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(71) Applicant(s)
Shengyi Technology Co., Ltd.

(72) Inventor(s)
He, Yueshan

(74) Agent / Attorney
Chrysiliou IP, 114-115/Biztek 20 Dale Street, Brookvale, NSW, 2100

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- (71) 申请人 (对除美国外的所有指定国): 广东生益科技股份有限公司 (GUANGDONG SHENGYI SCI. TECH CO., LTD) [CN/CN]; 中国广东省东莞市松山湖科技产业园区北部工业园工业西路 5 号, Guangdong 523000 (CN)。
- (72) 发明人; 及
- (75) 发明人/申请人 (仅对美国): 何岳山 (HE, Yueshan) [CN/CN]; 中国广东省东莞市松山湖科技产业园区北部工业园工业西路 5 号, Guangdong 523000 (CN)。
- (74) 代理人: 深圳市德力知识产权代理事务所 (COMIPS INTELLECTUAL PROPERTY OFFICE); 中国广东省深圳市深南中路新闻大厦 1 号楼 3 楼 307 室, Guangdong 518027 (CN)。
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(54) Title: HALOGEN-FREE RESIN COMPOSITION AND METHOD FOR PREPARATION OF COPPER CLAD LAMINATE WITH SAME

(54) 发明名称: 无卤树脂组合物以及使用其制作覆铜板的方法

(57) Abstract: A halogen-free resin composition and a method for preparation of copper clad laminate with the same, wherein based on total parts by weight of solid components, the halogen-free resin composition comprises a reactive allyl phenoxy cyclotriphosphazene or vinyl phenoxy cyclotriphosphazene of 5-50 parts, a thermosetting resin of 15-85 parts, a crosslinking curing agent of 1-35 parts, a crosslinking curing accelerant of 0-5 parts and a filler of 0-100 parts. In the present invention, the reactive allyl phenoxy cyclotriphosphazene or vinyl phenoxy cyclotriphosphazene having very low water absorption is introduced into the thermosetting resin, satisfying both the halogen-free and antilflaming requirements and improving the electrical properties of the system, and making it possible to prepare the halogen-free high-frequency high-speed substrate material. The resulted copper clad laminate satisfies the halogen-free requirement, and has advantages such as excellent resistance to heat and moisture, and low dielectric loss.

(57) 摘要: 一种无卤树脂组合物以及使用其制作覆铜板的方法, 以固体组分总重量份计算, 该无卤树脂组合物包括组分及其含量为: 反应型烯丙基苯氧基环三磷腈或乙烯基苯氧基环三磷腈 5-50 份、热固性树脂 15-85 份、交联固化剂 1-35 份、交联固化促进剂 0-5 份、及填料 0-100 份。本发明通过在热固性树脂中引入吸水率极低的反应型烯丙基苯氧基环三磷腈或乙烯基苯氧基环三磷腈, 既能满足无卤阻燃要求, 又能改良体系电性能, 使得高频高速基板材料无卤化成为可能, 所制得的覆铜板满足无卤要求, 具有优异的耐热性和耐湿性, 介电损耗低等优点。

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HALOGEN-FREE RESIN COMPOSITION AND METHOD FOR PREPARATION OF COPPER CLAD LAMINATE WITH SAME

Technical field

- 5 The present invention relates to a resin composition, especially a halogen-free resin composition and method for preparation of copper clad laminate with same.

Background art

- 10 Recently, electronic devices are continuously developed with the development of integrated technology, bonding technology and mounting technology of semiconductor devices used in wiring of electronic devices, high-density electronic packages and high-density printed circuit board. Especially, a rapid development thereof occurs in electronic devices using
15 broadband, *e.g.* mobile communication devices.

- As a component of such electronic devices, printed circuit board develops in a direction of a higher degree of multilayer printed circuit boards and more precise wiring. In order to increase the signal transmission speed to a
20 level desired for speeding up the information processing, an effective way is to reduce the dielectric constant of the used materials. In order to reduce the transmission loss, an effective way is to use the materials having a lower dielectric loss tangent (dielectric loss).

