

FAR-INFRARED EMITTING CATHODE RAY TUBE ✓

The present invention relates to a far-infrared emitting cathode ray tube, and more particularly to a far-infrared emitting cathode ray tube which includes far-infrared emitting material therein and thus emits far-infrared rays available for a human body.

While continuing the development toward information-oriented society, people are suffering from a pathological phenomenon, i.e., so-called visual display terminal (VDT) syndrome, caused by electromagnetic wave and working environment. The electromagnetic wave is weak in its intensity, but produces anxiety in users as its side effect appears as such electromagnetic-wave troubles as bloodshot eyes, amblyopia, headache, muscular pain, chronic fatigue, languor, skin roughness, etc. The most favorable method is to shield such electromagnetic waves. However, many difficulties actually occur including high cost and the like, which inhibits the adoption of this method. Accordingly, a cathode ray tube has been recognized as harmful to human body except in its function of transmitting information.

A far-infrared ray is a form of electromagnetic wave located at a wavelength range from $5\mu\text{m}$ to $1,000\mu\text{m}$ (the standard wavelength region of far-infrared rays is not critically defined. This range is selected to exclude near-infrared rays in this invention), and its various biological effects known until now comprise activation of hydrogen oxide molecule, maturation, freshness preservation, growth acceleration, energy saving, sterilization, deodorization, etc. Also, in actual life, the far-infrared ray is employed in

saunas, refrigerators, microwave ovens, and the like. The effect of the far-infrared ray cannot be represented by numerical and logical results obtained by means of scientific measuring instruments, but can be indirectly perceived. More specifically, a far-infrared lamp light promotes the growth in plants, a far-infrared sauna lets a man sweat even under low temperature, a far-infrared refrigerator keeps meat and fish fresh for a long time, a far-infrared emitting article makes cigarette and liquor taste pleasant once exposed thereto, far-infrared ray shortens the painting process when applying it, etc.

It is explained that the influence of the far-infrared ray on the human body is classified into two effects, i.e., thermal effect and non-thermal effect. The thermal effect indicates that heat energy absorbed by the skin affects peripheral tissue, deep tissue of a living organism, or the overall living organism by means of the circulation of blood flowing through a peripheral blood vessel. The non-thermal effect indicates that light quantum corresponding to specific wavelength of the far-infrared ray stimulates the receptors of the cell membrane outside or inside skin, and a far-infrared signal of the specific wavelength is transmitted into cells, thereby activating the cells. Consequently, the far-infrared ray functions as heat energy supply source in the thermal effect, and light quantum supply source in the non-thermal effect. Through these effects, such as direct thermal effect as an infrared ray, indirect effect by activating water constituting a part of the body, and non-thermal effect for stimulating nerve receptors for sensing variously warmth, coolness, pain which are located in the skin at a depth around $100\mu\text{m}$, the far-infrared ray provides acceleration of the blood circulation, speedy release of waste

matter, etc.

The effect of the far-infrared ray with respect to water (which constitutes most of the human body) will be described in detail below. FIG. I of the accompanying drawings is a graphical representation showing the transmittance of water in accordance with wavelength. Here, it can be noted that water absorbs light whose wavelength is near $3\mu\text{m}$ or longer than $6\mu\text{m}$. In a water molecule (H_2O , H-O-H), stretching vibration of the O-H bonding between oxygen and hydrogen occurs between $2.5\text{-}3.5\mu\text{m}$, and bending vibration occurs between $10\text{-}14\mu\text{m}$. Thus, when light of these wavelengths is observed, the absorption of the light accelerates the vibration of the water. In other words, by externally supplying light of these wavelengths, water molecules are activated so as to be arranged in the ideal bonding structure.

The wavelength of a human body can be calculated by taking body temperature, in accordance with the relation between absolute temperature and wavelength which is called as a Wien formula, is defined as: $\lambda = 2897/T$ (where T denotes an absolute temperature and λ denotes a wavelength in micrometers). Namely, if 309.5 (i.e. $273 + 36.5$) is substituted for T , the wavelength becomes $9.36\mu\text{m}$ which is within the far-infrared range.

FIGs. 2A and 2B of the accompanying drawings are graphical representations respectively showing a spectral transmittance (2A) and a spectral reflectance (2B) of skin according to each wavelength. The infrared wavelength emitted from a human's skin is within the range of $3\text{-}50\mu\text{m}$, and especially, the wavelength within the range of $8\text{-}14\mu\text{m}$ which occupies approximately 46% of the total emitted energy. The energy emitted to the skin

is transmitted, reflected or absorbed, and therefore, the amount of the energy absorbed by the skin can be calculated considering FIGs. 2A and 2B. For example, almost all of the energy within 8 to $14\mu\text{m}$ region of which transmittance and reflectance are low, is absorbed by the skin. Thus, upon supplying energy in this range, a living body mostly consisting of water is easily activated, which is displayed by the early blossom of flowers, a shortened period and early maturation of a hatching egg, prolonged life of cut flowers, etc.

Several products using the far-infrared effect are practically presented, e.g., a far-infrared refrigerator disclosed in Japanese Laid-Open Publication Nos. Hei 2-57883 and Hei 2-309169, a far-infrared emitting bathtub disclosed in Japanese Laid-Open Publication No. Hei 2-164365, etc.

Metal oxides such as alumino-silicates ($\text{Al}_2\text{O}_3\text{-SiO}_2$), cordierites ($\text{MgO-Al}_2\text{O}_3\text{-SiO}_2$), zircons ($\text{ZrO}_2\text{-SiO}_2$), carbons, ferric oxide (Fe_2O_3), manganese dioxide (MnO_2), cupric oxide (CuO), tricobalt tetraoxide (Co_3O_4), nickel monoxide (NiO), chromic oxide (Cr_2O_3), lithium oxide (Li_2O), zinc oxide (ZnO), bismuth oxide (Bi_2O_3), barium oxide (BaO), titanium oxide (TiO_2), boron oxide (B_2O_3), sodium oxide (Na_2O), potassium oxide (K_2O), phosphorus pentoxide (P_2O_5), molybdenum sesquioxide (Mo_2O_3), and calcium oxide (CaO) are examples of far-infrared emitting materials.

