INSTRUCTIONS

(a) If Convention application insert "Convention"

(b) Delete one

(c) Insert FULL name(s) of applicant(s)

(d) Insert FULL address(es) of applicant(s)

(e) Delete one

(f) Insert TITLE of invention

(g) Insert "complete" or "provisional" or "PCT Patent"

(h) Insert number, country and filing date for basic application

(i) Insert date of signing

(j) Signature of applicant(s)

(k) Corporate seal if any

Note: No legalization or other witness required
APPLICATION FOR A STANDARD/PETTY PATENT

I/We MOBIL OIL CORPORATION

of 150 East 42nd Street, New York, New York, United States of America

hereby apply for the grant of a Patent for an invention entitled

ACTIVITY ENHANCEMENT OF HIGH SILICA ZEOLITES BY THERMAL TREATMENT WITH ALKALINE ALUMINATE

which is described in the accompanying complete specification.

(Note: The following applies only to Convention applications)

Details of basic application(s)

<table>
<thead>
<tr>
<th>Application No.</th>
<th>Country</th>
<th>Filing Date</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
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</tbody>
</table>

Address for Service: PHILLIPS ORMONDE AND FITZPATRICK

Patent and Trade Mark Attorneys
367 Collins Street
Melbourne, Australia 3000

Dated 21 July, 1983

PHILLIPS ORMONDE AND FITZPATRICK
Attorneys for:
MOBIL OIL CORPORATION
AUSTRALIA
Patents Act

DECLARATION FOR A PATENT APPLICATION

In support of the application made by

MOBIL OIL CORPORATION

(hereinafter called "applicant(s) for a patent"

invention entitled "TREATMENT OF ZEOLITES"

of additon applicable

the patent application entitled

Edward H. Valance
Associate Patent Counsel
Authorized Attorney in Fact

To: The Commissioner of Patents

PHILLIPS ORMONE & FITZPATRICK
Patent and Trade Mark Attorneys
367 Collins Street
Melbourne, Australia
The process is carried out by treating the zeolite as crystallized or after calcination with the above described solution at elevated temperatures ranging up to 300°C, preferably to 160°C, for periods of time ranging from 1 hour to 30 days, preferably from 1 to 5 days. The hydrothermal conditions may require superatmospheric pressures at above 100°C to maintain adequate ionizing medium to maintain the metallate ion. Autogenous pressure can be maintained by autoclave or the like. Ordinarily pressures of 100 kpa to 1000 kpa are satisfactory.

Following the treatment, the zeolite may be further processed into its catalytically active form by conventional techniques, such as base exchange with appropriate cations such as hydrogen, ammonium, rare earth, and mixtures thereof. The zeolite is then advantageously calcined by heating to a temperature in the range of 200-600°C in an atmosphere such as air, nitrogen, etc., and atmospheric, subatmospheric, or superatmospheric pressure for between about 1 and 48 hours. The zeolite can, if desired, be incorporated in a matrix by techniques well known in the art for use as a catalyst. Conventional binder matrices include inorganic oxides, such as silica, alumina, silica-alumina, etc.
Claim

1. A method for increasing the catalytic activity, alpha, of a crystalline zeolite having a constraint index of 1 to 12 and a lattice silicon/non-silicon atomic ratio of at least 20 which comprises contacting the zeolite for a period of 1 hour to 30 days at a temperature of at least 100°C with an aqueous solution having a pH of 9 to 12 containing an ion of the formula M(OH)_4 where M is Al, B, Fe, Cr or Ga, separating the zeolite from the solution and converting it to the protonated form.
Complete Specification for the invention entitled:

**ACTIVITY ENHANCEMENT OF HIGH SILICA ZEOLITES BY THERMAL TREATMENT WITH ALKALINE ALUMINATE**

The following statement is a full description of this invention, including the best method of performing it known to applicant(s):
TREATMENT OF ZEOLITES

The invention relates to the treatment of zeolites to enhance their activity.

The silica-to-alumina mole ratio of a zeolite is often variable; for example, zeolite X can be synthesized with a silica-to-alumina ratio of from 2 to 3; zeolite Y from 3 to about 6. In some zeolites, the upper limit of silica-to-alumina ratio is unbounded, as in the case of ZSM-5 wherein the silica-to-alumina mole ratio is at least 5. US-A-3,941,871 discloses ZSM-5 essentially free of aluminum. US-A-4,061,724, 4,073,865 and 4,104,294 describe microporous crystalline silicas or organosilicates wherein the aluminum content present is at impurity levels.