- 25 While the electronic technology rapidly develops, people are pursuing environmental protection. However, the conventional high-frequency high speed materials basically use halides, antimonides and the like to achieve the object of flame retardance. When the halide-containing copper foil laminate is on fire and combusts, it will emit a great amount of smoke and
30 an undesirable odor, as well hydrogen halide gas having a great toxicity and a strong causticity, which not only pollute the environment, but also endanger human health. Currently, the epoxy resins corresponding to the phosphorous-containing phenanthrene type compound DOPO or ODOPB are generally and industrially used to achieve the common FR-4 and the
35 flame retardance. The phosphorous-containing phenanthrene type

compound DOPO or ODOPB, however, still has a greater water absorption and a great effect on the dielectric constant and dielectric loss tangent.

Contents of the invention

- 5 The object of the present invention lies in providing a halogen-free resin composition,

Another object of the present invention lies in providing a method for preparing copper clad laminate by using the halogen-free resin
10 composition, said method being simple, wherein the produced copper clad laminate has the advantages of halogen-free, excellent heat resistance and moisture resistance and low dielectric loss.

In order to achieve said objects, the present invention provides a
15 halogen-free resin composition, based on the total weight parts of the solid components, comprising

from 5 to 50 parts by weight of reactive allylphenoxy-
cyclotriphosphazene or vinylcyclotriphosphazene;

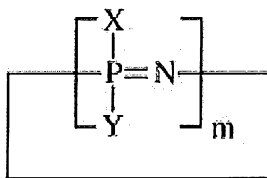
from 15 to 85 parts by weight of a thermosetting resin;

20 from 1 to 35 parts by weight of a crosslinking curing agent;

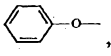
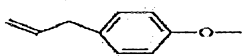

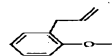
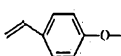

from 0 to 5 parts by weight of a crosslinking curing accelerator; and

from 0 to 100 parts by weight of a packing.

Said reactive allylphenoxy cyclotriphosphazene or vinylcyclo-
25 triphosphazene has the following structural formula (I)



30

wherein m represents an integer of 3-25; X is phenoxy , in an amount of 0-2m-1; Y is selected from the group consisting of p-allylphenoxy , m-allylphenoxy , o-allyl-phenoxy , p-vinylphenoxy  and m-vinylphenoxy , in an amount of 1-2m; X+Y=2m.

35

Said thermosetting resin comprises one or more of the following resins

epoxy resins, including bisphenol A epoxy resin, bisphenol F epoxy resin, dicyclopentadiene epoxy resin, triphenolic epoxy resin,
5 biphenyl-type epoxy resin, naphthol epoxy resin, and phosphorous-containing epoxy resin;

benzoxazine resin, including bisphenol A-type benzoxazine resin, bisphenol F-type benzoxazine resin, dicyclopentadiene-type benzoxazine resin and phenolphthalein-type benzoxazine resin;

10 cyanate resin, including bisphenol A-type cyanate resin, dicyclopentadiene-type cyanate resin and phenolic aldehyde-type cyanate resin;

bismaleimide resin, including 4,4'-diphenylmethane bismaleimide resin, allyl-modified diphenylmethane bismaleimide resin;

15 reactive polyphenylene oxide resin, having a number average molecular weight of 1000-7000, wherein the reactive groups is hydroxyl or double bond; and

C5 hydrocarbon resin consisting of vinylbutylbenzene resin having a molecular weight of 11000, vinylpolybutadiene resin with polar groups
20 and a copolymer in which butadiene is grafted to styrene with maleic anhydride.

Said crosslinking curing agent is selected from the group consisting of dicyandiamide, aromatic amine, anhydride, phenols, fulminuric triene ester
25 and phosphorous-containing phenolic aldehyde, or combinations thereof.

Said crosslinking curing accelerator is selected from the group consisting of 2-methylimidazole, 2-ethyl-4-methylimidazole, 2-phenylimidazole, tri-(dimethylaminomethyl), hexamethylene tetramine, dicumyl peroxide,
30 tert-butyl peroxybenzoate, 2,5-di(2-ethylcaproyl peroxy)-2,5-dimethylhexane, acetopyruvic salts, and caprilate, or combinations thereof.

Said packing is one or more selected from the group consisting of aluminium hydroxide, magnesium hydroxide, zeolite, wollastonite, silicon
35 dioxide, magnesia, calcium silicate, calcium carbonate, clay, talc and mica.

