Title: METHOD FOR PRODUCING ANHYDROUS CALCIUM NITRATE POWDER

Abstract: The invention relates to a method for producing an anhydrous powder having a calcium nitrate content of between 92 and 99.9 weight %, a water content of between 0.1 and 8 weight %; and a particle size of between 0.05 and 1.5 mm, wherein the method comprises the step of subjecting a calcium nitrate solution having a water content of between 70 and 15 weight % and a calcium nitrate content of between 30 and 85 weight % to a drying step in an industrial turbo-dryer, resulting in the anhydrous calcium nitrate powder. The invention furthermore relates to such an anhydrous calcium nitrate powder and the use of an industrial turbo-dryer to produce such anhydrous calcium nitrate powder.
METHOD FOR PRODUCING ANHYDROUS CALCIUM NITRATE POWDER

Field of the invention
The invention relates to a method for producing anhydrous calcium nitrate powder. Calcium nitrate is abbreviated as CN and has a structure formula of Ca(N0₃)₂.

The invention furthermore relates to an anhydrous calcium nitrate powder, more specifically produced by the abovementioned method.

The invention also relates to the use of an industrial turbo-dryer to produce such anhydrous calcium nitrate powder.

Background of the invention
Calcium nitrate has become a large and important product in the fertilizer field and also for many other technical applications such as waste water treatment, the concrete industry, etc.

Solid crystalline calcium nitrate is commonly found as hydrates such as Ca(N0₃)₂-4H₂O, Ca(N0₃)₂-3H₂O and Ca(N0₃)₂-2H₂O. Ca(N0₃)₂-4H₂O is a very common salt with a melting point of 42°C. It contains 68 - 69 weight % of calcium nitrate and about 30 weight % of water.

The traditional method for making calcium nitrate tetra hydrate [Ca(N0₃)₂-4H₂O] is by means of crystallization of a solution high in calcium nitrate which is cooled down below the saturation point of the calcium nitrate tetra hydrate. The solution is then filtered through which wet calcium nitrate tetra hydrate crystals are deposited on a filter. The crystals are removed from the filter and dried under vacuum at a temperature below 40°C.

Prilled or granulated calcium nitrate is commercially available. The normal composition of this material is between 77 and 80 weight % of Ca(N0₃)₂, between 6 and 8 weight % of ammonium nitrate (AN) and between 14 and 17 weight % of
crystallization water. Therein, the salt AN*5CN* 10H₂O, abbreviated as 1:5:10, constitutes a major part.

Prilled or granulated calcium nitrate products are produced by forming particles (e.g. prills, granules) of a melt of calcium nitrate, ammonium nitrate and water. Typically, such a melt has a water content of between 15 and 18 weight % of water and between 5 and 8 weight % of ammonium nitrate in addition to the calcium nitrate. The ammonium nitrate and the water are necessary to form a melt around 100 °C and 110 °C. The ammonium nitrate content is necessary to make the melt solidify quickly. If the ammonium nitrate is removed, the solidification process proceeds so slowly that normal prilling or granulation methods cannot be applied.

In WO 2004/039722, a method is described for the production of nitrate containing products from undercooling melts. More specifically, a cooling belt is used for solidification of the particles. This method is amongst others used to produce calcium nitrate in the form of prills, granules or pastilles. This procedure is not applicable to make pure calcium nitrate without crystal water since the melting point of pure anhydrous calcium nitrate is more than 500 °C.

For some applications however, there exists the need for pure, anhydrous calcium nitrate, i.e. calcium nitrate which is free of ammonium nitrate or any other salts and free of crystal water.