It is an object of the present invention to provide a far-infrared emitting cathode ray tube which emits far-infrared ray available for the human body to make use of information equipment in favorable surroundings even though electromagnetic waves, pointed to as one cause of the VDT syndrome,

cannot be completely shielded.

According to one aspect of the present invention, there is provided a far-infrared emitting cathode ray tube, wherein far-infrared emitting material is included in at least one of a phosphor layer, a black matrix, an outer graphite layer, and a shadow mask, thereby emitting far-infrared rays.

In embodiments of the far-infrared emitting cathode ray tube, the far-infrared emitting material is included in or coated on at least one of the phosphor layer, the black matrix and the outer graphite layer, e.g. inside the phosphor layer, inside or on an upper portion of the black matrix layer, inside or on an upper portion of the outer graphite layer, and the outer surface of the graphite layer on the shadow mask.

Particularly, embodiments of the present invention provide: a cathode ray tube wherein a far-infrared emitting material having not more than $5\mu\text{m}$ average diameter is formed on the upper portion of the black matrix to a thickness of $0.2\text{-}100\mu\text{m}$; a cathode ray tube wherein the far-infrared emitting material having not more than $5\mu\text{m}$ average diameter is included within the black matrix layer as much as 1-100wt%, based upon the total amount of the mixture with the graphite; a cathode ray tube wherein the far-infrared emitting material is not more than 50wt% with respect to the amount of the phosphor in the phosphor layer; a cathode ray tube wherein the average diameter of the far-infrared emitting material having a diameter of $0.05\text{-}2\mu\text{m}$ is coated to a thickness of about $0.1\text{-}10\mu\text{m}$ on the outer surface of the shadow mask toward the phosphor layer; and a cathode ray tube wherein the far-infrared emitting

material is included in the outer graphite layer which is formed on the outer surface of the funnel of the cathode ray tube.

The far-infrared emitting material is selectively included in each portion of the cathode ray tube, or included in all portions as circumstances require.

As for the far-infrared emitting materials, at least one selected from the group consisting of aluminium oxide (Al_2O_3), silicon dioxide (SiO_2), magnesium oxide (MgO), zirconium oxide (ZrO_2), carbons, ferric oxide (Fe_2O_3), manganese dioxide (MnO_2 , MnO), cupric oxide (CuO), tricobalt tetraoxide (Co_3O_4), nickel monoxide (NiO), chromic oxide (Cr_2O_3), lithium oxide (Li_2O), zinc oxide (ZnO), bismuth oxide (Bi_2O_3), barium oxide (BaO), titanium oxide (TiO_2), boron oxide (B_2O_3), sodium oxide (Na_2O), potassium oxide (K_2O), phosphorus pentoxide (P_2O_5), molybdenum sesquioxide (Mo_2O_3), calcium oxide (CaO) and a complex thereof can be employed.

Embodiments of the present invention will now be described, by way of example, with reference to the accompanying drawings, in which:

FIG. 1 is a graphical representation illustrating the transmittance of water

versus wavelength;

FIGs. 2A and 2B are graphical representations respectively illustrating the spectral transmittance and spectral reflectance of the skin versus wavelength;

FIGs. 3A to 3I illustrate a process for manufacturing a screen according to Example 1 of the present invention;

FIGs. 4A and 4B illustrate a spectral analysis (4A) of far-infrared emitting material components used in Example 1, and spectral emittance (4B) thereof versus wavelength;

FIG. 5 is a sectional view showing a portion of the screen manufactured by example 1 of the present invention;

FIGs. 6A and 6B illustrate a spectral analysis (6A) of far-infrared emitting material components used in Example 2, and spectral emittance (6B) thereof versus wavelength;

FIG. 7 is a graphical representation illustrating luminance of a conventional screen (a) and screens according to Examples 1 and 2 (b and c), respectively;

FIG. 8 is a photograph of a screen manufactured by Example 3 of the present invention;

FIG. 9 is a photograph of roses used for demonstrating the far-infrared emitting characteristic of a far-infrared emitting cathode ray tube manufactured by Example 3 of the present invention;

FIGs. 10A and 10B illustrate a spectral analysis (10A) of far-infrared emitting material components used in Example 4, and spectral emittance (10B)

thereof versus wavelength;

FIG. 11 is a photograph of a screen manufactured by Example 4 of the present invention;

FIGs. 12, 13 and 14 are graphs illustrating the luminous spectrum of green, blue and red phosphor screens manufactured by Example 5 of the present invention, respectively;

FIGs. 15A and 15B are photographs of roses, carnations and onions used for demonstrating the far-infrared emitting characteristic of a far-infrared emitting cathode ray tube manufactured by Example 5 of the present invention;

FIG. 16 is a view for illustrating a method for manufacturing a shadow mask according to Example 7 of the present invention;

FIG. 17 is a view for illustrating a method for manufacturing a shadow mask according to Example 8 of the present invention;

FIG. 18 shows a cross-sectional view of a portion of a funnel of a cathode ray tube manufactured according to Example 9 of the present invention;

FIG. 19 illustrates the experimental method used for demonstrating the effect of emitting far-infrared rays in the far-infrared emitting cathode ray tube manufactured by Example 9 of the present invention, and

FIG. 20 shows a cross-sectional view of a portion of a funnel of a cathode ray tube manufactured according to Example 10 of the present invention.

Even though any material capable of emitting far-infrared rays may be

adopted in the following examples, the most preferred methods will be given as an example hereinbelow since the additive amount or application method of the materials varies to suit the characteristics and limitations of the applied articles.

According to one embodiment of the present invention, a far-infrared emitting cathode ray tube has a screen which includes a black matrix layer covering the inner surface of a panel in a predetermined pattern, a phosphor layer arranged between stripes of the black matrix layer and emitting various colors by filtering the electromagnetic wave, and an aluminum layer for enhancing the overall luminance of the panel and strengthening the electrical potential of the phosphor screen, wherein a far-infrared emitting material layer is installed on the black matrix layer.

The above-described far-infrared emitting screen is manufactured in accordance with the following steps. The black matrix layer of a predetermined pattern is first formed on the inner surface of the panel. A photoresist layer is formed by coating, exposing and developing a photoresist on the black matrix layer, and then a far-infrared emitting material layer is formed by coating far-infrared emitting material on the photoresist layer. Successively, after coating, exposing, and developing a photoresist for forming a phosphor, phosphor slurry is coated, exposed, and developed, thereby forming a phosphor layer between the stripes of the black matrix layer.

