



(86) Date de dépôt PCT/PCT Filing Date: 2011/03/05  
(87) Date publication PCT/PCT Publication Date: 2011/12/29  
(45) Date de délivrance/Issue Date: 2018/05/01  
(85) Entrée phase nationale/National Entry: 2012/12/20  
(86) N° demande PCT/PCT Application No.: EP 2011/001098  
(87) N° publication PCT/PCT Publication No.: 2011/160733  
(30) Priorité/Priority: 2010/06/25 (EP10167305.1)

(51) Cl.Int./Int.Cl. *B01J 13/16* (2006.01)  
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(54) Titre : PROCEDE DE PRODUCTION DE MICROCAPSULES  
(54) Title: PROCESS FOR PRODUCING MICROCAPSULES

**(57) Abrégé/Abstract:**

The application describes a process for producing microcapsules which contain a shell made of polyurea and which surround in their interior a water-insoluble oil, where the shell is obtained by the reaction of two structurally different diisocyanates in emulsion form.

## (12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau(43) International Publication Date  
29 December 2011 (29.12.2011)(10) International Publication Number  
**WO 2011/160733 A1**(51) International Patent Classification:  
**B01J 13/16** (2006.01)

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PE, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(21) International Application Number:  
**PCT/EP2011/001098**(22) International Filing Date:  
5 March 2011 (05.03.2011)(25) Filing Language:  
English(26) Publication Language:  
English(30) Priority Data:  
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(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

## Declarations under Rule 4.17:

— of inventorship (Rule 4.17(iv))

## Published:

— with international search report (Art. 21(3))

WO 2011/160733 A1

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(57) Abstract: The application describes a process for producing microcapsules which contain a shell made of polyurea and which surround in their interior a water-insoluble oil, where the shell is obtained by the reaction of two structurally different diisocyanates in emulsion form.

**„Process for producing microcapsules“**

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The application relates to a process for producing microcapsules.

Microcapsules are powders or particles which consist of 10 a core and a wall material surrounding the core, wherein the core is a solid, liquid or gaseous substance which is surrounded by the solid, generally polymeric, wall material. They may be solid, i.e. consist of a single material. Microcapsules have on 15 average a diameter from 1 to 1000 µm.

A multitude of shell materials is known for producing microcapsules. The shell can consist either of natural, semisynthetic or synthetic materials. Natural shell materials are, for example, gum arabic, agar agar, agarose, maltodextrins, alginic acid or its salts, e.g. sodium alginate or calcium alginate, fats and fatty acids, cetyl alcohol, collagen, chitosan, lecithins, gelatin, albumin, shellac, polysaccharides, such as 20 starch or dextran, polypeptides, protein hydrolyzates, sucrose and waxes. Semisynthetic shell materials are inter alia chemically modified celluloses, in particular cellulose esters and cellulose ethers, e.g. cellulose acetate, ethyl cellulose, hydroxypropyl- 25 cellulose, hydroxypropylmethylcellulose and carboxymethylcellulose, and also starch derivatives, in particular starch ethers and starch esters. Synthetic shell materials are, for example, polymers such as 30 polyacrylates, polyamides, polyvinyl alcohol or polyvinylpyrrolidone.

Depending on the type of shell material and the production process, microcapsules are formed in each

case with different properties as far as diameter, size distribution and physical and/or chemical properties are concerned.

5 There is therefore a continuing need to develop novel production processes in order to be able to provide microcapsules with tailored properties.

10 A first subject matter of the present application is therefore directed to a process for producing microcapsules which contain a shell and a core made of a water-insoluble liquid material, where an aqueous solution of a protective colloid and a solution of a mixture of at least two structurally different at least difunctional isocyanates (A) and (B) in a 15 water-insoluble liquid are brought together until an emulsion is formed, to which an at least difunctional amine is then added, and which is then heated to temperatures of at least 60°C until the microcapsules are formed, wherein the isocyanate (B) is selected from the anionically modified 20 isocyanates or from polyethylene oxide-containing isocyanates or mixtures of these types and the isocyanate (A) is uncharged, but is not a polyethylene-containing isocyanate, with the proviso that during the production of the 25 microcapsules the weight ratio between the isocyanates (A) and (B) is in the range from 10:1 to 1:10.

30 A second object matter of the present application is directed to a perfume-free microcapsule comprising a liquid core of a water-insoluble liquid, and a shell of a reaction product of at least two different at least difunctional isocyanates (A) and (B), where (A) is uncharged and is not a polyethylene oxid-containing isocyanate and where the isocyanate (B) must be an anionically modified isocyanate, and an at least

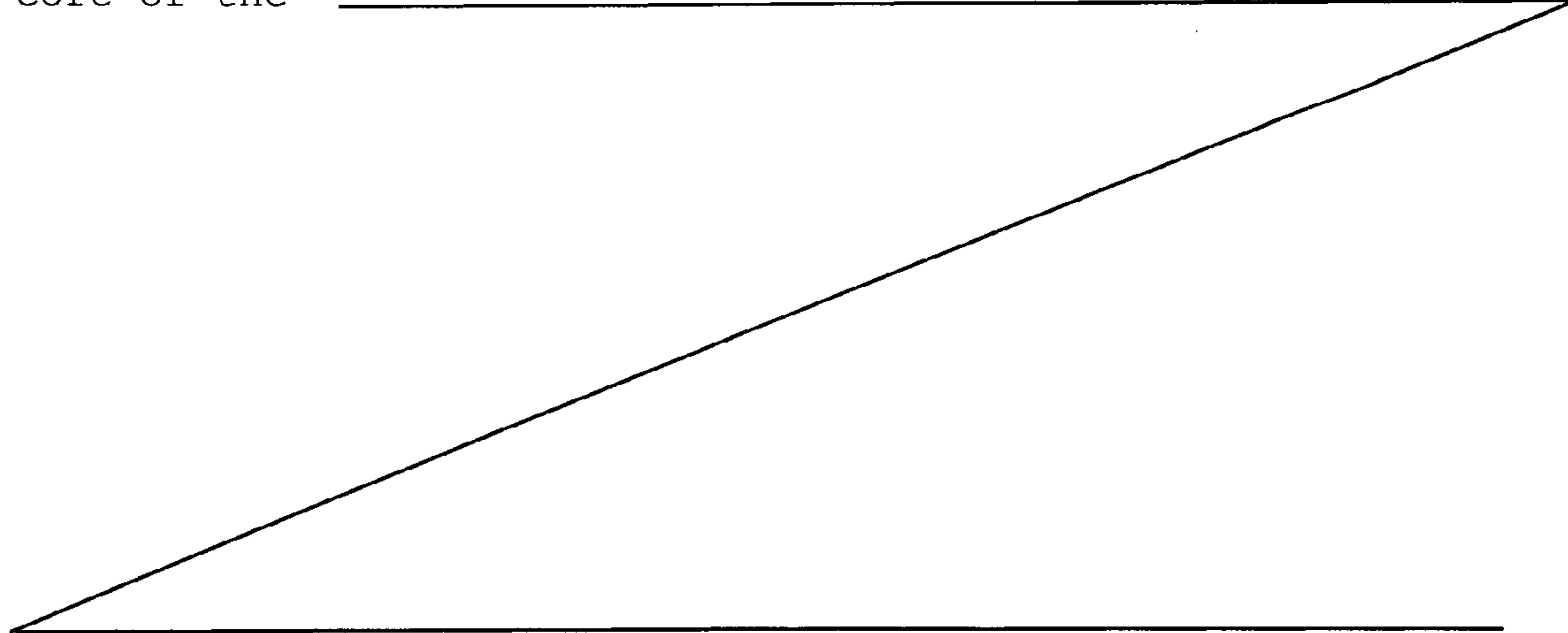
difunctional amine, with the proviso that during the production of the microcapsules the weight ratio between the isocyanates (A) and (B) is in the range from 10:1 to 1:10.

5 A third subject matter of the present application is directed to the use of microcapsules as defined herein for finishing of textiles, papers or nonwovens and for use in cosmetic, pharmaceutical, laundry and/or cleaning compositions.

10 The process has the advantage that microcapsules of a pre-given size or size distribution can be produced in a targeted manner, it being possible here to produce in particular relatively small microcapsules with diameters from 10 to 15 60 µm. Moreover, capsules with greater mechanical stability are obtained. Here, in particular those capsules are obtained the shells of which have only a low permeability to the liquid ingredients.

20 In principle, an aqueous solution of the protective colloid is always produced, and for this the isocyanates (A) and (B) are dissolved in the water-insoluble liquid which later forms the core of the

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microcapsules; the amine components are then added and the mixture is heated until an emulsion is formed. The temperature for the reaction of the isocyanates with the amine components must be at least 60°C, but better 5 70°C, but preferably 75 to 90°C and in particular 85 to 90°C, in order to ensure sufficiently rapid reaction progress.

Here, it may be preferred to increase the temperature in stages (e.g. in each case by 10°C) until then, 10 following completion of the reaction, the dispersion is cooled to room temperature (21°C). The reaction time typically depends on the amounts and temperatures used. Usually, however, the elevated temperature for forming the microcapsules is established between ca. 60 minutes 15 to 6 h or up to 8 h.

According to the present teaching, the addition of the amine also preferably takes place with the input of energy, e.g. by using a stirring apparatus.

20 In order to form an emulsion in the present process, the respective mixtures are emulsified by processes known to the person skilled in the art, e.g. by introducing energy into the mixture through stirring 25 using a suitable stirrer until the mixture emulsifies. The pH is preferably adjusted using aqueous bases, preference being given to using sodium hydroxide solution (e.g. 5% strength by weight).

30 It is essential to the process that at least two structurally different isocyanates (A) and (B) are used. These can be added in the form of a mixture or separately from one another in the process to the aqueous premix (1) containing the protective colloid 35 and are then emulsified and reacted with the amine. It is also conceivable to meter in both mixtures of (A) and (B), and also the individual isocyanates (A) and (B) separately at different times.

In one preferred embodiment, the process is carried out as follows:

- 5 (a) a premix (I) is prepared from water and a protective colloid;
- (b) this premix is adjusted to a pH in the range from 5 to 12;
- 10 (c) a further premix (II) is prepared from the liquid, water-insoluble material together with the isocyanates (A) and (B);
- (d) the two premixes (I) and (II) are brought together until an emulsion is formed and
- (e) the at least difunctional amine is then metered into the emulsion from step (d) and
- 15 (f) the emulsion is then heated to temperatures of at least 60°C until the microcapsules are formed.