Further, the present invention provides a method for preparing a copper clad laminate with the halogen-free resin composition above, comprising

5 Step 1: dissolving from 5 to 50 parts by weight of reactive allylphenoxycyclotriphosphazene or vinylphenoxycyclotriphosphazene in a solvent until complete dissolution at room temperature or moderate temperature;

10 Step 2: adding from 15 to 85 parts by weight of the corresponding thermosetting resin, from 1 to 35 parts by weight of a crosslinking curing agent, from 0 to 5 parts by weight of a crosslinking curing accelerator and from 0 to 100 parts by weight of a packing into said solution, homogeneously stirring to obtain a liquid cement;

Step 3: homogeneously coating with said liquid cement E-glass cloth having a flattening surface, and then baking to a B-stage prepreg;

15 Step 4: cutting the B-stage prepreg to a suitable size according to the press size, stacking several prepregs regularly, a copper foil being placed over and under the prepregs, pressing in a vacuum hot press to obtain a copper clad laminate.

20 The solvent is one or more selected from benzene and ketone solvents; said moderate temperature is $\leq 80^{\circ}\text{C}$.

During the step 3 of baking the B-stage prepreg, the temperature is set up according to the boiling point of the solvent used therein, and ranges from 25 85 to 175°C , and the baking lasts 5-20 min.

In step 4, the pressing process is carried out by the bench pressing method, comprising the specific steps of increasing from room temperature to 150°C within 15 min and remaining for 30 min, then increasing to 180°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h and post-processing at $200\text{-}245^{\circ}\text{C}$ and remaining for 1-5 h.

The present invention has the following beneficial effects. The reactive allyl phenoxy cyclotriphosphazene or vinyl phenoxy cyclotriphosphazene having very low water absorption is introduced into the thermosetting resin, satisfying both the halogen-free and antifiame requirements and improving the electrical properties of the system (decreasing and stabilizing Dk (dielectric constant) and Df (dielectric loss factor)), and making it possible to prepare the halogen-free high-frequency high-speed substrate material. The resulted copper clad laminate satisfies the halogen-free requirement, and has advantages such as excellent resistance to heat and moisture, and low dielectric loss.

Embodiments

The halogen-free resin composition of the present invention comprises, based on the total weight parts of the solid components, comprising

from 5 to 50 parts by weight of reactive allylphenoxy-cyclotriphosphazene or vinylcyclotriphosphazene;

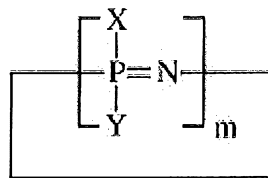
from 15 to 85 parts by weight of a thermosetting resin;

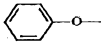

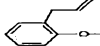
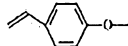
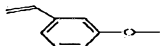
from 1 to 35 parts by weight of a crosslinking curing agent;

from 0 to 5 parts by weight of a crosslinking curing accelerator; and

from 0 to 100 parts by weight of a packing.

Said reactive allylphenoxy cyclotriphosphazene or vinylcyclotriphosphazene has the following structural formula (I)



wherein m represents an integer of 3-25; X is phenoxy in an amount of 0-2m-1; Y is selected from the group , , , , and , in an amount of 1-2m; X+Y=2m.

Said reactive allylphenoxy cyclotriphosphazene or vinylcyclo-

triphosphazene is a phosphazene compound having very low water absorption. By reacting the reactive groups therein with the specific thermosetting resin, the present invention may satisfy the halogen-free and antifi
5 any effect on the excellent dielectric constant and dielectric loss tangent performance.

Said thermosetting resin comprises one or more of the following resins

10 epoxy resins, including bisphenol A epoxy resin, bisphenol F epoxy resin, dicyclopentadiene (DCPD)epoxy resin, triphenolic epoxy resin, biphenyl-type epoxy resin, naphthol epoxy resin, and phosphorous-containing epoxy resin; benzoxazine resin, including bisphenol A-type benzoxazine resin, bisphenol F-type benzoxazine resin, dicyclopentadiene-type benzoxazine resin and phenolphthalein-type
15 benzoxazine resin; cyanate resin, including bisphenol A-type cyanate resin, dicyclopentadiene-type cyanate resin and phenolic aldehyde-type cyanate resin; bismaleimide resin, including 4,4'-diphenylmethane bismaleimide resin, allyl-modified diphenylmethane bismaleimide resin; reactive polyphenylene oxide resin, having a number average molecular weight of
20 1000-7000, wherein the reactive groups is hydroxyl or double bond; and C5 hydrocarbon resin consisting of vinylbutylbenzene resin having a molecular weight of 11000, vinylpolybutadiene resin with polar groups and a copolymer in which butadiene is grafted to styrene with maleic anhydride.

