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(54) **SEMICONDUCTOR DEVICE**

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(57) **ABSTRACT**

A semiconductor device is extremely reliable because a sealant thereof is difficult to deteriorate even when a SiC semiconductor element is energized. The semiconductor device is produced by sealing a SiC semiconductor element **11** mounted on a multilayer substrate **12** and electrically conductive connection members **14** and **18** with a sealant **20** containing an ultraviolet light absorbent.

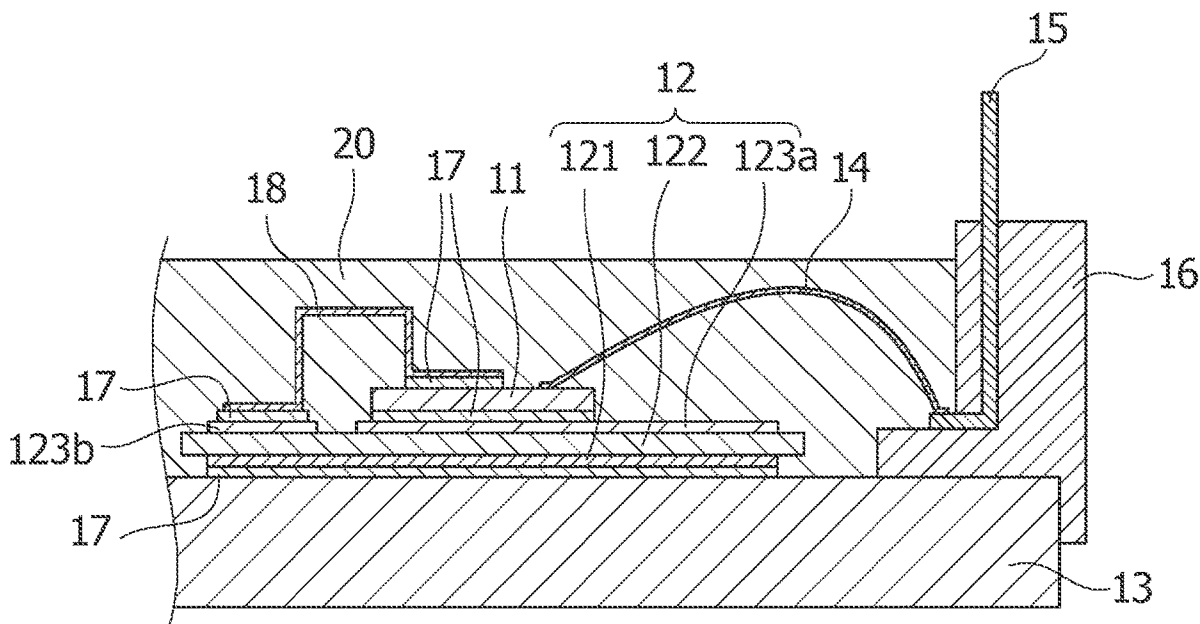


FIG.1

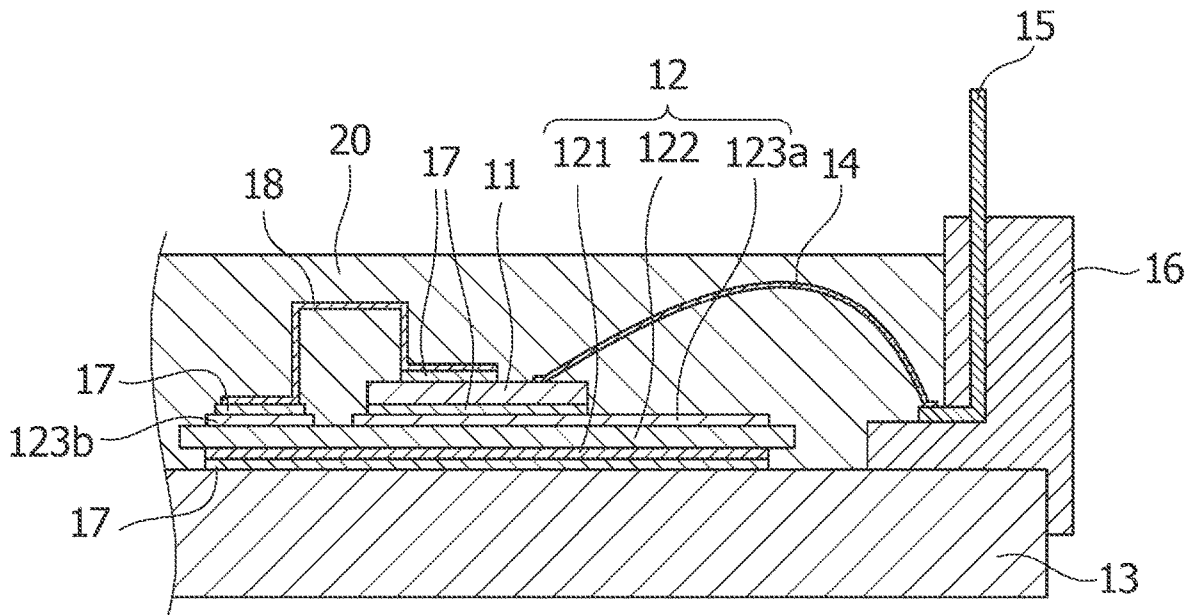


FIG.2

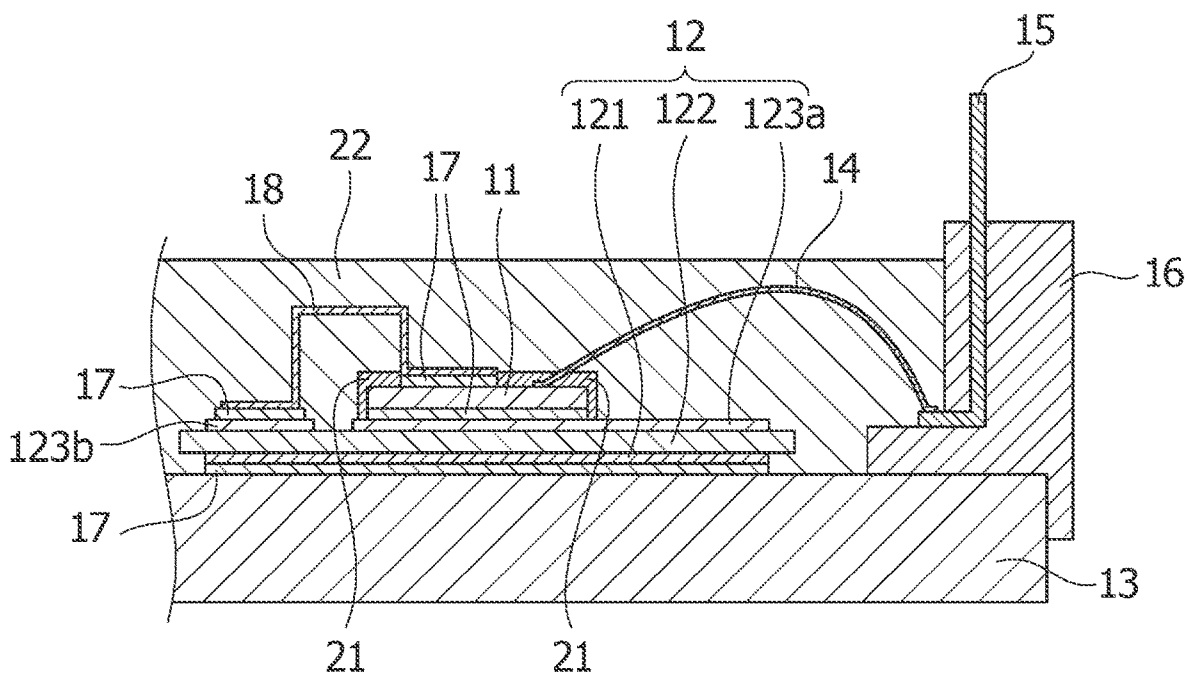
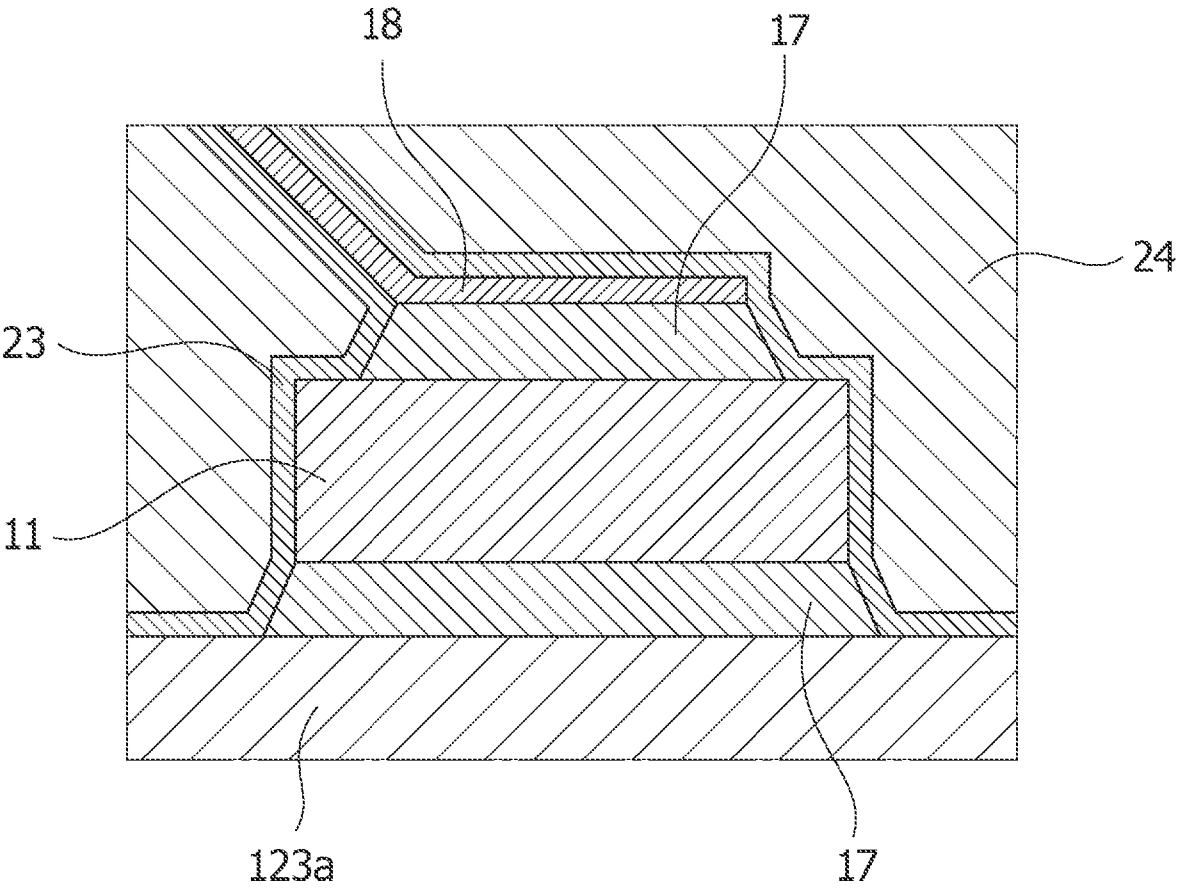


FIG.3



**SEMICONDUCTOR DEVICE**

**CROSS-REFERENCE TO RELATED APPLICATION**

**[0001]** This application claims priority from Japanese Patent Application No. 2021-118288, filed on Jul. 16, 2021, which is incorporated herein by reference in their entirety.

