Novel lignin materials containing compositions

The present invention primarily relates to a preparation comprising or consisting of 40 - 99.9 wt.-% of lignin, preferably unmodified lignin, 0.1 - 50 wt.-% of minerals, preferably 0.1 - 30 wt.-%, 0 - 25 wt.-% of one or more mono- and oligomeric carbohydrates, preferably 0.1 - 20 wt.-%, and 0 - 15 wt.-% of one or more solvents, in particular water, preferably 0.1 - 4 wt.-%. Furthermore, the present invention relates to a fertilizer comprising or consisting of such a preparation and to the use of such a preparation as fertilizer, as ingredient for a fertilizer or for producing a fertilizer.
The present invention relates to a preparation containing lignin, which is obtainable by a method for recovering lignin (and, optionally, minerals) from a slurry after the bulk of cellulose and hemicellulose has been removed from a lignocellulosic biomass. The invention further relates to a fertilizer comprising or consisting of such a preparation and to the use of such a preparation as a fertilizer or for producing a fertilizer.

Pulping processes have previously been used to separate cellulose from lignin and other components of lignocellulose containing materials. For example, various types of inorganic chemicals in water have been used to modify lignin to render it water soluble. In the last decade various methods have been proposed to produce regenerated cellulose and hemicellulose while keeping the more or less unmodified lignin in solution and hence separate the cellulose/hemicellulose and the lignin by means of a solid/liquid separation. The solution system of the resulting slurry consists of a cellulose/hemicellulose solvent and a cellulose/hemicellulose precipitant and contains besides the lignin usually a substantial part of the minerals introduced into the process.

Recovering the lignin as well as the minerals from pulping processes is desirable for various reasons. First of all lignin is one of the few abundant providers of aromatic structures in nature rendering it valuable for chemical industry. Current research efforts focus on how unmodified lignin may be used. One example is the oxidation to vanillin. New investigations have shown that it is possible to produce carbon fibers in a reasonable quality and with a high market value from lignin, but the carbohydrate and sulfur content has to be very low. Secondly, lignin is the precursor for humus, the main factor controlling the water, carbon and mineral household of soil, providing a substantial value to agriculture.

The importance of carbon- and mineral cycle closure has been recognized again in the last decades. And thirdly, lignin may act as a carbon dioxide sink, because it is degraded only very slowly and if kept anoxic and dry it is stable. Under suitable conditions it is converted to coal over time.

Furthermore, it is desirable to completely remove and/or recover the solvents and precipitants used in pulping processes as they may be expensive and/or ecologically harmful, which is detrimental to further uses of the products obtained from a pulping process.

A preferred method for digesting a biomass comprising lignin together with cellulose and/or hemicellulose comprises the following steps (cf. for example WO 2010/043424 A1):

First, the biomass is contacted with a solvent for dissolving the cellulose and/or the hemicellulose. This is preferably done at a temperature below 120 °C and below a pressure of 300 kPa for less than 4 hours. Secondly, the dissolved cellulose and/or hemicellulose are precipitated by contacting it/them with a precipitant to form a compact solid comprising cellulose and/or hemicellulose as well as possibly residual lignin. The precipitant may be selected from the group consisting of alkanes, ethers, and esters or a mixture of these and the precipitation is preferably performed at a temperature below 120 °C and at a pressure of below 300 kPa for less than 30 min. Thirdly, the precipitate may be contacted with a solvent for dissolving the lignin in the precipitate. This step can also be performed when precipitating the cellulose and/or hemicellulose.

The ratio of the solvent for dissolving the cellulose and/or the hemicellulose to the biomass may be about 5:1 (weight/weight), but can also be greater. The volumetric ratio of the solvent for dissolving the cellulose and/or the hemicellulose and the precipitant for precipitating cellulose and/or hemicellulose may range between 1:1 to 1.3 (volume/volume).

