Insecticide composition for application to a textile material or plastics material selected from the group consisting of fibers, fabric, knitgoods, nonwovens, netting material, foils, and tarpaulins, which composition comprises a mixture including at least one insecticide and/or at least one repellent, and at least one binder; an impregnated textile material or plastics material comprising at least one insecticide and/or at least one repellent, and at least one binder; processes for impregnation of a textile material or plastics material and a process for coating of a textile material or plastics material.
COMPOSITION FOR IMPREGNATION OF FIBERS, FABRICS AND NETTINGS IMPARTING A PROTECTIVE ACTIVITY AGAINST PESTS

[0001] The present invention relates to an insecticide composition for application to a textile material or plastics material selected from the group consisting of fibers, fabric, knitgoods, nonwovens, netting material, foils, and tarpaulins, which composition comprises a mixture including at least one insecticide and/or at least one repellent, and at least one binder; an impregnated textile material or plastics material comprising at least one insecticide and/or at least one repellent; processes for impregnation of a textile material or plastics material and a process for coating of a textile material or plastics material.

[0002] Infectious diseases cause huge damages by debilitating or even killing humans and animals in many countries, especially in tropical countries. Many of these diseases (e.g., malaria, dengue, and yellow fever, lymphatic filariasis, and leishmaniasis) are transmitted by insects. Since many medical methods like vaccination or medical treatments are either impossible or too expensive or have been rendered ineffective due to spreading resistance against drugs, efforts have been concentrated on controlling the transmitting insects. Methods to control these insects comprise treating surfaces of huts and houses, air spraying and impregnation of curtains and bednets. The latter treatment is up to now mostly done by dipping the textile material into emulsions or dispersions of insecticides or spraying them onto the nets. Since this provides only a loose adhesion of the insecticide molecules on the surface of the fibers, this treatment is not wash-permanent and has to be repeated after each washing. Studies have proven long-lasting insecticide-treated nets (LLINs) to be more reliable in preventing carrier-borne diseases in comparison with conventional nets, which have to be re-impregnated with insecticide after each washing. Experience shows, however, that washed nets are not re-treated in many cases leaving them without any biological activity. WHO, UNICEF and global relief organisations therefore recommend pre-treated, long-lasting insecticide-treated nets which are wash-permanent as an effective means for the containment of deadly tropical diseases, especially malaria and dengue fever. This is not only comfortable for the user but gives him an economic advantage as well, saving the costs for the repeated impregnation. It is an ecological advantage as well, since the permanent treatment is done under controlled conditions in textile finishing plants.

[0003] WO 01/37662 discloses impregnated nettings, or fabrics for insect or tick killing and/or repellent of an insect or tick comprising an insecticide and/or a repellent, and a film forming component reducing the wash off and degradation of the insecticide component from the netting or fabric by forming a water- and optionally an oil-resistant film. The film forming component preferably comprises one or more components selected from paraffin oil or wax derivatives, silicon derivatives, silicon oils or wax derivatives, and perfluorocarbon derivatives. The netting or fabric is impregnated by adding a solution or a water emulsion of an insecticide and/or repellent and a film forming component successively (in two steps) or in one process step. According to the specification of WO 01/37662 is the insecticide and/or repellent dissolved in an organic solvent in the process for impregnation of a fabric or a netting.

[0004] WO 03/034823 discloses an insecticide composition for application to a fabric material, which composition comprises a mixture including an insecticide, a copolymeric binder, that, after drying and while the fabric material is dry, imparts hydrophobicity to the insecticide, and a dispersing agent, that, after application of the composition to a fabric and upon wetting the fabric, reduces the hydrophobicity imparted to the insecticide by the binder to permit limited insecticide release. The copolymeric binder is prepared as a copolymer emulsion that is derived by an emulsion polymerization technique from monomers selected from at least one of the groups including a) vinyl esters of aliphatic acid having 1 to 18 carbon atoms, such as vinyl acetate and vinyl versatate; b) acrylic and methacrylic esters of an alcohol having 1 to 18 carbon atoms, such as butyl acrylate, 2-ethylhexylacrylate, and methyl acrylate; and c) mono- and di-ethylenically unsaturated hydrocarbons, such as styrene, and aliphatic diens, such as butadiene. The preferred copolymeric binder is prepared by emulsion polymerisation of two different monomers. The insecticide composition of WO 03/034823 is applied to the fabric or netting by dipping, spraying, brushing, and the like. According to the examples the insecticides have to be dissolved in organic solvents before applying the insecticide composition to a fabric material.

[0005] U.S. Pat. No. 5,631,072 discloses the manufacture of fabric intended to be made into washable garments, more specifically to the placement of an insecticide such as permethrin in the fabric by impregnation with polymeric binders and a cross-linking agent, or by surface coating with a polymeric binder and a thickening agent to improve the efficiency as an insect repellent and retention of the permethrin in the fabric as an effective insecticide through successive washings of the garments. According to the examples, suitable binders are acrylic binders and polyvinylacetate binders, which are not further specified. The amount of insecticide in the solutions for impregnation of the fabric is very high (1250 mg insecticide per m²).

[0006] It is an object of the present invention to provide an insecticide composition for application to a textile material or plastics material, wherein the insecticide is not washed out and in which the bioavailability of the insecticide for killing insects is maintained after multiple washes.

[0007] According to the present invention there is provided an insecticide composition for application to a textile material or plastics material which composition comprises a mixture including

[0008] a) at least one insecticide and/or at least one repellent as component A, and

[0009] b) at least one acrylic binder as component B1 obtainable by emulsion polymerisation of the following components:

[0010] b1a) n-butyl acrylate as component B1A,

[0011] b1b) at least one monomer of formula I as component B1B

(1)
(0012) wherein

(0013) R₁, R₂ and R₃ are independently selected from C₁- to C₁₀-alkyl which may be linear or branched, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, tert-butyl, n-pentyl, i-pentyl, sec-pentyl, neo-pentyl, 1,2-dimethylpropyl, i-amyl, n-hexyl, i-hexyl, sec-hexyl, n-heptyl, n-octyl, 2-ethylhexyl, n-nonyl, n-decyl, preferably C₂- to C₆-alkyl, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, tert-butyl; substituted or unsubstituted aryl, preferably substituted or unsubstituted C₁₀- to C₁₅-aryl, more preferably substituted or unsubstituted C₆-aryl, for example phenyl or tolyl;

(0014) R₁ and R₂ may further be H;

(0015) except of R³=n-butyl, when R₁ and R₂ are H.

(0016) Preferably R₁ is H or methyl. R₂ is preferably H; R³ is preferably methyl, ethyl or 2-ethylhexyl.

(0017) More preferably R₁ is H or methyl, R₂ is H and R³ is methyl, ethyl or 2-ethylhexyl.

(0018) Most preferably the monomer of formula I is selected from the group consisting of 2-ethylhexylacrylate, methacrylate, methylmethacrylate and ethylacrylate.

(0019) b1c) at least one monomer of formula II as component B₁C

(0020) wherein

(0021) R⁴, R⁵, R⁶ and R⁷ are independently selected from the group consisting of H, C₁- to C₁₀-alkyl which may be linear or branched, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, tert-butyl, n-pentyl, i-pentyl, sec-pentyl, neo-pentyl, 1,2-dimethylpropyl, i-amyl, n-hexyl, i-hexyl, sec-hexyl, n-heptyl, n-octyl, 2-ethylhexyl, n-nonyl and n-decyl; preferably R⁴, R⁵, R⁶ and R⁷ are selected from the group consisting of H, C₁- to C₆-alkyl, which may be linear or branched, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl and tert-butyl; substituted or unsubstituted aryl, preferably substituted or unsubstituted C₁₀- to C₁₅-aryl, more preferably substituted or unsubstituted C₆-aryl, for example phenyl or tolyl;

(0022) more preferably R⁴ is H or methyl, R⁵, R⁶ and R⁷ are preferably independent of each other H;

(0023) most preferably R⁴ is H or methyl and R⁵, R⁶ and R⁷ are H;

(0024) b1d) optionally at least one monomer of formula III as component B₁D

\[
\begin{align*}
\text{(III)}
\end{align*}
\]

(0025) wherein

(0026) R⁸ and R⁹ are independently selected from the group consisting of H, C₁- to C₁₀-alkyl which may be linear or branched, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, tert-butyl, n-pentyl, i-pentyl, sec-pentyl, neo-pentyl, 1,2-dimethylpropyl, i-amyl, n-hexyl, i-hexyl, sec-hexyl, n-heptyl, n-octyl, 2-ethylhexyl, n-nonyl and n-decyl; preferably R⁸ and R⁹ are selected from the group consisting of H, C₁- to C₆-alkyl, which may be linear or branched, for example methyl, ethyl, n-propyl, i-propyl, iso-propyl, n-butyl, i-butyl, sec-butyl and tert-butyl; substituted or unsubstituted aryl, preferably substituted or unsubstituted C₁₀- to C₁₅-aryl, more preferably substituted or unsubstituted C₆-aryl, for example phenyl or tolyl;

(0027) most preferably R⁸ and R⁹ are H;

(0028) X is selected from the group consisting of H, OH, NH₂, OR¹OH, glycidyl, hydroxypropyl,

(0029) groups of the formula

(0030) wherein

(0031) R¹⁰ is selected from the group consisting of C₁- to C₁₀-alkyl which may be branched or linear, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, tert-butyl, n-pentyl, i-pentyl, sec-pentyl, neo-pentyl, 1,2-dimethylpropyl, i-amyl, n-hexyl, i-hexyl, sec-hexyl, n-heptyl, n-octyl, 2-ethylhexyl, n-nonyl, n-decyl; preferably C₁- to C₆-alkyl, which may be branched or linear, for example methyl, ethyl, n-propyl, isopropyl, n-butyl, iso-butyl, sec-butyl and tert-butyl; substituted or unsubstituted aryl, preferably substituted or unsubstituted C₁₀- to C₁₅-aryl, more preferably substituted or unsubstituted C₆-aryl, for example phenyl or tolyl;
R\(^{11}\) is selected from the group consisting of C\(_1\) to C\(_{10}\) alkylene, for example methylene, ethylene, propylene, butylene, pentylene, hexylene, heptylene, octylene, nonylene, decylene; preferably C\(_1\) to C\(_2\) alkylene, for example methylene, ethylene, propylene, butylene;

substituted or unsubstituted arylenes, preferably substituted or unsubstituted C\(_6\) - to C\(_{10}\) arylene, more preferably substituted or unsubstituted C\(_6\)-arylene, for example phenylene; most preferably X is acetoacetyl;

further monomers which are copolymerizable with the monomers mentioned above selected from:

1. polar monomers, preferably, (meth)acrylic nitrile and/or methyl-(meth)acrylate as component B1E1; and/or
2. non polar monomers, preferably styrene and/or a methylstyrene as component B1E2;

and/or

at least one polyurethane as component B2, obtainable by reaction of the following components:

1. at least one diisocyanate or polyisocyanate as component B2A, preferably aliphatic, cycloaliphatic, alicyclic and/or aromatic isocyanates, more preferably diisocyanates, which are optionally biuretized and/or isocyanurized, most preferably 1-isocyanato-3,3,5-trimethyl-5-isocyanatomethylene cyclohexane (IPDI) and hexamethylene diisocyanate-1,6 (HMDI);

2. at least one diol, triol or polyol as component B2B, preferably aliphatic, cycloaliphatic and/or alicyclic diols having 2 to 14, preferably 4 to 10 carbon atoms, more preferably 1,6-hexanediol or neopentyl glycol;

3. optionally further components as component B2C, preferably adipic acid or carbonyl diimidazole (CDI); and

4. optionally further additives as component B2D.

The insecticide composition of the present invention may be in form of a solid or an aqueous formulation, wherein the aqueous formulation is preferred.

The insecticide composition of the present application provides wash resistance while permitting continuous release of the insecticide and/or repellent. It was found by the inventors that insecticide compositions comprising components A as well as B1A, B1B and B1C and/or B2 provide a very good wash resistance while permitting a continuous release of the insecticide and for repellent at a controlled rate.

In the context of the present invention a textile material or plastics material is a material selected from the group consisting of fibers, fabric, knitgoods, nonwovens, netting material, foils, and tarpaulins.