25 Said crosslinking curing agent is selected from the group consisting of dicyandiamide, aromatic amine, anhydride, phenols, fulminuric triene ester and phosphorous-containing phenolic aldehyde, or combinations thereof.

30 Said crosslinking curing accelerator is selected from the group consisting of 2-methylimidazole, 2-ethyl-4-methylimidazole, 2-phenylimidazole, DMP-30 (tri-(dimethylaminomethyl)), hexamethylene tetramine, dicumyl peroxide, tert-butyl peroxybenzoate, 2,5-di(2-ethylcaproyl peroxy)-2,5-dimethylhexane, acetoacetic salts, and caprilate, or combinations thereof.

35 Said packing is one or more selected from the group consisting of

aluminium hydroxide, magnesium hydroxide, zeolite, wollastonite, silicon dioxide, magnesia, calcium silicate, calcium carbonate, clay, talc and mica. The amount of said inorganic packing may be adjusted according to the use purpose, and may be better from 0 to 100 parts by weight, most
5 preferably from 25 to 100 parts by weight, based on the total amount of 100 parts by weight of the organic solids of the components in said halogen-free resin composition.

The aforesaid halogen-free resin composition is used to prepare prepregs, copper clad laminates and the like, wherein the fabricated prepregs include
10 the substrate material and the halogen-free resin composition coated thereon; the substrate material is glass cloth, preferably E-glass cloth; the fabricated copper clad laminates include several stacked prepregs, and copper foils on one or both sides of the stacked prepregs. The resulted
15 copper clad laminate satisfies the halogen-free requirement, and has advantages such as excellent resistance to heat and moisture, and low dielectric loss.

The method for preparing a copper clad laminate with the halogen-free resin composition above comprises the following steps:

20 Step 1: dissolving from 5 to 50 parts by weight of reactive allylphenoxycyclotriphosphazene or vinylphenoxycyclotriphosphazene in a phenol or ketone solvent until complete dissolution at normal temperature or moderate temperature, wherein the solvent may be one or more phenol or ketone solvents; said normal temperature is the normal
25 temperature in the prior art; the moderate temperature represents the temperature range of $\leq 80^{\circ}\text{C}$;

Step 2: adding from 15 to 85 parts by weight of the corresponding thermosetting resin, from 1 to 35 parts by weight of a crosslinking curing agent, from 0 to 5 parts by weight of a crosslinking curing accelerator and
30 from 0 to 100 parts by weight of a packing into said solution, homogeneously stirring to obtain a liquid cement;

Step 3: homogeneously coating with said liquid cement E-glass cloth having a flattening surface, and then baking to a B-stage prepreg, wherein during the process of baking the B-stage prepreg, the temperature is set up
35 according to the boiling point of the solvent used therein, and ranges from

85 to 175°C, and the baking lasts 5-20 min;

Step 4: cutting the B-stage prepreg to a suitable size according to the press size, stacking several prepreps regularly, a copper foil being placed over and under the prepreps, pressing in a vacuum hot press to obtain a copper clad laminate. In such step, the pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising the specific steps of increasing from room temperature to 150°C within 15 min and remaining for 30 min, then increasing to 180°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h and post-processing at 200-245°C and remaining for 1-5 h.

As for the fabricated copper clad laminate above, the dielectric constant, glass transition temperature, peeling strength, combustibility resistance, hygroscopicity and punching property thereof are measured, and further detailedly explained and described by the following examples.

The present invention is detailedly explained with the examples as follows, but not limited within the scope of the examples.

Example 1

A solvent, hexachlorocyclotriphosphazene, allylphenol (containing o-allylphenol as the primary ingredient), and an acid-binding agent were added to a reaction device, stirred, protected by feeding nitrogen gas, and heated to 50-120 °C. A catalyst was added to the reaction device to react for 5-16 h, then cool to room temperature and leach. The filtrate was distilled under pressure to evaporate the solvent to obtain a brown powder or viscous product. After 30 g of said product was dissolved in an organic solvent, 50 g of DCPD epoxy resin (which was HP-7200H (DIC), and had an equivalent of 275-280), 20 g of DCPD phenolic resin (which was Nihon Petro-chemical DPP-600 M), a suitable amount of imidazole and 2,5-(di(2-ethylcaproyl peroxy)) were added, homogeneously stirred to obtain a liquid cement. A flattening E-glass cloth having an area of 300x300 cm and a smooth surface was coated with said liquid cement, baked in

an oven at 155°C for 7 min to obtain a prepreg. 5 prepregs the rough edges of which were cut off were stacked, and copper foils having a thickness of 35 µm were attached on and under the prepregs and pressed in a vacuum hot press to obtain a copper clad laminate. The pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising increasing from room temperature to 150 °C within 15 min and remaining for 30 min, then increasing to 180°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h. The basic performances of the produced copper clad laminate are stated in Table 1 blow.

