[4-(4-chlorophenylsulfonyl)phenoxy]phenyl)thiourea;
1-(2-chlorobenzoyl)-3-(3-methoxy-4-[4-(4-chlorophenylsulfonyl)phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-[4-chlorophenylthio]phenylsulfonyl)-3-methyl-phenoxy]phenyl)urea;
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-4-
[4-chlorophenylsulfonyl]phenylsulfonyl)phenoxylphenyl)
urea;
1-(2,6-difluorobenzoyl)-3-(3,5-dichloro-4-[4-(4-
chlorophenoxysulfonyl)phenoxylphenyl]urea;
1-(2,6-difluorobenzoyl)-3-(3,5-dichloro-4-[4-(4-
chlorophenylsulfonyloxy)phenoxylphenyl]urea;
1-(2,6-difluorobenzoyl)-3-(2,5-dimethyl-3-chloro-4-
[4-(4-chlorophenoxysulfonyl)phenoxylphenyl]urea;
1-(2-chlorobenzoyl)-3-(2,5-dimethyl-3-chloro-4-[4-
(4-chlorophenoxysulfonyl)phenoxylphenyl]urea;
1-(2,6-difluorobenzoyl)-3-(3-chloro-4-[4-(2-chloro-
4-[4-chlorophenylsulfonlyl]phenylsulfonyl)phenoxylphenyl)
urea;
1-(2,6-difluorobenzoyl)-3-(3-methyl-4-[4-t-
butylsulfinylphenylsulfonlyl)phenoxylphenyl]urea;
1-(2,6-difluorobenzoyl)-3-(3-methyl-4-[4-(4-
methylthiophenylsulfonyl)phenoxylphenyl]urea; and
1-(2,6-difluorobenzoyl)-3-(3-methyl-4-[4-(4-methyl-
sulfonylphenylsulfonlyl)phenoxylphenyl]urea.

U.S. Patent Application Serial No. 672,007,
filed November 15, 1984, incorporated herein by
reference, discloses and claims biphenylloxy and
biphenylalkoxy aryl acyl urea compounds which may
be used in the method of this invention. Such
biphenylloxy and biphenylalkoxy aryl acyl urea
compounds can be prepared by the methods described
in the application, and are available from Union
Carbide Agricultural Products Company, Inc.,
Research Triangle Park, North Carolina.
Illustrative of the biphenylloxy and
biphenylalkoxy aryl acyl urea compounds of U.S.
Patent Application Serial No. (D-14121) which may be
useful in the method of this invention include the following:

1-(2,6-difluorobenzoyl)-3-[3-methyl-5-chloro-4-(2-phenylphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3,5-dimethyl-4-(2-phenyl-4-bromophenoxy)phenyl]urea;
1-(2-chlorobenzoyl)-3-[3,5-dimethyl-4-(2-phenyl-4-bromophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-methylphenoxy)phenyl]urea;
1-(2-trifluoromethoxybenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-methylphenoxy)phenyl]urea;
1-(2-chlorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-trifluoromethylphenoxy)phenyl]urea;
1-(2-chlorobenzoyl)-3-[3-trifluoromethyl-4-(2-phenyl-4-methylphenoxy)phenyl]urea;
1-(2-nitrobenzoyl)-3-[3-methoxy-4-(2-phenyl-4-ethoxycarbonylphenoxy)phenyl]urea;
1-(2-chlorobenzoyl)-3-[3-ethoxycarbonyl-4-(2-phenyl-4-bromophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-cyanophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3,5-dichloro-4-[4-(2-phenyl-4-chlorophenyl)butoxy]phenyl]urea;
1-(2-chlorobenzoyl)-3-[3-carboxy-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3,5-dichloro-4-[alpha, alpha-dimethyl-2-phenyl-4-chlorobenzyl]oxy]phenyl]urea;
1-(2-chlorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
WO 86/03941

- 97 -

1-(2-chloro-6-fluorobenzoyl)-3-[2,5-dimethyl-3-chloro-2-(2-phenylphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3,5-dichloro-4-(2-phenylphenylmethoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenylphenoxy)phenyl]urea;
1-(2-chlorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenylphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3,5-dichloro-4-(alpha-methyl-(2-phenylbenzyloxy))phenyl]urea;
1-(2-chlorobenzoyl)-3-[3,5-dichloro-4-(alpha-methyl-(2-phenylbenzyloxy))phenyl]urea;
1-(2-trifluoromethylbenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2-methoxybenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2-ethoxybenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenylphenoxy)phenyl]urea;
1-(2-fluorobenzoyl)-3-[3-methoxy-5-chloro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3-trifluoromethoxy-4-(2-phenyl-4-methoxyphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-trifluoromethoxyphenoxy)phenyl]urea;
1-(2-fluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenyl-4-cyanophenoxy)phenyl]urea;
1-(2-bromo-4-phenylphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3,5-dichloro-4-(2-phenyl-4-nitrophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(3-phenylphenoxy)phenyl]urea;
1-(2,6-dichlorobenzoyl)-3-[3,5-dichloro-4-(2-phenyl-4-carboxyphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-bromo-4-phenylphenoxy)phenyl]urea;
1-(2-chlorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenylphenoxy)phenyl]thiourea;
1-(2-fluorobenzoyl)-3-[3-nitro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2-fluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-phenylphenoxy)phenyl]thiourea;
1-(2-fluorobenzoyl)-3-(2,5-dimethyl-3-chloro-4-(2-phenylphenoxy)phenyl)urea;
1-(2-ethoxybenzoyl)-3-[3-chloro-5-ethyl-4-(2-phenyl-4-tert-butylphenoxy)phenyl]urea;
1-(2-isopropylbenzoyl)-3-[2,5-dimethyl-3-carboxy-4-(2-phenyl-4-carboxyphenoxy)phenyl]urea dissodium salt;
1-(2,6-difluorobenzoyl)-3-[3-cyano-5-chloro-4-(2-phenyl-4-chlorophenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[3-isopropyl-5-methyl-4-(2-phenyl-4-ethoxyphenoxy)phenyl]urea;
1-(2,6-difluorobenzoyl)-3-[2-methyl-5-chloro-4-(2-phenylphenoxy)phenyl]urea; and
1-(2,6-difluorobenzoyl)-3-[2,5-dimethyl-3-chloro-4-(2-(4-chlorophenoxy)phenyl]urea.

U.S. Patent Application Serial No. 480,697, filed March 31, 1983, incorporated herein by reference, discloses and claims bicyclooxyaryl thiourea compounds which may be used in the method of this invention. Such bicyclooxyaryl thiourea compounds can be prepared by the methods described in the application, and are available from Union Carbide Agricultural Products Company, Inc., Research Triangle Park, North Carolina. Illustrative of the bicyclooxyaryl thiourea compounds of U.S. Patent Application Serial No.
480,697 which may be useful in the method of this invention include the following:

1-[4-(4-chloro-1-naphthoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-2,3-dichloro-5-methylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[2-chloro-4-(4-chloro-1-naphthoxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-2,5-dichloro-5-methylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-3-methylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[5-chloro-4-(4-chloro-1-naphthoxy)-2-methylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-fluorobenzoyl)thiourea;
1-[4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[5-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2-methylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl)thiourea;
1-[3-chloro-2,5-dimethyl-4-[4-(N,N-dimethylamino]-5,6,7,8-tetrahydro-1-naphthoxy)phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-2,5-dimethyl-4-[4-(N,N-dimethylamino)-1-naphthoxy]-phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3,5-dichloro-4-[4-(N,N-dimethylamino)-1-naphthoxy]phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(4-methoxy-1-naphthoxy)phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3-chloro-4-(1,6-dibromo-2-naphthoxy)phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(1,6-dibromo-2-naphthoxy)-3-methylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[3,5-dichloro-(1,6-dibromo-2-naphthoxy)phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(6-bromo-2-naphthoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(4-chloro-1-naphthoxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-(3,5-dichloro-4-(2,2-dimethyl-2,3-dihydro-7-benzofuranyloxy)phenyl)-3-(2,6-difluorobenzoyl) thiourea;
1-[2-(4-chloro-1-naphthoxy)-3-methyl-5-pyridyl]-3-(2,6-difluorobenzoyl)thiourea; and
1-[2-(4-methoxy-1-naphthoxy)-3-chloro-5-pyridyl]-3-(2,6-difluorobenzoyl)thiourea.

Other urea compounds disclosed in the art and encompassed within formula I may also be used in
the method of this invention. Such urea compounds including the preparation thereof are described in
the following U.S. patents: 4,271,171; 4,085,226; 4,139,636; 4,310,548; 4,262,020; 4,323,579;
4,234,600; 4,321,276; 4,310,694; 4,089,975; 4,380,641; 4,366,155; 4,405,552; 4,264,605;
4,293,552; 4,338,257; 4,164,581; 4,212,870; 4,276,309; 4,310,530; 4,173,637; 4,344,951;
4,321,388; 4,350,706; 3,992,553; 4,041,177; 4,275,077; 3,748,356; 3,933,908; 3,989,842;
4,160,037; 4,336,264; 4,083,977; 4,148,977; 4,160,902; 4,148,902; 4,166,124; 4,160,834;
4,064,267; 4,005,223; 4,123,449; 4,068,002; 4,194,005; 4,399,152; and 4,013,717.

Other such urea compounds encompassed within formula I which may be used in the method of
this invention are described in the following patents and published applications: EP 42533; EP
14675; German DE 3,235,419; German DE 3,219,200; German DE 3,041,947; EP 77759; EP 40179; German DE
3,241,138; EP 79311; EP 71279; EP 42732; Belgian 868,228; United Kingdom 2,028,813; German DE
2,726,684; United Kingdom 1,580,876; Japan 5 7128
673; Japan 5 8039 657; Japan Kokai 81 15272; Japan 5
Illustrative of other urea compounds which may be useful in the method of this invention include the following:

1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(4-chlorophenoxy)-3,5-dichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(4-chlorophenoxy)-3,5-dichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-fluorobenzoyl)urea;
1-[4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2-fluorobenzoyl)urea;
1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-(4-chlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(3,5-dichloro-2,4-difluorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(1,1,2,2-tetrafluoroethoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(4-trifluoromethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-trifluoromethoxyphenyl)-3-(2-chlorobenzoyl)urea;
1-[5-(4-bromophenyl)-6-methyl-2-pyrazinyl]-3-(2,6-dichlorobenzoyl)urea;
1-[5-(4-bromophenyl)-6-methyl-2-pyrazinyl]-3-(2-chlorobenzoyl)urea;
1-(4-trifluoromethyl-2-thiazoyl)-3-(2,6-dichlorobenzoyl)urea;
1-(2-fluoro-4-iodophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(2,4-difluorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-chloro-2-fluorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-(4-bromo-2-fluorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[4-(1-phenylethoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(1-phenylethoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3-chloro-4-(1-phenylpropoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-(4-benzoyloxyphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dichloro-4-benzoyloxyphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dimethyl-4-benzoyloxyphenyl)-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(4-chlorobenzoyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3-chloro-4-(5-trifluoromethyl-6-chloro-2-pyridyloxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3-methoxycarbonyl-4-(3-chloro-5-(trifluoromethyl)-2-pyridyloxy)-5-methylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dichloro-4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[2,5-dimethyl-4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(2,2-dichlorocyclopropylmethoxy)-phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dimethyl-4-(2,2-dichlorocyclopropylmethoxy)-phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dimethyl-4-(2,2-dichlorocyclopropylmethoxy)-phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[5-chloro-2-pyridinyl-N-oxide]-3-(2,6-dichlorobenzoyl)urea;
1-[4-(1,2-dichlorovinyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(1,1,2,2-tetrafluoroethoxy)phenyl]-3-(3,5-dichlorobenzoyl)urea;
1-[3-methyl-4-(1,1,2,2-tetrafluoroethoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(1,1,2-trifluoro-2-chloroethylthio)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[6-methyl-5-(alpha,alpha,alpha-trifluoro-m-tolyl)-2-pyrazinyl]-3-(2-chlorobenzoyl)urea;
1-[5-(4-methoxyphenyl)-6-methyl-2-pyrazinyl]-3-(2-chlorobenzoyl)urea;
1-(5-bromo-2-pyridinyl)-3-(2,6-dichlorobenzoyl)urea;
1-(5-(trifluoromethyl)-2-pyridinyl)-3-(2,6-dichlorobenzoyl)urea;
1-(5-chloro-2-pyridinyl)-3-(2,6-difluorobenzoyl)urea;
1-(6-(4-chlorophenylthio)-3-pyridinyl)-3-(2,6-difluorobenzoyl)urea;
1-(6-(3,5-dichlorophenoxy)-3-pyridinyl)-3-(2,6-dimethoxybenzoyl)urea;
1-(6-(3-(trifluoromethyl)-4-chlorophenylthio)-3-pyridinyl)-3-(2,6-dimethoxybenzoyl)urea;
1-(6-(3,5-bis(trifluoromethyl)phenylthio)-3-pyridinyl)-3-(2,6-dichlorobenzoyl)urea;
1-(6-(2,4-dichlorobenzoyloxy)-3-pyridinyl)-3-(2,6-difluorobenzoyl)urea;
1-(6-[2-chloro-5-(trifluoromethyl)phenoxyl]-3-pyridinyl)-3-(2,6-difluorobenzoyl)urea;
1-(6-(2,4-dichlorobenzoyloxy)-3-pyridinyl)-3-(2-fluoro-6-methoxybenzoyl)urea;
1-[6-(3,5-dichlorophenoxy)-3-pyridinyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-(6-[4-chloro-3-(trifluoromethyl)phenoxyl]-3-pyridinyl)-3-(2-fluoro-6-methoxybenzoyl)urea;
1-[5-(4-chlorophenyl)-2-pyridyl]-3-(2-chloro-6-fluorobenzoyl)urea;
1-[5-(2,4-dichlorophenyl)-2-pyridyl]-3-(2,6-difluorobenzoyl)urea;
1-[5-(4-bromophenyl)-4,6-dimethyl-2-pyridyl]-3-(2,6-difluorobenzoyl)urea;
1-[6-chloro-5-(4-chlorophenyl)-2-pyridyl]-3-(2,6-difluorobenzoyl)urea;
1-[6-bromo-5-(4-bromophenyl)-2-pyridyl]-3-(2,6-difluorobenzoyl)urea;
1-[5-(4-t-butylphenyl)-3-isoxazolyl]-3-(2-chlorobenzoyl)urea;
1-[4-bromo-5-(4-(trifluoromethyl)phenyl)-3-isoxazolyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-chloro-5-(4-(trifluoromethyl)phenyl)-3-isoxazolyl]-3-(2,6-difluorobenzoyl)urea;
1-[1-(3,5-bis(trifluoromethyl)phenyl)-5-chloro-1,6-dihydro-6-oxopyridazin-4-yl]-3-(2-chlorobenzoyl)urea;
1-[1-t-butyl-5-chloro-1,6-dihydro-6-oxopyridazin-4-yl]-3-(2,6-difluorobenzoyl)urea;
1-[1-(3,5-bis(trifluoromethyl)phenyl)-5-bromo-1,6-dihydro-6-oxopyridazin-4-yl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(1,1,2,2-tetrafluoroethoxy)phenyl]-3-(3,5-dichloro-4-pyridinecarboxyl)urea;
1-[3-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-4-methylphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3-(3-chloro-5-trifluoromethyl-2-pyridyloxy)-4-chlorophenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3-(4-(trifluoromethylphenoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-ethynlyphenyl]-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-ethynlyphenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-[3,5-dichloro-4-(N,N-diallylamino)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(N-i-propyl-N-allylamino)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dibromo-4-(N-allyl-N-methylamino)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3'-chloroallyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-alloxyphenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(1',2'-dichlorovinylxoxy)phenyl]-3-(2-fluorobenzoyl)urea;
1-[3,5-dichloro-4-(3'-bromomallylthio)phenyl]-3-(2-fluorobenzoyl)urea;
1-[4-trifluoromethylthiophenyl]-3-(2,6-difluorobenzoyl)urea;
1-(4-trifluoromethoxyphenyl)-3-(2,6-difluorobenzoyl)urea;
1-[3-chloro-4-(2-chloro-1,1,2-trifluoroethoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3-(trifluoromethyl)-4-bromophenyl]-3-(2-chlorobenzoyl)urea;
1-[3,4-(tetrafluoroethylenedioxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(3,4-(difluoromethylenedioxy)phenyl)-3-(2-fluorobenzoyl)urea;
1-[4-(t-butoxycarbonylphenyl)-3-(2-chlorobenzoyl)urea;
1-[3-chloro-4-(t-butoxycarbonyl)phenyl]-3-(2-chlorobenzoyl)urea;
1-[4-(1-hydroxy-1-trifluoromethyl-2,2,2-trifluoroethyl)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[4-(1-hydroxy-1-trifluoromethyl-2,2-trifluoroethyl)phenyl]-3-(4-chloroethenoyl)urea;
1-[2,5-dimethyl-4-(1-hydroxy-1-trifluoromethyl-2,2,2-trifluoromethyl)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(2-chloro-4-(trifluoromethylsulfonyl)phenoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3-trifluoromethyl-5-chloro-2-pyridyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(5-chloro-2-pyrimidinyloxy)phenyl]-3-(2,6-dichlorobenzoyl)urea;
1-[3,5-dichloro-4-(5-chloro-2-pyrimidinyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)phenyl]-3-(2,6-dimethoxybenzoyl)urea;
1-[2,6-dichloro-3-pyridyl]-3-(2,6-difluorobenzoyl)urea;
1-[5-chloro-2-benzothiazolyl]-3-(2,6-difluorobenzoyl)urea;
1-[5-bromo-2-benzoxazolyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(4-chlorophenoxy)phenyl]-3-(2-hydroxycarbonylbenzoyl)urea;
1-[2-ethyl-4-(ethoxycarbonyl)-5-oxazolyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3-chloro-5-(2,2-dichloro-1,1,2-trifluoroethyl)-2-pyridyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(2,3,4,5-tetrachlorophenyl)-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(N-(2-methoxyethyl)-N-methylamino)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(N-(3-methoxypropyl)-N-methylamino)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(N-n-decyl-N-methylamino)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(N-n-hexyl-N-methylamino)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(2,3,4,5-tetrafluorophenyl)-3-(2-chloro-6-fluorobenzoyl)urea;
1-(2,3,4,5-tetrafluorophenyl)-3-(2-chlorobenzoyl)urea;
1-(2,3,4,5-tetrafluorophenyl)-3-(2,6-difluorobenzoyl)thiourea;
1-(2,3,4,5-tetrachlorophenyl)-3-(2,6-difluorobenzoyl)thiourea;
1-[3,5-dichloro-4-(5-chloro-2-pyrimidinyloxy)phenyl]-3-(2,6-difluorobenzoyl)thiourea;
1-[(3,5-dichloro-4-(2-bromo-4-trifluoromethylsulfonyl-phenoxy)phenyl)-3-(2-chlorobenzoyl)thiourea;
1-[(3,4-(difluoromethylenedioxy)phenyl)-3-(2-chlorobenzoyl)thiourea;
1-[(4-chloro-3-(4-(trifluoromethyl)phenyl)-5-isoxazolyl)-3-(2-chlorobenzoyl)thiourea;
1-[(3,5-dichloro-4-(3-chloro-5-trifluoromethyl-2-pyridyloxy)phenyl)-3-(2,6-difluorobenzoyl)thiourea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-3,5-dichloro-phenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-3,5-dichloro-phenyl)-3-(2,6-difluorobenzoyl)urea;
.1-[(4-(2-chloro-4-trifluoromethylphenoxy)phenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)phenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenyl)-2,5-dimethyl-phenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenyl)-2,5-dimethyl-phenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-2,3,5-trimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-2,3,5-trimethylphenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-2,3,6-trimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-2,3,6-trimethylphenyl)-3-(2-chlorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-2,3,5,6-tetramethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-[(4-(2-chloro-4-trifluoromethylphenoxy)-2,3,5,6-tetramethylphenyl)-3-(2-chlorobenzoyl)urea;
1-(3-chloro-4-(2-chloro-4-trifluoromethylphenoxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3-chloro-4-(2-chloro-4-trifluoromethylphenoxy)-2,5-dimethylphenyl)-3-(2-chlorobenzoyl)urea; 
1-(3,5-dichloro-4-(4-trifluoromethylphenoxy)phenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3-chloro-2,5-dimethyl-4-(4-trifluoromethylphenoxy)phenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(4-trifluoromethylphenoxy)-2,3,5-trimethylphenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3,5-dichloro-4-(2,4-dichlorophenoxy)phenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(4-(2-bromo-4-chlorophenoxy)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(4-(2-bromo-4-chlorophenoxy)-3,5-dichlorophenyl)-3-(2-fluorobenzoyl)urea; 
1-(4-(2,4-dichlorophenoxy)phenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(4-(2,4-dichlorophenoxy)phenyl)-3-(2-chlorobenzoyl)urea; 
1-(4-(4-chlorophenoxy)-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3-chloro-2,5-dimethyl-4-(4-nitrophenoxy)phenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3-chloro-2,5-dimethyl-4-(4-nitrophenoxy)phenyl)-3-(2-chlorobenzoyl)urea; 
1-(3-chloro-2,5-dimethyl-4-(3,5-dichloro-2-pyridyloxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3-chloro-4-(2-chloro-4-nitrophenoxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea; 
1-(3-chloro-4-(2-chloro-4-nitrophenoxy)-2,5-dimethylphenyl)-3-(2-chlorobenzoyl)urea; 
1-(3-chloro-4-(3,5-dichloro-2-pyridyloxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-((3,5-dichloro-2-pyridyloxy)-2,5-dimethylphenyl)-3-(2-chlorobenzoyl)urea;
1-(3-chloro-4-((3-chloro-5-trifluoromethyl-1-pyridyloxy)-2,5-dimethylphenyl)-3-(2,6-difluorobenzoyl)urea;
1-(3-chloro-4-((3-chloro-5-trifluoromethyl-1-pyridyloxy)-2,5-dimethylphenyl)-3-(2-chlorobenzoyl)urea;
1-(3-chloro-4-((3-chloro-5-trifluoromethyl-1-pyridyloxy)-2,5-dimethylphenyl)-3-(2,6-dichlorobenzoyl)urea;
1-(2,6-difluorobenzoyl)-3-[4-(7,9-dimethyl dibenzofuranyl)]urea;
1-(2,6-difluorobenzoyl)-3-[4-(9-chlorodihydrobenz(f,f)oxepinyl)]urea;
1-(2,6-difluorobenzoyl)-3-[4-(2,10-dichlorodihydro-7-fluoro-3-methyl-dibenz(f,f)oxepinyl)]urea;
1-(2,6-difluorobenzoyl)-3-[4-(10-chloro-6,7-dihydro-8-fluoro-dibenz(b,g)oxocinyl)]urea;
1-[3,5-dichloro-4-(4-nitrophenoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dichloro-4-(4-nitrophenoxy)phenyl)-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(2-chloro-4-nitrophenoxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(4-cyanophenoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2,4,6-trifluorobenzoyl)urea;
1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2,4-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3,5-dichloro-2-pyridyloxy)phenyl]-3-(2,5-difluorobenzoyl)urea;
1-[3,5-dichloro-4-(3-chloro-5-trifluormethyl-2-pyridyloxy)phenyl]-3-(2,4,6-trifluorobenzoyl)urea;
1-[2,5-dimethyl-3-chloro-4-(2-bromo-4-chlorophenoxy)phenyl]-3-(2-chloro-4-fluorobenzoyl)urea;
1-[2,5-dimethyl-3-chloro-4-(2-bromo-4-chlorophenoxy)phenyl]-3-(2-chloro-5-fluorobenzoyl)urea;
1-[2,5-dimethyl-3-chloro-4-(2-bromo-4-chlorophenoxy)phenyl]-3-(2,4,6-trifluorobenzoyl)urea;
1-[3,5-dichloro-4-(3-chloro-4-cyanophenoxy)phenyl]-3-(2-chlorobenzoyl)urea;
1-(3,5-dichloro-4-(3-chloro-5-bromo-2-pyridyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
1-(3,5-dichloro-4-(3,5-dibromo-2-pyridyloxy)phenyl]-3-(2,6-difluorobenzoyl)urea;
3-P-chlorophenyl-5-(2,6-difluorobenzoyl)-2,3,5,6-tetrahydro-4H-1,2,5-oxadiazinone-4;
1-[4-chlorophenyl]-3-(2,6-difluorobenzoyl)-hydantion; and
1-[4-chlorophenyl]-3-(2,6-difluorobenzoyl)-parabanic acid.

It is appreciated that the particular urea compounds listed hereinabove are illustrative of urea compounds which may be used in the method of this invention. The method of this invention is not to be construed as being limited only to the use of these urea compounds; but rather, the method of this invention includes those urea compounds encompassed within formula 1 hereinabove.
The term "parasites" as used in the specification and claims is meant to encompass all endoparasites and ectoparasites of warm-blooded animals as well as pests that breed in the manure of the animals.

In particular, representative parasites which may be controlled by the method of this invention include members of the Arthropoda, including mites of the suborders Mesostigmata, Sarcoptiformes, Trombidiformes and Onychochopalpida; sucking and biting lice of the orders Anoplura and Mallophaga; ticks of the families Ixodidae and Argasidae; fleas of the families Pulicidae, Ceratophyllidae, and others; *Cimex* and other Hemiptera; Triatoma and other Heteroptera; and myiasis-related fly larvae and blood sucking adults (including mosquitoes) of the suborders Brachycera, Cyclorrhapha and Nematocera. Representative also are helminths included in the Nematoda (Strongylida, including but not limited to Strongyloidea, Ancylostomatoidea, Trichostrongyloidea and Metastrongyloidea; Ascarida [Ascaris]; Filariina, such as but not limited to *Onchocerca* and *Dirofilaria*; Rhabditida; and Trichinellida); Cestoidea, especially Cyclophyllidea, and Trematoda, including Strigeatoidea such as *Schistosoma*; Echinostomida such as *Fasciola*; and Plagiorchiida such as *Paragonimus*. Other parasites which may be controlled by compounds represented by generic formula 1 include Acanthocephala such as *Macracanthorhynchus*, *Onicola* or *Moniliformis*, and Pentastomida, especially *Linguatula*; and Protozoa, especially Coccidia such as *Eimeria* and *Plasmodium*,
Piroplasmea such as *Babesia*; Toxoplasmea such as *Trypanosoma*; Trichomonadidae such as *Trichomonas* and Entamoebidae such as *Entamoeba*.

Illustrative of specific parasites of various host animals which may be controlled by the method of this invention include arthropods such as mites (mesostigmatids, itch, mange, scabies, chiggers), ticks (soft-bodied and hard-bodied), lice (sucking, biting), fleas (dog flea, cat flea, oriental rat flea), true bugs (bed bugs, kissing bugs), bloodsucking adult flies (horn fly, horse fly, stable fly, black fly, deer fly, louse fly, tsetse fly, punkies, mosquitoes), and parasitic fly maggots (bot fly, blow fly, screwworm, cattle grub, fleeceworm); helminths such as nematodes (threadworm, lungworm, hookworm, whipworm, nodular worm, stomach worm, round worm, pinworm, heartworm), cestodes (tapeworms) and trematodes (liver fluke, blood fluke); protozoa such as coccidia, trypanosomes, trichomonads, amoebas and plasmodia; acanthocephalans such as thorny-headed worms; and pentastomids such as tongueworms.

Also, the acyl urea compounds of formula 1 may be administered to a vertebrate definitive host for control of arthropod intermediate hosts which feed on the feces of the vertebrate host. For example, an arthropod intermediate host of helminths and other disease and nuisance-causing organisms, which arthropod host feeds on the feces of a vertebrate host treated with one or more acyl urea compounds of formula 1, can be controlled by the method of this invention.
The method of parasite control of this invention is of the systemic type. The acyl urea compounds of formula I are useful parasiticides for the systemic control of endoparasites and ectoparasites which feed on living tissues of warm-blooded animals. These acyl urea compounds have the ability to permeate the living tissues of a host animal to which one of the acyl urea compounds is administered. Endoparasites and ectoparasites which consume blood or other living tissues of the host animal ingest or contact the acyl urea compound with which the tissue is permeated, and are thereby killed. It is believed that the blood is an agency through which the acyl urea compound is dispersed through the host animal. However, as illustrated in the working examples hereinafter, systemic control is observed from topical (pour-on) applications of acyl urea compounds of formula I, thereby indicating that the acyl urea compounds permeate other tissues as well as blood. The acyl urea compounds of formula I are also effective for the control of pests which breed in the manure of warm-blooded animals.