In GB 392,531, a process is disclosed for the preparation of non-caking fertilizers containing calcium nitrate, in which calcium nitrate solutions are evaporated to a sandy-pasty consistency in which they contain about 90 - 95 % of calcium nitrate (calculated as anhydrous calcium nitrate), and then converting the concentrated product without any substantial further concentration into granular form by mechanical disaggregation in bulk at a temperature substantially above atmospheric of the order of 50 to 100 °C. In example 1, an amount of calcium nitrate tetra hydrate is melted in a kneading machine adapted to be heated, where after this melt is further evaporated under vacuum to a content of calcium nitrate of 94 %, while the temperature is raised to 90 °C. The melt, at first thinly liquid, becomes rapidly
thickened and immediately afterwards changes over to a sandy-pasty mass. This mass is then cooled to 80°C and can easily be brought into a granular form by mechanical disaggregation at 80 °C.

The disadvantage of the method as described in GB 392,531 however is that it gives rise to salt deposits on equipment and a flaked, dusty product. Therefore, this method is not well suited for large-scale production of calcium nitrate.

CS 151689 B1 relates to production of powdered calcium nitrate solutions dried with hot air, the solution having a concentration of 15 to 70 weight% calcium nitrate dried in a stream of hot air with an inlet temperature of 130-500 °C and an outlet temperature of at least 80 °C.

WO 2007/012951 A1 relates to a method of producing anhydrous calcium nitrate, anhydrous magnesium nitrate or a mixture thereof, which includes providing a solution of calcium nitrate, magnesium nitrate or a mixture thereof, and removing water therefrom in a batch working pulse combustion drier.

A disadvantage of the methods from CS 151689 B1 and WO 2007/012951 A1 is that the calcium nitrate powder produced can be too wet/big or too dry/dusty.

The aim of the present invention is to form an anhydrous calcium nitrate having high to very high CN concentrations (also denominated as "pure" anhydrous calcium nitrate), that can be mixed with other powdery systems, therewith avoiding segregation. A further aim of the present invention is to form an anhydrous calcium nitrate powder with a determined target particle size between 0.05 and 1.5 mm, preferred water content between 0.1 and 8 weight% and calcium nitrate content between 92 and 99.9 weight%. A further aim of the present invention is to form a calcium nitrate powder wherein the moister content is within a value suitable for handling. A further aim of the present invention is provide a method for producing calcium nitrate powder wherein clogging of particles during the process is avoided. A further aim of the present invention is to provide such a highly concentrated anhydrous calcium nitrate that can dissolve quickly.
Summary of the invention

According to a first aspect of the present invention, a method is provided for producing an anhydrous calcium nitrate powder wherein the method comprises the step of subjecting a calcium nitrate solution having a water content of between 70 and 15 weight % of water and a calcium nitrate content of between 30 and 80 weight % to a drying step, wherein the calcium nitrate solution is subjected to a drying step in an industrial turbo-dryer, resulting in the anhydrous calcium nitrate powder having

- a calcium nitrate content of between 92 and 99.9 weight %;
- a water content of between 0.1 and 8 weight %; and
- a particle size of between 0.05 and 1.5 mm.

An industrial turbo-dryer is usually used to reduce the moisture content of various types of forestry wastes such as bagasse, coir, jute, pith and others. For instance a turbo-dryer (also called turbo-concentrator) as described in European patent 0749772 and manufactured by the company VOMM Impianti e Processi, Rozzano (MI) can be used. This machine basically comprises a cylindrical tubular body (also called a drum), having a horizontal axis and closed at opposite ends. These ends are provided with openings for the introduction of a liquid mixture to be treated and a stream of dry air travelling in the same direction. Furthermore, a heating jacket for heating the internal wall of the tubular body to a predetermined temperature, and a bladed rotor rotatable supported in the cylindrical tubular body. The circumferential speed of the bladed rotor varies between 30 and 50 m/s.

In an advantageous method according to the invention, the drying step is performed at a temperature of between 200 and 300 °C, more preferably between 240 and 280 °C.

In order to avoid the risk of crystallization and not to spend too much energy to evaporate the water, in a favorable method according to the invention, the calcium nitrate solution has a water content of between 45 and 50 weight % and a calcium nitrate content of between 55 and 50 weight %.
In a preferred method according to the invention, the anhydrous calcium nitrate powder has a calcium nitrate content of between 96 and 99.9 weight % and a water content of between 0.1 and 4 weight %, and most preferably a calcium nitrate content of between 95 and 99.9 weight % and a water content of between 0.1 and 5 weight %.