**BACKGROUND OF THE INVENTION**

**Field of the Invention**

**[0002]** The present invention relates to semiconductor devices. In particular, the present invention relates to a highly reliable semiconductor device that is less likely to have deterioration in a resin sealing SiC semiconductor elements even when SiC semiconductor elements are used.

**Description of Related Art**

**[0003]** Power semiconductor modules are used in a wide range of fields that require efficient power conversion. For example, the use of power semiconductor modules is expanding in the field of power electronics, such as industrial equipment, electric vehicles, and home appliances. Such a power semiconductor module contains switching elements and diodes, and silicon (Si) semiconductor elements or silicon carbide (SiC) semiconductor elements are used as the elements.

**[0004]** In particular, SiC semiconductor elements have about 10 times higher dielectric breakdown field strength than Si semiconductor elements, and therefore, enable a power device having a high withstand voltage of 600 V to several thousand volts to be fabricated with a higher impurity concentration and a thinner drift layer than Si semiconductor elements. Accordingly, SiC semiconductor elements are being used in practice in high withstand voltage devices with very low on-resistance per unit area.

**[0005]** Conventionally, resins containing a variety of functional materials have been known as sealants for semiconductor elements. In particular, resins containing ultraviolet light absorbent are known for the purpose of sealing optical semiconductor elements (for example, see Patent Documents 1 to 4).

**PATENT DOCUMENTS**

- [0006]** Patent Document 1: Japanese Patent Laid-Open No. 2006-328231
- [0007]** Patent Document 2: Japanese Patent Laid-Open No. 2003-309229
- [0008]** Patent Document 3: Japanese Patent Laid-Open No. 2004-277697
- [0009]** Patent Document 4: Japanese Patent Laid-Open No. 2009-99784

**SUMMARY OF THE INVENTION**

**[0010]** When a voltage is applied to a SiC semiconductor element for activation, the element emits ultraviolet light (UV) having wavelengths of about 360 nm to about 400 nm, exhibiting a peak at 393 nm (primary wavelength), which is equivalent to the SiC band gap. It has emerged that such UV can degrade the sealing resin for SiC semiconductor ele-

ments. More specifically, a degraded sealing resin has cracking around the chip, reducing the reliability in terms of power cycle or the like.

**[0011]** The present inventors found, through extensive research, that providing the sealant of a power semiconductor module including a SiC semiconductor element with an ultraviolet light absorbent in a specific manner can prevent the deterioration of the sealing resin and prevent the module from decreased reliability, thus completing the invention.

**[0012]** According to an embodiment, the present invention relates to a semiconductor device including a SiC semiconductor element mounted on a multilayer substrate and an electrically conductive connection member; and a sealant containing an ultraviolet light absorbent, which seals the SiC semiconductor element and the electrically conductive connection member.

**[0013]** In the semiconductor device, preferably, the sealant includes a thermosetting resin sealing layer. The thermosetting resin sealing layer preferably contains a thermosetting resin base material, a curing agent, an inorganic filler, and an ultraviolet light absorbent.

**[0014]** In an embodiment of the semiconductor device, the thermosetting resin sealing layer is preferably formed of a layer containing the thermosetting resin base material, the curing agent, the inorganic filler, and the ultraviolet light absorbent.

**[0015]** In an embodiment of the semiconductor device, the thermosetting resin sealing layer preferably includes a first sealing layer covering surroundings of the SiC semiconductor element, and a second sealing layer covering the first sealing layer. The first sealing layer preferably contains the thermosetting resin base material, the curing agent, the inorganic filler, and the ultraviolet light absorbent.

**[0016]** In an embodiment of the semiconductor device, the sealant preferably includes a primer layer covering the SiC semiconductor element and the electrically conductive connection member, and a thermosetting resin sealing layer covering the primer layer. The primer layer preferably contains an ultraviolet light absorbent.

**[0017]** In the semiconductor device, the ultraviolet light absorbent is preferably one, two or more selected from the group consisting of benzotriazole derivatives, benzophenone derivatives, and triazine derivatives.

**[0018]** In the semiconductor device including the thermosetting resin, the thermosetting resin base material preferably includes an epoxy resin base material.

**[0019]** In the semiconductor device including the primer layer, the primer layer preferably contains one or more resins selected from the group consisting of polyamide resins, polyimide resins, and polyamide-imide resins.

**[0020]** The present invention can provide a highly reliable semiconductor device in which ultraviolet light emitted from the SiC semiconductor element is absorbed to prevent deterioration of the resin forming the sealant.

**BRIEF DESCRIPTION OF THE DRAWINGS**

**[0021]** FIG. 1 is a conceptual sectional view of a semiconductor device according to a first embodiment of the present invention, illustrating the sectional structure of the semiconductor device;

**[0022]** FIG. 2 is a conceptual sectional view of a semiconductor device according to a second embodiment of the present invention, illustrating the sectional structure of the semiconductor device; and

[0023] FIG. 3 is a conceptual sectional view of a semiconductor device according to a third embodiment of the present invention, illustrating the sectional structure of the semiconductor device.

#### DESCRIPTION OF PREFERRED EMBODIMENTS

[0024] Embodiments of the present invention will now be described with reference to the drawings. The present invention is, however, not limited to the embodiments described below.

[0025] An embodiment of the present invention relates to a semiconductor device. The semiconductor device according to the present embodiment includes a SiC semiconductor element mounted on a multilayer substrate and an output terminal connected by an electrically conductive connection member; and a sealant which seals these members. The sealant contains an ultraviolet light absorbent.

[0026] The sealant used herein is an insulating resin in contact at least with a SiC semiconductor element, typically with the SiC semiconductor element, an output terminal, an electrically conductive connection member, and a laminated substrate, to insulate and seal these members. Hence, the concept of the sealant disclosed herein includes both a primer layer mainly containing a thermoplastic resin and an insulating thermosetting resin layer mainly containing a thermosetting resin. However, the present invention also encompasses semiconductor devices that include a sealant mainly containing a thermosetting resin layer without including a primer layer.

[0027] In the semiconductor device of the present invention, the sealant may include one or two or more different resin layers, and at least one of the resin layers is an ultraviolet light absorption layer containing an ultraviolet light absorbent that can absorb the ultraviolet light emitted from the SiC semiconductor element. The ultraviolet light absorbent may be an organic material that can absorb 390 nm ultraviolet light. In particular, the ultraviolet light absorbent is preferably a material that can absorb 30% or more of the ultraviolet light having a wavelength of 370 nm emitted from the SiC semiconductor element. The ultraviolet light absorption (%) of such a specific material can be measured with an ultraviolet-visible spectrophotometer. For example, a predetermined amount of an ultraviolet light absorbent is dispersed in toluene, and the transmittance of the dispersion liquid may be measured using a cell with a width of 1 cm (a length in the direction light transmits). The measurement wavelength may be varied in the range of 290 nm to 410 nm to estimate the ultraviolet light absorption (%) at 370 nm.

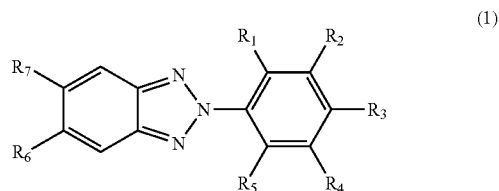
[0028] Specifically, compounds that can absorb 30% or more of 370 nm ultraviolet light include benzotriazole derivatives with no phenyl substituents, benzotriazole derivatives represented by formula (1) presented below, benzophenone derivatives represented by formula (2) presented below, and triazine derivatives represented by formula (3) presented below.

[0029] The benzotriazole derivative with no phenyl substituents may be a benzotriazole in which the H at the 1 or 3 position is substituted by OH and the H at the 5- or 6-position by a fluoromethyl group. Preferably, the fluoromethyl group is a trifluoromethyl group.

[0030] The benzotriazole derivative represented by formula (1) has a structure in which a phenyl group is bound to

the N atom at the 2-position of the triazole ring, having substituents  $R_1$  to  $R_5$  on the phenyl group and substituents  $R_6$  and  $R_7$  on the triazole ring.

[Formula 1]



In formula (1),

[0031]  $R_1$  and  $R_5$  each independently represent H or OH,

[0032]  $R_2$  and  $R_4$  each independently represent H, OH, a linear or branched, saturated or unsaturated, monovalent hydrocarbon group with 1 to 6 carbon atoms, or a linear or branched, saturated, or unsaturated monovalent hydrocarbon group with 1 to 6 carbon atoms in which a hydrogen atom is substituted by a phenyl group,

[0033]  $R_3$  represents H, OH, or a linear or branched, saturated alkoxy group with 1 to 10 carbon atoms, and

[0034]  $R_6$  and  $R_7$  each independently represent H or Cl.

[0035] One or two or more H atoms constructing substituents  $R_1$  to  $R_7$  may be substituted by Cl.

[0036] When either  $R_1$  or  $R_5$  is OH in formula (1), the other is preferably not OH.

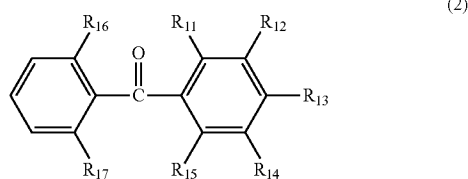
[0037] When  $R_2$  and/or  $R_4$  is a monovalent hydrocarbon group, the monovalent hydrocarbon group may be methyl, ethyl, propyl, n-butyl, or tert-butyl, and it is preferably a tert-butyl group, which is bulkier, among these. More preferably, both  $R_2$  and  $R_4$  are tert-butyl. When  $R_2$  and/or  $R_4$  is a linear or branched, saturated or unsaturated, monovalent hydrocarbon group with 1 to 6 carbon atoms in which a hydrogen atom is substituted by a phenyl group, the hydrocarbon group is preferably a propyl group in which a H atom is substituted by a phenyl group.

