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DESCRIPTION

FIELD OF THE INVENTION

[0001] This invention relates to opacifying particles used in coating compositions. In particular, this invention relates to opacifying particles with internal voids produced in a multi-stage polymerization process.

BACKGROUND OF THE INVENTION

[0002] Scattering pigments are commonly used in coating compositions to achieve opacity in both aqueous systems and non-aqueous systems. Titanium dioxide is commonly used as an opacifying pigment. One limitation in the use of titanium dioxide in aqueous coating compositions is that some of the pigment tends to aggregate. Any aggregation of pigment reduces the efficiency with which it can scatter light. This problem has been minimized in the past by use of surfactants and various dispersion techniques or low cost extenders added to the coating composition to create space between the individual titanium dioxide pigment particles. In other applications, pigment particles have been encapsulated within polymer to create a physical barrier between individual pigment particles and exhibit better dispersion within a polymer system due to the greater compatibility of the polymer enclosed particle than with the polymer system than the pigment particle itself.

[0003] More recently, the cost for producing titanium dioxide (TiO_2) has increased. Increased cost continues to drive an effort to minimize the amount of titanium dioxide used in coating compositions. One option for extending the amount of TiO_2 used in the coating composition is to include TiO_2 pigment particles incorporated into hollow particles that also include air voids. These types of particles exhibit enhanced scattering compared to similarly sized TiO_2 particles.

[0004] Document CN 102 311 556 discloses titanium dioxide particles which are covered by hollow polymeric microspheres. The hollow particles are first prepared and are then coated on titanium dioxide via a sol/gel process, so that a TiO_2 particle is coated with a multitude of hollow spheres.

SUMMARY OF THE INVENTION

[0005] The present invention includes a method of preparing an aqueous dispersion of particles at least partially encapsulated by a polymer, the method comprising dispersing inorganic particles in an aqueous phase; functionalizing at least a portion of the surfaces of the inorganic particles with a coupling agent, the coupling agent having pendent reactive groups;

polymerizing the reactive groups on the functionalized particles with a first polymerizable component to produce inner polymer shells surrounding the inorganic particles; polymerizing a second polymerizable component to form outer polymer shells at least partially encapsulating the inner polymer shells; and producing voids within the outer polymer shells, such that the inorganic particles are received within the voids. Also included in the present invention is an aqueous dispersion of particles comprising inorganic particles at least partially encapsulated by polymerized inner shells, wherein at least a portion of the surfaces of the particles comprise a coupling agent having pendent reactive groups, and wherein the inner shells are prepared from a reaction of the coupling agent pendent reactive groups with ethylenically unsaturated monomers and outer polymer shells at least partially encapsulating the inner polymer shells, wherein the outer polymer shells are at least partially spaced apart from the inner polymer shells, thereby defining voids within the outer polymer shells.

BRIEF DESCRIPTION OF THE DRAWINGS

[0006]

FIG. 1 is a transmission electron microscope (TEM) image of particles produced according to one embodiment of the present invention;

FIG. 2a is a photograph of a coating composition containing particles of the present invention and prior art particles, in a wet state; and

FIG. 2b is a photograph of a coating composition containing particles of the present invention and prior art particles, in a dry state.

DETAILED DESCRIPTION OF THE INVENTION

[0007] For purposes of the following detailed description, it is to be understood that the invention may assume various alternative variations and step sequences, except where expressly specified to the contrary. Any numerical value inherently contains certain errors necessarily resulting from the standard variation found in their respective testing measurements.

[0008] The opacifying particles of the present invention include multi-staged opacifying particles including inorganic particles that are encapsulated by polymeric shells. As used herein, the term "encapsulated" encompasses at least partially encapsulated or fully encapsulated unless indicated otherwise. At least a portion of the surfaces of the inorganic particles is functionalized with a coupling agent having pendant reactive groups. A polymer layer is produced on the functionalized particle surfaces by reaction of the coupling agent pendant reactive groups with ethylenically unsaturated monomers, yielding seed-core particles.

