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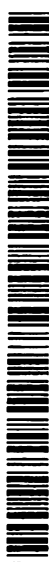


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(54) Title: **STORAGE-STABLE AERATED GEL COMPOSITION AND A PROCESS FOR PRODUCING IT**

(57) Abstract: A storage-stable aerated gel composition comprises 30 to 97 % by weight of water, 0.2 to 5 % by weight of a gelling agent selected from xanthan gum, sodium alginate and neutralised carboxyvinyl polymer and 2 to 5 % by weight of a fine particulate, hydrophobic silicone-treated silica having a surface area of from 80 to 300 m²/g. The composition is in the form of fine particles of an aqueous gel containing the water and gelling agent, the surfaces of which fine particles are coated with a coating of the finely particulate hydrophobic silica. The gel composition is prepared by mixing, under high shear conditions, the water and the fine particulate silica and then adding, to the resulting mixture, the gelling agent and mixing under high shear conditions.

STORAGE-STABLE AERATED GEL
COMPOSITION AND A PROCESS FOR PRODUCING IT

The present invention relates to storage-stable aerated gel compositions. It also relates to a process for making such compositions. In particular, the gel compositions contain a particular hydrophobic silica, water and gelling agent.

Aqueous dispersions of silica can be prepared into a state known generally in the prior art as "dry water". In fact, "dry water" is known in two forms. The first form can be produced by absorbing aqueous liquids onto hydrophilic material to form a material which exists as free-flowing powder or granules. The second form can be produced by coating finely divided aqueous liquids with powdered hydrophobic material, such as metal oxides. Each liquid particle in this second form of "dry water" is separated from the next by a hydrophobic metal oxide coating and by air spaces. Very high speeds of, for example, over 6000 rpm, and mixing times of 15 minutes are typically required. This second form is, however, thermodynamically unstable and, when produced, tends to break down after a relatively short period of time.

The present invention is based on the discovery that stable aerated gels analogous to this second form of "dry water" can be produced.

The present invention provides a storage-stable aerated gel composition comprising 30 to 97% by weight of water, 0.2 to 5% by weight of a gelling agent selected from xanthan gum, sodium alginate and neutralised carboxyvinyl polymer and 2 to 5% by weight of a fine particulate, hydrophobic silicone-treated silica having a surface area of from 80 to 300 m²/g which said composition is in the form of fine particles of an aqueous gel containing the water and gelling agent, the surfaces of which fine particles are coated with a coating of the finely particulate hydrophobic silica.

The present invention further provides a process for producing a storage-stable aerated gel composition which comprises the steps of mixing, under high shear conditions, from 2 to 5% by weight of a finely particulate,

hydrophobic silicone-treated silica having a surface area of from 80 to 300 m²/g and from 30 to 97% by weight of water, adding to the resulting mixture from 0.2 to 5% by weight of a gelling agent selected from xanthan gum, sodium alginate and neutralised carboxyvinyl polymer and mixing the gelling agent with the silica-water mixture under conditions of high shear.

A discovery on which the present invention is based lies in the use of certain gelling agents which, when added to a premix formed by mixing the water and a specific type of hydrophobic silica under high shear conditions and then mixed with the premix also under high shear conditions, give a storage-stable aerated gel composition. Many conventional gelling agents which are ordinarily used to gel aqueous systems do not produce storage-stable gel compositions according to the present invention. The reasons for this are not, at present, understood. Furthermore, by adjusting the components used in the present invention the structure of the gel composition produced can be varied from a free-flowing pulverulent fluid composition which may have applications using conventional spray or dusting equipment to a non-flowing composition which may have applications as a caulking material.

Where the words "comprises" or "comprising" are used herein, it is intended that these may have the meaning "includes" and "including", respectively, to the extent that the presence of one or more other materials is not excluded.

The composition of the present invention comprises a fine particulate hydrophobic silicone-treated silica having a surface area of from 80 to 300 m²/g. By the term "fine particulate", as applied to the hydrophobic silica, it is meant that the hydrophobic silica will typically have an average particle size of less than 40µm. The silica used is one that has been rendered hydrophobic by surface treatment using one or more organosilicon compounds to produce, on the silicon dioxide surface, silicone groups. The technique of hydrophobicizing silica in this way is well-known and such silicone-treated silica is available commercially. We have found that good results are obtained by using hydrophobic silica marketed under the name CAB-O-SIL

("CAB-O-SIL" is a trademark of Cabot Corporation), preferably CAB-O-SIL TS720. However, other silicone-treated silicas can also be used in the present invention if they have a surface area within the range of from 80 to 300 m²/g. The hydrophobic silica may also be one that has been surface treated to produce siloxane, as well as, silicone groups attached to the silicon dioxide surface.

