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PROCESS FOR PREPARING FINE
POWDER POLYUREA AND GREASES THEREFROM

FIELD OF THE INVENTION

[0001] The present invention relates to the manufacture of polyurea powder and the manufacture of greases therefrom.

BACKGROUND OF THE INVENTION

[0002] Industrial lubricating greases are homogeneous products of semi-liquid to solid consistency. Essentially, they consist of a dispersion of a thickener in a liquid lubricant or base oil. In general, the thickener is a significant determinant of the properties of the greases.

[0003] Polyurea compounds are among the thickeners used in making greases. Often the polyurea compounds are prepared directly in the base oil by the reaction of amines with isocyanates.

[0004] One method to test the performance of greases is cone penetration and prolonged working of lubricating greases through ASTM D217. A cone of specified weight is allowed to fall into a lubricating grease sample at 25°C. The depth of the cone, in tenths of a millimeter, identifies the consistency of the grease. With the use of Table 1, the NLGI grade of the grease is identified from the 60 stroke worked penetration.

Table 1: NLGI Classification Scale

NLGI Grade	ASTM Worked Penetration
000	445-475
00	400-430
0	355-385
1	310-340
2	265-295
3	220-250
4	175-205
5	130-160
6	85-115

[0005] This test can be used to determine the mechanical stability of a grease through prolonged working, such as 10,000 or more double strokes using the motorized grease worker. While cone penetrations are typically conducted at 25°C, measurements can be carried out at other temperatures. About 300 grams of grease are required to conduct the ASTM D217 test. ASTM method D1403, DIN 51 804, and IP 310 describe cone penetration equipment commonly referred to as 1/2 and 1/4 scale devices for use when less than 300 grams of grease are available.

[0006] Dropping point of lubricating greases is used to determine high temperature structural grease properties related to the thickener. In ASTM D2265, the dropping point of a lubricating grease is the temperature at which the thickener can no longer hold the base oil. Some of the reasons oil can no longer be held are that the thickener has melted or the oil has become so thin it is not held by the thickener. Grease is placed in a small cup and heated in an oven-like device. When a drop of oil falls from the lower opening, the dropping point of the grease is calculated using the temperatures in the oven and inside the cup. Soap or polymer

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thickened greases demonstrate a dropping point while inorganic thickeners such as clay or graphite may not have a dropping point.

[0007] In U.S. Patent 5,314,982 there is disclosed a process for making polyurea greases by first making a dry polyurea compound. Then the compound is pulverized to give powders having particles in the 100 to 400 micron range. Thereafter, a paste of the powder and base oil is heated, cooled and homogenized in a high pressure homogenizer at pressures of 400 to 1500 bar.

[0008] In U. S. Patent 6,498,130 B2 a grease having low noise characteristics is made by shearing a base oil and thickener to reduce the thickener particles below about 500 microns in size. In this instance homogenization is achieved at about 2000 psi.

[0009] Both the above mentioned patents show the desirability of using polyurea of small particle size. They also illustrate that the conditions under which the polyurea particles are prepared impact subsequent processing conditions such as the homogenization step in grease forming.

[0010] User demand for polyurea greases has been increasing steadily; however, manufacturing such greases has been more difficult and expensive when compared to the manufacture of other greases.

[0011] One of the difficulties in manufacturing polyurea greases is consistently obtaining on an industrial scale substantially uniformly fine polyurea powder that is readily dispersible in a lubricating base oil.

[0012] Accordingly, one object of the present invention is to provide a process for making fine polyurea powder without the need for shearing or pulverizing large polyurea particles.

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[0013] Another object of the invention is to provide a process for making fine powdered polyurea compounds which can be practiced on an industrial scale and that can be homogenized under standard grease homogenizing conditions.

[0014] Yet another object of the invention is to provide an improved method for preparing polyurea greases thus greatly reducing the risk (hazards) associated in manufacturing with neat amines or isocyanates.

[0015] These and other objects of the invention will become more apparent from the following description.

SUMMARY OF THE INVENTION

[0016] Broadly stated, polyurea compounds are prepared by reacting amines and isocyanates in the presence of a liquid diluent in a high-pressure impingement mixing device under conditions sufficient to produce polyurea compounds having the consistency of a powder and in which diluent is dispersed.

[0017] An embodiment of the invention comprises passing the powder formed through a containment zone having means for monitoring and controlling the process.

[0018] Other embodiments and aspects will be clear from the following detailed description.

BRIEF DESCRIPTION OF THE DRAWING

[0019] Figure 1 is a diagram of the machine device used to make polyurea.