At least one outer polymer shell encapsulates the inner shells with the particles received therein. The outer polymer shells are spaced from the inner particle shells, thereby defining voids within the outer polymer shell. As such, the opacifying particles of the present invention include outer polymer shells that encapsulate a void as well as inorganic particles that are encapsulated by polymerized inner shells. The core of the multi-staged opacifying particles includes inorganic particles, specifically pigment particles such as TiO_2 . Other suitable inorganic particles that may be received within the core include ZnO , ZnO_2 , PbO , ZnSi , ZnSiO_4 , ZnS or ZnSe_2 . Each of these materials has a high refractive index (e.g., over 1.65).

[0009] In one embodiment, the inorganic particles are encapsulated in an inner shell. The particle-inner shell structure is referred to herein as a seed-core particle, onto which subsequent polymer layers are deposited. To produce the seed-core particles, the inorganic particles are treated with a coupling agent to functionalize at least a portion of the surface of the inorganic particles with a coupling agent bearing pendant reactive groups. In one embodiment, the coupling agent may include a methoxysilane, such as methacryloxypropyltrimethoxy silane, vinyltrimethoxysilane, vinyltriethoxysilane, methacryloxypropyl methyltrimethoxysilane, methacryloxypropyl methyltriethoxysilane, methacryloxypropyl triethoxysilane, or acryloxypropyl trimethoxysilane. The coupling agent of the present invention includes reactive pendant groups, such as reactive groups comprising ethylenic unsaturation. The ethylenic unsaturation of the pendant reactive groups of the coupling agent are then reacted with a first polymerizable component also comprising ethylenic unsaturation. In another embodiment, the coupling agent may instead include a hydrophobic organic component (without ethylenic unsaturation) to which monomers for producing the core polymer shell adsorb and are subsequently polymerized to form the core polymer shell. Suitable monoethylenically unsaturated monomers useful for making the core polymer include monoethylenically unsaturated monomers containing acid-functionality such as monomers containing at least one carboxylic acid group including (meth)acrylic acid ("meth)acrylic" herein meaning acrylic or methacrylic), (meth)acryloxypropionic acid, itaconic acid, aconitic acid, maleic acid, fumaric acid, crotonic acid, monomethyl maleate, monomethyl fumarate, and monomethyl itaconate. Acid precursors such as anhydrides, for example maleic anhydride, are included. Acrylic acid and methacrylic acid are particularly suited for use as the first polymerizable component. Other suitable first polymerization components for producing the seed-core particles include nonpolymeric compounds containing at least one carboxylic acid group including C_6 - C_{12} aliphatic or aromatic monocarboxylic acids and dicarboxylic acids, such as benzoic acid, m-toluic acid, p-chlorobenzoic acid, o-acetoxybenzoic acid, azelaic acid, sebacic acid, octanoic acid, cyclohexanecarboxylic acid, lauric acid and monobutyl phthalate.

[0010] The core polymer shell may be made in a single stage or step of sequential polymerization or be made by a plurality of steps in sequence. In one embodiment, a water soluble free radical initiator is utilized in the aqueous emulsion polymerization. Suitable water soluble free radical initiators include ammonium persulfate, hydrogen peroxide, tert-butyl peroxide, alkylidene metals such as sodium potassium and lithium persulfate, and mixtures of such an initiator with a reducing agent such as a sulfite. The temperature may be in the range

of 10°C to 100°C. In the case of persulfate systems, the temperature is preferably in the range of 60°C to 90°C.

[0011] A nonionic or anionic emulsifier may be used, either alone or together. Examples of suitable nonionic types of emulsifiers include tert-octylphenoxyethylpoly(39)-ethoxyethanol, polyethylene glycol 2000 monooleate, ethoxylated castor oil, block copolymers of propylene oxide and ethylene oxide, polyoxyethylene(20) sorbitan monolaurate and nonylphenoxyethylpoly(40)ethoxyethanol. Examples of suitable anionic emulsifiers include sodium lauryl sulfate, sodium dodecylbenzenesulfonate, sodium lauryl ether sulfate, sodium alpha-olefin (C₁₄-C₁₆) sulfonate, ammonium or alkali metal salts of sulfosuccinate derivatives, ammonium or alkali metal salts of fatty acids such as stearic acid, linseed oil fatty acid, and coconut oil fatty acid, ammonium or alkali metal salts of phosphate esters of ethoxylated nonylphenol and tert-octylphenoxyethoxypoly(39)ethoxyethyl sulfate.