The biomass may be placed in a suitable container, in particular a reactor. As lignocellulose containing biomass, wood (hard or soft), forest trimmings, leaves, recycled paper, waste paper, paper waste, corn stover, corn fiber, wheat straw, rice straw, rye straw, sugarcane bagasse, sugar beet cuttings, switchgrass, bamboo, pulp and/or similar organic materials can be used. The lignocellulose containing material may be reduced in particle size to pieces of 1 cm or less and further treated, e.g. it may be washed and/or its water content is changed by means known to a person of skill in the art.

The solvent for dissolving the cellulose and/or the hemicellulose which is added to the biomass in the reactor can be an inorganic acid, an inorganic hydroxide, an ionic solution, an organic acid and water or a mixture of any of these (Klemm, 1998: Klemm, D., Philipp, B., Heinze, T., Heinze, U, Wagenknecht, U Comprehensive Cellulose Chemistry Vol. I, Weinheim, VCH-Wiley 1998).

The mixture of biomass and solvent may be stirred at a temperature between 30°C and 90°C, but the temperature need not to be constant. The pressure is at normal ambient pressure, but may be varied. After 10 to 90 minutes, at least 60 % of the cellulose and hemicellulose are dissolved, depending on the kind of biomass and the solvent used. The incubation time may be extended such that practically all of the cellulose and hemicellulose in the reactor is dissolved.

After completion of the cellulose/hemicellulose digestion as described above, the reaction is stopped using a precipitant for precipitating the cellulose and hemicellulose, thereby forming a compact solid (precipitate). A precipitant may be chosen from alkanes, ethers and/or esters or a mixture of these, which do not swell or only modestly swell cellulose. The liquid retention value (LRV) for the precipitant may be smaller than the LRV of water for the cellulose used, wherein the LRV is defined according to Klemm, 1998, page 237, as follows:
The volumetric ratio of the solvent and the precipitant is 1:1 to 1:3, reducing the solvent amount needed for this and the following step. For this step, the conditions of temperature and pressure are considered of minor importance. Therefore temperature and pressure may be maintained from the previous step. Subsequently, the solid and the liquid phase are separated from each other.

After the precipitate containing the bulk of the cellulose and hemicellulose is removed, the remaining slurry contains the solvent for dissolving the cellulose and/or the hemicellulose (from now on called the "cellulose solvent"), the precipitant for precipitating the cellulose and/or hemicellulose (from now on called the "cellulose precipitant") as well as lignin and minerals, both of which are for the most part dissolved. The mixture may also comprise residual cellulose and/or hemicellulose as well as hydrolysation products thereof.

Further aspects of a preferred method for digesting a biomass comprising lignin together with cellulose and/or hemicelluloses can be derived from WO 2010/043424 A1, the content of which is herewith incorporated in its entirety.

A primary objective of the present invention was to provide a preparation containing lignin, preferably together with minerals, which can be obtained in a cheap and energy efficient way.

A further objective of the present invention was to provide a preparation containing lignin, preferably together with minerals, which comprises minimal residual amounts of solvents and/or carbohydrates.

Yet another objective of the present invention was to provide a preparation containing lignin, preferably together with minerals, which comprises a suitable amount of nutrients to be used as a fertilizer or to produce a fertilizer.

The objectives of the present invention as set out above are met by a preparation comprising or consisting of:

- 40 - 99.9 wt.-% of lignin, preferably unmodified lignin,
- 0.1 - 50 wt.-% of minerals, preferably 0.1 - 30 wt.-%
- 0 - 25 wt.-% of one or more mono- and oligomeric carbohydrates, preferably 0.1 - 20 wt.-% and
- 0 - 5 wt.-% of one or more solvents, in particular water, preferably 0.1 - 4 wt.-%

preferably

- 60 - 99 wt.-% of lignin,
- 1 - 20 wt.-% of minerals,
- 0 - 10 wt.-% of mono. and oligomeric carbohydrates, preferably 0.1 - 5 wt.- %, and
- 0 - 3 wt.-% of one or more solvents, preferably 0.1 - 2 wt.-%

particularly preferably

- 70 - 99 wt.-% of lignin,
- 1 - 10 wt.-% of minerals,
- 0 - 3 wt.-% of mono. and oligomeric carbohydrates, preferably 0.1 - 2 wt.-% and
- 0 - 0.5 wt.-%, preferably 0.1 - 0.5 wt.-% of one or more solvents.