It may be advantageous to adjust the pH in step (b) to values from 8 to 12. Of suitability here are aqueous bases, preferably aqueous sodium hydroxide solution. The formation of the emulsion in step (d), but also step (e) is preferably ensured by using a suitable stirrer.

25

Another likewise preferred embodiment envisages that

- (a) a premix (I) is prepared from water and a protective colloid;
- (b) this premix is adjusted to a pH in the range from 5 to 12;
- 30 (c) a further premix (II) is prepared from a water-insoluble material that is liquid at 21°C together with the isocyanate (A);
- (d) an emulsion is formed from the premixes (I) and (II) by stirring and to this
- 35 (e) is added the second isocyanate (B), and then the pH of the emulsion is adjusted to a value from 5 to 10;

- (f) and then the at least difunctional amine is metered into the emulsion from step (e) and
- (g) then heated to temperatures of at least 60°C until the microcapsules are formed.

5

In this procedure, the isocyanates (A) and (B) are added separately to the protective colloid before the addition of the amine and the reaction to give the microcapsules takes place. The formation of the 10 emulsion - like the mixing in step (e) also takes place here preferably by using a stirring apparatus.

The pH in step (e) is preferably adjusted to values from 7.5 to 9.0. For step (b), the value can likewise 15 be adjusted from 8 to 12. Of suitability for this purpose are in particular aqueous bases, preferably aqueous sodium hydroxide solution.

#### Microcapsules

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Within the context of the present teaching, the microcapsules have a shell made of a reaction product of at least two different, at least difunctional isocyanates with amines, preferably with polyamines. 25 The reaction is a polycondensation between the isocyanates and the amines, which leads to a polyurea derivative.

The microcapsules are present in the form of aqueous 30 dispersions, the weight fraction of these dispersions in the capsules being preferably between 15 and 45% by weight and preferably 20 to 40% by weight. The microcapsules have an average diameter in the range from 1 to 500 µm and preferably from 1 to 50 µm or from 35 5 to 25 µm.

The microcapsules contain water-insoluble liquid or solid contents, e.g. an oil. The fraction of this oil can vary in the range from 10 to 95% by weight, based

on the weight of the capsules, where fractions from 70 to 90% by weight may be advantageous. As a result of the process, capsules are obtained which typically have core/shell ratios (w/w) from 20:1 to 1:10, preferably 5 from 5:1 to 2:1 and in particular from 4:1 to 3:1.

The microcapsules which are produced by the present process are preferably free from formaldehyde.

10 Protective colloid

During the reaction between the isocyanates and the amines, a protective colloid must be present. This is preferably a polyvinylpyrrolidone (PVP). Protective colloids are polymer systems which, in suspensions or 15 dispersions, prevent a clumping together (agglomeration, coagulation, flocculation) of the emulsified, suspended or dispersed substances. During solvation, protective colloids bind large amounts of water and in aqueous solutions produce high viscosities 20 depending on the concentration. Within the context of the process described herein, the protective colloid may also have emulsifying properties. The aqueous protective colloid solution is likewise preferably prepared with stirring.

25 The protective colloid may be, but does not have to be, a constituent of the capsule shell, with amounts from 0.1 to at most 15% by weight, but preferably in the range from 1 to 5% by weight and in particular from 1.5 to 3% by weight, based on the weight of the capsules, 30 being possible here.

Isocyanates

Isocyanates are N-substituted organic derivatives (R-N=C=O) of isocyanic acid (HNCO) tautomeric in the 35 free state with cyanic acid. Organic isocyanates are compounds in which the isocyanate group (-N=C=O) is bonded to an organic radical. Polyfunctional isocyanates are those compounds with two or more isocyanate groups in the molecule.

According to the invention, at least difunctional, preferably polyfunctional, isocyanates are used, i.e. all aromatic, alicyclic and aliphatic isocyanates are 5 suitable provided they have at least two reactive isocyanate groups.

The suitable polyfunctional isocyanates preferably contain on average 2 to at most 4 NCO groups. 10 Preference is given to using diisocyanates, i.e. esters of isocyanic acid with the general structure  $O=C=N-R-N=C=O$ , where R' here is aliphatic, alicyclic or aromatic radicals.

Suitable isocyanates are, for example, 1,5-naphthylene 15 diisocyanate, 4,4'-diphenylmethane diisocyanate (MOI), hydrogenated MDI (H12MDI), xylene diisocyanate (XDI), tetramethylxylo diisocyanate (TMXDI), 4,4'-diphenyl- 20 dimethylmethane diisocyanate, di- and tetraalkyl-diphenylmethane diisocyanate, 4,4'-dibenzyl diisocyanate, 1,3-phenylene diisocyanate, 1,4-phenylene diisocyanate, the isomers of tolylene diisocyanate (TDI), optionally in a mixture, 1-methyl-2,4-diisocyanatocyclohexane, 1,6-diisocyanato-2,2,4-trimethylhexane, 1,6-diisocyanato-2,4,4-trimethylhexane, 1-iso- 25 cyanatomethyl-3-isocyanato-1,5,5-trimethylcyclohexane, chlorinated and brominated diisocyanates, phosphorus-containing diisocyanates, 4,4'-diisocyanatophenyl-perfluoroethane, tetramethoxybutane 1,4-diisocyanate, butane 1,4-diisocyanate, hexane 1,6-diisocyanate (HDI), 30 dicyclohexylmethane diisocyanate, cyclohexane 1,4-diisocyanate, ethylene diisocyanate, phthalic acid bisisocyanatoethyl ester, also polyisocyanates with reactive halogen atoms, such as 1-chloromethylphenyl 2,4-diisocyanate, 1-bromomethylphenyl 2,6-diisocyanate, 35 3,3-bischloromethyl ether 4,4'-diphenyldiisocyanate. Sulfur-containing polyisocyanates are obtained, for example, by reacting 2 mol of hexamethylene diisocyanate with 1 mol of thiodiglycol or dihydroxydihexyl sulfide. Further suitable diisocyanates are

trimethylhexamethylene diisocyanate, 1,4-diisocyanato-butane, 1,2-diisocyanatododecane and dimer fatty acid diisocyanate.

5 One essential feature of the present process is the obligatory use of two structurally different isocyanates (A) and (B).

10 Suitable isocyanates of type (A) are at least difunctional compounds (i.e. compounds containing at least two isocyanate groups  $-N=C=O$ ).

15 Typical representatives may be hexamethylene diisocyanate (HDI), or derivatives thereof, e.g. HDI biuret (commercially available e.g. as Desmodur N3200), HDI trimers (commercially available as Desmodur N3300) or else dicyclohexylmethane diisocyanates (commercially available as Desmodur W). Toluene 2,4-diisocyanate or diphenylmethane diisocyanate is likewise suitable.

20 The second isocyanate of type (B) is structurally different from the isocyanate of type (A) and specifically the isocyanate of type (B) must either be an anionically modified isocyanate or a polyethylene oxide-containing isocyanate (or any desired mixtures of 25 these two isocyanate types).

25 The anionically modified isocyanates are known per se. Preferably, these isocyanates of type (B) contain at least two isocyanate groups in the molecule. One or more sulfonic acid radicals are preferably present as 30 anionic groups. Preferably, isocyanates of type (B) are selected which are oligomers, in particular trimers, of hexane 1,6-diisocyanate (HDI). Commercial products of these anionically modified isocyanates are known, for example, under the brand Bayhydur (Bayer), e.g. 35 Bayhydur XP.

Polyethylene oxide-containing isocyanates (with at least two isocyanate groups) are also known and are described, e.g. in US 5,342,556. Some of these

isocyanates are self-emulsifying in water, which may be advantageous within the context of the present process since it may be possible to dispense with a separate emulsifying step.

5

The weight ratio of the two isocyanates (A) and (B) is adjusted preferably in the range from 10:1 to 1:10, but in particular in the range from 5:1 to 1:5 and in particular in the range from 3:1 to 1:1.

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It is also possible to use mixtures of different isocyanates of types (A) and (B). Besides the isocyanates (A) and (B), further isocyanates can also additionally be used in the process according to the 15 invention.

Preferably, however, only anionically modified isocyanates are used as component (B) in the present process.

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#### Amines

At least difunctional amines, but preferably polyethyleneimines (PEI), are used as further component in the process according to the invention. 25 Polyethyleneimines are generally polymers in the main chains of which there are NH groups which are separated from one another in each case by two methylene groups:



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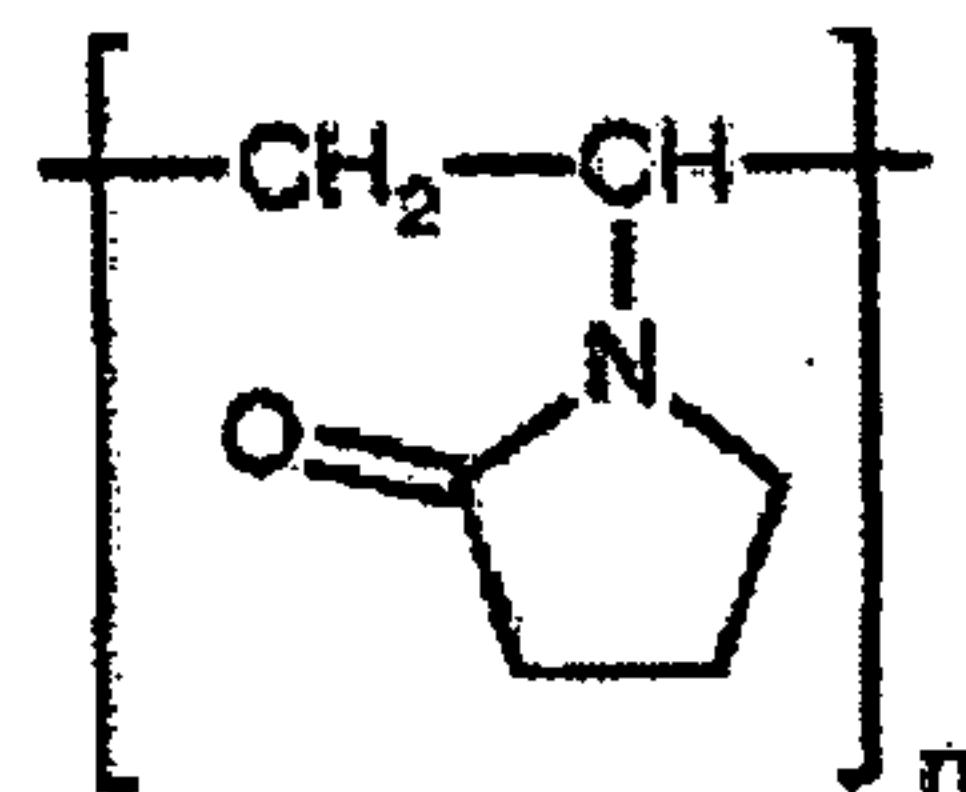
Polyethyleneimines belong to the polyelectrolytes and the complexing polymers. Short-chain, linear polyethyleneimines with a correspondingly high fraction of primary amino groups, i.e. products of the general 35 formula  $\text{H}_2\text{N}-[\text{CH}_2-\text{CH}_2-\text{NH}]_n\text{H}$  ( $n = 2$ : diethylenetriamine;  $n = 3$ : triethylenetetramine;  $n = 4$ :

tetraethylenepentamine) are sometimes called polyethyleneamines or polyalkylenepolyamines.