[0020] Figure 2 is a microphotograph of a polyurea particle made by impingement processing washed free of lubricating medium.

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DETAILED DESCRIPTION OF THE INVENTION

[0021] In the practice of the invention primary amines and isocyanates are reacted to form polyurea compounds.

[0022] The amines include aliphatic, alicyclic, aromatic amines and mixtures thereof. Examples of such mono amines include pentylamine, hexylamine, heptylamine, octylamine, dodecylamine, cyclohexylamine, benzylamine, aniline, diamnes and the like.

[0023] Suitable isocyanates are selected from the group consisting of polyisocyanate, monoisocyanate, and diisocyanate, and any combination thereof. Preferably, a diisocyanate is used. Furthermore, the isocyanate component may be aliphatic or aromatic and preferably is selected from an aromatic diisocyanate or mixtures of aromatic diisocyanates. Examples of such diisocyanates are phenylene diisocyanate, toluene diisocyanate, methylene diphenylene diisocyanate and the like. Typically, the amines and isocyanates are reacted in mole ratios of about 1:1 to about 2:1 and greater.

[0024] In the invention an amine and isocyanate is reacted in the presence of a liquid diluent in a high-pressure impingement mixing device such as impingement device 10 shown in Figure 1.

[0025] The liquid diluent employed in the practice of the present invention is one which is inert to the amines and/or isocyanates and is compatible with the components of any end product to be formed from the polyurea. Thus, in instances where the polyurea product is to be used as a grease thickener, a lubricating base oil suitable to grease formulations is a preferred diluent. Additionally, it is preferred that the amine or polyisocyanate be readily soluble in the diluent. In general, it is preferred to dissolve the amine in a base oil and preferably naphthenic, paraffinic, PAO, PAG and ester fluids can be used singularly or in combination as a diluent for

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either component individually or simultaneously. Also, any carbon base solvents that are compatible with either of the two components can be used as a diluent.

[0026] The amount of diluent used is not absolutely critical and will depend to a certain extent upon the solubility of the reactant, e.g., the amine in the diluent. Indeed, in the present invention, it is preferred to dissolve the amine in the diluent. Again, the diluent is not subjected to just amine alone but can be combined with the isocyanate. At least sufficient diluent is used to dissolve (disperse) the amine, and preferably, the amount of diluent will be sufficient to provide a solution having about the same density as that of the isocyanate or polyisocyanate – diluent combination. In the case where a naphthenic oil is used as diluent for the amine, the weight ratio of oil to amine will be about 1:1 to about 1:3.

[0027] Experience has shown that in the absence of diluent, the impingement device plugs almost immediately, and the polyurea formed typically has a significant amount of unreacted isocyanate groups, a result which is not desirable.

[0028] Referring to the Figure 1, isocyanate contained in vessel 11 is fed via line 12 to impingement device 10, while primary amine and diluent contained in vessel 14 is fed via line 15 to impingement device 10.

[0029] Means (not shown) are provided for regulating the flow of reactants from their respective vessels through orifices in the impingement device 10 (mix chamber) where the components are impinged and reacted.

[0030] The reactants are fed to impingement device 10 (mix chamber) under conditions sufficient to produce a polyurea compound having the consistency of a powder and, in which diluent is dispersed. The fluid composition of the solution or emulsion can be from 0.5:1 to 4:1 by weight of diluent to component. As presented in Laser Light Scattering analysis performed for particle size determination, the

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polyurea compound will comprise particles between 700 to 10 microns with a standard size being 200 microns.

[0031] Conditions that impact the size of the particles produced include the reactor orifice size (dimensions of the mix chamber), flow rates, pressure, individual component temperature and reactant residence time.

[0032] The flow rates of the reactants are typically in the mole ratio range described above.

[0033] The reactor orifice size for the amine and diluent and isocyanates typically will be different from that for the diisocyanate and generally will be chosen for facilitating the metering of the reactants in the appropriate mole ratio. Typically, the reactor orifice diameter will range from about 0.030 to about 0.150 inches; however, the reactor orifice diameter for the amine/diluent feed generally will be larger than that for the isocyanate.

[0034] The pressure at which the reactants are fed into the impingement device typically will be above 500 psig and preferably 1000 psig to 1800 psig, while the temperature at which they are fed may range from about 0°C to about 100°C and preferably 24°C to 55°C. The reactant residence time is a function of reactant feed rate and reactor volume, i.e., specified design of the mix chamber, the adjustment of which is well within the skill of a router in the art.

