(54) Title: COATING METAL PARTICLES

(57) Abstract: A method for preparing particles to retain a charge such that the particles are rendered electrostatically or electrophoretically mobile. The method involves coating the particles with a coating medium which facilitates attachment of a charge director material, and contacting the particles with the coating medium thereon with a charge director medium to impart a positive or negative charge thereto and thereby render the particles electrostatically or electrophoretically mobile. Electrostatically and electrophoretically mobile particles for use in an electrostatic or electrokinetic deposition process. The particles include a coating medium and a charge director on particle bodies.
COATING METAL PARTICLES

FIELD OF THE INVENTION
The present invention relates to a method for preparing particles including powders and larger particles for patterning by electrostatic or electrokinetic means, such as patterning of metal particles in the manufacture of electronic devices.

BACKGROUND OF THE INVENTION
In a variety of applications it is desirable to apply organic or inorganic particles, such as metal particles and metal powders, to substrates in a predetermined pattern. One such application is in the field of printed wiring boards, where solder paste containing metal powder has been applied to a substrate through a stainless steel or electroformed stencil which defines a pattern, or by screen printing.

Many industrial solder and other metal powders are not stable at room temperature for prolonged periods in that they are subject to oxidation.

SUMMARY OF THE INVENTION
Among the several aspects of the invention is to provide particles for electrokinetic or electrostatic deposition in a pattern on a substrate. Another aspect is to provide metal particles which are resistant to oxidation and stable over prolonged storage periods at room temperature.

Briefly, therefore, the invention is directed to a method for preparing particles to retain a charge such that the particles are rendered electrostatically or electrokinetically mobile. The method comprise coating
particles with a coating medium which facilitates attachment of a charge director material, wherein the coating medium is selected from the group consisting of anionic species, non-ionic species, cationic species, acids, bases, and mixtures thereof, to yield particles with the coating medium thereon; and contacting the particles with the coating medium thereon with a charge director medium to impart a positive or negative charge thereto and thereby render the particles electrostatically or electrokinetically mobile.

In another aspect the invention is directed to electrostatically or electrokinetically mobile particles for use in an electrostatic or electrokinetic deposition process. The particles comprise the particle bodies; a coating medium on the particle bodies which facilitates attachment thereto of a charge director material, wherein the coating medium is selected from the group consisting of anionic species, non-ionic species, cationic species, acids, bases, and mixtures thereof; and a charge director material on the coating medium wherein the charge director material has a positive or negative charge and thereby renders the particle bodies electrostatically or electrokinetically mobile.

Other aspects and features of the invention will be in part apparent, and in part described hereafter.

DETAILED DESCRIPTION OF THE INVENTION

In accordance with this invention, organic or inorganic particles are coated in order to render them amenable to treatment with a charge director, so that the particles can thereby be rendered electrokinetically or electrostatically mobile. In one preferred embodiment, the particles are a powder selected from among the
metals/alloys Sn, Sn/Pb, Sn/Ag/Cu, Sn/Ag, Cu, Sn/Ag/Bi, Sn/Bi, Sn/Pb/Bi or others.

The particle size is typically between about 0.5 microns and 1 mm. In one embodiment the particles are 5 powder particles having an average diameter between about 0.5 and about 50 microns. In another embodiment they are powder particles have an average diameter between about 1 and about 100 microns. One preferred powder is a Type 6 powder with a size distribution primarily between about 10 and about 20 microns. The particles are alternatively mid-sized particles having an average particle size between about 50 and about 500 microns. In a further alternative, they are relatively larger spheres of inorganic material having an average size greater than about 100 microns, such as between 100 microns and 1 mm. These are, for example, solder metal spheres. For certain applications, organometallic particles may be used.

In carrying out the invention, the particles are coated with a material selected from among ionic surfactant 20 materials, non-ionic materials such as non-ionic polymers, organic acids and/or bases, and mixtures thereof. Depending on how the coating is physically carried out, the coating medium may also contain a solvent. That is, if coating is accomplished by immersion in a solution, spraying with a 25 solution, or exposure to a solution in a fluidized bed, a solvent is needed. On the other hand, if the coating is accomplished by a plasma coating process or the like, no solvent is required.