The insecticide composition of the present invention comprises in general 0.001 to 95% by weight preferably 0.1 to 45% by weight, more preferably 0.5 to 30% by weight, based on the weight of the insecticide composition, of at least one insecticide and/or repellent.

The insecticide composition preferably comprises the following components, based on the solids content of the composition:

a. 0.1 to 45% by weight, preferably 0.5 to 30% by weight more preferably 1 to 25% by weight of at least one insecticide and/or at least one repellent (component A), and

b. preferably 70 to 98% by weight, more preferably 70 to 90% by weight based on the polyurethane of at least one acrylic binder (component B1) as defined above, comprising:

1. preferably 15 to 80% by weight more preferably 20 to 70% by weight based on the acrylic binder of at least one monomer of formula I (component B1A);

2. preferably 10 to 90% by weight, more preferably 15 to 65% by weight based on the acrylic binder of at least one monomer of formula II (component B1C);

3. preferably 0 to 5% by weight, more preferably 1 to 4% by weight, more preferably 0.2 to 3% by weight based on the acrylic binder of at least one monomer of formula III (component B1D);

4. further monomers which are copolymerizable with the monomers mentioned (component B1E) above selected from:

b. preferably 0 to 30% by weight, more preferably 0 to 25% by weight, more preferably 5 to 20% by weight based on the acrylic binder of at least one polar monomer, preferably (meth)acrylic nitrile and/or methyl-(meth)acrylate (component B1E1); and/or

c. preferably 0 to 40% by weight, preferably 0 to 30% by weight, more preferably 5 to 20% by weight based on the acrylic binder of at least one non polar monomer, preferably styrene and/or a-methylstyrene (component B1E2).

and/or

b. preferably 70 to 98% by weight, more preferably 75 to 90% by weight based on the polyurethane of at least one diisocyanate or polyisocyanate (component B2) as defined above, comprising:

b. preferably 70 to 98% by weight, more preferably 75 to 90% by weight based on the polyurethane of at least one diisocyanate or polyisocyanate (component B2).
B2A), preferably aliphatic, cycloaliphatic, araliphatic and/or aromatic isocyanates, more preferably diisocyanates, which are optionally biuretized and/or isocyanurized, most preferably 1-isocyanato-3,3,5-trimethyl-5-isocyanatomethylene cyclohexane (IPDI) and hexamethylene diisocyanate-1,6 (HMDI);

[0060] b2b) 10 to 90% by weight, preferably 12 to 85% by weight, more preferably 15 to 65% by weight based on the polyurethane of at least one diol, triol or polyol (component B2B), preferably aliphatic, cycloaliphatic and/or araliphatic diols having 2 to 14, preferably 4 to 10 carbon atoms, more preferably 1,6-hexanediol or neopentyl glycol;

[0061] b2c) 0 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 1 to 5% by weight based on the polyurethane of further components (component B2C), preferably adipic acid or carbonyl dimidazole (CDI) and

[0062] b2d) 0 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 0.5 to 5% by weight based on the polyurethane of further additives (component B2D)

[0063] wherein the sum of the components is 100% by weight of solids content of the insecticide composition.

[0064] The aim of the invention is to control a variety of pests, such as ticks, cockroaches, bed bugs, mites, fleas, lice, fleas, lice, fleas, lice, and very low mammalian toxicity. Suitable insecticides and/or repellents are known by a person skilled in the art.

[0065] Treated foils or tarpaulins can be used on all human premises which are permanently or temporarily inhabited such as refugee camps.

[0070] The insecticide composition of the present invention is particularly suitable for application to polyester nettings as used for mosquito nets.

[0071] The insecticide composition of the present invention may be applied to textile materials or plastics materials before their formation into the desired products, i.e. while still a yarn or in sheet form, or after formation of the desired products.

[0072] Insecticide and/or Repellent (Component A)

[0073] Preferably, the insecticide and/or repellent is an insecticide and/or repellent with a fast paralyzing or killing effect of the insect and very low mammalian toxicity. Suitable insecticides and/or repellents are known by a person skilled in the art.

[0074] Preferred insecticides and/or repellents are mentioned below:

[0075] pyrethroid compounds such as

[0076] Ethofenprox: 2-(4-ethoxyphenyl)-2-methylpropyl-3-phenoxynbenzyl ether,

[0077] Fenvalerate: (RS)-alpha-cyano-3-phenoxynbenzyl (RS)-2-(4-chlorophenyl)-3-methylbutyrate,

[0078] Esfenvalerate: (S)-alpha-cyano-3-phenoxynbenzyl (S)-2-(4-chlorophenyl)-3-methylbutyrate,

[0079] Fenpropatrin: (RS)-alpha-cyano-3-phenoxynbenzyl 2,2,3,3-tetramethylcyclopropane-carboxylate,

[0080] Cypermethrin: (RS)-alpha-cyano-3-phenoxynbenzyl (1RS)-cis, trans-3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropene-carboxylate, alpha-Cypermethrin,

[0081] Permethrin: 3-phenoxynbenzyl (1RS)-cis, trans-3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropene-carboxylate,

[0082] Cyhalothrin: (RS)-alpha-cyano-3-phenoxynbenzyl (Z)-(1RS)-cis-3-(2-chloro,3,3,3-trifluoroprop-1-enyl)-2,2-dimethylcyclopropane-carboxylate,

[0083] Deltamethrin: (S)-alpha-cyano-3-phenoxynbenzyl (1R)-cis-3-(2,2-dibromovinyl)-2,2-dimethylcyclopropene-carboxylate,

[0084] Cycloprothrin: (RS)-alpha-cyano-3-phenoxynbenzyl (RS)-2,2-dichloro-4-(4-ethoxyphenyl)cylopropene-carboxylate,

[0085] Fluvalinate: alpha-cyano-3-phenoxynbenzyl N-(2-chloro-alpha, alpha, alpha, alpha-trifluorop-toly)-D-valinate,

[0086] Bifenthrin: (2-methylbenzophenyl-3,3-ethylmethyl)(Z)-(1RS)-cis-3-(2-chloro,3,3,3-trifluoroprop-1-enyl)-2,2-dimethylcyclopropene-carboxylate,

[0087] 2-methyl-2-(4-bromodifluoromethoxyphenyl)-propyl (3-phenoxynbenzyl)ether,
Tralomethrin: \((S)-\text{alpha-cyano-3-phenoxybenzyl (1R-cis)}(1'RS)(1', 2', 2', \text{2-tetrabromomethyl})-2,2\text{-dimethylcyclopropanecarboxylate},\)

Silaflofen: \(4\text{-ethoxyphenyl}(3-(4\text{-fluoro-3-phenoxyphenyl})\text{propyl})\)-dimethylsilane,

D-fenothrin: 3-phenoxycarbanilide (IR-cis), trans-chrysanthemate,

Cyphenothrin: (RS)-\(\text{alpha-cyano-3-phenoxybenzyl (1R-cis), trans-chrysanthemate, D-resmethrin: 5-benzyl-3-furylmethyl (1R-cis), trans-chrysanthemate,}\)

Actinathrin: (S)-\(\text{alpha-cyano-3-phenoxycarbonylbenzyl (1R-cis(Z))-2,2\text{-dimethylcyclopropanecarboxylate,}}\)

Cyfluthrin: (RS)-\(\text{alpha-cyano-4-fluoro-3-phenoxycarbonylbenzyl 3-(2,2-dichlorovinyl)-2,2\text{-dimethylcyclopropanecarboxylate,}}\)

Tetramethrin: \(3,4,5,6\text{-tetrahydrophthalimidomethyl (IRS-cis), trans-chrysanthemate,}\)

Allethrin: (RS)-3-allyl-2-methyl-4-oxocyclopent-2-enyl (1RS-cis), trans-chrysanthemate,

Prallethrin: (S)-2-methyl-4-oxo-3-(2-propynyl)cyclopent-2-enyl (1R-cis), trans-chrysanthemate,

Empenthrin: (RS)-1-ethylthio-2-methyl-2-penteny (1R-cis), trans-chrysanthemate,

Imiprothrin: 2,5-dioxo-3-(prop-2-yny)imidazolidin-1-ylmethyl (1R-cis), trans-2,2-dimethyl-3-(2-methyl-1-propenyl)cyclopropanecarboxylate,

D-flamethrin: 5-(2-propynyl)-furfuryl (1R-cis), trans-chrysanthemate, and 5-(2-propynyl)furfuryl 2,2,3,3-tetramethylcyclopropanecarboxylate;

Carbamate compounds such as

Alyanycarb: S-methyl-N\[N\text{-methyl-N-2-ethoxy-carbonylethyl}][\text{aminothio}][\text{carbamoyl}][\text{thioacetimide},

Bendiocarb: 2,2-dimethyl-1,3-benzodioxol-4-y1methylcarbamate,

Carbaryl: (1-naphthyl N-methylcarbamate,

Isocarbox: 2-(1-methylthio)phenyl methylcarbamate,

Carbosulfan: 2,3 dihydro-2,2-dimethyl-7-benzofuranyl[dibutylamino]thio)methylcarbamate,

Fenoxycode: Ethyl\[2-(4-phenoxycarbonyl)ethyl]carbamate,

Indoxacarb: Methyl-7-chloro-22,3,4\text{-tetrachloro-2-6(4-trifluoromethoxyphenyl)]

Propoxur: 2-isopropoxyphenol methylcarbamate,

Dirimicarb: Dimethyl N,N-thiobis((methy1imino)carbonyloxyl)bisethanamidothioate,

Methomyl: S-methyl N-(methylcarbamoyl)oxothioacetimidate,

Ethiofencarb: 2-((ethylthio)methyl)phenyl methylcarbamate,

Fenothiocarb: S-(4-phenoxybutyl)-N,N-dimethyl thiocarbamate,

Cartap: \(S,S'(2\text{-dimethylamino})\text{trimethylencarbamothioate)\)

Fenobucarb: 2-sec-butylphénylméthyl carba
date,

XMC: 3,5-dimethylphenyl-methyl carbamate,

Xylylcarb: 3,4-dimethylphenylmethylcarbamate,

organophosphorous compounds such as

Fenitrothion: O,O-dimethyl O-(4-nito-m-tolyl)phosphorothioate,

Diazinon: O,O-diyethyl-O-(2-isopropyl-6-methyl-4-pyrimidinyl)phosphorothioate,

Pyridaphenthion: O-(1,6-dihydro-6-oxo-1-phenylpyrazin-3-yl) O,O-diyethyl phosphorothioate,

Pirimiphos-Ethyl: O,O-diyethyl O-(2-diethylaminol6-methylpyrimidinyl)phosphorothioate,

Pirimiphos-Methyl: O-[2-(diethylamino)6-methyl-4-pyrimidinyl]O,O-dimethyl phosphorothioate,

Etriphos: O-6-ethoxy-2-ethylpyrimidin-4-yl-O-dimethyl phosphorothioate,

Fenthion: O,O-dimethyl O-[3-methyl-4-(methylthio)phenyl phosphorothioate,

Phoxim: 2-(4-chloroxyphosphinothioyl)iminomethylacetimidire,

Chlorpyrifos: O,O-diyethyl-O-(3,5,6-trichloro-2-pyryl)phosphorothioate,

Chlorpyrifos-methyl: O,O-dimethyl O-(3,5,6-trichloro-2-pyridyl)phosphorothioate,

Cyanophos: O,O-dimethyl O-(4-cyanophenyl)phosphorothioate,

Pyraclofos: (R,S)-[4-chlorophenyl]-pyrazol-4-yl-O-ethyl-S-n-propyl phosphorothioate,