A preparation according to the invention contains lignin as major component and can efficiently be obtained without the need to modify the lignin chemically.

Preferably, the preparation according to the invention contains minerals in an amount suitable to provide nutrition value for an agricultural application.

Further preferably, the amount of mono- and oligomeric carbohydrates in a preparation according to the invention is low so that a minimal amount of CO₂ is generated when the preparation is subjected to combustion. Advantageously,
a preparation according to the invention can therefore be used as a clean energy source.

[0022] It is furthermore preferred that the amount of solvents in a preparation according to the invention is low, so that the release of ecologically harmful and toxic compounds into the environment is minimal when a preparation according to the invention is used as energy source or in an agricultural application.

[0023] A preparation according to the invention may comprise one or more components selected from the group hydrolysis products of cellulose and/or hemicellulose, in particular xylose.

[0024] Among the hydrolysis products of cellulose and/or hemicellulose, which may be present in a preparation according to the present invention, xylose is in general the most abundant.

[0025] A preparation according to the invention may comprise a total amount of solvent(s) of less than 3 wt.-% preferably less than 2 wt.-%, particularly preferably less than 0.5 wt.-%, based on the total weight of the preparation, or it may not comprise any solvent at all.

[0026] It is preferred, that a preparation according to the invention contains a minimal amount of solvents, especially ecologically harmful or toxic (organic) solvents, which may release CO₂ or other undesirable compounds into the environment. A preparation according to the present invention may therefore safely be used as an energy source or for agricultural applications.

[0027] Furthermore, a preparation according to the invention may comprise a total carbohydrate content of less than 3 wt.-%, preferably less than 1 wt.-%, particularly preferably less than 0.5 wt.-%.

[0028] Preferably, the content of (residual) carbohydrate(s) in a preparation according to the invention is low, so that the amount of CO₂, which can be generated from the preparation is minimized. By reducing the carbohydrate content, an environmentally safe application of the preparation can be facilitated.

[0029] In a preferred preparation according to the invention, one, more or all of the minerals is/are selected from the group consisting of salts comprising or consisting of K⁺, Na⁺, Ca²⁺, Mg²⁺, Cl⁻, PO₄³⁻ and SO₄²⁻.

[0030] Advantageously, the mineral composition naturally contained in typical pulping slurry provides a suitable nutritional value for an agricultural application of a preparation according to the invention and the application contributes to close the mineral cycle. Thus, by isolating the lignin together with the minerals from pulping slurry, a suitable mineral composition is obtained without additional enrichment or removal of certain mineral components.

[0031] In a further preferred preparation according to the invention the content of

- N is in the range of 0.1 - 15 wt.-%, preferably in the range of 0.1 - 5 wt.-%, particularly preferably in the range of 0.1 - 3 wt.-%, and/or

- S is in the range of 0.1 - 5 wt.-%, preferably in the range of 0.1 - 3 wt.-%, particularly preferably in the range of 0.1 - 1 wt.-%,

based on the total weight of the preparation.

[0032] Furthermore, in a preferred preparation according to the invention the content of P is in the range of 0.1 - 10 wt.-%, preferably in the range of 0.1 - 5 wt.-%, particularly preferably in the range of 0.1 - 3 wt.-% based on the total weight of the preparation.

[0033] A preparation according to the invention advantageously provides the major essential nutrients for agricultural application in a suitable amount without the need of further additions.

[0034] A preparation according to the invention is obtained or obtainable by a method for isolating lignin, preferably unmodified lignin, and, optionally, minerals from a mixture, in particular a pulping slurry, in particular a pulping slurry preferably a mixture of a cellulose solvent and a pulping slurry obtained by a method as described in WO 2010/043424 A1, containing lignin, a cellulose solvent, a cellulose precipitant, minerals and optionally (residual) cellulose and/or hemicellulose and/or hydrolysation products thereof, comprising or consisting of the following steps:

i) removing the cellulose precipitant from the mixture,

ii) contacting the mixture with a lignin precipitant, preferably at a temperature below 50 °C, to obtain a precipitate,

iii) removing the precipitate obtained in step ii) from the mixture,

iv) optionally washing the precipitate with a lignin precipitant,

v) removing the lignin precipitant from the precipitate,

vi) optionally separating the cellulose solvent and the lignin precipitant from the remaining mixture obtained in step iii),
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with the proviso that in case the cellulose precipitant is the same as the lignin precipitant used in step ii) or contains one or more components of the mixture used as lignin precipitant in step ii), step i) is optional only, i.e. step i) can be omitted.