Among the surfactant materials which are suitable for 30 coating particles according to this invention are the following cationic materials:

Stearamidopropyl diethanolamine available from Inolex Chemical Co. under the trade designation Lexamine 22.
C16-C18 monoalkyl amines such as hydrogenated tallowalkylamine available from Akzo Noel under the trade designation Armeen HT.

N-Myristoyl sarcosine available from Hampshire Chemical Corporation under the trade designation Hamposyl M.

N-Stearoyl sarcosine available from Hampshire Chemical Corporation under the trade designation Hamposyl S.

Mixture of N-Myristoyl and N-Stearoyl available from Hampshire Chemical Corporation under the trade designation Hamposyl MS.

N-(1,3-dimethylbutyl)-N-phenylenediamine available from Crompton Corporation under the trade designation Flexzone 7P (antioxidant 4020).

N, N'-hexane-1,6-diylbis(3-3(3,5-di-tert-butyl-4-hydroxyphenylproionamide)) available from Ciba under the trade designation Irganox 1098.

Amino-based silicone copolyol surfactant available from Lambent under the trade designation Lambent Amine PD.

Stearamidopropyl diethanolamine available from Inoless Chemical Co. under the trade designation Lexamine 22.

Amides, Polyamide and Polyamide resins for example Versamid 940 polyamide resin available from Cognis.

Also among the surfactant materials which are suitable for coating particles according to this invention are the following anionic materials:

N-Laurylhexamethylenediametriacetic acid available from Hampshire Chemical Corporation under the trade designation LED3A Acid.

Glyceryl Stearate SE containing potassium and free glycerol.
Behenic Acid (docosanoic acid) available from Croda under trade the designation Crodacid B.

Fatty alcohol/phosphate ester/ethoxylate mixture [alcohols C16-18 / 1-hexadecanol hydrogen phosphate / polyoxyethylene (1) cetyl ether phosphate] available from Croda under the trade designation Crodaphos CES.

Additional suitable materials include the following polymeric materials:

Styrene acrylic acid copolymer available under the designation Joncryl from S.C. Johnson Co.

Ethylene acrylic acid co-polymers A-C 540, 540A, 580, 5120, 5180 available from Allied Signal Corporation.

Palmitic acid, stearic acid and other fatty acids and mixture thereof.

Propylene acrylic acid copolymer.

Rosins and modified rosins for example Gum rosin, hydrogenated rosins, fumaric and maleic acid adducts of rosins etc.

The foregoing coating materials render the coated particles amenable to treatment with a charge director, so that the particles can be treated with a charge director to render them electrokinetically or electrostatically mobile. This coating provides a surface characterized as, for example, a very hydroxyl surface or a very acidic surface with which a subsequently applied charge director can react.

In selecting a coating material, if the particles are metal particles to be used as a solder, it is preferable to select one which provides a fluxing function in the eventual powder when it is used as a solder.
Among the organic acids which are suitable for coating particles according to this invention are aliphatic and aromatic acids, mono, di, tricarboxylic acids and also amino acids, sulfonic, sulfamic, phophinic, phosphonic acids etc., including the following: mono and dicarboxylic acids with a C backbone length between C₁ and C₂₀ such as adipic acid, citric acid, ascorbic acid, succinic acid, malic acid, malonic acid.

Among the organic bases which are suitable for coating particles according to this invention are the following: saturated and unsaturated, aliphatic and aromatic amines for example primary, secondary, tertiary amines, ethyl amine, tertiary butyl amine, propylamines, alkanolamines, alkylamines, Anisidine, N-phenylbenzylamine, N-Phenyldiethanolamine, Diphenylguanidine, 2-Aminobenzotrifluoride, etc.

The polar organic solvent, when used, is selected from among those solvents in which the coating compositions are soluble. These solvents include aromatic and aliphatic compositions; for example, alcohols such as methanol, ethanol, isopropanol, butanol, other alcohols, ketones, glycols, glycol ethers, esters, toluene, benzene, and mixtures thereof.