[0035] In one embodiment the fine powdered polyurea formed in impingement device 10 falls through a containment zone 16 as shown in the Figure 1. The containment zone 16 may be and preferably is equipped with an antistatic device 17. The finely divided powdered polyurea then may be collected on a conveyor belt system 18 for delivery to hopper 19 from which it can be automatically fed to packaging material 20, such as, bags, boxes or the like.

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[0036] In another embodiment of the invention, containment zone 16 is provided with product monitoring and control means 21. Such monitoring means may include an infrared spectrometer for detecting the presence of free isocyanate groups and a microscopy device for determining the size of the particles. Preferably, the monitoring means are operably connected to a computerized control device that functions to adjust processing conditions as necessary to produce particles of the requisite size.

[0037] The finely divided particulate polyurea powder, comprising from 5 to 15% of the total weight of base grease, is particularly suitable for forming greases and is mixed with a base lubricating oil, comprising from 85 to 95% of weight of base grease. The base lubricating fluid and polyurea powder are together placed in a conventional grease kettle or grease contactor at ambient temperature where it is dispersed, sheared (milled), mixed and heated to a top temperature in the range of about 150°C to 175°C, cooled to a specified temperature then milled (homogenized) to form a finished homogenous grease.

[0038] Optionally, typical grease additives such as extreme pressure additives, rust inhibitors, antiwear compounds and the like may be added to the mixture before milling.

[0039] In yet another embodiment of the invention, the finely powdered polyurea, rather than packaged, is fed directly to a grease kettle for mixing with a base oil and forming the mixture into a grease.

EXAMPLES

[0040] In the following Examples an air purge gun impingement device sold by Contrast Equipment Co., Kansas City, MO, was used. The mixing chambers used were obtained from Glas-Craft Inc., Indianapolis, IN, and had either flat, rectangular or round dispensing patterns. Fourier Transform Infrared Spectrometer

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("FTIR") analysis of the reaction was conducted on appropriate product samples to determine whether the product contained free isocyanate. The FTIR allows evaluating a product in process by monitoring the spectral peaks. The standard ninhydrine test was used to determine if a product contained unreacted amine.

Comparative Example 1:

[0041] In this comparative example the reaction chamber had a flat mixing chamber. The amine used was cyclohexyl amine, and the diisocyanate used was methylene diphenylene diisocyanate. The orifice diameter for the reactants were 0.071 inches and 0.042 inches respectively. The reactants were at ambient temperature (21°C), and each was fed to the reaction chamber at 700 psig. Within 5 seconds the chamber clogged.

Comparative Example 2:

[0042] The procedure of Comparative Example 1 was followed except a round mix chamber was used, and the pressure for each component was 1,000 psig. After about 3 seconds the chamber became clogged.

Example 1:

[0043] In this example a (round mix chamber) was employed. The orifice diameter for the amine feed was 0.109 inches, while the orifice diameter for the diisocyanate was 0.052 inches. The cyclohexyl amine was dissolved in a naphthenic oil having a kinematic viscosity @ 40°C of 143.21 cSt. The weight ratio of oil to amine was 1:1.2. Both the amine/oil feed and the diisocyanate feed were at 21°C. Then each were fed to the impingement device reaction chamber at 1000 psig. The resultant product had the consistency of a finely divided polyurea powder in which the diluent oil was dispersed. FTIR analysis failed to show a diisocyanate peak, and a ninhydrine test failed to show free amine establishing complete reaction. A sample of the product was mixed with an equal amount by

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weight of an oil typically used in formulating greases. The oil was readily incorporated in the product polyurea without any signs of separation.

Example 2:

[0044] In this example a round mix chamber was used. The amine used was tallow amine which was dissolved in the naphthenic oil at a 2:1 oil to amine weight ratio. The diisocyanate was at 21°C while the amine/oil feed was heated to 32°C. Both feeds were fed to the reactor at 1800 psig. The resultant polyurea product had a powdery consistency. FTIR analysis showed a diisocyanate peak at 2270⁻¹ cm of about 27% transmittance. As with the product of Example 1, this product blended very easily with a grease lubricating oil without any sign of oil separation.

[0045] Another attempt was made to manufacture completely reacted powder by using the same round mix chamber, the oil to amine ratio was changed to 1:1.5 and the feed pressure to the mix chamber was changed to 1000 psi. As a result, FTIR analysis failed to show a diisocyanate peak, and a ninhydrine test failed to show free amine establishing complete reaction. A sample of the product was mixed with an equal amount by weight of an oil typically used in formulating greases. The oil was readily incorporated in the product polyurea without any signs of separation.