Acephate: O,S-dimethyl acetylphosphoramo-midothioate,
Azamethiphos: S-(6-chloro-2,3-dihydro-oxo-1,3-oxazolyl[4,5-b]pyridine-3-ylmethyl phosphorothioate,
Malathion; O,O-dimethyl phosphorodithioate ester of diethyl mercaptosuccinate,
Temephos: O,O[(thiodi-4-1-phenylene) O,O O,O-tetramethyl phosphorodithioate,
Dimethoate: (O,O-dimethyl S-(α-methylcarbamoyl)ethyl]phosphorodithioate,
Formothion: S-[2-formylmethalamino]-2-oxoethyl]-O,O-dimethyl phosphorodithioate,
Phenthoate; O,O-dimethyl S-(α-hydroxy-carbamoyl)benzal]phosphorodithioate;
Insecticides with a stereiling effect on adult mosquitoes such as
1-(α,α-dichloro-α-cyclopropylbenzylidenamino-oxo)-p-tolyl]-3(2,6-difluorobenzyl)urea,
Diflubenzuron: N-((3,5-dichloro-4(1,2,2-tetrafluoroethoxy)phenylamino)carbonyl)2,6 difluoro benzamid,
Triflumuron: 2-Chloro-N-((4-(trifluoromethoxy)phenyl)-amino)carbonyl]benzamide, or a triazin such as N-cyclopropyl-1,3,5-triazine-2,4,6-triamine; and

Lambda-cyhalothrin:
α-cyano-3-phenoxybenzyl-3-(2-chloro-3,3,3-trifluoroprop-1-enyl)-2,2-dimethylcyclopropane carbonate, as a 1:1 mixture of (Z)-(1R,3R), R-ester and (Z)-(1S,3S), S-ester;
The repellent is selected from N,N-Diethyl-meta-toluidine (DEET), N,N-diethylphenylacetamide (DEPA), 1-(3-cyclobexan-1-yl-carbonyl)-2-methylpiperine, (2-hydroxymethylcyclohexyl) acetic acid lactone, 2-ethyl-i,3-hexandiol, indalone, Methyl-neodecanamide (MND), a pyrethroid not used for insect control such as (α)-3-allyl-2-methyl-4-oxocyclopent-2-(α)-enyl-(α)-trans-chrysanthemate (Essiothrin), a repellent derived from or identical with plant extracts like limonene, eugenol, (+)-Eucamol (1), (−)-1-epi-eucamol or crude plant extracts from plants like Eucalyptus maculata, Virex roundifolia, Cymbopogon martini, Cymbopogon citratus (lemon grass), Cymbopogon nardus (citronella).

Preferred insecticides and/or repellents of the insecticide composition of the present invention may be either one of a single insecticide and/or repellent or a mixture of insecticides and/or repellents selected from the group of insecticides and/or repellents that are suitable for application to a fabric material or a netting. This group of insecticides and/or repellents may include synthetic pyrethroids such as those known in the trade as alphacpermethrin, cyfluthrin, deltamethrin, ethofenprox and permethrin, other pyrethroids such as that known in the trade as bifenthrine and non-pyrethroids such as that known in the trade as carbosulphame.

If the insecticides and repellants mentioned above have one or more chiral centers in their molecules, they may be applied as racemates, pure enantiomers or diastereomers or in chirally or diastereomically enriched mixtures.

The insecticide and/or repellent mentioned in the present invention may also be included in the insecticide composition as one of a water-based insecticide and/or repellent concentrate or a solvent, preferably an organic solvent, based insecticide and/or repellent concentrate or a concentrate based on a mixture of water and a solvent, preferably an organic solvent. Water-based concentrates may be in the form of suspensions or dispersions comprising suitable dispersing agents if necessary or in the form of emulsions comprising emulsifiers, solvents and co-solvents if appropriate. Nanoparticle insecticidal formulations may be obtained by dissolving solid solutions of insecticides in a polar organic solvent, e.g. poly vinyl pyrrolidone (PVP). The concentration of the insecticide and/or repellent in the water based or solvent based concentrates is in general between 0.5 to 60%, preferably 1 to 40%, more preferably 3 to 20%.

The insecticide and/or repellent forming part of the insecticide composition of the present invention may also be selected from other groups suitable for different applications.

The particle size of the insecticide and/or repellent in water-based suspensions or dispersions is in general between 50 nm to 20 μm, preferably 50 nm to 8 μm, more preferably 50 nm to 4 μm, most preferably 50 nm to 500 nm.

Acrylic Binder (Component B1)

The acrylic binder is preferably obtainable by emulsion polymerization of

b1a) 10 to 90% by weight, preferably 15 to 80% by weight, more preferably 20 to 70% by weight of component B1A;
b1b) 10 to 90% by weight, preferably 12 to 85% by weight, more preferably 15 to 65% by weight of component B1B;
b1c) 1 to 5% by weight of component B1C;
b1d) 0 to 5% by weight preferably 1 to 4% by weight, more preferably 0.2 to 3% by weight of component B1D;
b1e) further monomers which are copolymerizable with the monomers mentioned above selected from
b1e1) 0 to 30% by weight, preferably 0 to 25% by weight, more preferably 5 to 20% by weight of component B1E1; and/or
b1e2) 0 to 40% by weight, preferably 0 to 30% by weight, more preferably 5 to 20% by weight of component B1E2;

wherein the sum of the components B1A, B1B, B1C and optionally B1D and B1E is 100% by weight.

The acrylic binder may comprise further additives as known by a person skilled in the art, for example film forming agents and plasticizers, e.g. adipate, phthalate, butyl diglycol, mixtures of diesters preparable by reaction of dicarboxylic acids and alcohols which may be linear or
branched. Suitable dicarboxylic acids and alcohols are known by a person skilled in the art.

[0163] The insecticide compositions comprising the specific binder as claimed in the present invention are wash resistant while permitting continuous release of the insecticide at a controlled rate, in order to provide the required bioavailability of the insecticide. It is not necessary to add for example a dispersing agent that, after application of the composition to a fabric and upon wetting of the fabric, reduces the hydrophobicity imparted to the insecticide by the binder to permit limited insecticide release. Preferably, the insecticide composition of the present invention does therefore not comprise a dispersing agent in addition to the acrylic binder.

[0164] Most preferably the acrylic binder is obtainable by emulsion polymerization of the following components:

[0165] b1a) 20 to 70% by weight of α-butylacrylate as component B1A;

[0166] b2b) 0 to 65% by weight of at least one monomer of formula I as component B1B

\[
\text{R}^2 \text{OR}^3 \quad \text{OR}^3 \quad \text{R}^1
\]

[0167] wherein

[0168] R¹ is H or methyl, R² is H and R³ is methyl, ethyl, or 2-ethylhexyl, as component B1B, most preferably component B1B is 2-ethylhexylacrylate, methylacrylate, methylnethacrylate or ethylacrylate;

[0169] b1c) 1 to 5% by weight of at least one monomer of formula II

\[
\text{R}^7 \text{N} \bigg| \text{R}^4 \text{OR}^5 \quad \text{OH} \quad \text{R}^4
\]

[0170] wherein R⁴ is H or methyl, R⁵, R⁶ and R⁷ each are H as component B1C;

[0171] b1d) 1 to 10% by weight, preferably 1 to 7% by weight, more preferably 2 to 5% by weight of at least one monomer of formula III

\[
\text{R}^8 \bigg| \text{R}^9 \text{X} \bigg| \text{OR}^{10}
\]

[0172] wherein R⁸ and R⁹ are H and X is H, OH, NH₂, OR¹OH, glycidyl or a group of the formula

\[
\text{O} \bigg| \text{R}^1 \text{C} \bigg| \text{O}
\]

[0173] wherein

[0174] R¹⁰ is selected from the group consisting of C₃ to C₁₀ alkyl which may be branched or linear, for example methyl, ethyl, n-propyl, i-propyl, butyl, isobutyl, sec-butyl, tert-butyl, n-pentyl, i-pentyl, sec-pentyl, neo-pentyl, 1,2-dimethylpropyl, i-amyl, n-hexyl, i-hexyl, sec-hexyl, n-heptyl, n-octyl, 2-ethylhexyl, n-nonyl, n-decyl; preferably C₁₋₃ to C₄₋₅ alkyl, which may be branched or linear, for example methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, sec-butyl and tert-butyl; substituted or unsubstituted aryl, preferably substituted or unsubstituted C₆₋₇ aryl, more preferably substituted or unsubstituted C₆₋₅ aryl, for example phenyl or tolyl;

[0175] R¹¹ is selected from the group consisting of C₁₋₃ to C₁₀₋₁₅ alkylene, for example methylene, ethylene, propylene, butylene, pentylene, hexylene, heptylene, octylene, nonylene, decylene; preferably C₁₋₃ to C₄₋₅ alkylene, for example methylene, ethylene, propylene, butylene; substituted or unsubstituted arylene, preferably substituted or unsubstituted C₆₋₇ arylene, more preferably substituted or unsubstituted C₆₋₅ arylene, for example phenylene;

[0176] as component B1D, most preferably X is acetacryl;

[0177] b1e) further monomers which are copolymerizable with the monomers mentioned above selected from

[0178] b1e1) 0 to 30% by weight, preferably 0 to 25% by weight, more preferably 5 to 20% by weight of component B1E1, preferably (meth-)acryl nitrile and/or methyl(meth)acrylate;

[0179] and/or

[0180] b1e2) 0 to 40% by weight, preferably 0 to 30% by weight, more preferably 5 to 20% by weight of component B1E2, preferably styrene and/or a-methylstyrene;

[0181] wherein the sum of components B1A, B1B, B1C and optionally B1D and B1E is 100% by weight.
The acrylic binder of the present invention is obtainable by emulsion polymerization of the monomers mentioned before. Suitable process conditions are known by a person skilled in the art.

The monomers are polymerized under usual conditions of temperature and pressure, i.e. at from atmospheric pressure to 10 bar and in general at temperatures of from 20 to 100°C, preferably 50 to 85°C, depending on the initiator used. Usually the polymerization is carried out in a stirred reaction vessel under an inert atmosphere.

The copolymerization is generally carried out in water. However, it is also possible to add before, within or after the polymerization process up to 80% by weight, relating to the aqueous phase, of a lower alcohol like methanol, ethanol or isopropanol or a lower ketone like acetone. Preferably the copolymerization is carried out in water without addition of further solvents.

The polymerization process may be carried out continuously or batch-wise, and it is possible to employ the usual methods of batch-wise polymerization, e.g. mixing all polymerization components at once or feeding emulsified monomers and catalysts from one or more metering vessels to a batch containing a portion of a monomer. It is possible to add polymer seed to the polymerization mixture to adjust the particle size of the emulsion polymers obtained.

The emulsion polymerization is preferably carried out in the presence at least one initiator which form radicals under the polymerization conditions. Suitable initiators are for example all common peroxy compounds or azo compounds.

Suitable peroxy compounds are for example alkali metal peroxodisulfates, ammonium persulfate; hydrogen peroxide; organic peroxy compounds, for example diacetyl peroxide, di-t-butyl peroxide, diacyl peroxide, diocanoyl peroxide, diene-conjugated peroxide, dilauroyl peroxide, dibenzoyl peroxide, bis(o-tolyl)peroxide, succinyl peroxide, terti-butyl peracetate, tert-butyl permalinate, tert-butyl peripivale, tert-butyl peroctoate, tert-butyl perenecanolate, tert-butyl perbenzoate, tert-butyl peroxide, tert-butyl hydroperoxide, cumene hydroperoxide, tert-butyl-peroxy-2-ethyhexanoate, and disopropyl peroxodicarboxformate. Further suitable initiators are azo compounds, for example azobis(isobutyronitrile), azobis(2-amidropropane)dihydrochloride, and 2,2'-azo-bis(2-methylbutyronitrile).

The initiators are added in usual amounts, for example in amount of 0.05 to 5% by weight, preferably 0.05 to 2% by weight, based on the total weight of monomers.

If the polymerization is carried out at low temperature, use may be made of conventional redox catalysts. For example, it is possible to use, in addition to the peroxide catalysts of the above kinds, from 0.05 to 2% by weight, based on the total of monomers, of reducing agents such as hydrazine, soluble oxidizable sulfuryl compounds such as alkali metal salts of hydrosulfurilates, sulfoxylates, thiocarboxylates, sulfites, and bisulfites, which may be optionally activated by the addition of traces of heavy metals, e.g. salts of Ce, Mo, Fe, and Cu, in the usual manner. Preferred redox catalysts are redox catalysts of acetone dinsulfite and organic peroxydes like tert-C4Hg—OOH; Na2S2O5 and organic peroxydes like tert-C4Hg—OOH; or HO—CH3SO2H and organic peroxydes like tert-C4Hg—OOH. Further preferred are redox catalysts like ascorbic acid and hydrogen peroxide.