In preparing a surfactant solution, the surfactant or polymer content in one embodiment is between about 0.05 and about 10% by weight, preferably between about 0.1 and about 1% by weight. The organic solvent content is between about 99.95 to 90% by weight, preferably between about 99.9 and about 99% by weight.

In preparing an organic acid or base solution, the acid or base content in one embodiment is between about 0.05 and about 10% by weight, preferably between about 0.5 and about 2% by weight. The organic solvent content is
between about 99.95 to 90% by weight, preferably between about 99.5 and about 98% by weight.

The particles are introduced into the coating mixture in an amount of up to about 99.9 wt%. In a preferred embodiment, the particles are introduced in an amount between about 85% and about 99wt%. The method of introduction is selected from among ultrasonication, fluidized bed coating, low-shear impeller mixing, ball milling, and bead milling.

After coating the particles, the supernatant is decanted off and the particles are washed at least once. In one embodiment the particles are washed between two and five times with a neutral high dielectric organic liquid such as isoparaffin distributed by Exxon Chemical of Houston, Texas under the trade name Isopar. This yields prepared solder particles ready for use in patterning. Also, due to an oxygen diffusion barrier created by the coating, the particles resist oxidation and are stable for prolonged storage periods at room temperature.

As an alternative to immersion coating of the particles as described above, they may be coated by other methods such as fluidized bed coating, spray coating, pan coating, rotary coating, roll coating, plasma coating, or solution coating.

The foregoing process yields particles which have an electrically insulative coating and which are capable of holding a static charge. For solder powder, the coating is capable of melting away at soldering temperatures, and in some instances the coating serves a fluxing function during soldering.

As noted above, the coating medium may contain a mixture of two or more of the species mentioned. In one example, the primary coating medium is anionic or cationic,
with a counter species (cationic or anionic) used for controlling the overall charge.

One especially preferred embodiment within the scope of those described above employs one of the several ionic species (anionic or cationic) bound to a metal particle surface. The ionic species is selected to complement the charge director which is to then be applied for purposes of rendering the particles electrostatically or electrokinetically mobile.

To use the particles in an electrostatic or electrokinetic process such as that disclosed in the U.S. Patent Application filed July 9, 2004 entitled DEPOSITION AND PATTERNING PROCESS, the entire disclosure of which is expressly incorporated herein by reference, the particles are introduced into a solution containing an electrolyte or polyelectrolyte, also known as a charge director. Examples of suitable electrolytes/polyelectrolytes include, but are not limited to, the following:

Polyvinyl lecithin solution available from HP Compaq

Phosphated monoacid diglycerides available under the trade name NPHOS

Sodium dodecyl sulfate (SDS)

Sodium lauryl sulfate

Cobalt nitrate

Perfluoro pentanoic, octanoic, decanoic acids and similar species

Perfluoro carboxylic acids (perfluoro sebacic, suberic, succinic acids)

Sodium and calcium fatty acid salts

Sulfates of ethoxylated alcohols

Ammonium lauryl sulfate
Sodium allyl sulfates and sulfonates

Barium petronate, optionally modified by perfusion with distilled water

Quaternary ammonia electrolytes and polyelectrolytes such as allyl methyl ammonium chloride, hexadecyl ammonium chloride, and dodecyl trimethyl ammonium chloride.

This thereby yields electrostatically or electrokinetically mobile particles for use in an electrostatic or electrokinetic deposition process. The particles comprise the particle bodies; the above-described coating medium on the particle bodies which facilitates attachment thereto of a charge director material, wherein the coating medium is selected from the group consisting of anionic species, non-ionic species, cationic species, acids, bases, and mixtures thereof; and the above-described charge director material on the coating medium wherein the charge director material has a positive or negative charge and thereby renders the particle bodies electrostatically or electrokinetically mobile.