Endoparasites and ectoparasites which feed on the living tissues of animals are killed by the acyl urea compounds of formula I. Those parasites which suck the host animal's blood, those which burrow into and feed on the animal's tissue, and those, like the larvae of bot flies, which enter a natural orifice of the host, attach to the mucous membranes, and feed therefrom may be all equally effectively killed.
It is understood that the parasites mentioned above are not confined to the single host animal with which each is generally identified. Certain parasites inhabit various hosts, although most parasites have a favorite host. For example, the mange mite attacks at least horses, hogs, mules, humans, dogs, cats, foxes, rabbits, sheep, and cattle. Horseflies freely attack horses, mules, cattle, hogs, dogs, and most other animals. Use of the acyl urea compounds of formula 1 kills parasites of the types described above growing in various host animals. For example, these acyl urea compounds may be effective in cats, goats, camels, and zoo animals.

The parasites controlled by the method of this invention normally affect warm-blooded animals. Representative of animals affected are humans and domestic animals such as cattle, horses, sheep, goats, poultry, swine, dogs, and cats.

The time, manner, and rates at which the acyl urea compounds are effectively administered may be varied over a wide range. The acyl urea compounds can be administered to the animals at rates from about 0.01 to about 1000 mg./kg of animal body weight. The best rate for killing a given parasite infesting a given animal must be determined individually, but it is generally found that in most cases the optimum rate is within the preferred range of from about 0.25 to 100 mg./kg. of animal body weight. The optimum rate for a given instance depends on such factors as the activity of the specific acyl urea compound employed, the method of administration, the level of infestation, the duration of control desired, the health of the animal to be treated, the susceptibility of the parasite of primary concern, the expense which can be borne by the animal, and the degree of control desired.
The acyl urea compounds of formula \(1\) are effective when administered at any time of year to animals of any age. It is possible to administer these acyl urea compounds to the animals continuously, as by constant feeding of a diet which contains one of the acyl urea compounds, and thus assure that all parasites which contact the treated animal will be killed. Such administration, however, may not be economical, and it will often be found best to administer the acyl urea compound at such times as to give the best return of parasite control for the compound expended. Certain parasites have a known active season when they attack animals. If such a parasite is of primary importance, the acyl urea compounds can be used only during the active season with assurance of year-round control of the parasite. Other parasites infest and bite animals essentially the year round. Control of such parasites can still be accomplished with relatively brief periods of administration by administering the acyl urea compounds to all the animals on a farm or in an area for a short period of time, such as for a few weeks. All the parasites of a generation are thus killed, and the animals can be expected to remain parasite-free for a considerable length of time.

Oral administration of an acyl urea compound of formula \(1\) may be performed by mixing the acyl urea compound in the animal's feed or drinking water, vitamin or mineral supplement, or by administering oral dosage forms such as drenches, tablets, bolus, salt block or capsules. Such oral means of administration are well known in the art.
Percutaneous administration may be conveniently accomplished by subcutaneous, dermal, intramuscular, and even intravenous injection of the injection devices as well as needle-less air-blast injection devices may be useful. Percutaneous administration is also meant to include surface treatment of the animal by dipping (drenching), pour-on, spraying, or dusting. As illustrated in the working examples hereinafter, systemic control is observed from surface treatments of the acyl urea compounds of formula 1.

Sustained action of the acyl urea compounds of formula 1 can be obtained by formulating the acyl urea compound in a matrix which will physically inhibit dissolution. The formulated matrix is injected into the body where it remains as a depot from which the compound slowly dissolves. Matrix formulations, now well known in the art, can be formulated in waxy semisolids such as vegetable waxes and high molecular weight polyethylene glycols.

Even more effective sustained action may be obtained by introducing into the animal an implant containing one of the acyl urea compounds of formula 1. Such implants are well known in veterinary art. Such administration can be highly economical and efficacious, because a properly designed implant maintains a constant concentration of the acyl urea compound in the tissues of the host animal. An implant can be designed to supply the acyl urea compound for several months, and can easily be inserted in the animal. No further handling of the animal or concern over the dosage is necessary after the insertion of the implant.
Percutaneous administration of formula 1 acyl urea compounds may be carried out in the ways usual in the animal veterinary art. For example, it may be practical to formulate an injectable suspension of the acyl urea compound as a fine powder, suspended in a formulation of physiologically–acceptable liquid carriers, surfactants, and suspending agents.

The liquid carriers can be, for example, a vegetable oil such as peanut oil, corn oil, or sesame oil, a glycol such as a polyethylene glycol, or water, depending on the acyl urea compound chosen.

Suitable physiologically acceptable adjuvants may be necessary to keep the acyl urea compound of formula 1 suspended. The adjuvants can be chosen from among the emulsifiers, such as salts of dodecylbenzene sulfate and toluenesulfonate, ethylene oxide adducts of alkylphenol, and oleate and laurate esters, and from the dispersing agents such as salts of naphthalenesulfonate, lignin sulfonate and fatty alcohol sulfates. Thickeners such as carboxymethyl cellulose, polyvinylpyrrolidone, gelatin and the alginates may also be used as adjuvants for injectable suspensions. Many classes of surfactants, as well as those which have been discussed above, serve to suspend the acyl urea compound. For example, lecithin and the polyoxyethylene sorbitan esters may be useful surfactants.

The acyl urea compounds used in the method of this invention can be administered to the animal directly in an undiluted form or the active compound
may be formulated with a suitable carrier prior to use. A suitable carrier for purposes of this invention is a carrier which will have no deleterious effect on the animal being treated or on the method or results of the invention. The carrier can be either solid or liquid depending upon the compound and method of administration selected by one skilled in the art. Suitable liquid carriers include water, N-methylpyrrolidone, vegetable oil, and other non-toxic liquid carriers with or without surfactants. Liquid concentrates can be prepared by dissolving one of the active compounds with a non-toxic solvent and dispersing the toxicants in water with the aid of suitable surface active emulsifying and dispersing agents. The choice and quantity of dispersing and emulsifying agents employed is dictated by the ability of the agent to facilitate dispersion of the toxicant. For a solid formulation such as a powder, tablet, paste, or the like, the active ingredient is dispersed in and on an appropriate carrier such as clay, talc, or dry molasses.

Formulations useful in the conduct of the method of this invention can also contain other optional ingredients such as stabilizers or other biologically active compounds, insofar as they do not impair or reduce the activity of the compounds towards the parasites and do not harm the animal being treated.

The formulation of veterinary additives in animal feed is an extremely well-known art. It is usual to formulate the compound first as a premix in
which the acyl urea compound of formula 1 is dispersed in a liquid or particulate solid carrier. The premix may conveniently contain from about 1 to 400 grams of compound per pound of carrier, depending on the desired concentration of the feed.

The premix is in turn formulated into feed by dispersing it in the feed mixture in a conventional mixer. The correct amount of acyl urea compound, and hence of premix, to mix in the feed can be easily computed by taking into account the weight of the animals, the approximate amount each animal eats per day, and the concentration of the compound in the premix.

Likewise, the amount of acyl urea compound to administer in the drinking water of animals can be computed by taking into account the animal's weight and the amount each animal drinks per day. It is convenient to use a suspendable formulation of the desired acyl urea compound. The formulation may be a suspension in the concentrated form which is mixed into the drinking water, or may be a dry preparation which is mixed with and suspended in the drinking water. In either event, the acyl urea compound should be in a finely-powdered form. The acyl urea compounds can easily be formulated into tablets and capsules according to the conventional methods known in the art. Drench formulations comprise the acyl urea compound dissolved or dispersed in an aqueous liquid mixture. It is convenient and efficacious to use a dispersion of the acyl urea compound made in the same way that the drinking water formulations discussed above are made.
It is appreciated that the acyl urea compounds of generic formula 1 may be used in combination with each other and/or with one or more biologically active materials, e.g., medicinal drugs, vertebrate toxicants, growth promoters, vitamins, minerals, etc., known in the veterinary art. Such combinations may be used for the known or other purpose of each ingredient and may provide a synergistic effect. The combinations may be administered according to the known or other methods for administering the biologically active materials, e.g., orally, implant, pour-on, injection, etc.

Illustrative of medicinal drugs known in the veterinary art which may be used in combination with the acyl urea compounds of formula 1 include, for example, the following: benzylbenzoate + lindane; benzylbenzoate; chloroform + rotenone; caraway oil + petrolatum; chlordane; chlorpyrifos; crotoxyphos + dichlorvos; coumaphos; diazinon; dichlorvos; dimethoate; dioxathion; famophos; fenthion; calcium polysulfide; lindane; malathion; methoprene; methoxychlor; benzylbenzoate + lindane + chlorobutanol + phenylmercuric chloride; naled; nicotine sulfate; phosmet; pyrethrins; synergized pyrethrins such as piperonyl butoxide and the like; stabilene, MGK 11 or MGK 326; stirofos + dichlorvos; ronnel; rotenone; stirofos; amicarbalide diisethionate; amprolium; decoquinate; carbarsone; dimetridazole; diminazine aceturate; ethidium bromide; furazolidone; homidium bromide; imidocarb dipropionate; isometamidium; lasalocid; nitarsone; pamaquine napthoate; phenamidine isethionate;
phenanthridinium; butynorotate + sulfanitran + dinosed + roxarsone; quinacrine hydrochloride; quinacrine hydrochloride + diiodohydroxyquin; quinapyramine sulfate; quinuronium sulfate; ronidazole; sulfamethazine; sulfaquinoxalin sodium; suramin; amphotericin B; griseofulvin; nystatin; benzalkonium chloride + chlordantoin; salicylic acid + benzoic acid; thiabendazole; albendazole; tolnaftate; neomycin + bactracin + polymyxin B sulfate + hydrocortisone acetate; chloroform + rotenone; triethylnolamine polypeptide oleate condensate + chlorobutanol + propylene glycol; ephedrine sulfate; hydrocortisone acetate + neomycin sulfate; 2-mercaptobenzothiazole + benzocaine + 2-chloro-4-phenlphenol; prednisolone acetate + sodium sulfacetamide + neomycin sulfate; nystatin + neomycin sulfate + thiostrepton + triamcinolone acetonide; resorcinol + zinc oxide + calamine + oil of cade + pyroligneous acid + zinc hydride; resorcin + bismuth subgallate + bismuth subnitrate + zinc oxide + calamine + oil of cade; penicillin G sodium; phenylephrine hydrochloride; phenylmercuric nitrate; dimethyl pthalate + cotton seed oil; thymol + ethyl alcohol; thiabendazol + neomycin sulfate + dexamethasone; albendazole + neomycin sulfate + dexamethasone; thiabendazole-piperazine phosphate; aracoline acetarsol; arecoline hydrobromide; arsenamide; bephenum embonate; bephenum hydroxynaphthoate; bunamide hydrochloride; N-butyl chloride; canbendazole; carbondisulfide; diethylcarbamazine citrate; dichlorophen; dichlorophen-toluene; disophenol; dithiazanine
iodide; fenbendazole; haloxon; hexylresorcinol; hygromycin; lead arsenate; levamisole; mebendazole; oxfendazole; niclosamide; piperazine-carbendisulfide complex; phenothiazine; phthalofyne; piperazine hexahydrate; piperazine salt of niclosamide, adipate, chloride, citrate, dihydrochloride, hexahydrate, phosphate, and sulfate; pyrantel embonate; pyrantel pamoate; pyrantel tartrate; sudadimethoxine-ormetoprin; styrylpyridinium-diethycarbamazine; themium closylate; thiophanate; tylosin; tetracycline; carbadox; lincomycin; flavomycin; bacitracin; virginiamycin; amprolium; ethopabate; robenidine; arprinocid; monensin; chlorotetracycllin; sulfa compounds; dibutyltin dialurate; lasalocid; salinomycin; narasin; diethylcarbanazine; phenothiazine; metronidazole; nitrothiazole; ipronidazole; sulfamethazine; amprolium; nitroturazone; diaiveridine + sulfaquinoxaline; sulfaguanidine; chloroquine; quinacrine; levamisole; ivermectin; avermectin; milbemycin; methyridine; crotamiton; ampicillin; lincomycin; cephalothrin; and amoxicillin trihydrate + clavulanate potassium.

Illustrative of vertebrate toxicants known in the veterinary art which may be used in combination with the acyl area compounds of formula 1 include, for example, the following: coumafuryl; pivalin; warfarin; dicumarol; coumachlor; coumatetralyl; chloropaciton; diphenacoun; bromethacoum; bromethalin; brodifacoum; bromodiolone + sulpha quinoxaline; bromodiolone; calciferol;
calciferol + any anticoagulant such as difenacoum; calcium carbide; calcium phosphide; alpha-chloralose; chlorophacinone; chloropicrin; cholecalciferol; coumafuryl; coumatetryl; crimidine; difenacoum; endrin; floroacetamide; lindane + teenazine + thiourea; acetylsalicylic acid + any anticoagulant such as difenacoum; pindone + warfarin; pyranocoumarin; scillirocide and toxic extracts of Urginea; strychine; thalium sulfate; and zinc phosphide. Such combinations may be administered, for example, as a bait or tracking powder in which an undesirable vertebrate host can be eliminated by the toxicant and, at the same time, any ectoparasites from the dead vertebrate host can be killed and prevented from moving to infect man or his domestic animals by the systemic action of the acyl urea compound.

Illustrative of growth promoters known in the veterinary art which may be used in combination with the acyl urea compounds of formula \( \text{I} \) include, for example, the following: 2,4-dihydro-6-phenyl-1H-s-triazolo[4,3-a][1,4]benzodiazepin-1-ones as described in U.S. Patent 3,786,149; 2,3-dibromopropanol as described in U.S. Patent 3,934,037; tertiary alcohols and their urea, oxyurea and carbamate derivatives as described in U.S. Patent Reissue 28,691; tropine benzohydryl ether or 3-quinuclidinyl benzohydryl ether as described in U.S. Patent 3,978,221; polynaloaldehyde aldol condensation products as described in U.S. Patent 3,755,601; nituroxazide as described in U.S. Patent 4,093,747;
phenylalkylsulfamide derivatives as described in U.S. Patent 3,860,723; quinoxalinedi-N-oxides as described in U.S. Patent 4,012,512; imines of quinoxalinedi-N-oxides as described in U.S. Patents 3,997,665, 3,983,235, 3,984,553, 3,984,550, 3,984,547 and 3,983,236; streptomycetes antibiotics including avermectin, ivermectin, milbemycin, valinomycin and the like as described in U.S. Patent Reissue 28,700 and U.S. Patents 4,083,964, 3,950,515, 4,073,884, and 3,790,667; mixtures of mocimycin and bacitracin as described in U.S. Patent 3,928,573; vancomycin, ristocetin and physiologically acceptable salts and esters thereof as described in U.S. Patent 3,928,571; dibenzazepines as described in U.S. Patent 3,812,255; hydroxyphenethanol amines as described in U.S. Patent 3,818,101; bromo FES derivatives and chloro FES derivatives as described in U.S. Patents 3,751,431 and 3,764,614; tetrahydroiminobenzol[G]thien-4-ylureas as described in U.S. Patent 4,060,627; pyrazoles as described in U.S. Patent 3,980,794; thiolcarbamates as described in U.S. Patent 3,781,440; iodonium salts as described in U.S. Patent 4,087,554; 3-nitro-pyrazole-gamma-carboxylic acid nitrofurfurylidene hydrazides as described in U.S. Patent 4,235,995; macrotrrolides such as nonactin, monactin, dinastin, trinactin, tetranaactin, polynactin and the like as described in DE 2835936 A 790222 79091; penicillin, cephalosporin and analogs thereof as described in DE 7932285A 790125 79051; and spiramycin and acid esters thereof as described in DE 861603A 780607 78241.
Illustrative of vitamins known in the veterinary art which may be used in combination with the acyl urea compounds of formula 1 include, for example, vitamins A, B, C, D, E, thiamine and the like.

Illustrative of essential minerals known in the veterinary art which may be used in combination with the acyl urea compounds of formula 1 include, for example, sodium, calcium, magnesium, copper, zinc, cobalt, phosphorous, sulfur, chlorine, iodine and the like.

It is understood that the above biologically active materials are merely representative of a wide variety of materials that can be combined with the acyl urea compounds of formula 1 for use in the method of this invention.

The inherent low mammalian toxicity of these compounds by all routes of entry makes them ideally suited for medical and veterinary parasite control. There may also be a reduced probability of pest resistance developing to this chemical class relative to compounds which are dependent upon cholinesterase-inhibition. When administered orally, by pour-on, by implant, or by injection, the costly expense of spray equipment and labor for spraying are eliminated. Problems associated with coverage and penetration of feathers or hair are also avoided.

The method of this invention including the preparation of acyl urea compounds of formula 1 utilized therein is illustrated by the following examples.
EXAMPLE I

Preparation of 1-[(3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl) urea

Part A: Preparation of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethyl-1-nitrobenzene

Into a solution containing 9.10 grams (51.1 mmol) of 4-chloro-1-naphthol and 10.12 grams (46.0 mmol) of 3-chloro-2,5-dimethyl-1-nitrobenzene in 22 milliliters of dimethylsulfoxide at room temperature and under a nitrogen atmosphere was added 8.89 grams (64.4 mmol) of anhydrous potassium carbonate. The resulting mixture was heated to a temperature of 130°C and maintained at that temperature for a period of 6 hours. The reaction mixture was then filtered and the filtrate was diluted with 20 milliliters of methanol and allowed to cool to room temperature. Crystallization occurred during this cooling period. The filtrate was further cooled in an ice bath, filtered and the collected crystals washed with cold methanol. Recrystallization from a hexane–ethyl acetate mixture afforded 7.92 grams of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethyl-1-nitrobenzene as a white solid having a melting point of 135°C-137°C.

Part B: Preparation of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylaniline

Into a solution containing 13.8 grams (38.1 mmol) of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethyl-1-nitrobenzene prepared in Part A in 400 milliliters of toluene was added 2.0 grams of a 5% platinum on carbon catalyst. The resulting
mixture was placed in a one liter rocking Parr hydrogenation vessel. Hydrogen was introduced into the vessel to a pressure of 120 psi. The hydrogen pressure was maintained between 100-120 psi until hydrogen uptake ceased. The contents of the reactor vessel were then removed and filtered. Removal of the solvents from the filtrate afforded 11.6 grams of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylaniline as a white solid having a melting point of 118°C-125°C.

Part C: Preparation of 1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea

Into a solution containing 3.0 grams (9.0 mmol) of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylaniline prepared in Part B in 25 milliliters of toluene which solution was warmed to a temperature of 50°C and placed under a nitrogen atmosphere was slowly added a solution containing 2.5 grams (13.8 mmol) of 2-chlorobenzoyl isocyanate in 5 milliliters of toluene. After stirring for 1 hour at a temperature of 50°C, the reaction mixture was cooled to room temperature, placed in an ice bath, and filtered through a Buchner funnel. The collected solid was dried under vacuum at a temperature of 50°C to afford 3.15 grams of 1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea as a white solid having a melting point of 187°C. Elemental analysis of the white solid indicated the following:
EXAMPLE II

Preparation of 1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl) urea

Into a solution containing 3.0 grams (9.0 mmol) of 3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylaniline prepared in Part B of Example I in 25 milliliters of toluene which solution was warmed to a temperature of 50°C and placed under a nitrogen atmosphere was slowly added a solution containing 2.5 grams (13.5 mmol) of 2,6-difluorobenzoyl isocyanate in 5 milliters of toluene. After stirring for one hour at a temperature of 50°C, the reaction mixture was cooled to room temperature, placed in an ice bath, and filtered through a Buchner funnel. The collected solid was dried under a vacuum at a temperature of 50°C to afford 2.89 grams of 1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea as a white powder having a melting point of 187°C. Elemental analysis of the white powder indicated the following:

Analysis: \( \text{C}_{26}\text{H}_{18}\text{Cl}_2\text{F}_2\text{N}_2\text{O}_3 \)
Calculated: C, 60.59; H, 3.52; N, 5.44
Found: C, 59.95; H, 3.28; N, 6.19

This compound is referred to hereinafter as Compound 2.
EXAMPLE III

Preparation of 1-[3-chloro-4-(2,4-dimethylphenoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea

Part A: Preparation of 4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chloronitrobenzene

Into a round bottom reaction flask equipped with a magnetic stirrer, condenser, thermometer and nitrogen inlet was added 50.0 grams (0.23 mol) of 2,5-dimethyl-3,4-dichloronitrobenzene, 38.5 grams (0.32 mol) of 2,4-dimethylphenol, 50 grams (0.36 mol) of potassium carbonate and 125 milliliters of dimethylformamide. The reaction mixture was heated to a temperature of 90°C-100°C and maintained at this temperature for a period of 72 hours. The reaction mixture was then cooled, filtered and concentrated to give a dark oil. The dark oil was partitioned between toluene (300 milliliters) and 4% sodium hydroxide (250 milliliters) and then separated. The organic layer was washed with water and brine, dried over Na₂SO₄ and concentrated to give a dark oil. The dark oil was triturated with hexane and a solid was formed. The solid was washed with cold hexane and vacuum dried to give 47.8 grams (0.16 mol) of 4-(2,4-dimethylphenoxy)2,5-dimethyl-3-chloronitrobenzene as a tan powder having a melting point of 78°C-80°C. Elemental analysis of the tan powder indicated the following:

Analysis: \( \text{C}_{16}\text{H}_{16}\text{ClNO}_3 \)
Calculated: C, 62.85; H, 5.27; N, 4.58
Found: C, 63.47; H, 5.35; N, 4.49
Part B: Preparation of 4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chloroaniline

Into a solution of 4-(2,4-dimethylphenoxy)-2,5-dimethylphenyl-3-chloronitrobenzene (25 grams, 81.8 mmol) prepared in Part A in toluene (250 milliliters) was added a 5% platinum on carbon catalyst (1.0 gram). The resulting mixture was placed in a one liter rocking Parr hydrogenation vessel. Hydrogen was introduced into the vessel to a pressure of 20 psi and maintained at this pressure for a period of 4.5 hours. The contents of the reactor vessel were then removed, filtered through celite and concentrated to give 24.5 grams of an oil. After high vacuum drying, the oil turned to a pinkish solid which was identified as 4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chloroaniline having a melting point of 86°C-88°C. Elemental analysis of the solid indicated the following:

Analysis: C₁₆H₁₈ClNO
Calculated: C, 69.69; H, 6.53; N, 5.08
Found: C, 70.02; H, 6.60; N, 5.68

Part C: Preparation of 1-[3-chloro-4-(2,4-dimethylphenoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea

Into a solution of 4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chloroaniline (1.75 grams, 6.3 mmol) prepared in Part B in 7 milliliters of toluene, which solution was warmed to a temperature of 50°C, was added 1.14 grams (6.2 mmol) of 2,6-difluorobenzoyl isocyanate in 1.5 milliliters of toluene. The reaction mixture was heated to a temperature of 90°C and maintained at this temperature for 0.5
hours. After cooling to room temperature, the reaction mixture was diluted with 20 milliliters of hexane and then cooled in an ice bath. A precipitate formed which was collected by filtration and washed with a cold solution of 50% hexane in toluene. After vacuum drying, a white solid (1.26 grams) which was identified as 1-[3-chloro-4-(2,4-dimethylphenoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea was obtained having a melting point of 183°C-185°C. Elemental analysis of the white solid indicated the following:

Analysis: \( \text{C}_2\text{H}_{11}\text{N}_2\text{O}_3\text{ClF}_2 \)

Calculated: C, 62.82; H, 4.61; N, 6.10

Found: C, 62.94; H, 4.88; N, 6.03

This compound is referred to hereinafter as Compound 3.

EXAMPLE IV
Preparation of 1-[4-(4-chloro-1-naphthoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea

Part A: Preparation of 4-(4-chloro-1-naphthoxy)-3,5-dichloronitrobenzene

Into a mixture of 3,4,5-trichloronitrobenzene (30.26 grams, 134.0 mmol), anhydrous potassium carbonate (28.02 grams, 203.0 mmol) and dimethylformamide (300 milliliters) was added solid 4-chloronaphthol (30.04 grams, 168.0 mmol). The mixture was stirred under a nitrogen atmosphere and heated overnight at a temperature of 105°C. The mixture was then allowed to cool and the dimethylformamide was evaporated under reduced pressure. The residue was dissolved in 1:1 ethyl acetate:ethyl ether (1.5 liters) and washed
sequentially with water, 5% NaOH solution, water and brine. The organic layer was dried over Na₂SO₄ and concentrated under reduced pressure to afford a crude product as a dark brown solid (48.73 grams). Decolorization with carbon and recrystallization (2x) from hexane:ethyl acetate afforded pure 4-(4-chloro-1-naphthoxy)-3,5-dichloronitrobenzene (15.84 grams, 42.98 mmol, 32%) as a yellow solid having a melting point of 121.5°C-123°C. This material was characterized by 'H-NMR spectroscopy as follows: 'H-NMR (CDCl₃) 8.70-8.10 (m, 2H), 8.38 (s, 2H), 7.90-7.50 (m, 2H), 7.33 (d, J=9Hz, 1H), 6.25 (d, J=9Hz, 1H).

Part B: Preparation of 4-(4-chloro-1-naphthoxy)-3,5-dichloroaniline

Into a 500 milliliter Parr bottle was charged 4-(4-chloro-1-naphthoxy)-3,5-dichloronitrobenzene (15.84 grams, 42.97 mmol) prepared in Part A and toluene (250 milliliters). The bottle was purged thoroughly with nitrogen and 5% platinum on carbon (0.50 grams) was added immediately following the purge. The reaction mixture was then hydrogenated for 1.5 hours on a rocking Parr hydrogenator at a pressure of 90-100 psi hydrogen at ambient temperature. The reaction mixture was filtered through a pad of celite and the filtrate concentrated under reduced pressure to afford pure 4-(4-chloro-1-naphthoxy)-3,5-dichloroaniline (13.99 grams, 41.31 mmol, 96%) as a white solid having a melting point of 145°C-147°C. Elemental analysis of the white solid indicated the following:
Part C: Preparation of 4-(4-chloro-1-naphthoxy)-2,3,5-trichloroaniline

Into a magnetically stirred solution of 4-(4-chloro-1-naphthoxy)-3,5-dichloroaniline (6.00 grams, 17.72 mmol) prepared in Part B in benzene (50 milliliters) was added solid N-chlorosuccinimide (2.84 grams, 21.20 mmol). The mixture was heated to reflux for 1 hour, cooled and filtered. The solvent was evaporated under reduced pressure and the crude product when subjected to flash column chromatography (3:1 hexane:ethyl acetate) afforded pure 4-(4-chloronaphthoxy)-2,3,5-trichloroaniline (3.70 grams, 9.92 mmol, 56%). Elemental analysis of the product indicated the following:

Analysis:  \( \text{C}_{16}\text{H}_{10}\text{Cl}_4\text{NO} \)
Calculated: C, 51.51; H, 2.43; N, 3.75
Found: C, 51.78; H, 2.49; N, 3.69

Part D: Preparation of 1-[4-(4-chloro-1-naphthoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea

Into a magnetically stirred solution of 4-(4-chloro-1-naphthoxy)-2,3,5-trichloroaniline (1.60 grams, 4.29 mmol) prepared in Part C in dichloromethane (20 milliliters) was added neat 2,6-difluorobenzoyl isocyanate (786 milligrams, 4.29 mmol). The heterogeneous mixture was stirred at room temperature for about 15 minutes and then chilled to a temperature of 0°C. The precipitate was collected on a fritted disc and washed with 3:1
hexane:ethyl acetate to afford pure
1-[4-(4-chloro-1-naphthoxy)-2,3,5-trichlorophenyl]-3-
(2,6-difluorobenzoyl) urea (1.9 grams, 5.41 mmol,
80%) as a white powder having a melting point of
213°C-215°C. Elemental analysis of the white powder
indicated the following:

<table>
<thead>
<tr>
<th>Analysis:</th>
<th>C&lt;sub&gt;24&lt;/sub&gt;H&lt;sub&gt;12&lt;/sub&gt;Cl&lt;sub&gt;4&lt;/sub&gt;F&lt;sub&gt;2&lt;/sub&gt;N&lt;sub&gt;2&lt;/sub&gt;O&lt;sub&gt;3&lt;/sub&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calculated:</td>
<td>C, 51.83; H, 2.17; N, 5.04</td>
</tr>
<tr>
<td>Found:</td>
<td>C, 51.41; H, 2.54; N, 4.91</td>
</tr>
</tbody>
</table>

This compound is referred to hereinafter as Compound 4.