[0012] Upon reaction of the first polymerizable component with the coupling agent pendant reactive groups, the inorganic particles are encapsulated in a core polymer shell, which may be hydrophilic, thereafter serving as seed-core particles. The phrase "seed-core particles" means an aqueous emulsion polymer dispersion that includes the inorganic particles encapsulated in core polymer shells. Thereafter, the seed-core particles are subsequently encapsulated in subsequent shells in emulsion polymerization processes by deposition thereon of polymerized products of one or more successive monomer charges introduced to a medium containing the dispersed seed-core particles in one or more subsequent stages.

[0013] At least one outer shell is deposited onto the seed-core particles to at least partially encapsulate the seed-core particles. In one embodiment, one or more of the outer shells are hydrophilic at least at some pH to enhance subsequent swelling thereof. The outer shells may be produced from nonionic monoethylenically unsaturated monomers such as styrene, α -methyl styrene, vinyltoluene, ethylene, vinyl acetate, vinyl chloride, vinylidene chloride, acrylonitrile, (meth)acrylamide, (C₁-C₂₀) alkyl or (C₃-C₂₀) alkenyl esters of (meth)acrylic acid, such as methyl (meth)acrylate, ethyl (meth)acrylate, butyl (meth)acrylate, 2-ethylhexyl (meth)acrylate, benzyl (meth)acrylate, lauryl (meth)acrylate, oleyl (meth)acrylate, palmityl (meth)acrylate, and stearyl (meth)acrylate.

[0014] Polyethylenically unsaturated monomers that can be used in preparing the outer shells include alkylene glycol diacrylates and dimethacrylates, such as for example, ethylene glycol diacrylate; ethylene glycol dimethacrylate; 1,3-butylene glycol diacrylate; 1,4-butylene glycol diacrylate propylene glycol diacrylate and triethylene glycol dimethylacrylate; 1,3-glycerol dimethacrylate; 1,1,1-trimethylol propane dimethacrylate; 1,1,1-trimethylol ethane diacrylate; pentaerythritol trimethacrylate; 1,2,6-hexane triacrylate; sorbitol pentamethacrylate; methylene bisacrylamide; methylene bis-methacrylamide; divinyl benzene; vinyl methacrylate; vinyl crotonate; vinyl acrylate; vinyl acetylene; trivinyl benzene; triallyl cyanurate; divinyl acetyl divinyl ethane; divinyl sulfide; divinyl ether; divinyl sulfone; diallyl cyanamide; ethylene glycol divinyl ether; diallyl phthalate; divinyl dimethyl silane; glycerol trivinyl ether; divinyl adipate; dicyclopentenyl (meth)acrylates; dicyclopentenylloxy (meth)acrylates; unsaturated esters of

glycol monodicyclopentenyl ethers; allyl esters of α,β -unsaturated mono- and dicarboxylic acids having terminal ethylenic unsaturation including allyl methacrylate; allyl acrylate; diallyl maleate; diallyl fumarate; and diallyl itaconate.

[0015] Polymerization of the at least one outer shell onto the seed-core particle may be formed in the same reaction vessel in which the seed-core particles were produced or the reaction medium containing the dispersed seed-core particles may be transferred to another reaction container. In one embodiment, the monomers used in producing each of the outer shell polymers are first combined in pre-emulsions which are then sequentially charged into the container containing the seed-core particles. In one embodiment, a plurality of outer shells are applied sequentially to the seed-core particles yielding a multistage polymer-encapsulated particle, with a final outer shell produced from nonionic monoethylenically unsaturated monomers and polyethylenically unsaturated monomers. In general, the seed-core particles include hydrophilic exterior surfaces, and the shells deposited thereon are progressively less hydrophilic, typically resulting in a hydrophobic outermost shell.

[0016] The outer most shell may be produced by hydrophobic, higher T_G monomers such as styrene, vinylchloride or methylmethacrylate and mixtures thereof, crosslinked with a suitable crosslinking agent such as ethylene glycol dimethacrylate.