In the processes according to the invention, 5 polyethyleneimines with a molecular weight of at least 500 g/mol, preferably from 600 to 30 000 or 650 to 25 000 g/mol and in particular from 700 to 5000 g/mol or 850 to 2500 g/mol, are preferably used.

10 Protective colloids

In the process according to the invention, PVP is used as protective colloid. PVP is the abbreviation for polyvinylpyrrolidones (also known as polyvidone). According to Römpf Chemie Lexikon, Online-edition 3.6, 15 2010, they are [poly(1-vinylpyrrolidin-2-ones)], i.e. polymers (vinyl polymers) which conform to the general formula:



Standard commercial polyvinylpyrrolidones have molar 20 masses in the range from ca. 2500-750 000 g/mol which are characterized by stating the K values and have - depending on the K value - glass transition temperatures from 130 to 175 °C. They are supplied as white, hygroscopic powders or as aqueous solution.

25 In the processes according to the invention, preference is given to using PVPs with a high molecular weight, i.e. more than 400 000 g/mol and preferably from 500 000 g/mol to 2 000 000 g/mol. It is furthermore preferred for the polyvinylpyrrolidones to have a K 30 value of more than 60, preferably more than 75 and in particular more than 80. A preferred range is between 65 and 90 for the K value.

Water-insoluble liquid material

The microcapsules produced using the process described above contain in the interior a material that is preferably water-insoluble and liquid at 21°C (i.e. at 21°C, a maximum of 10 g of the material can be 5 dissolved in 1 l of water). This includes all types of hydrophobic water-insoluble liquids, and any blends thereof. Excluded are any fragrances or perfumes as such materials.

This material is also referred to hereinbelow as "oil".  
10 These oils must be able, preferably without auxiliaries, to dissolve the isocyanates in order to be able to use them in the present process. Should an oil not ensure adequate solubility of the isocyanates, there is the option of overcoming this disadvantage by 15 using suitable solubility promoters.

Besides the aforementioned oils, the microcapsules can also have further, optionally liquid or solid, ingredients which are dissolved, dispersed or emulsified in the oil in the microcapsules.

20 The phrase "oil" in the context of the present invention encompasses all kinds of oil bodies or oil components, in particular vegetable oils like e.g. rape seed oil, sunflower oil, soy oil, olive oil and the like, modified vegetable oils e.g. alkoxylated sunflower or soy oil, synthetic (tri)glycerides like e.g. technical mixtures of mono, di and triglycerides of C6-C22 fatty acids, fatty acid alkyl esters e.g. methyl or ethyl esters of vegetable oils (Agnique® ME 25 18 RD-F, Agnique® ME 18 SD-F, Agnique® ME 12C-F, Agnique® ME1270, all products of Cognis GmbH, Germany) fatty acid alkyl esters based on said C6-C22 fatty acids, mineral oils and their mixtures. Examples 30 illustrating the nature of suitable hydrophobic carriers without limiting the invention to these examples are: Guerbet alcohols based on fatty alcohols 35 having 6 to 18, preferably 8 to 10, carbon atoms, esters of linear C6-C22-fatty acids with linear or branched C6-C22-fatty alcohols or esters of branched

C6-C 13-carboxylic acids with linear or branched C6-C 22-fatty alcohols, such as, for example, myristyl myristate, myristyl palmitate, myristyl stearate, myristyl isostearate, myristyl oleate, myristyl behenate, myristyl erucate, cetyl myristate, cetyl palmitate, cetyl stearate, cetyl isostearate, cetyl oleate, cetyl behenate, cetyl erucate, stearyl myristate, stearyl palmitate, stearyl stearate, stearyl isostearate, stearyl oleate, stearyl behenate, stearyl erucate, isostearyl myristate, isostearyl palmitate, isostearyl stearate, isostearyl isostearate, isostearyl oleate, isostearyl behenate, isostearyl oleate, oleyl myristate, oleyl palmitate, oleyl stearate, oleyl isostearate, oleyl oleate, oleyl behenate, oleyl erucate, behenyl myristate, behenyl palmitate, behenyl stearate, behenyl isostearate, behenyl oleate, behenyl behenate, behenyl erucate, erucyl myristate, erucyl palmitate, erucyl stearate, erucyl isostearate, erucyl oleate, erucyl behenate and erucyl erucate. Also suitable are esters of linear C6-C22-fatty acids with branched alcohols, in particular 2-ethylhexanol, esters of C18-C38- alkylhydroxy carboxylic acids with linear or branched C6-C 22-fatty alcohols, in particular Dioctyl Malate, esters of linear and/or branched fatty acids with polyhydric alcohols (such as, for example, propylene glycol, dimerdiol or trimertriol) and/or Guerbet alcohols, triglycerides based on C6-C10-fatty acids, liquid mono-/di-/triglyceride mixtures based on C6-C18-fatty acids, esters of C6-C22-fatty alcohols and/or Guerbet alcohols with aromatic carboxylic acids, in particular benzoic acid, esters of C2- C12-dicarboxylic acids with linear or branched alcohols having 1 to 22 carbon atoms or polyols having 2 to 10 carbon atoms and 2 to 6 hydroxyl groups, vegetable oils, branched primary alcohols, substituted cyclohexanes, linear and branched C6-C22-fatty alcohol carbonates, such as, for example, dicaprylyl carbonate (Cetiol® CC), Guerbet carbonates, based on fatty alcohols having 6 to 18, preferably 8 to 10, carbon

atoms, esters of benzoic acid with linear and/or branched C6-C22-alcohols, linear or branched, symmetrical or asymmetrical dialkyl ethers having 6 to 22 carbon atoms per alkyl group, such as, for example, 5 dicaprylyl ether, ring-opening products of epoxidized fatty acid esters with polyols, silicone oils (cyclomethicones, silicone methicone grades, etc.), aliphatic or naphthenic hydrocarbons, such as, for example, squalane, squalene or dialkylcyclohexanes, 10 and/or mineral oils.

Within the context of the present invention, preferred oils are, Guerbet alcohols based on fatty alcohols having 6 to 18, preferably 8 to 10, carbon atoms, 15 esters of linear C6-C22-fatty acids with linear or branched C6-C22-fatty alcohols or esters of branched C6-C13-carboxylic acids with linear or branched C6-C22-fatty alcohols, such as e.g. myristyl myristate, myristyl palmitate, myristyl stearate, myristyl 20 isostearate, myristyl oleate, myristyl behenate, myristyl erucate, cetyl myristate, cetyl palmitate, cetyl stearate, cetyl isostearate, cetyl oleate, cetyl behenate, cetyl erucate, stearyl myristate, stearyl palmitate, stearyl stearate, stearyl isostearate, 25 stearyl oleate, stearyl behenate, stearyl erucate, isostearyl myristate, isostearyl palmitate, isostearyl stearate, isostearyl isostearate, isostearyl oleate, isostearyl behenate, isostearyl oleate, oleyl myristate, oleyl palmitate, oleyl stearate, oleyl 30 isostearate, oleyl oleate, oleyl behenate, oleyl erucate, behenyl myristate, behenyl palmitate, behenyl stearate, behenyl isostearate, behenyl oleate, behenyl behenate, behenyl erucate, erucyl myristate, erucyl palmitate, erucyl stearate, erucyl isostearate, erucyl 35 oleate, erucyl behenate and erucyl erucate.

Also preferred oils are esters of linear C6-C22-fatty acids with branched alcohols, in particular 2-ethylhexanol, esters of C18-C38-

alkylhydroxycarboxylic acids with linear or branched C6-C22-fatty alcohols, linear or branched C6-C22-fatty alcohols, in particular dioctyl malates, esters of linear and/or branched fatty acids with polyhydric 5 alcohols (such as e.g. propylene glycol, dimerdiol or trimertriol) and/or Guerbet alcohols, triglycerides based on C6-C10-fatty acids, liquid mono-/di-/triglyceride mixtures based on C6-C18-fatty acids, esters of C6-C22-fatty alcohols and/or Guerbet 10 alcohols with aromatic carboxylic acids, in particular benzoic acid, esters of C2-C12-dicarboxylic acids with linear or branched alcohols having 1 to 22 carbon atoms or polyols having 2 to 10 carbon atoms and 2 to 6 hydroxyl groups, vegetable oils, branched primary 15 alcohols, substituted cyclohexanes, linear and branched C6-C22-fatty alcohol carbonates, such as e.g. dicaprylyl carbonate (Cetiol TM CC), Guerbet carbonates based on fatty alcohols having 6 to 18, preferably 8 to 10, carbon atoms, esters of benzoic acid with linear 20 and/or branched C6-C22-alcohols (e.g. Finsolv TM TN), linear or branched, symmetrical or asymmetrical dialkyl ethers having 6 to 22 carbon atoms per alkyl group, such as e.g. dicaprylyl ether (Cetiol TM OE), ring-opening products of epoxidized fatty acid esters with 25 polyols, silicone oils (cyclomethicones, silicon methicone types etc.) and/or aliphatic or naphthenic hydrocarbons, such as e.g. squalane, squalene or dialkylcyclohexanes.