**EXAMPLE V**

**Preparation of 2-(2-chlorobenzoyl)-3-
[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethyl-
phenyl]-1,1-dimethylguanidine**

**Part A: Preparation of 1-(2-chlorobenzoyl)-3-[3-
chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-
S-methylisothiourea**

Into a solution containing 1.2 grams of
1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethyl-
phenyl]-3-(2-chlorobenzoyl)thiourea and 200
milliliters of dry tetrahydrofuran under an
atmosphere of nitrogen and cooled to 10°C was
added 6.3 milliliters (6.4 grams) of 1,8-diaza-
bicyclo[5.4.0]undec-7-ene. The resulting solution
was warmed to room temperature and stirred for 20
minutes. After cooling the resultant mixture to a
temperature of 10°C, 2.5 milliliters (5.7 grams)
of methyl iodide was added. After 5 minutes at
10°C, the mixture was warmed to room temperature,
stirred for 2 hours, and then poured into 500
milliliters of ethyl acetate. This solution was
MICROCOOP® RESOLUTION TEST CHART
NATIONAL BUREAU OF STANDARDS
STANDARD REFERENCE MATERIAL 1010a
(ANSI and ISO TEST CHART No. 2)
washed twice with water and the organic layer dried over anhydrous sodium sulfate. Removal of the solvents afforded 21.14 grams of an orange solid. Recrystallization of the orange solid using a hexane-ethyl acetate mixture afforded 11.29 grams of 1-(2-chlorobenzoyl)-3-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-S-methylisothiourea as white crystals having a melting point of 165°C-168°C. Elemental analysis of the white crystals indicated the following:

Analysis: C_{27}H_{21}Cl_{3}N_{2}O_{2}S
Calculated: C, 59.62; H, 3.89; N, 5.15
Found: C, 59.87; H, 3.89; N, 5.02

Part B: Preparation of 2-(2-chlorobenzoyl)-3-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-1,1-dimethylquandine.

A mixture containing 5.0 grams of 1-(2-chlorobenzoyl)-3-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-S-methylisothiourea prepared in Part A, 25 milliliters of tetrahydrofuran, 10 milliliters of ethanol and 50 milliliters of a 40% aqueous dimethylamine solution was refluxed in a nitrogen atmosphere for 5 hours. An additional 25 milliliters of 40% aqueous dimethylamine solution was added and reflux was continued for an additional 16 hours. After cooling to room temperature, the mixture was poured into 200 milliliters of water. This aqueous mixture was extracted twice with methylene chloride and twice with ethyl acetate. The combined extracts were dried over anhydrous sodium sulfate. Removal of the solvents afforded 3.97 grams of a crude material.
Chromatography of the crude material on silica gel using 9:1 ethyl acetate:methanol for elution afforded 1.87 grams of 2-(2-chlorobenzoyl)-3-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-1,1-dimethylguanidine as an amphomorous solid having a melting point of 85°C-97°C. Elemental analysis of the amphomorous solid indicated the following:

Analysis: \( \text{C}_{28}\text{H}_{24}\text{Cl}_3\text{N}_3\text{O}_2 \)
Calculated: C, 62.18; H, 4.47; N, 7.77
Found: C, 62.10; H, 4.22; N, 8.00

This compound is referred to hereinafter as Compound 49.

**EXAMPLE VI**

**Preparation of 1-(4-tert-butyltrimethylsiloxy-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea**

Into a homogeneous solution of 1-(3,5-dichloro-4-hydroxyphenyl)-3-(2,6-difluorobenzoyl)urea (3.13 grams, 8.66 mmol) in dimethylformamide (25 milliliters) was added triethylamine (1.75 grams, 17.3 mmol), and the resulting mixture was stirred for 5 minutes. Tertiary-butyldimethylsilanol chloride (3.92 grams, 26.0 mmol) was then added, and a thick precipitate formed immediately. This heterogeneous mixture was then diluted with ethyl acetate (200 milliliters), washed with saturated sodium bicarbonate (twice with 150 milliliters) and then washed with a brine solution (150 milliliters). The organic layer was dried over sodium sulfate, filtered, and concentrated under reduced pressure to give a brown solid. The solid was washed with cold ether and vacuum dried to afford 1-(4-tert-butyltrimethyl-
siloxyl-3,5-dichlorophenyl)-3-(2,6-difluorobenzoyl)urea (2.71 grams, 5.70 mmol) having a melting point of 184°C-186°C. Elemental analysis of the product indicated the following:

Analysis: C$_{20}$H$_{22}$Cl$_2$F$_2$N$_2$O$_3$S
Calculated: C, 50.52; H, 4.66; N, 5.89
Found: C, 50.47; H, 4.71; N, 5.69

This compound is referred to hereinafter as Compound 45.

EXAMPLE VII

In a manner similar to that employed in the preceding examples, other urea compounds were prepared. Compounds 5, 6, 7, 14, 18, 21, 22, 23, 29, 32, 33, 34, 39, 43, 48, 68, 78 and 85 were prepared in a manner similar to that employed in Examples I and II for the preparation of Compounds 1 and 2 respectively. Compounds 8, 10, 12, 15, 24, 28, 55, 56, 57, 58, 61, 62, 63, 71, 73, 74, 76, 77, 87, 89, 90, 91, 92, 93 and 98 were prepared in a manner similar to that employed in Example III for the preparation of Compound 3. Compound 79 was prepared in manner similar to that employed in Example V for the preparation of Compound 49. Compounds 25, 37, and 41 were prepared in a manner similar to the procedure described in U.S. Patent Application Serial No. 480,697, filed March 31, 1983. Compound 19 was prepared in a manner similar to the procedure described in U.S. Patent 4,521,426. Compound 26 was prepared in a manner similar to the procedure described in U.S. Patent Application Serial No. 712, 197, filed March 15, 1985. Compounds 30, 53, 64, 94, 95, 96, 97 and 99
were prepared in a manner similar to the procedure described in U.S. Patent Application Serial No. 781,382, filed September 30, 1985. Compounds 69 and 72 were prepared in a manner similar to the procedure described in U.S. Patent Application Serial No. 672,007, filed November 15, 1984. Compound 65 was prepared in a manner similar to the procedure described in U.S. Patent Application Serial No. 686,735, filed December 27, 1984.

The following compounds were prepared in a manner similar to that described in the indicated publication: Compound 16 (U.S. Patent 3,748,356); Compound 36 (U.S. Patent 3,748,356); Compound 47 (U.S. Patent 3,748,356); Compound 17 (U.S. Patent 4,275,077); Compound 20 (U.S. Patent 4,275,077); Compound 27 (U.S. Patent 4,457,943); Compound 31 (U.S. Patent 4,139,636); Compound 9 (U.S. Patent 4,041,177); Compound 88 (U.S. Patent 4,005,223); Compound 35 (EP 72,438); Compound 54 (EP 69,288); Compound 59 (EP 72,438); Compound 46 (U.S. Patent 4,212,870); Compound 75 (U.S. Patent 4,162,330); Compound 80 (U.S. Patent 4,162,330); Compound 11 (U.S. Patent 4,173,637); Compound 13 (U.S. Patent 4,173,637); and Compound 60 (Japan 56 092857).

The following intermediate anilines and aromatic amines were prepared in a manner similar to that described in the indicated publication, and were then reacted with the appropriate benzoyl isocyanate according to the procedure employed in Example II above: 5-amino-3-methyl-1,2,4-oxadiazole described in U.S. Patent 3,917,478 (Compound 82); 2-amino-5-(1-naphthyl) methylthio-1,3,4-thiadiazole...
described in German Patent 1,079,060 (Compound 38); 5-(4-nitrophenyl)-3-trifluoromethyl-1,2,4-oxadiazole described in U.S. Patent 3,917,478 (Compound 50) in which the reduction conditions of Example I, Part B above were used to prepare the corresponding aniline; 1-chloro-4-nitronaphthalene described in Bassilios, Bull. Soc. Chim. Fr., 1951, 651 in which catalytic reduction (Example I, Part B above) afforded 1-amino-4-chloronaphthalene (Compound 40); 1-chloro-4-nitronaphthalene was converted into 1-amino-4-(2,4-dimethylphenoxy) naphthalene according to U.S. Patent Application Serial No. 495,331, filed May 20, 1983 (Compound 42); 9-nitroanthracene was reduced using the procedure described in Example I, Part B above (Compound 44); 1-amino-4-chloro-5,6,7,8-tetrahydronaphthalene described in Press and Hoffman, Org. Prep. Proc. Int., 1982, 14, 204 and Simonetta Beltrane, Gazz. Chim. Ital., 1958, 88, 769 (Compound 70); 3,5-dichloro-4-[4-chloro-1-naphthoxy]butoxy] aniline and 4-[1-(1-naphthyl)ethoxy] aniline described in U.S. Patent Application Serial No. 430,368, filed September 30, 1982 (Compounds 66 and 81); 1,1-bis-(4-chlorophenyl)methylamine and alpha-methyl-4-chlorobenzylamine described in Cram and Guthrie, J. Am. Chem. Soc., 1966, 88, 5760 (Compounds 83 and 84); and alpha-amino-4-chlorophenylacetonitrile described in Stevens et al., J. Chem. Soc., 1931, 2568 (Compound 86). Compounds 51 and 52 were prepared from commercially available anilines or amines according to the procedure employed in Example II above.
Compound 67 was prepared according to the procedure described in Example II above. The aniline used in this procedure was prepared as follows: A mixture containing 32.81 grams of 3-chloro-2,5-dimethyl-4-(2,4-dimethylphenoxy)aniline, 36.40 grams of methyl bromoacetate, 16.44 grams of anhydrous potassium carbonate and 240 milliliters of toluene was heated to reflux for 66 hours. The reaction mixture was then cooled to room temperature, diluted with 700 milliliters of ethyl acetate, and washed with 400 milliliters of water and 400 milliliters of brine solution. The organic layer was dried over anhydrous sodium sulfate and the solvents removed under reduced pressure. The residue was triturated with hexane to afford 28.32 grams of methyl N-[3-chloro-2,5-dimethyl-4-(2,4-dimethylphenoxy)phenyl] glycinate as an off white solid. $^1$H-NMR(CDCl$_3$): 2.12(s, 3H), 2.28(s, 3H), 2.31(s,3H), 2.37(s,3H), 3.81(s,3H), 3.97(s,2H), 6.17(d, $J=8$Hz ,1H), 6.29(s,1H), 6.78(d of d; $J=8$, 2Hz ,1H), 7.01(broad d, $J=2$Hz ,1H).

The structure and analytical data for Compounds 1 through 99, which compounds are used in the examples hereinafter for controlling certain endoparasites and ectoparasites of warm-blooded animals, are set forth in Table 1 below:
### Table 1
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>N</td>
</tr>
<tr>
<td>1</td>
<td><img src="image1" alt="Structure 1" /></td>
<td>60.17</td>
<td>3.13</td>
</tr>
<tr>
<td>2</td>
<td><img src="image2" alt="Structure 2" /></td>
<td>60.59</td>
<td>3.52</td>
</tr>
<tr>
<td>3</td>
<td><img src="image3" alt="Structure 3" /></td>
<td>62.62</td>
<td>4.61</td>
</tr>
<tr>
<td>4</td>
<td><img src="image4" alt="Structure 4" /></td>
<td>51.83</td>
<td>2.17</td>
</tr>
</tbody>
</table>

- Note: The images of the structures are not included in the text. Please refer to the original document for visual representation.
### TABLE 1
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>1</td>
<td><img src="image1.png" alt="Structure 1" /></td>
<td>60.17</td>
<td>3.73</td>
</tr>
<tr>
<td>2</td>
<td><img src="image2.png" alt="Structure 2" /></td>
<td>60.59</td>
<td>3.52</td>
</tr>
<tr>
<td>3</td>
<td><img src="image3.png" alt="Structure 3" /></td>
<td>62.82</td>
<td>4.61</td>
</tr>
<tr>
<td>4</td>
<td><img src="image4.png" alt="Structure 4" /></td>
<td>51.63</td>
<td>2.17</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td><img src="image" alt="Structure 5" /></td>
<td>Calculated: 55.25, 2.31, 5.37</td>
<td>Found: 55.56, 2.46, 5.36</td>
</tr>
<tr>
<td>6</td>
<td><img src="image" alt="Structure 6" /></td>
<td>Calculated: 64.93, 3.98, 5.83</td>
<td>Found: 63.93, 3.92, 6.14</td>
</tr>
<tr>
<td>7</td>
<td><img src="image" alt="Structure 7" /></td>
<td>Calculated: 62.03, 4.41, 5.57</td>
<td>Found: 60.03, 4.15, 5.49</td>
</tr>
<tr>
<td>8</td>
<td><img src="image" alt="Structure 8" /></td>
<td>Calculated: 58.43, 4.09, 5.68</td>
<td>Found: 58.72, 4.18, 5.68</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
</table>
| 5        | ![Structure 5](image1) | Calculated: C 55.25, H 2.51, N 5.31  
                         Found: C 55.56, H 2.46, N 5.36 | 237-239 |
| 6        | ![Structure 6](image2) | Calculated: C 64.93, H 2.90, N 5.03  
                         Found: C 65.93, H 3.92, N 6.14 | 206 |
| 7        | ![Structure 7](image3) | Calculated: C 62.03, H 4.01, N 5.57  
                         Found: C 60.03, H 4.15, N 5.49 | 224-226 |
| 8        | ![Structure 8](image4) | Calculated: C 58.43, H 4.09, N 5.66  
                         Found: C 58.72, H 4.18, N 5.68 | 221-222.5 |
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C     H     N</td>
</tr>
<tr>
<td>9</td>
<td><img src="image" alt="Structure 9" /></td>
<td>49.81 2.30 6.72</td>
</tr>
<tr>
<td>10</td>
<td><img src="image" alt="Structure 10" /></td>
<td>50.93 2.35 5.94</td>
</tr>
<tr>
<td>11</td>
<td><img src="image" alt="Structure 11" /></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td><img src="image" alt="Structure 12" /></td>
<td>53.60 3.75 5.21</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)

Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>9</td>
<td><img src="image" alt="Structure 9" /></td>
<td>49.01</td>
<td>2.30</td>
</tr>
<tr>
<td>10</td>
<td><img src="image" alt="Structure 10" /></td>
<td>50.93</td>
<td>2.35</td>
</tr>
<tr>
<td>11</td>
<td><img src="image" alt="Structure 11" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td><img src="image" alt="Structure 12" /></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Possible error in the structure of Compound 12.
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>13</td>
<td><img src="image1" alt="Structure" /></td>
<td>45.00</td>
<td>1.79</td>
</tr>
<tr>
<td>14</td>
<td><img src="image2" alt="Structure" /></td>
<td>64.31</td>
<td>3.67</td>
</tr>
<tr>
<td>15</td>
<td><img src="image3" alt="Structure" /></td>
<td>52.07</td>
<td>3.63</td>
</tr>
<tr>
<td>16</td>
<td><img src="image4" alt="Structure" /></td>
<td>H NMR (DMSO - D6): 11.54(s, 1H), 10.33 (s, 1H), 7.00 - 7.90 (m, 7H).</td>
<td>229</td>
</tr>
</tbody>
</table>

PCT/US88/002545
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>13</td>
<td><img src="image1.png" alt="Structure 13" /></td>
<td>C: 45.00, H: 1.79, N: 8.28</td>
<td>229-230</td>
</tr>
<tr>
<td>14</td>
<td><img src="image2.png" alt="Structure 14" /></td>
<td>C: 64.31, H: 3.67, N: 6.00</td>
<td>233</td>
</tr>
<tr>
<td>15</td>
<td><img src="image3.png" alt="Structure 15" /></td>
<td>C: 52.07, H: 3.03, N: 9.61</td>
<td>180-190</td>
</tr>
<tr>
<td>16</td>
<td><img src="image4.png" alt="Structure 16" /></td>
<td>C: 78.17, H: 3.49, N: 5.92</td>
<td>229</td>
</tr>
</tbody>
</table>

**Elemental Analysis**

- **Calculated**
  - C
  - H
  - N
- **Found**
  - C
  - H
  - N

**Notes:**
- $^1$H NMR (DMSO-d$_6$): 11.34 (s, 1H), 10.33 (s, 1H), 7.00 - 7.90 (m, 7H).
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
</table>
| 11 | ![Structure 11](image1.png) | Calculated: C 59.15, H 2.90, N 5.75  
Found: C 59.22, H 2.90, N 5.00 | 234-236 |
| 12 | ![Structure 12](image2.png) | Calculated: C 57.65, H 2.71, N 5.38  
Found: C 57.46, H 2.69, N 5.30 | 215-217 |
| 13 | ![Structure 13](image3.png) | Calculated: C 56.58, H 7.68, N 8.61  
Found: C 56.09, H 7.44, N 8.43 | 211.5-212.5 |
| 14 | ![Structure 14](image4.png) | Calculated: C 63.65, H 3.34, N 6.19  
Found: C 63.40, H 3.20, N 6.15 | 214 |
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>17</td>
<td><img src="image1" alt="Structure 1" /></td>
<td>59.15</td>
<td>2.90</td>
</tr>
<tr>
<td>18</td>
<td><img src="image2" alt="Structure 2" /></td>
<td>57.65</td>
<td>2.71</td>
</tr>
<tr>
<td>19</td>
<td><img src="image3" alt="Structure 3" /></td>
<td>56.58</td>
<td>2.68</td>
</tr>
<tr>
<td>20</td>
<td><img src="image4" alt="Structure 4" /></td>
<td>63.65</td>
<td>3.34</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acp1 Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>21</td>
<td><img src="image" alt="Structure 21" /></td>
<td>63.69</td>
<td>5.34</td>
</tr>
<tr>
<td>22</td>
<td><img src="image" alt="Structure 22" /></td>
<td>59.89</td>
<td>3.22</td>
</tr>
<tr>
<td>23</td>
<td><img src="image" alt="Structure 23" /></td>
<td>59.42</td>
<td>3.99</td>
</tr>
<tr>
<td>24</td>
<td><img src="image" alt="Structure 24" /></td>
<td>64.74</td>
<td>5.43</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
</table>
| 21       | ![Structure 21](image1) | Calculated: C 63.69, H 5.34, N 7.96  
            Found: C 63.71, H 5.31, N 7.83  
            | 219       |                    |
| 22       | ![Structure 22](image2) | Calculated: C 59.09, H 3.22, N 5.59  
            Found: C 60.89, H 3.38, N 5.77  
            | 238       |                    |
| 23       | ![Structure 23](image3) | Calculated: C 59.42, H 3.99, N 5.54  
            Found: C 59.56, H 4.08, N 5.48  
            | 210-212   |                    |
| 24       | ![Structure 24](image4) | Calculated: C 54.74, H 5.43, N 5.59  
            Found: C 64.70, H 5.73, N 5.65  
<pre><code>        | 192-194   |                    |
</code></pre>
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C     H     N</td>
<td>C     H     N</td>
</tr>
<tr>
<td>25</td>
<td><img src="image" alt="Structure 25" /></td>
<td>58.76 3.42 5.27</td>
<td>60.03 2.54 5.13</td>
</tr>
<tr>
<td>26</td>
<td><img src="image" alt="Structure 26" /></td>
<td>55.25 2.53 0.79</td>
<td>55.19 2.51 0.58</td>
</tr>
<tr>
<td>27</td>
<td><img src="image" alt="Structure 27" /></td>
<td>44.12 1.39 7.35</td>
<td>44.08 1.46 7.38</td>
</tr>
<tr>
<td>28</td>
<td><img src="image" alt="Structure 28" /></td>
<td>50.22 3.06 5.32</td>
<td>50.16 2.14 5.30</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>25</td>
<td><img src="image1.png" alt="Structure 1" /></td>
<td>58.76</td>
<td>3.42</td>
</tr>
<tr>
<td>26</td>
<td><img src="image2.png" alt="Structure 2" /></td>
<td>55.25</td>
<td>2.53</td>
</tr>
<tr>
<td>27</td>
<td><img src="image3.png" alt="Structure 3" /></td>
<td>44.12</td>
<td>1.59</td>
</tr>
<tr>
<td>28</td>
<td><img src="image4.png" alt="Structure 4" /></td>
<td>50.22</td>
<td>3.06</td>
</tr>
</tbody>
</table>
TABLE 1 (cont.)
Representative Acril Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>C</strong>   <strong>H</strong>  <strong>N</strong></td>
<td><strong>C</strong>   <strong>H</strong>  <strong>N</strong></td>
</tr>
<tr>
<td>29</td>
<td><img src="image" alt="Structure 29" /></td>
<td>56.65  2.93  5.08</td>
<td>56.40  3.03  5.18</td>
</tr>
<tr>
<td>30</td>
<td><img src="image" alt="Structure 30" /></td>
<td>57.27  2.96  5.14</td>
<td>56.07  3.05  4.91</td>
</tr>
<tr>
<td>31</td>
<td><img src="image" alt="Structure 31" /></td>
<td>56.01  3.57  5.52</td>
<td>59.21  3.03  5.09</td>
</tr>
<tr>
<td>32</td>
<td><img src="image" alt="Structure 32" /></td>
<td>56.65  3.57  5.52</td>
<td>59.21  3.03  5.09</td>
</tr>
</tbody>
</table>

$^1$H NMR (DMSO - d$_6$): \(d = 10.40\) (br s, 1H), 10.05 (br s, 1H), 7.14 - 8.03 (m, 8H).
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>29</td>
<td><img src="image" alt="Structure 29" /></td>
<td><strong>Elemental Analysis</strong>&lt;br&gt;Calculated: C 56.65, H 2.93, N 5.08&lt;br&gt;Found: C 56.40, H 3.03, N 5.18</td>
<td>202-210</td>
</tr>
<tr>
<td>30</td>
<td><img src="image" alt="Structure 30" /></td>
<td><strong>Elemental Analysis</strong>&lt;br&gt;Calculated: C 57.27, H 2.96, N 5.14&lt;br&gt;Found: C 57.82, H 3.05, N 4.91</td>
<td>235-238</td>
</tr>
<tr>
<td>31</td>
<td><img src="image" alt="Structure 31" /></td>
<td><strong>Elemental Analysis</strong>&lt;br&gt;Calculated: C 56.81, H 3.57, N 5.52&lt;br&gt;Found: C 59.21, H 3.05, N 5.09</td>
<td>200-214</td>
</tr>
<tr>
<td>32</td>
<td><img src="image" alt="Structure 32" /></td>
<td><strong>Elemental Analysis</strong>&lt;br&gt;Calculated: C 56.81, H 3.57, N 5.52&lt;br&gt;Found: C 59.21, H 3.05, N 5.09</td>
<td>200-214</td>
</tr>
</tbody>
</table>

**1H NMR (DMSO - d6):**<br>7.14 (br s, 1H), 10.05 (br s, 1H), 11.05 (br s, 1H), 10.05 (br s, 1H), 11.05 (br s, 1H).
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>33</td>
<td><img src="image" alt="Structure 33" /></td>
<td>C: 54.60 H: 2.74 N: 4.24</td>
<td>112-119</td>
</tr>
<tr>
<td>34</td>
<td><img src="image" alt="Structure 34" /></td>
<td>C: 55.41 H: 2.71 N: 5.38</td>
<td>262</td>
</tr>
<tr>
<td>35</td>
<td><img src="image" alt="Structure 35" /></td>
<td>C: 53.45 H: 4.50 N: 12.11</td>
<td>103</td>
</tr>
<tr>
<td>36</td>
<td><img src="image" alt="Structure 36" /></td>
<td>C: 55.48 H: 3.41 N: 8.62</td>
<td>111-115</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>33</td>
<td><img src="image" alt="Structure 33" /></td>
<td>54.60</td>
<td>2.74</td>
</tr>
<tr>
<td>35</td>
<td><img src="image" alt="Structure 35" /></td>
<td>55.41</td>
<td>2.71</td>
</tr>
<tr>
<td>35</td>
<td><img src="image" alt="Structure 35'" /></td>
<td>55.48</td>
<td>3.41</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point°C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>31</td>
<td><img src="image31" alt="Structure 31" /></td>
<td>C: 56.75 H: 4.16 N: 5.51 C: 56.64 H: 4.09 N: 5.47</td>
<td>183.5-184</td>
</tr>
<tr>
<td>38</td>
<td><img src="image38" alt="Structure 38" /></td>
<td>C: 55.75 H: 3.09 N: 12.27 C: 55.40 H: 3.16 N: 12.22</td>
<td>204-205</td>
</tr>
<tr>
<td>40</td>
<td><img src="image40" alt="Structure 40" /></td>
<td>C: 59.92 H: 3.07 N: 7.16 C: 59.91 H: 3.22 N: 7.62</td>
<td>201-210</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>------------------</td>
</tr>
<tr>
<td>37</td>
<td><img src="image" alt="Structure 37" /></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td></td>
<td></td>
<td>56.75</td>
<td>4.16</td>
</tr>
<tr>
<td>38</td>
<td><img src="image" alt="Structure 38" /></td>
<td>55.25</td>
<td>3.09</td>
</tr>
<tr>
<td>39</td>
<td><img src="image" alt="Structure 39" /></td>
<td>63.09</td>
<td>4.19</td>
</tr>
<tr>
<td>40</td>
<td><img src="image" alt="Structure 40" /></td>
<td>59.92</td>
<td>3.07</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point°C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>41</td>
<td><img src="image1" alt="Structure 41" /></td>
<td>Calculated: C 60.13, H 4.38, N 6.09</td>
<td>Found: C 61.01, H 4.55, N 6.93</td>
</tr>
<tr>
<td>42</td>
<td><img src="image2" alt="Structure 42" /></td>
<td>Calculated: C 69.94, H 4.51, N 6.27</td>
<td>Found: C 68.61, H 4.60, N 6.21</td>
</tr>
<tr>
<td>43</td>
<td><img src="image3" alt="Structure 43" /></td>
<td>Calculated: C 61.43, H 4.01, N 5.30</td>
<td>Found: C 61.20, H 4.00, N 5.42</td>
</tr>
<tr>
<td>44</td>
<td><img src="image4" alt="Structure 44" /></td>
<td>Calculated: C 70.20, H 3.74, N 7.44</td>
<td>Found: C 61.62, H 3.70, N 7.55</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>41</td>
<td><img src="image1" alt="Structure 41" /></td>
<td>60.12</td>
<td>4.38</td>
</tr>
<tr>
<td>42</td>
<td><img src="image2" alt="Structure 42" /></td>
<td>69.94</td>
<td>4.51</td>
</tr>
<tr>
<td>43</td>
<td><img src="image3" alt="Structure 43" /></td>
<td>61.43</td>
<td>4.01</td>
</tr>
<tr>
<td>44</td>
<td><img src="image4" alt="Structure 44" /></td>
<td>70.20</td>
<td>3.74</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>------------------</td>
</tr>
<tr>
<td>45</td>
<td><img src="image" alt="Structure 45" /></td>
<td>Calculated: C 50.52, H 4.66, N 5.89; Found: C 50.47, H 4.71, N 5.69</td>
<td>184-186</td>
</tr>
<tr>
<td>46</td>
<td><img src="image" alt="Structure 46" /></td>
<td>Calculated: C 43.30, H 2.23, N 11.65; Found: C 43.29, H 2.20, N 11.46</td>
<td>210-212</td>
</tr>
<tr>
<td>47</td>
<td><img src="image" alt="Structure 47" /></td>
<td>Calculated: C 52.33, H 2.63, N 8.13; Found: C 52.35, H 2.56, N 8.90</td>
<td>238-240</td>
</tr>
<tr>
<td>48</td>
<td><img src="image" alt="Structure 48" /></td>
<td>Calculated: C 55.93, H 3.43, N 5.01; Found: C 55.10, H 3.63, N 5.02</td>
<td>163-166</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point°C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>45</td>
<td><img src="image1" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td><img src="image2" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>46</td>
<td><img src="image3" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>47</td>
<td><img src="image4" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>48</td>
<td><img src="image5" alt="Structure" /></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>------------------</td>
</tr>
<tr>
<td>49</td>
<td><img src="image" alt="Compound 49 Structure" /></td>
<td>C: 62.19, H: 4.41, N: 7.76</td>
<td>65-77</td>
</tr>
<tr>
<td>50</td>
<td><img src="image" alt="Compound 50 Structure" /></td>
<td>C: 49.52, H: 2.20, N: 13.59</td>
<td>237-240</td>
</tr>
<tr>
<td>51</td>
<td><img src="image" alt="Compound 51 Structure" /></td>
<td>C: 55.48, H: 3.41, N: 8.42</td>
<td>165-168</td>
</tr>
<tr>
<td>52</td>
<td><img src="image" alt="Compound 52 Structure" /></td>
<td>C: 55.20, H: 3.03, N: 7.98</td>
<td>194-197</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C     H     N</td>
<td>C     H     N</td>
</tr>
<tr>
<td>49</td>
<td>![Structure 49]</td>
<td>62.17  4.47  7.76</td>
<td>62.10  4.22  8.00</td>
</tr>
<tr>
<td>50</td>
<td>![Structure 50]</td>
<td>49.52  2.20  13.59</td>
<td>49.49  2.26  13.61</td>
</tr>
<tr>
<td>51</td>
<td>![Structure 51]</td>
<td>55.48  3.41  8.62</td>
<td>55.72  3.37  8.64</td>
</tr>
<tr>
<td>52</td>
<td>![Structure 52]</td>
<td>64.21  3.83  7.60</td>
<td>65.20  4.43  7.54</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>53</td>
<td><img src="image" alt="Structure 1" /></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td></td>
<td></td>
<td>47.70</td>
<td>3.16</td>
</tr>
<tr>
<td>54</td>
<td><img src="image" alt="Structure 2" /></td>
<td>48.27</td>
<td>2.13</td>
</tr>
<tr>
<td>55</td>
<td><img src="image" alt="Structure 3" /></td>
<td>50.21</td>
<td>3.06</td>
</tr>
<tr>
<td>56</td>
<td><img src="image" alt="Structure 4" /></td>
<td>52.89</td>
<td>3.66</td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>53</td>
<td><img src="image1" alt="structure" /></td>
<td>41.10</td>
<td>3.16</td>
</tr>
<tr>
<td>54</td>
<td><img src="image2" alt="structure" /></td>
<td>48.27</td>
<td>2.13</td>
</tr>
<tr>
<td>55</td>
<td><img src="image3" alt="structure" /></td>
<td>50.21</td>
<td>3.04</td>
</tr>
<tr>
<td>56</td>
<td><img src="image4" alt="structure" /></td>
<td>52.89</td>
<td>3.66</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point°C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>-------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>57</td>
<td><img src="image" alt="Structure 57" /></td>
<td>Calculated: C 50.72, H 3.72, N 4.92</td>
<td>Found: C 50.41, H 3.74, N 4.64</td>
</tr>
<tr>
<td>58</td>
<td><img src="image" alt="Structure 58" /></td>
<td>Calculated: C 53.75, H 3.94, N 5.22</td>
<td>Found: C 55.16, H 4.12, N 5.02</td>
</tr>
<tr>
<td>59</td>
<td><img src="image" alt="Structure 59" /></td>
<td>Calculated: C 41.55, H 2.15, N 3.31</td>
<td>Found: C 41.00, H 2.24, N 3.11</td>
</tr>
<tr>
<td>60</td>
<td><img src="image" alt="Structure 60" /></td>
<td>Calculated: C 55.88, H 4.10, N 5.43</td>
<td>Found: C 56.65, H 3.66, N 4.41</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td>57</td>
<td><img src="image1" alt="Structure 57" /></td>
<td>Calculated &lt;br&gt; C: 50.72&lt;br&gt;H: 3.12&lt;br&gt;N: 4.92</td>
<td>Found &lt;br&gt; C: 50.41&lt;br&gt;H: 3.74&lt;br&gt;N: 4.64</td>
</tr>
<tr>
<td>58</td>
<td><img src="image2" alt="Structure 58" /></td>
<td>Calculated &lt;br&gt; C: 53.75&lt;br&gt;H: 3.94&lt;br&gt;N: 5.22</td>
<td>Found &lt;br&gt; C: 55.16&lt;br&gt;H: 4.12&lt;br&gt;N: 5.02</td>
</tr>
<tr>
<td>59</td>
<td><img src="image3" alt="Structure 59" /></td>
<td>Calculated &lt;br&gt; C: 47.55&lt;br&gt;H: 2.19&lt;br&gt;N: 8.31</td>
<td>Found &lt;br&gt; C: 47.80&lt;br&gt;H: 2.24&lt;br&gt;N: 8.11</td>
</tr>
<tr>
<td>60</td>
<td><img src="image4" alt="Structure 60" /></td>
<td>Calculated &lt;br&gt; C: 55.88&lt;br&gt;H: 4.10&lt;br&gt;N: 5.43</td>
<td>Found &lt;br&gt; C: 56.65&lt;br&gt;H: 3.66&lt;br&gt;N: 5.41</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>61</td>
<td><img src="image1.png" alt="Structure 61" /></td>
<td>54.09</td>
<td>3.35</td>
</tr>
<tr>
<td>62</td>
<td><img src="image2.png" alt="Structure 62" /></td>
<td>54.02</td>
<td>2.12</td>
</tr>
<tr>
<td>63</td>
<td><img src="image3.png" alt="Structure 63" /></td>
<td>40.55</td>
<td>2.17</td>
</tr>
<tr>
<td>64</td>
<td><img src="image4.png" alt="Structure 64" /></td>
<td>585.9510</td>
<td>HRMS</td>
</tr>
</tbody>
</table>
**TABLE 1 (Cont.)**
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>61</td>
<td><img src="image" alt="Structure 61" /></td>
<td>C: 54.89, H: 3.35, N: 5.82</td>
<td>140-154</td>
</tr>
<tr>
<td>65</td>
<td><img src="image" alt="Structure 65" /></td>
<td>C: 51.03, H: 2.12, N: 5.31</td>
<td>214-216</td>
</tr>
<tr>
<td>63</td>
<td><img src="image" alt="Structure 63" /></td>
<td>C: 51.92, H: 2.83, N: 5.15</td>
<td>163-166</td>
</tr>
<tr>
<td>64</td>
<td><img src="image" alt="Structure 64" /></td>
<td>C: 48.58, H: 2.77, N: 5.14</td>
<td>195.5-199.5</td>
</tr>
</tbody>
</table>