[0017] The amount of polymer deposited to form the plurality of outer shells is generally such as to provide an overall size of the opacifying particle of 100 to 2500 nanometers, such as 200 to 1500 nanometers, in unswollen condition (that is, before any neutralization to raise the pH to 6 or higher). In one embodiment where the opacifying particles include three outer shells, each of the first and second outer shells are at least 15%, preferably at least 20%, by weight of the total of all three outer shell polymers.

[0018] Voids are produced in the multistage polymer-encapsulated particle by adding a suitable swelling agent to which the final hydrophobic shell polymer is permeable. For example, the outer shells may be swollen with a conjugate base and a solvent, which permeates through the shell polymers. The swollen multistaged polymer-encapsulated particles are then dried. The composition of the polymer of the core polymer shell and at least the hydrophilic outer shells may be selected to allow for swelling thereof while the composition of the outermost shell may be selected so as to be permeable to an aqueous or gaseous volatile or fixed basic swellant for the hydrophilic polymers of the internal shells and provide stability to a final dispersion of the opacifying particles.

[0019] The hydrophilic polymers of the multistage polymer-encapsulated particle are swollen when the particles are subjected to a basic swelling agent that permeates the outermost shell to at least partially neutralize (to a pH of at least 6 to 10) the hydrophilic-functionality of the seed-core polymer and at least one hydrophilic outer shell and thereby to cause swelling by hydration of both the core polymer shell and at least one hydrophilic outer shell. Nonlimiting examples of swelling agents include volatile bases such as ammonia, ammonium hydroxide, and volatile lower aliphatic amines, such as morpholine, trimethylamine, and triethylamine; and

fixed or permanent bases such as potassium hydroxide, lithium hydroxide, zinc ammonium complex, copper ammonium complex, silver ammonium complex, strontium hydroxide, and barium hydroxide. When in an aqueous dispersion, swelling of the core polymer shell and at least one hydrophilic shell results in a net flow of water into the particles, thereby swelling the hydrophilic polymers. When the swelling agent is neutralized (e.g. by addition of a neutralizing agent such as hydrogen peroxide) and the particles are dried, the hydrophilic polymers shrink. The shrinkage of the core polymer shell and at least one outer shell develops a void between the hydrophilic shells and the hydrophobic outer shell. The extent of the voids depends in part upon the resistance of the shells to restoration to their previous sizes. By "void" it is meant a polymer-free space, typically filled with water in the aqueous dispersion and with air in the dried hollow sphere pigment. After swelling, the outermost shell is crosslinked to yield opacifying particles comprising a polymer encapsulated pigment (TiO_2) received within a seed-core shell, with voids between the outermost shell and the polymer encapsulated pigment. In one embodiment, the polymer encapsulated pigment is free to move within the outer shell (may be referred to as "rattling" within the outer shell) or the polymer encapsulated pigment particles may adhere to an inner surface of the outer shell. In either such an arrangement, the presence of air surrounding or partially surrounding the polymer encapsulated pigment particle results in refractive index contrast that enhances the hiding ability of the particles.

[0020] In one embodiment, the opacifying particles are provided in an aqueous dispersion of opacifying particles. As described above, the opacifying particles comprises inorganic particles at least partially encapsulated by polymerized inner shells, wherein at least a portion of the surfaces of the particles comprise a coupling agent having pendent reactive groups, and wherein the inner shells are prepared from a reaction of the coupling agent pendent reactive groups with ethylenically unsaturated monomers and further encapsulated by outer polymer shells at least partially encapsulating the inner polymer shells, wherein the outer polymer shells are at least partially spaced apart from the inner polymer shells, thereby defining voids within the outer polymer shells.

[0021] The opacifying particles and aqueous dispersion thereof according to the present invention are particularly suited for use in coating compositions containing a waterborne resinous binder and may also be solvent based. It has been found that the opacifying particles of the present invention provide similar or enhanced hiding in coatings compositions, such as paints, and improved hiding in wet paint, as compared to traditional hiding pigments (TiO_2).

