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(54) Title: MODIFIED NANOPARTICLES

(57) Abstract: The invention relates to modified nanoparticles based on pyrogenic silica which are produced by treating pyrogenic silica with compounds of the general formula (I) $\text{Me}(\text{OR})_4$, and/or of the general formula (II) $\text{Me}(\text{OCOR})_4$, wherein R¹ means an alkyl, aryl and/or aralkyl residue and Me means zirconium and/or titanium and to coating compositions containing the modified nanoparticles.



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Title of the Invention

Modified Nanoparticles

Field of the Invention

5 The invention relates to modified nanoparticles based on pyrogenic silica, to a process for the *production of the modified nanoparticles and to coating compositions with improved scratch resistance which* contain the modified nanoparticles.

Description of Related Art

10 Improving the mechanical stability, such as, scratch resistance, hardness and abrasion resistance, of coatings has long been a central task, in particular for automotive top coats. It is known to improve the above-stated properties by incorporating nanoparticles into clear coats or
15 top coats. The technical challenge here is to incorporate the necessary quantity of nanoparticles into the coatings in such a manner that the general range of properties of the coatings remains unaffected. For example, it should be ensured that properties such as, optical quality (transparency in clear coats or colour shade in pigmented coatings),
20 rheology, flow and adhesion are not negatively affected by the use of nanoparticles.

 EP 1 216 278, EP 1 195 416 and DE 10239424 describe, for example, differently structured and functionalised nanoparticles and the use thereof in coatings to impart scratch resistance.

25 WO 03102089 furthermore describes chemically modified nanoparticles, which are present as a dispersion in a film-forming binder. The nanoparticles are here modified by compounds of the general formula $\text{Si}(\text{OR})_3\text{-(CH}_2)_n\text{-Z}$, wherein Z represents a long-chain alkyl group, a fluorocarbon group or a silane group with at least 2 methyl groups. The
30 group Z is intended to ensure that the modified nanoparticles exhibit lower

compatibility towards the binders than do the unmodified particles and so preferably congregate at the surface of the coatings containing them.

DE 102 41 510 describes compositions prepared from agglomerated nanoparticle powders and organic binders. The
5 nanoparticles are here treated with compounds of the general formulae $\text{Si}(\text{OR}')_n\text{R}_{4-n}$, $\text{SiCl}_n\text{R}_{n-4}$, $(\text{R}'_m\text{R}''_{m-3}\text{Si}_2)\text{NH}$, $\text{Ti}(\text{OR}')_n\text{R}_{4-n}$ and $\text{Zr}(\text{OR}')_n\text{R}_{4-n}$, wherein R is a functional group which is attached directly via a C atom to the silicon, titanium or zirconium. In particular, the functional groups are those with unsaturated double bonds which, once the nanoparticles have
10 been incorporated into the unsaturated monomers forming the binder, are polymerised with the monomers and give rise to a crosslinked nanocomposite.

EP 1 166 283 describes coating compositions for metallic conductors, in particular wires, which exhibit improved partial discharge
15 resistance and which give rise to flexible coatings on the wires. These compositions contain particles based on an element-oxygen network with reactive and optionally, unreactive functions on the surface, wherein the unreactive functions are attached via the oxygen of the network.

It has not, however, proved possible hitherto to provide
20 nanoparticles or processes for incorporating nanoparticles into coatings which are capable of satisfactorily solving the above-stated technical problem.

There is accordingly still a requirement for nanoparticles which may be incorporated into coatings in a sufficient quantity to achieve improved
25 scratch resistance and hardness without having to make relatively large concessions with regard to other essential coating properties.

Summary of the Invention

The invention relates to modified nanoparticles based on pyrogenic
30 silica which are produced by treating pyrogenic silica with compounds of

the general formula I $\text{Me}(\text{OR}^1)_4$ and/or the general formula II $\text{Me}(\text{OCOR}^1)_4$. R^1 here means an alkyl, aryl and/or aralkyl residue. Me means zirconium and/or titanium. Me and R^1 can be each selected independently in formula I and II.

5 A preferred embodiment of the present invention starts from nanoparticles based on hydrophobic, silanised pyrogenic silica.

The invention accordingly also relates to modified nanoparticles based on silanised pyrogenic silica which are produced by treating silanised pyrogenic silica with compounds of the general formula I
10 $\text{Me}(\text{OR}^1)_4$ and/or the general formula II $\text{Me}(\text{OCOR}^1)_4$. R^1 and Me have the above-stated meaning.