The far-infrared emitting material is at least one of SiO_2 , Al_2O_3 , MgO , Na_2O , P_2O_5 , K_2O , ZrO_2 , CaO and TiO_2 ; the preferable average particle diameter of these is not more than $5\mu\text{m}$.

The thickness of the far-infrared emitting material layer is 0.2-100 μ m, and preferably 0.5-50 μ m.

Hereinbelow, the above-stated method for manufacturing the far-infrared emitting screen will be described in detail with reference to FIGs. 3A through 3I.

EXAMPLE 1

A panel 1 is cleansed using hydrofluoric acid and distilled water according to a conventional method prior to drying it. Then, a photoresist composed of polyvinyl alcohol, sodium bichromate, acryl emulsion, etc. is coated on the inner surface of panel 1, and thus a photoresist pattern is formed by drying, exposing and developing. After covering and drying a graphite on the photoresist pattern, a black matrix 2 pattern is formed by separating this photoresist pattern, using an etching solution (FIG. 3A).

After covering a negative-type photoresist 3 on black matrix 2 pattern (FIG. 3B), the photoresist on the panel is exposed to light and developed through a shadow mask, so that the photoresist layer on panel 1 is removed and an unexposed resist layer is left on black matrix 2 (FIG. 3C). After this, a far-infrared emitting material 4 composed of at least one of silicon dioxide (SiO_2), aluminium oxide (Al_2O_3), magnesium oxide (MgO), sodium oxide (Na_2O), phosphorus pentoxide (P_2O_5), potassium oxide (K_2O), zirconium dioxide (ZrO_2), calcium oxide (CaO) and titanium oxide (TiO_2) and having not more than a 5 μ m average diameter, is sprayed on the upper portion of black matrix 2 and the resist pattern, using a spray gun whose nozzle has a diameter of 0.4mm (FIG. 3D). Therefore, a far-infrared

emitting material layer 4 is formed on the upper portion of resist layer 3 on black matrix 2 (FIG. 3E). Then, via the sequence of an ordinary slurry process, i.e. photoresist coating (FIG. 3F), exposure and development (FIG. 3G), phosphor slurry coating (FIG. 3H), and an exposure and development, a first phosphor layer 5 is formed (FIG. 3I). Finally, after forming a second and third phosphor layer via the same common slurry process, a screen for a cathode ray tube having the black matrix, far-infrared emitting material layer, phosphor layer, and the aluminum layer is formed through the sequential steps of filming composition coating, aluminum deposition, and baking.

The thickness of the far-infrared emitting layer is between $0.2\mu\text{m}$ to $100\mu\text{m}$, and preferably is the range of $0.5\mu\text{m}$ to $50\mu\text{m}$. If the layer's thickness is greater than $100\mu\text{m}$, manufacture of the phosphor layer is difficult and the material powder separates easily. On the other hand, if the thickness is below $0.2\mu\text{m}$, the effects of far-infrared ray generation and luminance increase are trivial.

FIGs. 4A and B are a spectral analysis (4A) of far-infrared emitting material components used in the first example, and a graph (4B) obtained by measuring spectral emittance thereof versus wavelength. In FIG. 4A, the peak of each far-infrared emitting material used can be seen. The spectral emission measuring graph shown in FIG. 4B is obtained by employing black body (at 40°C) as a reference, measuring with an FT-IR detector of Bruker Co., and detecting by DTGS. From FIG. 4B, it can be noted that the emission of far-infrared light is intense at wavelengths shorter than $25\mu\text{m}$.

FIG. 5 is a sectional view which illustrates a portion of the screen

manufactured by the above example of the present invention. As shown in the drawing, graphite layer 2 and phosphor layer 5 are formed on the inner surface of a panel 1, and far-infrared emitting material layer 4 whose color is off-white is formed on the graphite layer. Also, an aluminum layer 6 is formed above phosphor layer 5 and far-infrared emitting material layer 4, spaced apart from them by a predetermined distance. Here, since far-infrared emitting material layer 4 is white or off-white, the far-infrared emitting material layer again reflects the light which has been reflected toward the graphite among emitted from the phosphor layer and thus absorbed thereinto, so that luminance is enhanced.

EXAMPLE 2

The same method as example 1 is employed except that a material consisting of SiO_2 , Al_2O_3 and MgO is used as the far-infrared emitting material to manufacture a screen by forming a far-infrared emitting material layer inside a panel and over a black matrix layer of a cathode ray tube.

FIGs. 6 are a spectral analysis table (6A) of far-infrared emitting material components used, and a graph (6B) obtained by measuring spectral emittance thereof versus wavelength. Here, each peak of each far-infrared emitting component can be seen, and it can be noted that the emission of far-infrared light is intense at wavelengths shorter than $25\mu\text{m}$.

As described above, when the cathode ray tube is manufactured with the far-infrared emitting material layer over the black matrix layer, the cathode ray tube emits far-infrared rays to give a user an activation effect provided

by the far-infrared rays.

The inventor of this invention has confirmed that the enhanced luminance effect can also be obtained by forming the far-infrared emitting material layer over the black matrix layer in accordance with the principle shown in FIG. 5. The luminance of a conventional screen (without a far-infrared emitting layer), and that of screens according to examples 1 and 2 (with the far-infrared emitting layer) are illustrated in <Table 1>. Here, additional data is included for the screens of examples 1 and 2.

<Table 1>

	conventional CRT	example 1	example 2
emitting layer material	N/A	SiO ₂ , Al ₂ O ₃ , MgO, Na ₂ O, P ₂ O ₅ , K ₂ O, CaO, TiO ₂	SiO ₂ , Al ₂ O ₃ , MgO
color of emitting layer	N/A	white	light gray
thickness of emitting layer (μm)	N/A	20	20
luminance (%)	100	110	105

FIG. 7 is a graphical representation illustrating luminance of the conventional screen (a) and the screens according to examples 1 and 2 (b and c), respectively. Here, the luminance of the screen according to an embodiment of the present invention is enhanced by approximately 5-10% as compared with that of the conventional screen.