30 Further suitable oils or oil constituents may be UV filter substances. Typical oil-soluble UV-B filters or broad-spectrum UV A/B filters are, for example, 3-benzylidene camphor or 3-benzylidene norcamphor and derivatives thereof, e.g. 3-(4-methylbenzylidene)- 35 camphor, 3-(4'-trimethylammonium)benzylidenebornan-2-one methylsulfate (Mexoryl SO), 3,3'-(1,4-phenylenedimethine)bis(7,7-dimethyl-2-oxobicycle-[2.2.1]heptane-1-methanesulfonic acid) and salts (Mexoryl SX), 3-(4'-sulfo)benzylidenebornan-2-one and

salts (Mexoryl SL), polymer of N-[(2 and 4)-[2-oxoborn-3-ylidene)methyl]benzyl]acrylamide (Mexoryl SW), 2-(2H-benzotriazol-2-yl)-4-methyl-6-(2-methyl-3-(1,3,3,3-tetramethyl-1-(trimethylsilyloxy)disiloxanyl)propyl)-5-phenol (Mexoryl SL), 4-aminobenzoic acid derivatives, preferably 2-ethylhexyl 4-(dimethylamino)benzoate, 2-octyl 4-(dimethylamino)benzoate and amyl 4-(dimethylamino)benzoate; esters of cinnamic acid, preferably 2-ethylhexyl 4-methoxycinnamate, propyl 10 4-methoxycinnamate, isoamyl 4-methoxycinnamate, 2-ethylhexyl 2-cyano-3,3-phenylcinnamate (octocrylene); esters of salicylic acid, preferably 2-ethylhexyl salicylate, 4-isopropylbenzyl salicylate, homomenthyl salicylate; derivatives of benzophenone, preferably 15 2-hydroxy-4-methoxybenzophenone, 2-hydroxy-4-methoxy-4'-methylbenzophenone, 2,2'-dihydroxy-4-methoxybenzophenone; esters of benzalmalonic acid, preferably di-2-ethylhexyl 4-methoxybenzmalonate; triazine derivatives, such as e.g. 2,4,6-trianilino(p-carbo-2'-ethyl-1'-hexyloxy)-1,3,5-triazine and 2,4,6-20 tris[p-(2-ethylhexyloxycarbonyl)anilino]-1,3,5-triazine (Uvinul T 150) or bis(2-ethylhexyl) 4,4'-(6-[4-((1,1-dimethylethyl)aminocarbonyl)phenylamino]-1,3,5-triazine-2,4-diyl)diimino]bisbenzoate (Uvasorb® HEB); 25 2,2-(methylenebis(6-(2H-benzotriazol-2-yl)-4-(1,1,3,3-tetramethylbutyl)phenol (Tinosorb M); 2,4-bis[4-(2-ethylhexyloxy)-2-hydroxyphenyl]-6-(4-methoxyphenyl)-1,3,5-triazine (Tinosorb S); propane-1,3-diones, such as e.g. 1-(4-tert-butylphenyl)-3-(4'-methoxyphenyl)-30 propane-1,3-dione; ketotricyclo(5.2.1.0)decane derivatives, dimethicodiethyl benzalmalonate (Parsol SLX).

Furthermore, liquid linear and/or branched and/or saturated or unsaturated hydrocarbons or any desired mixtures thereof can be used as oils within the context of the present invention. These may be e.g. alkanes having 4 to 22, preferably 6 to 18, carbon atoms, or any desired mixtures thereof. Also of suitability are

the unsaturated hydrocarbons having 4 to 22 carbon atoms, or unsaturated hydrocarbons of identical carbon number, and any desired mixtures of these hydrocarbons. Cyclic hydrocarbons and aromatics, e.g. toluene and 5 mixtures thereof may also be oils within the context of the present invention. Also suitable are silicone oils. Any desired mixtures of all of the specified core materials are also suitable.

10 It is also possible for other liquid, preferably water-insoluble materials, such as thickeners, silicone defoamers, oil soluble corrosion inhibitors and similar additives, like extreme pressure additives, yellow metal deactivators and the like, dyes or oil-soluble 15 medicaments, emollients, odor absorbing compounds, cosmetic oil phases, film forming additive, pearlizer, vitamins, dyes, biocides to be used and be present in the microcapsules. Any desired mixtures of these further materials may also be present in the 20 microcapsules. In cases where such material is not oil-soluble, additives may be used for dispersing or emulsifying it. Otherwise, many actives, as for example biocides or dyes often only available as blends with an oily solvent. Those compositions are also useful in the 25 context of the present invention. Most preferred is the use of biocides, emollients, dyes, and UV-filters in the microcapsules of the present invention.

#### Biocides

30 A biocide is a chemical substance capable of killing different forms of living organisms used in fields such as medicine, agriculture, forestry, and mosquito control. Usually, biocides are divided into two sub-groups:  
35

- pesticides, which includes fungicides, herbicides, insecticides, algicides, moluscicides, miticides and rodenticides, and

- antimicrobials, which includes germicides, antibiotics, antibacterials, antivirals, antifungals, antiprotozoals and antiparasites.

5 Biocides can also be added to other materials (typically liquids) to protect the material from biological infestation and growth. For example, certain types of quaternary ammonium compounds (quats) can be added to pool water or industrial water systems to act  
10 as an algicide, protecting the water from infestation and growth of algae.

Pesticides: The U.S Environmental Protection Agency (EPA) defines a pesticide as "any substance or mixture 15 of substances intended for preventing, destroying, repelling, or mitigating any pest". A pesticide may be a chemical substance or biological agent (such as a virus or bacteria) used against pests including insects, plant pathogens, weeds, mollusks, birds, 20 mammals, fish, nematodes (roundworms) and microbes that compete with humans for food, destroy property, spread disease or are a nuisance. In the following examples, pesticides suitable for the agrochemical compositions according to the present invention are given:

25 Fungicides: A fungicide is one of three main methods of pest control - the chemical control of fungi in this case. Fungicides are chemical compounds used to prevent the spread of fungi in gardens and crops. Fungicides 30 are also used to fight fungal infections. Fungicides can either be contact or systemic. A contact fungicide kills fungi when sprayed on its surface. A systemic fungicide has to be absorbed by the fungus before the fungus dies. Examples for suitable fungicides, 35 according to the present invention, encompass the following species: (3-ethoxypropyl)mercury bromide, 2-methoxyethyl-mercury chloride, 2-phenylphenol, 8-hydroxyquinoline sulfate, 8-phenylmercurioxy-quinoline, acibenzolar, acylamino acid fungicides, acypetacs,

aldimorph, aliphatic nitrogen fungicides, allyl alcohol, amide fungicides, ampropylfos, anilazine, anilide fungicides, antibiotic fungicides, aromatic fungicides, aureofungin, azaconazole, azithiram, 5 azoxystrobin, barium polysulfide, benalaxy, l benalaxy-M, benodanil, benomyl, benquinox, bentaluron, benthiavalicarb, benzalkonium chloride, benzamacril, benzamide fungicides, benzamorf, benzanilide fungicides, benzimidazole fungicides, benzimidazole 10 precursor fungicides, benzimidazolylcarbamate fungicides, benzohydroxamic acid, benzothiazole fungicides, bethoxazin, binapacryl, biphenyl, bitertanol, bithionol, blasticidin-S, Bordeaux mixture, boscalid, bridged diphenyl fungicides, bromuconazole, 15 bupirimate, Burgundy mixture, buthiobate, butylamine, calcium polysulfide, captafol, captan, carbamate fungicides, carbamorph, carbanilate fungicides, carbendazim, carboxin, carpropamid, carvone, Cheshunt mixture, chinomethionat, chlobenthiazone, 20 chloraniformethan, chloranil, chlorfenazole, chlorodinitronaphthalene, chloroneb, chloropicrin, chlorothalonil, chlorquinox, chlozolinate, ciclopirox, climbazole, clotrimazole, conazole fungicides, conazole fungicides (imidazoles), conazole fungicides 25 (triazoles), copper(II) acetate, copper(II) carbonate, basic, copper fungicides, copper hydroxide, copper naphthenate, copper oleate, copper oxychloride, copper(II) sulfate, copper sulfate, basic, copper zinc chromate, cresol, cufraneb, cuprobam, cuprous oxide, 30 cyazofamid, cyclafuramid, cyclic dithiocarbamate fungicides, cycloheximide, cyflufenamid, cymoxanil, cpendazole, cyproconazole, cyprodinil, dazomet, DBCP, debacarb, decafentin, dehydroacetic acid, dicarboximide fungicides, dichlofluanid, dichlone, dichlorophen, 35 dichlorophenyl, dicarboximide fungicides, dichlozoline, diclobutrazol, diclocymet, diclomezine, dicloran, diethofencarb, diethyl pyrocarbonate, difenoconazole, diflumetorim, dimethirimol, dimethomorph, dimoxystrobin, diniconazole, dinitrophenol fungicides,

dinobuton, dinocap, dinocton, dinopenton, dinosulfon,  
 dinoterbon, diphenylamine, dipyrithione, disulfiram,  
 ditalimfos, dithianon, dithiocarbamate fungicides,  
 DNOC, dodemorph, dodicin, dodine, DONATODINE,  
 5 drazoxolon, edifenphos, epoxiconazole,  
 etaconazole, etem, ethaboxam, ethirimol, ethoxyquin,  
 ethylmercury 2,3-dihydroxypropyl mercaptide,  
 ethylmercury acetate, ethylmercury bromide,  
 ethylmercury chloride, ethylmercury phosphate,  
 10 etridiazole, famoxadone, fenamidone, fenaminosulf,  
 fenapanil, fenarimol, fenbuconazole, fenfuram,  
 fenhexamid, fenitropan, fenoxanil, fenpiclonil,  
 fenpropidin, fenpropimorph, fentin, ferbam, ferimzone,  
 fluazinam, fludioxonil, flumetover, flumorph,  
 15 fluopicolide, fluoroimide, fluotrimazole,  
 fluoxastrobin, fluquinconazole, flusilazole,  
 flusulfamide, flutolanil, flutriafol, folpet,  
 formaldehyde, fosetyl, fuberidazole, furalaxyd,  
 furametpyr, furamide fungicides, furanilide fungicides,  
 20 furcarbanil, furconazole, furconazole-cis, furfural,  
 furmecyclox, furopghanate, glyodin, griseofulvin,  
 guazatine, halacrinate, hexachlorobenzene,  
 hexachlorobutadiene, hexachlorophene, hexaconazole,  
 hexylthiofos, hydrargaphen, hymexazol, imazalil,  
 25 imibenconazole, imidazole fungicides, iminoctadine,  
 inorganic fungicides, inorganic mercury fungicides,  
 iodomethane, ipconazole, iprobenfos, iprodione,  
 iprovalicarb, isoprothiolane, isovaledione,  
 kasugamycin, kresoxim-methyl, lime sulphur, mancopper,  
 30 mancozeb, maneb, mebenil, mecarbinzid, mepanipyrim,  
 mepronil, mercuric chloride, mercuric oxide, mercurous  
 chloride, mercury fungicides, metalaxyl, metalaxyl-M,  
 metam, metazoxolon, metconazole, methasulfocarb,  
 methfuroxam, methyl bromide, methyl isothiocyanate,  
 35 methylmercury benzoate, methylmercury dicyandiamide,  
 methylmercury pentachlorophenoxyde, metiram,  
 metominostrobin, metrafenone, metsulfovax, milneb,  
 morpholine fungicides, myclobutanil, myclozolin, N-  
 (ethylmercury)-p-toluenesulphonanilide, nabam,