It has surprisingly been found that, by using the nanoparticles according to the invention, increased quantities of active nanoparticles may be incorporated into coatings. It has proved possible to achieve a
15 considerable improvement in the dispersion and deagglomeration of the nanoparticles. This results in improved mechanical stability, in particular, improved scratch resistance of the coatings applied from these compositions. The existing advantageous properties of coatings which have not been modified with the nanoparticles are retained. The
20 nanoparticles according to the invention exhibit virtually neutral behaviour in the coating. The effect on the rheological behaviour of the coatings is minimal, the transparency of resultant clear coats is very good, the colour of pigmented coatings is unchanged and the surface structure of the coating is not impaired. There is also no discernible negative influence on
25 the development of hardness.

Detailed Description of the Embodiments

The invention will be explained in greater detail below.

Pyrogenic silica is here taken to mean highly disperse synthetic
30 silicas which have been produced by flame hydrolysis or with the

assistance of the arc process (for definition c.f. also RÖMPP Lexikon, Lacke und Druckfarben [Coatings and Printing Inks], 1998, p. 323). The production of pyrogenic silica is known to the person skilled in the art. Pyrogenic silica is obtainable as a commercial product.

5 According to the invention, nanoparticles based on pyrogenic silica, preferably based on silanised pyrogenic silica, are modified by treatment with the above-stated zirconium and/or titanium compounds of the general formula I and/or of the general formula II, wherein R^1 and Me have the above-stated meaning.

10 R^1 has the following meaning:

R^1 may be an alkyl residue, the alkyl residue comprising an optionally substituted linear or branched alkyl residue with 1-20, preferably with 1-12, particularly preferably with 1-6 C atoms. The alkyl residue may be substituted with any desired organic groups, for example, with acid
15 groups, hydroxyl groups and amino groups.

R^1 may likewise be an aryl residue, the aryl residue comprising aromatic hydrocarbon residues such as, for example, phenyl and naphthyl residues.

R^1 may also be an aralkyl residue, the aralkyl residue comprising
20 alkyl residues substituted by aryl groups, such as, benzyl residues, phenylalkyl residues with 1-10 C atoms in the alkyl residue, for example, phenylethyl residues. The alkyl residues may here also be substituted in the above-stated manner.

The residues R^1 in the general formula I and II preferably comprise
25 identical residues R^1 in one formula, but combinations of different residues R^1 in one formula may also be present.

R^1 is preferably a lower alkyl residue with 1-6 C atoms, such as, for example, a methyl, ethyl, propyl, isopropyl, butyl, isobutyl, pentyl or hexyl residue.

Zirconium and/or titanium compounds of the general formula I are preferably used.

Preferred compounds of general formula I are tetrapropyl titanate, tetrapropyl zirconate, tetrabutyl titanate, tetrabutyl zirconate, tetrapentyl titanate and tetrapentyl zirconate.

Examples of compounds of general formula II are zirconium and titanium esters of organic acids.

The modification of the nanoparticles with the zirconium and/or titanium compounds of the general formula I and/or II will now be described in greater detail. Both here and below, the above-described zirconium and/or titanium compounds of the general formula I and/or II will be designated compounds of the general formula I and/or II.

One possible production method involves initially introducing the compounds of the general formula I and/or II to be used for modification in one or more organic solvents. Organic solvents which may be used are conventional organic solvents for coatings, such as, for example, glycol ethers, such as, ethylene glycol dimethyl ether; propylene glycol dimethyl ether; glycol ether esters, such as, ethylene glycol monoethyl ether acetate, ethylene glycol monobutyl ether acetate, 3-methoxy-n-butyl acetate, diethylene glycol monobutyl ether acetate, methoxypropyl acetate, esters, such as, butyl acetate, isobutyl acetate, amyl acetate; ketones, such as, methyl ethyl ketone, methyl isobutyl ketone, cyclohexanone, isophorone, aromatic hydrocarbons (for example, with a boiling range of 136-180°C) and aliphatic hydrocarbons. The compounds of the general formula I and/or II may here be mixed with the organic solvent(s) by stirring, for example, at 10-100°C, in a suitable unit. The nanoparticles or nanoparticle agglomerates may then be sprinkled into the mixture obtained in this manner, preferably with stirring and in small portions. This may proceed for example, at 10-100°C over a few minutes to several hours.