In a preferred method according to the invention, the anhydrous calcium nitrate powder has a particle size between 0.1 and 1 mm, and most preferably between 0.08 and 1 mm.

In order to obtain an anhydrous, "pure" calcium nitrate powder, in a preferred method according to the invention, the calcium nitrate solution is purified before being subjected to the drying step, for instance by filtration.

According to a further aspect of the invention, anhydrous calcium nitrate powder is provided having
- a calcium nitrate content of between 92 weight % and 99.9 weight %;
- a water content of between 0.1 weight % and 8 weight %; and
it has a particle size of between 0.05 and 1.5 mm,
wherein the calcium nitrate content and water content add up to 100 weight %.

More specifically, the anhydrous calcium nitrate according to the invention is produced by a method according to the invention as described above.

According to a final aspect of the present invention, the use of an industrial turbo-dryer to produce anhydrous calcium nitrate powder having
- a calcium nitrate content of between 92 weight % and 99.9 weight %;
- a water content of between 0.1 weight % and 8 weight %; and
- a particle size of between 0.05 and 1.5 mm;
starting from a calcium nitrate solution having a water content of between 70 and 15 weight % and a calcium nitrate content of between 30 and 85 weight % is provided.
More preferably, this use implements the method using an industrial turbo-drier according to the invention as described above.

**Detailed description of the invention**

The present invention relates to a method to produce anhydrous calcium nitrate powder by subjecting a calcium nitrate solution to a drying step which is performed in an industrial turbo-drier which is a continuous operating process. In order to obtain a pure end product, the calcium nitrate solution preferably is purified by filtering it.

The industrial turbo-drier consists of a horizontal drum with heating walls. Preferably, oil is used as the heating medium. In the longitudinal direction of the drum, a rotating shaft with blades has been mounted. The blades are extending only 5 mm from the wall of the drum. During operation, the shaft rotates with a high speed and creates a strong turbulence. By varying the angle of the blades, the retention time of the material in the drum can be varied. The liquid calcium nitrate solution is fed into the drum by warm air, and anhydrous calcium nitrate powder is extracted from the air stream leaving the drum by using a cyclone.

The temperature applied in the industrial dryer is preferably between 200 and 300 °C, and more preferably between 240 and 280 °C.

The starting calcium nitrate solution has a water content of between 15 and 70 weight % and a calcium nitrate content of between 30 and 85 weight %, and more preferably a water content of between 45 and 50 weight % and a calcium nitrate content of between 55 and 50 weight %, and most preferably approximately 50 weight % and a calcium nitrate content of approximately 50 weight %.

Subjecting this calcium nitrate solution to a turbo-dryer as disclosed above, allows an almost complete removal of the calcium nitrate dissolved in the calcium nitrate solution, resulting in an anhydrous calcium nitrate powder having

- a calcium nitrate content of between 92 and 99.9 weight %; and
- a water content of between 0.1 and 8 weight %; and
- a particle size of between 0.05 and 1.5 mm.

The obtained anhydrous calcium nitrate powder more preferably has a calcium nitrate content of between 96 and 99.9 weight % and a water content of between 0.1 and 4 weight %, and most preferably a calcium nitrate content of between 95 and 99.9 weight % and a water content of between 0.1 and 5 weight %.

In order to measure the particle size, the sample can be segregated by particle size. In particle segregation, particulate solids tend to segregate by virtue of differences in the size, and also physical properties such as volume, density, shape and other properties of particles of which they are composed. In the present example the particles obtained were segregated in a sieve shaker using a stack of woven wire mesh sieves having mesh sizes of 2 mm, 1.5 mm, 1 mm, 0.5 mm, 0.1 mm and 0.05 mm, respectively. Additional sieves can be used to obtain additional segregation.