Advantageously, in step i) of the method described above, the cellulose precipitant is removed from the mixture, while the lignin and minerals largely remain dissolved. The method provides an energy efficient, cheap and easy way to obtain a preparation according to the invention.

The present invention further relates to a fertilizer comprising or consisting of a preparation as described above.

The application of a preparation as fertilizer is ecologically very valuable as it contributes to close the carbon and mineral cycle.

Therefore the present invention furthermore relates to the use of a preparation as described above as fertilizer, as ingredient for a fertilizer or for producing a fertilizer.

Moreover, further uses of a preparation according to the invention include the production of carbon fiber or polyurethanes as well as an energy source in a combustion process.

The following examples are added to illustrate the subject-matter of the invention, without limiting the scope of protection.

Example 1: Method for digesting lignocelluloses containing biomass using a cellulose solvent and a cellulose precipitant (to obtain a pulping slurry)

Lignocellulose containing biomass in the form of 50 g of coarsely grained wheat straw was stirred with 300 ml 80 % H₃PO₄ as cellulose solvent at 55°C for 45 min to the complete solution of the straw. The black viscous medium was cooled to approx. 40°C and then mixed vigorously with 600 ml of pre-cold MtB ether (methyl-tert. butylether) as a precipitant to give a compact precipitate of cellulose/hemicellulose. The solid phase was removed from the slurry.

Example 2: Method for digesting lignocelluloses containing biomass using a cellulose solvent and a cellulose precipitant in combination with a lignin solvent (to obtain a pulping slurry)

Lignocellulose containing biomass in the form of 30 g solid biogas digestate were mixed with 200 ml 80 % H₃PO₄ as cellulose solvent and stirred at 50 °C for 40 min. The solution is precipitated with 1 l pre-cold mixture of ethanol/acetone as precipitant with a ratio of 3: 1. The solid phase (the precipitate) was separated by centrifugation and the supernatant was collected.

Example 3: Method to obtain a preparation according to the invention from a pulping slurry obtained according to Example 1

i) The organic cellulose precipitant was removed under reduced pressure of 400 mbar from the collected supernatant of example 1 to give a black viscous solution, which was cooled to 40°C.

ii) 10 ml of the black solution was mixed vigorously with 50 ml ethyl acetate, which readily formed a brown precipitate.

iii) The brown precipitate was filtered of. The filtrate was collected for later treatment.

iv) The solids were suspended in another 50 ml ethyl acetate and filtered again. The filtrate was collected for later use in a step ii).

v) The brown solids were dried at 65 °C.

vi) The ethyl acetate was removed from the phosphoric acid by distillation leaving approx. 8 ml.

The regenerated acid of step vi) was able to dissolve 1.5g of straw.

The carbohydrate and ash content of the solids of step v) were determined as well as the element composition of to confirm that the brown precipitate was lignin.

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Example 4: Method to obtain a preparation according to the invention from a pulping slurry obtained according to Example 2

[0046]

i) The organic cellulose precipitant was removed under reduced pressure of 500 mbar from the collected supernatant of example 2 to give a black viscous solution, which was cooled to 40°C.

ii) 20 ml of the black solution was mixed vigorously with 40 ml di-propyl ether, which readily formed a brown precipitate.

iii) The brown precipitate was filtered off. The filtrate was collected for later treatment.

iv) The solids were suspended in another 40 ml di-propyl ether and filtered again. The filtrate was collected for later use in a step ii). The brown solids were washed with 20 ml hot water.

v) The brown solids were dried at 65 °C.

vi) The di-propyl ether was removed from the phosphoric acid by distillation leaving approx. 17 ml.