The initiator may be added completely at the beginning of the polymerization, but it is also possible to add the initiator in the course of the emulsion polymerization process in a continuous or stepwise way. The way of adding the initiator is known in the art.

The polymerization process is carried out until a conversion of at least 90% by weight of the monomers is reached. For removal of the residual monomer at the end of the emulsion polymerization initiator may be added for chemical deoxidation.

The emulsion polymerization is carried out by adding emulsifiers or mixtures of emulsifiers known in the art. The emulsifiers generally used are ionic (anionic or cationic) and/or non-ionic emulsifiers such as polyglycol ethers, sulfonated paraffin hydrocarbons, higher alkylsulfates such as oleyl amine, laurylsulfate, alkali metal salts of fatty acids such as sodium steareate and sodium oleate, sulfuric acid esters of fatty alcohols, ethoxylated C12-15-alkylphenols, usually having from 5 to 30 ethylene oxide radicals, and their saponification products, and also sulfosuccinimide acid esters. The emulsifier or mixtures of emulsifiers are usually employed in an amount of 0.05 to 7% by weight, preferably 0.5 to 4% by weight, based on the total weight of monomers.

In some cases there is added a co-solvent or a mixture of co-solvents to the emulsifiers. Preferred co-solvents are aliphatic C1- to C30-alcohols which are linear or branched, alicyclic C2- to C30-alcohols and mixtures thereof. Examples are n-butanol, n-hexanol, cyclohexanol, 2-ethylhexanol, i-octanol, n-octanol, n-decanol, n-dodecanol, stearyl alcohol, oleyl alcohol or cholesterol. Further possible co-solvents are alkane diol, ethylene glycol alkyl ethers, N-alkyl pyrrolidones, and N-alkyl and N,N-dialkyl amides like ethylene glycol monobutyl ether, diethyl glycol monoethyl ether, tetraethylen glycol dimethyl ether, N-methyl pyrrolidone, N-hexyl pyrrolidone, di-ethyl amide or N-oxacyl amide acid. The co-solvents or mixture of co-solvent is added in an amount of 0 to 20% by weight, preferably 1 to 5% by weight.

In many cases use is also made of a protective colloid, examples of which are polyvinyl alcohol, partially saponified polyvinyl acetates, cellulose derivatives, copolymers of methyl acrylate with acrylic amide and methacrylic amide or vinyl pyrrolidone polymers in amounts of from 0.5 to 10% by weight and in particular 1.0 to 5% by weight of the weight of the monomers.

Further, it is possible to add in general up to 10% by weight, preferably 0.05 to 5% by weight of mono- or di-olefinically unsaturated monomers containing reactive or cross-linking groups. Examples of such monomers are in particular the amides of α,ω-olefinically unsaturated C3-5-carboxylic acids, particularly acrylic amides, methacrylic amides and maleic diamides, and their N-methylol derivatives such as N-methacryl acrylamide, N-methyl methacrylic amide, N-alkoxy methyl amides of α,ω-monomethacrylic unsaturated C3-5-carboxylic acids such as N-methoxy methacrylamide and N-n-butoxymethylacrylamide, vinyl sulfonic amide, monoesters of acrylic and methacrylic
acids with alkanediols such as glycol, butanediol-1,4, hexane diol-1,6, and 3-chloropropanediol-1,2, and also allyl and methallyl esters of α,β-olefinically unsaturated mono- and di-carboxylic acids such as diallyl maleate, dimethyl allyl fumarate, allyl acrylate and allyl methacrylate, diallyl phthalate, diallyl terephthalate, p-di-vinyl benzene, methylene-bis-acrylamide and ethylene glycol di-allyether.

The solids content of the aqueous dispersions of polymers obtained in the emulsion polymerization is usual 15 to 75% by weight, preferably 25 to 50% by weight. To obtain high space time yields of the reactor dispersions having a high solids content are preferred. To obtain solids contents of more than 60% by weight a bi- or polymodal particle distribution should be adjusted, because otherwise it is not possible to handle the dispersion, because of the high viscosity. New particle generations (for obtaining bi- or polymodal particle size distributions) are for example formed by addition of seed (EP-A-0810 S31), addition of an excess of emulsifier or addition of mini-emulsions. The formation of new particle generations may be carried out at any time and is depending on the desired particle size distribution for a low viscosity.

The molecular weight of the non crosslinked emulsion polymers obtained is in general 40,000 to 250,000 (determined by GPC). The molecular weight is usually controlled by the use of conventional chain stoppers in conventional amounts. Conventional chain stoppers are for example sulforganic compounds.

The acrylic binder of the present invention is obtained in form of its aqueous dispersion and is preferably employed in the insecticide compositions of the present invention in form of the aqueous dispersion.

Polyurethane (Component B)

The polyurethane is preferably obtainable by reaction of the following components;

b2a) 55 to 99% by weight, preferably 70 to 98% by weight, more preferably 75 to 90 by weight based on the polyurethane of at least one diisocyanate or polyisocyanate (component B2a), preferably aliphatic, cycloaliphatic, aliphatic and/or aromatic Isocyanates, more preferably diisocyanates, which are optionally biuretized and/or isocyanurized, more preferably alkylene diisocyanates having from 4 to 12 carbon atoms in the alkyne unit, like 1,12-dodecane diisocyanate, 2-ethylhexamethylene diisocyanate-1,4, 2-methylpentamethylene diisocyanate-1,5, tetramethylene diisocyanate-1,4, lysine diisocyanate (LID), hexamethylene diisocyanate-1,4, cyclohexane-1,3- and/or 1,4-diisocyanate, 2,4 and 2,6-hexahydro-toluylene diisocyanate as well as the corresponding isomeric mixtures 4,4',2,2' and 2,4'-dicyclohexylmethane diisocyanate as well as the corresponding mixtures, 1-isocyanato-3,3,5-trimethyl-5-isocyanatomethylene cyclohexane (IPDI), 2,4- and 2,6-toluylene diisocyanate, 4,4', 2,4 and 2,2'-diphenylmethane diisocyanate (monomeric MDI), polyphenylpolyisocyanate polyisocyanate (polymeric MDI) and/or mixtures comprising at least 2 of the isocyanates mentioned before; further esters, ureas, aliphosphate-, carbodiimid-, ureidone- and urethane groups comprising di- and/or polyisocyanates may be used; most preferably 1-isocyanato-3,3,5-trimethyl-5-isocyanatomethylene cyclohexane (IPDI) and hexamethylene diisocyanate-1,6 (HMDI);

b2b) 10 to 90% by weight, preferably 12 to 85% by weight, more preferably 15 to 65% by weight based on the polyurethane of at least one diol, triol or polyol (component B2B), preferably aliphatic, cycloaliphatic and/or aliphatic diols having 2 to 14, preferably 4 to 10 carbon atoms, more preferably polyols, selected from the group consisting of polyethers, e.g. polytetrahydrofurane, polyethers, polyethyetherpolys, hydroxy group containing polyacetales and hydroxyl group containing aliphatic polycarbonates or mixtures of at least 2 of the polyols mentioned before. Preferred are polyetherols and/or polyethers. The hydroxyl number of the polyhydroxy compounds is in general from 20 to 850 mg KOH/g and preferably 25 to 80 mg KOH/g. Further, diols and/or triols having a molecular weight of from in general 60 to <400, preferably from 60 to 300 g/mol are employed. Suitable diols are aliphatic, cycloaliphatic and/or aliphatic diols having from 2 to 14, preferably 4 to 10 carbon atoms, e.g. ethylene glycol, propane diol-1,3, decane diol-1,10, -o-, m-, p-dihydroxy-cyclohexane, diethylene glycol, dipropylene glycol and preferably butane diol-1,4, neopentyl glycol, hexane diol-1,6 and bis-(2-hydroxy-ethyl)hydroquinone, triols, like 1,2,4-, 1,3,5-trihydroxy-cyclohexane, glycerine and trimethyl propane and mixtures of low molecular hydroxyl groups containing polyalkylene oxides based on ethylene oxide and/or 1,2-propylene oxide and the diols and/or triols mentioned before;

b2c) 0 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 1 to 5% by weight based on the polyurethane of further components (component B2C), preferably adipic acid or carbonyl diimide (CDI); and

b2d) 0 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 0.5 to 5% by weight based on the polyurethane of further additives (component B2D);

wherein the sum of the components B2A, B2B, B2C and B2D is 100% by weight

The polyurethanes are prepared by methods known in the art. Further, additives as known by a person skilled in the art may be used in the process for preparing the polyurethanes.

Insecticide Compositions

Depending on the use of the final product the insecticide composition of the present invention may further comprise one or more components selected from water, preservatives, detergents, fillers, impact modifiers, anti-foaming agents, blowing agents, clarifiers, nucleating agents, coupling agents, conductivity-enhancing agents (antistats), stabilizers such as anti-oxidants, carbon and oxygen radical scavengers and peroxide decomposing agents and the like, flame retardants, mould release agents, agents having UV protective properties, optical brighteners, spreading agents, anti-blocking agents, anti-migrating agents, foam-forming
agents, anti-soiling agents, thickeners, further biocides, wetting agents, plasticizers and film forming agents, adhesive or anti-adhesive agents, optical brightening (fluorescent whitening) agents, pigments and dye-stuffs.

[0209] The insecticide compositions of the present invention may be aqueous compositions comprising water or dry compositions, e.g. compositions which do not comprise water. Preferably, the insecticidal compositions are aqueous compositions, preferably comprising 0.1 to 45% by weight, more preferably 1 to 25% by weight of water, based on the total of the components in the insecticidal compositions of the present invention except of water.

[0210] Suitable anti-foam agents are for example silicon anti-foam agents. Suitable UV-protecting agents for protecting UV-sensitive insecticides and/or repellents are for example para-aminobenzoic acids (PABA), octylmethoxy-sinameth, stilbenes, styryl or benzotriazole derivatives, benzoazol derivatives, hydroxy-substituted benzophenones, salicylates, substituted triazines, cinnamic acid derivatives (optionally substituted by 2-cyano groups), pyrazoline derivatives, 1,1'-biphenyl-4,4-bis-2-(methoxyphenyl)eth- nyl or other UV protecting agents. Suitable optical brighteners are dihydroquinoline derivatives, 1,3-diaryl pyrazoline derivatives, pyrenes, napthalic acid imides, 4,4'-dipy- styryl, biphénylene, 4,4'-diamino-2,2'-stilbene disulphonic acids, cumarin derivatives and benzoxazole, benzoxazol or benzimidazole systems which are linked by —CH═CH-bridges or other fluorescent whitening agents.

[0211] Typical pigments used in the insecticidal compositions of the present invention are pigments which are used in pigment dyeing or printing processes or are applied for the coloration of plastics and are known by a person skilled in the art.

[0212] Pigments may be inorganic or organic by their chemical nature. Inorganic pigments are mainly used as white pigments (e.g., titanium dioxide in the form of rutile or anatas, ZrO, chalk) or black pigments (e.g., carbon black). Colored inorganic pigments may be used as well but are not preferred because of potential toxicologic hazards. For imparting color, organic pigments or dye-stuffs are preferred. Organic pigments may be mono or disazo, naphthol, benzimidazolone, (thio) indigoid, dioxazine, quinacridone, phthalocyanine, isoindolinone, perylene, perinone, metal complex or diketo pyrrolo pyrrole type pigments. Pigments may be used in powder or liquid form (i.e., as a dispersion). Preferred pigments are Pigment Yellow 83, Pigment Yellow 138, Pigment Orange 34, Pigment Red 170, Pigment Red 146, Pigment Violet 19, Pigment Violet 23, Pigment Blue 15/1, Pigment Blue 15/3, Pigment Green 7, Pigment Black 7. Other suitable pigments are known to a person skilled in the art.