As described above, by forming the far-infrared emitting layer over the

black matrix, far-infrared rays are emitted from the cathode ray tube and, at the same time, luminance can be enhanced.

Another embodiment of the present invention, will now be described in which the far-infrared emitting material is contained in the black matrix layer.

The far-infrared emitting material is composed of at least one of SiO_2 , Al_2O_3 , MgO , Na_2O , P_2O_5 , K_2O , ZrO_2 , CaO and TiO_2 , preferably having not more than a $5\mu\text{m}$ average diameter. Also, the addition or additive amount of graphite can be determined in accordance with the overall color, and the addition of conductivity and the light-absorbing characteristic.

In order to introduce the far-infrared emitting material in the black matrix layer, the far-infrared emitting material is mixed with graphite solution generally used for forming a black matrix, by being subjected to a ball-mill, if necessary, to become consistent. The additive amount of the far-infrared emitting material is 1-100wt%, preferably 10-100wt%, when the total amount of the mixture is used as a reference. It is observed that, if the mixed amount of the far-infrared emitting material is increased, the far-infrared emission is increased. Also, the far-infrared emitting effect can be obtained when the content of the far-infrared emitting material is less than 10%, though it is slight.

The prepared mixture having the far-infrared emitting characteristic is applied on the inner surface of the screen as dots or stripes according to the same steps as the conventional black matrix manufacturing steps. That is, the panel is cleansed using distilled water, hydrofluoric acid, etc. Then, a photoresist is coated

and dried, thereby forming a photoresist pattern via an exposure and development process. Thereafter, a black matrix pattern is formed by coating and etching a black matrix mixture containing far-infrared emitting material. Finally, the screen is completed by carrying out common phosphor slurry coating, filming composition coating, aluminum deposition, and baking processes.

Upon operating the cathode ray tube, since 20% of the electron beam energy emitted from the electron gun is converted into light, and 80% becomes heat, the far-infrared rays are emitted externally, while increasing the temperature and the emission of far-infrared rays within the cathode ray tube. As expressed by the Stefan-Boltzmann formula (defined as: $W=5.67 \times 10^{-12}T^4$, where W is the total emitted energy W/cm^2 , and T is the absolute temperature K), the emitted energy amount is in proportion to 4-square of the absolute temperature, and thus the higher the applied temperature is, the higher the obtained emitting efficiency is.

Hereinbelow, a preferred embodiment of a cathode ray tube wherein the far-infrared emitting material is introduced into the black matrix, will be described in detail.

EXAMPLE 3

An applied solution composed of 2.5wt% far-infrared emitting material containing SiO_2 , Al_2O_3 , MgO , Na_2O , P_2O_3 , K_2O , and TiO_2 , having an average diameter of $3\mu m$, 2.5wt% graphite for black matrix having an average diameter of $1\mu m$, 0.2wt% sulphonic surface active agent, 0.2wt% kasil, 2wt% alumina-based dispersant, and a remainder of distilled water, is subjected to a ball-mill for 24

hours for consistent dispersion, and is formed as a black matrix pattern on the inner surface of the panel. Then, a screen is manufactured in accordance with the conventional method.

FIG. 8 is a photograph taking the external view of the screen manufactured as above, wherein black lines are a black matrix pattern including the far-infrared emitting material.

To demonstrate the far-infrared emitting effect of the cathode ray tube having the far-infrared emitting material, the following experiment was performed.

Four roses being in bloom to the same degree were put into similar flasks (A, B, C and D), respectively. Then, flask A was placed at a distance of 10cm from a conventional television operating without the far-infrared emitting material, and flasks B, C and D were also placed at distance of 10cm from a television operating with the far-infrared emitting material according to example 3 of the present invention. The above was left in place for 100 hours. Throughout, other parameters were controlled, such as, the light intensity emitted from the two kinds of televisions was kept constant, and external light was blocked.

FIG. 9 is a photograph taken after the elapse of 100 hours, showing the relative states of the roses. As shown in the photograph, the rose of flask A is already withered and dried while its petals are brittle, but those of flasks B, C and D are fresh and moist due to sufficient supply of water. It can be considered that this is because the far-infrared rays emitted from the television according to an embodiment of the present invention activates the water in the flasks B, C and D and the inherent water of the roses, so that the withering of the roses is inhibited.

EXAMPLE 4

An applied solution composed of 5wt% far-infrared emitting material containing SiO_2 , Al_2O_3 , MgO , CaO and TiO_2 , having an average diameter of $1\mu\text{m}$, 0.1wt% non-ionic surfactant, 0.1wt% graphite for black matrix having an average diameter of $1\mu\text{m}$, 0.1wt% kasil, 1wt% $\text{Al}(\text{OH})_3$ dispersant, ammonium hydroxide NH_4OH having an amount capable of controlling pH at 10-11, and a remainder of distilled water, is prepared. This solution is applied on the inner surface of the panel, thereby forming a black matrix pattern. Then, a screen is manufactured in accordance with the conventional method.

FIGs. 10A and 10B are an analysis table (10A) of far-infrared emitting material component used in the foregoing, and a graph (10B) obtained by measuring spectral emittance over a given band of wavelengths. In the figures the presence of each component and its high emission intensity in the far-infrared region can be confirmed.

In FIG. 11, a photograph taking the external view of the screen manufactured by Example 4 of the present invention. Here, since there is no graphite in the black matrix layer, this layer appears as white.

The black matrix layer of the screen manufactured according to the example 4 is composed of the far-infrared material only, and cannot function as its own. Therefore, a drawback of degrading color purity of the image exists, but there is a benefit. A high far-infrared emitting effect can be obtained, provided that the black matrix layer according to this example is adopted for a cathode ray tube wherein the above drawback is negligible.

A further embodiment of the present invention provides a cathode ray tube which comprises a phosphor layer composed of phosphors as dots or stripes for emitting white or colored light by collision of electron beams, and formed on the inner surface of a panel, wherein a far-infrared emitting material is included in the phosphor layer.

A far-infrared emitting material included in the phosphor layer is composed of SiO_2 , Al_2O_3 , MgO , Co_3O_4 , TiO_2 , CaO , K_2O , and Na_2O , as well as transition oxide metals other than these. Preferably, the average particle diameter of these is $0.5\text{-}4\mu\text{m}$, and the maximum particle diameter is $8\mu\text{m}$.