[0047] The regenerated acid of step vi) was able to dissolve 2.5 g of straw.

[0048] The carbohydrate and ash content of the solids of step v) were determined.

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Elementary analysis:

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<tr>
<td>N</td>
<td>1%</td>
</tr>
<tr>
<td>S</td>
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Claims

1. Preparation comprising or consisting of

- 40 - 99.9 wt.-% of lignin, preferably unmodified lignin,
- 0.1 - 50 wt.-% of minerals, preferably 0.1 - 30 wt.-%
- 0 - 25 wt.-% of one or more mono- and oligomeric carbohydrates, preferably 0.1 - 20 wt.-%, and
- 0 - 15 wt.-% of one or more solvents, in particular water, preferably 0.1 - 4 wt.-%,

preferably

- 60 - 99 wt.-% of lignin,
- 1 - 20 wt.-% of minerals,
- 0 - 10 wt.-% of mono. and oligomeric carbohydrates, preferably 0.1 - 5 wt.-%, and
- 0 - 3 wt.-% of one or more solvents, preferably 0.1 - 2 wt.-%

particularly preferably

- 70 - 99 wt.-% of lignin,
- 1 - 10 wt.-% of minerals,
- 0 - 3 wt.-% of mono. and oligomeric carbohydrates, preferably 0.1 - 2 wt.-% and
- 0 - 0.5 wt.-%, preferably 0.1 - 0.5 wt.-% of one or more solvents.

2. Preparation according to claim 1, wherein the preparation further comprises one or more components selected from the group hydrolysation products of cellulose and/or hemicellulose, in particular xylose.

3. Preparation according to claim 1 or 2, wherein the preparation comprises a total amount of solvent(s) of less than 3 wt.-% preferably less than 2 wt.-%, particularly preferably less than 0.5 wt.-%, based on the total weight of the preparation, or does not comprise any solvent at all.

4. Preparation according to one of the previous claims, wherein the preparation comprises a total carbohydrate content of less than 3 wt.-%, preferably less than 1 wt.-%, particularly preferably less than 0.5 wt.-%.

5. Preparation according to one of the previous claims, wherein one, more or all of the minerals is/are selected from the group consisting of salts comprising or consisting of \( K^+ \), \( Na^+ \), \( Ca^{2+} \), \( Mg^{2+} \), \( Cl^- \), \( PO_4^{3-} \) and \( SO_4^{2-} \).

6. Preparation according to one of the previous claims, wherein the content of

- \( N \) is in the range of 0.1 - 15 wt.-%, preferably in the range of 0.1-5 wt.-%, particularly preferably in the range of 0.1 - 3 wt.-%, and/or
- \( S \) is in the range of 0.1 - 5 wt.-%, preferably in the range of 0.1 -3 wt.-%, particularly preferably in the range of 0.1 - 1 wt.-%,

based on the total weight of the preparation.

7. Preparation according to one of the previous claims, wherein the preparation is obtained/obtainable by a method for isolating lignin, preferably unmodified lignin, and, optionally, minerals from a mixture, in particular a pulping slurry (preferably a mixture or, respectively, a pulping slurry obtained by a method as described in WO 2010/043424 A1), containing lignin, a cellulose solvent, a cellulose precipitant, minerals and optionally (residual) cellulose and/or hemicellulose and/or hydrolysation products thereof, comprising or consisting of the following steps:

i) removing the cellulose precipitant from the mixture,
ii) contacting the mixture with a lignin precipitant, preferably at a temperature below 50 °C, to obtain a precipitate,
iii) removing the precipitate obtained in step ii) from the mixture,
iv) optionally washing the precipitate with a lignin precipitant,
v) removing the lignin precipitant from the precipitate,
vi) optionally separating the cellulose solvent and the lignin precipitant from the remaining mixture obtained in step iii),

with the proviso that in case the cellulose precipitant is the same as the lignin precipitant used in step ii) or
contains one or more components of the mixture used as lignin precipitant in step ii), step i) is optional only, i.e. step i) can be omitted.