Example 2

A solvent, hexachlorocyclotriphosphazene, allylphenol (containing o-allylphenol as the primary ingredient), and an acid-binding agent were added to a reaction device, stirred, protected by feeding nitrogen gas, and heated to 50-120 °C. A catalyst was added to the reaction device to react for 5-16 h, then cool to room temperature and leach. The filtrate was distilled under pressure to evaporate the solvent to obtain a brown powder or viscous product. After 30 g of said product was dissolved in an organic solvent, 40 g of DCPD benzoxazine (which was MT36000 (Huntsman)), 20 g of DCPD epoxy resin (which was HP-7200H (DIC) and had an equivalent of 275-280), 10 g of styrene/maleic anhydride (said anhydride being EP-30, Sartomer), a suitable amount of imidazole and 2,5-(di(2-ethylcaproyl peroxy)) were added, homogeneously stirred to obtain a liquid cement. A flattening E-glass cloth having an area of 300x300 cm and a smooth surface was coated with said liquid cement, baked in an oven at 155°C for 7 min to obtain a prepreg. 5 prepregs the rough edges of which were cut off were stacked, and copper foils having a thickness of 35 µm were attached on and under the prepregs and pressed in a vacuum hot press to obtain a copper clad laminate. The pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising increasing from room temperature to 150 °C within 15 min and remaining for 30 min, then increasing to 190°C within 5 min and remaining for 2 h, finally decreasing to room temperature

within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h. The basic performances of the produced copper clad laminate are stated in Table 1 blow.

5

Example 3

A solvent, hexachlorocyclotriphosphazene, allylphenol (containing o-allylphenol as the primary ingredient), and an acid-binding agent were added to a reaction device, stirred, protected by feeding nitrogen gas, and heated to 50-120
10 °C. A catalyst was added to the reaction device to react for 5-16 h, then cool to room temperature and leach. The filtrate was distilled under pressure to evaporate the solvent to obtain a brown powder or viscous product. After 30 g of said product was dissolved in an organic solvent, 30 g of DPCP cyanate (which was LONZA-Primaset BADCy), 20 g of 4,4'-diphenylmethane bismaleimide resin,
15 20 g of DCPD epoxy resin (which was HP-7200H (DIC) and had an equivalent of 275-280), and a suitable amount of aluminum acetopyruvate were added, homogeneously stirred to obtain a liquid cement. A flattening E-glass cloth having an area of 300x300 cm and a smooth surface was coated with said liquid cement, baked in an oven at 155°C for 7 min to obtain a prepreg. 5 prepregs the
20 rough edges of which were cut off were stacked, and copper foils having a thickness of 35 µm were attached on and under the prepregs and pressed in a vacuum hot press to obtain a copper clad laminate. The pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising increasing
25 from room temperature to 150°C within 15 min and remaining for 30 min, then increasing to 210°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h. The basic
30 performances of the produced copper clad laminate are stated in Table 1 blow.

Example 4

A solvent, hexachlorocyclotriphosphazene, allylphenol (containing
35 o-allylphenol as the primary ingredient), and an acid-binding agent were added to

a reaction device, stirred, protected by feeding nitrogen gas, and heated to 50-120 °C. A catalyst was added to the reaction device to react for 5-16 h, then cool to room temperature and leach. The filtrate was distilled under pressure to evaporate the solvent to obtain a brown powder or viscous product. After 30 g of said product was dissolved in an organic solvent, 50 g of reactive polyphenylene oxide resin (which was MX9000, SABIC), 20 g of DCPD epoxy resin (which was HP-7200H (DIC) and had an equivalent of 275-280), a suitable amount of imidazole and 2,5-(di(2-ethylcaproyl peroxy)) were added, homogeneously stirred to obtain a liquid cement. A flattening E-glass cloth having an area of 300x300 cm and a smooth surface was coated with said liquid cement, baked in an oven at 155°C for 7 min to obtain a prepreg. 5 prepregs the rough edges of which were cut off were stacked, and copper foils having a thickness of 35 µm were attached on and under the prepregs and pressed in a vacuum hot press to obtain a copper clad laminate. The pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising increasing from room temperature to 150 °C within 15 min and remaining for 30 min, then increasing to 190°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h. The basic performances of the produced copper clad laminate are stated in Table 1 blow.