Thereafter, the particles are dried by pan drying, spray drying, or other suitable means. The function of the electrolyte or polyelectrolyte is to facilitate maintenance of electrical charge so the particles will respond in desired manner in the presence of an electrical field.

In preparation for printing, the particles are combined and wetted with a liquid having suitable dielectric properties. One example of such a liquid is the Isopar product. Another example is a product available from 3-M Corporation under the trade name Fluorinert. The combined particles and liquid ranges from about 2% by weight particles up to about 90% by weight particles. The particles constitute from about 2 to about 60% of the mixture on a volume basis.
When introducing elements of the present invention or
the preferred embodiment(s) thereof, the articles "a",
"an", "the" and "said" are intended to mean that there are
one or more of the elements. The terms "comprising",
"including" and "having" are intended to be inclusive and
mean that there may be additional elements other than the
listed elements.

As various changes could be made in the above without
departing from the scope of the invention, it is intended
that all matter contained in the above description and
shown in the accompanying drawings shall be interpreted as
illustrative and not in a limiting sense. The foregoing
relates to a limited number of embodiments that have been
provided for illustration purposes only. It is intended
that the scope of invention is defined by the appended
claims and there are modifications of the above embodiments
that do not depart from the scope of the invention.
WHAT IS CLAIMED IS:

1. A method for preparing particles to retain a charge such that the particles are rendered electrostatically or electrokinetically mobile, the method comprising:

coating the particles with a coating medium which facilitates attachment of a charge director material, wherein the coating medium is selected from the group consisting of anionic species, non-ionic species, cationic species, acids, bases, and mixtures thereof, to yield particles with the coating medium thereon; and

contacting the particles with the coating medium thereon with a charge director medium to impart a positive or negative charge thereto and thereby render the particles electrostatically or electrokinetically mobile.

2. The method of claim 1 wherein the particles are solder metal particles.

3. The method of claim 1 wherein:

the particles are solder metal particles; and

coating the particles with a coating medium comprises coating the particles with a cationic material.

4. The method of claim 1 wherein:

the particles are solder metal particles; and

coating the particles with a coating medium comprises coating the particles with a material selected from the group consisting of the following: a) stearamidopropyl diethanolamine, b) C16-C18 monoalkyl amines, c) hydrogenated tallowalkylamine, d) N-Myristoyl sarcosine, e)
N-Stearoyl sarcosine, f) N-(1,3-dimethylbutyl)-N-phenylenediamine, g) N, N'-hexane-1,6-diylbis(3-3(3,5-di-tert-butyl-4-hydroxyphenylproionamide)), h) amino-based silicone copolyol surfactants, and i) mixtures thereof.

5. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises coating the particles with a material selected from the 5 group consisting of amides, polyamides, and polyamide resins.

6. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises coating the particles with an anionic material.

7. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises coating the particles with a material selected from the 5 group consisting of a) N-Laurylethlenediametriacetic acid, b) Glyceryl Stearate SE containing potassium and free glycerol, c) Stearamidopropyl diethanolamine, d) Behenic Acid (docosanoic acid), e) a fatty alcohol/phosphate ester/ethoxylate mixture, and f) mixtures thereof.
8. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a polymeric material.

9. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of a) styrene acrylic acid copolymer, b)
ethylene acrylic acid co-polymers, c) palmitic acid, d)
stearic acid, e) propylene acrylic acid copolymer, and f)
mixtures thereof.

10. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of rosins and modified rosins.

11. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of gum rosin, hydrogenated rosins, fumaric
adducts of rosins, and maleic acid adducts of rosins.
12. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with an organic acid material.

13. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of a) aliphatic acids, b) aromatic acids,
c) monocarboxylic acids, d) dicarboxylic acids, e) tricarboxylic acids, f) amino acids, g) sulfonic acids, h)
sulfamic acids, i) phophinic acids, j) phosphonic acids,
and k) mixtures thereof.

14. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of mono and dicarboxylic acids with a C
backbone length between C$_1$ and C$_{20}$.

15. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of adipic acid, citric acid, ascorbic
acid, succinic acid, malic acid, malonic acid, and mixtures
thereof.
16. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with an organic base material.

17. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of aliphatic and aromatic amines.

18. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of a) primary amines, b) secondary amines,
c) tertiary amines, and d) mixtures thereof.

19. The method of claim 1 wherein:
the particles are solder metal particles; and
coating the particles with a coating medium comprises
coating the particles with a material selected from the
5 group consisting of a) ethyl amine, b) tertiary butyl
amine, c) propylamines, d) alkanolamines, e) alkylamines,
f) anisidine, g) N-phenylbenzylamine, h) N-i) phenyldiethanolamine, j) diphenylguanidine, k) 2-
Aminobenzotri fluoride, and l) mixtures thereof.
20. Electrostatically or electrokinetically mobile particles for use in an electrostatic or electrokinetic deposition process, the particles comprising:

   a) coating medium on the particle bodies which facilitates attachment thereto of a charge director material, wherein the coating medium is selected from the group consisting of anionic species, non-ionic species, cationic species, acids, bases, and mixtures thereof; and

   b) a charge director material on the coating medium wherein the charge director material has a positive or negative charge and thereby renders the particle bodies electrostatically or electrokinetically mobile.

21. The particles of claim 20 wherein the particle bodies are solder metal particle bodies.

22. The particles of claim 20 wherein the coating medium comprises a cationic material.

23. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of the following: a) stearamidopropyl diethanolamine, b) C16-C18 monoalkyl amines, c) hydrogenated tallowalkylamine, d) N-Myristoyl sarcosine, e) N-Stearoyl sarcosine, f) N-(1,3-dimethylbutyl)-N-phenylenediamine, g) N, N'-hexane-1,6-diylbis(3-3(3,5-0d-tet-butyl-4-hydroxphenylproionamide)), h) amino-based silicone copolyol surfactants, and i) mixtures thereof.
24. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of amides, polyamides, and polyamide resins.

25. The particles of claim 20 wherein the coating medium comprises an anionic material.

26. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of a) N-Laurylethlenediametriacetic acid, b) Glyceryl Stearate SE containing potassium and free 5 glycerol, c) Stearamidopropyl diethanolamine, d) Behenic Acid (docosanoic acid), e) a fatty alcohol/phosphate ester/ethoxylate mixture, and f) mixtures thereof.

27. The particles of claim 20 wherein the coating medium comprises a polymeric material.

28. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of a) styrene acrylic acid copolymer, b) ethylene acrylic acid co-polymers, c) palmitic acid, d) stearic acid, e) propylene acrylic acid copolymer, and f) mixtures thereof.

29. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of rosins and modified rosins.
30. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of gum rosin, hydrogenated rosins, fumaric adducts of rosins, and maleic acid adducts of rosins.

31. The particles of claim 20 wherein the coating medium comprises an organic acid material.

32. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of a) aliphatic acids, b) aromatic acids, c) monocarboxylic acids, d) dicarboxylic acids, e) tricarboxylic acids, f) amino acids, g) sulfonic acids, h) sulfamic acids, i) phophinic acids, j) phosphonic acids, and k) mixtures thereof.

33. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of mono and dicarboxylic acids with a C backbone length between C₁ and C₂₀.

34. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of adipic acid, citric acid, ascorbic acid, succinic acid, malic acid, malonic acid, and mixtures thereof.

35. The particles of claim 20 wherein the coating medium comprises an organic base material.
36. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of aliphatic and aromatic amines.

37. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of a) primary amines, b) secondary amines, c) tertiary amines, and d) mixtures thereof.

38. The particles of claim 20 wherein the coating medium comprises a material selected from the group consisting of a) ethyl amine, b) tertiary butyl amine, c) propylamines, d) alkanolamines, e) alkylamines, f) anisidine, g) N-phenylbenzylamine, h) N-i) phenyldiethanolamine, j) diphenylguanidine, k) 2-Aminobenzotrifluoride, and l) mixtures thereof.

39. The particles of each of claims 22 through 38 wherein the particle bodies comprise solder metal particle bodies.