[0022] The following examples are presented to demonstrate the general principles of the invention. The invention should not be considered as limited to the specific examples presented. All parts are by weight unless otherwise indicated.

EXAMPLES

Example 1: Seed-core particles

[0023] TiO₂ pigment particles were coated with alkali-swellable polymer shells as follows. Deionized (DI) water (800g), 150g TiO₂ slurry (TRONOX® CR-826 available from TRONOX Limited, 2.5g Disponil A 1080 (an emulsifier, available from BASF), 2.5g Disponil FES-993 (an emulsifier, available from BASF) and 3g 30% ammonia aqueous solution were first charged into a four necked round bottom flask fitted with a thermocouple, mechanical stirrer, and condenser, and then heated up to 80°C under an N₂ blanket. Once the temperature reached 80°C, a mixture of 12g of Silquest A174NT (methacryloxypropyltrimethoxy silane), 20g methyl methacrylate (MMA) and 1g of Sipomer PAM 200 (an emulsifier, available from Rhodia) were charged into the flask over 30 minutes. After 30 minutes holding, a mixture of 1.5g of ammonium persulfate and 18g of DI water was charged into the flask all at once and held at 30 minutes. After holding was completed, a pre-emulsion of 90g of DI water, 2g of SR 550 (methoxy polyethylene glycol (350) monomethacrylate, available from Sartomer), 5.0g of Disponil A 1080, 6g of Disponil FES 993, 45g of methacrylic acid (MAA) and 115g of MMA, was charged into the flask over 2 hours. After the charge was completed, it was held at 80°C for an additional 1 hour before being cooled down to room temperature.

Example 2

[0024] Opacifying particles with TiO₂ in the center were prepared using the seed-core particles produced in Example 1 as follows. DI water (4340 g) was first charged into a four necked round bottom flask fitted with a thermocouple, mechanical stirrer, and condenser, and then heated up to 85°C under N₂ blanket. Once the temperature reached 85°C, a mixture of 4.4g of ammonium persulfate and 62.5g of DI water was charged into the flask, and then 720g of the seed-core latex of Example 1 was charged into the flask. The mixture was held at 77°C for 10 minutes.

[0025] Three shells were formed over the seed-core particles by first charging into the flask (over 30 minutes) a first monomer charge of a pre-emulsion of 160g of DI water, 3.8g of sodium dodecylbenzenesulfonate (SDBS), 9.4g of MAA, 62.5g of MMA, 1.8g of ethyleneglycol dimethacrylate (EGDMA) and 87.5g of styrene. Immediately after the first monomer charge was completed, a second monomer charge of a pre-emulsion of 531g of DI water, 4.7g of ammonium persulfate, 6.9g of SDBS, 12.0g of MAA, 47g of EGDMA and 1375g of styrene, was charged over 90 minutes into the flask. After that, the mixture was heated up to 90°C, and held at 90°C for 15 minutes. After holding, a third monomer charge of a pre-emulsion of 375g of DI water, 3.75g of SDBS, 31.2g of Disponil A1080, 63g of EGDMA, and 250g of styrene, was charged into the flask over 10 minutes.

[0026] The seed-core and first shell were swollen by neutralization with a base. A mixture of 106g of 30% ammonia solution and 281g of DI water was charged into the flask over 15 minutes, and held at 90°C for 45 minutes to swell the seed core and shells of the latex

particles. The latex was cooled down to 77°C, and a mixture of 6.0g of t-butyl hydroperoxide (70%) and 62.0g of DI water was charged into the flask all at once. A mixture of 6.0g of ascorbic acid and 120.0g of DI water was then charged over 15 minutes to polymerize the styrene in the shells.

[0027] The resultant latex was further centrifuged at 2500 rpm for 15 minutes to separate the voided opacifying particles from the dispersion. The voided structure, with the TiO₂ pigment particles in air voids (nanorattles containing TiO₂), was confirmed by TEM (Fig. 1). Further analysis indicated that TiO₂/polymer weight ratio is 1:2.4 in the separated opacifying particles.