It is, however, also possible initially to prepare a suspension of the nanoparticles in one or more organic solvents and then to add the compounds of the general formula I and/or II.

The resultant mixture may then be predispersed in a suitable unit, for example, a high-speed mixer, and dispersed down to a desired particle size in a suitable manner, for example, in a bead mill, by means of a jet disperser or by means of ultrasound. Dispersion is performed, for example, down to a primary particle size of 1-200 nm, preferably of 1-70 nm.

The nanoparticles are treated, for example, with 1-60 wt.%, preferably with 1-40 wt.%, most preferred with 1-20 wt.%, of the compounds of the general formula I and/or II, relative to the quantity of nanoparticles. The quantity of compounds of the general formula I and/or II to be used here vitally depends on the selected nanoparticles.

The preparation of the modified nanoparticles in the organic phase as described above is preferably used to prepare nanoparticles modified with compounds of the general formula I.

The nanoparticles based on pyrogenic silica to be used comprise conventional particles known to the person skilled in the art in the "nanometre" size range, for example, with an average primary particle size of 1 to 200 nm, preferably of 1 to 100 nm, which may contain functional groups, in particular hydroxyl groups, on the particle surface. Usable nanoparticles based on pyrogenic silica are obtainable as commercial products, for example, various grades under the name Aerosil[®] from Degussa, for example, Aerosil[®] R 90, R 130, R 150, R200, R300, R380, R812, R 9200, R 8200, under the name(s) Cab-O-Sil[®] LM 150, M-5, PTG, TS-610 and TS-530 from Cabot Corporation and under the name HDK N 20, HDK H 18, HDK T 40 and HDK T 30 from Wacker.

As has already been mentioned above, the nanoparticles to be modified according to the invention may, in a preferred embodiment,

already be silanised. On silanisation, a proportion of the OH groups of the silica particles is reacted with appropriate compounds, so rendering the nanoparticles water-repellent. Various options are known for silanisation. For example, the hydroxyl groups of the pyrogenic silica may be reacted
5 with chlorosilanes or also with alkyl (di)silazanes. Another option involves, as described in EP 0 672 731, reacting the hydroxyl groups of the pyrogenic silica with compounds of the general formula $(OR)_3SiC_nH_{2n+1}$, wherein $n = 10$ to 18 and R means an alkyl residue with 1 to 4 C atoms. Such silanised nanoparticles based on pyrogenic silica are likewise
10 commercially obtainable, for example, under the name Aerosil[®] R 711, R 805, R 972, R 974, R 202, R 812, R 7200, R 8200 and R 9200 from Degussa, HDK H 15, HDK H 18, HDK H 20 and HDK H 30 from Wacker and Cab-O-Sil[®] TS 610, TS 530 and TS 720 from Cabot Corporation.

In a further preferred embodiment according to the invention, the
15 nanoparticles may be modified with the compounds of the general formula I and/or II in combination with the treatment with silane compounds. The silane compounds may, for example, comprise compounds of the general formula $Si(OR^2)_nR^3_{4-n}$ with $n = 1, 2, 3$ or 4 , wherein R^2 has the meaning of R^1 , and R^3 represents any desired organic group, which is attached
20 directly to the silicon via a carbon atom. If n is 1 or 2, R^3 can be identical or different, if n is 2,3 or 4 R^2 can be identical or different. Preferably silane compounds with $n=4$ are used, wherein R^2 has the above stated meaning and can be identical or different.

Treatment with the silane compounds may proceed in parallel with
25 or preferably after treatment with the compounds of the general formula I and/or II. In any event, such treatment should proceed before the grinding process.

One embodiment according to the invention which is particularly preferred is that in which the obtained modified nanoparticles are
30 additionally incorporated into specific functional compounds and are then jointly predispersed and ground together with these functional compounds.

It is assumed that, when the nanoparticles are modified according to the invention with compounds of the general formula I and/or II, not all functional groups of these compounds, e.g., $-OR^1$ groups, are bound to the surface of the nanoparticles and free groups $-OR^1$ are still retained. It is
5 further assumed that, with the assistance of these still free groups, the nanoparticles may be bound to specific functional compounds, whether by chemical reaction or coordinative attachment.