Because of the extremely low aluminum content of such highly siliceous zeolites, their ion exchange capacity is not as great as materials with a higher aluminum content. Therefore, when these materials are contacted with an acidic solution and thereafter are processed in a conventional manner, they are not as catalytically active as their higher-aluminum counterparts.

According to the present invention a method of increasing the catalytic activity, alpha, of a crystalline zeolite having a constraint index of 1 to 12 and a lattice silicon/non-silicon atomic ratio of at least 20 comprises contacting the zeolite for a period of 1 hour to 30 days at a temperature of at least 100°C with an aqueous solution having a pH of 9 to 12 containing an ion of the formula M(OH)₄ where M is Al, B, Fe, Cr or Ga, separating the zeolite from the solution and converting it to the protonated (i.e., hydrogen or hydronium) form.

Typical zeolites to which the invention may be applied are ZSM-5, -11, -12, -23, -35, -38 and -48, defined respectively by the x-ray data set forth in US-A-3,702,886, 3,709,979, 3,832,649, 4,076,842, 4,016,245 and 4,046,859 and EP-A-15132. Both aluminosilicate and other, e.g., boro-, chromo-, ferro- and gallosilicate, forms of such zeolites are contemplated as starting materials, as are their effectively pure silica forms.

The invention may, from another aspect, be regarded
as zeolite activation by contacting high silica zeolite
under hydrothermal conditions with an alkaline metatellate
solution comprising at least one tetrahedrally bound
hydroxylated metatellate at a pH of about 9 to 12, and main-
taining the zeolite in contact with the solution at elevated
temperature under conditions to introduce tetrahedrally
coordinated metatellate into the crystalline zeolite.

The technique is particularly advantageous for trea-
ting acid ZSM-5 type zeolites having a silica:alumina mole
ratio greater than 50:1, in which case alkaline solutions
containing aluminate anion may be employed under hydrother-
mal conditions at elevated temperatures. Nitrogenous bases,
such as quaternary ammonium or amine compounds are preferred
to maintain the aluminum ion in the desired state for trea-
ting the zeolites.

The novel process of this invention permits the
preparation of highly siliceous zeolites which have all
the desirable properties inherently possessed by such high
silica materials yet have an acid cracking activity (alpha
value) which heretofore has only been manifested by mater-
ials having a higher aluminum content.

The preferred starting material for the process
of this invention is a high silica containing ZSM-5. The
process is simple in nature and easy to carry out although
the results obtained therefrom are dramatic. Nitrogenous
bases and/or alkali metal cations can be employed to achieve
a pH of at least 7. Preferably a pH greater than 9 up to
about 13 is maintained with organic cations, such as amines
or quaternary ammonium compounds. Being amphoteric, the
ions of the metal M can be supplied by any suitable organic
or inorganic salt such as chloride, sulfate, nitrate, acet-
tate, etc. However, at a moderately alkaline pH, (eg. pH
9 to 12) a stable M(OH)$^4^+$ ion becomes available for
hydrothermal treatment. This tetrahedral hydroxylated metatellate
ion is uniquely adapted for enhancing catalytic cracking
activity. The hydroxylated metatellates which are tetrahed-
rally coordinated in the zeolite crystalline structure ac-
cording to the invention are Ga(OH)$^4^+$, Fe(OH)$^4^+$,
Al(OH)$^4^+$, Cr(OH)$^4^+$ and B(OH)$^4^+$. 
The organic nitrogen-containing cation utilized in the hydrothermal solution can be a tetraalkylammonium cation, such as tetraethylammonium, tetrapropylammonium, tetrabutylammonium, methyl triethylammonium, methyl tripropylammonium, as well as mixtures thereof. The nitrogenous base can include various primary, secondary or tertiary amines especially normal alkyl amines such as n-propylamine, n-butylamine. It is advantageous to carry out the treatment at a pH of at least 7 up to about 13, and most preferably at a pH of 9 to 12. In addition to the organic bases, pH control can be accomplished by adding a suitable inorganic base such as sodium or ammonium hydroxide, to a solution of organic nitrogen-containing cation and aluminum ion.

A particularly preferred embodiment, in the case where aluminum is to be introduced into the lattice, is to use an alkali metal aluminate as the source of aluminum ion. This anion species may be prepared in situ by adding a base such as sodium hydroxide to an aqueous solution of an aluminum salt such as aluminum sulfate, or by dissolving sodium aluminate directly.