[0038] When  $R_3$  is an alkoxy group, the alkoxy group is preferably an octoxy group.

[0039] Specific examples of the benzotriazole derivatives include, but are not limited to, 2-(2-hydroxy-5-methylphenyl)benzotriazole, 2-(5-tert-butyl-2-hydroxyphenyl)benzotriazole, 2-(2-hydroxy-3-tert-butyl-5-methylphenyl)-5-chlorobenzotriazole, 2-(2-hydroxy-3,5-di-tert-butylphenyl)benzotriazole, 2-(2-hydroxy-3,5-di-tert-butylphenyl)-5-chloro-2H-benzotriazole, 2-(2-hydroxy-4-octoxyphenyl)benzotriazole, 2-(2H-benzotriazole-2-yl)-6-(1-methyl-1-phenylethyl)-4-(1,1,3,3-tetramethylbutyl)phenol, 1-hydroxy-6-(trifluoromethyl)benzotriazole, and 2-[2-hydroxy-3,5-bis( $\alpha,\alpha$ -dimethylbenzyl)phenyl]-2H-benzotriazole.

**[0040]** Benzophenone derivatives are represented by the following formula (2):

[Formula 2]



In formula (2),

**[0041]**  $R_{11}$ ,  $R_{12}$ ,  $R_{13}$ ,  $R_{14}$ ,  $R_{15}$ ,  $R_{16}$ , and  $R_{17}$  are each independently selected from H, OH, and linear or branched, saturated or unsaturated, hydrocarbon groups with 1 to 3 carbon atoms.

**[0042]** One or more of  $R_{11}$  to  $R_{17}$  is selected from OH, linear or branched alkyl groups with 1 to 3 carbon atoms, and linear or branched, saturated or unsaturated, alkoxy groups with 1 to 3 carbon atoms. One or two or more H atoms forming these substituents may be substituted by Cl.

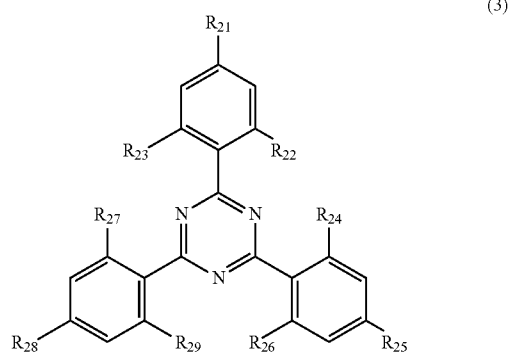
**[0043]** Preferably, in formula (2), either  $R_{11}$  and  $R_{15}$  is OH, and/or either  $R_{16}$  or  $R_{17}$  is OH. In this instance, preferably, both  $R_{11}$  and  $R_{15}$  are not OH at the same time and both  $R_{16}$  and  $R_{17}$  are not OH at the same time.

**[0044]** Preferably, either  $R_{13}$  or  $R_{14}$  is an alkyl group or an alkoxy group, and the other is H or OH. The alkyl group is preferably a methyl group. Also, the alkoxy group is preferably a methoxy group or an allyloxy group.

**[0045]** Specific examples of benzophenone derivatives include, but are not limited to, 2-hydroxy-4-methylbenzophenone, 2-hydroxy-4-methoxybenzophenone, 4-(allyloxy)benzophenone, and 2,2'-dihydroxy-4-methoxybenzophenone.

**[0046]** The triazine derivative represented by formula (3) has a structure in which a substituted or unsubstituted phenyl group is bound to each of the three carbon atoms of 1,3,5-triazine.

[Formula 3]



In formula (3),

$R_{21}$ ,  $R_{22}$ ,  $R_{23}$ ,  $R_{24}$ ,  $R_{25}$ ,  $R_{26}$ ,  $R_{27}$ ,  $R_{28}$ , and  $R_{29}$  are each independently selected from H, OH, linear or branched alkoxy groups with 1 to 8 carbon atoms, and linear or

branched alkyl groups with 1 to 3 carbon atoms, provided that one or more substituents of  $R_{21}$  to  $R_{29}$  are selected from OH, linear or branched alkoxy groups with 1 to 8 carbon atoms and linear or branched alkyl groups with 1 to 3 carbon atoms. One or two or more H atoms forming these substituents may be substituted by Cl.

**[0047]** Preferably, at least one, more preferably two, or all three, of one substituent of  $R_{23}$  and  $R_{22}$ , one substituent of  $R_{24}$  and  $R_{26}$ , and one substituent of  $R_{27}$  and  $R_{29}$  is OH. In this instance, preferably, both  $R_{23}$  and  $R_{22}$  are not OH at the same time, both  $R_{24}$  and  $R_{26}$  are not OH at the same time, and both  $R_{27}$  and  $R_{29}$  are not OH at the same time.

**[0048]** Preferably, at least one of  $R_{21}$ ,  $R_{25}$ , and  $R_{28}$  is a linear or branched alkyl group with 1 to 3 carbon atoms. More preferably, this alkyl group is a methyl group. Furthermore, at least one of  $R_{21}$ ,  $R_{25}$ , and  $R_{28}$  is preferably an alkoxy group. More preferably, two or all of  $R_{21}$ ,  $R_{25}$ , and  $R_{28}$  are alkoxy groups. Still more preferably, the alkoxy group is a butoxy group or an octyloxy group. One or two of  $R_{21}$ ,  $R_{25}$ , and  $R_{28}$  may be an alkyl group, and the rest may be an alkoxy group.

**[0049]** Specific examples of triazine derivatives include, but are not limited to, 2-(2,4-dihydroxyphenyl)-4,6-diphenyl-1,3,5-triazine, 2,4,6-tris(2,4-dihydroxyphenyl)-1,3,5-triazine, 2-(2,4-dihydroxyphenyl)-4,6-bis(2,4-dimethylphenyl)-1,3,5-triazine, 2-(2-hydroxy-4-methoxyphenyl)-4,6-diphenyl-1,3,5-triazine, 2,4-bis(2,4-dimethylphenyl)-6-(2-hydroxy-4-n-octyloxyphenyl)-1,3,5-triazine, 2,4,6-tris(4-butoxy-2-hydroxyphenyl)-1,3,5-triazine, and 2,4-bis(2-hydroxy-4-butoxyphenyl)-6-(2,4-bis-butoxyphenyl)-1,3,5-triazine.

**[0050]** The benzotriazole, benzophenone, and triazine derivatives described above contain hydrophobic alkyl or phenyl groups and are therefore highly dispersible in the matrix resin (thermosetting resin or thermoplastic resin) of the ultraviolet light absorption layer. Many of these ultraviolet light absorbents are unlikely to react with the matrix resin such as epoxy resin, phenol resin, or maleimide resin. The ultraviolet light absorbent in the ultraviolet light absorption layer also holds particles or the like separate from the matrix resin even if the ultraviolet light absorbent melts after curing. Alkoxy group-containing derivatives are highly reactive and likely to bind to the inorganic filler or any other inorganic material contained in the ultraviolet light absorption layer, or Cu or the like contained in the members that may contact the ultraviolet light absorption layer.

**[0051]** The ultraviolet light absorbent can be dispersed in the matrix resin (thermosetting resin or thermoplastic resin) of the ultraviolet light absorption layer. In view of the dispersibility, the average particle size of the ultraviolet light absorbent may be 10 nm to 5000 nm and is preferably 50 nm to 500 nm. In particular, the ultraviolet light absorbent of a benzotriazole derivative may have an average particle size of 10 nm to 1000 nm, preferably 50 nm to 200 nm. Benzotriazole derivatives and triazine derivatives do not form covalent bonds with the matrix resin and have high melting points. Such an ultraviolet light absorbent holds particles separate from the matrix resin even if the ultraviolet light absorbent melts after curing. Accordingly, the particle size of the ultraviolet light absorbent can be controlled to be smaller than that of the inorganic filler to increase the dispersibility.

**[0052]** In the present invention, the ultraviolet light sealing layer can be provided in several ways in the sealant. The

present invention will be further described in detail with reference to a first, a second, and a third embodiment in which the ultraviolet light sealing layer is disposed in different ways.

#### First Embodiment

**[0053]** In the semiconductor device according to a first embodiment, the sealant includes a thermosetting resin sealing layer. The thermosetting resin sealing layer consists of a layer containing a thermosetting resin base material, a curing agent, an inorganic filler, and an ultraviolet light absorbent.

**[0054]** FIG. 1 is a conceptual sectional view of a power semiconductor module that is an implementation of the semiconductor device according to the first embodiment of the present invention. The illustrated power semiconductor module has a multilayer structure in which a SiC semiconductor element **11** and a laminated substrate **12** are bonded together on a heat sink **13** with a bonding layer **17**. A case **16** having an external terminal **15** therein is bonded to the heat sink **13**. An electrode of the SiC semiconductor element **11** and an electrode of the laminated substrate **12** are connected to each other by a lead frame **18** acting as an electrically conductive connection member. The SiC semiconductor element **11** and the external terminal **15** are connected by an aluminum wire **14**. A thermosetting resin sealing layer **20** is filled on the SiC semiconductor element **11** and laminated substrate **12**, the lead frame **18**, and electrically conductive connection members including the aluminum wire **14** in such a manner as to contact with these members. Hence, the sealant consists essentially of the thermosetting resin sealing layer **20**, and the thermosetting resin sealing layer **20** functions as an ultraviolet light absorption layer.

**[0055]** The SiC semiconductor element **11** is an insulated gate bipolar transistor (IGBT) chip, a diode chip, or any other power chip. A semiconductor element used may be a combination of Si devices and wide gap semiconductor devices, such as GaN devices, diamond devices, or ZnO devices. For example, a hybrid module using a Si-IGBT and a SiC Schottky barrier diode (SiC-SBD) may be used. The number of the SiC semiconductor elements **11** to be mounted may be one or two or more.