[0213] Typical dye-stuffs which may be used in the present invention are vat dyes, cationic dyes and disperse dyes in powder or liquid form. Vat dyes may be used as pigments or following the vatting (reduction) and oxidation procedure. Using the vat pigment form is preferred. Vat dyes may be of the indanthrone type, e.g. C.I. Vat Blue 4, 6 or 14; or of the flavanthrone type, e.g. C.I. Vat Yellow 1; or of the pyran-throne type, e.g. C.I. Vat Orange 2 and 9; or of the isoben- zanthrone (isoviolanthrone) type, erg. C.I. Vat Violet 1; or of the dibenzanthrone (violanthrone) type, e.g. C.I. Vat Blue 16, 19, 20 and 22, C.I. Vat Green 1, 2 and 9, C.I. Vat Black 9; or of the anthraquinone carbazole type, e.g. C.I. Vat Orange 11 and 15, C.I. Vat Brown 1, 3 and 44, C.I. Vat Green 8 and C.I. Vat Black 27; or of the benzanthrone acridone type, e.g. C.I. Vat Green 3 and 13 and C.I. Vat Black 25; or of the anthraquinone oxazole type, e.g. C.I. Vat Red 10; or of the perylene tetra carbonic acid diimide type, e.g. C.I. Vat Red 23 and 32; or amidazole derivatives, e.g. C.I. Vat Yellow 46; or amino triazine derivatives, e.g. C.I. Vat Blue 66. Other suitable vat dyes are known by a person skilled in the art.

[0214] Typical disperse and cationic dye-stuffs are known by a person skilled in the art.

[0215] In a further embodiment the insecticidal compositions of the present invention are insecticide compositions as mentioned before comprising at least one pigment and or at least one dyestuff. The insecticidal compositions of the present invention preferably comprise 10 to 200% by weight, more preferably 20 to 150% by weight of the pigment and/or dyestuff relating to the total weight of the solids content of the insecticide and/or repellent.

[0216] Textile material or plastics material to be Impregnated according to the present invention and to be use of a composition as described in the present invention may be impregnated locally when the composition is delivered in the form a kit comprising the ingredients of the insecticidal composition in a handy form. In a further embodiment the present invention therefore relates to an insecticidal composition as described in the present invention which is provided as a kit for impregnation by the end-user. In a preferred embodiment the kit is adapted for preparing a solution or emulsion by adding water. The ingredients of the kit may accordingly be in form of a dry composition such as a powder, a capsule, a tablet, or an effervescent tablet. In a further embodiment, the kit comprises an emulsion wherein water is added by the end-user. The emulsion may be a micro-emulsion, which is generally very stable. The emulsion may be embodied in a capsule.

[0217] The kit comprises at least the following ingredients:

[0218] a) at least one insecticide and/or at least one repellent, and

[0219] b) at least one acrylic binder as described in the present invention;

[0220] and/or

[0221] b) at least one polyurethane as described in the present invention.

[0222] Preferred insecticides and/or repellents as well as preferred acrylic binders and preferred polyurethanes are already mentioned in the present invention. The kit may obtain further ingredients as mentioned above, especially one or more compounds selected from preservatives, detergents, stabilizers, agents having UV-protecting properties, optical brighteners, spreading agents, anti-migrating agents, foam-forming agents, wetting agents, anti-soiling agents, thickeners, further biocides, plasticizers, adhesive agents, pigments and dye-stuffs. Preferred kits comprise beside the insecticide and/or repellent and the acrylic binder and/or the polyurethane at least one pigment and/or at least one dye-stuff. Preferred pigments and dye-stuffs are mentioned before.
[0223] In a further embodiment the present invention relates to an impregnated textile material or plastics material for insect killing and/or repellence of an insect comprising

[0224] a) at least one insecticide and/or at least one repellent, and

[0225] b1) at least one acrylic binder as disclosed in the present invention;

[0226] and/or

[0227] b2) at least one polyurethane as disclosed in the present invention.

[0228] Preferred insecticides and/or repellents and preferred acrylic binders and polyurethanes are mentioned before. Materials of the textile material or plastics material and preferred textile material or plastics material are also mentioned before.

[0229] A typical amount of insecticide and/or repellent in the impregnated textile material or plastics material is from 0.01 to 10% (dry weight) of the (dry) weight of the textile material or plastics material depending on the insecticidal efficiency of the insecticide respectively the efficiency of the repellent. A preferred amount is between 0.05 and 7% by weight of the textile material or plastics material depending on the insecticide and/or repellent. For a pyrethroid like deltamethrin or alphacypermethrin, the preferred amounts are between 0.1 and 3.5% of the weight of the textile material or plastics material. For a pyrethroid like permethrin or ethofenprox, the preferred amount is from 0.1 to 6%.

[0230] A typical amount for the acrylic binder and/or the polyurethane is from 0.001 to 10% by weight (dry weight) of the (dry) weight of the textile material or plastics material.

As a rule, the higher amount the insecticide of the specific type to be added, the higher the concentration of the acrylic binder and/or the polyurethane so that the ratio between insecticide and acrylic binder and/or the polyurethane is approximately constant with a value depending on the insecticidal and migratory ability of the insecticide. Preferred amounts of acrylic binders and/or polyurethane are from 0.1 to 5% by weight, more preferably 0.2 to 3% by weight of the (dry) weight of the textile material or plastics material.

[0231] In a further embodiment, the impregnated textile material or plastics material according to the present invention further comprises one or more components selected from preservatives, detergents, stabilizers, agents having UV-protecting properties, optical brighteners, spreading agents, anti-migrating agents, foam-forming agents, wetting agents, anti-soiling agents, thickeners, further biocides, plasticizers, adhesive agents, pigments and dyestuffs. Suitable examples of the components mentioned above are known by a person skilled in the art.

[0232] In a further embodiment of the present invention the impregnated textile material or plastics material comprises besides the at least one insecticide and/or repellent and the at least one acrylic binder and/or polyurethane as described before and/or at one pigment and/or at least one dyestuff. The amount of the at least one pigment is in general from 0.05 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 0.2 to 3.5% by weight of the (dry) weight of the textile material or plastics material. The amount of the at least one dyestuff is in general from 0.05 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 0.2 to 3.5% by weight of the (dry) weight of the textile material or plastics material. The textile material or plastic material comprises preferably either at least one pigment or at least one dyestuff. Suitable pigments and dyestuffs are mentioned before.

[0233] Process for Impregnation of a Textile Material or Plastics Material

[0234] In a further embodiment, the present invention relates to a process for impregnation of a textile material or plastics material comprising

[0235] i) forming an aqueous formulation comprising at least one insecticide and/or at least one repellent and at least one acrylic binder anchor at least one polyurethane as defined in the present invention and optionally further ingredients;

[0236] ii) applying the aqueous formulation to the textile material or plastics material by

[0237] iia) passing the textile material or plastics material through the aqueous formulation;

[0238] or

[0239] iib) bringing the textile material or plastics material in contact with a roller that is partly or fully dipped into the aqueous formulation and drawing the aqueous formulation to the side of the textile material or plastics material in contact with the roller;

[0240] or

[0241] iic) double-side coating of the textile material or plastics material

[0242] or

[0243] iid) spraying the aqueous formulation onto the textile material or plastics material;

[0244] or

[0245] iie) applying the aqueous formulation in form of a foam;

[0246] or

[0247] iif) submerging the textile material or plastics material into the aqueous formulation;

[0248] or

[0249] iig) brushing the aqueous formulation onto or into the textile material or plastics material;

[0250] or

[0251] iih) pouring the aqueous formulation onto the textile material or plastics material;

[0252] iii) optionally removing surplus aqueous formulation; and

[0253] iv) drying and/or curing the textile material or plastics material.

[0254] In the context of the present invention an aqueous formulation may be a solution, an emulsion or a suspension/dispersion.
The aqueous formulation preferably comprises the insecticide composition as disclosed in the present invention, which is preferably employed in form of an aqueous formulation.

In the context of the present invention "impregnation" is a process for applying the insecticide composition. This process may include a process for curing the applied insecticide composition to achieve a coating onto the textile material or plastics material, if desired. An "impregnated textile material or plastics material" is a material onto which the insecticide composition is applied. The "impregnated textile material or plastics material" may be coated by curing the applied insecticide composition, if desired.

Suitable textile materials or plastics materials are mentioned before. Preferred insecticides and/or repellents used are also defined before as well as preferred acrylic binders and/or polyurethanes.

It is one advantage of the present invention that the impregnation is carried out in an aqueous formulation. It is not necessary to add any organic solvents. In a preferred embodiment of the present invention the treatment bath is a an aqueous formulation which does not comprise any further solvents, especially no organic solvents.

It is advantageous to avoid the use of organic solvents because the textile material or plastics material of the present invention may be used in close contact to the human body. The textile material or plastics material impregnated by the method of the present invention does not comprise any organic solvent residue which is beneficial both from a human toxicological and an ecological point of view.

The aqueous formulation employed in the impregnation process may further comprise one or more ingredients selected from the group consisting of preservatives, detergents, stabilisers, agents having UV-protecting properties, spreading agents, anti-migrating agents, foaming agents, wetting agents, anti-soiling agents, thickeners, further biocides, plasticisers, adhesive agents, pigments and dyestuffs.

In a further embodiment the aqueous formulations employed for impregnation of the textile material or plastics material comprise in addition to the at least one insecticide and/or repellent the at least one acrylic binder and/or at least one polyurethane at least one pigment and/or at least one dyestuff. These aqueous formulations are suitable for impregnation of the textile material or plastics material with at least one insecticide and/or repellent and additionally coloring the textile material or plastics material at the same time. Many of the textile materials or plastics materials which are impregnated with at least one insecticide and/or repellent are preferably colored. With the process of the present invention it is possible to color and impregnate the textile material or plastics material with an insecticide and/or repellent at the same time. The method of the present invention is therefore very economical, because the coloring and impregnation with an insecticide and/or repellent is carried out in one step.

In a further embodiment of the present invention the present invention therefore relates to a process for impregnation of a textile material or plastics material as described before, wherein the dyeing of the textile material or plastics material is carried out simultaneously with the impregnation of the textile material or plastics material, wherein an aqueous formulation is formed further comprising at least one dyestuff and/or at least one pigment.

Suitable amounts of pigments are in general 0.01 to 20% by weight, preferably 0.1 to 10% by weights, more preferably 0.2 to 5% by weight, based on the weight of the aqueous formulation used for impregnation. Suitable dyestuffs are employed in general in an amount of from 0.01 to 20% by weight, preferably 0.1 to 10% by weight, more preferably 0.2 to 5% by weight, based on the weight of the aqueous formulation used for impregnation.

Step i) Forming an Aqueous Formulation Comprising an Insecticide and/or Repellent and at Least One Acrylic Binder and/or at Least One Polyurethane and Optionally Further Ingredients

The aqueous formulation is formed by mixing all ingredients necessary for impregnation of the textile material or plastics material with water. The aqueous formulation is generally formed at temperatures of from 10 to 70°C, preferably 15 to 50°C, more preferably 20 to 40°C. Suitable aqueous formulations comprise an insecticide composition for application to a textile material or plastics material as described in the present invention.

Step ii) Applying the Aqueous Formulation Comprising an Insecticide and/or Repellent and at Least One Acrylic Binder and/or at Least One Polyurethane and Optionally Further Ingredients to the Textile Material or Plastics Material

The insecticide composition is applied by passing the textile material or plastics material through the aqueous formulation. This step is known by a person skilled in the art as padding. In a preferred embodiment the textile material or plastics material is completely submerged in the aqueous treatment liquor (aqueous formulation) either in a trough containing the liquor or passed through the treatment bath (aqueous formulation) which is held between two horizontally oriented rollers.

In accordance with the invention, the textile material or plastics material may either be passed through the aqueous formulation or the aqueous formulation may be passed through the textile material or plastics material. These processes are preferred for impregnating open-width material which is later tailored into nets. For small-scale production or re-impregnating of non-treated nets, use of a simple hand-held roller might be sufficient.

Step iii)
Knife-/doctor-blade-coating systems are for example knife-over-air-systems, knife-over-roller systems, knife-over-table systems or knife-over-rubber-belt systems. Further knife coating systems are for example comminutor or Mayerbar knife systems.

Roller-coating systems are for example kiss-coating systems with one, two, three or more rollers, reverse-roll-coater systems and raster roll systems. In these roller-coating systems at least one roller is partly dipped into the aqueous formulation thus applying the aqueous formulation to the side of the textile material or plastics material in contact with the roller (kiss-rolling).