Comparison Example 1

A solvent, hexachlorocyclotriphosphazene, vinylphenol (a 50/50 mixture of p-vinylphenol and m-vinylphenol) and an acid-binding agent were added to a reaction device, stirred, protected by feeding nitrogen gas, and heated to 50-120 °C. A catalyst was added to the reaction device to react for 5-16 h, then cool to room temperature and leach. The filtrate was distilled under pressure to evaporate the solvent to obtain a brown powder or viscous product. After 30 g of said product was dissolved in an organic solvent, 30 g of DPCP cyanate (which was LONZA-Primaset BADCy), 20 g of 4,4'-diphenylmethane bismaleimide resin, 20 g of DCPD epoxy resin (which was HP-7200H (DIC)) were added. The prepregs the rough edges of which were cut off were stacked, and copper foils having a thickness of 35 µm were attached on and under the prepregs and pressed

in a vacuum hot press to obtain a copper clad laminate. The pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising increasing from room temperature to 150°C within 15 min and remaining for 30 min, then increasing to 190°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h. The basic performances of the produced copper clad laminate are stated in Table 1 blow.

Comparison Example 2

A solvent, hexachlorocyclotriphosphazene, vinylphenol (a 50/50 mixture of p-vinylphenol and m-vinylphenol) and an acid-binding agent were added to a reaction device, stirred, protected by feeding nitrogen gas, and heated to 50-120 °C. A catalyst was added to the reaction device to react for 5-16 h, then cool to room temperature and leach. The filtrate was distilled under pressure to evaporate the solvent to obtain a brown powder or viscous product. After 30 g of said product was dissolved in an organic solvent, 50 g of reactive polyphenylene oxide resin (which was MX9000, SABIC), 20 g of DCPD epoxy resin (which was HP-7200H (DIC) and had an equivalent of 275-280), a suitable amount of imidazole and 2,5-(di(2-ethylcaproyl peroxy)) were added, homogeneously stirred to obtain a liquid cement. A flattening E-glass cloth having an area of 300x300 cm and a smooth surface was coated with said liquid cement, baked in an oven at 155°C for 7 min to obtain a prepreg. 5 prepregs the rough edges of which were cut off were stacked, and copper foils having a thickness of 35 µm were attached on and under the prepregs and pressed in a vacuum hot press to obtain a copper clad laminate. The pressing process is carried out by the bench pressing method (step-by-step temperature-increasing and pressure-increasing), comprising increasing from room temperature to 150 °C within 15 min and remaining for 30 min, then increasing to 190°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h. The basic performances of the produced copper

clad laminate are stated in Table 1 blow.

Table 1 Property Evaluation

	Example 1	Example 2	Example 3	Example 4	Com.Exp. 1	Com.Exp .2
Glass transition temperature Tg °C, DMA)	160-170	170-185	195-225	195-215	200-230	190-220
Peeling strength (1/2OZ, N/mm)	>1.4	>1.4	>1.0	>1.0	>1.0	>1.0
Combustibility (1.60mm)	V-1	V-0	V-0	V-0	V-0	V-0
Dip-soldering resistance (stratification)	O	Δ	O	O	O	O
Hygroscopicity	0.11	0.10	0.13	0.11	0.12	0.10
Dielectric constant (RC50, 1GHz)	4.0	4.2	3.8	3.6	3.7	3.5
Dielectric loss (RC50, 1GHz)	0.010	0.009	0.005	0.003	0.005	0.003
T-300/min	>60	>60	>120	>120	>120	>120
Punching property	O	Δ	O	O	Δ	O

5 The aforesaid properties are measured by the following methods.

(1) Glass transition temperature (Tg): measured according to the differential scanning calorimetry (DSC) as stipulated under IPC-TM-650 2.4.25.

10 (2) Combustibility resistance: measured according to UL 94 vertical firing method.

(3) Dip-soldering resistance: the sample (the substrate material of 100x100 mm) which remains in pressure processing device at 121°C and 105 Kpa
15 for 2 hours is impregnated in a solder bath heated to 260°C for 20 seconds to visually observe (h1) whether there is stratification; (h2) whether there

is white plague or wrinkling, wherein the symbol O in the table represents unchanged; the symbol Δ represents white plague; the symbol x represents stratification.