[0046] An important aspect regarding prior art is that the structure of the particles are comparatively different. The prior art formulation and manufacturing of the polyurea powder results in the structure of polyurea particles being solid, compact and vary in size. U.S. Patent No. 5,314,982 and U.S. Patent Application No. 2006/0052261 are examples of the prior art. In comparison, polyurea particles made by impingement processing also vary in size but, the structure of the particle itself is different. The polyurea particles manufactured, by virtue of the processing described in this patent, are porous and sponge like (foamy cells) in their

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construction. The lubricating fluid used in the dilution of the amine, as explained in the examples presented, is encapsulated or rather occupy the interstices of the particle as shown in Figure 2. As observed during impingement processing, there is no oil pooling or separation of the polyurea powder from oil during or after the powder is made.

[0047] The polyurea powder is less dense than commercially available prior art polyurea. The polyurea preferably has a density less than 6.5 lbs/gal, more preferably less than 6 lbs/gal and most preferably has a density of less than 5.75 lbs/gal. In addition, the new polyurea powder has a larger surface area than prior art polyurea. Preferably, the polyurea powder has a specific surface area of more than 20 m²/g (measured by Hg porosimetry), more preferably 26 m²/g and most preferably more than 32 m²/g.

[0048] A shelf life study was conducted to monitor the integrity of a typical batch of impinged polyurea powder over a 12 month period. The observations made from the shelf life study are as follows: in general, the polyurea powder still maintained its original integrity after 12 months of storage, there was sign of excessive caking and no indication visibly or microscopically of oil leaching from the stored powder.

[0049] Regarding grease processing with the impinged preformed polyurea powder, the visibly large particles are easily broken apart, or fragmented, by the shearing devices, mills and homogenizers, used in grease processing. Thus, over time during the base grease processing prior to top temperature, the mass becomes thicker and more grease like as time and temperature increase over the course of the proscribed manufacturing process.

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[0050] Grease made with polyurea using the new method has improved properties. Table 2 shows the different properties of the two polyurea greases. Sample A is a grease made with conventional polyurea and higher pressures. Sample B is a grease made with new polyurea and with new method. As shown in Table 2, Sample A made with the old prior art method of higher cook temperature and higher homogenization pressure ("Hog") exhibits poor penetration properties. Sample B shows improved properties using the new method of producing grease.

TABLE 2: Properties of the Polyurea Grease

SAMPLE	A	B
TEST		
Hog pressure	@ 6000 psig	1500 psig
Cook temp, °F	400	320
Thickener content, %	12%	12%
Appearance	smooth dark brown	smooth light brown
Penetration		
Unworked	300	294
Worked (60x)	324	292
Extended (10K)	400	100k = 353
Points change (60X to ext)	76	65
Dropping Point, °C	254	269

CLAIMS

1. A polyurea particle having interstices in which lubricating oil is encapsulated, said particle having the consistency of a powder and a density of less than 6.5 lbs/gal.

2. A polyurea particle having interstices in which lubricating oil is encapsulated, said particle having the consistency of a powder and a specific surface area of more than 20 m²/g.

3. A polyurea particle having interstices in which lubricating oil is encapsulated, said particle having a particle size that is at least 10 microns and less than 700 microns, a density of less than 6.5 lbs/gal, a specific surface area of more than 20 m²/g, and the consistency of a powder.

4. A polyurea particle having interstices in which lubricating oil is encapsulated, said particle having a particle size that is at least 10 microns and less than 700 microns, a density of less than 5.75 lbs/gal, a specific surface area of more than 32 m²/g, and the consistency of a powder.

5. A polyurea grease comprising a polyurea particle having interstices in which lubricating oil is encapsulated, said particle having a particle size that is at least 10 microns and less than 700 microns, a density of less than 6.5 lbs/gal, a specific surface area of more than 20 m²/g, and the consistency of a powder, wherein the polyurea particle is blended with a lubricating oil.

6. A polyurea grease according to claim 5 wherein the polyurea particle has a density of less than 5.75 lbs/gal and a specific area of more than 32 m²/g.

7. A polyurea grease according to claims 5 or 6 wherein the polyurea particle and lubricating oil are mixed, heated to a top temperature of about 150°C to 175°C, cooled and homogenized.

8. The polyurea particle of claim 1, wherein the lubricating oil is chosen from the group consisting of naphthenic, paraffinic, PAO, PAG and ester fluids, alone or in any combination thereof.