The anhydrous calcium nitrate powder more preferably has a particle size between 0.1 and 1 mm, and most preferably between 0.08 and 1 mm.

**Example**

2000 litres of a purified calcium nitrate solution having a water content and a calcium nitrate content of 50 weight %, for example a purified Nutriox® solution, commercialised by Yara, having a density of 1.482 kg at 22°C and a pH of 6.0, was pumped into a pilot turbo-dryer commercialised by the Italian firm VOMM. The feeding rate of the drum of this turbo-dryer was between 70 and 80 litre per hour. By applying a temperature of approximately 270° in the VOMM turbo-dryer, an anhydrous calcium nitrate powder was obtained at a rate of 35 to 40 kg per hour. The obtained anhydrous calcium nitrate powder was collected and analysed. Results of this analysis can be found in Table 1 below.
The obtained anhydrous calcium nitrate powder has a calcium content of 23.5 - 24 weight % of calcium and around 16.5 weight % of nitrogen.

X-ray diffraction of the obtained calcium nitrate powder shows that the powder mainly consists of pure dry \( \text{Ca(N}_3\text{O}_3\text{)}_2 \) with traces of \( \text{Ca(N}_3\text{O}_3\text{)}_2\cdot2\text{H}_2\text{O} \) and \( \text{Ca(N}_3\text{O}_3\text{)}_2\cdot3\text{H}_2\text{O} \). No \( \text{CaO} \) and \( \text{Ca(OH)}_2 \) were detected. It is however likely that some ppm of these latter components were formed, being responsible for the pH increase.

The obtained anhydrous calcium nitrate powder dissolves quickly in water and forms a clear solution up to 10 - 15 weight % calcium nitrate. On further increasing the calcium nitrate in the solution, the calcium nitrate solution becomes weakly turbid, this probably due to the presence of some ppm \( \text{CaO} \) and \( \text{Ca(OH)}_2 \).

In order to store the anhydrous calcium nitrate powder according to the invention for a longer period of at least a year, this anhydrous calcium nitrate powder must be
stored in well-sealed, vapour tight bags. The anhydrous calcium nitrate powder according to the invention can also be compacted into granules by using various compaction machines.
CLAIMS

1. A method for producing an anhydrous powder wherein the method comprises the step of subjecting a calcium nitrate solution having a water content of between 70 and 15 weight % of water and a calcium nitrate content of between 30 and 80 weight % to a drying step, CHARACTERISED IN THAT the calcium nitrate solution is subjected to a drying step in an industrial turbo-dryer, resulting in the anhydrous calcium nitrate powder having
   - a calcium nitrate content of between 92 and 99.9 weight %;
   - a water content of between 0.1 and 8 weight %; and
   - a particle size of between 0.05 and 1.5 mm.

2. A method according to claim 1, wherein the drying step is performed at a temperature of between 200 and 300 °C.

3. A method according to claim 2, characterized in that the drying step is performed a temperature of between 240 and 280 °C.

4. A method according to claim 1 or 2, characterized in that the calcium nitrate solution has a water content of between 45 and 50 weight % and a calcium nitrate content of between 55 and 50 weight %.

5. A method according to any one of claims 1 to 4, wherein the anhydrous calcium nitrate powder has a content of calcium nitrate between 95 and 99.9 weight % and a water content of between 0.1 and 5 weight %.

6. A method according to claim 5, wherein the anhydrous calcium nitrate powder has a content of calcium nitrate between 96 and 99.9 weight % and a water content of between 0.1 and 4 weight %.
7. A method according to claim any one of claims 1 to 6, wherein the anhydrous calcium nitrate powder has a particle size between 0.1 and 1 mm.

8. A method according to claim 7, wherein the anhydrous calcium nitrate powder has a particle size of between 0.08 and 1 mm.

9. A method according to any one of claims 1 to 8, wherein the calcium nitrate solution is purified before being subjected to the drying step.