Screen-printing systems are for example rotary-screen printing systems and flat-screen printing systems. With these applications methods a dot coating or a full-surface coating can be applied to the textile material or the plastics material, for example by using an additional whisker-blade behind the rotary-screen.

Step iic)

It is further possible to apply the aqueous formulations to the textile material or plastics material by double-side coating application methods, for example double-side knife-coating systems, foulard with two air-knives or foulard with squeezing rollers.

By using the double-side-knife-coating system or by passing two times through a one-side-only coating system, it is possible to apply the aqueous formulation only to the surfaces of both sides of the textile material or plastics material and thus to reduce the quantity of the aqueous formulations needed for a certain effect.

Step iid) and Step iie)

It is further possible to apply the aqueous formulation by spraying the solution or emulsion onto the textile material or plastics material. Further it is possible, to apply the emulsion in the form of a foam which is applied to the textile material or plastics material. A foam comprises less water than the solution or emulsion mentioned above. The drying process may therefore be very short.

Step iif), iig) and Step iii)

It is further possible to apply the aqueous formulation on the textile material or plastics material by submerging the textile material or plastics material into the aqueous formulation, brushing the aqueous formulation onto or into the textile material or plastics material, or pouring the aqueous formulation onto the textile material or plastics material. Said methods are known by a person skilled in the art.

Impregnation of the fabric material or netting in step iia), iib), iic), iid), iie), iif), iig), or iii) is carried out at temperatures of in general from 10 to 70 °C, preferably 15 to 50 °C, more preferably 20 to 40 °C.

Step iii) Optionally Removing the Surplus Aqueous Formulation

The surplus aqueous formulation is usually removed by squeezing the textile material or plastics material, preferably by passing the textile material or plastics material rollers as known in the art, preferably by means of doctor blade, thus achieving a defined liquor uptake. The squeezed-off liquor is usually re-used.

The surplus aqueous formulation may alternatively be removed by centrifuging or vacuum suction.

Step iv) Drying and/or Curing the Textile Material or Plastics Material

The drying is in general carried out temperatures below 200 °C. Preferred temperatures are from 50 to 170 °C, more preferably from 70 to 150 °C. The temperature choice is a function of the evaporation temperature and mobility of the insecticide in the formulation.

It should be noted that the drying process may be a passive drying as the process may be carried out in rather hot climates. An active drying process would normally be performed during high scale processing.

After or simultaneously to the drying, the impregnated textile material or plastics material is optionally finally cured and/or fixed at. A person skilled in the art knows how to carry out a curing and/or fixation. The curing process in general carried out at a temperature which may be higher than the drying temperature. Preferred temperatures for curing are 70 to 170 °C, more preferably 80 to 150 °C. Drying and curing can be advantageously be performed during one single process, e.g. in stenter with different compartments which can be heated to different temperatures. If a reactive crosslinking agent is used temperatures may be lower.

The curing process may also include or consist of passing the textile material or plastics material by a heated surface under pressure such as an iron or a heated roller. During drying processes and curing the textile material or plastics material is preferably mechanically fixed in a way to prevent change of the form e.g. shrinkage or dimensional deformation. Further, is prevented that the insecticide and/or repellent is washed out.

The acrylic binder and the polyurethane may advantageously be amplified with a fixative agent for improved attachment of the insecticide and/or repellent on the textile material or plastics material. The fixative agent may comprise free isocyanate groups.

As described above, the solution or emulsion may further comprise one or more components selected from water, preservatives, detergents, fillers, impact modifiers, anti-fogging agents, blowing agents, clarifiers, nucleating agents, coupling agents, conductivity-enhancing agents (antistats), stabilizers such as anti-oxidants, carbon and oxygen radical scavengers and peroxide decomposing agents and the like, flame retardants, mould release agents, agents having UV protecting properties, spreading agents, anti-blocking agents, anti-migrating agents, foam-forming agents, anti-soiling agents, thickeners, further biocides, wetting agents, plasticizers, adhesive or anti-adhesive agents, optical brightening (fluorescent whitening) agents, pigments and dyestuffs.

The process may also involve using the kit as described before, accordingly, the impregnation process may be carried out by the end-user in a low-scale process. The present invention therefore relates to a process for impregnating a fabric material or netting as described
before, wherein the impregnating composition is provided as a kit for impregnation by the end-user.

[0294] In a further embodiment of the present invention the impregnation process which comprises applying an insecticide composition as described before may also take place before the fibers are woven or knitted.

[0295] In a further embodiment the invention relates to a process for coating a textile material or plastics material by applying a composition comprising at least one insecticide and/or at least one repellant and at least one acrylic binder and/or at least one polyurethane as defined in the present invention to the textile material or plastics material. The coating is preferably carried out in a doctor-blade process. The process conditions are known by a person skilled in the art.

[0296] Preferred compositions for coating a textile material or plastics material and preferred further ingredients of the composition are already mentioned above.

[0297] In a further embodiment the present invention relates to an exhaust process for impregnation of a textile material or plastics material comprising the steps

[0298] i) placing the textile material or plastics material in an aqueous bath optionally comprising further additives contained in a pressure-proof vessel;

[0299] ii) adding an aqueous formulation comprising at least one insecticide and/or repellent;

[0300] iii) heating the aqueous bath to a temperature of 100 to 140°C and keeping the temperature for 20 to 120 minutes;

[0301] and

[0302] iv) cooling and draining the bath, and rinsing and drying the impregnated textile material or plastics material.

[0303] It was found by the inventors that the insecticide and/or repellent is not washed out and the bioavailability of the insecticide and/or repellent for killing insects is maintained after multiple washes by carrying out the impregnation of a textile material or plastics material by the process mentioned above, even in absence of a binder or a polyurethane. The aqueous bath and the aqueous formulation comprising at least one insecticide and/or repellent preferably do not comprise a binder or a polyurethane.

[0304] Suitable textile materials and plastics materials are the textile materials and plastics materials described before. Suitable insecticides and/or repellents are also mentioned before.

[0305] Suitable further additives are preferably selected from preservatives, detergents, fillers, impact modifiers, anti-fogging agents, blowing agents, clarifiers, nucleating agents, coupling agents, conductivity-enhancing agents (anistats), stabilizers such as antioxidants, carbon and oxygen radical scavengers and peroxide decomposing agents and the like, flame retardants, mould release agents, agents having UV protecting properties, optical brighteners, spreading agents, anti-blocking agents, anti-migrating agents, foam-forming agents, anti-soiling agents, thickeners, further biocides, wetting agents, plasticizers and film-forming agents, adhesive or anti-adhesive agents, optical brightening (fluorescent whitening) agents, pigments and dyes.

[0306] Preferred further additives from the group mentioned above are already mentioned before.

[0307] The present invention therefore preferably relates to a process as mentioned above, wherein the aqueous bath comprises as further additives one or more components selected from preservatives, detergents, fillers, impact modifiers, anti-fogging agents, blowing agents, clarifiers, nucleating agents, coupling agents, conductivity-enhancing agents (anistats), stabilizers such as anti-oxidants, carbon and oxygen radical scavengers and peroxide decomposing agents and the like, flame retardants, mould release agents, agents having UV protecting properties, optical brighteners, spreading agents, anti-blocking agents, anti-migrating agents, foam-forming agents, anti-soiling agents, thickeners, further biocides, wetting agents, plasticizers and film-forming agents, adhesive or anti-adhesive agents, optical brightening (fluorescent whitening) agents, pigments and dyes.

[0308] The insecticide and/or repellent forming part of the aqueous formulation added in step ii) also may be in form of a water-based insecticide and/or repellent concentrate or a solvent, preferably an organic solvent, based insecticide and/or repellent concentrate or a concentrate based on a mixture of water and a solvent, preferably an organic solvent. Water-based concentrates may be in the form of suspensions or dispersions comprising suitable dispersing agents if necessary or in the form of emulsions comprising emulsifiers, solvents and co-solvents if appropriate. Nanoparticle insecticidal formulations may be obtained by dissolving solid solutions of insecticides in a polar organic solvent, e.g. polyvinyl pyrrolidone (PVP). The concentration of the insecticide and/or repellent in the water based solvent based concentrates is in general between 0.5 to 60%, preferably 1 to 40%, more preferably 3 to 20%.

[0309] The particle size of the insecticide and/or repellent in water-based suspensions or dispersions is in general between 50 nm to 20 μm, preferably 60 nm to 8 μm, more preferably 50 nm to 4 μm, most preferably 50 nm to 500 nm.

[0310] The aqueous formulation which is added in step ii) comprising at least one insecticide and/or repellent preferably comprises 0.1 to 45% by weight of water, preferably 1 to 25% by weight of water, based on the total of the components in the insecticide except of water.

[0311] Step i)

[0312] The aqueous bath is contained in a pressure-proof vessel. Suitable pressure-proof vessels are known by a person skilled in the art.

[0313] The liquor ratio, which is the weight-ratio between the amount of the textile material or plastics material and the amount of liquid of the aqueous bath, is preferably from 1:3 to 1:50, more preferably from 1:5 to 1:30, most preferably 1:20 (i.e. 1 kg textile material or plastics material in 20 liters of the aqueous bath).

[0314] The pH of the aqueous is preferably set slightly acidic, preferably from 3 to 6, more preferably from 4 to 5. Suitable additives for setting the pH to the desired value are known by a person skilled in the art.
Step ii) The addition of the aqueous formulation comprising at least one insecticide and/or repellent may be carried out by any method known by a person skilled in the art. Suitable aqueous formulations comprising at least one insecticide and/or repellent are mentioned before.

The aqueous bath is heated to a temperature of 100 to 140°C, and the temperature is kept for 20 to 120 minutes. Preferably the aqueous bath is heated to a temperature of 110 to 130°C, more preferably 120 to 130°C. The temperature of the aqueous bath is preferably kept for 20 to 90 minutes, more preferably for 30 to 60 minutes. The aqueous bath may be heated by any suitable means known by a person skilled in the art.

Step iv) The aqueous bath is cooled and drained. Further, the impregnated textile material or plastics material is rinsed and dried. Preferably, the aqueous bath is cooled to 90 to 50°C, more preferably to 80 to 60°C. The draining of the bath is carried out by any method known by a person skilled in the art. After the draining the impregnated textile material or plastics material is rinsed, preferably with warm and/or cold water. Finally the impregnated textile material or plastics material obtained is dried by any method known by a person skilled in the art.

The exhaust processor impregnation of a textile material or plastics material of the present invention is therefore preferably carried out as follows:

The textile material or plastics material is placed in an aqueous bath contained in a pressure-proof vessel in a liquor ratio of in general 1:3 to 1:50, preferably from 1:5 to 1:30, more preferably 1:20 (i.e. 1 kg netting in 20 liters). The pH is set slightly acidic (in general 3-6, preferably 4-5). The aqueous formulation of the insecticide and/or repellent is added preferably as a suspension or emulsion and the bath was heated to 100 to 140°C, preferably 100 to 130°C, more preferably 120 to 130°C. The temperature was kept for 20-120 minutes, preferably 20 to 90 minutes, more preferably 30 to 60 minutes. Then the treatment liquor is cooled to 90 to 50°C, more preferably to 80 to 60°C and subsequently the bath is drained. The treated material is rinsed with warm and cold water, respectively and is dried.

In a further embodiment the present invention relates to an impregnated textile material or plastics material obtainable by an exhaust process by comprising the steps i) to iv) of the impregnation process are described in detail above.

EXAMPLES

Material

Commercial netting material made from PET (polyethylene terephthalate) fibers, complying with WHO Roll Back Malaria “Specifications for netting materials” (Geneva, 2001) is employed. The netting material is treated with Alpha-Cypermethrin using different treatment processes, as indicated below. Dosages of Alpha-Cypermethrin are 25, 100, 200 mg/m², respectively. Samples A are unwashed, samples B are washed by the “Montpellier method”, samples C have been washed once, samples D have been washed 3 times, and samples E have been washed 5 times according to the procedure given below.