The relative proportion of metallate ion and organic nitrogen-containing cation present in the solution is not narrowly critical and usually ranges from 1 to 150 grams of equivalent metal salt per liter of solution and 1 to 200 grams of organic compound per liter of solution. The most preferred solvent is water for reasons of economy and ease of operation; however, various cosolvents may be employed within the inventive concept.

The amount of solution utilized to treat the zeolites is not critical and the solution to zeolite ratio can vary from 1 to 100 grams of solution per gram of zeolite. The amount of solution will vary as a function of its reaction kinetics concentration and desired enhancement of activity.

The process is carried out by treating the zeolite as crystallized or after calcination with the above described solution at elevated temperatures ranging up to 300°C, preferably to 160°C, for periods of time ranging from 1 hour to 30 days, preferably from 1 to 5 days. The hydrothermal conditions may require superatmospheric pressures.
at above 100°C to maintain adequate ionizing medium to maintain the metallate ion. Autogenous pressure can be maintained by autoclave or the like. Ordinarily pressures of 100 kpa to 1000 kpa are satisfactory.

Following the treatment, the zeolite may be further processed into its catalytically active form by conventional techniques, such as base exchange with appropriate cations such as hydrogen, ammonium, rare earth, and mixtures thereof. The zeolite is then advantageously calcined by heating to a temperature in the range of 200-600°C in an atmosphere such as air, nitrogen, etc., and atmospheric, subatmospheric, or vacuum for between about 1 and 48 hours. The zeolite can, if desired, be incorporated in a matrix by techniques well known in the art for use as a catalyst. Conventional binder matrices include inorganic oxides, such as silica, alumina, silica-alumina, etc.

The following examples illustrate the invention.

**EXAMPLE 1**

ZSM-5 of silica/alumina mole ratio 500 (A), 1600 (B) and >30,000 (C) was treated with a solution prepared from 2.55 grams of sodium hydroxide, 10.0 grams of tetrathylammonium bromide and 7.2 grams of Al₂SO₄·14H₂O and 115 grams of water under hydro-treating conditions: 200°C (212°F) for 6 hours at 1 atmosphere pressure. The solution to zeolite weight ratio was 3.4 parts of solution per part of zeolite. The zeolite was calcined in nitrogen at 540°C (1000°F) for 3 hours prior to treatment. Zeolite C was used as crystallized without calcination.

The three zeolites were then processed into their active form by calcination in nitrogen at 540°C (1000°F) followed by ammonium exchange with an aqueous solution of ammonium nitrate at ambient temperatures to remove sodium and/or excess aluminum ions. Finally, all three were air calcined at a temperature of 540°C (1000°F) for three hours. The three zeolites, before and after treatment, were evaluated for hydrocarbon cracking activity (alpha). (This test is described in Journal of Catalysis, Volume 4, pages 522-529, August 1965). The results obtained are shown...
ium can be further conventional cations thereof. Atmospheres about incorporated for use inorgan.

A), 1600 prepared of tetrapro-

It can be seen that the process of the invention resulted in a marked increase of activity. Thus, the ZSM-5 having a silica-to-alumina ratio of 500 had an alpha value of 1.0 prior to activation in accordance with the novel process of this invention, and thereafter its activity was raised to a value of 27. The effects of the alpha enhancement are more pronounced as the silica-to-alumina increases.

<table>
<thead>
<tr>
<th>Material</th>
<th>A</th>
<th>B</th>
<th>C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zeolite</td>
<td>ZSM-5</td>
<td>ZSM-5</td>
<td>ZSM-5</td>
</tr>
<tr>
<td>SiO2/Al2O3</td>
<td>500:1</td>
<td>1600:1</td>
<td>&gt;30,000:1</td>
</tr>
<tr>
<td>Form</td>
<td>Acid</td>
<td>Acid</td>
<td>Acid (as calcined)</td>
</tr>
<tr>
<td>cis(Untreated acid form)</td>
<td>10</td>
<td>1.7</td>
<td>0.1</td>
</tr>
<tr>
<td>cis(Treated acid form)</td>
<td>27</td>
<td>15</td>
<td>2.3</td>
</tr>
<tr>
<td>Enhancement</td>
<td>2.7</td>
<td>8.8</td>
<td>23</td>
</tr>
</tbody>
</table>

The following comparative examples demonstrate that the simple addition of aluminum either by impregnation or exchange cannot achieve the activity enhancement of high silica zeolites.