natamycin, nitrostyrene, nitrothal-isopropyl, nuarimol, OCH, octhilinone, ofurace, organomercury fungicides, organophosphorus fungicides, organotin fungicides, orysastrobin, oxadixyl, oxathiin fungicides, oxazole 5 fungicides, oxine copper, oxpoconazole, oxycarboxin, pefurazoate, penconazole, pencycuron, pentachlorophenol, penthiopyrad, phenylmercuriurea, phenylmercury acetate, phenylmercury chloride, phenylmercury derivative of pyrocatechol, phenylmercury 10 nitrate, phenylmercury salicylate, phenylsulfamide fungicides, phosdiphen, phthalide, phthalimide fungicides, picoxystrobin, piperalin, polycarbamate, polymeric dithiocarbamate fungicides, polyoxins, polyoxorim, polysulfide fungicides, potassium azide, 15 potassium polysulfide, potassium thiocyanate, probenazole, prochloraz, procymidone, propamocarb, propiconazole, propineb, proquinazid, prothiocarb, prothioconazole, pyracarbolid, pyraclostrobin, pyrazole fungicides, pyrazophos, pyridine fungicides, 20 pyridinitril, pyrifenox, pyrimethanil, pyrimidine fungicides, pyroquilon, pyroxichlor, pyroxyfur, pyrrole fungicides, quinacetol, quinazamid, quinconazole, quinoline fungicides, quinone fungicides, quinoxaline fungicides, quinoxyfen, quintozone, rabenzazole, 25 salicylanilide, silthiofam, simeconazole, sodium azide, sodium orthophenylphenoxide, sodium pentachlorophenoxide, sodium polysulfide, spiroxamine, streptomycin, strobilurin fungicides, sulfonanilide fungicides, sulfur, sultopen, TCMTB, tebuconazole, 30 tecloftalam, tecnazene, tecoram, tetriconazole, thiabendazole, thiadifluor, thiazole fungicides, thicyofen, thifluzamide, thiocarbamate fungicides, thiochlorfenphim, thiomersal, thiophanate, thiophanate-methyl, thiophene fungicides, thioquinox, thiram, 35 tiadinil, tioxymid, tivedo, tolclofos-methyl, tolnaftate, tolylfluanid, tolylmercury acetate, triadimefon, triadimenol, triamiphos, triarimol, triazbutil, triazine fungicides, triazole fungicides, triazoxide, tributyltin oxide, trichlamide,

tricyclazole, tridemorph, trifloxystrobin, triflumizole, triforine, triticonazole, unclassified fungicides, undecylenic acid, uniconazole, urea fungicides, validamycin, valinamide fungicides, 5 vinclozolin, zarilamid, zinc naphthenate, zineb, ziram, zoxamide and their mixtures.

10 . Herbicides: An herbicide is a pesticide used to kill unwanted plants. Selective herbicides kill specific targets while leaving the desired crop relatively unharmed. Some of these act by interfering with the growth of the weed and are often based on plant hormones. Herbicides used to clear waste ground are 15 nonselective and kill all plant material with which they come into contact. Herbicides are widely used in agriculture and in landscape turf management. They are applied in total vegetation control (TVC) programs for maintenance of highways and railroads. Smaller 20 quantities are used in forestry, pasture systems, and management of areas set aside as wildlife habitat. In the following, a number of suitable herbicides are compiled:

25 o 2,4-D, a broadleaf herbicide in the phenoxy group used in turf and in no-till field crop production. Now mainly used in a blend with other herbicides that act as synergists, it is the most widely used herbicide in the world, third most commonly used in the United 30 States. It is an example of synthetic auxin (plant hormone).

o Atrazine, a triazine herbicide used in corn and sorghum for control of broadleaf weeds and grasses. It is still used because of its low cost and because it 35 works as a synergist when used with other herbicides, it is a photosystem II inhibitor.

o Clopyralid, a broadleaf herbicide in the pyridine group, used mainly in turf, rangeland, and for control of noxious thistles. Notorious for its ability to

persist in compost. It is another example of synthetic auxin.

- o Dicamba, a persistent broadleaf herbicide active in the soil, used on turf and field corn. It is another 5 example of synthetic auxin.
- o Glyphosate, a systemic nonselective (it kills any type of plant) herbicide used in no-till burndown and for weed control in crops that are genetically modified to resist its effects. It is an example of a EPSPs 10 inhibitor.
- o Imazapyr, a non-selective herbicide used for the control of a broad range of weeds including terrestrial annual and perennial grasses and broadleaved herbs, woody species, and riparian and emergent aquatic 15 species.
- o Imazapic, a selective herbicide for both the pre- and post-emergent control of some annual and perennial grasses and some broadleaf weeds. Imazapic kills plants by inhibiting the production of branched chain amino 20 acids (valine, leucine, and isoleucine), which are necessary for protein synthesis and cell growth.
- o Metoalachlor, a pre-emergent herbicide widely used for control of annual grasses in corn and sorghum; it has largely replaced atrazine for these uses.
- 25 o Paraquat, a nonselective contact herbicide used for no-till burndown and in aerial destruction of marijuana and coca plantings. More acutely toxic to people than any other herbicide in widespread commercial use.
- 30 o Picloram, a pyridine herbicide mainly used to control unwanted trees in pastures and edges of fields. It is another synthetic auxin.
- o Triclopyr.

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Insecticides: An insecticide is a pesticide used against insects in all developmental forms. They include ovicides and larvicides used against the eggs and larvae of insects. Insecticides are used in

agriculture, medicine, industry and the household. In the following, suitable insecticides are mentioned:

- o Chlorinated insecticides such as, for example, 5 Camphechlor, DDT, Hexachloro-cyclohexane, gamma-Hexachlorocyclohexane, Methoxychlor, Pentachlorophenol, TDE, Aldrin, Chlordane, Chlordcone, Dieldrin, Endosulfan, Endrin, Heptachlor, Mirex and their mixtures;
- o Organophosphorus compounds such as, for example, Acephate, Azinphos-methyl, Ben-sulide, Chlorethoxyfos, Chlorpyrifos, Chlorpyriphos-methyl, Diazinon, Dichlorvos (DDVP), Dicrotophos, Dimethoate, Disulfoton, Ethoprop, Fenamiphos, Fenitrothion, Fenthion, 15 Fosthiazate, Malathion, Methamidophos, Methidathion, Methyl-parathion, Mevinphos, Naled, Omethoate, Oxydemeton-methyl, Parathion, Phorate, Phosalone, Phosmet, Phostebupirim, Pirimiphos-methyl, Profenofos, Terbufos, Tetrachlorvinphos, Tribufos, Trichlorfon and 20 their mixture;
- o Carbamates such as, for example, Aldicarb, Carbofuran, Carbaryl, Methomyl, 2-(1-Methylpropyl)phenyl methylcarbamate and their mixtures;
- o Pyrethroids such as, for example, Allethrin, 25 Bifenthrin, Deltamethrin, Permethrin, Resmethrin, Sumithrin, Tetramethrin, Tralomethrin, Transfluthrin and their mixtures;
- o Plant toxin derived compounds such as, for example, Derris (rotenone), Pyrethrum, Neem 30 (Azadirachtin), Nicotine, Caffeine and their mixtures.

Rodenticides: Rodenticides are a category of pest control chemicals intended to kill rodents. Rodents are difficult to kill with poisons because their feeding habits reflect their place as scavengers. They would eat a small bit of something and wait, and if they do not get sick, they would continue eating. An effective rodenticide must be tasteless and odorless in lethal

concentrations, and have a delayed effect. In the following, examples for suitable rodenticides are given:

5 o Anticoagulants are defined as chronic (death occurs after 1 - 2 weeks post ingestion of the lethal dose, rarely sooner), single-dose (second generation) or multiple dose (first generation) cumulative rodenticides. Fatal internal bleeding is caused by  
10 lethal dose of anticoagulants such as brodifacoum, coumatetralyl or warfarin. These substances in effective doses are antivitamins K, blocking the enzymes K1-2,3-epoxide-reductase (this enzyme is preferentially blocked by 4-hydroxycoumarin/4-  
15 hydroxythiacaoumarin derivatives) and K1-quinone-reductase (this enzyme is preferentially blocked by indandione derivatives), depriving the organism of its source of active vitamin K1. This leads to a disruption of the vitamin K cycle, resulting in an inability of  
20 production of essential blood-clotting factors (mainly coagulation factors II (prothrombin), VII (proconvertin), IX (Christmas factor) and X (Stuart factor)). In addition to this specific metabolic disruption, toxic doses of 4-hydroxycoumarin/4-  
25 hydroxythiacaoumarin and indandione anticoagulants are causing damage to tiny blood vessels (capillaries), increasing their permeability, causing diffuse internal bleedings (haemorrhagias). These effects are gradual; they develop in the course of days and are not  
30 accompanied by any nociceptive perceptions, such as pain or agony. In the final phase of intoxication the exhausted rodent collapses in hypovolemic circulatory shock or severe anemia and dies calmly. Rodenticidal anticoagulants are either first generation agents (4-  
35 hydroxycoumarin type: warfarin, coumatetralyl; indandione type: pindone, diphacinone, chlorophacinone), generally requiring higher concentrations (usually between 0.005 and 0.1%), consecutive intake over days in order to accumulate the

lethal dose, poor active or inactive after single feeding and less toxic than second generation agents, which are derivatives of 4-hydroxycoumarin (difenacoum, brodifacoum, bromadiolone and flocoumafen) or 4-5 hydroxy-1-benzothiin-2-one (4-hydroxy-1-thiacoumarin, sometimes incorrectly referred to as 4-hydroxy-1-thiocoumarin, for reason see heterocyclic compounds), namely difethialone. Second generation agents are far more toxic than first generation agents, they are 10 generally applied in lower concentrations in baits (usually in the order of 0.001 - 0.005%), and are lethal after single ingestion of bait and are effective also against strains of rodents that have become 15 resistant against first generation anticoagulants; thus the second generation anticoagulants are sometimes referred to as "superwarfarins". Sometimes, anticoagulant rodenticides are potentiated by an antibiotic, most commonly by sulfaquinoxaline. The aim of this association (e.g. warfarin 0.05% + 20 sulfaquinoxaline 0.02%, or difenacoum 0.005% + sulfaquinoxaline 0.02% etc.) is that the antibiotic/bacteriostatic agent suppresses intestinal/gut symbiotic microflora that represents a source of vitamin K. Thus the symbiotic bacteria are 25 killed or their metabolism is impaired and the production of vitamin K by them is diminished, an effect which logically contributes to the action of anticoagulants. Antibiotic agents other than sulfaquinoxaline may be used, for example co- 30 trimoxazole, tetracycline, neomycin or metronidazole. A further synergism used in rodenticidal baits is that of an association of an anticoagulant with a compound with vitamin D-activity, i.e. cholecalciferol or ergocalciferol (see below). A typical formula used is, 35 e. g., warfarin 0.025 - 0.05% + cholecalciferol 0.01%. In some countries there are even fixed three-component rodenticides, i.e. anticoagulant + antibiotic + vitamin D, e. g. difenacoum 0.005% + sulfaquinoxaline 0.02% + cholecalciferol 0.01%. Associations of a second-

generation anticoagulant with an antibiotic and/or vitamin D are considered to be effective even against the most resistant strains of rodents, though some second generation anticoagulants (namely brodifacoum 5 and difethialone), in bait concentrations of 0.0025 - 0.005% are so toxic that no known resistant strain of rodents exists and even rodents resistant against any other derivatives are reliably exterminated by application of these most toxic anticoagulants.