Also, in order to sustain the luminous efficiency of the phosphor, the desirable color of the far-infrared emitting material is white or off-white, and a highly purified metal oxide (not mixed with Fe, Co and Ni which decrease the luminous characteristic of the phosphor) is used as the far-infrared material.

To introduce the above-described far-infrared emitting material into the phosphor layer, the material is mixed with phosphor slurry, and this mixture is applied to the conventional manufacturing process of the phosphor layer. At this time, the additive amount of the far-infrared emitting material is not more than 50wt%, preferably 1-40wt% of the amount of the phosphor. If the additive amount of the far-infrared emitting material is increased, luminance is lowered even though there is no difference in the luminous spectrum and luminous color coordinates. Accordingly, an additive amount of more than 50wt% is not preferable.

Here, in the case of the screen for a color cathode ray tube, which has a phosphor layer obtained by forming color phosphor layers on the inside the panel

via several processes of separately prepared red, green and blue luminous phosphor slurry such as coating, drying, exposing and developing, the phosphor layer is manufactured in accordance with a conventional process by mixing the far-infrared emitting material with each color slurry. Here, since the luminous characteristic of the ZnS-based phosphor such as the green luminous phosphor like ZnS:Cu,Au,Al, ZnS:Cu,Al, (Zn,Cd)S:Cu,Al, and blue luminous phosphor like ZnS:Ag is deteriorated by impurities such as Fe, Co and Ni, the purity of the materials (especially of the far-infrared emitting material) must be carefully supervised, so as not to contain the impurities.

A color cathode ray tube with the phosphor layer including the far-infrared emitting material will be given as example. Here, a case in which far-infrared emitting material is included in all color phosphor layers will be described in detail.

EXAMPLE 5

To begin with, 85g of ZnS:Cu,Au,Al as a green phosphor, a 15g far-infrared emitting material mixture obtained by mixing SiO₂, Al₂O₃, MgO, CaO and TiO₂ in the same ratio, 200g of distilled water, 70g of polyvinyl alcohol, 4.1g of sodium bichromate, and 50ml of surface active agent are mixed to prepare the green phosphor slurry. At this time, the average particle diameter of the phosphor is 7 μ m, and that of the far-infrared emitting material is 1.5 μ m.

In the analysis table of the far-infrared emitting material illustrated in FIG. 10A, each peak of the components is depicted. Also, the graph obtained by

measuring the spectral emittance of the far-infrared emitting material shown in FIG. 10B illustrates the fact that the emission efficiency is high in the range below $25\mu\text{m}$.

By using the green phosphor slurry, a green phosphor pattern is formed according to the conventional process.

The luminous spectrum of the green phosphor layer is represented in FIG. 12, wherein the inherent luminous spectrum of the green phosphor has no difference from that of the conventional phosphor.

Next, 70g of ZnS:Ag as the blue luminous phosphor having cobalt blue ($\text{CoO} \cdot \text{Al}_2\text{O}_3$) attached thereto, a 10g far-infrared emitting material mixture composed of SiO_2 , Al_2O_3 and MgO which are mixed in the same ratio, 150g of distilled water, 80g of polyvinyl alcohol, 3.8g of sodium bichromate, and 8ml of surface active agent are all mixed to produce the blue phosphor slurry. At this time, the average particle diameter of the phosphor is $5\mu\text{m}$ and that of the far-infrared emitting material is $1\mu\text{m}$.

In the analysis table of the far-infrared emitting material illustrated in FIG. 6A, each peak of the components is depicted. Also, the graph obtained by measuring the spectral emittance of the far-infrared emitting material shown in FIG. 6B illustrates the fact that the emission efficiency is high in the range below $25\mu\text{m}$.

By using the blue phosphor slurry, a blue phosphor pattern is formed according to the conventional process.

The luminous spectrum of the blue phosphor layer having the far-infrared

emitting material is represented in FIG. 13, wherein the inherent luminous spectrum of the blue phosphor has no difference from that of the conventional phosphor.

Lastly, 90g of $Y_2O_3S:Eu$ having the red luminous phosphor having Fe_2O_3 attached thereto, a 10g far-infrared emitting material mixture composed of SiO_2 , Al_2O_3 , MgO , Na_2O , K_2O and TiO_2 mixed in the same ratio, 100g of distilled water, 50g of polyvinyl alcohol, 3.7g of sodium bichromate, and 30ml of surface active agent are all mixed to produce the red phosphor slurry. At this time, the average particle diameter of the phosphor is $6\mu m$ and that of the far-infrared emitting material is $2\mu m$.

In the analysis table of the far-infrared emitting material illustrated in FIG. 4A, each peak of the components is depicted. Also, the graph obtained by measuring the spectral emittance of the far-infrared emitting material shown in FIG. 4B illustrates the fact that the emission efficiency is high in the range below $25\mu m$.

By applying the red phosphor slurry on the inner surface of the panel, a red phosphor pattern is formed according to the conventional processes such as drying, exposing and developing.

The luminous spectrum of the red phosphor layer having the far-infrared emitting material is represented in FIG. 14, wherein the inherent luminous spectrum of the red phosphor has no difference from that of the conventional phosphor.

The chromaticity on color coordinates and luminance of each phosphor layer

are measured to examine the phosphor characteristic of each color phosphor pattern mixed with the far-infrared emitting material, which is manufactured according to the above-described method. This data is shown in <Table 2>, along with the chromaticity of the conventional phosphor layer without the far-infrared emitting material. The relative luminance is the value when the luminance of the conventional phosphor is assumed as 100%.

<Table 2>

	conventional phosphor		mixed phosphor		rel. luminance (%)	luminance dif. (%)
	X coordinate	Y coordinate	X coordinate	Y coordinate		
red	0.652	0.341	0.650	0.342	95.7	-4.3
green	0.313	0.610	0.314	0.607	93.0	-7
blue	0.152	0.058	0.152	0.055	97.3	-2.7

As shown in <Table 2>, and in FIGs. 12, 13 and 14, there is no significant difference in the luminous spectrum and color coordinates when mixing the far-infrared emitting material, though the luminance is slightly lowered. This is because the far-infrared emitting material without the luminous characteristic is mixed in the phosphor layer, which results in degrading the luminous efficiency accordingly. Therefore, the amount of the far-infrared emitting material included in the phosphor slurry must not exceed 50% of the maximum amount of the phosphor, and a favorable body color thereof is also white or off-white for maintaining the luminance efficiency of the phosphor.