**[0056]** The laminated substrate **12** may be composed of an insulation substrate **122**, a first electrically conductive plate **121** with a predetermined form (pattern) formed on one of the main surfaces of the insulation substrate, and second electrically conductive plates **123a** and **123b** formed on the other main surface. A material excellent in electrical insulation and thermal conductivity may be used as the insulation substrate **122**. Examples of the material of the insulation substrate **122** include  $\text{Al}_2\text{O}_3$ , AlN, and SiN. For usage requiring high withstand voltage, in particular, materials satisfying both electrical insulation and thermal conductivity are preferred, and AlN and SiN may be used, but they are not limited to these materials. The first electrically conductive plate **121** and the second electrically conductive plates **123a** and **123b** may be made of a metal material having excellent workability, such as Cu or Al. For rust prevention, the electrically conductive plates may be made of Ni-plated Cu or Al or the like. The electrically conductive plates **121**, **123a**, and **123b** may be disposed on the insulation substrate **122** by direct copper bonding or active metal brazing. In the illustrated embodiment, the two second electrically conduc-

tive plates **123a** and **123b** are disposed in a discontinuous manner on the insulation substrate **122**. One (**123a**) of the second electrically conductive plate may function as an electrode to be connected to the SiC semiconductor element **11**, and the other (**123b**) may function as an electrode to be connected to the lead frame **18**.

**[0057]** The lead frame **18** is an electrically conductive connection member connecting the SiC semiconductor element **11**, the second electrically conductive plate **123b**, and the like to each other. More specifically, the lead frame **18** is bonded to an electrode (surface electrode) of the SiC semiconductor element **11** by a bonding layer **17** made of a solder material or the like. The lead frame **18** is also bonded to the second electrically conductive plate **123b** and other wiring members by a bonding layer **17** made of a solder material or the like. The lead frame **18** may be a metal, such as copper or an alloy containing copper. The lead frame **18** may be provided with a Ni or Ni alloy layer or a Cr or Cr alloy layer on the surface thereof by plating. In this instance, the thickness of the Ni or Ni alloy layer or the Cr or Cr alloy layer may be about not more than 20  $\mu\text{m}$ . The electrically conductive connection member connecting the SiC semiconductor element **11** to the second electrically conductive plate **123b** and the like may be a lead terminal having a pin shape or a wire, such as an aluminum or copper wire, used for wire bonding, instead of the lead frame.

**[0058]** The heat sink **13** may be made of a thermally conductive metal, such as copper or aluminum. Also, the heat sink **13** may be coated with Ni or an Ni alloy for rust prevention. A heat sink used here may be a cooling device with water cooling or air cooling capability.

**[0059]** The bonding layer **17** may be formed by using a lead-free solder. For example, an Sn—Ag—Cu, Sn—Sb, Sn—Sb—Ag, Sn—Cu, Sn—Sb—Ag—Cu, Sn—Cu—Ni, Sn—Ag, or similar solder may be used, but the lead-free solder is not limited to these. Alternatively, a bonding layer used here may be formed by using a bonding material containing sintered silver nanoparticles or other fine metal particles.

**[0060]** The case **16** may be made of a thermoplastic resin, such as polyphenylene sulfide (PPS) or polybutylene terephthalate (PBT).

**[0061]** In the present embodiment, the SiC semiconductor element **11**, the laminated substrate **12**, and the electrically conductive connection members including the lead frame **18** and the aluminum wire **14** may be referred to as members to be sealed. The space over the members to be sealed is filled with the sealant formed of the thermosetting resin sealing layer **20**. Thus, the thermosetting resin sealing layer **20** contacts the SiC semiconductor element **11**, the laminated substrate **12**, and the electrically conductive connection members to cover the surroundings of these members to be sealed. The thickness of the thermosetting resin sealing layer **20** varies depending on the shape or the like of the module and may be about 0.1 mm to 2 mm.

**[0062]** The thermosetting resin sealing layer **20** is a cured product of a thermosetting resin composition containing a thermosetting resin base material, a curing agent, an ultraviolet light absorbent, and an inorganic filler, and optionally, a curing accelerator and other additives.

**[0063]** Examples of the thermosetting resin base material include, but are not limited to, epoxy, phenol, and maleimide resins which have heat resistance and high insulation. Among others, epoxy resins having at least two epoxy

groups in the molecule is particularly preferable. Such epoxy resins are dimensionally stable, resistant to water and chemicals, and highly electrically insulative. Specifically, an aliphatic epoxy resin, an alicyclic epoxy resin, or a mixture thereof is preferably used.

**[0064]** An aliphatic epoxy resin refers to an epoxy compound in which the carbon atom bound directly to the epoxy group is a carbon atom forming an aliphatic hydrocarbon. Therefore, provided that the same holds true, even a compound having an aromatic ring in the backbone is also an aliphatic epoxy resin. Examples of aliphatic epoxy resins include, but are not limited to, bisphenol A epoxy resin, bisphenol F epoxy resin, bisphenol AD epoxy resin, biphenyl epoxy resin, naphthalene epoxy resin, cresol novolac epoxy resin, and trifunctional or more polyfunctional epoxy resins. These epoxy resins may be used singly or in combination of two or more. Naphthalene epoxy resin and trifunctional or more polyfunctional epoxy resins have high glass transition temperatures and are accordingly referred to as highly heat-resistant epoxy resins. Using such a highly heat-resistant epoxy resin can enhance heat resistance.

**[0065]** An alicyclic epoxy resin refers to an epoxy compound in which the two carbon atoms of the epoxy group are members forming an alicyclic compound. Alicyclic epoxy resins include, but are not limited to, monofunctional epoxy resin, bifunctional epoxy resin, and trifunctional or higher polyfunctional epoxy resins. Alicyclic epoxy resins may also be used singly or in combination of two or more. Mixing an alicyclic epoxy resin with an acid anhydride curing agent and curing the mixture forms a product with a high glass transition temperature. Accordingly, mixing an aliphatic epoxy resin with an alicyclic epoxy resin can form a product with a higher heat resistance.

**[0066]** The thermosetting resin base material used in the thermosetting resin sealing layer **20** may be a mixture of the above-described aliphatic epoxy resins and alicyclic epoxy resins. Aliphatic epoxy resin and alicyclic epoxy resin may be mixed in any proportion and may be, but is not limited to, in a mass ratio of about 2:8 to 8:2 or about 3:7 to 7:3. In a preferred embodiment, the thermosetting resin base material used in the thermosetting resin sealing layer **20** is a mixture of bisphenol A epoxy resin and alicyclic epoxy resin in a mass ratio of 1:1 to 1:4.

**[0067]** The thermosetting resin sealing layer **20** preferably contains 0.1 parts to 10 parts by mass, more preferably 1 part to 5 parts by mass, of an ultraviolet light absorbent relative to 100 parts by mass of the matrix resin of the thermosetting resin sealing layer **20** in view of reliability. The matrix resin used in the present embodiment contains the thermosetting resin base material and the curing agent, but not the inorganic filler. In general, the ultraviolet light absorbent can be uniformly dispersed in the thermosetting resin sealing layer **20** in the above-mentioned range of content. However, the concentration of the ultraviolet light absorbent may vary among portions of the thermosetting resin sealing layer **20** and is preferably set to be higher in the vicinity of the semiconductor element **11**. In the first embodiment, the thermosetting resin sealing layer **20** preferably contains an ultraviolet light absorbent having one or more OH groups selected from the ultraviolet light absorbents represented by general formulas (1), (2), and (3). This produces an additional effect that when the thermosetting resin sealing layer

**20** comes into contact with the electrically conductive plate **123a** of the laminated substrate **12** and other members, the adhesion can be enhanced.

**[0068]** The inorganic filler essentially contained in the thermosetting resin sealing layer **20** may be a metal oxide or metal nitride having a high thermal conductivity and a low linear expansion coefficient. Examples of such an inorganic filler include, but are not limited to, fused silica, silica (silicon oxide), alumina, aluminum hydroxide, titania, zirconia, aluminum nitride, talc, clay, mica, and glass fiber. The inorganic filler can increase the thermal conductivity of the thermosetting resin sealing layer **20** and reduce the thermal expansion coefficient. The thermosetting resin sealing layer **20** is preferably electrically insulative. Accordingly, the inorganic filler is also preferably insulative. Inorganic fillers may be used singly or in combination of two or more as a mixture. The inorganic filler may be a microfiller, a nanofiller, or a mixture of two or more inorganic fillers of different particle size and/or material. Preferably, the inorganic filler has an average particle size of about 0.2  $\mu\text{m}$  to 20  $\mu\text{m}$ . The amount of the inorganic filler added in the thermosetting resin sealing layer **20** is preferably 100 parts to 600 parts by mass, more preferably 200 parts to 400 parts by mass, relative to 100 parts by mass of the matrix resin. When the amount of the inorganic filler added is less than 100 parts by mass, the thermal expansion coefficient of the thermosetting resin sealing layer **20** is likely to increase, causing peeling and cracking. In contrast, when the amount of the inorganic filler is greater than 600 parts by mass, the composition has an increased viscosity and can be difficult to extrude.

**[0069]** The curing agent essentially contained in the thermosetting resin sealing layer **20** is reactive to the thermosetting resin base material, preferably epoxy resin base material, to cure the base material and is not otherwise limited. Preferably, an acid anhydride-based curing agent is used. Examples of the acid anhydride-based curing agent include aromatic acid anhydrides, such as phthalic anhydride, pyromellitic anhydride, and trimellitic anhydride. Other examples include cyclic aliphatic acid anhydrides, such as tetrahydrophthalic anhydride, methyltetrahydrophthalic anhydride, hexahydrophthalic anhydride, methylhexahydrophthalic anhydride, and methylnadac anhydride; and aliphatic acid anhydrides, such as succinic anhydride, polyadipic anhydride, polysebacic anhydride, and polyazelaic anhydride. Preferably, the curing agent is added in a proportion of about not less than 50 parts and not greater than 170 parts by mass, more preferably about not less than 80 parts and not greater than 150 parts by mass, relative to 100 parts by mass of the epoxy resin base material. When the proportion of the curing agent is less than 50 parts by mass, the thermosetting resin cannot have sufficient crosslinks, resulting in reduced glass transition temperature. When the proportion of the curing agent exceeds 170 parts by mass, the thermosetting resin may deteriorate in moisture resistance, heat resistance, and heat stability. In an embodiment using a bisphenol A epoxy resin independently or using a mixture of a bisphenol A epoxy resin and any of the above-cited highly heat-resistant epoxy resins, it may be preferable not to use the curing resin from the viewpoint of increasing the heat resistance. The highly heat-resistant epoxy resin may be added in a proportion of, for example, not less than 10% and not more than 50% by mass, preferably not less than 10% and not more than 25% by mass,



relative to 100% by mass in total of the thermosetting resin base materials. In such a range, the heat resistance is preferably increased without increasing the viscosity.

[0070] The thermosetting resin composition forming the thermosetting resin sealing layer 20 may optionally contain a curing accelerator. Examples of the curing accelerator include imidazole and its derivatives, tertiary amines, boric acid esters, Lewis acids, organic metal compounds, and organic acid metal salts, and any one of them can be added if necessary. Preferably, the curing accelerator is added in a proportion of about 0.01 part or more and 50 parts or less by mass, more preferably about 0.1 parts or more and 20 parts or less by mass, relative to 100 parts by mass of the thermosetting resin base material.