10 Vitamin K1 has been suggested and successfully used as an antidote for pets or humans, which/who were either accidentally or intentionally (poison assaults on pets, suicidal attempts) exposed to anticoagulant poisons. In addition, since some of these poisons act 15 by inhibiting liver functions and in progressed stages of poisoning, several blood-clotting factors as well as the whole volume of circulating blood lacks, a blood transfusion (optionally with the clotting factors present) can save a person's life who inadvertently 20 takes them, which is an advantage over some older poisons.

o Metal phosphides have been used as a means of killing rodents and are considered single-dose fast 25 acting rodenticides (death occurs commonly within 1-3 days after single bait ingestion). A bait consisting of food and a phosphide (usually zinc phosphide) is left where the rodents can eat it. The acid in the digestive system of the rodent reacts with the phosphide to 30 generate the toxic phosphine gas. This method of vermin control has possible use in places where rodents are resistant to some of the anticoagulants, particularly for control of house and field mice; zinc phosphide baits are also cheaper than most second-generation 35 anticoagulants, so that sometimes, in cases of large infestation by rodents, their population is initially reduced by copious amounts of zinc phosphide bait applied, and the rest of the population that survived the initial fast-acting poison is then eradicated by

prolonged feeding on anticoagulant bait. Inversely, the individual rodents that survived anticoagulant bait poisoning (rest population) can be eradicated by pre-baiting them with nontoxic bait for a week or two (this 5 is important to overcome bait shyness, and to get rodents used to feeding in specific areas by offering specific food, especially when eradicating rats) and subsequently applying poisoned bait of the same sort as used for pre-baiting until all consumption of the bait 10 ceases (usually within 2-4 days). These methods of alternating rodenticides with different modes of action provides a factual or an almost 100% eradication of the rodent population in the area if the acceptance/palatability of bait is good (i.e., rodents 15 readily feed on it).

o Phosphides are rather fast acting rat poisons, resulting in that the rats are dying usually in open areas instead of the affected buildings. Typical 20 examples are aluminum phosphide (fumigant only), calcium phosphide (fumigant only), magnesium phosphide (fumigant only) and zinc phosphide (in baits). Zinc phosphide is typically added to rodent baits in amounts of around 0.75-2%. The baits have a strong, pungent 25 garlic-like odor characteristic for phosphine liberated by hydrolysis. The odor attracts (or, at least, does not repulse) rodents, but has a repulsive effect on other mammals; birds, however (notably wild turkeys), are not sensitive to the smell and feed on the bait 30 thus becoming collateral damage.

o Hypercalcemia. Calciferols (vitamins D), cholecalciferol (vitamin D3) and ergocalciferol (vitamin D2) are used as rodenticides, which are toxic 35 to rodents for the same reason that they are beneficial to mammals: they are affecting calcium and phosphate homeostasis in the body. Vitamins D are essential in minute quantities (few IUs per kilogram body weight daily, which is only a fraction of a milligram), and

like most fat soluble vitamins they are toxic in larger doses as they readily result in the so-called hypervitaminosis, which is, simply said, poisoning by the vitamin. If the poisoning is severe enough (that is, if the dose of the toxicant is high enough), it eventually leads to death. In rodents consuming the rodenticidal bait it causes hypercalcemia by raising the calcium level, mainly by increasing calcium absorption from food, mobilising bone-matrix-fixed calcium into ionised form (mainly monohydrogencarbonate calcium cation, partially bound to plasma proteins,  $[\text{CaHCO}_3^+]$ ), which circulates dissolved in the blood plasma, and after ingestion of a lethal dose the free calcium levels are raised sufficiently so that blood vessels, kidneys, the stomach wall and lungs are mineralised/calcified (formation of calcificates, crystals of calcium salts/complexes in the tissues thus damaging them), leading further to heart problems (myocard is sensitive to variations of free calcium levels that are affecting both myocardial contractibility and excitation propagation between atrias and ventriculas) and bleeding (due to capillary damage) and possibly kidney failure. It is considered to be single-dose, or cumulative (depending on concentration used; the common 0.075% bait concentration is lethal to most rodents after a single intake of larger portions of the bait), sub-chronic (death occurring usually within days to one week after ingestion of the bait). Applied concentrations are 0.075% cholecalciferol and 0.1% ergocalciferol when used alone. There is an important feature of calciferols toxicology which is that they are synergistic with anticoagulant toxicants. This means that mixtures of anticoagulants and calciferols in the same bait are more toxic than the sum of toxicities of the anticoagulant and the calciferol in the bait so that a massive hypercalcemic effect can be achieved by a substantially lower calciferol content in the bait and vice-versa. More pronounced

anticoagulant/hemorrhagic effects are observed if calciferol is present. This synergism is mostly used in baits low in calciferol because effective concentrations of calciferols are more expensive than 5 effective concentrations of most anticoagulants. The historically very first application of a calciferol in rodenticidal bait was, in fact, the Sorex product Sorexa® D (with a different formula than today's Sorexa® D) back in the early 1970's, containing 10 warfarin 0.025% + ergocalciferol 0.1%. Today, Sorexa® CD contains a 0.0025% difenacoum + 0.075% cholecalciferol combination. Numerous other brand products containing either calciferols 0.075 - 0.1% (e. g. Quintox®, containing 0.075% cholecalciferol) alone, 15 or a combination of calciferol 0.01 - 0.075% with an anticoagulant are marketed.

Miticides, moluscicides and nematicides: Miticides are pesticides that kill mites. Antibiotic miticides, 20 carbamate miticides, formamidine miticides, mite growth regulators, organochlorine, permethrin and organophosphate miticides all belong to this category. Molluscicides are pesticides used to control mollusks, such as moths, slugs and snails. These substances 25 include metaldehyde, methiocarb and aluminium sulfate. A nematicide is a type of chemical pesticide used to kill parasitic nematodes (a phylum of worm). A nematicide is obtained from a neem tree's seed cake; which is the residue of neem seeds after oil 30 extraction. The neem tree is known by several names in the world but was first cultivated in India since ancient times.

Antimicrobials: In the following examples, 35 antimicrobials suitable for agrochemical compositions according to the present invention are given. Bactericidal disinfectants mostly used are those applying

- o active chlorine (i.e., hypochlorites, chloramines, dichloroisocyanurate and trichloroisocyanurate, wet chlorine, chlorine dioxide, etc.),
- o active oxygen (peroxides such as peracetic acid, 5 potassium persulfate, sodium perborate, sodium percarbonate and urea perhydrate),
- o iodine (iodpovidone (povidone-iodine, Betadine), Lugol's solution, iodine tincture, iodinated nonionic surfactants),
- 10 o concentrated alcohols (mainly ethanol, 1-propanol, called also n-propanol and 2-propanol, called isopropanol and mixtures thereof; further, 2-phenoxyethanol and 1- and 2-phenoxypropanols are used),
- o phenolic substances (such as phenol (also called 15 "carbolic acid"), cresols (called "Lysole" in combination with liquid potassium soaps), halogenated (chlorinated, brominated) phenols, such as hexachlorophene, triclosan, trichlorophenol, tribromophenol, pentachlorophenol, Dibromol and salts thereof),
- 20 o cationic surfactants such as some quaternary ammonium cations (such as benzalkonium chloride, cetyl trimethylammonium bromide or chloride, didecyldimethylammonium chloride, cetylpyridinium chloride, benzethonium chloride) and others, non-quaternary compounds such as chlorhexidine, glucoprotamine, octenidine dihydrochloride, etc.),
- 25 o strong oxidizers such as ozone and permanganate solutions;
- 30 o heavy metals and their salts such as colloidal silver, silver nitrate, mercury chloride, phenylmercury salts, copper sulfate, copper oxide-chloride etc. Heavy metals and their salts are the most toxic and environmentally hazardous bactericides and, therefore, their use is strongly suppressed or forbidden; further, also
- 35 o properly concentrated strong acids (phosphoric, nitric, sulfuric, amidosulfuric, toluenesulfonic acids) and

o alcalis (sodium, potassium, calcium hydroxides) between pH < 1 or > 13, particularly below elevated temperatures (above 60°C) kill bacteria.

5 As antiseptics (i.e., germicide agents that can be used on human or animal body, skin, mucoses, wounds and the like), few of the above mentioned disinfectants can be used under proper conditions (mainly concentration, pH, temperature and toxicity toward man/animal). Among  
10 them, important are

- o Some properly diluted chlorine preparations (e. g. Daquin's solution, 0.5% sodium or potassium hypochlorite solution, pH-adjusted to pH 7 - 8, or 0.5 - 1% solution of sodium benzenesulfochloramide (chloramine B)), some
- o iodine preparations such as iodopovidone in various galenics (ointments, solutions, wound plasters), in the past also Lugol's solution,
- 20 o peroxides as urea perhydrate solutions and pH-buffered 0.1 - 0.25% peracetic acid solutions,
- o alcohols with or without antiseptic additives, used mainly for skin antiseptis,
- o weak organic acids such as sorbic acid, benzoic acid, lactic acid and salicylic acid
- 25 o some phenolic compounds such as hexachlorophene, triclosan and Dibromol, and
- o cation-active compounds such as 0.05 - 0.5% benzalkonium, 0.5 - 4% chlorhexidine, 0.1 - 2% octenidine solutions.