The hydrophobic silica is used in an amount of from 2 to 5% by weight based on the total weight of the composition. The use of greater than 5% by weight of the hydrophobic silica results in a gel composition which is excessively dusty. The use of such a composition may give rise to a greater nuisance dust risk. Preferably, the amount of hydrophobic silica will be in the range of from 3 to 4% by weight of the composition.

The water used may typically be tap water although purified grades may be appropriate for some applications. The water will normally be used at ambient temperature since there appears to be no advantage in using heated or cooled water in the performance of the invention. The water generally will form from 30 to 97% by weight of the total composition. Preferably, however, the amount of water will be from 90 to 97% by weight to ensure the formation of aerated gel compositions of good consistency and improved stability.

As mentioned above, the gelling agent is one or more selected from xanthan gum, sodium alginate and neutralised carboxyvinyl polymers, such as carboxypolymethylene neutralised with triethanolamine. These gelling agents are included in an amount of from 0.2 to 5% by weight. The use of an amount greater than 5% by weight of the gelling agent results in a gel composition having an excessively high gel strength. Preferably, from 0.5% to 2% by weight of the gelling agent will be used depending on the desired stability and structure of the gel composition. Other hydrophilic or hydrophobic additives known to those skilled in the art may be incorporated to modify the physical or biological properties of the composition.

The process for producing the gel compositions involves mixing the water with the silica under high shear conditions typically for a few minutes, for example 2 to 5 minutes. The mixing at this stage in the process must be

carried out under high shear conditions, i.e., conditions which cause the water to be finely fragmented into minute droplets which become dispersed within the finely particulate hydrophobic silica such that the surfaces of the water droplets become coated with the hydrophobic silica particles. The term "high shear" is, of course, well-known to the person skilled in the art of mixing or blending and whether or not a particular mixing apparatus is capable of mixing aqueous compositions under high shear conditions will be known to one skilled in the art. This may be achieved by using standard high speed mixers, typically using a mixing speed of at least 2000 rpm and generally from 2000 to 3000 rpm. After the hydrophobic silica and the water have been mixed to create a dispersion of fine droplets of water in the silica, the gelling agent is added and mixing at high speed is continued for several minutes until the gelling agent has been thoroughly incorporated into the liquid phase. It is preferred in the present invention to add the gelling agent after the silica and water have been thoroughly mixed together. If the gelling agent is added before the silica, mixing requires more energy and the homogeneity and stability of the resulting gel composition can be compromised.

The storage-stable aerated gel compositions of the present invention can be used in sprayable formulations produced for domestic, veterinary, agricultural or horticultural applications.

The invention will now be illustrated by the following examples in which the compositions in Examples 1 to 3 and 5 to 7 were mixed using an IKA RE166 high speed mixer having a radial flow toothed disc mixer head and the composition in Example 4 was mixed using a larger scale Torrance high speed mixer having a radial flow toothed disc mixer head.

EXAMPLE 1

Cold tap water 95.5g was mixed with 3g of CAB-O-SIL TS720 (silica) at 2800 rpm on the laboratory mixer for 2 minutes. "Dry water" was not formed. Powdered xanthan gum 1.5g was added and the mixer speed increased to 5500 rpm for a further 3 minutes. A free-flowing aerated gel was formed with

a density of 0.6g/ml. No water separated on storage at laboratory ambient (max/min 30/8°C) at 24 weeks.

EXAMPLE 2

Cold tap water 96.5g was mixed with 3g of CAB-O-SIL TS720 at 2800 rpm on the laboratory mixer for 2 minutes. "Dry water" did not form. Powdered xanthan gum 0.5g was added and the mixer speed increased to 5500 rpm for a further 3 minutes. A free-flowing aerated gel was formed similar in appearance to that formed in Example 1. The gel composition formed in this Example could be syringed through a 0.26mm internal diameter needle but blocked a 0.21mm internal diameter needle.

EXAMPLE 3

Using a larger mixing vessel with the laboratory mixer 1930g of cold tap water, 60g of CAB-O-SIL TS720 and 10g of powdered xanthan gum were mixed by the same process as described in Example 2. The product was identical to that produced in Example 2. A 1 litre sample was transferred to a 2 litre PET bottle and held under 1.7 bar pressure. This sample remained stable with no water separation for greater than 31 days stored at ambient temperature. Some water separation was noted after 59 days under continuous pressure. The sample was easily re-homogenised by 5 vigorous inversions and no further water separated until 16 hours later.