*Calculated vs. Found.*

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>61</td>
<td><img src="image" alt="Structure 61" /></td>
<td>C: 54.89, H: 3.35, N: 5.82</td>
<td>140-154</td>
</tr>
<tr>
<td>65</td>
<td><img src="image" alt="Structure 65" /></td>
<td>C: 51.03, H: 2.12, N: 5.31</td>
<td>214-216</td>
</tr>
<tr>
<td>63</td>
<td><img src="image" alt="Structure 63" /></td>
<td>C: 51.92, H: 2.83, N: 5.15</td>
<td>163-166</td>
</tr>
<tr>
<td>64</td>
<td><img src="image" alt="Structure 64" /></td>
<td>C: 48.58, H: 2.77, N: 5.14</td>
<td>195.5-199.5</td>
</tr>
</tbody>
</table>

*High Resolution Mass Spectroscopy.*
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>55</td>
<td><img src="image1" alt="Structure" /></td>
<td><img src="image2" alt="Calculated" /></td>
<td>201-209</td>
</tr>
<tr>
<td>56</td>
<td><img src="image3" alt="Structure" /></td>
<td><img src="image4" alt="Calculated" /></td>
<td>204.2-206.9</td>
</tr>
<tr>
<td>57</td>
<td><img src="image5" alt="Structure" /></td>
<td><img src="image6" alt="Calculated" /></td>
<td>186-190</td>
</tr>
<tr>
<td>58</td>
<td><img src="image7" alt="Structure" /></td>
<td><img src="image8" alt="Calculated" /></td>
<td>220-225</td>
</tr>
</tbody>
</table>
### Provisional Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>H</td>
<td>N</td>
<td>C</td>
</tr>
<tr>
<td>65</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>66</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>67</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>68</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td>61</td>
<td><img src="image1.png" alt="Structure 1" /></td>
<td>62.00</td>
<td>3.97</td>
</tr>
<tr>
<td>70</td>
<td><img src="image2.png" alt="Structure 2" /></td>
<td>59.26</td>
<td>4.14</td>
</tr>
<tr>
<td>11</td>
<td><img src="image3.png" alt="Structure 3" /></td>
<td>63.99</td>
<td>5.12</td>
</tr>
<tr>
<td>12</td>
<td><img src="image4.png" alt="Structure 4" /></td>
<td>61.49</td>
<td>3.44</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td>9</td>
<td><img src="image" alt="Structure 9" /></td>
<td><strong>C</strong> 63.80 <strong>H</strong> 3.97 <strong>N</strong> 5.52</td>
<td><strong>C</strong> 64.03 <strong>H</strong> 4.74 <strong>N</strong> 5.86</td>
</tr>
<tr>
<td>10</td>
<td><img src="image" alt="Structure 10" /></td>
<td><strong>C</strong> 59.26 <strong>H</strong> 4.14 <strong>N</strong> 7.67</td>
<td><strong>C</strong> 59.27 <strong>H</strong> 4.39 <strong>N</strong> 7.41</td>
</tr>
<tr>
<td>11</td>
<td><img src="image" alt="Structure 11" /></td>
<td><strong>C</strong> 63.99 <strong>H</strong> 5.12 <strong>N</strong> 6.78</td>
<td><strong>C</strong> 63.93 <strong>H</strong> 5.35 <strong>N</strong> 6.50</td>
</tr>
<tr>
<td>12</td>
<td><img src="image" alt="Structure 12" /></td>
<td><strong>C</strong> 61.49 <strong>H</strong> 3.44 <strong>N</strong> 5.31</td>
<td><strong>C</strong> 61.57 <strong>H</strong> 3.51 <strong>N</strong> 5.27</td>
</tr>
</tbody>
</table>
MICROCOPY RESOLUTION TEST CHART
NATIONAL BUREAU OF STANDARDS
STANDARD REFERENCE MATERIAL 1010b
(ANSI and ISO TEST CHART No. 2)
<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
</tr>
</thead>
<tbody>
<tr>
<td>73</td>
<td><img src="image1" alt="Structure" /></td>
<td>Calculated</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>60.26</td>
</tr>
<tr>
<td>74</td>
<td><img src="image2" alt="Structure" /></td>
<td>52.03</td>
</tr>
<tr>
<td>75</td>
<td><img src="image3" alt="Structure" /></td>
<td>52.31</td>
</tr>
<tr>
<td>76</td>
<td><img src="image4" alt="Structure" /></td>
<td>64.05</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
</tr>
<tr>
<td>73</td>
<td><img src="image1" alt="Structure 73" /></td>
<td>Calculated: C 60.26, H 0.61, N 9.56</td>
</tr>
<tr>
<td>74</td>
<td><img src="image2" alt="Structure 74" /></td>
<td>Calculated: C 60.22, H 4.85, N 9.17</td>
</tr>
<tr>
<td>75</td>
<td><img src="image3" alt="Structure 75" /></td>
<td>Calculated: C 52.24, H 3.54, N 6.49</td>
</tr>
<tr>
<td>76</td>
<td><img src="image4" alt="Structure 76" /></td>
<td>Calculated: C 64.00, H 5.68, N 9.86</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point ($^\circ$C)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>C</strong></td>
<td><strong>H</strong></td>
</tr>
<tr>
<td>71</td>
<td><img src="image1" alt="Structure 71" /></td>
<td>57.83</td>
<td>4.85</td>
</tr>
<tr>
<td>78</td>
<td><img src="image2" alt="Structure 78" /></td>
<td>46.30</td>
<td>2.11</td>
</tr>
<tr>
<td>79</td>
<td><img src="image3" alt="Structure 79" /></td>
<td>57.15</td>
<td>4.49</td>
</tr>
<tr>
<td>80</td>
<td><img src="image4" alt="Structure 80" /></td>
<td>51.14</td>
<td>2.52</td>
</tr>
</tbody>
</table>
**TABLE 1 (Cont.)**

Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C     H     N</td>
<td>C     H     N</td>
</tr>
<tr>
<td>11</td>
<td><img src="image1" alt="Structure 1" /></td>
<td>57.83 4.85 4.81</td>
<td>58.02 4.90 4.77</td>
</tr>
<tr>
<td>78</td>
<td><img src="image2" alt="Structure 2" /></td>
<td>46.30 2.11 4.90</td>
<td>47.34 2.18 5.01</td>
</tr>
<tr>
<td>79</td>
<td><img src="image3" alt="Structure 3" /></td>
<td>57.15 4.49 12.45</td>
<td>57.06 4.49 12.36</td>
</tr>
<tr>
<td>80</td>
<td><img src="image4" alt="Structure 4" /></td>
<td>51.14 2.52 7.01</td>
<td>50.31 2.55 6.51</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>------------------</td>
</tr>
<tr>
<td>01</td>
<td><img src="image1" alt="Structure" /></td>
<td>C: 69.94, H: 4.51, N: 6.77</td>
<td>212.5-216</td>
</tr>
<tr>
<td>02</td>
<td><img src="image2" alt="Structure" /></td>
<td>C: 46.91, H: 2.85, N: 19.85</td>
<td>151-154</td>
</tr>
<tr>
<td>03</td>
<td><img src="image3" alt="Structure" /></td>
<td>C: 51.49, H: 3.74, N: 7.50</td>
<td>181-183</td>
</tr>
<tr>
<td>04</td>
<td><img src="image4" alt="Structure" /></td>
<td>C: 57.94, H: 3.74, N: 6.43</td>
<td>196.5-199</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)
**Representative Acyl Urea Compounds**

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td>C</td>
<td>H</td>
</tr>
<tr>
<td><strong>81</strong></td>
<td><img src="image1" alt="" /></td>
<td>69.94</td>
<td>4.51</td>
</tr>
<tr>
<td><strong>82</strong></td>
<td><img src="image2" alt="" /></td>
<td>46.81</td>
<td>2.85</td>
</tr>
<tr>
<td><strong>83</strong></td>
<td><img src="image3" alt="" /></td>
<td>51.49</td>
<td>3.74</td>
</tr>
<tr>
<td><strong>84</strong></td>
<td><img src="image4" alt="" /></td>
<td>57.94</td>
<td>3.74</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>------------------</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>C</strong></td>
<td><strong>H</strong></td>
</tr>
<tr>
<td>85</td>
<td><img src="image1" alt="Structure" /></td>
<td>55.78</td>
<td>3.24</td>
</tr>
<tr>
<td>86</td>
<td><img src="image2" alt="Structure" /></td>
<td>54.94</td>
<td>2.88</td>
</tr>
<tr>
<td>87</td>
<td><img src="image3" alt="Structure" /></td>
<td>47.16</td>
<td>2.69</td>
</tr>
<tr>
<td>88</td>
<td><img src="image4" alt="Structure" /></td>
<td>46.49</td>
<td>1.95</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>C</strong></td>
<td><strong>H</strong></td>
</tr>
<tr>
<td>05</td>
<td><img src="image1.png" alt="Structure" /></td>
<td>55.70</td>
<td>3.24</td>
</tr>
<tr>
<td>06</td>
<td><img src="image2.png" alt="Structure" /></td>
<td>54.94</td>
<td>2.88</td>
</tr>
<tr>
<td>07</td>
<td><img src="image3.png" alt="Structure" /></td>
<td>47.16</td>
<td>2.69</td>
</tr>
<tr>
<td>08</td>
<td><img src="image4.png" alt="Structure" /></td>
<td>46.89</td>
<td>1.95</td>
</tr>
</tbody>
</table>
**TABLE 1 (Cont.)**
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Calculated</td>
<td>Found</td>
</tr>
<tr>
<td></td>
<td></td>
<td>C     H     N</td>
<td>C     H     N</td>
</tr>
<tr>
<td>89</td>
<td><img src="image1.png" alt="Structure" /></td>
<td>62.08 5.01 5.57</td>
<td>62.00 5.14 5.50</td>
</tr>
<tr>
<td>90</td>
<td><img src="image2.png" alt="Structure" /></td>
<td>58.42 4.08 5.67</td>
<td>58.16 4.19 5.70</td>
</tr>
<tr>
<td>91</td>
<td><img src="image3.png" alt="Structure" /></td>
<td>59.94 4.15 5.67</td>
<td>60.71 4.12 5.97</td>
</tr>
<tr>
<td>92</td>
<td><img src="image4.png" alt="Structure" /></td>
<td>60.82 4.67 8.85</td>
<td>62.66 4.71 8.12</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point °C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>89</td>
<td><img src="image" alt="Structure" /></td>
<td>C: 62.06, H: 5.01, N: 5.37</td>
<td>105-106</td>
</tr>
<tr>
<td>90</td>
<td><img src="image" alt="Structure" /></td>
<td>C: 58.42, H: 4.08, N: 5.67</td>
<td>157-159</td>
</tr>
<tr>
<td>91</td>
<td><img src="image" alt="Structure" /></td>
<td>C: 60.11, H: 4.12, N: 5.97</td>
<td>182-186</td>
</tr>
<tr>
<td>92</td>
<td><img src="image" alt="Structure" /></td>
<td>C: 62.66, H: 4.71, N: 8.12</td>
<td>125-130</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>93</td>
<td><img src="image1" alt="Structure" /></td>
<td>C: 51.83, H: 3.16, N: 5.49</td>
<td>161.5-167</td>
</tr>
<tr>
<td>94</td>
<td><img src="image2" alt="Structure" /></td>
<td>C: 47.86, H: 2.84, N: 4.65</td>
<td>170-173</td>
</tr>
<tr>
<td>95</td>
<td><img src="image3" alt="Structure" /></td>
<td>C: 47.33, H: 2.34, N: 5.01</td>
<td>213-214.5</td>
</tr>
<tr>
<td>96</td>
<td><img src="image4" alt="Structure" /></td>
<td>C: 47.16, H: 2.89, N: 5.00</td>
<td>177-180</td>
</tr>
</tbody>
</table>

Representative Acyl Urea Compounds

- **Structure**: Chemical structures of the compounds are shown.
- **Elemental Analysis**:
  - **Calculated**: C, H, N values for the compound.
  - **Found**: Observed C, H, N values.
- **Melting Point**: Melting temperature range for each compound.
TABLE 1 (Cont.)
Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>93</td>
<td><img src="image" alt="Structure 93" /></td>
<td>C: 51.83  H: 3.16  N: 5.49</td>
<td>C: 58.62  H: 3.60  N: 5.96</td>
</tr>
<tr>
<td>94</td>
<td><img src="image" alt="Structure 94" /></td>
<td>C: 47.86  H: 2.84  N: 4.65</td>
<td>C: 49.73  H: 3.43  N: 4.40</td>
</tr>
<tr>
<td>95</td>
<td><img src="image" alt="Structure 95" /></td>
<td>C: 47.33  H: 2.34  N: 5.01</td>
<td>C: 47.06  H: 2.32  N: 4.71</td>
</tr>
<tr>
<td>96</td>
<td><img src="image" alt="Structure 96" /></td>
<td>C: 47.16  H: 2.69  N: 5.00</td>
<td>C: 47.30  H: 2.62  N: 5.10</td>
</tr>
</tbody>
</table>
### TABLE 1 (Cont.)

Representative Acyl Urea Compounds

<table>
<thead>
<tr>
<th>Compound</th>
<th>Structure</th>
<th>Elemental Analysis</th>
<th>Melting Point °C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><img src="image1.png" alt="Structure 1" /></td>
<td><strong>Calculated</strong></td>
<td><strong>Found</strong></td>
</tr>
<tr>
<td>97</td>
<td><img src="image1.png" alt="Structure 1" /></td>
<td>C: 52.14  H: 3.23  N: 5.28</td>
<td>C: 51.04  H: 3.22  N: 5.25</td>
</tr>
<tr>
<td>99</td>
<td><img src="image3.png" alt="Structure 3" /></td>
<td>C: 49.54  H: 3.35  N: 4.44</td>
<td>C: 49.57  H: 3.35  N: 4.46</td>
</tr>
<tr>
<td>Compound</td>
<td>Structure</td>
<td>Elemental Analysis</td>
<td>Melting Point°C</td>
</tr>
<tr>
<td>----------</td>
<td>-----------</td>
<td>--------------------</td>
<td>-----------------</td>
</tr>
<tr>
<td>97</td>
<td><img src="image" alt="Structure 97" /></td>
<td><strong>Calculated</strong> C: 52.14  H: 3.23  N: 5.28</td>
<td><strong>Found</strong> C: 51.04  H: 3.22  N: 5.25</td>
</tr>
<tr>
<td>98</td>
<td><img src="image" alt="Structure 98" /></td>
<td><strong>Calculated</strong> C: 62.81  H: 6.61  N: 6.10</td>
<td><strong>Found</strong> C: 62.53  H: 4.44  N: 6.04</td>
</tr>
<tr>
<td>99</td>
<td><img src="image" alt="Structure 99" /></td>
<td><strong>Calculated</strong> C: 49.54  H: 3.35  N: 4.44</td>
<td><strong>Found</strong> C: 49.57  H: 3.35  N: 4.40</td>
</tr>
</tbody>
</table>
EXAMPLE VIII
Systemic Control of Northern Fowl Mite
(Ornithonyssus sylviarum)

White Leghorn laying hens housed in commercial egg laying cages were preinfested with northern fowl mites 3 weeks prior to being fed the treated feed. The feed was prepared by dissolving Compound 1 (technical grade) in soybean oil and blending the solution with a standard poultry ration. An untreated blank was prepared by blending soybean oil alone with the ration. Twenty birds received feed treated with 500 ppm of Compound 1 and 20 birds received the soybean oil (control). Birds were maintained on the rations for 5 consecutive weeks. Mite counts on each bird were made at weekly intervals, and the number of mites expressed according to the standard index: 0=no mites, 1=1 to 10 mites, 2=11 to 50 mites, 3=50 to 100 mites, 4=100 to 500 mites, 5=500 to 1,000 mites, 6=1,000 to 10,000 mites, 7=10,000 mites.

Results presented in Table 2 show that mite populations on birds receiving feed treated with Compound 1 declined rapidly with most birds free of mites after 3 weeks of feeding. Untreated birds remained heavily infested throughout the trial. At the end of 5 weeks, mite-infested control birds maintained on untreated feed were switched to feed treated with the title compound. As before, mite-free birds were produced in 3 weeks (Table 3).
## Table 2

**Effect of Compound 1 as a Feed Additive (500 ppm) for Systemic Control of Northern Fowl Mite**

*Average mite index/bird* at weeks on additive

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Pretreatment</th>
<th>1 Wk.</th>
<th>2 Wk.</th>
<th>3 Wk.</th>
<th>4 Wk.</th>
<th>5 Wk.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 1</td>
<td>4.20</td>
<td>4.15</td>
<td>2.15</td>
<td>0.65</td>
<td>0.10</td>
<td>0.0</td>
</tr>
<tr>
<td>Untreated</td>
<td>3.75</td>
<td>4.44</td>
<td>6.10</td>
<td>4.40</td>
<td>4.30</td>
<td>2.50</td>
</tr>
</tbody>
</table>

*Mean value based on 20 birds/treatment.*

## Table 3

**Reduction in Mite Population on Previously Untreated Birds Switched Onto Feed-Additive (500 ppm)**

*Average mite index/bird* after being switched onto additive

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Pretreatment</th>
<th>1 Wk.</th>
<th>2 Wk.</th>
<th>3 Wk.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 1</td>
<td>2.50</td>
<td>1.15</td>
<td>0.1</td>
<td>0.9</td>
</tr>
</tbody>
</table>

*Mean value based on 20 birds/treatment.

"Untreated control birds from Table 1."
When the experiment was repeated using Compound 1 at a lower feed additive concentration (50 ppm), a similar result occurred (Table 4). Treated feed for this trial was prepared by mixing dry formulated wettable powder directly with the feed. The dry formulated wettable powder was prepared by uniformly mixing the following ingredients (percentages are all by weight):

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 1</td>
<td>30%</td>
</tr>
<tr>
<td>Aluminum silicate (Englehard ASP-172)</td>
<td>54%</td>
</tr>
<tr>
<td>ICI MX 2026 (a nonionic/ionic surfactant blend, ICI Americas, Inc.)</td>
<td>8%</td>
</tr>
<tr>
<td>Talc clay</td>
<td>5%</td>
</tr>
<tr>
<td>Ethylene oxide condensate of hydrophobic bases (Tergitol NP)</td>
<td>2%</td>
</tr>
<tr>
<td>Dioctyl sodium sulfosuccinate (Aerosol OTB)</td>
<td>1%</td>
</tr>
</tbody>
</table>

This mixture was then finely pulverized and remixed by mechanical or air-attrition grinding to obtain the desired wettable powder.
Table 4

Effect of Compound 1 as a Feed Additive (50 ppm) for Systemic Control of Northern Fowl Mite

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Pretreatment</th>
<th>1 Wk.</th>
<th>2 Wk.</th>
<th>3 Wk.</th>
<th>4 Wk.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 1</td>
<td>3.00</td>
<td>1.50</td>
<td>0.08</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Untreated</td>
<td>2.83</td>
<td>2.00</td>
<td>2.33</td>
<td>2.58</td>
<td>2.58</td>
</tr>
</tbody>
</table>

* Mean value based on 12 birds/treatment.

EXAMPLE IX

Systemic Control of Northern Fowl Mite *(Ornithonyssus sylviarum)*

The experiment described in Example VIII was conducted using Compound 2 in concentrations of 50 ppm and 5 ppm. Treated feed for this trial was prepared by mixing dry formulated wettable powder of Compound 2 directly with the feed. The dry formulated wettable powder was prepared by uniformly mixing the following ingredients (percentages are all by weight):

- Compound 2: 26.5%
- Kaolin clay: 61.2%
- Alkylaryl sulfonate: 4.0%
  *(Nacconol 90F)*
- Silica clay: 4.0%
  *(Zeolex 7)*
Dioctyl sodium sulfosuccinate (Aerosol OTB)
Sodium naphthalene formaldehyde condensate (Morwet D425)
Citric acid

This mixture was then finely pulverized and remixed by mechanical or air-attrition grinding to obtain the desired wettable powder.

Similar results as occurred in Example VIII were observed (Table 5) using the same mite index as in Example VIII. After 4 weeks the birds receiving Compound 2 in their feed were virtually free of mites even at 5 ppm of Compound 2. Untreated birds remained heavily infested with mites throughout the test period.

Table 5

Effect of Compound 2 as a Feed Additive for Systemic Control of Northern Fowl Mite

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Pretrt</th>
<th>1 wk</th>
<th>2 wk</th>
<th>3 wk</th>
<th>4 wk</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2 50</td>
<td>5.0</td>
<td>4.8</td>
<td>1.1</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 2 5</td>
<td>5.1</td>
<td>4.7</td>
<td>3.4</td>
<td>1.1</td>
<td>0.6</td>
</tr>
<tr>
<td>UNTREATED</td>
<td>-</td>
<td>4.1</td>
<td>4.4</td>
<td>4.3</td>
<td>3.4</td>
</tr>
</tbody>
</table>

* Mean value based on twelve birds/treatment.
EXAMPLE X

Systemic Control of Northern Fowl Mite on Chickens

White Leghorn pullets (20 weeks old at beginning of this experiment) of the Hy-Line strain were preinfested with northern fowl mites 3 weeks prior to being fed the treated feed. The infested birds were housed two per standard commercial cage with two cages per treatment (four birds) and the untreated controls. Each group of birds was provided with the corresponding treated feed continuously during the 4 weeks of each test. The feed was prepared by suspending one of the compounds (technical grade) identified in Tables 6 and 7 below in ethanol and then spraying the solution onto a standard poultry ration to provide 200 ppm of active ingredient (by weight). An untreated blank was prepared by blending ethanol alone with the ration. Mite counts on each bird were made at weekly intervals, and the number of mites expressed according to the standard index described hereinabove. The results are set for in Tables 6 and 7 below.
### Table 6
Systemic Control of Northern Fowl Mite on Chickens

Average mite index/bird* at weeks on additive

<table>
<thead>
<tr>
<th>Feed Additive</th>
<th>Pretreatment</th>
<th>1 wk</th>
<th>2 wk</th>
<th>3 wk</th>
<th>4 wk</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 5</td>
<td>5.0</td>
<td>2.2</td>
<td>0.0</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 6</td>
<td>5.0</td>
<td>6.2</td>
<td>6.0</td>
<td>4.8</td>
<td>1.0</td>
</tr>
<tr>
<td>Compound 7</td>
<td>4.0</td>
<td>2.8</td>
<td>1.5</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 8</td>
<td>4.2</td>
<td>6.0</td>
<td>0.8</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 9</td>
<td>4.8</td>
<td>6.0</td>
<td>4.0</td>
<td>1.5</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 10</td>
<td>4.8</td>
<td>6.2</td>
<td>6.5</td>
<td>6.0</td>
<td>3.2</td>
</tr>
<tr>
<td>Compound 11</td>
<td>3.8</td>
<td>3.0</td>
<td>0.8</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 17**</td>
<td>3.3</td>
<td>3.3</td>
<td>--</td>
<td>3.5</td>
<td>2.8</td>
</tr>
<tr>
<td>Compound 20**</td>
<td>3.5</td>
<td>3.7</td>
<td>--</td>
<td>2.2</td>
<td>1.5</td>
</tr>
<tr>
<td>Compound 21</td>
<td>5.8</td>
<td>6.2</td>
<td>4.5</td>
<td>2.2</td>
<td>0.2</td>
</tr>
<tr>
<td>Untreated</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control #1</td>
<td>4.8</td>
<td>6.0</td>
<td>6.5</td>
<td>6.5</td>
<td>6.3</td>
</tr>
<tr>
<td>Untreated</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control #2</td>
<td>4.3</td>
<td>5.5</td>
<td>6.3</td>
<td>6.8</td>
<td>7.0</td>
</tr>
</tbody>
</table>

* Mean value based on four birds/treatment.
** Administered at concentration of 250 ppm.
Table 7

Systemic Control of Northern Fowl Mite on Chickens

Average mite index/bird* at weeks on additive

<table>
<thead>
<tr>
<th>Feed Additive</th>
<th>Pretreatment</th>
<th>1 wk</th>
<th>2 wk</th>
<th>3 wk</th>
<th>4 wk</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 4</td>
<td>7.0</td>
<td>7.0</td>
<td>4.2</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 12</td>
<td>7.0</td>
<td>7.0</td>
<td>4.5</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 13</td>
<td>7.0</td>
<td>7.0</td>
<td>6.5</td>
<td>0.2</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 19</td>
<td>7.0</td>
<td>7.0</td>
<td>5.8</td>
<td>6.0</td>
<td>6.5</td>
</tr>
<tr>
<td>Compound 27</td>
<td>7.0</td>
<td>7.0</td>
<td>7.0</td>
<td>6.0</td>
<td>5.2</td>
</tr>
<tr>
<td>Compound 29</td>
<td>7.0</td>
<td>6.8</td>
<td>3.0</td>
<td>0.0</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 30</td>
<td>7.0</td>
<td>7.0</td>
<td>6.8</td>
<td>5.8</td>
<td>5.8</td>
</tr>
<tr>
<td>Untreated</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control #1</td>
<td>7.0</td>
<td>7.0</td>
<td>7.0</td>
<td>7.0</td>
<td>6.8</td>
</tr>
<tr>
<td>Untreated</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control #2</td>
<td>6.8</td>
<td>6.8</td>
<td>6.0</td>
<td>5.8</td>
<td>5.3</td>
</tr>
</tbody>
</table>

* Mean value based on four birds/treatment.