Bactericidal antibiotics kill bacteria; bacteriostatic antibiotics only slow down their growth or reproduction. Penicillin is a bactericide, as are cephalosporins. Aminoglycosidic antibiotics can act in both a bactericidic manner (by disrupting cell wall precursor leading to lysis) or bacteriostatic manner (by connecting to 30s ribosomal subunit and reducing translation fidelity leading to inaccurate protein

synthesis). Other bactericidal antibiotics according to the present invention include the fluoroquinolones, nitrofurans, vancomycin, monobactams, co-trimoxazole, and metronidazole. The preferred biocides are selected 5 from the group consisting of oxyfluorfen, glyphosate, tebucanozol, desmedipham, phenmedipham, ethofumesat and their mixtures.

#### Emollients

10 The microcapsules could also contain emollients. An emollient is a material that softens, soothes, supplies, coats, lubricates, moisturizes, or cleanses the skin. An emollient typically accomplishes several of these objectives such as soothing, moisturizing, and 15 lubricating the skin. Suitable emollients are mostly selected from the oils as described above. Emollients useful in the present invention can be petroleum-based, fatty acid ester type, alkyl ethoxylate type, fatty acid ester ethoxylates, fatty alcohol type, 20 polysiloxane type, or mixtures of these emollients.

#### Dyes

The microcapsules may also contain dyes, preferably any dyes suitable and approved for cosmetic purposes. 25 Examples include cochineal red A (C.I. 16255), patent blue V (C.I. 42051), indigotin (C.I. 73015), chlorophyllin (C.I. 75810), quinoline yellow (C.I. 47005), titanium dioxide (C.I. 77891), indanthrene blue RS (C.I. 69800) and madder lake (C.I. 58000). These 30 dyes are normally used in concentrations of 0.001 to 0.1% by weight, based on the mixture as a whole.

Besides the beforementioned compounds the microcapsules 35 of the present invention may also contain any desired blends of oils, as well as blends of oil and water in emulsified form. Any kind of emulsion (water-in-oil or oil-in-water, or multiple emulsions) is possible.

For this purpose emulsifiers are needed: The microcapsules according to the present invention might also contain one or more emulsifier. Suitable emulsifiers are, for example, nonionic surfactants from 5 at least one of the following groups: products of the addition of 2 to 30 mol ethylene oxide and/or 0 to 5 mol propylene oxide onto linear C<sub>sub.6-22</sub> fatty alcohols, onto C<sub>sub.12-22</sub> fatty acids, onto alkyl phenols containing 8 to 15 carbon atoms in the alkyl 10 group and onto alkylamines containing 8 to 22 carbon atoms in the alkyl group; alkyl oligoglycosides containing 8 to 22 carbon atoms in the alkyl group and ethoxylated analogs thereof; addition products of 1 to 15 mol ethylene oxide onto castor oil and/or hydrogenated castor oil; addition products of 15 to 60 mol ethylene oxide onto castor oil and/or hydrogenated castor oil; partial esters of glycerol and/or sorbitan with unsaturated, linear or saturated, branched fatty acids containing 12 to 22 carbon atoms and/or 20 hydroxycarboxylic acids containing 3 to 18 carbon atoms and addition products thereof onto 1 to 30 mol ethylene oxide; partial esters of polyglycerol (average degree of self-condensation 2 to 8), polyethylene glycol (molecular weight 400 to 5,000), trimethylolpropane, 25 pentaerythritol, sugar alcohols (for example sorbitol), alkyl glucosides (for example methyl glucoside, butyl glucoside, lauryl glucoside) and polyglucosides (for example cellulose) with saturated and/or unsaturated, linear or branched fatty acids containing 12 to 22 30 carbon atoms and/or hydroxycarboxylic acids containing 3 to 18 carbon atoms and addition products thereof onto 1 to 30 mol ethylene oxide; mixed esters of pentaerythritol, fatty acids, citric acid and fatty alcohol and/or mixed esters of fatty acids containing 6 35 to 22 carbon atoms, methyl glucose and polyols, preferably glycerol or polyglycerol, mono-, di- and trialkyl phosphates and mono-, di- and/or tri-PEG-alkyl phosphates and salts thereof, wool wax alcohols, polysiloxane/polyalkyl/polyether copolymers and

corresponding derivatives, block copolymers, for example Polyethyleneglycol-30 Dipolyhydroxystearate; polymer emulsifiers, for example Pemulen types (TR-1, TR-2) of Goodrich; polyalkylene glycols and glycerol 5 carbonate and ethylene oxide addition products.

It is likewise possible for the ingredients to migrate from the core of the microcapsules (i.e. the oil and/or further materials present in the core) into the shell.

10

The invention further provides aqueous dispersions comprising 5 to 50% by weight, based on the total weight of the dispersion, preferably from 15 to 40% by weight, of microcapsules which can be produced by the 15 above process. A further preferred range is between 20 and 35% by weight. These aqueous dispersions are preferably obtained directly from the process described above.

20

The microcapsule dispersions which are obtained by the present process can be used in a large number of different applications, depending on the type of oil. Preference is given to using the microcapsules for the finishing of all kind of nonwovens, like wipes (for 25 example wet wipes or dry wipes for cosmetic or cleaning purposes), but also for finishing papers (including wallpapers, toilet paper or papers for books and newsletters), for finishing diapers or sanitary napkins and similar hygienic products or textiles e.g. in order 30 to finish the papers or textiles with a dye or an insecticide, or in cosmetic compositions, e.g. for producing sunscreen compositions which comprise the UV filter in the form of the microcapsules. Another use pertains to finishing diapers or sanitary napkins and 35 similar hygienic products. Furthermore the microcapsules may be used in massage oils or crèmes or personal lubricants, and suppositories, e.g. to provide this products with antiinflammatory actives.

The present invention further provides perfume-free microcapsules, preferably formaldehyde-free, and containing a liquid core of a water-insoluble liquid or a hydrophobic material, and a shell of a reaction product of at least two different at least difunctional isocyanates (A) and (B), where the isocyanate (B) must be an anionically modified isocyanate or a polyethylene oxide containing isocyanate or mixtures of the types, and an at least difunctional amine, with the proviso that during the production of the microcapsules the weight ratio between the isocyanates (A) and (B) is in the range from 10:1 to 1:10. Preferably, the aforementioned weight ratios can be adjusted, where the ratio from 3:1 to 1:1 may be attributed particular importance.

These microcapsules preferably have diameters from 1 to 50  $\mu\text{m}$  and preferably diameters from 2 to 45  $\mu\text{m}$ . They may be present in the form of an aqueous dispersion, where the fraction of the capsules can be 1 to 90% by weight, but preferably 5 to 50% by weight.

**Examples**

5 Six microcapsule dispersions were produced using the process according to the invention. For the comparison, a microcapsule dispersion without the addition of an anionically modified isocyanate (B) was prepared and in each case the particle size of the capsules was  
10 determined:

**Particle size determination**

The particle size determinations specified in the examples were carried out by means of static laser 15 diffraction. The stated  $d_{50}$  and  $d_{99}$  values are based on the volume distribution of the particles.

**Comparative example (without isocyanate type (B) - not according to the invention)**

20 Microcapsules were produced as follows using only one isocyanate type (A): a premix (I) was prepared from 50 g of PVP K90 and 1160 g of water and adjusted to a pH of 10.0 using aqueous sodium hydroxide solution (5% strength by weight). Premix II was prepared from 500 g 25 of Myritol<sup>®</sup> 318 (caprylic/capric triglyceride) and 90 g of Desmodur<sup>®</sup> W (dicyclohexylmethane diisocyanate). The two premixes were combined and emulsified with the help of a Mig stirrer for 30 minutes at room temperature at a speed of 1000 rpm. The pH of the emulsion was then 30 adjusted to 8.5 using aqueous sodium hydroxide solution (5% strength by weight). Then, at room temperature and with stirring at 1000 rpm, a solution of 40 g of Lupasol<sup>®</sup> PR8515 (polyethyleneimine) in 160 g of water was added over the course of 1 minute. The reaction 35 mixture was then subjected to the following temperature program: heating to 60°C in 60 minutes, maintaining this temperature for 60 minutes, then 60 minutes at 70°C, 60 minutes at 80°C, and finally 60 minutes at 85°C. The mixture was then cooled to room temperature,

giving the desired microcapsule dispersion with a fraction of nonvolatile components of 34% and a particle size distribution according to the following values:  $d_{50} = 26 \mu\text{m}$  and  $d_{90} = 53 \mu\text{m}$ .

5

Example 1:

Using two different isocyanates of types (A) and (B), microcapsules were prepared as follows: a premix (I) was prepared from 50 g of PVP K90 and 1169 g of water 10 and adjusted to a pH of 10.0 using aqueous sodium hydroxide solution (5% strength by weight). Premix II was prepared from 500 g of Myritol® 318 (caprylic/capric triglyceride), 58 g of Desmodur® W (dicyclohexylmethane diisocyanate) and 39 g of Bayhydur® XP 2547 (anionic HDI 15 oligomer). These two premixes were combined and emulsified with the help of a Mig stirrer for 30 minutes at room temperature at a speed of 1000 rpm. The pH of the emulsion was then adjusted to 8.5 using aqueous sodium hydroxide solution (5% strength by 20 weight). Then, at room temperature and with stirring at 1000 rpm, a solution of 37 g of Lupasol® PR8515 (polyethyleneimine) in 147 g of water was added over the course of 1 minute. The reaction mixture was then subjected to the following temperature program: heating 25 to 60°C in 60 minutes, maintaining this temperature for 60 minutes, then 60 minutes at 70°C, 60 minutes at 80°C and finally 60 minutes at 85°C. The mixture was then cooled to room temperature, giving the desired microcapsule dispersion with a fraction of nonvolatile 30 components of 34% and a particle size distribution according to the following values:  $d_{50} = 6 \mu\text{m}$  and  $d_{90} = 10 \mu\text{m}$ .