[0071] The thermosetting resin composition forming the thermosetting resin sealing layer 20 may optionally contain other additives as long as the properties thereof are not inhibited. Such additives include, but are not limited to, a flame retardant, a pigment for coloring the resin, and a plasticizer or silicone elastomer to enhance crack resistance. Such optionally added constituents and their content can be appropriately determined by those skilled in the art according to the specifications required for the semiconductor device and/or the thermosetting resin sealing layer 20.

[0072] The process for producing the illustrated power semiconductor module will now be described. After bonding the heat sink 13, the laminated substrate 12, and the SiC semiconductor element 11 together and attaching the case 16 to the heat sink 13, the lead frame 18 is bonded, and wire bonding process is performed using the aluminum wire 14. Then, a thermosetting resin composition to form the thermosetting resin sealing layer 20 is injected into the case 16, followed by heat curing. The heat curing may be conducted, for example, in two stages. In an embodiment using an epoxy resin as the thermosetting resin base material, the thermosetting resin composition may be heated at 90° C. to 120° C. for 1 to 2 hours to be brought into a semi-cured state (preliminary curing). Subsequently, the semi-cured composition is further heated at 175° C. to 185° C. for 1 to 2 hours (full curing). The full curing may be performed by continuously increasing temperature from the temperature for the preliminary curing. The temperature and time for the heat curing are not limited to specific values, and the two-stage curing is not necessarily required. For preliminary curing, preferably, the temperature is slowly increased from room temperature at a rate of 0.5° C./min to 2° C./min. This manner increases the dispersibility of the ultraviolet light absorbent, thus increasing the reliability of the resulting module.

[0073] In a modification of the illustrated power semiconductor module, the sealant may further include a primer layer. The primer layer may be disposed at the interface between the thermosetting resin sealing layer 20 and the members to be sealed including the laminated substrate, the semiconductor element, and the electrically conductive connection members. The primer layer can relieve stress at the interface between the thermosetting resin sealing layer 20 and the sealed members and may be preferably used from the viewpoint of ensuring adhesion. The primer layer may be made of a resin containing polyamide, polyimide, or polyamide-imide. Resins containing polyamide, polyimide, or polyamide-imide are referred to as primer material. Polyimide is a general term for polymer compounds containing imide bonds. A polymer having a cyclic imide structure is

preferable. Examples of the polyimide that can be used include, but are not limited to, PMDA/4,4'-ODA (copolymer of pyromellitic dianhydride and 4,4'-oxydianiline), BPDA/PDA (copolymer of 3,3',4,4'-biphenyltetracarboxylic dianhydride and p-phenylenediamine), and BPDA/4,4'-ODA (copolymer of 3,3',4,4'-biphenyltetracarboxylic dianhydride and 4,4'-oxydianiline). Polyamide is a general term for polymer compounds containing amide bonds, and polyamide-imide is a general term for polymer compounds containing both imide bonds and amide bonds.

[0074] In the modification of the first embodiment, the primer layer may contain, but need not contain, an ultraviolet light absorbent. The primer layer may be formed to any thickness, provided that the primer layer functions to enhance adhesion. For the primer layer containing no ultraviolet light absorbent, the thickness may be, for example, about 1 μm to 20 μm and is preferably 1 μm to 10 μm. The primer layer with such a thickness can relieve stress and enhance adhesion, thus being advantageous. The primer layer may be provided to cover the entirety of the surfaces of the SiC semiconductor element 11, laminated substrate 12, lead frame 18, and aluminum wire 14, illustrated in FIG. 1. The primer layer may contain silica or any other inorganic filler to, for example, control the linear expansion coefficient. In this instance, the inorganic filler may have an average particle size of 10 nm to 1000 nm and may be added in a range of 0.5 parts to 3 parts by mass relative to 100 parts by mass of the primer layer material. The inorganic filler in such ranges does not inhibit the effect of a primer to relieve stress.

[0075] In another modification of the first embodiment, the sealant may further include one or more additional resin sealing layers other than the thermosetting resin sealing layer. For example, the sealant may include a thermoplastic resin layer which can cover the air-contacting surface of the thermosetting resin sealing layer 20. In this modification, a primer layer may or may not be provided.

[0076] In a process for producing the power semiconductor module including the primer layer, the primer layer is formed after assembling the members to be sealed and before injecting the thermosetting resin composition to form the thermosetting resin sealing layer 20. The primer layer may be formed over the entire surfaces of the members to be sealed depicted in FIG. 1, including the SiC semiconductor element 11, the lead frame 18, the laminated substrate 12, the aluminum wire 14, and the case 16 by, for example, spray coating, dip coating, or using a dispenser. After the formation of the primer layer, the resulting structure is preferably heated in a nitrogen inert gas oven in stages at 70° C. to 100° C. for about 60 min to 80 min and at 200° C. to 220° C. for 60 min to 80 min. This heating operation heats the Cu of the lead frame 18 and evaporates the solvent to solidify the primer. After the primer layer is thus formed, the resulting structure can be subjected to insulation and encapsulation with the thermosetting resin sealing layer 20 in the same manner as the process for producing the power semiconductor module depicted in FIG. 1. When optional additional resin sealing layers are provided, they may be formed in ordinary manners.

[0077] The structure of the illustrated power semiconductor module is merely an example and is not intended to limit the present invention. For example, any electrically conductive connection members may be used, and implant pins may be used. Only the lead frame or the wire may be the

electrically conductive member. In a modification of the semiconductor module in which the electrically conductive connection members include an implant pin, when the primer layer is provided, it may cover the surface of the implant pin. In a modification of the semiconductor module in which only the wire is the electrically conductive connection member, when the primer layer is provided, it may cover the surface of the wire.

**[0078]** The power semiconductor module may be a caseless module including no case. The caseless power semiconductor module (not shown) may include an implant pin and a printed substrate bonded to the implant pin, instead of the lead frame and the aluminum wire depicted in FIG. 1, and the members to be sealed, including these connection members are sealed with the thermosetting resin sealing layer. The printed substrate may be a polyimide or epoxy film provided with a Cu or Al electrically conductive layer or the like on the film. The implant pin may be a copper pin including copper. The Cu or Al of the implant pin and of the electrically conductive layer of the printed substrate may be plated with Ni or the like for rust prevention. The printed substrate and implant pin electrically connect the semiconductor elements one to another or the semiconductor element to the laminated substrate. The implant pin and the laminated substrate or the semiconductor element may be bonded with a solder bonding layer therebetween. The implant pin can be led out from the laminated substrate out of the thermosetting resin sealing layer and thus connected to an external connection terminal. For producing such a caseless power semiconductor module, the member to be sealed, including the laminated substrate, the semiconductor element, the implant pin, and the printed substrate, are assembled, and optionally a primer layer is formed over the laminated substrate, the semiconductor element, the implant pin, and the printed substrate by, for example, spray coating. The assembly of those members is placed in an appropriate mold, and the mold is filled with the thermosetting resin composition for forming the thermosetting resin sealing layer, followed by curing. Such a sealant may be formed by, but is not limited to, vacuum injection, transfer molding, liquid transfer molding, potting, or any other forming technique.

**[0079]** In the present embodiment, substantially the entirety of the thermosetting resin sealing layer acts as the ultraviolet light absorption layer. The present embodiment is advantageous for preventing the deterioration of the sealing resin by ultraviolet light without increasing the number of process steps.

#### Second Embodiment

**[0080]** In the semiconductor device according to a second embodiment, the sealant includes a thermosetting resin sealing layer. The thermosetting resin sealing layer includes a first sealing layer covering the surroundings of the SiC semiconductor element, and a second sealing layer covering the first sealing layer. The first sealing layer (ultraviolet light absorption layer) contains a thermosetting resin base material, a curing agent, an inorganic filler, and an ultraviolet light absorbent. Hence, in the second embodiment, one or some of the layers defining the thermosetting resin sealing layer function as an ultraviolet light absorption layer.

**[0081]** FIG. 2 is a conceptual sectional view of a power semiconductor module that is an implementation of the semiconductor device according to the second embodiment

of the present invention. In the illustrated power semiconductor module, a first sealing layer 21 is formed in contact with a SiC semiconductor element 11, a laminated substrate 12, a lead frame 18, electrically conductive connection members including an aluminum wire 14, and other members to be sealed, and a second sealing layer 22 is formed around the first sealing layer 21 and the members to be sealed. In other words, the sealant is formed of a thermosetting resin sealing layer, and the thermosetting resin sealing layer includes the first sealing layer 21 and the second sealing layer 22. The SiC semiconductor element 11 is bonded onto the second electrically conductive plate 123a of the laminated substrate 12 with a bonding layer 17 made of a solder material or the like therebetween. Since ultraviolet light is emitted mainly through one or more lateral faces of the SiC element, the first sealing layer 21 covers at least the one or more lateral faces of the SiC semiconductor element 11 and may cover the top face of the SiC semiconductor element 11. Also, the first sealing layer 21 may be disposed to cover one or more exposed ends of the bonding layer 17 and contact the second electrically conductive plate 123a. In a structure in which the first sealing layer 21 contacts the second electrically conductive plate 123a, the contact portion with the second electrically conductive plate 123a preferably has a length of 0.5 mm to 1.0 mm in view of adhesion. The length of the contact portion refers to the length from the exposed end of the bonding layer 17 in contact with the second electrically conductive plate 123a to the end of the first sealing layer 21 along the second electrically conductive plate 123a. Also, the first sealing layer 21 may be applied to contact the insulation substrate 122 of the laminated substrate 12.

**[0082]** In the embodiment illustrated in FIG. 2, typically, only the first sealing layer 21 is in contact with the SiC semiconductor element 11, and the second sealing layer 22 does not contact the SiC semiconductor element 11. The first sealing layer 21 may have the same composition as the thermosetting resin sealing layer 20 of the first embodiment. The thickness of the first sealing layer 21 is preferably 2 μm to 1000 μm and more preferably 100 μm to 1000 μm. The thickness mentioned here is the thickness of the first sealing layer 21 when covering the lateral and top faces of the SiC semiconductor element 11. In view of reliability, the first sealing layer 21 preferably contains 0.1 parts to 10 parts by mass, more preferably 1 part to 5 parts by mass, of an ultraviolet light absorbent relative to 100 parts by mass of the matrix resin of the first sealing layer 21. The matrix resin used in the present embodiment refers to a concept which contains the thermosetting resin base material and the curing agent, but not the inorganic filler. In general, the ultraviolet light absorbent is uniformly dispersed in the first sealing layer 21 in the above-mentioned content range. However, the concentration of the ultraviolet light absorbent may be varied among portions of the first sealing layer 21 and is preferably higher in the vicinity of the semiconductor element 11. Alternatively, the ultraviolet light absorbent may be added in the same amount as in the first embodiment.