EXAMPLE XI

Pass-through Effects on Pests in Poultry Manure

Concurrent with monitoring feed additive effects of Compounds 1 and 2 on northern fowl mite, manure from the treated and untreated hens of Examples VIII and IX was collected to determine its toxicity to developing larvae of the house fly (*Musca domestica*). House fly eggs used in these tests were from a laboratory colony established from field-caught flies within a year of the test.

Manure collections were made by suspending plastic covered boards below the cages at intervals throughout the feeding studies of Examples VIII and IX. Samples were then placed in plastic cups with screen lids and inoculated with 50 fly eggs per cup. Adult fly emergence was monitored over a
period of 10-30 days. Also measured was the residual toxicity of manure after treated birds were returned to untreated feed (flushing effect).

Table 8 presents the results of the treated feed pass-through tests. Manure from birds fed Compound 1 or Compound 2 in their feed was highly toxic to house fly larvae. Few adults emerged from the droppings of the treated birds whereas a substantial percentage of adults emerged from the manure of the untreated birds.

Table 8

<table>
<thead>
<tr>
<th>Treatment</th>
<th>(ppm)</th>
<th>3</th>
<th>6</th>
<th>8</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 1</td>
<td>50</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Compound 2</td>
<td>50</td>
<td>0.5</td>
<td>0.8</td>
<td>0.5</td>
</tr>
<tr>
<td>Compound 2</td>
<td>5</td>
<td>2.6</td>
<td>1.6</td>
<td>3.2</td>
</tr>
<tr>
<td>UNTREATED</td>
<td>-</td>
<td>35.9</td>
<td>34.8</td>
<td>31.0</td>
</tr>
</tbody>
</table>

*Fifty house fly eggs initially added/replicate; 10 reps/treatment.

Table 9 presents the results when birds were returned to untreated feed after previously maintained on the feed additive for four weeks. Manure from birds fed rations treated with Compound 1 remained toxic to fly larvae 2-5 days after the birds were returned to untreated feed.
Table 9

Residual Effect of Compound 1 Feed Additive (50 ppm) on Fly Development Following Return of Birds to Untreated Feed**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Mean number emerging adult flies* (days following removal of additive)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>-1</td>
</tr>
<tr>
<td>Compound 1</td>
<td>0</td>
</tr>
<tr>
<td>CONTROL</td>
<td>34.1</td>
</tr>
</tbody>
</table>

*Fifty housefly eggs added per replicate; 10 reps/treatment.

**Birds previously maintained on the feed additive for four weeks.

EXAMPLE XII

Pass-through Effects on Pests in Poultry Manure

Concurrent with monitoring feed additive effects of the compounds in Example X on northern fowl mite, manure from the treated and untreated chickens of Example X was collected to determine its toxicity to developing larvae of the house fly (Musca domestica). House fly eggs used in these tests were from a laboratory colony established from field-caught flies within a year of the test.

Manure collections were made by suspending plastic covered boards below the cages at intervals throughout the feeding studies of Example X. Samples were then placed in plastic cups with screen lids and inoculated with 50 fly eggs per cup. Adult fly emergence was monitored over a period of 10-30 days.
Table 10 presents the results of the treated feed pass-through tests. Manure from birds fed certain compounds identified in Table 10 in their feed was highly toxic to house fly larvae. Few adults emerged from the droppings of the treated birds whereas a substantial percentage of adults emerged from the manure of the untreated birds.
### Table 10

**Effects of Certain Compounds Used as Feed Additives on House Fly Development in Manure from Treated Birds**

<table>
<thead>
<tr>
<th>Feed Additive</th>
<th>Percent Inhibition of Adult Fly Emergence Relative to Untreated Control*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 3</td>
<td>99.7</td>
</tr>
<tr>
<td>Compound 4</td>
<td>100.0</td>
</tr>
<tr>
<td>Compound 5</td>
<td>65.3</td>
</tr>
<tr>
<td>Compound 6</td>
<td>73.1</td>
</tr>
<tr>
<td>Compound 7</td>
<td>73.8</td>
</tr>
<tr>
<td>Compound 8</td>
<td>77.8</td>
</tr>
<tr>
<td>Compound 9</td>
<td>99.4</td>
</tr>
<tr>
<td>Compound 10</td>
<td>100.0</td>
</tr>
<tr>
<td>Compound 11</td>
<td>100.0</td>
</tr>
<tr>
<td>Compound 12</td>
<td>100.0</td>
</tr>
<tr>
<td>Compound 13</td>
<td>100.0</td>
</tr>
<tr>
<td>Compound 16</td>
<td>100</td>
</tr>
<tr>
<td>Compound 18</td>
<td>20.6</td>
</tr>
<tr>
<td>Compound 19</td>
<td>65.9</td>
</tr>
<tr>
<td>Compound 21</td>
<td>90.7</td>
</tr>
<tr>
<td>Compound 22</td>
<td>34.4</td>
</tr>
<tr>
<td>Compound 23</td>
<td>87.1</td>
</tr>
<tr>
<td>Compound 24</td>
<td>92.4</td>
</tr>
<tr>
<td>Compound 25</td>
<td>31.5</td>
</tr>
<tr>
<td>Compound 26</td>
<td>75.1</td>
</tr>
<tr>
<td>Compound 27</td>
<td>100</td>
</tr>
<tr>
<td>Compound 28</td>
<td>98.0</td>
</tr>
<tr>
<td>Compound 29</td>
<td>87.1</td>
</tr>
<tr>
<td>Compound 30</td>
<td>12.3</td>
</tr>
</tbody>
</table>

*Based on two weekly manure collections; 6 replicates per treatment, fifty house fly eggs added per replicate.

### EXAMPLE XIII

**Systemic Control of Poultry Lice (Menacanthus stramineus)**

White leghorn laying hens housed in commercial egg laying cages were preinfested with lice by adding approximately 50 lice in all stages.
of development to each bird 3 weeks prior to testing. There were 8 birds each (2 per cage, 4 cages) for the treatment and for the control.

The treated feed was prepared by mixing 1.135g of Compound 2 (technical grade) with 1 lb. of standard 16% protein laying mash. The 1 lb. of treated feed was then mixed with 99 lbs of laying mash to give a final concentration of Compound 2 in the feed equal to 25 ppm. The control birds were fed the standard laying mash minus Compound 2. The feed, treated or untreated, was provided continuously during the 5 weeks of the test.

Lice populations on each bird were estimated by a standard procedure pretreatment and at weekly intervals after the birds were receiving the treated feed. The lice populations per bird were estimated by recording the total number of lice (nymphs and adults) observed upon parting the feathers in 5 locations (vent area, left thigh, right thigh, left side of breast, right side of breast). Thus, the index in Table 11 is the "no. of lice per 5 feather parts per bird".

As shown in Table 11, the lice population on the birds fed Compound 2 at 25 ppm rapidly declined. The lice population was extremely low after 2 weeks of feeding, and nearly eliminated after 3 weeks of feeding. No lice could be found on any birds after 4 and 5 weeks of feeding. Also, no clusters of lice eggs appeared on the feathers.

Concurrently, there was a gradual increase in the lice population on the untreated control birds. On these birds the lice population increased
3-4 fold during the 5 weeks of the test. Also, there were many lice egg masses on the feathers of these birds.

Lice on chickens are normally very difficult to control but when provided in as little as 25 ppm in the feed of the birds, Compound 2 completely controlled infestations of chicken lice in about 3 weeks.
Table 11

Effect of Compound 2 as a Feed Additive at 25 ppm (technical sprayed on 1 lb. feed aliquot) on Populations of Chicken Lice (Menacanthus stramineus)

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Total No. Lice Per 5 feather Parts Per Bird</th>
<th>Mean</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pretreatment</td>
<td></td>
</tr>
<tr>
<td>Compound 2</td>
<td>47 20 22 74 14 31 21 52</td>
<td>35.1</td>
</tr>
<tr>
<td>Untreated</td>
<td>17 48 64 87 13 11 20 87</td>
<td>43.4</td>
</tr>
<tr>
<td></td>
<td>1 wk. Posttreatment</td>
<td></td>
</tr>
<tr>
<td>Compound 2</td>
<td>41 34 10 56 7 16 29 31</td>
<td>28.0</td>
</tr>
<tr>
<td>Untreated</td>
<td>115 130 100 140 66 10 210 55</td>
<td>103.2</td>
</tr>
<tr>
<td></td>
<td>2 wk. Posttreatment</td>
<td></td>
</tr>
<tr>
<td>Compound 2</td>
<td>9 2 0 15 1 0</td>
<td>3.6</td>
</tr>
<tr>
<td>Untreated</td>
<td>165 200 135 110 13 45 85 145</td>
<td>112.2</td>
</tr>
<tr>
<td></td>
<td>3 wk. Posttreatment</td>
<td></td>
</tr>
<tr>
<td>Compound 2</td>
<td>2 0 0 0 0 0 0 0</td>
<td>0.2</td>
</tr>
<tr>
<td>Untreated</td>
<td>160 135 200 190 37 85 85 210</td>
<td>137.7</td>
</tr>
<tr>
<td></td>
<td>4 wk. Posttreatment</td>
<td></td>
</tr>
<tr>
<td>Compound 2</td>
<td>0 0 0 0 0 0 0 0</td>
<td>0</td>
</tr>
<tr>
<td>Untreated</td>
<td>250 250 155 225 26 110 145 185</td>
<td>168.2</td>
</tr>
<tr>
<td></td>
<td>5 wk. Posttreatment</td>
<td></td>
</tr>
<tr>
<td>Compound 2</td>
<td>0 0 0 0 0 0 0 0</td>
<td>0</td>
</tr>
<tr>
<td>Untreated</td>
<td>202 185 155 105 45 77 131 185</td>
<td>135.6</td>
</tr>
</tbody>
</table>
EXAMPLE XIV

Systemic Control of Cattle Lice

Systemic and direct contact action were evaluated against cattle lice on cattle naturally infested with cattle lice. Compound 1 was administered orally in the feed, as a wipe-on (pour-on) salve along the backline and as a full-coverage contact spray. Compounds 2 and 3 were administered orally in feed and as a wipe-on (pour-on) salve.

Cattle on feed supplements received 8.85 mg ai/kg body wt/day by blending dry wettable powder formulations into a premix of ground grain, minerals, and salt. The dry wettable powder formulations for Compound 1 and 2 were prepared as described in Example VIII and IX respectively. The dry wettable powder formulation for Compound 3 was prepared by uniformly mixing the following ingredients (percentages are all by weight):

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 3</td>
<td>25.0%</td>
</tr>
<tr>
<td>Aluminum Silicate (Englehard ASP-172)</td>
<td>64.5</td>
</tr>
<tr>
<td>Synthetic Silica (Zeolux 7)</td>
<td>2.0</td>
</tr>
<tr>
<td>Citric Acid</td>
<td>0.5</td>
</tr>
<tr>
<td>Sodium Naphthalene</td>
<td>2.0</td>
</tr>
<tr>
<td>Formaldehyde Condensate (Morwet D425)</td>
<td></td>
</tr>
<tr>
<td>Alkylaryl Sulfonate (Nacconol 90F)</td>
<td>4.0</td>
</tr>
<tr>
<td>Dioctyl Sodium</td>
<td>2.0</td>
</tr>
<tr>
<td>Sulfoosuccinate (Aerosol OTB)</td>
<td></td>
</tr>
</tbody>
</table>

This mixture was then finely pulverized and remixed by mechanical, or air-attrition grinding to obtain the desired wettable powder.
Wipe-on formulations were rubbed into the hair along the backline at a rate of 4 oz. form./animal (35 mg ai/kg) (single application). These formulations were prepared by mixing the following ingredients together in a ball mill to produce the following suspensions (percentages are all by weight):

<table>
<thead>
<tr>
<th>Compound</th>
<th>I</th>
<th>II</th>
<th>III</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>5.0%</td>
<td>1.9%</td>
<td>5.0%</td>
</tr>
<tr>
<td>Carageen-based flow</td>
<td>4.5</td>
<td>2.0</td>
<td>4.5</td>
</tr>
<tr>
<td>modifier*</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dioctyl sodium</td>
<td>0.4</td>
<td>0.4</td>
<td>0.4</td>
</tr>
<tr>
<td>sulfosuccinate</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cotton seed oil</td>
<td>90.1</td>
<td>96.1</td>
<td>90.1</td>
</tr>
</tbody>
</table>

*A finely divided, solidified vegetable oil derivative which when dispersed by strong shearing action in non-aqueous media, swells to form a thixotropic gel structure.

Compound 1 was additionally applied as a whole-animal contact spray. The spray was prepared by mixing together the following ingredients in a ball mill (all percentages by weight):
This mixture is then diluted with water and applied to the animal at 0.03 lb ai/gal (1 gal/animal). Cattle averaged 430 lb/head. One animal was assigned per treatment with the exception of the control group which contained 2 animals.

Parasite loads were measured before and after treatment at 6 weekly intervals. Separate louse counts were taken from head, shoulder and sides, and tail-rump areas of each animal. The hair was parted using a 5-inch blade along which the number of lice/inch were counted.

Louse counts taken from head, shoulder, side, rump, and tail areas of treated cattle confirm that the compounds moved systemically throughout the animal following ingestion or localized penetration in the area of the pour-on (Table 8). Excellent control of both chewing (Bovicola bovis) and sucking (Linognathus vituli) lice was achieved within 2 - 3 weeks of initial treatment by all routes of administration. After 4 weeks the cattle administered Compound 1 or 2 were virtually lice-free whereas the untreated cattle remained heavily infested.
### Table 12
Systemic Control of Cattle Lice with Compounds 1 and 2 Administered as Feed Additives, Pour-ons and Contact Sprays.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Compound 1</th>
<th>Compound 2</th>
<th>Compound 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>FA</td>
<td>.35</td>
<td>.35</td>
<td>.35</td>
</tr>
<tr>
<td>PO</td>
<td>.35</td>
<td>33</td>
<td>35</td>
</tr>
<tr>
<td>SP</td>
<td>.35</td>
<td>18</td>
<td>15</td>
</tr>
<tr>
<td>Untreated</td>
<td>#1 .35</td>
<td>#2 .35</td>
<td>#3 .35</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Mean number lice per inch (weeks posttreatment)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Treatment</td>
</tr>
<tr>
<td>-----------</td>
</tr>
<tr>
<td>Compound 1 FA</td>
</tr>
<tr>
<td>Compound 1 PO</td>
</tr>
<tr>
<td>Compound 1 SP</td>
</tr>
<tr>
<td>Compound 2 FA</td>
</tr>
<tr>
<td>Compound 2 PO</td>
</tr>
<tr>
<td>Compound 3 FA</td>
</tr>
<tr>
<td>Compound 3 PO</td>
</tr>
<tr>
<td>Untreated #1 .35</td>
</tr>
<tr>
<td>Untreated #2 .35</td>
</tr>
</tbody>
</table>

- 90% *Bovicola bovis; 10% Linognathus vituli*
- Composite counts from all locations; three 5-inch parts/location
- FA = Feed additive; PO = Pour-on; SP = spray.

**EXAMPLE XV**
Systemic Control of Cattle Nematodes
Systemic action was evaluated against cattle nematodes on cattle naturally infested with internal roundworms. Compound 3 was administered
orally in the feed and as a wipe-on (pour-on) salve along the backline.

Cattle on feed supplements received 8.85 mg ai/kg body wt/day by blending dry wettable powder formulations into a premix of ground grain, minerals, and salt. The dry wettable powder formulation for Compound 3 was prepared as described in Example XIV.

Wipe-on formulations were rubbed into the hair along the backline at a rate of 4 oz. form./animal (35 mg ai/kg body wt.) (single application). The wipe-on formulation was prepared as described in Example XIV.

Cattle averaged 430 lb/head. One animal was assigned per treatment with the exception of the control group which contained 2 animals.

Parasite loads were measured before and after treatment at 6 weekly intervals. Intestinal roundworms (primarily strongyles) were monitored using a standard salt fecal floatation technique. Walnut-size portions of manure collected from treated animals were placed in paper cups three-fourths full of a saturated salt solution. After stirring, the resultant slurry was filtered through cheese cloth into a shell vial and a coverslip placed over the top. After 20 minutes, the coverslip was removed, placed on a slide and the total number of worm eggs per slide counted under a light microscope. Nematode egg counts from fecal samples are summarized in Table 13. After 4 weeks, the cattle administered Compound 3 exhibited significantly reduced nematode egg counts in comparison with the untreated cattle.
Table 13

Systemic Control of Cattle Nematodes with Compound 3 Administered as Feed Additives and Pour-ons

Number of Nematode Eggs/Fecal Float (weeks posttreatment)

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Pretrt.</th>
<th>1 wk</th>
<th>2 wk</th>
<th>3 wk</th>
<th>4 wk</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 3 FA</td>
<td>127</td>
<td>34</td>
<td>6</td>
<td>42</td>
<td>59</td>
</tr>
<tr>
<td>Compound 3 PO</td>
<td>93</td>
<td>140</td>
<td>51</td>
<td>26</td>
<td>48</td>
</tr>
<tr>
<td>Untreated #1</td>
<td>400</td>
<td>400</td>
<td>400</td>
<td>400</td>
<td>400</td>
</tr>
<tr>
<td>Untreated #2</td>
<td>6</td>
<td>10</td>
<td>3</td>
<td>220</td>
<td>201</td>
</tr>
</tbody>
</table>

(a) FA = Feed Additive; PO = Pour-on

EXAMPLE XVI

Systemic Control of Sarcoptic Mange in Pigs

Systemic action was evaluated against the mite *Sarcoptes scabiei* in pigs infested with the mite. Compound 2 was administered orally in the feed or as a wipe-on (pour-on) salve.

Pigs on feed supplements received 6 mg ai/kg body wt/day by blending dry wettable powder formulations into the feed. The dry wettable powder formulation for Compound 2 was prepared as described in Example IX.

Wipe-on formulations were rubbed into the back or other area of the animal at a rate of 35 mg ai/kg body wt in a single application. The wipe-on formulations were prepared by mixing the following
ingredients together in a ball mill to produce a suspension (percentages are all by weight):

<table>
<thead>
<tr>
<th>Compound 2</th>
<th>1.9%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carageenen-based flow modifier*</td>
<td>2.0</td>
</tr>
<tr>
<td>Dioctyl sodium sulfosuccinate</td>
<td>0.4</td>
</tr>
<tr>
<td>Petroleum crop oil</td>
<td>96.1</td>
</tr>
</tbody>
</table>

*A finely divided, solidified vegetable oil derivative which when dispersed by strong shearing action in non-aqueous media, swells to form a thixotropic gel structure.

Parasitic loads were measured before and after treatment at 3 weekly intervals. Excellent systemic control of the mite was achieved within 1 week of initial treatment by both routes of administration (Table 14).

Table 14

**Systemic Control of Sarcoptic Mange in Pigs with Compound 2 Administered as Feed Additive and Pour-on**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Pretreatment</th>
<th>1 Wk</th>
<th>2 Wk</th>
<th>3 Wk</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2 FA</td>
<td>32.8</td>
<td>0.5</td>
<td>1.5</td>
<td>0.0</td>
</tr>
<tr>
<td>Compound 2 PO</td>
<td>62.8</td>
<td>1.3</td>
<td>4.3</td>
<td>0.0</td>
</tr>
<tr>
<td>UNTREATED</td>
<td>25.0</td>
<td>16.8</td>
<td>22.5</td>
<td>11.0</td>
</tr>
</tbody>
</table>

*Based on one scraping/animal; 4 animals/treatment
EXAMPLE XVII
Systemic Control of Rodent Bot Fly

Systemic activity was evaluated against rodent bot fly on Peramiscus mice infested with rodent bot fly (Cuterebra fontinella). Larval development of this parasite takes place entirely within the body of the host. All compounds identified in Table 15 were administered orally in the feed as part of the daily ration. There were fifteen (15) mice each for the treatments and for the controls. A separate control was used for each treatment. The treated diet was prepared by pulverizing standard laboratory rat chow pellets, dry-incorporating the technical material and repelletizing. The final feeding concentration of each treatment was 200 ppm. Infestation was accomplished by manually placing 3-4 newly hatched bot fly larvae into the eyes of each mouse. Effectiveness was determined by periodically examining the inguinal region of the mouse for breathing holes cut by the migrating bot fly larvae. Further larval development and emergence from the body of the host was also monitored and recorded. All compounds were offered continuously throughout the experiment as well as during a 5-7 day period prior to infestation with the parasite.

The results are set forth in Table 15 below.
Table 15
Systemic Effect on Rodent Bot Fly

<table>
<thead>
<tr>
<th>Feed Additive</th>
<th>Percent Mortality*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 3</td>
<td>100</td>
</tr>
<tr>
<td>Compound 4</td>
<td>100</td>
</tr>
<tr>
<td>Compound 5</td>
<td>90</td>
</tr>
<tr>
<td>Compound 6</td>
<td>97</td>
</tr>
<tr>
<td>Compound 7</td>
<td>80</td>
</tr>
<tr>
<td>Compound 8</td>
<td>94</td>
</tr>
<tr>
<td>Compound 9</td>
<td>63</td>
</tr>
<tr>
<td>Compound 10</td>
<td>100</td>
</tr>
<tr>
<td>Compound 11</td>
<td>100</td>
</tr>
<tr>
<td>Compound 12</td>
<td>100</td>
</tr>
<tr>
<td>Compound 13</td>
<td>100</td>
</tr>
<tr>
<td>Compound 14</td>
<td>93</td>
</tr>
<tr>
<td>Compound 15</td>
<td>100</td>
</tr>
<tr>
<td>Compound 16</td>
<td>54</td>
</tr>
<tr>
<td>Compound 17</td>
<td>10</td>
</tr>
<tr>
<td>Compound 18</td>
<td>48</td>
</tr>
<tr>
<td>Compound 19</td>
<td>22</td>
</tr>
<tr>
<td>Compound 20</td>
<td>100</td>
</tr>
<tr>
<td>Compound 21</td>
<td>10</td>
</tr>
<tr>
<td>Compound 22</td>
<td>20</td>
</tr>
<tr>
<td>Compound 23</td>
<td>38</td>
</tr>
</tbody>
</table>

* Based on the number of air holes cut in treated mice in comparison with air holes cut in untreated mice for each treatment.

The results from Table 15 show a systemic effect as indicated by the major reduction in the number of air holes cut in treated mice.

EXAMPLE XVIII
Systemic Effect Against Flea Reproduction on Cats

Systemic activity was evaluated against fleas on cats infested with fleas (*Ctenocephalides* spp.). Treated feed for this experiment was prepared by mixing dry formulated wettable powder of the compounds identified in Tables 16 and 17.
directly with the feed. The dry formulated wettable powder was prepared by uniformly mixing the following ingredients (percentages are all by weight):

- Compound 2 or 8 25.0%
- Aluminum Silicate (Englehard ASP-172) 64.5%
- Synthetic Silica (Zeolex 7) 2.0%
- Citric Acid 0.5%
- Sodium Naphthalene Formaldehyde Condensate (Morwet D425) 2.0%
- Alkylaryl Sulfonate 4.0%
- (Nacconol 90F)
- Dioctyl Sodium Sulfosuccinate 2.0%
- (Aerosol OTB)

This mixture was then finely pulverized and remixed by mechanical or air-attrition grinding to obtain the desired wettable powder. The wettable powders were mixed into a portion of the daily ration (10 mg ai/kg body wt/day) and administered orally to the cats which were housed individually in galvanized wire cages. Flea infestations were established and maintained by infesting the test animals with adult fleas prior to and at weekly intervals after beginning treatment. Fifty, 7 to 14 day old, unfed adult fleas were placed along the dorsal midline of each cat. Three cats were used per treatment, each cat receiving the feed supplements for one month.

Treatment effects on adult fleas were assessed by combing them from the cats each week. To determine possible reproductive effects on the
subsequent generation, eggs from these adults were collected on plywood boards suspended beneath the cage of each cat. Flea eggs were swept from the boards with a soft nylon brush, transferred to larval rearing media and held for development through one generation.

Table 16
Systemic Effect on Progeny of Adult Fleas Feeding on Cats

<table>
<thead>
<tr>
<th>Feed Additive</th>
<th>-1</th>
<th>0</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>5</th>
<th>7</th>
<th>14</th>
<th>21</th>
<th>28</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2</td>
<td>328</td>
<td>335</td>
<td>3</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Compound 8</td>
<td>257</td>
<td>237</td>
<td>13</td>
<td>20</td>
<td>3</td>
<td>0</td>
<td>9</td>
<td>1</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Untreated</td>
<td>404</td>
<td>235</td>
<td>191</td>
<td>343</td>
<td>407</td>
<td>275</td>
<td>266</td>
<td>129</td>
<td>262</td>
<td>59</td>
</tr>
</tbody>
</table>

*Counts based on 3 egg sweepings/treatment/sample (one sweeping per cat).

Table 17
Systemic Effect on Progeny of Adult Fleas Feeding on Cats Which Received Single Oral Dose**

<table>
<thead>
<tr>
<th>Feed Additive</th>
<th>0</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2</td>
<td>544</td>
<td>29</td>
<td>1</td>
<td>11</td>
<td>32</td>
</tr>
<tr>
<td>Untreated</td>
<td>169</td>
<td>248</td>
<td>321</td>
<td>484</td>
<td>314</td>
</tr>
</tbody>
</table>

*Counts based on 3 eggs sweepings/treatment/sample (one sweeping per cat).

**Single dose of 10 mg ai/kg body wt. of Compound 2 administered on day zero.
Although adult fleas feeding on the treated animals were not directly affected, the results from Tables 16 and 17 show that development of their offspring was significantly inhibited. This effect was particularly dramatic after 2 days of receiving the feed supplement, i.e., no F₁ progeny developed through to the adult stage. Examination of the larval rearing containers indicated that the lethal effect on the progeny of exposed adults occurred very early in development; probably within the egg or shortly after hatch. The results from Table 17 indicate that a single 10 mg/kg body weight dose administered orally to cats also provides excellent control (98% nonviable eggs) for 3 days. Roughly 90% nonviable eggs were still being produced 5 days after treatment.

EXAMPLE XIX
Systemic Control of Black Blowfly in Mice

Male Swiss-Webster mice weighing 30 grams or more were administered the compounds identified in Table 18 via gavage. Four mice were used for each treatment and the controls. The compounds were formulated by mixing technical grade samples of each compound identified in Table 18 with Tween 20K to a 4% concentration. The mice were dosed at a concentration of 400 mg active ingredient/kg body weight. The mice used in the positive control (bromophos) were dosed at a concentration of 100 mg active ingredient/kg body weight. The mice were euthanized 16 hours following treatment and the rear legs excised. Twenty first instar black blowfly larvae (Phormia regina (Meigen)) were introduced to
the excised mouse leg muscle tissue. Blowfly development was monitored (i) 48 hours after introduction to muscle tissue (as a measure of direct larval mortality), (ii) 5 days after introduction to muscle tissue (as a measure of inhibition of pupal development), and (iii) through to adult emergence of the control group (approximately 10-12 days after pupation). Percent mortality was determined by observation of the number of surviving larvae, pupae or adults in treated mice in comparison with the untreated controls.