5 (4) Hygroscopicity: measured according to the method of IPC-TM-650 2.6.2.1.

(5) Dielectric Loss Factor: measured according to the resonance method using strip lines and IPC-TM-650 2.6.2.1.

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(6) Punching property: the basis material having a thickness of 1.60 mm is placed on a punch die having a certain figure for punching to visually observe (h1) no bright border at the hole edge; (h2) bright border at the hole edge; (h3) cleavage of the hole edge, which are respectively
15 represented with O, Δ and x.

According to the results above, it can be seen that the present invention can satisfy both the halogen-free and antifiaming requirements (halogen content falling within the scope required by the JPCA halogen-free requirements) without
20 not reducing the dielectric performance, excellent resistance to heat, and better processability.

The aforesaid examples are not used to limit the content of the composition of the present invention. Any tiny amendment, equivalent
25 change or modification to the aforesaid examples according to the technical essence of the present invention and the weight part or content of the composition still falls within the scope of the technical solution of the present invention.

Claims

1. A halogen-free resin composition for preparing a copper clad laminate, comprising, based on the total weight parts of the solid components,

from 5 to 50 parts by weight of reactive allylphenoxycyclotriphosphazene or vinylcyclotriphosphazene;

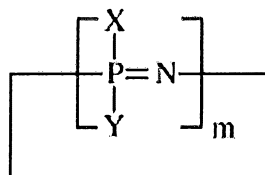
from 15 to 85 parts by weight of an epoxy resin and a thermosetting resin;

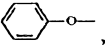
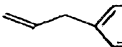
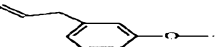
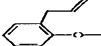
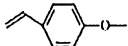
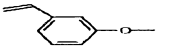
from 1 to 35 parts by weight of a crosslinking curing agent;

from 0 to 5 parts by weight of a crosslinking curing accelerator;
and

from 0 to 100 parts by weight of a packing;

wherein said reactive allylphenoxycyclotriphosphazene or vinylcyclotriphosphazene has the following structural formula (I)



wherein m represents an integer of 3-25; X is phenoxy , an amount of 0-2m-1; Y is selected from the group consisting of p-allylphenoxy , m-allylphenoxy , o-allyl-phenoxy , p-vinylphen  and m-vinylphenoxy , in an amount of 1-2m; X+Y=2m;

wherein said epoxy resin is selected from the group consisting of bisphenol A epoxy resin, bisphenol F epoxy resin, dicyclopentadiene epoxy resin, triphenolic epoxy resin, biphenyl-type epoxy resin, naphthol epoxy resin, and phosphorous-containing epoxy resin; and

wherein the thermosetting resin is selected from the group consisting of the following resins:

benzoxazine resin, including bisphenol A-type benzoxazine

resin, bisphenol F-type benzoxazine resin, dicyclopentadiene-type benzoxazine resin and phenolphthalein-type benzoxazine resin;

cyanate resin, including bisphenol A-type cyanate resin, dicyclopentadiene-type cyanate resin and phenolic aldehyde-type cyanate resin;

bismaleimide resin, including 4,4'-diphenylmethane bismaleimide resin, allyl-modified diphenylmethane bismaleimide resin;

reactive polyphenylene oxide resin, having a number average molecular weight of 1000-7000, wherein the reactive groups is hydroxyl or double bond; and

C5 hydrocarbon resin consisting of vinylbutylbenzene resin having a molecular weight of 11000, vinylpolybutadiene resin with polar groups and a copolymer in which butadiene is grafted to styrene with maleic anhydride.

2. The halogen-free resin composition for preparing a copper clad laminate according to claim 1, wherein said crosslinking curing agent is selected from the group consisting of dicyandiamide, aromatic amine, anhydride, phenols, fulminuric triene ester and phosphorous-containing phenolic aldehyde, or combinations thereof.

3. The halogen-free resin composition for preparing a copper clad laminate according to claim 1, wherein said crosslinking curing accelerator is selected from the group consisting of 2-methylimidazole, 2-ethyl-4-methylimidazole, 2-phenylimidazole, tri-(dimethylaminomethyl), hexamethylene tetramine, dicumyl peroxide, tert-butyl peroxybenzoate, 2,5-di(2-ethylcaproyl peroxy)-2,5- dimethylhexane, acetopyruvic salts, and caprilate, or combinations thereof.