**[0083]** The first sealing layer 21 contacts part of the electrically conductive member, such as the lead frame 18, connected to the top face of the SiC semiconductor element 11 and may or may not cover the entirety of the lead frame 18. The lead frame and other electrically conductive members are typically made of copper or aluminum. Such material does not transmit ultraviolet light emitted from the

SiC semiconductor element **11** and need not be covered. In the second embodiment, the first sealing layer **21** preferably contains an ultraviolet light absorbent having one or more OH groups selected from the ultraviolet light absorbents represented by formulas (1), (2), and (3). This produces an additional effect that when the first sealing layer **21** comes into contact with the electrically conductive plate **123a** of the laminated substrate **12** and other members, the adhesion can be enhanced.

**[0084]** The second sealing layer **22** contains a thermosetting resin base material, a curing agent, and an inorganic filler. The ultraviolet light absorbent may or may not be contained, and if it is contained, the amount is smaller than in the first sealing layer **21**. The thermosetting resin base material and the curing agent in the second sealing layer **22** are preferably the same as those in the first sealing layer **21**. Thus, the linear expansion coefficients of the first sealing layer **21** and the second sealing layer **22** can be the same level, preventing separation at the interface between the first sealing layer **21** and the second sealing layer **22**. For the same reason, the type and content of the inorganic filler of the second sealing layer **22** are preferably the same as those of the first sealing layer **21**.

**[0085]** In the second embodiment, a primer layer (not shown) may be provided between the first sealing layer **21** and the members to be sealed including the SiC semiconductor element **11** or between the second sealing layer **22** and the laminated substrate and other members to be sealed. The composition and thickness of the primer layer may be the same as in the first embodiment.

**[0086]** The semiconductor device of the second embodiment may be produced in the same manner as in the first embodiment, except for the step of forming the sealant. In the second embodiment, the first sealing layer **21** is formed, optionally followed by preliminary curing, and then the second sealing layer **22** is formed. After the preliminary curing and full curing of the first sealing layer **21**, the second sealing layer **22** may be formed, followed by preliminary curing and full curing. However, in view of the adhesion between the first sealing layer **21** and the second sealing layer **22**, the second sealing layer **22** is preferably formed after the preliminary curing of the first sealing layer **21** and is then subjected to preliminary curing and full curing. The first sealing layer **21** may be formed by, but is not limited to, spray coating or using a dispenser. The semiconductor device of the second embodiment can be modified in the same manner as in the modifications of the first embodiment.

**[0087]** In the semiconductor device according to the second embodiment of the present invention, a thermosetting resin sealing layer partially containing an ultraviolet light absorbent is disposed around the SiC semiconductor element. This structure enables the ultraviolet light absorbent to efficiently absorb ultraviolet light even in a small amount and protects the sealing resin with adhesion at the interfaces maintained. Ultraviolet light absorbents are difficult to react with thermosetting resin, notably epoxy resin. When the layer containing an ultraviolet light absorbent is large, the dispersion of the ultraviolet light absorbent may not be uniform. In the semiconductor device according to the second embodiment of the present invention, the layer containing an ultraviolet light absorbent (first sealing layer) is thin and small. Accordingly, a necessary and sufficient amount of ultraviolet light absorbent can probably be provided around the SiC semiconductor element.

### Third Embodiment

**[0088]** The semiconductor device according to a third embodiment includes a SiC semiconductor element mounted on a laminated substrate and an electrically conductive connection member, and a sealant containing an ultraviolet light absorbent, which seals the SiC semiconductor element and the electrically conductive connection member. The sealant includes a primer layer and a thermosetting resin sealing layer. The primer layer contains the ultraviolet light absorbent.

**[0089]** FIG. 3 is a fragmentary sectional view of a power semiconductor module that is an implementation of the semiconductor device according to the third embodiment of the present invention, illustrating a SiC semiconductor element and its vicinity. In the illustrated power semiconductor module, a primer layer **23** is disposed in contact with a SiC semiconductor element **11**, a second electrically conductive plate **123a**, a lead frame **18**, and other members to be sealed, and a thermosetting resin sealing layer **24** covers the primer layer **23**. More specifically, the primer layer **23** covers the SiC semiconductor element **11**, the laminated substrate, the lead frame **18**, and other members with wiring members to be sealed. Hence, the sealant includes the primer layer **23** and the thermosetting resin sealing layer **24**, and the primer layer **23** functions as an ultraviolet light absorption layer.

**[0090]** In the embodiment illustrated in FIG. 3, typically, only the primer layer **23** is in contact with the SiC semiconductor element **11**, and the thermosetting resin sealing layer **24** does not contact the SiC semiconductor element **11**. The primer layer **23** contains an ultraviolet light absorbent in addition to the composition of the primer layer described in a modification of the first embodiment. The primer layer **23** preferably contains 0.1 parts to 5 parts by mass, more preferably 0.5 parts to 2.0 parts by mass, of an ultraviolet light absorbent relative to 100 parts by mass of the matrix resin of the primer layer. The matrix resin in the primer layer used in the present embodiment contains a primer material but not the inorganic filler. The primer layer **23** may contain an inorganic filler to control the linear expansion coefficient but, preferably, contains no inorganic filler. Adding an inorganic filler in the primer layer reduces the ultraviolet light absorbent content by the amount of the inorganic filler added and may reduce the effect of relaxing stress. The ultraviolet light absorbent contained in the primer layer **23** is preferably a compound having one or more OH groups selected from the ultraviolet light absorbents represented by formulas (1), (2), and (3). Such an ultraviolet light absorbent can enhance the adhesion of the primer layer **23** to the electrically conductive plate **123a** of the laminated substrate **12**.

**[0091]** The thickness of the primer layer **23** is preferably 5  $\mu\text{m}$  to 200  $\mu\text{m}$  and more preferably 10  $\mu\text{m}$  to 100  $\mu\text{m}$ . The primer layer with such a thickness can absorb ultraviolet light emitted from the SiC semiconductor element **11** effectively to prevent the deterioration of the thermosetting resin sealing layer **24** while maintaining the effect of relaxing stress.

**[0092]** The thermosetting resin sealing layer **24** contains a thermosetting resin base material, a curing agent, and an inorganic filler. The ultraviolet light absorbent may or may not be contained, and if contained, the amount is smaller than in the primer layer **23**. The thermosetting resin base material, the curing agent, and the inorganic filler in the thermosetting resin sealing layer **24** may be selected from

those of the thermosetting resin sealing layer 20 described in the first embodiment. Also, the thermosetting resin sealing layer 24 may contain optional components added to the thermosetting resin sealing layer 20 of the first embodiment. [0093] In the semiconductor device according to the third embodiment of the present invention, the primer layer contains an ultraviolet light absorbent. This structure enables the ultraviolet light absorbent to efficiently absorb ultraviolet light even in a small amount and protects the sealing resin with adhesion at interfaces maintained, without increasing the number of basic process steps. Since the layer containing an ultraviolet light absorbent (primer layer) is thinner than the layer containing an ultraviolet light absorbent of the first embodiment (thermosetting resin sealing layer), a necessary and sufficient amount of ultraviolet light absorbent can be probably provided around the SiC semiconductor element as in the second embodiment.

EXAMPLES

[0094] The present invention will now be further described in detail with reference to Examples. The present invention is, however, not limited to the following Examples.

1. Production of Semiconductor Devices

[0095] Power semiconductor modules according to the first, the second, and the third embodiments of the present invention were produced as Examples 1, 2, and 3, respectively, for evaluating the reliability of the modules. The reliability was estimated from the  $T_j$  power cycle resistance ( $T_j$  P/C resistance). Power cycling tests in which one cycle consisted of energized operation for 2 s and a pause for 9 s was conducted at 40° C. to 185° C. ( $\Delta T_j$ : 145° C.), and the number of cycles was counted until an electrical property reached an abnormal value. More specifically, the power semiconductor modules were energized from 40° C. to 185° C. For grasping the effect of ultraviolet light, the modules were examined under conditions in which the temperature of the modules was higher than usual (175° C.) and the current flowing the semiconductor element was higher. For satisfactory reliability, at least 50 kilocycles are required.

[0096] Each thermosetting resin used in the thermosetting resin sealing layer 20 of Example 1, the first sealing layer 21 and the second sealing layer 22 of Example 2, and the thermosetting resin sealing layer 24 of Example 3 was epoxy

resin. Epoxy resin ME-276 (produced by Pelnox Ltd.) was used as the epoxy resin base material. To 100 parts by mass of this base material, 121 parts by mass of MV-138 (produced by Pelnox Ltd.) was added as an acid anhydride-based curing agent. Spherical silica (produced by AGC Inc.) having an average particle size of 10  $\mu\text{m}$  was added as the inorganic filler in a proportion of 270 parts by mass relative to 100 parts by mass in total of the epoxy resin base material and the curing agent. In the sealing layer acting as the ultraviolet light absorption layer in Examples 1 and 2, the ultraviolet light absorbent presented in Table 1 was added in the proportion presented in the Table 1 to the matrix resin. Each content, or proportion, is represented in parts by mass relative to 100 parts by mass of the matrix resin.

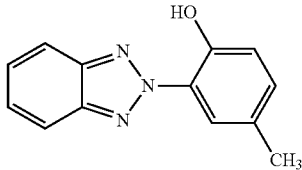
[0097] In Examples 1 and 2, the primer layer was formed of HIMAL HL 1200F (Showa Denko Materials Co., Ltd.) mainly containing polyamide over the laminated substrate, the semiconductor element, the lead frame, the wire, and the case. In Examples 1 and 2, no ultraviolet light absorbent was added into HIMAL HL1200F. In contrast, in Example 3, the ultraviolet light absorbent was added with the content presented in Table 1 into the matrix resin made of HIMAL HL1200F. The thickness of the primer layer is presented in Table 1.