EXAMPLE 4

Cold tap water 77.2kg was mixed with 2.4kg of CAB-O-SIL TS720 on the production mixer at 2500 rpm for 1 minute. "Dry water" did not form. Powdered xanthan gum 0.4kg was added and mixing continued at the same speed for a further 10 minutes. The product was a free-flowing aerated gel as in Examples 1 to 3. Six further 80kg batches were made by the same process and recipe without variation in the product. A sample of this product 8.6kg was packed into a polypropylene bucket with a tightly fitting lid. This sample was transported in a car for 2022 miles over a duration of 6 weeks. No water

separation occurred during this period. At 14 weeks there was 4.7% m/m water separation. Further samples from these production batches were applied as a "wet dust" through a GLORIA 2010 Knapsack sprayer at 2 to 4 bar pressure fitted with a TEEJET 65030E brass nozzle. The nozzle filter was removed for these applications. A total of 115 litres of dry gel were applied through this knapsack sprayer without blockage. Further samples were applied as a "wet dust" through a STIHL SR400 motorised mistblower fitted with a pressure pump. The tank filter was removed and the standard adjustable spray nozzle was retained for these applications. A total of 250 litres of dry gel was applied through this mistblower at a nozzle setting of 3 to 4 without blockage.

EXAMPLE 5

Cold tap water 76.5g, propylene glycol 20g and 3g of CAB-O-SIL TS720 were mixed on the laboratory mixer at 2800 rpm for 2 minutes. "Dry water" did not form. Powdered xanthan gum 0.5g was added and the mixing speed increased to 5500 rpm for a further 3 minutes. The product was a free-flowing aerated gel of a wetter consistency than in Examples 1 to 4. No water separation occurred after 5 days ambient storage.

EXAMPLE 6

Cold tap water 96.3g and CARBOPOL 980[®] (registered trademark of BF Goodrich Co.) 0.5g (a carboxyvinyl polymer) were mixed on the laboratory mixer at 2800 rpm for 1 minute. CAB-O-SIL TS720 3g was added and mixed at the same speed for a further 2 minutes. "Dry water" did not form. Triethanolamine 0.2g was added to neutralise the CARBOPOL and the mixing speed increased to 5500 rpm for a further 3 minutes. A thick pliable aerated dry gel was formed that could be easily extruded through a 30ml plastic syringe. No water separation occurred after 13 weeks ambient storage.

EXAMPLE 7

Cold tap water 96g and 3g of CAB-O-SIL TS720 were mixed on the laboratory mixer at 2800 rpm for 2 minutes. "Dry water" did not form. Sodium alginate, high viscosity grade, 1g (a polysaccharide) was added and the mixing speed increased to 5500 rpm for a further 3 minutes. A free-flowing aerated gel was initially formed which thickened to a non-flowing gel after 5 days. The gel remained pliable and could easily be extruded from a 30ml plastic syringe. No water separation occurred after 6 days ambient storage.

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The reference to any prior art in this specification is not, and should not be taken as, an acknowledgment or any form or suggestion that that prior art forms part of the common general knowledge in Australia.



CLAIMS

1. A storage-stable aerated gel composition comprising 30 to 97% by weight of water, 0.2 to 5% by weight of a gelling agent selected from xanthan gum, sodium alginate and neutralised carboxyvinyl polymer and 2 to 5% by weight of a fine particulate, hydrophobic silicone-treated silica having a surface area of from 80 to 300 m²/g which said composition is in the form of fine particles of an aqueous gel containing the water and gelling agent, the surfaces of which fine particles are coated with a coating of the finely particulate hydrophobic silica.
2. A composition according to claim 1 comprising from 90 to 97% by weight of water.
3. A composition according to claim 1 or claim 2 comprising from 3 to 4% by weight of the silica.
4. A process for producing the storage-stable aerated gel composition which comprises the steps of mixing, under high shear conditions, from 2 to 5% by weight of a fine particulate, hydrophobic silicone-treated silica having a surface area of from 80 to 300 m²/g and from 30 to 97% by weight of water, adding to the resulting mixture from 0.2 to 5% by weight of a gelling agent selected from xanthan gum, sodium alginate and neutralised carboxyvinyl polymer and mixing the gelling agent with the silica-water mixture under conditions of high shear.
5. A process according to claim 4, wherein the water is used in an amount of from 90 to 97% by weight.
6. A process according to either claim 4 or claim 5, wherein the silica is used in an amount of from 3 to 4%.

7. A process according to any one of claims 4 to 6, wherein the gelling agent is xanthan gum.
- 5 8. A process according to any one of claims 4 to 7, wherein the water and the silica are mixed together at a mixing speed in the range of from 2000 to 3000 rpm.
9. A process according to any one of claims 4 to 8, wherein the gelling agent is mixed with the premix of silica and water at a mixing speed of
- 10 from 5000 to 6000 rpm.
10. A composition according to claim 1 substantially as hereinbefore described with reference to the Examples.
- 15 11. A process according to claim 4 substantially as hereinbefore described with reference to the Examples.

20 DATED this 15th day of March, 2004

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