Example 2:

35 Using two different isocyanates of types (A) and (B), microcapsules were produced as follows: a premix (I) was prepared from 50 g of PVP K90 and 1169 g of water and adjusted to a pH of 10.0 using aqueous sodium hydroxide solution (5% strength by weight). Premix II

was prepared from 500 g of Myritol® 318 (caprylic/capric triglyceride) and 58 g of Desmodur® W (dicyclohexylmethane diisocyanate). These two premixes were combined and preemulsified by stirring. Then, 39 g 5 of Bayhydur® XP 2547 (anionic HDI oligomer) were added to this preemulsion and the mixture was emulsified with the help of a Mig stirrer for 30 minutes at room temperature at a speed of 1000 rpm. The pH of the emulsion was then adjusted to 8.5 using aqueous sodium hydroxide solution (5% strength by weight). Then, at 10 room temperature and with stirring at 1000 rpm, a solution of 37 g of Lupasol® PR8515 (polyethyleneimine) in 147 g of water was added over the course of 1 minute. The reaction mixture was then subjected to 15 the following temperature program: heating to 60°C in 60 minutes, maintaining this temperature for 60 minutes, then 60 minutes at 70°C, 60 minutes at 80°C and finally 60 minutes at 85°C. The mixture was then cooled to room temperature, giving the desired 20 microcapsule dispersion with a fraction of nonvolatile components of 34% and a particle size distribution according to the following values:  $d_{50} = 9 \mu\text{m}$  and  $d_{90} = 16 \mu\text{m}$ .

25 Example 3:

Using two different isocyanates of types (A) and (B), microcapsules were produced as follows: a premix (I) was prepared from 40 g of PVP K90 and 1146 g of water and adjusted to a pH of 10.0 using aqueous sodium 30 hydroxide solution (5% strength by weight). Premix II was prepared from 500g of Myritol® 318 (caprylic/capric triglyceride) and 94 g of Desmodur® W (dicyclohexylmethane diisocyanate). These two premixes were combined and preemulsified by stirring. 20 g of Bayhydur® XP 2655 35 (anionic HDI oligomer) were then added to this preemulsion and the mixture was emulsified with the help of a Mig stirrer for 30 minutes at room temperature at a speed of 1000 rpm. The pH of the emulsion was then adjusted to 8.5 with aqueous sodium

hydroxide solution (5% strength by weight). Then, at room temperature and with stirring at 1000 rpm, a solution of 80 g of Lupasol® G 100 (polyethyleneimine) in 120 g of water was added over the course of 5 1 minute. The reaction mixture was then subjected to the following temperature program: heating to 60°C in 60 minutes, maintaining this temperature for 60 minutes, then 60 minutes at 70°C, 60 minutes at 80°C and finally 60 minutes at 85°C. The mixture was then 10 cooled to room temperature, giving the desired microcapsule dispersion with a fraction of nonvolatile components of 35% and a particle size distribution according to the following values:  $d_{50} = 4 \mu\text{m}$  and  $d_{90} = 8 \mu\text{m}$ .

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Example 4:

Using two different isocyanates of types (A) and (B), microcapsules were produced as follows: a premix (I) was prepared from 50 g of PVP K90 and 1134 g of water 20 and adjusted to a pH of 10.5 using aqueous sodium hydroxide solution (5% strength by weight). Premix II was prepared from 300 g of Eutanol® G (octyldodecanol), 300 g of Fitoderm (squalane) and 45 g of Desmodur® N 3300 (HDI trimer). These two premixes were combined 25 and preemulsified by stirring. 31 g of Bayhydur® XP 2547 (anionic HDI oligomer) were then added to this preemulsion and the mixture was emulsified with the help of a Mig stirrer for 30 minutes at room temperature at a speed of 1000 rpm. The pH of the 30 emulsion was then adjusted to 8.7 using aqueous sodium hydroxide solution (5% strength by weight). Then, at room temperature and with stirring at 1000 rpm, a solution of 28 g of Lupasol® PR8515 (polyethyleneimine) in 112 g of water was added over the course of 35 1 minute. The reaction mixture was then subjected to the following temperature program: heating to 60°C in 60 minutes, maintaining this temperature for 60 minutes, then 60 minutes at 70°C, 60 minutes at 80°C and finally 60 minutes at 85°C. The mixture was then

cooled to room temperature, giving the desired microcapsule dispersion with a fraction of nonvolatile components of 38% and a particle size distribution according to the following values:  $d_{50} = 8 \mu\text{m}$  and  $d_{90} = 20 \mu\text{m}$ .

Example 5:

Using two isocyanates of type (A) and one of type (B), microcapsules were produced as follows: a premix (I) was prepared from 50 g of PVP K90 and 1130 g of water and adjusted to a pH of 10.0 using aqueous sodium hydroxide solution (5% strength by weight). Premix II was prepared from 500g of Myritol® 318 (caprylic/capric triglyceride), 50 g of Desmodur® W (dicyclohexylmethane diisocyanate), 50 g of Desmodur® N 3300 (HDI trimer) and 30 g of Bayhydur® XP 2547 (anionic HDI oligomer). These two preemulsions were combined and emulsified with the help of a Mig stirrer for 30 minutes at room temperature at a speed of 1000 rpm. The pH of the emulsion was then adjusted to 8.5 using aqueous sodium hydroxide solution (5% strength by weight). Then, at room temperature and with stirring at 1000 rpm, a solution of 38 g of Lupasol® FG (polyethyleneimine) in 152 g of water was added over the course of 1 minute. The reaction mixture was then subjected to the following temperature program: heating to 60°C in 60 minutes, maintaining this temperature for 60 minutes, then 60 minutes at 70°C, 60 minutes at 80°C and finally 60 minutes at 85°C. The mixture was then cooled to room temperature, giving the desired microcapsule dispersion with a fraction of nonvolatile components of 33% and a particle size distribution according to the following values:  $d_{50} = 8 \mu\text{m}$  and  $d_{90} = 14 \mu\text{m}$ .

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Example 6:

Microcapsules were produced analogously to example 1 by using, instead of the 500 g of Myritol® 318, an oil body comprising 350 g of Eusolex 2292 (octyl

methoxycinnamate) and 150 g of Eusolex 9020 (butyl-methoxydibenzoylmethane). This gave a microcapsule dispersion with a fraction of nonvolatile components of 34% and a particle size distribution according to the 5 following values:  $d_{50} = 4 \mu\text{m}$  and  $d_{90} = 20 \mu\text{m}$ .

### Claims

1. A process for producing microcapsules which contain a shell and a core of a liquid water-insoluble material,

5 where an aqueous solution of a protective colloid and a solution of a mixture of at least two structurally different at least difunctional diisocyanates (A) and (B) in a water-insoluble liquid are brought together until an emulsion is formed, to which an at least difunctional amine is then added and which is then heated to temperatures of at least 60°C until the microcapsules are formed, wherein the isocyanate (B) is selected from the anionically modified isocyanates and the isocyanate (A) is uncharged and is not a polyethylene oxide-containing isocyanate, with the proviso that during the production of the microcapsules the weight ratio between the isocyanates (A) and (B) is in the range from 10:1 to 1:10.

10 2. The process as claimed in claim 1, wherein a polyvinylpyrrolidone is used as protective colloid.

15 3. The process as claimed in claim 1 or 2, wherein the isocyanate (A) is selected from the group consisting of hexane 1,6-diisocyanate, hexane 1,6-diisocyanate biuret or oligomers of hexane 1,6-diisocyanate and dicyclohexanemethylene diisocyanate.

20 4. The process as claimed in claim 3, wherein the isocyanate (A) is selected from the group consisting of trimers of hexane 1,6-diisocyanate.

5. The process as claimed in any one of claims 1 to 4, wherein the isocyanate (B) is selected from the group of anionically modified diisocyanates which contain at least one sulfonic acid group in the molecule.

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6. The process as claimed in claim 5, wherein the sulfonic acid group is an aminosulfonic acid group.

7. The process as claimed in any one of claims 1 to 6, 10 wherein the at least difunctional amine used is a polyethyleneimine.

8. The process as claimed in any one of claims 1 to 7, 15 wherein the weight ratio between the isocyanates (A) and (B) is in the range from 10:1 to 1:10.

9. The process as claimed in claim 8, wherein the weight ratio between the isocyanates (A) and (B) is in the range from 5:1 to 1:5.

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10. The process as claimed in claim 8, wherein the weight ratio between the isocyanates (A) and (B) is in the range from 3:1 to 1:1.

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11. The process as claimed in any one of claims 1 to 10, wherein the core-shell ratio w/w of the microcapsules is 20:1 to 1:10.

30 12. The process as claimed in claim 11, wherein the core-shell ratio w/w of the microcapsules is 5:1 to 2:1.

13. The process as claimed in claim 11, wherein the core-shell ratio w/w of the microcapsules is 4:1 to 3:1.

14. The process as claimed in any one of claims 1 to 13, wherein

- 5 (a) a premix (I) is prepared from water and a protective colloid;
- (b) the premix (I) is adjusted to a pH in the range from 5 to 12;
- 10 (c) a further premix (II) is prepared from the liquid, water-insoluble material together with the isocyanates (A) and (B);
- (d) the two premixes (I) and (II) are brought together until an emulsion is formed and
- (e) the at least difunctional amine is then metered into the emulsion from step (d) and
- 15 (f) the emulsion is then heated to temperatures of at least 60°C until the microcapsules are formed.

15. The process as claimed in claim 14, wherein the pH in process step (b) is adjusted to 8 to 12.

20 16. The process as claimed in any one of claims 1 to 15, wherein

- (a) a premix (I) is prepared from water and a protective colloid;
- 25 (b) the premix (I) is adjusted to a pH in the range from 5 to 12;
- (c) a further premix (II) is prepared from a water-insoluble material that is liquid at 21°C together with the isocyanate (A);
- 30 (d) an emulsion is formed from the premixes (I) and (II) by stirring and to this
- (e) is added the second isocyanate (B), and then the pH of the emulsion is adjusted to a value from 5 to 10;

- (f) and then the at least difunctional amine is metered into the emulsion from step (e) and
- (g) then heated to temperatures of at least 60°C until the microcapsules are formed.

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17. The process as claimed in claim 16, wherein the pH in step (e) is adjusted to 7.5 to 9.0.

18. A perfume-free microcapsule comprising a liquid core of a water-insoluble liquid, and a shell of a reaction product of at least two different at least difunctional isocyanates (A) and (B), where (A) is uncharged and is not a polyethylene oxide-containing isocyanate and where the isocyanate (B) must be an anionically modified isocyanate, and an at least difunctional amine, with the proviso that during the production of the microcapsules the weight ratio between the isocyanates (A) and (B) is in the range from 10:1 to 1:10.

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20. 19. The microcapsule as claimed in claim 18, which has a diameter from 1 to 50  $\mu\text{m}$ .

25. 20. The microcapsule as claimed in claim 18 or 19, which is present in the form of an aqueous dispersion.

21. The microcapsule as claimed in any one of claims 18 to 20, which is free from formaldehyde.

30. 22. The use of microcapsules as claimed in any one of claims 18 to 21 for finishing of textiles, papers or nonwovens.

23. The use of microcapsules as claimed in any one of claims 18 to 21 in cosmetic, pharmaceutical, laundry and/or cleaning compositions.

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