The functional compounds comprise mono- and/or polyfunctional monomeric, oligomeric and/or polymeric compounds which contain
10 functional groups which are intended to be capable of binding to those reactive groups, e.g., OR^1 groups, of the nanoparticles which are still present. Suitable functional groups which may be considered are any desired organic groups with Lewis basicity, such as, for example, OH, COOH, amino, thiol, carbamate, imido, epoxide, isocyanate and/or ester
15 groups. Correspondingly, functionalised oligomeric or polymeric binders and/or other correspondingly functionalised compounds may be used. For example, hydroxy-functional or carboxy-functional binders and/or other hydroxy- or carboxy-functional compounds, for example, mono- or polyhydric longer-chain linear or branched alcohols, for example, mono- or
20 polyhydric aliphatic alcohols with 5 to 20 C atoms may be used.

Functionalised compounds which may particularly advantageously be used are such correspondingly functionalised binders, for example, hydroxy-functional binders, which are also a constituent or preferably the main constituent of the coating binder system into which the modified
25 nanoparticles are to be incorporated. Stable masterbatches containing nanoparticles are obtained, which may be directly incorporated into a coating system without any problems. Improved deagglomeration and thus dispersion of the nanoparticles in the coating system is obtained.

The nanoparticle masterbatches may, for example, be produced by
30 mixing and predispersing the modified nanoparticles with the functionalised compounds, in particular the functionalised binders, in a

high-speed mixer and then continuing dispersing in suitable units, for example, by means of a bead mill. The functionalised compounds, in particular the functionalised binders, may here initially be introduced and the modified nanoparticles added thereto or the modified nanoparticles
5 may initially be introduced and the functionalised compounds, in particular the functionalised binders, added thereto. At variance with these variants, it is, however, also possible, to perform the treatment/modification according to the invention of the nanoparticles with the compounds of the general formula I and/or II in the presence of the functionalised
10 compounds. The functionalised compounds can be present as organic solution or organic dispersion.

The nanoparticles and the functionalised compounds, in particular, the functionalised binders, may here be used in a ratio by weight of functionalised compound: modified nanoparticles of 99:1, preferably of
15 80:20 as an upper limit. The lower limit of the ratio by weight may be considered to be the ratio by weight of functionalised compound: modified nanoparticles corresponding to the critical pigment volume concentration (CPVC), i.e., the quantity of functionalised compound which is just sufficient adequately to wet a specific quantity of nanoparticles. It may, for
20 example, comprise a ratio by weight of functionalised compound: modified nanoparticles of 80:20 to 3:20.

Ideally, the functionalised compounds, in particular the functionalised binders, are here used in a quantity and the conditions selected such that as far as possible all the reactive groups still present on
25 the nanoparticle surface may be consumed by reaction with the corresponding functional groups of the binder. Any unreacted functional groups of the binder which are optionally still present may then, in the presence of an appropriate crosslinking agent in the coating composition, react with the reactive groups of the crosslinking agent and so participate
30 in the crosslinking process.

Dispersion of the nanoparticle masterbatches proceeds down to a desired particle size of for example, 1-200 nm, preferably 1-70 nm.

In the absence of water, the resultant nanoparticle masterbatches exhibit excellent storage stability within a wide temperature range, for example, from -15 to 80°C.

Self-evidently, all the above-described embodiments according to the invention may be used alone or combined with one another in any desired manner.

The present invention also relates to a process for the production of modified nanoparticles, comprising the following steps:

- I) providing nanoparticles based on pyrogenic silica, in particular silanised pyrogenic silica,
- II) treating the pyrogenic silica, in particular the silanised pyrogenic silica, with compounds of the general formula I $\text{Me}(\text{OR}^1)_4$ and/or of the general formula II $\text{Me}(\text{OCOR}^1)_4$, wherein R^1 means an alkyl, aryl and/or aralkyl residue and Me means zirconium and/or titanium,
- III) optionally, treating the pyrogenic silica, in particular the already silanised pyrogenic silica, with silane compounds of the general formula $\text{Si}(\text{OR}^2)_n\text{R}^3_{4-n}$ with $n = 1, 2, 3$ or 4 , wherein R^2 has the meaning of R^1 , and R^3 is any desired organic group, which is attached directly to the silicon atom via a carbon atom, jointly with the treatment in step II or after the treatment in step II and
- IV) optionally, incorporating the nanoparticles into functional monomeric, oligomeric and/or polymeric compounds, which contain functional groups with Lewis basicity, wherein modified nanoparticles obtained in step II or III may be incorporated into the functional compounds or wherein modification of the starting nanoparticles proceeds in step II and/or III in the presence of the functional compounds.

The functionalised compounds and the functional groups comprise those already stated above.