[0098] In Comparative Example 1, a power semiconductor module was produced in the same manner as in Example 1, except that the thermosetting resin sealing layer contained no ultraviolet light absorbent and no primer layer was provided. In Comparative Example 2, a power semiconductor module was prepared in the same manner as in Example 3, except that the primer layer contained no ultraviolet light absorbent. Table 1 presents the compound as the ultraviolet light absorbent, the thickness ( $\mu\text{m}$ ) of the ultraviolet light absorption layer, and the ultraviolet light absorbent content (parts by mass), and the measurement results of P/C resistance (kilocycles). The units of physical quantities in Tables 2 to 7 below are the same as those in Table 1. The presence of an ultraviolet light absorption layer increased the reliability 1.5 times to twice compared to known structures with no ultraviolet light absorption layer. Comparison among Examples 1 to 3 suggests that the reliability of the modules according to the second and third embodiments was slightly higher than that of the first embodiment. This is probably because of a uniform distribution of the ultraviolet light absorbent around the SiC semiconductor element, which emits ultraviolet light.

TABLE 1

Examples and Comparative Examples					
	Example 1	Example 2	Example 3	Comparative Example 1	Comparative Example 2
Structure	First Embodiment	Second Embodiment	Third Embodiment	None	None
Primer layer	HIMAL layer provided	HIMAL layer provided	HIMAL layer provided	None	HIMAL layer provided
Thickness	10 $\mu\text{m}$	10 $\mu\text{m}$	10 $\mu\text{m}$		10 $\mu\text{m}$
Ultraviolet light absorption layer	Entire thermosetting resin sealing layer	First sealing layer	Primer layer	None	None
Ultraviolet light absorption layer thickness	5000	200	10	None	None
Ultraviolet light absorbent	2-(2-Hydroxy-5-methylphenyl)benzotriazole	←	←	None	None
CAS Number	2440-22-4	←	←	None	None

TABLE 1-continued

Examples and Comparative Examples					
	Example 1	Example 2	Example 3	Comparative Example 1	Comparative Example 2
Structural formula		←	←	None	None
Content	1	←	←	None	None
P/C Resistance	75	100	100	50	50

[0099] Next, power semiconductor modules of Examples 4 to 11 were prepared in the same manner as in Example 2, except that the thickness and the ultraviolet light absorbent content of the first sealing layer (ultraviolet light absorption layer) according to the second embodiment in Example 2 were varied. The primer layer was formed with the same composition and the same thickness as in Example 2. The results are shown in Table 2. The power semiconductor modules having a first sealing layer with a thickness of 100  $\mu\text{m}$  to 1000  $\mu\text{m}$  exhibited satisfactory results. Also, the modules with ultraviolet light absorbent content of 0.1 parts

to 5 parts by mass exhibited twice higher reliability than the cases not provided with any ultraviolet light absorption layer. When the ultraviolet light absorbent content was increased to 7 parts by mass, the reliability tended to be slightly degraded compared to the cases of lower ultraviolet light absorbent content. The modules with a higher ultraviolet light absorbent content and a larger ultraviolet light absorption layer thickness exhibited slightly higher reliability than the known power semiconductor modules but slightly lower reliability than the case that exhibited the highest reliability.

TABLE 2

Example	2	4	5	6	7	8	9	10	11
Structure	Second Embodiment	←	←	←	←	←	←	←	←
First sealing layer thickness	200	100	500	1000	100	100	750	100	750
Content	1	←	←	←	0.1	5	5	7	7
P/C Resistance	100	100	100	100	90	100	80	60	60

[0100] Next, power semiconductor modules of Examples 12 to 19 were prepared in the same manner as in Example 3, except that the thickness and the ultraviolet light absorbent content of the primer layer containing an ultraviolet light absorbent (ultraviolet light absorption layer) according to the third embodiment in Example 3 were varied. The results are shown in Table 3. The primer layer thicknesses presented in Table 3 are represented in  $\mu\text{m}$ . The results suggest that Table 3 shows that the primer layer with an ultraviolet light absorbent content of 0.1 parts by mass or more was effective regardless of whether the primer layer thickness is large or small, and notably, an ultraviolet light absorbent content of 1 part to 5 parts by mass led to high reliability.

TABLE 3

Example	3	12	13	14	15	16	17	18	19
Structure	Third Embodiment	←	←	←	←	←	←	←	←
Primer layer thickness	10	5	10	100	5	5	200	5	200
Content	1	←	←	←	0.1	5	5	7	7
P/C Resistance	100	100	100	100	80	100	100	60	60

[0101] Next, power semiconductor modules of Examples 20 to 27 were prepared in the same manner as in Example 2, except that varying benzotriazole derivatives were used as the ultraviolet light absorbent to estimate the difference in the effect (of preventing deterioration by ultraviolet light to increase the reliability) among the ultraviolet light absorbents. The thickness of the first sealing layer (ultraviolet light absorption layer) and the thickness and composition of the primer layer were the same as those in Example 2. The

results are shown in Tables 4 and 5. The results suggest that the ultraviolet light absorbents represented by formula (1) containing an alkyl group as R<sub>2</sub> or R<sub>4</sub> were notably good. The butyl group, which is bulky, is more preferable than the methyl group, which has a low molecular weight, as R<sub>2</sub> or R<sub>4</sub>. More preferably, the number of butyl groups is greater. In other words, it is preferable for R<sub>2</sub> and R<sub>4</sub> to include bulky functional groups. Also, the results suggest that R<sub>3</sub> includes an octoxy group, which is an alkoxy group.

TABLE 4

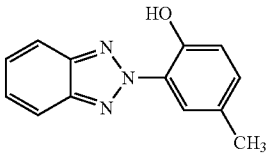
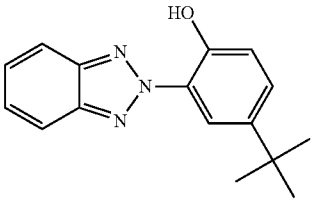
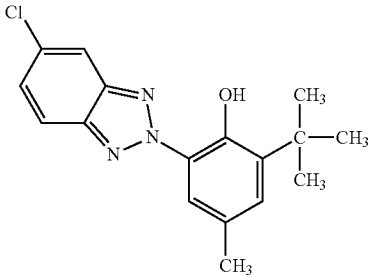
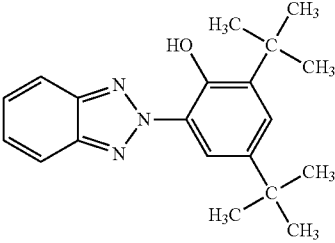
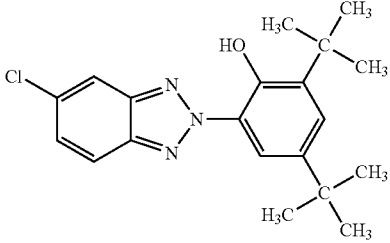
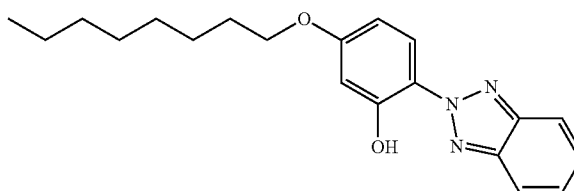
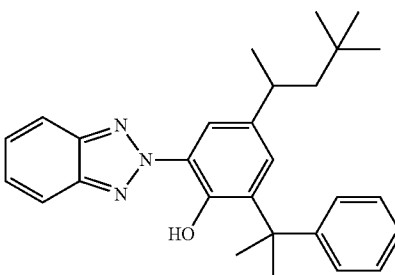
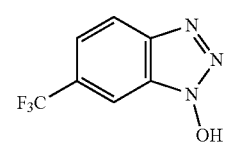
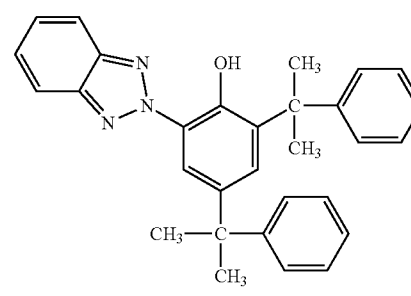
	Example		
	2	20	21
Structure	Second Embodiment	←	←
Ultraviolet light absorbent	2-(2-Hydroxy-5-methylphenyl)benzotriazole	2-(5-tert-Butyl-2-hydroxyphenyl)benzotriazole	2-(2-Hydroxy-3-tert-butyl-5-methylphenyl)-5-chlorobenzotriazole
CAS Number	2440-22-4	3147-76-0	3896-11-5
Structural formula			
First sealing layer thickness	200	←	←
Content	1	←	←
P/C Resistance	100	110	110
		Example	
		22	23
Structure	←	←	←
Ultraviolet light absorbent	←	2-(2-Hydroxy-3,5-di-tert-butylphenyl)benzotriazole	2-(2-Hydroxy-3,5-di-tert-butylphenyl)-5-chloro-2H-benzotriazole
CAS Number	←	3846-71-7	3864-99-1
Structural formula	←		
First sealing layer thickness	←	←	←
Content	←	←	←
P/C Resistance	←	120	120

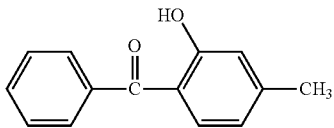
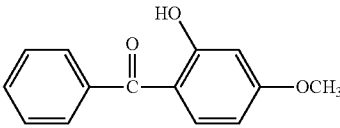
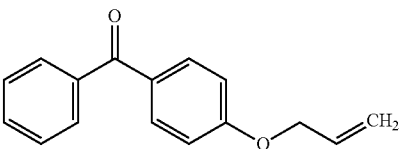
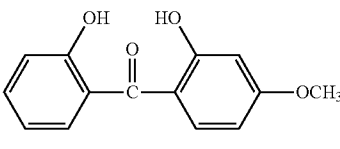
TABLE 5

	Example	
	24	25
Structure	Second Embodiment	←
Ultraviolet light absorbent	2-(2-Hydroxy-4-octoxyphenyl)benzotriazole	2-(2H-benzotriazole-2-yl)-6-(1-methyl-1-phenylethyl)-4-(1,1,3,3-tetramethylbutyl)phenol
CAS Number	3147-77-1	73936-91-1
Structural formula		
First sealing layer thickness	200	←
Content	1	←
P/C Resistance	130	130
	Example	
	26	27
Structure	←	←
Ultraviolet light absorbent	1-Hydroxy-6-(trifluoromethyl)benzotriazole	2-[2-hydroxy-3,5-bis(α,α-dimethylbenzyl)phenyl]-2H-benzotriazole
CAS Number	26198-21-0	70321-86-7
Structural formula		
First sealing layer thickness	←	←
Content	←	←
P/C Resistance	80	130

[0102] Next, power semiconductor modules of Examples 28 to 31 were prepared in the same manner as in Example 2, except that varying benzophenone derivatives were used as the ultraviolet light absorbent. The thickness of the first sealing layer (ultraviolet light absorption layer) and the thickness and composition of the primer layer were the same

as those in Example 2. The results are shown in Table 6. The results suggest that the ultraviolet light absorbents represented by formula (2) containing an alkyl group or an alkoxy group as  $R_{13}$  or  $R_{14}$  increased the P/C resistance, and notably, containing an alkoxy group was advantageous. Also, the results suggest that the methoxy group and the allyloxy group are preferable as the alkoxy group.