Binder

Preparation of Polymer Dispersions

General Procedure:

250 g of water and 3 g of a styrene seed (33% by weight) having a medium particle diameter of 30 nm are heated to 85°C and 5% by weight of feed 2 are added. After 10 min. addition of feed 1 comprising the monomers mentioned below and feed 2 is started.

Feed 2 comprises 3.0 g sodiumperoxodisulfate dissolved in 39.9 g of water.

The composition of feed 1 is listed in table 1.

Feed 1 and 2 are added in 3 h, and it was polymerized for further 0.5 h.

**TABLE 1**

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[0338] The amount of initiator sodium peroxodisulfate is 0.3 parts by weight, the emulsifier comprises 0.4 parts by weight of Dowfax 2A1 (Dow) and 0.6 parts by weight of Lumiten IRA (BASF AG), relating to 100 parts by weight of the monomer composition of table 1.

[0339] Abbreviations:
MMA: methyl methacrylate
S: styrene
AN: acrylic nitril
EA: ethyl acrylate
EHA: 2-ethylhexylacrylate
BA: n-butyl acrylate
Fl: copolymerizable benzophenone having an acrylic group
GMA: glycidylmethacrylate
BMA-Ac: bitandiolmonoacrylate acetylacetate
Amol: N-methyloxyl acrylamide
MAMol: N-methyloxyl methacrylamide
HPMA: hydroxypropyl methacrylate
AS: acrylic acid
AM: acrylic amide
Dowfax 2A1:

[0355] Lumiten IRA;

Examples A1 and A8

[0356] Aqueous polymer dispersions comprising Fl-1 polymerizable photo initiator which is later useful as crosslinking agent is a photo initiator of formula

\[
\text{O} \quad \begin{array}{c}
\text{C}_8 \text{H}_7 \text{O} \\
\text{SO}_3 \text{Na}
\end{array}
\]

[0357] wherein

[0358] \( R^0 \) is an organic radical having from 1 to 30 carbon atoms,

[0359] \( R^0 \) is H or a methyl group, and

[0360] \( R^{10} \) is a phenyl group which is optionally substituted or a \( C_1- \) to \( C_2- \) alkyl group.

[0361] Method

[0362] “Montpellier washing procedure” (as described in the annex WHO PVC, Mar. 7, 2002 “Evaluation of wash resistance of long-lasting insecticidal nets”): Net samples (B) are washed individually in beakers containing 0.5 L deionised water and 2 g/L soap (pH 10-11) at 30°C, in a water bath shaken for 10 minutes in clean water at 155 movements per minute.

[0363] Since there are no industry norms for washing procedures mimicking hand washes, we adapted a standard laundry process by using the IEC detergent test formulation (which contains anionic, non-ionic detergents, zeolithes, inorganic builder but omitting the bleaching system) for 30 min at 60°C, in metal vessels which are moved (tumbled) and heated. This method imparts more severe conditions than the hand wash relative to temperature, emulsifying and dispersing effects but exerts probably less friction on the nets.

[0364] Testing procedure: Two bioassays are used with 4 replicate tests done on each piece: the 3 min bioassay in which there is forced contact at laboratory-reared Anopheles mosquitoes with treated netting for 3 min and 24-h holding period before scoring mortality (as “% mortality”); The results are obtained using a plastic cone whose open basis
was covered with the different bed net samples, 11 mosquitoes are introduced. After 3 min contact time with the treated nets they are removed and their mortality rate and the median time to knockdown (MTKD) bioassay in which knockdown time is scored is determined after 24 h.

The MTKD is the time it takes for the middle mosquito (the 6th mosquito of the 11 exposed) to be knocked down.

Each sample is tested 4 times with 11 mosquitoes each. A higher mortality rate usually corresponds with a lower MTKD, indicating a faster action of the insecticide (or the formulation used).

The long-established standard for ITN testing is with the use of WHO cones (WHOPES 96.1): clear plastic cone structures (1.1 cm diameter) with a flat flange around the bottom edge and a hole at the apex. The general procedure is to secure the cone to the treated netting, flange side to the net, and introduce 5 mosquitoes into the cone and the hole plugged with cotton or a stopper. For inherent insecticidal activity, the insects are left in the cone resting on the netting for three minutes and then removed to a holding container supplied with sugar water. Mosquitoes from four cones are pooled to give 20 or more insects per holding container. Knockdown (KD) data can then be collected from the pooled mosquitoes at predetermined times out to 24-hours. For speed of knockdown, the mosquito can be left in the cone and the time to KD recorded for each individual mosquito until the 6th of 11 insects (median) goes down. Each KD mosquito is removed as it goes down to prevent recounting that insect if it once again flies. All mosquitoes are then held as described previously for a 24-hour KD count.

The following series have been completed:

1. An insecticide composition for application to a textile material or plastics material which composition comprises a mixture including:
   a) at least one insecticide and/or at least one repellent as component A, and
   b1) at least one acrylic binder as component B1 obtainable by emulsion polymerisation of the following components:
      b1a) n-butyl acrylate as component B1A,
      b1b) at least one monomer of formula I as component B1B

   wherein
   R₁, R₂ and R₃ are independently selected from C₁₋ to C₁₀₋-alkyl which may be linear or branched; substituted or unsubstituted aryl;
   R₁ and R₂ may further be H;
   except of R₃-n-butyl, when R₁ and R₂ are H;
   b1c) at least one monomer of formula II as component B1C

   wherein
   R₅, R₆, R₇ and R₈ are independently selected from the group consisting of H, C₁₋ to C₁₀₋-alkyl which may be linear or branched; substituted or unsubstituted aryl;

   Results

Bioassay results are given for the following compositions given in table 2 for the not washed nets:

<table>
<thead>
<tr>
<th>Composition</th>
<th>Insecticide concentration</th>
<th>Liquor up-take</th>
<th>Drying</th>
<th>Curing</th>
<th>KD rate after 60 min</th>
<th>Mortality after 24 hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>A 17</td>
<td>Alpha- cypermethrine 200 mg/mm² = 0.7% on weight of netting</td>
<td>100%</td>
<td>60 s at</td>
<td>2 minutes</td>
<td>100%</td>
<td>100%</td>
</tr>
<tr>
<td></td>
<td>Untreated netting</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
b1d) optionally at least one monomer of formula III as component B1D

![Formula III](image)

wherein

\[ R^8 \text{ and } R^9 \text{ are independently selected from the group consisting of } \text{H, } \text{C}_1 \text{ to } \text{C}_{10} \text{-alkyl which may be linear or branched, substituted or unsubstituted aryl; } \]

\[ X \text{ is selected from the group consisting of } \text{H, } \text{OH, } \text{NH, } \text{OR', OH, glycidyl, hydroxypropyl, } \]

b2c) optionally further components as component B2C; and

b2d) optionally further additives as component B2D.

2. The insecticide composition as claimed in claim 1, wherein the acrylic binder is obtainable by emulsion polymerization of the following components:

b1a) 10 to 90% by weight, preferably 15 to 80% by weight, more preferably 20 to 70% by weight of component B1A;

b1b) 10 to 90% by weight, preferably 12 to 85% by weight, more preferably 15 to 65% by weight of component B1B;

b1c) 1 to 5% by weight of component B1C;

b1d) 0 to 5% by weight, preferably 1 to 4% by weight more preferably 0.2 to 3% by weight of component B1D;

b1e) further monomers which are copolymerizable with the monomers mentioned above selected from

b1e1) 0 to 30% by weight, preferably 0 to 25% by weight, more preferably 5 to 20% by weight of component B1E1; and/or

b1e2) 0 to 40% by weight, preferably 0 to 30% by weight, more preferably 5 to 20% by weight of component B1E2;

wherein the sum of the components B1A, B1B, B1C and optionally B1D and B1E is 100% by weight.

3. The insecticide composition as claimed in claim 1, wherein the polyurethane is obtainable by reaction of the following components:

b2a) 55 to 99% by weight, preferably 70 to 98% by weight, more preferably 75 to 90 by weight based on the polyurethane of component B2A;

b2b) 10 to 90% by weight, preferably 12 to 85% by weight, more preferably 15 to 65% by weight based on the polyurethane of component B2B;

b2c) 0 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 1 to 5% by weight based on the polyurethane of component B2C; and

b2d) 0 to 10% by weight, preferably 0.1 to 5% by weight, more preferably 0.5 to 5% of component B2D;


4. The insecticide composition as claimed in claim 1, wherein the insecticide is selected from pyrethroid compounds such as

Ethofenprox: 2-(4-ethoxyphenyl)-2-methylpropyl-3-phenoxybenzyl ether,

Fenvalerate: (RS)-alpha-cyano-3-phenoxybenzyl (RS)-2-(4-chlorophenyl)-3-methylbutyrate,

Esfenvalerate: (S)-alpha-cyano-3-phenoxybenzyl (S)-2-(4-chlorophenyl)-3-methylbutyrate,

Fenpropathrin: (RS)-alpha-cyano-3-phenoxybenzyl 2,2,3, 3-tetramethylcyclopropane-carboxylate,
Cypermethrin: (RS)-alpha-cyano-3-phenoxybenzyl (1RS)-cis, trans-3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropanecarboxylate, alpha-Cypermethrin, Permethrin: 3-phenoxybenzyl (1RS)-cis, trans-3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropanecarboxylate, Cyhalothrin: (RS)-alpha-cyano-3-phenoxybenzyl (Z)-(1RS)-cis-3-(2-chloro-3,3,3-trifluoroprop-1-enyl)-2,2-dimethylcyclopropanecarboxylate, Deltamethrin: (S)-alpha-cyano-3-phenoxybenzyl (1R)-cis-3-(2,2-dibromovinyl)-2,2-dimethylcyclopropanecarboxylate, Cycloprothrin: (RS)-alpha-cyano-3-phenoxybenzyl (RS)-2,2-dichloro-1-(4-ethoxyphenyl)cyclopropanecarboxylate, Fluvalinate: alpha-cyano-3-phenoxybenzyl N-(2-chloroalpha, alpha, alpha, alpha-trifluoro-p-toly) D-valinate, Bifenthrin: 2-methyl biphenyl-3-ylmethyl(1R)-cis-3-(2-chloro-3,3,3-trifluoro-1-propenyl)-2,2-dimethylcyclopropanecarboxylate, 2-methyl-2-(4-bromodifluoromethoxyphenyl)propyl (3-phenoxybenzyl) ether, Tralomethrin: (S)-alpha-cyano-3-phenoxybenzyl (1R)-cis-(1'R,2'R,2'R,3'R,tetradibromoethyl)-2,2-dimethylcyclopropanecarboxylate, Silafluufen: 4-ethoxyphenyl(3-(4fluorophenyl)propyl)dimethylsiloxane, D-fenothrin: 3-phenoxybenzyl (1R)-cis, trans-2-chrysanthemate, Cyphenothrin: (RS)-alpha-cyano-3-phenoxybenzyl (1R)-cis, trans-chrysanthemate, D-resmethrin: 5-benzyl-3-furylmethyl (1R)-cis, trans-chrysanthemate, Acrinathrin: (S)-alpha-cyano-3-phenoxybenzyl (1R)-cis(Z)-2,2-dimethyl-3-(oxo-3(1,1,1,3,3,3-hexafluoropropyloxy)propenyl)cyclopropanecarboxylate, Cyfluthrin: (RS)-alpha-cyano-4-fluoro-3-phenoxybenzyl 3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropanecarboxylate, Tefluthrin: 2,3,5,6-tetrafluoro-4-methylbenzyl (1RS)-cis(Z)-3-(2-chloro-3,3,3-trifluoro-prop-1-enyl)-2,2-dimethylcyclopropanecarboxylate, Transfluthrin: 2,3,5,6-tetrafluorobenzyl (1R)-trans-3-(2,2-dichlorovinyl)-2,2-dimethylcyclopropanecarboxylate, Tetramethrin: 3,4,5,6-tetrahydrothaphthalimidomethyl (1RS)-cis, trans-chrysanthemate, Allethrin: (RS)-3-allyl-2-methyl-4-oxocyclopent-2-enyl (1RS)-cis, trans-chrysanthemate, Prallethrin: (S)-2-methyl-4-oxo-3-(2-propynyl)cyclopent-2-enyl (1R)-cis, trans-chrysanthemate, Empenthrin: (RS)-1-ethynyl-2-methyl-2-pentenyl (1R)-cis, trans-chrysanthemate, Imiprothrin: 2,5-dioxo-3-(prop-2-ynyl)imidazolidin-1-ylmethyl (1R)-cis, trans-2,2-dimethyl-3-(2-methyl-1-propenyl)-cyclopropanecarboxylate, D-lametrin: 5-(2-propynyl)-furfuryl (1R)-cis, trans-chrysanthemate, and 5-(2-propynyl)furfuryl 2,2,3,3-tetramethylcyclopropanecarboxylate;