Table 18
Systemic Control of Black Blowfly in Mice

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Larvae</th>
<th>Pupae</th>
<th>Adults</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 3</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 4</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 5</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 6</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 8</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 9</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 10</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 11</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 12</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 13</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 14</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 15</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 16</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 21</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 23</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 24</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 26</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 27</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 28</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 31</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 32</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 33</td>
<td>0</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>Compound</td>
<td>0</td>
<td>0</td>
<td>10</td>
</tr>
<tr>
<td>----------</td>
<td>---</td>
<td>---</td>
<td>----</td>
</tr>
<tr>
<td>Compound 35</td>
<td>0</td>
<td>0</td>
<td>45</td>
</tr>
<tr>
<td>Compound 36</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 37</td>
<td>0</td>
<td>0</td>
<td>35</td>
</tr>
<tr>
<td>Compound 38</td>
<td>0</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>Compound 39</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 40</td>
<td>0</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>Compound 41</td>
<td>*</td>
<td>*</td>
<td>80</td>
</tr>
<tr>
<td>Compound 42</td>
<td>0</td>
<td>*</td>
<td>80</td>
</tr>
<tr>
<td>Compound 43</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 44</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 45</td>
<td>0</td>
<td>*</td>
<td>40</td>
</tr>
<tr>
<td>Compound 46</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 47</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 48</td>
<td>0</td>
<td>*</td>
<td>55</td>
</tr>
<tr>
<td>Compound 49</td>
<td>0</td>
<td>--</td>
<td>45</td>
</tr>
<tr>
<td>Compound 50</td>
<td>0</td>
<td>*</td>
<td>65</td>
</tr>
<tr>
<td>Compound 51</td>
<td>0</td>
<td>*</td>
<td>20</td>
</tr>
<tr>
<td>Compound 52</td>
<td>0</td>
<td>*</td>
<td>70</td>
</tr>
<tr>
<td>Compound 53</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 54</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 55</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 56</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 57</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 58</td>
<td>0</td>
<td>*</td>
<td>35</td>
</tr>
<tr>
<td>Compound 59</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 60</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 61</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 62</td>
<td>100</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Compound 63</td>
<td>0</td>
<td>*</td>
<td>90</td>
</tr>
<tr>
<td>Compound 64</td>
<td>0</td>
<td>*</td>
<td>65</td>
</tr>
<tr>
<td>Compound 65</td>
<td>0</td>
<td>0</td>
<td>65</td>
</tr>
<tr>
<td>Compound 66</td>
<td>0</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>Compound 67</td>
<td>0</td>
<td>0</td>
<td>20</td>
</tr>
<tr>
<td>Compound 68</td>
<td>0</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>Compound 69</td>
<td>0</td>
<td>0</td>
<td>25</td>
</tr>
<tr>
<td>Compound 70</td>
<td>0</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>Compound 71</td>
<td>*</td>
<td>*</td>
<td>70</td>
</tr>
<tr>
<td>Compound 72</td>
<td>0</td>
<td>0</td>
<td>70</td>
</tr>
<tr>
<td>Compound 73</td>
<td>0</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>Compound 74</td>
<td>0</td>
<td>0</td>
<td>35</td>
</tr>
<tr>
<td>Compound 75</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound 76</td>
<td>0</td>
<td>0</td>
<td>20</td>
</tr>
<tr>
<td>Compound 77</td>
<td>0</td>
<td>*</td>
<td>65</td>
</tr>
<tr>
<td>Compound 78</td>
<td>*</td>
<td>*</td>
<td>50</td>
</tr>
<tr>
<td>Compound 79</td>
<td>0</td>
<td>0</td>
<td>75</td>
</tr>
<tr>
<td>Compound 80</td>
<td>0</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>Compound 81</td>
<td>*</td>
<td>100</td>
<td>--</td>
</tr>
<tr>
<td>Compound</td>
<td>Effect</td>
<td>Effect</td>
<td>Effect</td>
</tr>
<tr>
<td>--------------</td>
<td>--------</td>
<td>--------</td>
<td>--------</td>
</tr>
<tr>
<td>82</td>
<td>0</td>
<td>0</td>
<td>15</td>
</tr>
<tr>
<td>83</td>
<td>0</td>
<td>0</td>
<td>30</td>
</tr>
<tr>
<td>84</td>
<td>0</td>
<td>0</td>
<td>40</td>
</tr>
<tr>
<td>85</td>
<td>*</td>
<td>*</td>
<td>100</td>
</tr>
<tr>
<td>86</td>
<td>0</td>
<td>0</td>
<td>10</td>
</tr>
<tr>
<td>87</td>
<td>*</td>
<td>*</td>
<td>100</td>
</tr>
<tr>
<td>88</td>
<td>*</td>
<td>100</td>
<td>---</td>
</tr>
<tr>
<td>89</td>
<td>*</td>
<td>100</td>
<td>---</td>
</tr>
<tr>
<td>90</td>
<td>*</td>
<td>100</td>
<td>---</td>
</tr>
<tr>
<td>91</td>
<td>100</td>
<td>--</td>
<td>---</td>
</tr>
<tr>
<td>92</td>
<td>100</td>
<td>--</td>
<td>---</td>
</tr>
<tr>
<td>93</td>
<td>*</td>
<td>100</td>
<td>---</td>
</tr>
<tr>
<td>94</td>
<td>*</td>
<td>100</td>
<td>---</td>
</tr>
<tr>
<td>95</td>
<td>*</td>
<td>*</td>
<td>70</td>
</tr>
<tr>
<td>96</td>
<td>100</td>
<td>--</td>
<td>---</td>
</tr>
<tr>
<td>97</td>
<td>*</td>
<td>100</td>
<td>---</td>
</tr>
<tr>
<td>98</td>
<td>*</td>
<td>*</td>
<td>95</td>
</tr>
<tr>
<td>99</td>
<td>0</td>
<td>0</td>
<td>5</td>
</tr>
<tr>
<td>Untreated Control</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Tween 20K</td>
<td>100</td>
<td>--</td>
<td>---</td>
</tr>
</tbody>
</table>

* Indicates some control of larvae, however, not complete control.

EXAMPLE XX
Systemic Control of Endoparasitic Cestodes in Mice

Male Swiss-Webster mice weighing 30 grams or more which had been orally infected with tapeworm (Hymenolepis nana) eggs were treated with the compounds identified in Table 19. Four mice were used for each treated host and control and two mice were used for each nontreated host. Yomesan was used as a positive control. The compounds were administered as feed additive rations for 18 consecutive days at a concentration of 1000 ppm. Feed additive rations were prepared by uniformly blending technical grade samples of each compound identified in Table 19 with pulverized mice chow.
(Purina brand Rat Chow #5001). Mice were necropsied after 18 days on the ration and tapeworm development was monitored. Tapeworm eggs found in the gut of the treated mice were introduced into a second group of noninfected and nontreated mice (2 mice per treatment). These mice were each necropsied 14 days after infestation, and again tapeworm development was monitored. Percent mortality was determined by observation of tapeworm development in the gut of mice in comparison with the untreated control, and is based on the number of infected mice in comparison with the number of uninfected mice. The results in Table 19 demonstrate a reduction in tapeworm offspring in the nontreated host. The results in Table 19 also demonstrate direct adult tapeworm developmental inhibition by compounds 56 and 64.

<table>
<thead>
<tr>
<th>TREATMENT</th>
<th>TREATED HOST</th>
<th>NON-TREATED HOST</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 5</td>
<td>0</td>
<td>50</td>
</tr>
<tr>
<td>Compound 10</td>
<td>0</td>
<td>50</td>
</tr>
<tr>
<td>Compound 13</td>
<td>0</td>
<td>50</td>
</tr>
<tr>
<td>Compound 15</td>
<td>0</td>
<td>50</td>
</tr>
<tr>
<td>Compound 56</td>
<td>50</td>
<td>--</td>
</tr>
<tr>
<td>Compound 64</td>
<td>75*</td>
<td>--</td>
</tr>
<tr>
<td>Untreated Control</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Yomesan</td>
<td>100</td>
<td>--</td>
</tr>
</tbody>
</table>

*Growth size stunted about 75 percent for surviving adult tapeworms (25 percent).
EXAMPLE XXI

Systemic Control of Cat Flea F₁ Progeny

Adult cats weighing 4.5-6.0 kg were treated with feed additive rations of the compounds identified in Table 20 for seven consecutive days. One cat was used for each treatment and the control. Feed additive rations were prepared by uniformly blending technical grade samples of each compound identified in Table 20 with the daily food to attain a 10 mg active ingredient/kg body weight dosage. Twenty-four hours after the cat hosts were given the feed additive ration for day number 1, each cat was infested with 100 adult cat fleas [Ctenocephalides felis (Bouche')] as 50:50 mixture of males and females. Cat flea ova were collected from pans suspended beneath each cage at various intervals (day numbers 4, 5 and 6), while the host cat was on the 7 day ration. Collected ova were cultured to determine hatch and development to adult stadium. The results are given in Table 20. Percent mortality was determined by observation of the collected ova which developed in comparison with the untreated control.
TABLE 20
Systemic Control of Cat Flea $F_1$-Progeny

<table>
<thead>
<tr>
<th>FEED ADDITIVE</th>
<th>PERCENT MORTALITY OF CAT FLEA $F_1$-PROGENY</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2</td>
<td>100</td>
</tr>
<tr>
<td>Compound 5</td>
<td>100</td>
</tr>
<tr>
<td>Compound 14</td>
<td>28</td>
</tr>
<tr>
<td>Compound 15</td>
<td>99.5</td>
</tr>
<tr>
<td>Compound 26</td>
<td>60.5</td>
</tr>
<tr>
<td>Compound 28</td>
<td>100</td>
</tr>
<tr>
<td>Untreated Control</td>
<td>0</td>
</tr>
</tbody>
</table>

EXAMPLE XXII
Systemic Control of Endoparasitic Nematodes in Mice

Male Swiss-Webster mice weighing 30 grams or more which had been orally infected with threadworm (Nematospiroides dubius) third stage larvae were treated with compounds identified in Table 21. Four mice were used for each treated host and the controls. Thiabendazole was used as a positive control. The compounds were administered as feed additive rations for 18 consecutive days at a concentration of 1000 ppm. Feed additive rations were prepared by uniformly blending technical grade samples of each compound identified in Table 21 with pulverized mice chow (Purina brand Rat Chow #5001). Mice were necropsied after 18 days on the ration and threadworm development was monitored. On days 12, 13 and 14 of the 18 day period, fecal pellets were collected from the mice and any threadworm eggs which had passed into the feces were cultured to third stage larvae ($F_1$). Culturing occurred in a clean petri dish at ambient temperature.
Development from egg to third stage larvae occurred in about 7-10 days. The results in Table 21 are based on four mice per treatment for the treated host and the controls. Percent mortality for the treated host was determined by observation of threadworm in the necropsied mice after 18 days. Percent mortality for F1 larvae was determined by observation of third stage larvae in the fecal material in comparison with the untreated control for days 12, 13 and 14 of the 18 day period. The results in Table 21 demonstrate a reduction in F1 threadworm larvae in comparison with the untreated control for certain of the compounds. The results in Table 21 also demonstrate stunted threadworm size development in the treated host after 18 days.

TABLE 21
Systemic Control of Endoparasitic Nematodes (Threadworm) in Mice

<table>
<thead>
<tr>
<th>TREATMENT</th>
<th>TREATED HOST (18 days)</th>
<th>LARVAE DEVELOPMENT (F1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Compound 2</td>
<td>0*</td>
<td>100</td>
</tr>
<tr>
<td>Compound 4</td>
<td>0*</td>
<td>100</td>
</tr>
<tr>
<td>Compound 5</td>
<td>0*</td>
<td>0</td>
</tr>
<tr>
<td>Compound 9</td>
<td>0*</td>
<td>0</td>
</tr>
<tr>
<td>Compound 11</td>
<td>0*</td>
<td>33</td>
</tr>
<tr>
<td>Compound 13</td>
<td>0*</td>
<td>0</td>
</tr>
<tr>
<td>Compound 15</td>
<td>0*</td>
<td>33</td>
</tr>
<tr>
<td>Compound 21</td>
<td>0*</td>
<td>0</td>
</tr>
<tr>
<td>Compound 24</td>
<td>0*</td>
<td>0</td>
</tr>
<tr>
<td>Compound 37</td>
<td>0**</td>
<td>0</td>
</tr>
<tr>
<td>Compound 54</td>
<td>0*</td>
<td>33</td>
</tr>
<tr>
<td>Untreated Control</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Thiabendazole</td>
<td>100</td>
<td>-</td>
</tr>
</tbody>
</table>

* Growth size stunted about 75 percent.
** Growth size stunted about 25 percent.
It will be understood that the parasite species employed in the above tests are merely representative of a wide variety of endoparasites and ectoparasites that can be controlled by the acyl urea compounds represented by the generic formula 1.

Although the invention has been illustrated by the preceding examples, it is not to be construed as being limited thereby; but rather, the invention encompasses the generic area as hereinbefore disclosed. Various modifications and embodiments can be made without departing from the spirit and scope thereof.
Claims

1. A method of controlling parasites of warm-blooded animals which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
&O \quad Y \\
R_1 - C - N - C - N - R_4 \\
&\quad R_2 \quad R_3
\end{align*}
\]

wherein:

- \( R_1 \) and \( R_4 \) are independently a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system, a bicyclic aromatic or nonaromatic ring system, a polycyclic aromatic or nonaromatic ring system, and a bridged ring system which may be saturated or unsaturated in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, poly haloalkenylthio, thiocyano, propargylthio, hydroxy imino, alkoxyimino, trialkylsilyloxy, aryldialkylsilyloxy, triarylsilyloxy, formamidino, alkylsulfamido, dialkylsulfoamido, alkoxy sulfonyl, poly haloalkoxy sulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts,
formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxycarbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfanyl, polyhaloalkylsulfanyl, alkylsulfonyl, polyhaloalkylsulfonyl, alkylsulfonamido, alkylcarbonylamino, polyhaloalkylsulfonamido, polyhaloalkylcarbonylamino, trialkylsilyl, arylalkylsilyl, triaryl silyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxycarbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynloxy, polyfluoralkanol, cyanalkylamino, semicarbazononemethyl, alkoxycarbonylhydrazononemethyl, alkoximinemethyl, unsubstituted or substituted arylaminocarbonyloxy, hydrazonemethyl, unsubstituted or substituted aryldiazonemethyl, a hydroxy group condensed with a mono-, di- or polysaccharide.

\[-X, = X, -X = R_5, = X-R_5,\]
\[\begin{array}{c}
Y_1 \\
\hline
Y_1
\end{array} \]
\[-X - R_5 , - P - Y_2R_6 , -Y_4 - P - Y_2R_6 \quad \text{or}\]
\[\begin{array}{c}
Y_3R_7 \\
Y_2R_6
\end{array} \]
\[\begin{array}{c}
Y_3R_7 \\
Y_2R_6
\end{array} \]
wherein $R_5$ is a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system, a bicyclic aromatic or nonaromatic ring system, a polycyclic aromatic or nonaromatic ring system, and a bridged ring system which may be saturated or unsaturated in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio, thiocyanato, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, arylalkylsilyloxy, triarylsilyloxy, formamidino, alkylsulfamido, dialkylsulfamido, alkoxy sulfonyl, polyhaloalkoxy sulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, nitro, cyano, hydroxy carbonyl and derivative salts, formamido, alky1, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkyl sulfinyl, polyhaloalkyl sulfinyl, alkyl sulfonyl, polyhaloalkyl sulfonyl, alkyl sulfonylamino, alkyl carbonylamino, polyhaloalkyl sulfonylamino, polyhaloalkyl carbonylamino, trialkylsilyl, arylalkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy,
dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonyl hydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5,\]

\[
\begin{array}{c}
Y_1 \\
\end{array}
\]

\[-X - R_5 , - P - Y_2 R_6 , -Y_4 - P - Y_2 R_6
\]

\[
\begin{array}{c}
Y_3 R_7 \\
Y_3 R_7
\end{array}
\]

or

\[
\begin{array}{c}
Y_2 R_6 \\
Y_2 R_6
\end{array}
\]

\[
\begin{array}{c}
Y_7 R_7 \\
Y_7 R_7
\end{array}
\]

Y_1 and Y_4 are independently oxygen or sulfur, Y_2 and Y_3 are independently oxygen, sulfur, amino or a covalent bond, R_6 and R_7 are independently hydrogen or substituted or unsubstituted alkyl, polyhaloalkyl, phenyl or benzyl in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonyl alkyl, alkoxy carbonyl alkyl thio, polyhaloalkenyl thio, thiocyanato, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, aryldialkylsilyloxy, triarylsilyloxy, foraminido, alkyl sulfamido, dialkyl sulfamido, alkoxy sulfonyl, polyhaloalkoxy sulfonyl, hydroxy, amino,
aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkenyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonyle, polyhaloalkylsulfonyle, alkylcarbonylamino, polyhaloalkylsulfonyle, polyhaloalkylcarbonylamino, trialkylsilyl, aryldialkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynyl, polyfluoroalkyl, cyanoalkyl amino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloximinomethyl, hydrazonomethyl, unsubstituted or substituted aryldihydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5,\]

\[-X - R_5, - P - Y_2R_6, -Y_4 - P - Y_2R_6\]

\[-Y_3R_7\]
or

\[ Y_2R_6 \quad \text{and} \quad Y_3R_7 \]

\[ X \]
is a covalent single bond or double bond, a substituted or unsubstituted heteroatom or substituted carbon atom, or a substituted or unsubstituted, branched or straight chain containing two or more carbon atoms or heteroatoms in any combination in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonyl alkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio, thiocyanato, propargylthio, hydroxy imino, alkoxyimino, trialkylsilyloxy, arylalkylsilyloxy, triarylsilyloxy, formamidino, alkylsulfamido, dialkylsulfamido, alkoxy sulfonylethyl, polyhaloalkoxysulfonylethyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamidino, alkyl, alkoxy, poly haloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or poly haloalkenyl; alkylthio, poly haloalkylthio, alkylsulfinyl, poly haloalkylsulfinyl, alkylsulfonylethyl, poly haloalkylsulfonyl, alkylsulfonylethyl, alkylcarbonylamino, poly haloalkylsulfonylethyl.
polyhaloalkylcarbonylamino, trialkysilyl, aryldialkysilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkeny1, alkenyloxy, alkynyl, alkynloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[ -X_1 = X, -X = R_5, = X - R_5, \]

\[ \begin{align*}
  & \quad Y_1 \\
  & \quad -X - R_5, -P - Y_2 R_6, -Y_4 - P - Y_2 R_6, \\
  \quad Y_3 R_7 & \quad Y_3 R_7 \\
  \text{or} \\
  \quad Y_2 R_6 & \quad Y_3 R_7 \\
  \quad Y_3 R_7 \\
\end{align*} \]

or \( R_4 \) is hydrogen;

\( Y \) is oxygen or sulfur; and

\( R_2 \) and \( R_3 \) are independently hydrogen, alkyl, substituted alkyl in which the permissible substituents are the same or different and are one or more halogen, alkoxy, alkylthio, or cyano; cycloalkyl, cycloalkeny1, substituted benzyl in which the permissible substituents are the same or different and are one or more halogen, hydroxy, nitro, cyano, alkyl, polyhaloalkyl, alkoxy or
polyhaloalkoxy; hydroxy, alkoxy, polyhaloalkoxy, alkylthio, polyhaloalkylthio, acyl, alkoxy carbonyl, alkoxythiocarbonyl, alkylsulfonyl, substituted phenylsulfonyl in which the permissible substituents are the same or different and are one or more halogen, nitro, cyano or polyhaloalkyl; substituted phenylthio in which the permissible substituents are the same or different and are one or more halogen, alkyl, nitro, cyano, polyhaloalkyl, or substituted or unsubstituted alkoxy carbonyl, alkenoxycarbonyl, alkynoxycarbonyl, cycloalkoxycarbonyl, phenoxy carbonyl, hydroxy carbonyl or the alkali metal salt or ammonium salt thereof, alkyl, alkoxy, alkylthio, cycloalkyl, cycloalkyloxy, cycloalkylthio, alkenyl, alkenylthio, alkanoyl, alkylsulfonyl, alkynyl, phenyl, phenoxy, phenylthio or amino; N-(alkylcarbonyl)-N-alkylaminothio, alkoxy carbonylthio, trialkylsilyl, dialkylarylsilyl, thiarylsilyl,

or \( R_2 \) and \( R_3 \) are independently

\[-X, \quad -X = R_5,\]

\[-X - R_5, \quad -P - Y_2 R_6, \quad -Y_4 - P - Y_2 R_6 \]

\[ Y_1 \]

\[ Y_1 \]
MICROCOPY RESOLUTION TEST CHART

DATE: J. BUREAU OF STANDARDS 1963-A
or R₂ and R₃ may be linked together to form a substituted or unsubstituted, heterocyclic ring system which may be saturated or unsaturated and in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonyl alkyl, alkoxy carbonyl alkyl thio, poly halo alk enyl thio, thiocyano, propargyl thio, hydroxy imino, alkoxy imino, trialkyl sily loxy, aryldialkyl silyloxy, triaryl silyloxy, for amidino, alkyl sulfamido, dialkyl sulfamido, alk oxysulfonyl, poly haloalk oxysulfonyl, hydroxy, amino, aminocarbonyl, alkylamidocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, poly halo alkyl, poly halo alk oxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxy alkyl, alkyl thio alkyl, alkyl, alkenyl, halo alkenyl or poly halo alkenyl; alkyl thio, poly halo alkyl thio, alkyl sulfinyl, poly halo alkyl sulfinyl, alkyl sulfon yl, poly halo alkyl sulfonyl, alkyl sulfon yl amino, alkyl carbonylamino, poly halo alkyl sulfonylamino, poly halo alkyl carbonylamino, trialkyl silyl, aryldialkyl silyl, triaryl silyl, sulfonic acid and
derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryl oxyiminomethyl, hydrazonomethyl, unsubstituted or substituted aryl hydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5,\]

\[
\begin{align*}
& Y_1 \quad Y_1 \\
& \quad Y_3R_7 \\
& \quad Y_3R_7 \\
& \quad Y_3R_7
\end{align*}
\]

2. The method of claim 1 wherein \( R_1, R_4 \) and \( R_5 \) are independently a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system having the formula
a bicyclic aromatic or nonaromatic ring system
having the formula selected from

\[
\begin{array}{c}
\text{A}_2 \text{B}_2 \text{A}_3 \\
\text{Z} \text{A}_3 \text{B}_3 \text{Z} \\
\end{array}
\]

and

\[
\begin{array}{c}
\text{A}_2 \text{B}_2 \text{A}_3 \\
\text{Z} \text{A}_3 \text{B}_3 \text{Z} \\
\end{array}
\]

a polycyclic aromatic or nonaromatic ring system
having the formula selected from

\[
\begin{array}{c}
\text{A}_4 \text{B}_4 \text{A}_5 \text{B}_5 \text{A}_6 \text{B}_6 \\
\text{Z} \text{Z} \text{Z} \\
\end{array}
\]

\[
\begin{array}{c}
\text{A}_4 \text{B}_4 \text{A}_5 \text{B}_5 \text{A}_6 \text{B}_6 \\
\text{Z} \text{Z} \text{Z} \\
\end{array}
\]

\[
\begin{array}{c}
\text{A}_4 \text{B}_4 \text{A}_5 \text{B}_5 \text{A}_6 \text{B}_6 \\
\text{Z} \text{Z} \text{Z} \\
\end{array}
\]
and a bridged ring system which may be saturated or unsaturated having the formula selected from

---

and
wherein:

- \( A_1 \) represents a ring-forming chain of atoms which together with \( B_1 \) forms a carbocyclic or heterocyclic ring system containing from 0 to 3 double bonds or from 0 to 2 triple bonds;
- \( B_1 \) represents a saturated or unsaturated carbon atom;
- \( A_2 \) and \( A_3 \) independently represent a ring-forming chain of atoms which together with \( B_2 \) and \( B_3 \) form a carbocyclic or heterocyclic ring system;
- \( B_2 \) and \( B_3 \) are independently a saturated or unsaturated carbon atom or a saturated nitrogen atom;
- \( A_4 \), \( A_5 \), \( A_6 \) and \( A_7 \) independently represent a ring-forming chain of atoms which together with \( B_4 \), \( B_5 \), \( B_6 \) and \( B_7 \) form a carbocyclic or heterocyclic ring system;
- \( B_4 \), \( B_5 \), \( B_6 \) and \( B_7 \) are independently a saturated or unsaturated carbon atom or a saturated nitrogen atom;
- \( A_8 \), \( A_9 \) and \( A_{10} \) independently represent a ring-forming chain of atoms which together with \( B_8 \), \( B_9 \), \( B_{10} \) and \( B_{11} \) form a carbocyclic or heterocyclic ring system;
- \( B_8 \), \( B_9 \) and \( B_{10} \) are independently a saturated or unsaturated carbon atom or a saturated nitrogen atom;
- \( B_{11} \) represents a saturated or unsaturated carbon atom, nitrogen atom or phosphorous atom;
- \( A_{11} \), \( A_{12} \) and \( A_{13} \) independently represent a ring-forming chain of atoms which
together with $B_{12}$ and $B_{13}$ form a carbocyclic or heterocyclic ring system;

$B_{12}$ and $B_{13}$ are independently a saturated carbon atom or a nitrogen atom; and

$Z$ is the same or different and is one or more of hydrogen, halogen, alkylcarbonyl,
alcoholcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio,
thiocyanate, propargylthio, hydroxylimino, alkoxyimino, trialkylsilyleoxy, arylalkylsilyleoxy,
triarylsilyleoxy, foraminido, alkylsulfamido, dialkylsulfamido, alkoxy sulfonyl,
polyhaloalkylsulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl,
dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl,
nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkylalkoxy, polyhaloalkyl,
polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkoxythioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkythio, polyhaloalkylthio, alkylsulfanyl, polyhaloalkylsulfanyl, alkylsulfonyleoxy, polyhaloalkylsulfonylamino, alkylcarbonylamino, polyhaloalkylsulfonyleoxy, polyhaloalkylcarbonylamino, trialkylsilyl, arylalkylsilyl, triaryl silyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkenyloxy, polyhaloalkenyloxy,
polyhaloalkynyl, polyhaloalkynyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arylydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5,\]

\[-X - R_5 , -P - Y_2 R_6 , -Y_4 - P - Y_2 R_6\]
\[Y_3 R_7 \quad Y_3 R_7\]

or

\[Y_2 R_6 \quad Y_3 R_7\]

wherein \( R_5, R_6, R_7, Y_1, Y_2, Y_3, Y_4 \) and \( X \) are as defined in claim 1.

3. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[O \quad Y\]
\[R_1 - C - N - C - N - R_4 - X - R_5\]
\[R_2 \quad R_3\]
wherein:

$R_1$ and $R_4$ are independently a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system, a bicyclic aromatic or nonaromatic ring system, a polycyclic aromatic or nonaromatic ring system, and a bridged ring system which may be saturated or unsaturated in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio, thiocyano, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, aryl dialkylsilyloxy, triarylsilyloxy, formamidino, alkyl sulfamido, dialkylsulfamido, alkoxy sulfonyl, polyhaloalkoxysulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxycarbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkyl sulfonyl, polyhaloalkylsulfonyl, alkyl sulfonlamino, alkyl carbonylamino, polyhaloalkylsulfonlamino, polyhaloalkylcarbonylamino, trialkylsilyl, aryl dialkylsilyl, triarylsilyl, sulfonic acid and
derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted arylhydrazonomethyl, unsubstituted or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

$$- P - Y_2 R_6, - Y_4 - P - Y_2 R_6$$

$$Y_3 R_7, Y_3 R_7$$

or

$$Y_2 R_6$$

$$Y_3 R_7$$

wherein $Y_1$ and $Y_4$ are independently oxygen or sulfur, $Y_2$ and $Y_3$ independently are oxygen, sulfur, amino or a convalent bond, and $R_6$ and $R_7$ are independently hydrogen or substituted or unsubstituted alkyl, polyhaloalkyl, phenyl or benzyl in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylo carbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio, thiocyano, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, aryldialkylsilyloxy, triarylsilyloxy, foramidino,
alkylsulfamido, dialkylsulfamido, alkoxy sulfonyl, polyhaloalkoxy sulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonyl, polyhaloalkylsulfonyl, alkylsulfonlamino, alkylcarbonylamino, polyhaloalkylsulfonlamino, polyhaloalkylcarbonylamino, trialkylsilyl, aryl dialkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalknyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonyl hydrazonomethyl, alkoxy iminomethyl, unsubstituted or substituted arloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted arly hydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

-X, = X, -X = R₅, = X-R₅,

-Y₁

-Y₂ R₆

-Y₃R₇

-Y₄

-Y₂ R₆

-Y₃ R₇
or

\[
\begin{align*}
Y_2R_6 & \\
Y_3R_7 \\
\end{align*}
\]

\(R_5\) is a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocyclic aromatic or nonaromatic ring system, a bicyclic aromatic or nonaromatic ring system, a polycyclic aromatic or nonaromatic ring system, and a bridged ring system which may be saturated or unsaturated in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkoxyalkylalkyl, alkoxyalkylalkylalkyl, alkoxyalkylalkylalkylthio, polyhaloalkenylthio, thiocyanato, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, aryldialkylsilyloxy, triarylalkylsilyloxy, foraminido, alkylsulfamido, dialkylsulfamido, alkoxyalkylsulfonyl, polyhaloalkoxysulfonyl, hydroxy, amino, alkoxyalkylalkyl, dialkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxyalkyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxyalkyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfamido, polyhaloalkylsulfonyl, alkylsulfamido, polyhaloalkylsulfonyl, alkoxyalkylsulfonyl, alkoxyalkylsulfonamido,
alkylcarbonylamino, polyhaloalkylsulfonylamino, polyhaloalkylcarbonylamino, trialkysilyl, arylalkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxyacylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynloxy, polyfluoroalkanol, cyberalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazononomethyl, alkoxyiminonomethyl, unsubstituted or substituted arylhydrazononomethyl, unsubstituted or substituted arylhydrazononomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[ -X, = X, \quad -X = R_5, = X-R_5, \]

\[ -X - R_5, \quad -P - Y_2R_6, \quad -Y_4 - P - Y_2R_6 \]

\[ \quad Y_3R_7 \quad Y_3R_7 \]

\[ \quad Y_2R_6 \quad Y_3R_7 \]

\[ X \text{ is a covalent single bond or double bond, a substituted or unsubstituted heteroatom or substituted carbon atom, or a substituted or unsubstituted, branched or straight chain containing two or more carbon atoms or heteroatoms in any combination in which the permissible substituents are the same or different and are one or more} \]
hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylothio, thiocyano, propargylthio, hydroxyimino, alkox yimino, trialkylsilyloxy, aryldialkylsilyloxy, triaryl silyloxy, formamido, alkylsulfamido, dialkylsulfamido, alkoxy sulfonyl, polyhaloalkoxy sulfonyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, poly haloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or poly haloalkenyl; alkylthio, polyhaloalkylthio, alkyl sulfinyl, polyhaloalkylsulfinyl, alkylsulfonyl, polyhaloalkylsulfonyl, alkylsulfonylamino, alkylcarbonylamino, polyhaloalkylsulfonylamino, polyhaloalkylcarbonylamino, trialkylsilyl, aryldialkylsilyl, triaryl silyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyl, alkynyl, polyhaloalkenyloxy, polyhaloalkynyl, polyfluoroalkanoyl, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted
or substituted arylhydrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5.\]

\[-X - R_5 , - P - Y_2 R_6 , -Y_4 - P - Y_2 R_6 \]

\[-Y_3 R_7 - Y_3 R_7 \]

or

\[-Y_2 R_6 \]

\[-Y_3 R_7 \]

\[Y\] is oxygen or sulfur; and

\[R_2\] and \[R_3\] are independently hydrogen, alkyl, substituted alkyl in which the permissible substituents are the same or different and are one or more halogen, alkoxy, alkylthio, or cyano; cycloalkyl, cycloalkenyl, substituted benzyl in which the permissible substituents are the same or different and are one or more halogen, hydroxy, nitro, cyano, alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy; hydroxy, alkoxy, polyhaloalkoxy, alkylthio, polyhaloalkylthio, acyl, alkoxyacarbonyl, alkoxythiocarbonyl, alkylsulfanyl, substituted phenylsulfonyl in which the permissible substituents are the same or different and are one or more halogen, nitro, cyano or polyhaloalkyl; substituted phenylthio in which the permissible substituents are the same or different and are one or more halogen, alkyl, nitro, cyano, polyhaloalkyl, or substituted or unsubstituted alkoxyacarbonyl, alkenoxyacarbonyl, alkynoxycarbonyl, cycloalkoxycarbonyl, phenoxyacarbonyl, hydroxycarbonyl or the alkali metal.
salt or ammonium salt thereof, alkyl, alkoxy, alkythio, cycloalkyl, cycloalkyloxy, cycloalkythio, alkenyl, alkenythio, alkanoyl, alkylsulfonyl, alkynyl, phenyl, phenoxy, phenylthio or amino; N-(alkylcarbonyl)-N-alkylaminothio, alkoxy carbonylthio, trialkylsilyl, dialkylalrylsilyl, thiarylsilyl,

\[
\begin{align*}
\text{Y}_1 & \quad \text{Y}_2 \quad \text{R}_6 \\
\text{Y}_3 & \quad \text{R}_7
\end{align*}
\]

or \( R_2 \) and \( R_3 \) may be linked together to form a substituted or unsubstituted, heterocyclic ring system which may be saturated or unsaturated and in which the permissible substituents are the same or different and are one or more hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, poly halo alkenylthio, thiocyanato, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, aryldialkylsilyloxy, triarylsilyloxy, formamidino, alkylsulfamido, dialkylsulfamido, alkoxy sulfonfyl, poly halo alkoxy sulfonfyl, hydroxy, amino, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbo
dialkylaminocarbo

alcohlythiocarbonyl, alkylnothyocarbonyl, 1, dialkylaminothyocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, poly halo alkoxo, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl,
alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkylthio, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonyl, polyhaloalkylsulfonyl, alkylsulfonylamino, alkylcarbonylamino, polyhaloalkylsulfonylamino, polyhaloalkylcarbonylamino, trialkylsilyl, arylalkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxycarbonylamino, alkylaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynyloxy, polyhaloalkenyloxy, polyhaloalkynyl, polyhaloalkynyloxy, polyfluoroalkanol, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonylhydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted aryloxyiminomethyl, hydrazonomethyl, unsubstituted or substituted aryldihyrazonomethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5,\]

\[-X - R_5, -P - Y_2R_6, -Y_4 - P - Y_2R_6\]

or

\[-Y_2R_6\]

\[-Y_2R_6, Y_3R_7\]

4. The method of claim 3 wherein \(R_1\), \(R_4\) and \(R_5\) are independently a substituted or unsubstituted, carbocyclic or heterocyclic ring system selected from a monocylic aromatic or nonaromatic ring system having the formula
a bicyclic aromatic or nonaromatic ring system having the formula selected from

\[ A_1 \]

\[ \begin{align*}
\ & B_1 \\
\ & Z
\end{align*} \]

and

\[ A_2 \]

\[ \begin{align*}
\ & B_2 \\
\ & Z
\end{align*} \]

a polycyclic aromatic or nonaromatic ring system having the formula selected from

\[ A_3 \]

\[ \begin{align*}
\ & B_3 \\
\ & Z
\end{align*} \]

\[ A_4 \]

\[ \begin{align*}
\ & B_4 \\
\ & Z
\end{align*} \]

\[ B_5 \]

\[ \begin{align*}
\ & B_5 \\
\ & Z
\end{align*} \]

\[ A_6 \]

\[ \begin{align*}
\ & B_6 \\
\ & Z
\end{align*} \]

\[ A_7 \]

\[ \begin{align*}
\ & B_7 \\
\ & Z
\end{align*} \]
and a bridged ring system which may be saturated or unsaturated having the formula selected from

A_{11} A_{12} A_{13}
A_{14} A_{15} A_{16}

wherein:

A_1 represents a ring-forming chain of atoms which together with B_1 forms a carbocyclic
or heterocyclic ring system containing from 0 to 3 double bonds or from 0 to 2 triple bonds;

$B_1$ represents a saturated or unsaturated carbon atom;

$A_2$ and $A_3$ independently represent a ring-forming chain of atoms which together with $B_2$ and $B_3$ form a carbocyclic or heterocyclic ring system;

$B_2$ and $B_3$ are independently a saturated or unsaturated carbon atom or a saturated nitrogen atom;

$A_4$, $A_5$, $A_6$ and $A_7$ independently represent a ring-forming chain of atoms which together with $B_4$, $B_5$, $B_6$ and $B_7$ form a carbocyclic or heterocyclic ring system;

$B_4$, $B_5$, $B_6$ and $B_7$ are independently a saturated or unsaturated carbon atom or a saturated nitrogen atom;

$A_8$, $A_9$ and $A_{10}$ independently represent a ring-forming chain of atoms which together with $B_8$, $B_9$, $B_{10}$ and $B_{11}$ form a carbocyclic or heterocyclic ring system;

$B_8$, $B_9$ and $B_{10}$ are independently a saturated or unsaturated carbon atom or a saturated nitrogen atom;

$B_{11}$ represents a saturated or unsaturated carbon atom, nitrogen atom or phosphorous atom;

$A_{11}$, $A_{12}$ and $A_{13}$ independently represent a ring-forming chain of atoms which together with $B_{12}$ and $B_{13}$ form a carbocyclic or heterocyclic ring system;

$B_{12}$ and $B_{13}$ are independently a saturated carbon atom or a nitrogen atom; and
Z is the same or different and is one or more of hydrogen, halogen, alkylcarbonyl, alkylcarbonylalkyl, alkoxyalkylalkyl, alkoxyalkylalkylthio, polyhaloalkenylthio, thiocyanato, propargylthio, hydroxyimino, alkoxyimino, trialkylsilyloxy, arylalkylsilyloxy, triarylsilyloxy, formamido, alkylsulfamido, dialkylsulfamido, alkoxy sulfonamide, polyhaloalkoxy sulfonamide, hydroxy, amino, aminocarbonyl, alkyaminocarbonyl, dialkyaminocarbonyl, aminothiocarbonyl, alkylaminothiocarbonyl, dialkylaminothiocarbonyl, nitro, cyano, hydroxycarbonyl and derivative salts, formamido, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, substituted amino in which the permissible substituents are the same or different and are one or two propargyl, alkoxyalkyl, alkylthioalkyl, alkyl, alkenyl, haloalkenyl or polyhaloalkenyl; alkylthio, polyhaloalkythio, alkylsulfanyl, polyhaloalkylsulfanyl, alkylsulfonamido, alkylaminosulfonamido, polyhaloalkylsulfonamido, polyhaloalkylsulfonamido, polyhaloalkylsulfonamido, trialkylsilyl, aryldialkylsilyl, triarylsilyl, sulfonic acid and derivative salts, phosphonic acid and derivative salts, alkoxy carbamino, alkyaminocarbonyloxy, dialkylaminocarbonyloxy, alkenyl, polyhaloalkenyl, alkenyloxy, alkynyl, alkynylthio, polyhaloalkenyloxy, polyhaloalkynyloxy, polyfluoroalkanoyl, cyanoalkylamino, semicarbazonomethyl, alkoxy carbonyl hydrazonomethyl, alkoxyiminomethyl, unsubstituted or substituted
aryloxyiminomethyl, hydrazonornethyl, unsubstituted or substituted aryihydrazonornethyl, a hydroxy group condensed with a mono-, di- or polysaccharide,

\[-X, = X, -X = R_5, = X-R_5,\]

\[-X - R_5 \leftrightarrow - P - Y_2 R_6 \quad -Y_4 - P - Y_2 R_6\]

\[Y_3 R_7\]

or

\[Y_2 R_6\]

\[Y_3 R_7\]

wherein \( R_5, R_6, R_7, Y_1, Y_2, Y_3, Y_4 \) and \( X \) are as defined in claim 3.

5. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

![Chemical Structure](image)

wherein:

- \( M \) represents a monocyclic aromatic ring system or a monocyclic heterocyclic ring system
containing up to 2 nitrogen atoms, and wherein the
rings can contain up to 2 nitrogen atoms, and
wherein the ring can contain up to four $X_1$
substituents wherein each $X_1$ individually can be
halogen, nitro, cyano, or alkyl, polyhaloalkyl,
alkoxy or polyhaloalkoxy of from 1 to 3 carbon atoms;
$Z_1$ and $Z_2$ individually are hydrogen or
alkyl of from 1 to 8 carbon atoms, or alkyl of from
1 to 8 carbon atoms which is substituted with at
least one of halogen, hydroxyl or alkoxy;
$Y_5$ is oxygen or sulfur;
$Y_6$ individually represents halogen, or
alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy of
from 1 to 3 carbon atoms, and $n'$ has a value of from
0 to 4; and

$B_{14}$ is a bicyclic fused ring system which
is attached to the oxygen through a carbocyclic ring
and wherein (a) at least one ring is a six-membered,
unsaturated carbocyclic ring which can contain up to
two $R_{16}$ and $R_{17}$ substituents wherein $R_{16}$ and
$R_{17}$ can be halogen, nitro, cyano, amino,
formamido, formamidino, phenylsulfenyl,
phenylsulfinyl, phenylsulfonyl, phenylsulfamido
wherein the phenyl ring optionally may be
substituted with one or more halogen, nitro, or
alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy of
from 1 to 3 carbon atoms, or $R_{16}$ and $R_{17}$
individually are alkyl, alkoxy, polyhaloalkyl,
polyhaloalkoxy, alkenyloxy, polyhaloalkenyloxy,
alkynyloxy, polyhaloalkynyloxy, alkenylsulfenyl,
polyhaloalkylsulfenyl, alkenylsulfinyl,
polyhaloalkylsulfinyl, alkylsulfenyl,
polyhaloalkylsulfonyl, mono- or di-alkylsulfamido, mono or di-alkylamino, alkylcarbonylamino, alkoxy carbonylamino, mono- or di-alkylamino-carbonyloxy of up to 6 carbon atoms, and (b) the second ring, hereinafter also referred to as A<sub>14</sub>, when it is not a five or six-membered carbocyclic can be a five or six-membered saturated or unsaturated heterocyclic ring which can contain in any combination carbonyl or one or two oxygen or sulfur, and up to two R<sub>18</sub> and R<sub>19</sub> substituents attached to unsaturated ring carbon atoms wherein R<sub>18</sub> and R<sub>19</sub> individually can be halogen, nitro, cyano, or alkyl, polyhaloalkyl, alkoxy, polyhaloalkoxy, alkylthio, arylthio, alkylsulfinyl, arylsulfinyl, alkyl- sulfonyl, arylsulfonyl, alkylsulfamido, arylsulfamido, alkoxy carbonylamino, or alkylcarbonylamino of from 1 to 6 carbon atoms, or can contain up to two R<sub>20</sub> and R<sub>21</sub> substituents attached to saturated carbon atoms of the chain wherein R<sub>20</sub> and R<sub>21</sub> individually can be alkyl or polyhaloalkyl of from 1 to 6 carbon atoms.

§. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein:

$X_2$ and $X_3$ are independently hydrogen, halogen, alkyl, haloalkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy;

$Z_{22}$ represents oxygen, azo, sulfur or sulfone;

$Y_7$ represents oxygen or sulfur;

$m'$ and $n'$ are independently 0 - 4;

$R_{22}$ independently represents hydrogen, halogen, alkyl, haloalkyl, polyhaloalkyl, alkoxy, nitro, hydroxycarbonyl, oxycarbonyl, alkenyl or alkynyl; and

$R_{23}$ independently represents hydrogen, halogen, alkyl, polyhaloalkyl, polyhaloalkoxy, alkylsulfonfonyl, alkoxy, alkylthio, dialkylamino, cyanop, nitro, haloalkylthio, trifluoromethylsulfonfonyl, alkenyl, alkynyl, thiocyano, alkylsulfinyl, arylthio, $CO_2R_{24}$, CONHR$_{24}$, arylsulfinyl or arylsulfonyl, wherein $R_{24}$ represents alkyl.

7. The method of controlling parasites of warm-blooded animals of claim 1 which comprises
orally or percutaneously administering to the animal a parasitically effective amount of a compound having the formula:

![Chemical Structure](image)

wherein:
- $X_4$ represents halogen;
- $X_5$ represents hydrogen or halogen;
- $X_6$ represents halogen or hydrogen;
- $Y_8$ is oxygen or sulfur;
- $R_{25}$, $R_{26}$ and $R_{27}$ are independently alkyl, halogen, polyhaloalkyl, alkoxy, nitro, cyano, alkylthio, polyhaloalkylthio or arylthio; and
- $R_{28}$, $R_{29}$, $R_{30}$ and $R_{31}$ are independently hydrogen, alkyl, halogen, nitro, alkoxy, polyhaloalkyl, polyhaloalkoxy, dialkylamino, or aryl.

8. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasitically effective amount of a compound having the formula:

![Chemical Structure](image)
wherein:

\[ X_7 \] and \[ X_8 \] are independently hydrogen, halogen, alkyl, haloalkyl, polyhaloalkyl, alkoxy, haloalkoxy or polyhaloalkoxy;

\[ R_{32} \] is independently hydrogen, halogen, alkyl, haloalkyl, polyhaloalkyl, alkoxy, haloalkoxy, polyhaloalkoxy, carboxylic acid, carboxylic acid salt or carboxylic acid ester;

\[ R_{33} \] is hydrogen, alkyl, polyhaloalkyl, hydroxy, amino or dialkylamino;

\[ R_{34} \] is independently hydrogen, halogen, alkyl, haloalkyl, polyhaloalkyl, alkoxy, haloalkoxy, polyhaloalkoxy, carboxylic acid, carboxylic acid salt, carboxylic acid ester, cyano or nitro;

\[ m' \] and \[ n' \] independently have a value of from 0 to 4, and \[ q'' \] has a value from 1 to 4; and

\[ Y_9 \] is oxygen or sulfur.

9. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
\text{[Formula Diagram]}
\end{align*}
\]

wherein:

\[ Y_{10} \] represents oxygen or sulfur;

\[ Z_3 \] represents oxygen, sulfur, sulfinyl, sulfonyl, unsubstituted or substituted amino in
which the permissible substituents are one or more hydrogen, alkyl, acyl or alkenyl:

\[ X_9 \text{ and } X_{10} \text{ are independently hydrogen, halogen, trifluoromethyl, alkyl or polyhaloalkyl;} \]

\[ R_{35} \text{ independently represents hydrogen, halogen, trifluoromethyl, alkyl, alkoxy, nitro, alkenyl, hydroxycarbonyl, alkoxy carbonyl or alkyl carbonyl;} \]

\[ R_{36} \text{ independently represents hydrogen, halogen, alkyl, cyano, nitro, trifluoromethyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, dialkylamino, alkylthio, polyhaloalkylthio, arylthio, or alkenyl optionally substituted with halogen; and} \]

\[ m' \text{ and } n' \text{ independently have a value of from 0 to 4, and } q'' \text{ has a value of from 1 to 4.} \]

10. The method of controlling parasites of warm-blooded animals of claim 1 which comprise orally or percutaneously administering to the animal a parasitically effective amount of a compound having the formula:

\[
\begin{align*}
\text{Y}_{11} & \text{ is oxygen or sulfur;} \\
\text{Y}_{12} & \text{ is CR}_{37} \text{ or nitrogen;} \\
\text{B}_{15} & \text{ is oxygen, sulfur, sulfinyl, sulfonyl, oxysulfonyl, sulfonyloxy or carbonyl;} \\
\end{align*}
\]

\[ (R_{38})_{n'} \text{ and } (R_{39})_{q} \]
X_{11} and X_{12} are independently hydrogen, halogen, alkoxy, alkyl, trifluoromethyl or polyhaloalkyl;

R_{37} and R_{38} are independently alkoxy, hydrogen, halogen, alkyl, trifluoromethyl, polyhaloalkyl or polyhaloalkoxy;

R_{39} is independently hydrogen, halogen, alkyl, trifluoromethyl, polyhaloalkyl, polyhaloalkoxy, alkoxy, alkylthio, alkylsulfinyl, alkylsulfonyl, or R_{39} can optionally be a substituted arylthio, a substituted arylsulfonyl, or a substituted arylsulfinyl, wherein said optional substituents are alkyl or halogen, and

m', n' and q independently have a value of from 0 to 4, and q" has a value of from 1 to 4.

11. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
X_{13} & \quad \text{is halogen, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy or nitro;} \\
X_{14} & \quad \text{is hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy or nitro;} \\
R_{40} & \quad \text{is hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, nitro, cyano,}
\end{align*}
\]
carboxylic acid, carboxylic acid salt or carboxylic acid ester;
m' has a value of from 0 to 4;
Y_{13} is oxygen or sulfur;
Y_{14} is alkylene or a covalent bond;
Z_{4} is substituted or unsubstituted

wherein the permissible substituents are one or more halo, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, nitro, cyano, carboxylic acid, carboxylic acid salt or carboxylic acid ester substituents which may be the same or different; and q'' has a value of from 1 to 4.

12. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasitcidally effective amount of a compound having the formula:

\[ \text{M1} \]

\[ \text{N} - \text{S} - \text{E} - \text{B}_{16} \]

wherein:

M_{1} represents a monocyclic aromatic ring system or a monocyclic heterocyclic ring system
containing up to two nitrogen atoms and \(M_1\) can contain up to four \(X_{15}\) substituents, each of which individually can be halogen, nitro, cyano or alkyl, polyhaloalkyl, alkoxy, or polyhaloalkoxy of from one to three carbon atoms;

\(Z_5\) and \(Z_6\) individually are hydrogen, alkyl or alkyl which may be substituted by at least one halogen, hydroxy or alkoxy;

\(E\) is a six membered carbocyclic aromatic ring or a five or six membered heterocyclic ring containing up to two oxygen, sulfur or nitrogen atoms or a combination thereof, and wherein this ring may contain up to four \(Y_{15}\) substituents wherein each \(Y_{15}\) individually can be halogen, cyano or alkyl, polyhaloalkyl, alkoxy, polyhaloalkyloxy, alkylsulfenyl, or polyhaloalkylsulfenyl of from one to six carbon atoms;

\(B_{16}\) is a bicyclic fused ring system which is attached to the oxygen through a carbocyclic ring and wherein (a) at least one ring is six-membered, unsaturated and carbocyclic and which contains the substituents \(R_{41}\) and \(R_{42}\) (b) the second ring; hereinafter also referred to as a \(A_{15}\), when it is not a five or six-membered saturated or unsaturated carbocyclic can be a five or six-membered saturated or unsaturated heterocyclic which can contain in any combination a carbonyl group or one or two oxygen or sulfur atoms and the substituents \(R_{43}\) and \(R_{44}\) attached to unsaturated ring carbons or the substituents \(R_{45}\) and \(R_{46}\) attached to saturated ring carbons;
R\textsubscript{41} and R\textsubscript{42} individually may be hydrogen, halogen, nitro, cyano, amino, formamido, formamidino, phenylsulfenyl, phenylsulfinyl, phenylsulfonyl, phenylsulfamido or phenyloxysulfonyl, wherein the phenyl ring optionally may contain one or more substituents, selected from halogen, nitro, or alkyl, polyhaloalkyl, alkoxy, or polyhaloalkoxy of from one to three carbon atoms, or R\textsubscript{41} and R\textsubscript{42} individually may be alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkenyloxy, polyhaloalkenyloxy, alkynyloxy, polyhaloalkynyloxy, alkylsulfenyl, alkylsulfonyl, alkylsulfenyl, polyhaloalkylsulfenyl, alkylsulfinyl, mono- or di-alkylsulfamido, mono- or di-alkylamino, alkylcarbonylamino, alkoxycarbonylamino, mono- or di-alkylaminocarbonyloxy, alkoxysulfonyl, polyhaloalkylsulfonyl of up to six carbons in each alkyl chain, morphilinosulfonyl or the group SO\textsubscript{3}Na;

R\textsubscript{43} and R\textsubscript{44} can individually be hydrogen, halogen, nitro, cyano, or alkyl, polyhaloalkyl, alkoxy, polyhaloalkoxy, alkylsulfinyl, alkylsulfenyl, alkylsulfonyl, alkoxycarbonylamino or alkylcarbonylamino of from one to eight carbons; and

R\textsubscript{45} and R\textsubscript{46} individually can be hydrogen, alkyl or polyhaloalkyl from one to six carbon atoms or phenyl which may be optionally substituted with up to three halogen, cyano, nitro or alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy of up to three carbon atoms.
13. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
&\text{wherein } M_2 \text{ represents a monocyclic aromatic ring system or a monocyclic heterocyclic ring system containing up to four } X_{16} \text{ substituents, each } X_{16} \text{ substituent is selected from the group of halogen, nitro, cyano or alkyl, polyhaloalkyl, alkoxy, polyhaloalkoxy, alkylsulfenyl or polyhaloalkylsulfenyl of from one to three carbon atoms;} \\
&Z_7 \text{ and } Z_8 \text{ individually are hydrogen, alkyl, or alkoxy, either or which may be substituted by at least one halogen, hydroxy or alkoxy;} \\
&Y_{17} \text{ is oxygen or sulfur;} \\
&E_2 \text{ is a five or six member heterocyclic ring containing up to two oxygen, sulfur or nitrogen atoms or a combination thereof, and wherein said ring may contain up to three } Y_{16} \text{ substituents, each } Y_{16} \text{ is selected from the group of halogen, cyano, or alkyl, polyhaloalkyl, alkoxy, polyhaloalkoxy, alkylsulfenyl or polyhaloalkylsulfenyl of from one to three carbon atoms;} \\
&B_{17} \text{ is a bicyclic fused ring system which is attached to the oxygen through a carbocyclic ring and wherein (a) at least one ring is six-membered,}
\end{align*}
\]
unsaturated and carbocyclic and which contains the substituents \( R_{47} \) and \( R_{48} \), (b) the second ring, \( R_{46} \), when it is not a five or six-membered saturated or unsaturated carbocyclic can be a five or six-membered saturated or unsaturated heterocyclic ring which can contain in any combination carbonyl or one or two oxygen or sulfur atoms and the substituents \( R_{49} \) and \( R_{50} \) attached to unsaturated ring carbons or the substituents \( R_{51} \) and \( R_{52} \) attached to the saturated ring carbon atoms, wherein \( R_{47} \) and \( R_{48} \) are selected from the group of hydrogen, halogen, nitro, cyano, amino, formamido, phenylsulfenyl, phenylsulfinyl, phenylsulfonyl, phenylsulfonate, phenylsulfamido, wherein the phenyl ring optionally may be substituted with one or more halogen, nitro, or alkyl, polyhaloalkyl, alkoxy, or polyhaloalkoxy of from one to three carbon atoms, or \( R_{47} \) and \( R_{48} \) individually may be alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkenyloxy, polyhaloalkenyloxy, alkynyloxy, polyhaloalkynyoxy, alkylsulfenyl, polyhaloalkylsulfenyl, alkylsulfinyl, polyhaloalkylsulfinyl, alkylsulfonate, polyhaloalkylsulfonate, polyhaloalkylsulfamido, mono- or dialkylsulfamido, mono- or di-alkylaminocarbonyloxy, alkylsulfonate, or polyhaloalkylsulfate of up to six carbon atoms;

\( R_{49} \) and \( R_{50} \) are selected from the group of hydrogen, halogen, nitro, cyano, or alkyl, polyhaloalkyl, alkoxy, polyhaloalkoxy, alkylthio, arylthio, alkylsulfenyl, arylsulfinyl, alkylsulfonyl, arylsulfonyl, alkylsulfamido, arylysulfamido, alkoxy carbonylamino or
alkylcarbonylamino of from one to eight carbon atoms; and

R₁ and R₂ are selected from the group of alkyl or polyhaloalkyl of from one to six carbon atoms.

14. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{array}{c}
\text{X}^{17} \quad \text{X}^{18} \\
\text{CONC-N} - \text{R}^{55} \\
\text{Y}^{53} \text{R}^{54}
\end{array}
\]

\[
\text{(X}^{18})^{q^*}
\]

wherein:

q" is a value of from 1 to 4;

X₁ and X₁₈ are independently hydrogen, halogen, trifluoromethyl, alkyl, alkoxy, polyhaloalkyl or polyhaloalkoxy;

Y₁₈ is oxygen or sulfur;

R₅₅ is a substituted or unsubstituted heterocyclic system selected from

\[
\begin{array}{c}
\text{Y}^{19}, \\
\text{X}^{19}, \\
\text{X}^{19}, \\
\text{Y}^{19}, \\
\text{Y}^{19}, \\
\text{Y}^{19}, \\
\text{Y}^{19}
\end{array}
\]
wherein $Y_{19}$ is the same or different and is one or more hydrogen, halogen, cyano, nitro, alkyl, branched alkyl, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, or an optionally substituted cycloalkyl, phenyl, biphenyl, benzyl, phenoxy, phenylthio, phenoxy benzyl, phenoxyphenyl, phenylthiobenzyl, phenylthiophenyl, phenylsulfinyl benzyl, phenylsulfinylphenyl, phenylsulfonyl benzyl, phenylsulfonylphenyl, naphthoxy, tetrahydronaphthoxy or naphthyl in which the permissible substituents are the same or different and are one or more halogen, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkyl, nitro or cyano; and

$R_{53}$ and $R_{54}$ are independently hydrogen, alkyl, acyl, alkoxy carbonyl or, when $Y_{18}$ is oxygen, $R_{53}$ and $R_{54}$ may be linked together to form a ring selected from

15. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein:

\[ X_{19}, X_{20}, X_{21}, X_{22} \text{ and } X_{23} \text{ are independently halogen, hydrogen, alkyl, alkoxy, nitro, cyano or polyhaloalkyl;} \]

\[ Y_{20} \text{ is oxygen or sulfur;} \]

\[ Y_{21}, Y_{22}, Y_{23} \text{ and } Y_{24} \text{ are independently hydrogen, halogen, trialkylsilyl, nitro, cyano, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, alkoxy carbonyl, alkylthio, substituted or unsubstituted cycloalkylalkoxy wherein the permissible substituents are the same or different and are one or more hydrogen, alkyl, aryl, haloaryl, alkylaryl or halogen; alkynyl, alkylcarbonyl, alkoxy carbonylalkyl, alkoxy carbonylalkylthio, polyhaloalkenylthio; a group selected from alkoxy, alkylthio, alkylsulfonyl, alkenyloxy or alkylsulfiny1, each of which may be substituted with one or more halogen atoms which may be the same or different; substituted or unsubstituted cycloalkoxy wherein the permissible substituents are the same or different and are one or more halogen, hydrogen, alkyl or aryl; polyfluoroalkanol, optionally substituted phenyl or \(-X-R_5\) wherein X and R_5 are as defined in claim 1, or \(Y_{21}, Y_{22}, Y_{23} \text{ and } Y_{24}\) may form, together with an adjacent group selected from \(Y_{21}', Y_{22}', Y_{23} \text{ and } Y_{24}'\) an alkylenedioxy group which may be substituted with one or more halogen atoms; when \(Y_{20}\) is oxygen, \(R_{56}\) and \(R_{57}\) are independently hydrogen, alkyl, substituted alkyl in which the permissible substituents are the same or different and are one or more halogen, alkoxy, alkylthio or cyano; substituted benzyl in which the
permissible substituents are the same or different and are one or more halogen, hydroxy, nitro, cyano, alkyl, polyhaloalkyl, alkoxy or polyhaloalkoxy; hydroxy, alkoxy, polyhaloalkoxy, alkylthio, polyhaloalkylthio, acyl, alkoxy carbonyl, alkysulfonyl, substituted phenylsulfonyl in which the permissible substituents are the same or different and are one or more halogen, nitro, cyano or alkyl; N-alkylcarbonyl-N-alkylaminothio,

\[
\begin{align*}
  \text{Y}_1 \quad & \quad \text{P} \quad \text{Y}_2 \quad \text{R}_6 \\
  \text{Y}_3 \quad & \quad \text{R}_7 
\end{align*}
\]

wherein \( \text{Y}_1, \text{Y}_2, \text{Y}_3, \text{R}_6 \) and \( \text{R}_7 \) are as defined in claim 1; or \( \text{R}_{56} \) and \( \text{R}_{57} \) may be linked together to form a heterocycle selected from

\[\text{Chemical Structures}\]
wherein \( R_{58} \) and \( R_{59} \) are independently hydrogen, halogen, substituted or unsubstituted alkyl, polyhaloalkyl, phenyl or benzyl; and when \( Y_{20} \) is sulfur, \( R_{56} \) is hydrogen and \( R_{57} \) is hydrogen or alkyl.

16. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
(x_{24})^{m'}(Y_{25})^{n'}
\]

wherein:

- \( X_{24} \) is halogen, hydrogen, alkyl or alkoxy;
- \( m' \) is a value of 1 to 5;
- \( Y_{25} \) is halogen, alkyl or trifluoromethyl;
- \( n' \) is a value of from 0 to 4;
- \( Z_9 \) is halogen,

wherein \( A_{17} \) and \( B_{18} \) are independently hydrogen, halogen, trifluoromethyl, cyano or nitro.
17. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
\text{C}_6 &\text{H}_4 - \text{N} - \text{C} - \text{N} - \text{S} \\
\text{C}_6 &\text{H}_4 - \text{CO}_2 \text{R}_{61}
\end{align*}
\]

wherein:

- \(X_{25}\) and \(X_{26}\) are independently hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, nitro or cyano;
- \(q\) is a value of from 1 to 4;
- \(R_{61}\) is substituted or unsubstituted alkyl, alkenyl, alkynyl, cycloalkyl or phenyl; a hydrogen atom or 1 equivalent of an alkali metal or optionally substituted ammonium cation; each \(X_{27}\) is independently halogen, cyano, nitro, hydroxycarbonyl or optionally substituted alkyl, alkoxy, alkylthio, cycloalkyl, cycloalkyloxy, cycloalkylthio, alkenyl, alkenylthio, alkanoyl, alkoxy carbonyl, alkylsulphonyl, alkynyl, phenyl, phenoxy, phenylthio or amino;
q is a value of from 0 to 4;

R₆₀ is halogen, alkyl, polynhaloalkyl, alkoxy, polyhaloalkoxy, alkenyloxy, polyhaloalkenyl, nitro, cyano or -X-R₅ wherein X and R₅ are as defined in claim 1; each Y₂₆ is independently halogen, nitro, cyano, alkyl, or polynhaloalkyl; and

n' is a value of from 0 to 4.

18. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
(x_{28})^n & - \begin{array}{c}
\text{N} \\
\text{H}
\end{array} - \begin{array}{c}
\text{O} \\
\text{K}
\end{array} - \begin{array}{c}
\text{N} \\
\text{H}
\end{array} - \begin{array}{c}
\text{C} \\
\text{H}
\end{array} - (x_{27} \cdot h') \\
\end{align*}
\]

and the acid addition salt thereof, wherein:

X₂₈ is independently halogen, hydrogen, alkoxy, alkyl, polyhaloalkoxy, polyhaloalkyl, nitro or cyano;

n" is a value of from 0 to 3, and n' is a value of from 0 to 4; and

Y₂₇ and Y₂₈ are independently hydrogen, halogen, alkoxy, alkyl, polyhaloalkoxy, polyhaloalkyl, nitro, cyano, alkenyl, alkenyloxy, polyhaloalkenyl, or polyhaloalkenyloxy.
19. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

\[
\begin{align*}
\text{(X}_{29}\text{)}^q \\
\text{R} \\
\text{N} \\
\text{N} \\
\text{X}_{29} \text{h'} \\
\end{align*}
\]

wherein:

- \(X_{29}\) and \(X_{30}\) are independently hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, nitro, or polyhaloalkoxy;
- \(X_{47}\) is halogen, hydrogen, hydroxy, alkoxy, or polyhaloalkoxy;
- \(Y_{30}\) is halogen, alkyl, polyhaloalkyl, alkoxy, polyhaloalkoxy, alkenyloxy, polyhaloalkenyloxy, nitro, cyano, or \(-X-R_5\) wherein \(X\) and \(R_5\) are as defined in claim 1;
- \(Y_{29}\) is independently halogen, nitro, cyano, alkyl, or polyhaloalkyl; and
- \(q'\) is a value of from 1 to 4 and \(n'\) is a value of from 0 to 4.

20. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein:

$X_{31}$ and $X_{32}$ are independently hydrogen, halogen, alkyl, alk'oxy, polyhaloalkyl, nitro or polyhaloalkoxy;

$Y_{31}$ is independently halogen, alkyl, alkoxy, polyhaloalkyl or polyhaloalkoxy;

$n$ is a value of from 0 to 2, and $q''$ is a value of from 1 to 4;

$Z_{11}$ is hydrogen, halogen or alkyl; and

$Z_{10}$ is halogen, alkyl optionally substituted with fluorine or chlorine; nitro or cyano; and wherein the pyridyloxy group is linked to the phenyl ring in the 3- or 4- position.

21. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

wherein:

$q''$ is a value of from 1 to 4;
X_{33} and X_{34} are independently hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, nitro or polyhaloalkoxy;

Y_{32} and Y_{33} are independently hydrogen, halogen, alkyl or polyhaloalkyl;

Z_{12} is hydrogen, alkyl, propargyl, alkenyl optionally substituted with chlorine, bromine, cycloalkyl or alkoxyalkyl; phenyl, naphthyl or tetrahydronaphthyl, all of which may be optionally substituted with halogen, alkyl, alkoxy, cyano, nitro, polyhaloalkyl or polyhaloalkoxy;

Z_{13} is alkyl, propargyl, alkenyl optionally substituted with halogen, cycloalkyl or alkoxyalkyl; and

wherein Z_{12} and Z_{13} can be combined to form a heterocyclic ring selected from pyrrol, piperidine or pyridone.

22. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasitically effective amount of a compound having the formula:
wherein:

\[ X_{36} \] is CH or nitrogen or oxide derivatives thereof;

\[ X_{35} \] is independently hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, nitro or polyhaloalkoxy;

\[ Y_{34} \] is oxygen or sulfur;

\[ R_{62} \] is hydrogen, hydroxy or alkyl;

\[ Y_{35} \] and \[ Y_{36} \] are independently halogen, alkyl, alkoxy, polyhaloalkyl or polyhaloalkoxy;

\[ A_{18}, A_{19}, A_{20} \] and \[ A_{21} \] are independently hydrogen, alkyl, alkenyl, polyhaloalkyl or halogener;

\[ m^{\prime\prime\prime} \] is a value of from 0 to 3, \[ n^{\prime\prime} \] is a value of from 1 to 3, and \[ n^{\prime} \] is a value of from 0 to 3 provided that either of \[ m^{\prime\prime\prime} \] and \[ n^{\prime} \] has a value of at least 1;

\[ B_{19} \] is CH or nitrogen or oxide derivatives thereof;

\[ Z_{15} \] is hydrogen, halogen, alkyl, haloalkyl, nitro, cyano, trifluoromethyl, polyhaloalkyl or polyhaloalkoxy; and

\[ Z_{14} \] is oxygen, sulfur, sulfoxide, sulfone, substituted or unsubstituted amino or a covalent bond.

23. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein:

- $X_{37}$ represents independently hydrogen, halogen, alkyl, polyhaloalkyl, polyhaloalkoxy, alkoxy, nitro or alkylthio;
- $q'$ is a value of from 1 to 5;
- $Y_{37}$ is oxygen or sulfur;
- $Y_{38}$ and $Y_{39}$ are independently hydrogen, halogen, alkyl, alkoxy, polyhaloalkoxy or polyhaloalkyl; and
- $Z_{16}$ is a heterocycle-containing moiety having the formula

![Chemical Structure]

wherein $A_{22}$ is CH or nitrogen, and $Z_{17}$ and $Z_{18}$ are independently hydrogen, halogen, trifluoromethyl, nitro, alkyl, cycloalkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy or phenyl optionally substituted with cyano, nitro, halogen, alkyl, alkoxy, polyhaloalkyl or polyhaloalkoxy; provided that $Z_{16}$ is attached at the 3- or 4- position.

24. The method of controlling parasites of warm-blooded animals of claim 1 which comprises
orally or percutaneously administering to the animal
a parasiticidally effective amount of a compound
having the formula:

\[
\begin{align*}
\text{wherein:} \\
X_{38} \text{ and } X_{39} \text{ are independently hydrogen, halogen, alkyl, alkoxy, poly haloalkyl, poly haloalkoxy or nitro;} \\
Y_{40} \text{ is oxygen or sulfur;} \\
Y_{41} \text{ is halogen, alkyl, cyano, poly haloalkyl, alkoxy or poly haloalkoxy;} \\
m''' \text{ is a value of from 0 to 2;} \\
n''' \text{ is a value of 0 or 1;} \\
A_{23} \text{ is a covalent bond, oxygen, sulfoxide, sulfone, oxyalkyl, thioalkylene, sulfinylalkylene, sulfonylalkylene, or sulfur;} \\
Z_{19} \text{ is halogen, trifluoromethyl, poly haloalkyl, poly haloalkoxy, poly haloalkythio, alkyl, nitro, cyano, alkoxy or alkylthio; and} \\
q \text{ is a value of from 0 to 4, and } q'' \text{ is a value of from 1 to 4.}
\end{align*}
\]

25. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein:

$q' \in \{1, 2, \ldots, 5\}$

$R_{63}$ is an unsubstituted or substituted heterocyclic aromatic ring system selected from

in which the permissible substituents are the same or different and are one or more halogen, hydrogen, polyhaloalkyl, alkoxy, polyhaloalkoxy or alkyl groups;

$Y_{42}$ is oxygen or sulfur;

$Y_{43}$ is independently selected from hydrogen, halogen, alkyl, polyhaloalkyl, phenyl optionally substituted with one or more hydrogen, halogen, alkyl, cyano, nitro or polyhaloalkyl; phenoxy, naphthoxy, pyridyloxy or tetrahydronaphthoxy, all optionally substituted by one or more halogen, alkyl, cyano, nitro, polyhaloalkyl, polyfluoroalkanol, alkoxy or polyhaloalkoxy; polyfluoroalkanol, alkoxy, polyhaloalkoxy, alkylcarbonyl, alkoxy carbonyl, alkylthio or dialkylamino.
26. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

and the acid addition salt thereof, wherein:

- \( q'' \) is a value of from 1 to 4, and \( q \) is a value of from 0 to 4;
- \( X_{40} \) and \( X_{41} \) are independently hydrogen, halogen, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy or nitro;
- \( Y_{44} \) is oxygen or sulfur;
- \( Y_{45} \) is the same or different and is one or more substituents selected from hydrogen, alkynyl, halogen, polyhaloalkyl, polyhaloalkoxy, trifluoromethyl, cyano, nitro, alkyl or alkoxy; and
- \( n'''' \) is a value of 0 or 1.

27. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:
wherein:

- $R_{64}$ and $R_{65}$ are independently halogen, hydrogen, alkyl, alkoxy, polyhaloalkyl, polyhaloalkoxy, or nitro;
- $Y_{46}$ is oxygen or sulfur;
- $R_{66}$ is halogen, alkyl, trifluoromethyl, polyhaloalkyl, polyhaloalkoxy or alkoxy;
- $m''$ is a value of from 0 to 3 and $q''$ is a value of from 1 to 4;
- $R_{68}$ is alkyl, cycloalkyl, hydrogen, alkylthio, cyano or alkoxy;
- $R_{67}$ is hydrogen or alkyl; and
- $A_{24}$ is alkyl, alkylthio or substituted alkyl with alkoxy, alkylthio, alkylsulfinyl or alkylsulfonyl or a ring system selected from

\[
\begin{align*}
\text{Z}_{20} & \quad \text{H}\text{on}^\prime
\end{align*}
\]

wherein $Z_{20}$ is hydrogen, halogen, alkyl, alkoxy, haloalkyl or haloalkoxy, and $n'$ is a value from 0 to 4; $A_{24}$ and $R_{68}$ can optionally be joined to form a ring system of from 5 to 7 atoms which may contain up to two oxygen, sulfur or nitrogen atoms and which may be optionally substituted with alkyl or halogen.

28. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasitically effective amount of a compound having the formula:
wherein:

$X_{43}$ is CH or nitrogen;

$X_{42}$ is independently halogen, alkyl or nitro;

$m^{**}$ is a value of from 0 to 3;

$Y_{47}$ and $Y_{48}$ are independently oxygen or sulfur;

$Y_{49}$ is hydrogen, hydroxy or optionally substituted alkyl;

$R_{70}$ is unsubstituted or substituted phenylene or heteroarylene with one or two nitrogen atoms in which the permissible substituents are one or more halogen, optionally halogenated alkyl or alkoxy;

$R_{71}$ is alkylene, alkenylene, alkylidene, cycloalkylene or cycloalkylidene optionally substituted with one or more halogen, alkyl which may be optionally substituted by halogen or hydroxy, phenyl, cyano, alkoxy carbonyl, hydroxy imino, alkoxy imino, alkenyl or alkynyl;
$R_{72}$ is unsubstituted or substituted phenyl or heteroaryl having from 1 to 3 nitrogen, oxygen or sulfur atoms in any combination which may be the same or different in which the permissible substituents are one or more halogen, nitro, cyano, optionally halogen-substituted alkyl, alkenyl, alkoxy, alkenyloxy, alkylthio, or alkylsulfonyl; and

$Z_{21}$ is oxygen, sulfur, sulfoxide, sulfone, amino or alkylimino, or

$Z_{21}$ and $R_{71}$ are together alkylideneamino.

29. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

![Chemical Structure](image)

wherein:

- $X_{44}$ is halogen;
- $X_{45}$ and $X_{46}$ are independently hydrogen or halogen provided that when $X_{45}$ is halogen then
$X_{46}$ is hydrogen, and when $X_{46}$ is halogen then
$X_{45}$ is hydrogen;
$Y_{49}$ is oxygen or sulfur;
$Y_{50}$ and $Y_{53}$ are independently hydrogen, halogen, optionally substituted alkyl, alkoxy, or alkylthio;
$Y_{51}$ is hydrogen, halogen, cyano, nitro or optionally substituted alkyl, alkylthio, alkylsulfonyl, alkoxy, naphthoxy, phenoxy, alkylcarbonyl, alkoxy carbonyl, alkoxy carbonylalkyl or alkoxy carbonylalkylthio, or $Y_{50}$ and $Y_{51}$ are together optionally substituted alkylenedioxy; and
$Y_{52}$ is hydrogen, halogen or optionally substituted alkyl, alkoxy or aryloxy.

30. The method of controlling parasites of warm-blooded animals of claim 1 which comprises orally or percutaneously administering to the animal a parasiticidally effective amount of a compound having the formula:

$$R_1 - X - R_4$$

wherein $R_1$, $R_4$ and $X$ are as defined in claim 1.

31. The method of claim 1 in which the compound is 1-[4-(4-chloro-1-naphthoxy)-3,5-dichloro-phenyl]-3-(2,6-difluorobenzoyl)urea.

32. The method of claim 1 in which the compound is 1-[4-(4-chloro-1-naphthoxy)-3-methyl-phenyl]-3-(2,6-difluorobenzoyl)urea.

33. The method of claim 1 in which the compound is 1-[4-(4-chloro-5,6,7,8-tetrahydro-1-
34. The method of claim 1 in which the compound is 1-[4-(4-chloro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea.

35. The method of claim 1 in which the compound is 1-[4-(4-dimethylamino-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea.

36. The method of claim 1 in which the compound is 1-[4-(2-bromo-4-chlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea.

37. The method of claim 1 in which the compound is 1-[4-(2,4-dimethylphenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea.

38. The method of claim 1 in which the compound is 1-[4-(2,4-dichlorophenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea.

39. The method of claim 1 in which the compound is 1-[4-(2-methyl-4-t-butylphenoxy)-2,5-dimethyl-3-chlorophenyl]-3-(2,6-difluorobenzoyl)urea.

40. The method of claim 1 in which the compound is 1-[2,5-dichloro-4-(alpha-cyano-4-fluorobenzyl)phenyl]-3-(2,6-difluorobenzoyl)urea.

41. The method of claim 1 in which the compound is 1-[3-chloro-4-(4-chloro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2-chlorobenzoyl)urea.
42. The method of claim 1 in which the compound is 1-[4-(4-chloro-1-naphthoxy)-2,3,5-trichlorophenyl]-3-(2,6-difluorobenzoyl)urea.

43. The method of claim 1 in which the compound is 1-[4-(4-chloro-1-naphthoxy)-3,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea.

44. The method of claim 1 in which the compound is 1-[3-chloro-4-(4-chloro-5,6,7,8-tetrahydro-1-naphthoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea.

45. The method of claim 1 in which the compound is 1-[3-chloro-4-(4-chloro-2,5-dimethylphenoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea.

46. The method of claim 1 in which the compound is 1-[4-(4-nitrophenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea.

47. The method of claim 1 in which the compound is 1-[4-(4-chlorophenoxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea.

48. The method of claim 1 in which the compound is 1-[4-(5-trifluoromethyl-3-chloro-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea.

49. The method of claim 1 in which the compound is 1-[3-chloro-4-(2,3-dimethyl-4-bromophenoxy)-2,5-dimethylphenyl]-3-(2,6-difluorobenzoyl)urea.
50. The method of claim 1 in which the compound is 1-[4-(3,5-dichloro-2-pyridyloxy)-3,5-dichlorophenyl]-3-(2,6-difluorobenzoyl)urea.

51. The method of claim 1 in which from about 0.01 mg/kg body wt. to about 1000 mg/kg body wt. of the compound is administered to the host animal.

52. The method of claim 1 in which from about 1 mg/kg body wt. to about 100 mg/kg body wt. of the compound is administered to the host animal.

53. The method of claim 1 in which the administration is oral.

54. The method of claim 53 in which the administration is as a feed additive.

55. The method of claim 1 in which the administration is percutaneous.

56. The method of claim 55 in which the administration is by injection.

57. The method of claim 55 in which the administration is by pour-on application.

58. The method of claim 55 in which the administration is by implant.

59. The method of claim 55 in which the administration is by spraying.

60. The method of claim 55 in which the administration is by dipping or drenching.
61. The method of claim 1 in which the compound is used in combination with one or more biologically active materials.

62. The method of claim 1 in which the compound is used in combination with one or more medicinal drugs.

63. The method of claim 1 in which the compound is used in combination with one or more vertebrate toxicants.

64. The method of claim 1 in which the compound is used in combination with one or more growth promoters.

65. The method of claim 1 in which the compound is used in combination with one or more vitamins.

66. The method of claim 1 in which the compound is used in combination with one or more minerals.
### INTERNATIONAL SEARCH REPORT

**International Application No:** PCT/US 85/02545

**I. CLASSIFICATION OF SUBJECT MATTER**

According to International Patent Classification (IPC) and according to National Classification (NPC)

**IPC:**
- A 01 N 47/34, 47/36, 47/44, 55/00
- A 61 K 31/17, 31/33

**II. FIELDS SEARCHED**

- Classification System: A 01 N
- Classification Symbols: A 61 K

**III. DOCUMENTS CONSIDERED TO BE RELEVANT**

<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of Document</th>
<th>Relevance to Claim No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>X, Y US, A, 4089975 (L.L. WADE et al.) 16 May 1978, see abstract</td>
<td>1-29,31-66</td>
<td></td>
</tr>
<tr>
<td>X, Y US, A, 4092421 (L.L. WADE et al.) 30 May 1978, see abstract</td>
<td>1-29,31-66</td>
<td></td>
</tr>
<tr>
<td>X, Y US, A, 4166107 (J.A. MILLER et al.) 28 August 1979, see abstract</td>
<td>1-29,31-66</td>
<td></td>
</tr>
<tr>
<td>X, Y DE, A, 3225942 (A. NATTERMAN) 12 January 1984, see claims 1,7 (diflubenzuron) and 8</td>
<td>1-29,31-66</td>
<td></td>
</tr>
<tr>
<td>X, Y EP, A, 0079311 (CIBA-GEIGY) 18 May 1983, see claims 1,10-14; page 7, line 2</td>
<td>1-29,31-66</td>
<td></td>
</tr>
</tbody>
</table>

**IV. CERTIFICATION**

**Date of the Actual Completion of the International Search:** 23rd April 1986

**Date of Mailing of this International Search Report:** 11 JUN 1986

**International Searching Authority:** EUROPEAN PATENT OFFICE

**Signature of Authorized Officer:** L. ROSSI

**Form PCT/ISA/2/10 (second sheet) (January 1985)**
### III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)

<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of Document, with indication, where appropriate, of the relevant passages</th>
<th>Relevant to Claim No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Y</td>
<td>EP, A, 0050321 (UNION CARBIDE) 28 April 1982, see claims 1,28</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>Y</td>
<td>EP, A, 0098158 (UNION CARBIDE) 11 January 1984, see claims 1,30</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>Y</td>
<td>EP, A, 0113247 (UNION CARBIDE) 11 July 1984, see claims 1,22</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>Y</td>
<td>EP, A, 0115210 (UNION CARBIDE) 8 August 1984, see claims 1,17</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>Y</td>
<td>EP, A, 0124282 (UNION CARBIDE) 7 November 1984, see claims 1,26</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>X</td>
<td>EP, A, 0025363 (ISHIHARA) 18 March 1981, see claims 1,8 and 9</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>X</td>
<td>EP, A, 0107214 (DUPHAR) 2 May 1984, see claims 3 and 4</td>
<td>1-29,31-66</td>
</tr>
<tr>
<td>X</td>
<td>EP, A, 0136745 (DUPHAR) 10 April 1985, see claims 1,4 and 6 (cited in the application)</td>
<td>1-29,31-66</td>
</tr>
</tbody>
</table>

---

Form PCT/ISA:210 (extra sheet) (January 1985)
This Annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on 05/06/86.

The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

<table>
<thead>
<tr>
<th>Patent document cited in search report</th>
<th>Publication date</th>
<th>Patent family member(s)</th>
<th>Publication date</th>
</tr>
</thead>
<tbody>
<tr>
<td>US-A- 4089975</td>
<td>16/05/78</td>
<td>None</td>
<td></td>
</tr>
<tr>
<td>US-A- 4092421</td>
<td>30/05/78</td>
<td>None</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>JP-A- 59021616</td>
<td>03/02/84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>AU-A- 1664683</td>
<td>12/01/84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>EP-A, B 0068297</td>
<td>05/01/83</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DE-A- 3125423</td>
<td>13/01/83</td>
</tr>
<tr>
<td></td>
<td></td>
<td>AU-A- 8444182</td>
<td>06/01/83</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CA-A- 1173746</td>
<td>04/09/84</td>
</tr>
<tr>
<td>EP-A- 0079311</td>
<td>18/05/83</td>
<td>GB-A, B 2110672</td>
<td>22/06/83</td>
</tr>
<tr>
<td></td>
<td></td>
<td>AU-A- 9028082</td>
<td>19/05/83</td>
</tr>
<tr>
<td></td>
<td></td>
<td>GB-A, B 2155475</td>
<td>25/09/85</td>
</tr>
<tr>
<td></td>
<td></td>
<td>JP-A- 57158753</td>
<td>30/09/82</td>
</tr>
<tr>
<td></td>
<td></td>
<td>US-A- 4426385</td>
<td>17/01/84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CA-A- 1190553</td>
<td>16/07/85</td>
</tr>
<tr>
<td></td>
<td></td>
<td>OA-A- 6E312</td>
<td>31/07/85</td>
</tr>
<tr>
<td></td>
<td></td>
<td>AU-A- 1637483</td>
<td>05/01/84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>JP-A- 59155352</td>
<td>04/09/84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>JP-A- 59130870</td>
<td>27/07/84</td>
</tr>
<tr>
<td></td>
<td></td>
<td>US-A- 4321388</td>
<td>23/03/82</td>
</tr>
<tr>
<td>EP-A- 0107214</td>
<td>02/05/84</td>
<td>AU-A- 1862083</td>
<td>08/03/84</td>
</tr>
</tbody>
</table>

For more details about this annex, see Official Journal of the European Patent Office, No. 12/82.
V. OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE

This International search report has not been established in respect of certain claims under Article 17(3) (a) for the following reasons:

1. Claim numbers ........., because they relate to subject matter not required to be searched by this Authority, namely:

2. Claim numbers ........., because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

   - See PCT, Art. 17.2.a(ii)

3. Claim numbers ........., because they are dependent claims and are not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING

This International Searching Authority found multiple Inventions in this International application as follows:

1. As all required additional search fees were timely paid by the applicant, this International search report covers all searchable claims of the International application.

2. As only some of the required additional search fees were timely paid by the applicant, this International search report covers only those claims of the International application for which fees were paid, specifically claims:

3. No required additional search fees were timely paid by the applicant. Consequently, this International search report is restricted to the Invention first mentioned in the claims; it is covered by claim numbers:

4. As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

☐ The additional search fees were accompanied by applicant's protest.
☐ No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (supplemental sheet (2)) (January 1985)
<table>
<thead>
<tr>
<th>Document Type</th>
<th>Number</th>
<th>Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>AU-A-</td>
<td>3250484</td>
<td>07/02/85</td>
</tr>
<tr>
<td>JP-A-</td>
<td>60120847</td>
<td>28/06/85</td>
</tr>
<tr>
<td>JP-A-</td>
<td>59059617</td>
<td>05/04/84</td>
</tr>
</tbody>
</table>

For more details about this index:
see Official Journal of the European Patent Office, No. 12/82