Moreover, since the average particle diameter of the phosphor is $5\text{-}8\mu\text{m}$ and its maximum particle diameter is $10\text{-}14\mu\text{m}$, the average and maximum particle diameters of the far-infrared emitting material is preferably below those of the phosphor. When the diameter is minimized as much as possible, such a non-luminous part of the screen which is not displayed after the subsequent formation of the phosphor layer is reduced and the mixing with the pigment and/or phosphor is even better.

As described above, the far-infrared emitting material is included in the phosphor slurry while manufacturing the phosphor layer of the cathode ray tube, so that this cathode ray tube can offer beneficial far-infrared ray to the user. The luminance is slightly degraded, but it is not significant.

To demonstrate the far-infrared emitting effect of the screen completed as above, the following experiment was carried out.

Four roses and four carnations being in bloom to the same degree were put into respective flasks. These were placed 20cm from a television having the conventional screen operating without the far-infrared emitting material, and a television having the screen manufactured according to the Example 5. These were left in place for 100 hours.

Additionally, five onions being in the same state of maturity were put into beakers of water such that their roots are 1cm below the water's surface. The beakers were positioned 10cm from the flasks for 100 hours in front of the turned on televisions, as were the flowers.

FIGs. 15A and 15B are photographs of the roses, carnations and onions. In

FIG. 15A, the leftmost rose and carnation were exposed to the television adopting the conventional screen, and the remaining three roses and carnations were exposed to the television adopting the screen which emits the far-infrared rays and manufactured according to the above example. In FIG. 15B, the leftmost onion was exposed to the television adopting the conventional screen, and the remaining four onions were exposed to the television adopting the screen which emits the far-infrared rays and manufactured according to the example. Here, all that were exposed to the far-infrared emitting cathode ray tube and thus under the influence of the far-infrared rays grow quickly, wither slowly, and maintain freshness, while the flowers exposed to the conventional cathode ray tube has withered and dried before blooming, and the onion grows slowly and has no buds.

In a still further embodiment of the present invention, a far-infrared emitting cathode ray tube comprises a color selecting electrode, i.e., shadow mask, which emits far-infrared rays by forming a far-infrared emitting layer on the outer surface of the shadow mask toward the phosphor layer.

The shadow mask is constructed such that apertures are formed in a metal plate, corresponding to the phosphor to allow the electron beams emitted from the electron gun to accurately collide with the phosphor. Only a portion of the electron beams emitted from the electron gun land on the phosphor layer having passed through the apertures, and the remaining 70-80% collide with the shadow mask. By this collision, electrical energy is instantaneously converted into heat energy, which augments the surface temperature of the shadow mask by up to roughly 80°C.

In this embodiment, the far-infrared emitting material is included in the shadow mask requiring several treatments due to the rise of temperature, so that the far-infrared rays can be emitted from the cathode ray tube which had emitted harmful electromagnetic waves.

The far-infrared emitting material is first mixed with solvent to prepare a coating solution. Then, this solution is coated on the surface of the shadow mask, using a spray coating, or using a powder flame or powdered plasma spray which is used for preventing doming of the shadow mask. When using the powder flame or plasma spray, a very securely fixed, far-infrared emitting layer can be manufactured.

The far-infrared emitting material is composed of at least one of aluminium oxide (Al_2O_3), silicon dioxide (SiO_2), magnesium oxide (MgO), zirconium oxide (ZrO_2), carbons, ferric oxide (Fe_2O_3), manganese dioxide (MnO_2), manganese monoxide (MnO), cupric oxide (CuO), tricobalt tetraoxide (Co_3O_4), nickel monoxide (NiO), chromic oxide (Cr_2O_3), titanium oxide (TiO_2), boron oxide (B_2O_3), sodium oxide (Na_2O), potassium oxide (K_2O), molybdenum sesquioxide (Mo_2O_3), calcium oxide (CaO), lithium oxide (Li_2O), zinc oxide (ZnO), bismuth oxide (Bi_2O_3), barium oxide (BaO), and an alloy of these, wherein a low heat-expansion material is preferable. The diameter of the far-infrared emitting material is in the range from $0.05\mu\text{m}$ to $2\mu\text{m}$. If the diameter is smaller than $0.05\mu\text{m}$, the far-infrared emitting efficiency is too low, but if greater than $2\mu\text{m}$, the apertures of the shadow mask are clogged. Thus, it is preferable that the above range is maintained, and the average diameter is set to about $0.4\mu\text{m}$.

Also, the thickness of the far-infrared emitting layer to be formed is within 0.1-10 μ m, preferably 3-5 μ m. The reason for setting the range is that the emittance effect is insufficient when thinner than 0.1 μ m, and the apertures are clogged when thicker than 10 μ m.

As described above, if the cathode ray tube is manufactured by adopting the shadow mask covered with the far-infrared emitting material on the outer surface thereof, the temperature of the shadow mask increases up to approximately 80°C due to the collision of the electron beams during cathode ray tube operation, which facilitates the emittance of the far-infrared rays.

Hereinbelow, a preferred embodiment will be described in detail with reference to examples.

EXAMPLE 6

30wt% of SiO₂, 15wt% of Al₂O₃, 30wt% of Li₂O, 10wt% of ZnO and 15wt% of ZrO₂ are mixed, thereby forming low heat-expansion far-infrared ceramic. Then, a far-infrared emitting mixture for coating the shadow mask is prepared by mixing 20wt% of the far-infrared ceramic, 20wt% of frit (lead-borate glass) consisting of 80wt% of PbO, 10wt% of B₂O₃ and 10wt% of SiO₂, 59wt% of distilled water, and 1wt% of a dispersant. This mixture is sprayed on the outer surface of the shadow mask toward the phosphor layer to a thickness of 5 μ m and dried, thereby forming the shadow mask coated with the far-infrared emitting material.