**EXAMPLE 2**

Four grams of type C low Al ZSM-5 (containing 50 ppm Al2O3) in its NH4 form are impregnated with solution of 0.03 g Al(NO3)3.9H2O in 2 g water. The wet zeolite mix is dried slowly, slugged and sized into 14/25 mesh. The resultant catalyst is activated at 540°C (1000°F) for three hours. The alpha value is 0.38.

**EXAMPLE 3**

Zeolite C catalyst is treated in ammonium form in a manner similar to those described in Example 2 except using 0.15 g Al(NO3)3.9H2O. The alpha value is 0.56.

**EXAMPLE 4**

Ammonium Zeolite C is treated in a manner similar to Example 2 except using 0.3 g Al(NO3)3.9H2O. The alpha value is 0.81.
Zeolite C is treated by impregnating 4 g of Type C low alumina ZSM-5 in its NH₄ form with 0.13 g of NaALO₂ dissolved in 2 g of H₂. The wet mix was dried at 110°C (230°F) for three hours and then calcined at 540°C (1000°F) for three hours. The Na content of the sample was reduced by NH₄ exchange. The sample was finally sized to 14/25 mesh and activated at 540°C (1000°F) for three hours again. The alpha activity of the sample was 0.1.

**EXAMPLE 6**

This catalyst is prepared in a similar manner as described in Example 5 except that the "as crystallized" low alumina ZSM-5 is used as base material. The alpha value was 0.43.

**EXAMPLE 7**

Four grams of "as synthesized" low alumina ZSM-5 is exchanged with a solution of 1.5 g Al(NO₃)₃.9H₂O in 20 ml of H₂O. After mixing for 2 hours the sample is filtered, washed and dried. The sample is then sized to 14/25 mesh and calcined at 1000°F for three hours and then exchanged with NH₄NO₃ solution to remove Na to 0.02% wt. The catalyst is finally activated in air at 540°C (1000°F) for three hours. The alpha activity of the catalyst is found to be 0.24.

**EXAMPLE 8**

This catalyst is treated in a similar manner to Example 7 except that 1.18 Al₂(OH)₅Cl is used instead of Al(NO₃)₃.9H₂O. The alpha activity of the catalyst was found to be 0.31.

The results obtained in Examples 2-8 are shown in Table 2, wherein it is demonstrated that the results obtained are distinctly inferior to the claimed invention as illustrated in Table 1.
Table 2

Aluminum Addition to High Silica Zeolites

<table>
<thead>
<tr>
<th>Example No.</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
<th>7</th>
<th>8</th>
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<tbody>
<tr>
<td>Base Material</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Zeolite</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>SiO₂/Al₂O₃</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Form</td>
<td>NH₄</td>
<td>NH₄</td>
<td>NH₄</td>
<td>NH₄</td>
<td>As Crystalized</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Treating Agent</td>
<td>Al(NO₃)₃</td>
<td>NaAlO₂</td>
<td>Al(NO₃)₃</td>
<td>Al₂(OH)₅Cl</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Method of Treat</td>
<td>Impregnation</td>
<td>EXCH</td>
<td>EXCH</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>%Al₂O₃ Incorporated (Based on Zeolite)</td>
<td>0.1</td>
<td>0.5</td>
<td>2</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
</tr>
<tr>
<td>Alpha of H Form Untreated</td>
<td>0.2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Treated</td>
<td>0.38</td>
<td>0.56</td>
<td>0.81</td>
<td>0.2</td>
<td>0.43</td>
<td>0.24</td>
<td>0.31</td>
</tr>
<tr>
<td>Example No.</td>
<td>2</td>
<td>3</td>
<td>4</td>
<td>5</td>
<td>6</td>
<td>7</td>
<td></td>
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<tr>
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<td>-----</td>
<td>-----</td>
<td>-----</td>
<td>-----</td>
<td></td>
</tr>
<tr>
<td>Base Material</td>
<td>Zeolite</td>
<td>ZSM-5</td>
<td>ZSM-5</td>
<td>ZSM-5</td>
<td>ZSM-5</td>
<td>ZSM-5</td>
<td></td>
</tr>
<tr>
<td>SiO₂/Al₂O₃</td>
<td>5</td>
<td>30,000</td>
<td>30,000</td>
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<td>30,000</td>
<td></td>
</tr>
<tr>
<td>Form</td>
<td>NH₄</td>
<td>NH₄</td>
<td>NH₄</td>
<td>NH₄</td>
<td>NH₄</td>
<td>NH₄</td>
<td></td>
</tr>
<tr>
<td>Treating Agent</td>
<td>Al(NO₃)₃</td>
<td>NaAlO₂</td>
<td>Al(NO₃)₃</td>
<td>NaAlO₂</td>
<td>Al(NO₃)₃</td>
<td>NaAlO₂</td>
<td></td>
</tr>
<tr>
<td>Method of Treat</td>
<td>Impregnation</td>
<td>EXCH</td>
<td>EXCH</td>
<td>EXCH</td>
<td>EXCH</td>
<td>EXCH</td>
<td></td>
</tr>
<tr>
<td>%Al₂O₃ Incorporated (Based on Zeolite)</td>
<td>0.1</td>
<td>0.5</td>
<td>2</td>
<td>0.5</td>
<td>0.5</td>
<td>0.5</td>
<td></td>
</tr>
<tr>
<td>Alpha of H Form Untreated</td>
<td>0.38</td>
<td>0.56</td>
<td>0.81</td>
<td>0.2</td>
<td>0.43</td>
<td>0.24</td>
<td></td>
</tr>
<tr>
<td>Treated</td>
<td>0.38</td>
<td>0.56</td>
<td>0.81</td>
<td>0.2</td>
<td>0.43</td>
<td>0.24</td>
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</tr>
</tbody>
</table>
The claims defining the invention are as follows:

1. A method for increasing the catalytic activity, \( \alpha \), of a crystalline zeolite having a constraint index of 1 to 12 and a lattice silicon/non-silicon atomic ratio of at least 20 which comprises contacting the zeolite for a period of 1 hour to 30 days at a temperature of at least 1000°C with an aqueous solution having a pH of 9 to 12 containing an ion of the formula \( M(OH)_4^- \) where \( M \) is Al, B, Fe, Cr or Ga, separating the zeolite from the solution and converting it to the protonated form.

2. A method according to claim 1 wherein pH is determined by the presence of alkali or nitrogenous base.

3. A method according to claim 2 wherein the nitrogenous base is a tetraalkylammonium compound.

4. A method according to claim 2 wherein the nitrogenous base is an alkylamine.

5. A method according to claim 2 wherein the nitrogenous base is tetraethylammonium, tetrabutyrammonium, methyltrihexammonium, methyltripropyrammonium, propylamine and/or butylamine.

6. A method according to any preceding claim wherein the ion \( M(OH)_4^- \) is derived from an organic or inorganic salt of \( M \).

7. A method according to any of claims 1 to 5 wherein the ion \( Al(OH)_4^- \) is derived from sodium aluminate.

8. A method according to claim 6 wherein the metal salt is present in a concentration of 1 to 150 gram equivalents per liter and a nitrogenous base is present in a concentration of 1 to 200 g/liter.

9. A method according to any preceding claim wherein the aqueous solution/zeolite weight ratio is from 1 to 100.

10. A method according to any preceding claim wherein the zeolite is calcined at a temperature of 200 to 600°C before the contacting.

11. A method according to any preceding claim wherein the contacting is carried out at a temperature of 100 to 300°C.

12. A method according to any preceding claim wherein the contacting is carried out at a temperature of 100 to
13. A method according to any preceding claim wherein the contacting is carried out at a pressure of 100 kpa to 10,000 kpa.

14. A method according to any preceding claim wherein the conversion of the zeolite to the protonated form is effected by hydrolysis or by ammonium exchange followed by calcination at 200 to 600°C.

15. A method according to any preceding claim wherein the non-silicon lattice element of the initial zeolite is other than aluminum.

16. A method according to any of claims 1 to 14 wherein the initial zeolite is an aluminosilicate having a silica/alumina ratio greater than 500, preferably greater than 1600.

17. A method according to any preceding claim in which the initial zeolite is one synthesized from a reaction mixture in which aluminum is present only as an impurity.

18. A method according to any preceding claim in which the zeolite has a constraint index of 1 to 12.

19. A method according to any of claims 1 to 17 in which the zeolite is ZSM-5, ZSM-11, ZSM-12, ZSM-23, ZSM-35, ZSM-38 or ZSM-48.

20. Use as a catalyst, or catalyst component, for the conversion of organic compounds of a zeolite obtained by the method claimed in any of claims 1 to 19.

DATED: 21 July, 1983

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MOBIL OIL CORPORATION
END