Example 3

[0028] The opacifying particles (nanorattles) from Example 2 were compared with a mixture of TiO₂ pigment particles and hollow particles in a coating for hiding. Opacifying particles from Example 2 (10g), which contained 8.7% TiO₂ and 20.7 % polymer, were mixed with 40g Rhoplex SG-30 from Dow Chemical, and then drawn down on a black LENETA chart with a 50 µm thick bar and dried at room temperature. Reflectance at 550 nm was measured with Oceanoptics USB2000 for comparison of hiding. In comparison, a first control paint was made by replacing the opacifying particles with a TiO₂ slurry of CR-826 while doubling the amount of TiO₂ by weight in the paint and a comparative paint was produced using the TiO₂ slurry with the pigment extender Ropaque Ultra (from The Dow Chemical Company). As shown in Table 1, the hiding of the paint with the opacifying particles from Example 2 was comparable to the control with double the amount of TiO₂ by weight percentage. In the comparative example, the total amount of TiO₂ was the same as in the paint with the opacifying particles and exhibited slightly improved hiding by the dried paint. However, as wet films, the opacifying particles provided significantly improved hiding over the comparative example as shown in Fig. 2.

Table 1

Test	Pigment particles	TiO ₂ wt % in paint	R% @550 nm
Control	TiO ₂ slurry	3.6	33.3
Example	Opacifying particles	1.8	36.6
Comparative Example	TiO ₂ slurry + Ropaque Ultra	1.8	48.7

[0029] Although the present invention has been described with reference to specific details of certain embodiments thereof, it is not intended that such details should be regarded as limitations upon the scope of the invention except insofar as they are included in the accompanying claims.

REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- [CN102311556 \[0004\]](#)

Patentkrav

1. Vandig dispersion af partikler, hvilken dispersion omfatter:

5 uorganiske partikler, der er mindst delvist indkapslede af polymeriserede indre skaller, hvor mindst en del af overfladerne af partiklerne omfatter et koblingsmiddel, der har vedhængende reaktive grupper, og hvor de indre skaller fremstilles ud fra en omsætning af koblingsmidlets vedhængende reaktive grupper med ethylenisk umættede monomerer; og

10 ydre polymerskaller, der mindst delvist indkapsler de indre polymere skaller, hvor de ydre polymerskaller er mindst delvist adskilte fra de indre polymerskaller, hvorved der defineres hulrum inden for de ydre polymerskaller.

2. Vandig dispersion ifølge krav 1, hvor koblingsmidlet omfatter et silankoblingsmiddel.

20 3. Vandig dispersion ifølge krav 1, hvor koblingsmidlets vedhængende reaktive grupper omfatter ethylenisk umættethed.

4. Vandig dispersion ifølge krav 1, hvor de ethylenisk umættede monomerer omfatter C₁-C₁₂-alkylestere af (meth)acrylsyre.

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5. Vandig dispersion ifølge krav 2, hvor silankoblingsmidlet omfatter en hydrolyserbar gruppe, som ved hydrolyse, omfatter en rest med frie hydroxylgrupper, der er til stede på overfladerne de uorganiske partikel.

30 6. Vandig dispersion ifølge krav 2, hvor de ydre polymerskaller omfatter polystyren.

7. Vandig dispersion ifølge krav 6, hvor de ydre skaller omfatter første ydre

skaller, som omfatter polystyren, og andre ydre skaller, der omfatter tværbundet polystyren, hvor de andre ydre skaller mindst delvist indkapsler de første ydre skaller.

5 **8.** Vandig dispersion ifølge krav 2, hvor de uorganiske partikler omfatter titandioxid.

9. Fremgangsmåde til fremstilling af en vandig dispersion af partikler, der er mindst delvist indkapslet af en polymer, hvilken fremgangsmåde omfatter:

10

at dispergere uorganiske partikler i en vandig fase;

at funktionalisere mindst en del af overfladerne af de uorganiske partikler med et koblingsmiddel, hvilket koblingsmiddel har vedhængende reaktive grupper;

15

at polymerisere de reaktive grupper på de funktionaliserede partikler med en første polymeriserbar komponent for at producere indre polymerskaller, der omgiver de uorganiske partikler;

20 at polymerisere en anden polymeriserbar komponent for at danne ydre polymerskaller, der mindst delvist indkapsler de indre polymerskaller; og

at producere hulrum i de ydre polymerskaller, således at de uorganiske partikler optages inden i hulrummene.