10. A method according to any of the previous claims wherein the calcium nitrate content and the water content add up to 100 weight %.

11. Anhydrous calcium nitrate powder, CHARACTERISED IN THAT it consists of

   - a calcium nitrate content of between 92 weight % and 99.9 weight %;
   - a water content of between 0.1 weight % and 8 weight %; and

   it has a particle size of between 0.05 and 1.5 mm,

   wherein the calcium nitrate content and the water content add up to 100 weight %.

12. Anhydrous calcium nitrate according to claim 11 produced by a method according to any one of claims 1 to 10.

13. Use of an industrial turbo-dryer to produce anhydrous calcium nitrate powder having

   - a calcium nitrate content of between 92 weight % and 99.9 weight %;
   - a water content of between 0.1 weight % and 8 weight %; and
   - a particle size of between 0.05 and 1.5 mm;

   starting from a calcium nitrate solution having a water content of between 70 and 15 weight % and a calcium nitrate content of between 30 and 85 weight %.

14. Use according to claim 13 implementing the method according to any one of the claims 1 to 10.
INTERNATIONAL SEARCH REPORT

A. CLASSIFICATION OF SUBJECT MATTER

INV. COFIL/44
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

COIF

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, BIOSIS, CHEM ABS Data, COMPENDEX, EMBASE, INSPEC, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

<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>X</td>
<td>GB 420 793 A (MONTEDISON SPA) 7 December 1934 (1934-12-07) 1,4-10, 13,14 1,2; example 1</td>
<td>2,3</td>
</tr>
<tr>
<td>X</td>
<td>GB 602 063 A (LONZA AG) 19 May 1948 (1948-05-19) page 1, line 10 - line 12 page 1, line 61 - line 75; example 1</td>
<td>11,12</td>
</tr>
<tr>
<td>Y</td>
<td>GB 254 939 A (BASF AG) 15 July 1926 (1926-07-15) page 1, line 17 - line 39</td>
<td>11,12</td>
</tr>
</tbody>
</table>

Further documents are listed in the continuation of Box C. See patent family annex.

Date of the actual completion of the international search: 13 August 2015

Date of mailing of the international search report: 24/08/2015

Name and mailing address of the ISA:
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016

Authorized officer: K Ling, Ruth
<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>A</td>
<td>EP 0 749 772 A1 (VOMM IMPIANTI &amp; PROCESSI SRL [IT]) 27 December 1996 (1996-12-27) cited in the application column 1, line 36 - line 57</td>
<td>1, 13</td>
</tr>
<tr>
<td>Patent document cited in search report</td>
<td>Publication date</td>
<td>Patent family member(s)</td>
</tr>
<tr>
<td>----------------------------------------</td>
<td>-----------------</td>
<td>--------------------------</td>
</tr>
<tr>
<td>GB 420793 A</td>
<td>07-12-1934</td>
<td>NONE</td>
</tr>
<tr>
<td>GB 602063 A</td>
<td>19-05-1948</td>
<td>BE 461733 A</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DE 849702 C</td>
</tr>
<tr>
<td></td>
<td></td>
<td>FR 919050 A</td>
</tr>
<tr>
<td></td>
<td></td>
<td>GB 602063 A</td>
</tr>
<tr>
<td></td>
<td></td>
<td>NL 67756 C</td>
</tr>
<tr>
<td>WO 2015067588 A1</td>
<td>14-05-2015</td>
<td>NONE</td>
</tr>
<tr>
<td>GB 254939 A</td>
<td>15-07-1926</td>
<td>NONE</td>
</tr>
<tr>
<td></td>
<td></td>
<td>BR 9602839 A</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CA 2179126 A1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DE 69617808 D1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>DE 69617808 T2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>EP 0749772 A1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>ES 2168435 T3</td>
</tr>
<tr>
<td></td>
<td></td>
<td>IT MI951323 A1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>US 5902520 A</td>
</tr>
</tbody>
</table>