8. Fertilizer comprising or consisting of a preparation according to one of the previous claims.

9. Use of a preparation according to one of the claims 1 to 7 as fertilizer, as ingredient for a fertilizer or for producing a fertilizer.

Amended claims in accordance with Rule 137(2) EPC.

1. Preparation comprising or consisting of

- 60 - 99 wt.-% of unmodified lignin,
- 1 - 20 wt.-% of minerals,
- 0 - 10 wt.-% of mono. and oligomeric carbohydrates, preferably 0.1 - 5 wt.-%, and
- 0 - 3 wt.-% of one or more solvents, preferably 0.1 - 2 wt.-% particularly preferably
- 70 - 99 wt.-% of lignin,
- 1 - 10 wt.-% of minerals,
- 0 - 3 wt.-% of mono. and oligomeric carbohydrates, preferably 0.1 - 2 wt.-% and
- 0 - 0.5 wt.-%, preferably 0.1 - 0.5 wt.-% of one or more solvents.

2. Preparation according to claim 1, wherein the preparation further comprises one or more components selected from the group hydrolysation products of cellulose and/or hemicellulose, in particular xylose.

3. Preparation according to claim 1 or 2, wherein the preparation comprises a total amount of solvent(s) of less than 3 wt.-% preferably less than 2 wt.-%, particularly preferably less than 0.5 wt.-%, based on the total weight of the preparation, or does not comprise any solvent at all.

4. Preparation according to one of the previous claims, wherein the preparation comprises a total carbohydrate content of less than 3 wt.-%, preferably less than 1 wt.-%, particularly preferably less than 0.5 wt.-%.

5. Preparation according to one of the previous claims, wherein one, more or all of the minerals is/are selected from the group consisting of salts comprising or consisting of K⁺, Na⁺, Ca²⁺, Mg²⁺, Cl⁻, PO₄³⁻ and SO₄²⁻.

6. Preparation according to one of the previous claims, wherein the content of

- N is in the range of 0.1 - 15 wt.-%, preferably in the range of 0.1-5 wt.-%, particularly preferably in the range of 0.1 - 3 wt.-%, and/or
- S is in the range of 0.1 - 5 wt.-%, preferably in the range of 0.1 -3 wt.-%, particularly preferably in the range of 0.1 - 1 wt.-%,

based on the total weight of the preparation.

7. Preparation according to one of the previous claims, wherein the preparation is obtained/obtainable by a method for isolating lignin, preferably unmodified lignin, and, optionally, minerals from a mixture, in particular a pulping slurry (preferably a mixture or, respectively, a pulping slurry obtained by a method as described in WO 2010/043424 A1), containing lignin, a cellulose solvent, a cellulose precipitant, minerals and optionally (residual) cellulose and/or hemicellulose and/or hydrolysation products thereof, comprising or consisting of the following steps:

i) removing the cellulose precipitant from the mixture,
ii) contacting the mixture with a lignin precipitant, preferably at a temperature below 50 °C, to obtain a precipitate,
iii) removing the precipitate obtained in step ii) from the mixture,
iv) optionally washing the precipitate with a lignin precipitant,
v) removing the lignin precipitant from the precipitate,
vi) optionally separating the cellulose solvent and the lignin precipitant from the remaining mixture obtained in step iii).
with the proviso that in case the cellulose precipitant is the same as the lignin precipitant used in step ii) or contains one or more components of the mixture used as lignin precipitant in step ii), step i) is optional only, i.e. step i) can be omitted.

8. Fertilizer comprising or consisting of a preparation according to one of the previous claims.

9. Use of a preparation according to one of the claims 1 to 7 as fertilizer, as ingredient for a fertilizer or for producing a fertilizer.
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The present search report has been drawn up for all claims

Place of search: Munich
Date of completion of the search: 26 September 2013
Examiner: Friedrich, Christof
This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on 26-09-2013. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

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For more details about this annex: see Official Journal of the European Patent Office, No. 12/82
REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

• WO 2010043424 A1 [0005] [0014] [0034]

Non-patent literature cited in the description