The present invention also relates to coating compositions which contain the above-described modified nanoparticles. In particular, the invention relates to coating compositions modified with nanoparticles, comprising

- A) at least one film-forming binder,
- B) optionally, at least one crosslinking agent for the binder,
- C) 0.5-40 wt.%, preferably 1-20 wt.%, relative to the quantity of film-forming binder A), of nanoparticles based on pyrogenic silica which are produced by treating pyrogenic silica with compounds of the general formula I $\text{Me}(\text{OR}^1)_4$ and/or of the general formula II $\text{Me}(\text{OCOR}^1)_4$, wherein R^1 means an alkyl, aryl and/or aralkyl residue and Me means zirconium and/or titanium, and
- D) optionally, organic solvents, water, pigments and fillers and conventional coating additives.

Selection of the film-forming binders A) and optionally present crosslinking agents B) in the coating compositions according to the invention is not subject to any particular restriction. Any desired binders conventionally usable in coatings production may be used. The binders A) may comprise, for example, polyesters, polyurethanes, (meth)acrylic copolymers, epoxy resins and mixtures as well as hybrids of the stated binders. The listing is, however, not intended to constitute a limitation. Polymers other than those stated may also be used. The binders A) may comprise functional groups. When functional groups are present, the binders preferably have at least two functional groups per molecule.

Non-limiting examples of functional groups which may be present in the binders A) are epoxy groups, carboxyl groups, hydroxyl groups, amino groups, isocyanate groups, olefinically unsaturated groups, alkoxy silane

groups. The functional groups may here also be present in blocked form, for example, as blocked hydroxyl groups, isocyanate groups or amino groups.

Production of the optionally functionalised binders A) is sufficiently
5 known to the person skilled in the art and requires no explanation.

The binders A) may be self- or externally crosslinking. They may, however, also comprise physically drying binders.

In addition to the binders A), the coating compositions according to the invention may contain crosslinking agents B) which are capable of
10 entering into a crosslinking reaction with functional groups of the binders A).

Selection of the crosslinking agents is guided by the functional groups present in the binders A), i.e., the crosslinking agents are selected such that they exhibit a reactive functionality complementary to the
15 functionality of the binders, wherein the functional groups may react with one another by free-radical polymerisation and/or by addition and/or condensation.

Examples of addition reactions between binders A) and crosslinking agents B) are the ring-opening addition of an epoxy group onto a carboxyl
20 group with formation of an ester group and a hydroxyl group, the addition of a hydroxyl and/or primary and/or secondary amino group onto an isocyanate group with formation of a urethane group and/or urea group, the addition of a primary and/or secondary amino group and/or CH-acidic group onto an alpha,beta-unsaturated carbonyl group, in particular
25 (meth)acryloyl group, the addition of a primary and/or secondary amino group onto an epoxy group. Examples of condensation reactions between (A) and (B) groups are the reaction of a hydroxyl and/or primary and/or secondary amino group with a blocked isocyanate group with formation of a urethane group and/or urea group and elimination of the blocking agent,
30 the reaction of a hydroxyl group with an n-methylol group with elimination

of water, the reaction of a hydroxyl group with an n-methylol ether group with elimination of the etherification alcohol, the transesterification reaction of a hydroxyl group with an ester group with elimination of the esterification alcohol, the transurethanisation reaction of a hydroxyl group with a
5 carbamate group with elimination of alcohol, the reaction of a carbamate group with an n-methylol ether group with elimination of the etherification alcohol. Examples of functional groups (A) and (B) capable of reacting by means of free-radical polymerisation are olefinically unsaturated groups, for example, vinyl groups, allyl groups, in particular (meth)acryloyl groups.

10 Providing that they are mutually compatible, two or more complementary functionalities may simultaneously be present in a binder curable by addition and/or condensation reactions, such that curing may proceed via two or more different reactions of the types stated above by way of example.

15 The coating compositions according to the invention contain 0.5-40 wt.%, preferably 1-20 wt.%, relative to the quantity of film-forming binder A), of the above-described modified nanoparticles based on pyrogenic silica. All the previously described possible embodiments of the nanoparticles are, of course, also included.

20 The modified nanoparticles may additionally, as already explained above, particularly preferably be incorporated into functionalised compounds. It is particularly advantageous here, as has also already been explained above, to use as the functionalised compounds correspondingly functionalised binders, for example, hydroxy-functional
25 binders, which are also a constituent or preferably the main constituent of the binder system of the coating composition. Functionalised binders which may accordingly be considered in principle are also those which have already been described above for use in the coating compositions.