Carbamate compounds such as Alanycarb: S-methyl-[N-methyl-N[2-ethoxy-carbonyl(amino)-thio]carbamoyl]thioacetimidate, Bendiocarb: 2,2-dimethyl-1,3-benzodioxol-4-yl-methyl carbamate, Carbaryl(1-naphthyl N-methylcarbamate, Isopropcarb: 2-(1-methylethyl)phenyl methylcarbamate, Carbosulfan: 2,3-dihydro-2,2-dimethyl-7-benzo furanyl [(dibutylamino)thio]methylcarbamate, Fenoxycarb: Ethyl[2-(4-phenoxyphenoxy)ethyl]carbamate, Indoxacarb: Methyl-7-chloro-22,3,4,5-tetrahydro-2-[methoxycarbonyl-(4-trifluoromethoxyphenyl)] Propoxur: 2-isopropoxyphenoxy methyl carbamate, Pirimicarb: 2-dimethylamino-5,6-dimethyl-4-pyrimidinyl-dimethyl carbamate, Thiodicarb: Dimethyl N,N-thiodiis(2-methyl-iminocarbonoyloxy) bisethanimidiothioate, Methomyl: S-methyl N-(methyl carbamoyloxy)thioacetamidate, Ethiofencarb: 2-(ethylthio)methylphenyl methyl carbamate, Fenothiocarb: S-(4-phenoxybutyl)-N,N-dimethyl thiocarbamate, Cartap: S,S'-2-(5 dimethylamino) trimethylene) bis (thiocarbamate) hydrochloride, Fenobucarb: 2-sec-butylyphenylmethyl carbamate, XMC: 3,5-dimethylphenyl methyl carbamate, Xylylcarb: 3,4-dimethylyphenylmethyl carbamate, organophosphorous compounds such as Fenitrothion: O,O-diethyl O-(4-nitro-m-tolyl)phosphorothioate, Diazinon: O,O-diethyl O-(2-isopropyl-6-methyl-4-pyrimidinyl) phosphorothioate, Pyridaphenthion: O-(1,6-dihydro-6-oxo-1-phenylpyrazidine-3-yl) O,O-diethyl phosphorothioate,
Chlorpyrifos: O,O-diethyl-O-(3,5,6-trichloro-2-pyridyl)phosphorothioate,
Chlorpyrifosmethyl: O,O-dimethyl, O-(3,5,6-trichloro-2-pyridinyl)phosphorothioate,
Cyano: O,O-dimethyl O-(4 cyanophenyl)phosphorothioate,
Pyraclofos: (R,S)-[4-cyano(phenyl)-pyrazolyl-4-yl]-ethyl-S-n-propyl phosphorothioate,
Acephate: O,S-dimethyl acetylphosphoroamidothioate,
Azamethiphos: S-(6-chloro-2,3-dihydro-oxo-1,3-oxazolo[4,5-b]pyridine-3methyl)phosphorothioate,
Malathion: O,O-dimethyl phosphoroathioate ester of diethyl mercaptosuccinate,
Temephos: O,O(thiodi-4-1-phenylene) O,O,O-tetramethyl phosphoroathioate,
Dimethoate: (O,O-dimethyl S-(n-methylcarbamoylthethyl)phosphorothioate,
Formothion: S[2-formylmethylamino]-2-oxoethyl]-O,O-dimethyl phosphorothioate,
Phenthato: O,O-dimethyl S(alpha ethoxycarbonylbenzal)-phosphorothioate;
Insecticides with a sterilising effect on adult mosquitoes such as
1-(alpha-(chloro-alpha-cyclopropylbenzylidenamino-oxy)p-tolyl)-3(2,6-difluorobenzoyl)urea,
Diflubenzuron: N-((3,5-dichloro-4-(1,1,2,2-tetrafluoroethoxy)phenylamino)carbonyl)2,6 difluoro benzamid,
Triflumuron: 5-Chloro-N-((4-(trifluoromethoxy)phenyl)amino)carbonyl)benzamide, or a triazin such as
N-cyclopropyl-1,3,5-triazine-2,4,6-triamin; and
cyano-3-phenoxophenylbenzyl-3-(2-chloro-3,3,3-trifluoroprop-1-enyl)-2,2-dimethylcyclopropane carboxylate, as a 1:1 mixture of (Z)-(1R,3R), R-ester and (Z)-(1S,3S), S-ester;
Lambda-cyhalothrine:
cyano-3-phenoxophenylbenzyl-3-(2chloro-3,3,3-trifluoroprop-1-enyl)-2,2-dimethylcyclopropane carboxylate, as a 1:1 mixture of (Z)-(1R,3R), R-ester and (Z)-(1S,3S), S-ester,
the repellent is selected from N,N-Diethyl-meta-toluamide (DEET), N,N-diethylphenylacetamide (DEPA),
1-(3-cyclohexan-1-yl-carbonyl)-2-methylpiperine,
(2-hydroxymethylcyclohexyl) acetic acid lactone, 2-ethyl-1,3-hexadiol, indalone, Methyleneeucamade (MND), a pyrethroid not used for insect control such as {1/(+/-)-3-allyl-2-methyl-4-oxocyclopent-2-/(+/-)enyl/(+/-)-trans-chrysantemate (Eshiothrin), a repellent derived from or identical with plant extracts like limonene, eugenol, (+)Eucamalol (1), (--)1-epi-eucamalol or crude plant extracts from plants like Eucalyptus maculata, Vetiver roundifolia, Cymbopogan martini, Cymbopogan citratus (lemon grass),Cymbopogan nardus (citronella).
5. The insecticide composition as claimed in any of claims 1 to 4, wherein the particle size of the insecticide and/or repellent is from 50 nm to 20 μm, preferably 50 nm to 8 μm, more preferably 50 nm to 4 μm, most preferably 50 nm to 500 nm.
6. The insecticide composition as claimed in claim 1, further comprising one or more component selected from water, preservatives, detergents, stabilizers, agents having UV-protecting properties, optical brighteners, spreading agents, anti-migrating agents, foam-forming agents, wetting agents, anti-soiling agents, thickeners, further biozides, plasticizers, adhesion agents, pigments and dyestuffs.
7. The insecticide composition as claimed in claim 1 comprising from about 0.001 to 95% by weight of the insecticide and/or repellent.
8. The insecticide composition as claimed in claim 1 which is provided as a kit for impregnation by the end-user.
9. The insecticide composition as claimed in claim 7 wherein the composition in the kit is adapted for preparing a solution or emulsion by adding water.
10. An impregnated textile material or plastics material for insect killing and/or repellence of an insect comprising
a) at least one insecticide and/or at least one repellent, and
b) at least one acrylic binder as claimed in claim 1; and/or
b) at least one polyurethane as claimed in claim 1.
11. The impregnated textile material or plastics material as claimed in claim 9 comprising an insecticide and/or repellent as defined in claim 4.
12. The impregnated textile material or plastics material as claimed in claim 9 further comprising one or more components selected from preservatives, detergents, stabilizers, agents having UV-protecting properties, optical brighteners, spreading agents, anti-migrating agents, foam-forming agents, wetting agents, anti-soiling agents, thickeners, further biozides, plasticizers, adhesives, pigments and dyestuffs.
13. The impregnated textile material or plastics material as claimed in claim 9 comprising from about 0.001 to 10% by weight of the textile material or plastics material of at least one insecticide and/or at least one repellent.
14. A process for impregnation of a textile material or plastics material comprising the steps
i) forming an aqueous formulation comprising at least one insecticide and/or at least one repellent and at least one acrylic binder and/or at least one polyurethane as defined in the present invention and optionally further ingredients;
ii) applying the aqueous formulation to the textile material or plastics material by
ia) passing the textile material or plastics material through the aqueous formulation; p2 or
ib) bringing the textile material or plastics material in contact with a roller that is partly or fully dipped into the aqueous formulation and drawing the aqueous formulation to the side of the textile material or plastics material in contact with the roller;
or
ii) double-side coating of the textile material or plastics material
or

ii) spraying the aqueous formulation onto the textile material or plastics material;

or

ii) applying the aqueous formulation in form of a foam;

or

iii) submerging the textile material or plastics material into the aqueous formulation;

or

iig) brushing the aqueous formulation onto or into the textile material or plastics material;

or

iib) pouring the aqueous formulation onto the textile material or plastics material;

iii) optionally removing surplus aqueous formulation; and

iv) drying and/or curing the textile material or plastics material.

15. The process as claimed in claim 13, wherein step iia) is carried out by completely submerging the textile material or plastics material in the aqueous formulation either in a trough containing the aqueous formulation or passing the textile material or plastics material through the aqueous formulation which is held between two horizontally oriented rollers.

16. The process as claimed in claim 13, wherein the insecticide and/or repellent is an insecticide and/or repellent as defined in claim 4.

17. The process as claimed in claim 13, wherein the aqueous formulation further comprises one or more ingredients selected from the group consisting of detergents, stabilizers, agents having UV-protecting properties, optical brighteners, spreading agents, anti-migrating agents, preservatives, foam-forming agents, wetting agents, thickeners, further biozides, plasticizers, adhesive agents, anti-soiling agents, pigments and dyestuffs.

18. The process as claimed in claim 13 wherein the impregnating composition is provided as a kit for impregnation by the end-user.

19. The process as claimed in claim 13, wherein the drying of the textile material or plastics material is carried out simultaneously with the impregnation of the textile material or plastics material, wherein an aqueous formulation is formed further comprising at least one dyestuff and/or at least one pigment.

20. A process for coating a textile material or plastics material by applying a composition comprising at least one insecticide and/or at least one repellent and at least one acrylic binder as defined in claim 1 and/or at least one polyurethane as defined in claim 1 to the textile material or plastics material.

21. The process as claimed in claim 19, wherein the composition further comprises one or more ingredients selected from the group consisting of detergents, stabilizers, agents having UV-protecting properties, optical brighteners, spreading agents, anti-migrating agents, preservatives, foam-forming agents, anti-soiling agents, wetting agents, thickeners, further biozides, plasticizers, adhesive agents, pigments and dyestuffs.

22. An exhaust process for impregnation of a textile material or plastics material comprising the steps

i) placing the textile material or plastics material in an aqueous bath optionally comprising further additives contained in a pressure-proof vessel;

ii) adding an aqueous formulation comprising at least one insecticide and/or repellent;

iii) heating the aqueous bath to a temperature of 100 to 140°C and keeping the temperature for 20 to 120 minutes;

and

iv) cooling and draining the bath, and rinsing and drying the impregnated textile material or plastics material.

23. The process as claimed in claim 21, wherein the aqueous bath comprises as further additives one or more components selected from preservatives, detergents, fillers, impact modifiers, anti-fogging agents, blowing agents, clarifiers, nucleating agents, coupling agents, conductivity-enhancing agents (antistats), stabilizers such as anti-oxidants, carbon and oxygen radical scavengers and peroxide decomposing agents and the like, flame retardants, mould release agents, agents having UV protecting properties, optical brighteners, spreading agents, anti-blocking agents, anti-migrating agents, foam-forming agents, anti-soiling agents, thickeners, further biozides, wetting agents, plasticizers and film forming agents, adhesive or anti-adhesive agents, optical brightening (fluorescent whitening) agents, pigments and dyestuffs.

24. The process as claimed in claim 21, wherein the aqueous formulation comprising at least one insecticide and/or repellent comprises 0.1 to 45% by weight of water, based on the total of the components in the insecticide except of water.

25. Impregnated textile material or plastics material obtainable by an exhaust process as claimed in claim 21.

* * * * *