EXAMPLE 7

35wt% of SiO_2 , 4wt% of Al_2O_3 , 55wt% of ZrO_2 , 4wt% of Fe_2O_3 and 2wt% of Bi_2O_3 are consistently mixed, thereby forming far-infrared emitting material. The average particle diameter of the far-infrared emitting material is $1\mu\text{m}$, and the maximum particle diameter is $1.5\mu\text{m}$. As shown in FIG. 16, the far-infrared emitting material is coated on the outer surface of the shadow mask to a thickness of $3\mu\text{m}$ by the powder flame spray. According to this method, far-infrared emitting material 4 is sprayed by high-pressure air and, simultaneously, oxygen and acetylene gases 8 are sprayed to flame, so that a compact far-infrared emitting layer is formed on shadow mask 7.

EXAMPLE 8

60wt% of SiO_2 , 20wt% of Al_2O_3 , 5wt% of Fe_2O_3 and 5wt% of MgO are consistently mixed, thereby forming far-infrared emitting powder. The far-infrared emitting powder thus obtained is coated on the outer surface of the shadow mask by plasma spray as shown in FIG. 17, thereby forming the far-infrared emitting layer to a thickness of $5\mu\text{m}$. According to the plasma spray, high-pressure argon or nitrogen gases 9 are sprayed to produce a high-voltage arc, which in turn creates plasma and simultaneously supplies far-infrared emitting powder 4. Thus, a compact far-infrared emitting layer is formed on the outer surface of shadow mask 7.

As described above, when the far-infrared emitting material layer is formed on the outer surface of the shadow mask, the temperature is increased to

approximately 80°C, so that an excellent far-infrared emitting effect can be obtained as compared with that forming the material in other parts.

In another embodiment of the present invention, a cathode ray tube comprises an envelope formed by a panel, a funnel and neck, wherein the far-infrared emitting material is included in the outer graphite layer covering the outer surface of the funnel, and thus emits the far-infrared rays.

The graphite formed on the outer surface of the funnel functions as a capacitor. In the cathode ray tube, an anode is supplied with 25KV which is supplied by rectifying the output of the flyback transformer. Since this output is obtained from a square wave input, the ripple is high. Generally, in a rectifier circuit, a capacitor is used in a smoothing element to eliminate the ripple. Here, the outer and inner graphite layers of the cathode ray tube serve as the capacitor.

In an office, the rear of a computer terminal often faces another person located behind it, and thus electromagnetic waves are likely to harm that person. However, in this embodiment, the far-infrared emitting material is included in the outer graphite layer to emit the far-infrared rays from the rear portion of the cathode ray tube.

On the other hand, as described above in view of the Stefan-Boltzmann formula, the emitted energy amount is in proportional to 4-square of the absolute temperature. This denotes that the applied temperature should be high to heighten the emission efficiency of the far-infrared rays and increase the amount of emitted energy. In the case of the terminal mounted with the cathode ray tube, the temperature of the outer graphite is increased to 45-55°C, after being turned on. Also, the larger the size of the cathode ray tube is, the higher the temperature

becomes, and under certain circumstances, up to about 70°C. Therefore, more far-infrared radiation can be emitted from the outer graphite, using the natural heat generated during the operation of the terminal.

A method for manufacturing the cathode ray tube including the far-infrared emitting material in the outer graphite layer will be described in detail.

EXAMPLE 9

15wt% of graphite for outer graphite layer, 15wt% of a mixture of SiO₂, Al₂O₃, MgO, CaO and TiO₂, 0.1wt% of kasil, 5wt% of the others, and the remainder of distilled water are mixed to form a composition for the outer graphite layer.

The overall composition is tinted gray since the introduced far-infrared emitting material is white. In accordance with the common method, the obtained composition is sprayed to coat the outer surface of the funnel, using a spray gun for coating under high pressure (more than 1.5kg/cm²).

FIG. 10A illustrates the spectral analysis of the far-infrared emitting material used in the above example, and FIG. 10B illustrates the spectral emittance graph versus wavelength.

FIG. 18 shows a cross-sectional view of a portion of a funnel of the cathode ray tube 10 manufactured according to the example 9 of the present invention, wherein the far-infrared emitting material 4 is consistently mixed with the outer graphite 12 on the funnel 11.

To demonstrate the far-infrared emitting effect of the cathode ray tube

having the outer graphite layer which has been manufactured by including the far-infrared emitting material, the following experiment was executed, which will be described with reference to FIG. 19.

Three roses in bloom to the same degree were put into respective flasks.

First, the cathode ray tube produced according to example 9 of the present invention was installed on a cabinet 13, one rose (A) is placed near a vent 14 of the cabinet, as shown in the drawing. Another rose (B) was placed on a conventional cathode ray tube of the same type as example 9, except that the far-infrared emitting material is omitted. The third rose (C) was placed so as not to be affected by electromagnetic waves from any cathode ray tube.

Of the above three roses, rose B stops blooming after 70 hours, and the tips of its petals began to turn black after 90 hours, while roses A and C are still in bloom. After 120 hours, the petals of rose B are completely scorched and its stalk is also dried. At this time, the petals of rose A begins to dry slightly. After 160 hours, the petals and stalk of rose B are dried up and mummified. At this time, rose A is drying, and rose C is beyond full bloom. The results obtained by measuring the temperature at each location after the elapse of 160 hours is shown in <Table 3>.

<Table 3>

classification	A	B	C
temperature of water in flask (°C)	37	37	26
surface temperature of vent in cabinet (°C)	40	40	N/A
surface temperature of outer graphite (°C)	43	43	N/A

Here, rose A under the influence of the far-infrared rays begins drying earlier than rose C in natural surroundings, because the water temperature in the flask is 37°C due to the heat emitted from the cathode ray tube as shown in the <Table 3>, and the harmful electromagnetic effects on rose A. However, when comparing roses A and B, it can be confirmed that far-infrared rays emitted from the outer graphite layer of the cathode ray tube are useful for a living body.

EXAMPLE 10

30wt% of a mixture of SiO₂, Al₂O₃ and MgO, 20wt% of kasil, and 50wt% of distilled water are mixed to form a composition for outer graphite layer.

After the above composition obtained according to the conventional method is coated over the outer graphite layer of a cathode ray tube by spray or brush, it is dried in an oven at 100°C for 30 minutes.