4. The halogen-free resin composition for preparing a copper clad laminate according to claim 1, wherein said packing is one or more selected from the group consisting of aluminium hydroxide, magnesium hydroxide, zeolite, wollastonite, silicon dioxide, magnesia, calcium silicate, calcium carbonate, clay, talc and mica.

5. A method for preparing a copper clad laminate with the halogen-free resin composition for preparing a copper clad laminate according to claim 1, comprising

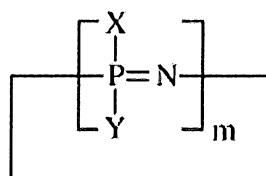
Step 1: dissolving from 5 to 50 parts by weight of reactive allylphenoxycyclotriphosphazene or vinylphenoxycyclotriphosphazene in a solvent until complete dissolution at room temperature or moderate temperature;

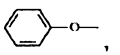
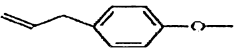
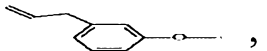
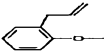
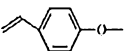
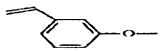
Step 2: adding from 15 to 85 parts by weight of an epoxy resin and a thermosetting resin, from 1 to 35 parts by weight of a crosslinking curing agent, from 0 to 5 parts by weight of a crosslinking curing accelerator and from 0 to 100 parts by weight of a packing into said solution, homogeneously stirring to obtain a liquid cement;

Step 3: homogeneously coating with said liquid cement E-glass cloth having a flattening surface, and then baking to a B-stage prepreg;

Step 4: cutting the B-stage prepreg to a suitable size according to the press size, stacking several prepreps regularly, a copper foil being placed over and under the prepreps, pressing in a vacuum hot press to obtain a copper clad laminate;

wherein said reactive allylphenoxycyclotriphosphazene or vinylcyclotriphosphazene has the following structural formula (I)



wherein m represents an integer of 3-25; X is phenoxy , an amount of 0-2m-1; Y is selected from the group consisting of p-allylphenoxy , m-allylphenoxy , o-allyl-phenoxy , p-vinylphen  and m-vinylphenoxy , in an amount of 1-2m; X+Y=2m;

wherein said epoxy resin is selected from the group consisting of bisphenol A epoxy resin, bisphenol F epoxy resin, dicyclopentadiene epoxy resin, triphenolic epoxy resin, biphenyl-type epoxy resin, naphthol epoxy resin, and phosphorous-containing epoxy resin; and

wherein the thermosetting resin is selected from the group consisting of the following resins:

benzoxazine resin, including bisphenol A-type benzoxazine resin, bisphenol F-type benzoxazine resin, dicyclopentadiene-type benzoxazine resin and phenolphthalein-type benzoxazine resin;

cyanate resin, including bisphenol A-type cyanate resin, dicyclopentadiene-type cyanate resin and phenolic aldehyde-type cyanate resin;

bismaleimide resin, including 4,4'-diphenylmethane bismaleimide resin, allyl-modified diphenylmethane bismaleimide resin;

reactive polyphenylene oxide resin, having a number average molecular weight of 1000-7000, wherein the reactive groups is hydroxyl or double bond; and

C5 hydrocarbon resin consisting of vinylbutylbenzene resin having a molecular weight of 11000, vinylpolybutadiene resin with polar groups and a copolymer in which butadiene is grafted to styrene with maleic anhydride.

6. The method for preparing a copper clad laminate according to claim 5, wherein the solvent is one or more selected from benzene and ketone solvents; said moderate temperature is $\leq 80^{\circ}\text{C}$.

7. The method for preparing a copper clad laminate according to claim 5, wherein, during the step 3 of baking the B-stage prepreg, the temperature is set up according to the boiling point of the solvent used therein, and ranges from 85 to 175°C, and the baking lasts 5-20 min.

8. The method for preparing a copper clad laminate according to claim 5, wherein, in the step 4, the pressing process is carried out by the bench pressing method, comprising the specific steps of increasing from room temperature to 150°C within 15 min and remaining for 30 min, then increasing to 180°C within 5 min and remaining for 2 h, finally decreasing to room temperature within 30 min; increasing the pressure from 0 to 0.6 Mpa within 1 min and remaining for 20 min, then increasing the pressure 1.0 Mpa within 1 min and remaining for 2.5 h and post-processing at 200-245°C and remaining for 1-5 h.