The nanoparticle masterbatches may here be produced directly
30 during base formulation of the coating composition and be incorporated into the coating composition, but they may, however, also be produced

and stored as semifinished products and then incorporated into finished coating compositions when required. Generally, the modified nanoparticles can be incorporated into the coating compositions as such or in the form of the above described masterbatches. Preferably, the nanoparticles and nanoparticle masterbatches are prepared in the organic phase, as described above, and then incorporated in a suitable manner into waterbased or solventbased coating compositions,

The coating compositions according to the invention may contain pigments and/or fillers and conventional coating additives in conventional coating quantities.

The coating compositions according to the invention may contain conventional coating organic solvents and/or water. Aqueous coating compositions may, for example, assume emulsion form. The emulsified state may here be achieved by the addition of external emulsifiers or the systems may contain groups having a self-emulsifying action in water, for example, ionic groups.

The coating compositions according to the invention may be formulated as one- or two-component coating systems, depending on the binders and crosslinking agents selected. Preferred examples of one- or two-component coating compositions are those which contain hydroxy-functional binders, such as, hydroxy-functional (meth)acrylic copolymers, polyester resins and/or polyurethane resins as binder and, as crosslinking agents, triazine-based components, for example, tris(alkoxycarbonylamino)triazine, which crosslink with the hydroxyl groups of the binders to form ether and/or ester groups, amino resins, in particular, melamine resins, and/or transesterification crosslinking agents and/or free or blocked polyisocyanates. Further preferred one- or two-component coating systems are those which contain epoxy-functional binders in combination with carboxy-functional crosslinking agents.

The coating compositions according to the invention may be used in single layer and multilayer coatings. When used in multilayer coatings,

they are preferably used for the production of the external coating layer of a multilayer structure. The external coating layer may comprise pigmented or unpigmented coating layers, which are produced, for example, from a pigmented top coat, a clear coat or a sealing coat.

5 The coating compositions may be applied by conventional application methods onto any desired substrates, for example, of metal, plastics, wood or glass. Examples of application methods are brushing, roller application, knife coating, dipping, but in particular spraying. After application, the coating layers applied from the coating compositions
10 according to the invention are dried or cured after an optional flash-off phase. Depending on the composition of the coating compositions according to the invention, drying/curing may proceed at room temperature or be forced at elevated temperatures, for example, 40 to 80°C, or by stoving (baking) at higher temperatures, for example, 80 to 220°C. In the
15 case of radiation-curable coating compositions according to the invention, curing proceeds by exposure to high-energy radiation, for example, UV radiation.

 The nanoparticles according to the invention, in particular the nanoparticles incorporated into functionalised compounds, such as,
20 functionalised binders, in the form of nanoparticle masterbatches, exhibit no compatibility problems between the nanoparticles and further binder or coating composition constituents, whether during production, storage or during or after the incorporation thereof into coating compositions or during or after application of the coating compositions. It has proved possible to
25 achieve an improvement in the dispersion and deagglomeration of the nanoparticles. The masterbatches and coating compositions containing the nanoparticles are stable in storage. Thanks to the use of specific compounds of the general formula I and/or II for modifying the nanoparticles, it has proved possible to minimise the risk of secondary or
30 competitive reactions of the nanoparticles modified according to the invention with one another, for example, before incorporation into the

stated functionalised compounds, and/or with the binders and/or other coating components. The coating layers applied from the coating compositions according to the invention have for example, very good surface qualities, in particular for example, elevated scratch resistance and
5 chemical resistance. These advantageous properties may be achieved without having to accept impairment of other important properties of the coating composition or of the coatings obtained therefrom, such as, for example, colour matching, transparency, rheological properties, flow, adhesion and development of hardness.

10 The nanoparticles according to the invention or coating compositions containing them may be used in any application where high quality coatings with good mechanical resistance, in particular elevated scratch resistance, are required. They may accordingly in particular be used in vehicle and industrial coating.

15

The invention will be further described by reference to the following examples.

Examples

Example 1:

20 Determining the optimal amount of $Ti(OPr)_4$ for modifying pyrogenic silica nano-particles

HDK® T30 from Wacker has been used as pyrogenic silica nano-particles. HDK® T30 is a synthetic, hydrophilic amorphous silica, produced
25 via flame hydrolysis.