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10. Fremgangsmåden ifølge krav 9, hvor trinnet af producere hulrum omfatter (i) at ekspandere de indre polymerskaller og ydre polymerskaller og (ii) at krympe de ekspanderede indre polymerskaller for derved at danne hulrum inde i de ekspanderede ydre polymerskaller.

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11. Fremgangsmåden ifølge krav 10, hvor de uorganiske partikler, der er modtaget inden for hulrummene er mindst delvist indkapslet af de indre

polymerskaller.

12. Fremgangsmåden ifølge krav 9, hvilken fremgangsmåde yderligere omfatter at tværbinde de ekspanderede ydre skaller.

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13. Fremgangsmåden ifølge krav 12, hvor den anden polymeriserbare komponent omfatter styren, methylnmethacrylat og/eller acrylonitril.

14. Fremgangsmåden ifølge krav 9, hvor de ekspanderede ydre skaller, koblingsmidlet, den første polymeriserbare komponent, og de uorganiske partikler er defineret som i ethvert af kravene 1-8.

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15. Overtrækssammensætning, der omfatter den vandige dispersion ifølge krav 1.

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DRAWINGS

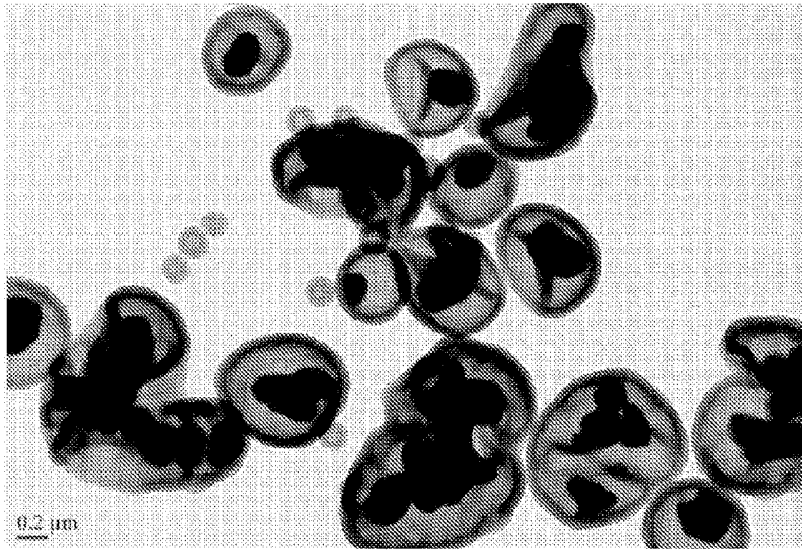


FIG. 1

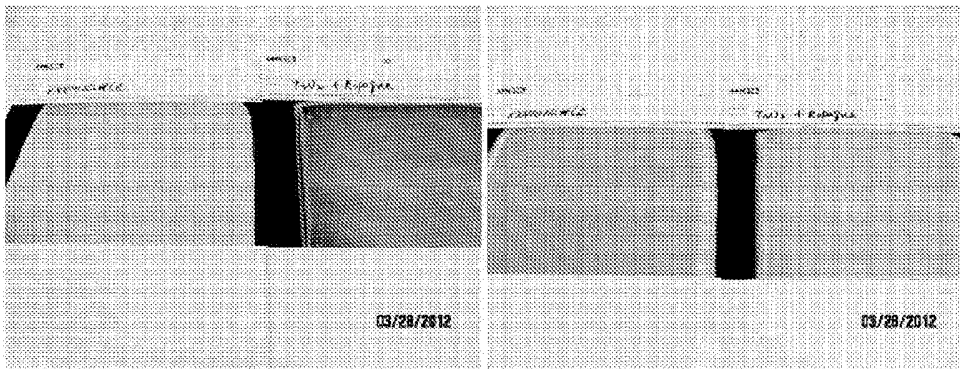


FIG. 2A

FIG. 2B