TABLE 6

	Example	
	28	29
Structure	Second Embodiment	←
Ultraviolet light absorbent	2-Hydroxy-4-methylbenzophenone	2-Hydroxy-4-methoxybenzophenone
CAS Number	3098-18-8	131-57-7
Structural formula		
First sealing layer thickness	200	←
Content	1	←
P/C Resistance	100	110
	Example	
	30	31
Structure	←	←
Ultraviolet light absorbent	4-(Allyloxy)benzophenone	2,2'-Dihydroxy-4-methoxybenzophenone
CAS Number	42403-77-0	131-53-3
Structural formula		
First sealing layer thickness	←	←
Content	←	←
P/C Resistance	120	110

[0103] Next, power semiconductor modules of Examples 32 to 38 were prepared in the same manner as in Example 2, except that varying triazine derivatives were used as the ultraviolet light absorbent. The thickness of the first sealing layer (ultraviolet light absorption layer) and the thickness and composition of the primer layer were the same as those

in Example 2. The results are shown in Tables 7 and 8. The results reveal that triazine derivatives containing an alkyl group in any of  $R_{21}$ ,  $R_{25}$ , and  $R_{28}$  of formula (3) are preferable, triazine derivatives containing an alkoxy group in any of  $R_{21}$ ,  $R_{25}$ , and  $R_{28}$  are more preferable, and that the number of alkoxy groups is preferably greater.

TABLE 7

	Example	
	32	33
Structure	Second Embodiment	←
Ultraviolet light absorbent	2-(2,4-Dihydroxyphenyl)-4,6-diphenyl-1,3,5-triazine	2,4,6-tris(2,4-dihydroxyphenyl)-1,3,5-triazine
CAS Number	38369-95-8	2125-23-7
Absorption wavelength nm	380	337



TABLE 7-continued

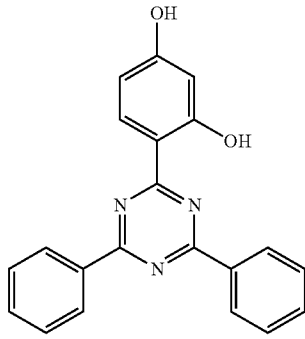
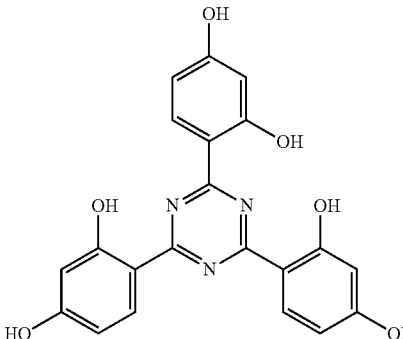
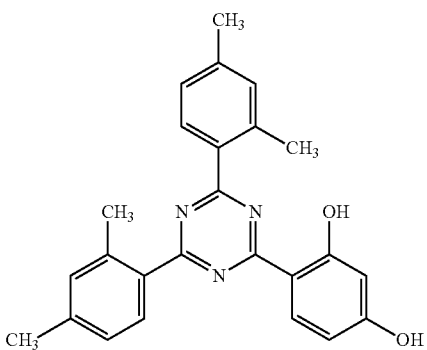
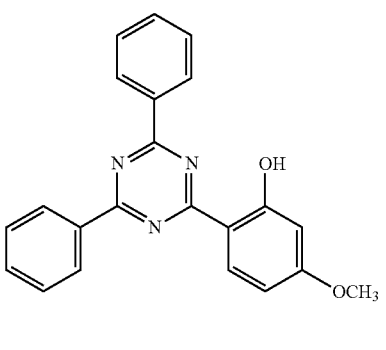
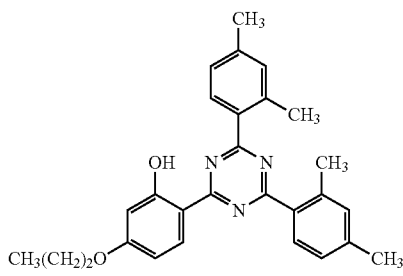
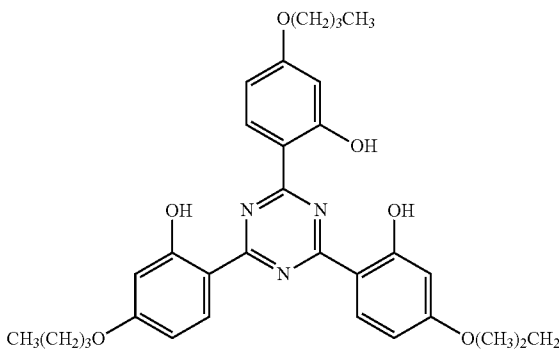
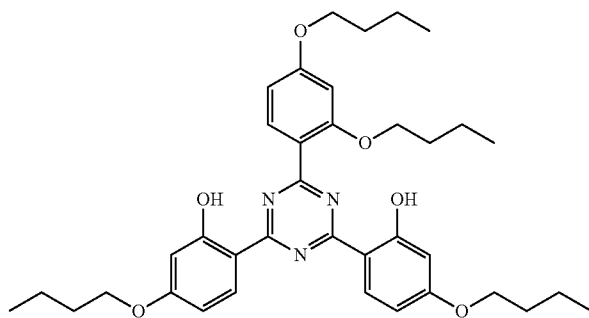
Structural formula		
		
First sealing layer thickness	200	←
Content	1	←
P/C Resistance	100	90
Example		
	34	35
Structure		
Ultraviolet light absorbent	2-(2,4-dihydroxyphenyl)-4,6-bis(2,4-dimethylphenyl)-1,3,5-triazine	← 2-(2-hydroxy-4-methoxyphenyl)-4,6-diphenyl-1,3,5-triazine
CAS Number	1668-53-7	106556-36-9
Absorption wavelength nm	380	340
Structural formula		
		
First sealing layer thickness	←	←
Content	←	←
P/C Resistance	110	110

TABLE 8

Example		
	36	37
Structure		
Ultraviolet light absorbent	Second Embodiment 2,4-Bis(2,4-dimethylphenyl)-6-(2-hydroxy-4-n-octyloxyphenyl)-1,3,5-triazine	← 2,4,6-Tris(4-butoxy-2-hydroxyphenyl)-1,3,5-triazine
CAS Number	2725-22-6	3135494
Absorption wavelength nm	380	380

TABLE 8-continued

Structural formula		
First sealing layer thickness	200	←
Content	1	←
P/C Resistance	120	130
		Example 38
Structure	Ultraviolet light absorbent	←
CAS Number		2,4-Bis(2-hydroxy-4-butoxyphenyl)-6-(2,4-bis-butoxyphenyl)-1,3,5-triazine
Absorption wavelength nm		208343-47-9 380
Structural formula		
First sealing layer thickness		←
Content		←
P/C Resistance		130

[0104] According to the present invention, a highly reliable semiconductor device with increased power cycle resistance can be provided by providing an ultraviolet light absorption layer in the semiconductor device as in the disclosed embodiments.

## REFERENCE SIGNS LIST

- [0105] 11 SiC semiconductor element
- [0106] 12 laminated substrate
- [0107] 121 electrically conductive plate
- [0108] 122 insulation substrate
- [0109] 123a, 123b electrically conductive plate
- [0110] 13 heat sink
- [0111] 14 aluminum wire
- [0112] 15 external terminal
- [0113] 16 case
- [0114] 17 bonding layer
- [0115] 18 lead frame
- [0116] thermosetting resin sealing layer (ultraviolet light absorption layer)

[0117] 21 first sealing layer (ultraviolet light absorption layer)

[0118] 22 second sealing layer

[0119] 23 primer layer (ultraviolet light absorption layer)

[0120] 24 thermosetting resin sealing layer

What is claimed is:

1. A semiconductor device comprising:

a SiC semiconductor element mounted on a multilayer substrate and an electrically conductive connection member; and

a sealant containing an ultraviolet light absorbent, which seals the SiC semiconductor element and the electrically conductive connection member.

2. The semiconductor device according to claim 1, wherein the sealant comprises a thermosetting resin sealing layer comprising a thermosetting resin base material, a curing agent, an inorganic filler, and the ultraviolet light absorbent.

3. The semiconductor device according to claim 2, wherein the thermosetting resin sealing layer consists of a layer comprising the thermosetting resin base material, the curing agent, the inorganic filler, and the ultraviolet light absorbent.

4. The semiconductor device according to claim 2, wherein the thermosetting resin sealing layer comprises: a first sealing layer covering surroundings of the SiC semiconductor element; and a second sealing layer covering the first sealing layer, and

wherein the first sealing layer comprises the thermosetting resin base material, the curing agent, the inorganic filler, and the ultraviolet light absorbent.

5. The semiconductor device according to claim 2, wherein the sealant comprises: a primer layer covering the SiC semiconductor element and the electrically conductive connection member; and the thermosetting resin sealing layer covering the primer layer, and wherein the primer layer comprises an ultraviolet light absorbent.

6. The semiconductor device according to claim 1, wherein the ultraviolet light absorbent is one, two or more selected from the group consisting of benzotriazole derivatives, benzophenone derivatives, and triazine derivatives.

7. The semiconductor device according to claim 2, wherein the thermosetting resin base material comprises an epoxy resin base material.

8. The semiconductor device according to claim 5, wherein the primer layer comprises one or more resins selected from the group consisting of polyamide resins, polyimide resins, and polyamide-imide resins.

9. The semiconductor device according to claim 2, wherein the ultraviolet light absorbent is one, two or more selected from the group consisting of benzotriazole derivatives, benzophenone derivatives, and triazine derivatives.

10. The semiconductor device according to claim 3, wherein the ultraviolet light absorbent is one, two or more selected from the group consisting of benzotriazole derivatives, benzophenone derivatives, and triazine derivatives.

11. The semiconductor device according to claim 4, wherein the ultraviolet light absorbent is one, two or more selected from the group consisting of benzotriazole derivatives, benzophenone derivatives, and triazine derivatives.

12. The semiconductor device according to claim 5, wherein the ultraviolet light absorbent is one, two or more selected from the group consisting of benzotriazole derivatives, benzophenone derivatives, and triazine derivatives.

\* \* \* \* \*