5 g of HDK® T30 were stirred in a beaker with 100 g butylacetate and 2,55 g $Ti(OPr)_4$ (98%, 2,55 g \pm 8,83 mmol, from Aldrich). After 10 min of stirring with a Dispermat at 200-2000 rpm the silica was filtered off by a commercially available 0,45 μm filter (Millex-LCR, Millipore). The filtrate

was investigated on excess of $Ti(OPr)_4$ by AAS (Atom Absorptions Spectroscopy).

The amount of titanate to modify the surface of the silica was investigated as 35,3 % by weight on silica.

5 Pr = Propyl

Preparation of titanate modified pyrogenic silica paste

35,7 g (98%, 35,7 g 0,123 mol) $Ti(OPr)_4$ were added to 371,0 g of a usual hydroxy-functional polyester binder (P) while stirring with a Dispermat at 100 rpm. 99,1 g HDK® T30 were added to the solution together with 300 g butylacetate. The mixture was stirred for 10 min at 100-1000 rpm.

The grinding process was done in a PM1 pear mill (Drais) with 1300 g of 1mm SAZ pearls (Garbe,Lahmeyer&Co. AG) at 2800 rpm at 60°C. After 3h the modified silica containing paste was obtained.

(P): 70 wt-% solution of a polyester polyol composed of neopentyl glycol, trimethylolpropane, hexahydrophthalic anhydride and coconut fatty acid, with an acid value of 13 mg KOH/g and a hydroxyl value of 136 mg KOH/g, a calculated hydroxyl functionality of 3.8 and a calculated molecular mass of 1500 in Solvesso® 100.

Comparative Example 1:

Preparation of an unmodified pyrogenic silica paste

470,6 g of a the hydroxy-functional polyester binder above were mixed with 71,3 g butylacetate and 58,14 g of HDK® T30. The mixing and grinding procedure was done under the same conditions as for the modified pyrogenic silica paste in Example 1.

Example 2:

Preparation of a clear coat containing the modified silica paste and comparative clear coat containing the unmodified silica paste

The titanate modified silica paste of Example 1 was incorporated into a usual 2K solvent-based polyester clear coat in such amounts, that a content of modified silica particles of 10 % is achieved, based on solid binder in the clear coat. The base clear coat was prepared by mixing the following ingredients:

Hydroxy-functional polyester (P)	70
Light stabilizer of the HALS type	1.4
UV absorber based on benzotriazole	1.4
Commercial levelling agent (silicone oil)	0.5
Ethoxypropyl acetate	9
Butyl diglycol acetate	1
Solvesso® 100	10.2
Solvesso® 150	2
Butyl acetate	4.5

10 Accordingly the unmodified silica paste of Comparative Example 1 was incorporated into the base clear coat in such amounts that a content

of silica particles of 10 % is achieved, based on solid binder in the clear coat.

The so obtained clear coat and the comparative clear coat (P) were mixed each with the following hardener solution in amounts that an equivalent ratio of OH : NCO of 1:1.2 results.

Hardener Solution: Mixture of 81 g of hexamethylene diisocyanate isocyanurate (Desmodur 3390; Bayer), 9.5 g of Solvesso® 100 and 9.5 g of butyl acetate

The clear coat and the comparative clear coat compositions obtained by mixing were then applied by electrostatic spraying in a dry layer thickness of 35 µm to test panels provided with an aqueous base coat layer of a usual black waterborne base coat (Base coat Brilliant Black of DuPont). After a 15 minute flash-off at room temperature, the clear coat coating layer was baked for 25 minutes at 140°C (object temperature).

15 Results

Table 1 gives results of technological tests carried out on the coatings.

Table 1: Technological results

		Clear coat according to invention	Comparative Clear coat
Appearance	Gloss (20°)	85	83
	Haze	10	13
Hardness	Fischer	129	127

	hardness (N/mm ²)		
Amtec	Initial gloss	85	83
(Amtec Kistler GmbH)	Gloss after strain	51	42
	Gloss after reflow	76,9	61,4

As can be seen in Table 1 gloss and haze values of the modified silica nanoparticle containing clear coat are superior to the comparative clear coat containing unmodified silica. Also technological values such as

5 Amtec test results are improved by using surface modified silica particles. In particular remarkably improved results have been achieved with regard to gloss after strain and after reflow.

Test Methods:

10 Haze and Gloss values measured by Micro-Haze plus (Byk-Gardner) according to DIN 67530 (Gloss) and DIN EN ISO 13803 (Haze)

Fischer hardness measured by Fischerscope H 100 (Fischer GmbH und CO.KG) according to DIN EN ISO 14577-1

15 Amtec: according to DIN 55668; Scratching was carried out using the laboratory-scale Amtec Kistler car wash (cf. Th. Klimmasch and Th. Engbert, Development of a uniform laboratory test method for assessing the car wash resistance of automotive top coats, in DFO Proceedings 32, pages 59 to 66, Technologie-Tage, Proceedings of the Seminar on 29.

and 30.4.97 in Cologne, Published by Deutsche Forschungsgesellschaft für Oberflächenbehandlung e.V., Adersstraße 94, 40215 Düsseldorf.

The residual gloss in % was measured in each case directly after strain and after one hour's reflow at an angle of illumination of 20°.

Claims

What is claimed is:

1. Modified nanoparticles comprising pyrogenic silica which are produced by treating pyrogenic silica with compounds of the general formula I $\text{Me}(\text{OR}^1)_4$ and/or of the general formula II $\text{Me}(\text{OCOR}^1)_4$; wherein R^1 is an alkyl, aryl and/or aralkyl residue and Me is zirconium and/or titanium.
2. Modified nanoparticles according to claim 1, wherein R^1 is a residue selected from a group consisting of optionally substituted linear or branched alkyl residue with 1-20 C atoms, phenyl residue, naphthyl residue, benzyl residue and phenylalkyl residue with 1-10 C atoms in the alkyl residue.
3. Modified nanoparticles according to claim 1, wherein R^1 is an alkyl residue with 1-6 C atoms.
4. Modified nanoparticles according to any one of claims 1 to 3, wherein silanised pyrogenic silica is used.
5. Modified nanoparticles according to any one of claims 1 to 4, wherein treatment of the nanoparticles with the compounds of the general formula I and/or II proceeds in combination with the treatment with silane compounds of the general formula $\text{Si}(\text{OR}^2)_n\text{R}^3_{4-n}$, wherein $n = 1, 2, 3$ or 4 , wherein R^2 has the meaning of R^1 and R^3 represents any desired organic group which is attached directly to the silicon via a carbon atom.
6. Modified nanoparticles according to any one of claims 1 to 5, wherein the nanoparticles are produced by treating pyrogenic silica with 1-60 wt.-% of compounds of the general formula I and/or II, relative to the quantity of the nanoparticles.
7. Modified nanoparticles according to any one of claims 1 to 6, wherein the modified nanoparticles are incorporated into mono- and/or polyfunctional monomeric, oligomeric and/or polymeric compounds.

8. Modified nanoparticles according to claim 7, wherein the modified nanoparticles are incorporated into hydroxy-functional and/or carboxy-functional binders and/or other hydroxy- or carboxy-functional compounds.
9. A coating composition comprising the modified nanoparticles according to any one of claims 1 to 8.
10. A coating composition according to claim 9 comprising the following components:
 - A) at least one film-forming binder,
 - B) optionally, at least one crosslinking agent for the binder,
 - C) 0.5-40 wt.%, relative to the quantity of film-forming binder A), of modified nanoparticles according to any one of claims 1 to 8, and
 - D) optionally, organic solvents, water, pigments, fillers and/or conventional coating additives.
11. A coating composition according to claim 10, comprising 1-20 wt.% of the modified nanoparticles C), relative to the quantity of film-forming binder A).
12. A coating composition according to any one of claims 9 to 11, wherein it comprises a transparent clear coat or a pigmented coat.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2006/032731

A. CLASSIFICATION OF SUBJECT MATTER			
INV.	C08K3/36 C09D7/12 ADD. C08L75/06	C08K5/057 C09D151/10	C08K5/098 C08K9/04 C09C1/30
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) C08F C08K C08L C09C C09D			
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 practical, search terms used) EPO-Internal, WPI Data, CHEM ABS Data			
C. DOCUMENTS CONSIDERED TO BE RELEVANT			
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<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C.		<input checked="" type="checkbox"/> See patent family annex.	
* Special categories of cited documents :			
"A" document defining the general state of the art which is not considered to be of particular relevance		"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	
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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)		"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.	
"O" document referring to an oral disclosure, use, exhibition or other means		"&" document member of the same patent family	
"P" document published prior to the international filing date but later than the priority date claimed			
Date of the actual completion of the international search 22 November 2006		Date of mailing of the international search report 28/11/2006	
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer Meiners, Christian	

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International application No

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C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

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