FIGs. 6 show the spectral analysis (FIG. 6A) of the far-infrared emitting material, and the graph (FIG. 6B) obtained by measuring spectral emittance of the far-infrared emitting material used in the above example.

FIG. 20 shows a cross-sectional view of a portion of a funnel of cathode ray tube 10 manufactured according to the above example of the present invention, wherein far-infrared emitting material layer 4 is formed over outer graphite layer 12 on funnel 11.

The cathode ray tube manufactured according to example 10 also has a far-infrared emitting effect similar to that of the cathode ray tube manufactured according to example 9.

In embodiments of the present invention as described above, far-infrared ray material is included in each part of the interior of cathode ray tube, so that far-infrared rays are emitted. Therefore, the cathode ray tube which has emitted electromagnetic waves harmful to living bodies emits far-infrared rays which are useful, thereby providing pleasant working surroundings to the user who operates the cathode ray tube.

Apart from the far-infrared emitting cathode ray tube described with reference to the examples according to the present invention, it will be apparent that many modifications and variations could be effected easily by one skilled in the art without departing from the scope of the novel concepts of the present invention, including monochrome and color cathode ray tube.

CLAIMS:

1. A far-infrared emitting cathode ray tube, wherein a layer of far-infrared emitting material (4) having not more than a $5\mu\text{m}$ average diameter is provided on a black matrix (2) to a thickness of from 0.2 to $100\mu\text{m}$.
2. A far-infrared emitting cathode ray tube as claimed in claim 1, wherein the thickness of said layer of far-infrared emitting material is from 0.5 to $50\mu\text{m}$.
3. A far-infrared emitting cathode ray tube, wherein a far-infrared emitting material (4) having an average diameter of not more than $5\mu\text{m}$ is included within a black matrix layer (2) of the cathode ray tube as much as 1-100wt%, based upon the total amount of the mixture with the graphite.
4. A far-infrared emitting cathode ray tube as claimed in claim 3, wherein the mixing amount of said far-infrared emitting material is from 10 to 100wt% based upon the total amount of the mixture with the graphite.
5. A far-infrared emitting cathode ray tube, wherein far-infrared emitting material (4) of not more than 50wt% with respect to the amount of phosphor is contained in a phosphor layer (5) of the cathode ray tube.
6. A far-infrared emitting cathode ray tube as claimed in claim 5, wherein said far-infrared emitting material is from 1 to 40wt% with respect



to the amount of the phosphor.

7. A far-infrared emitting cathode ray tube as claimed in claim 5 or 6, wherein the average diameter of said far-infrared emitting material is between $0.5\mu\text{m}$ and $4\mu\text{m}$, and the maximum diameter thereof is $8\mu\text{m}$.
8. A far-infrared emitting cathode ray tube as claimed in claims 5, 6 or 7, wherein said far-infrared emitting material is a metal oxide prevented from being mixed with any one of iron (Fe), cobalt (Co) and nickel (Ni) as an impurity.
9. A far-infrared emitting cathode ray tube as claimed in any of claims 5 to 8, wherein the colour of said far-infrared emitting material is white.
10. A far-infrared emitting cathode ray tube as claimed in any preceding claim, wherein said far-infrared emitting material is composed of at least one selected from the group consisting of aluminium oxide (Al_2O_3), silicon dioxide (SiO_2), magnesium oxide (MgO), zirconium oxide (ZrO_2), carbons, ferric oxide (Fe_2O_3), manganese dioxide (MnO_2), cupric oxide (CuO), tricobalt tetraoxide (Co_3O_4), nickel monoxide (NiO), chromic oxide (Cr_2O_3), lithium oxide (Li_2O), zinc oxide (ZnO), bismuth oxide (Bi_2O_3), barium oxide (BaO), titanium oxide (TiO_2), boron oxide (B_2O_3), sodium oxide (Na_2O), potassium oxide (K_2O), phosphorus pentoxide (P_2O_5), molybdenum sesquioxide



(Mo_2O_3), calcium oxide (CaO) and a complex thereof.

11. A far-infrared emitting cathode ray tube, wherein far-infrared emitting material (4) having a diameter of from 0.05 to $2\mu\text{m}$ and a thickness of from about 0.1 to $10\mu\text{m}$ on the outer surface of a shadow mask (7) of the cathode ray tube toward the phosphor layer.
12. A far-infrared emitting cathode ray tube as claimed in claim 11, comprising a compact far-infrared emitting layer formed by coating said far-infrared emitting material on the outer surface of said shadow mask by means of a powder flame spray or plasma spray.
13. A far-infrared emitting cathode ray tube, wherein an outer graphite layer (12) on the outer surface of a funnel (11) of the cathode ray tube comprises far-infrared emitting material.
14. A far-infrared emitting cathode ray tube as claimed in claim 13, wherein said far-infrared emitting material is consistently dispersed in said outer graphite layer.
15. A far-infrared emitting cathode ray tube as claimed in claim 13 or 14, wherein said far-infrared emitting material is coated on said outer graphite layer.



16. A far-infrared emitting cathode ray tube, wherein at least one of a phosphor layer (5), a black matrix (2), an outer graphite layer (12), and a shadow mask (7), includes a far-infrared emitting material for emitting far-infrared rays.

17. A far-infrared emitting cathode ray tube as claimed in any of claims 11 to 16, wherein said far-infrared emitting material is composed of at least one selected from the group consisting of silicon dioxide (SiO_2), aluminium oxide (Al_2O_3), magnesium oxide (MgO), zirconium oxide (ZrO_2), carbons, ferric oxide (Fe_2O_3), manganese dioxide (MnO_2), manganese monoxide (MnO), cupric oxide (CuO), tricobalt tetraoxide (Co_3O_4), nickel oxide (NiO), chromic oxide (Cr_2O_3), titanium oxide (TiO_2), boron oxide (B_2O_3), sodium oxide (Na_2O), potassium oxide (K_2O), molybdenum sesquioxide (Mo_2O_3), calcium oxide (CaO), lithium oxide (Li_2O), zinc oxide (ZnO), bismuth oxide (Bi_2O_3), barium oxide (BaO), and a complex thereof.

ABSTRACT

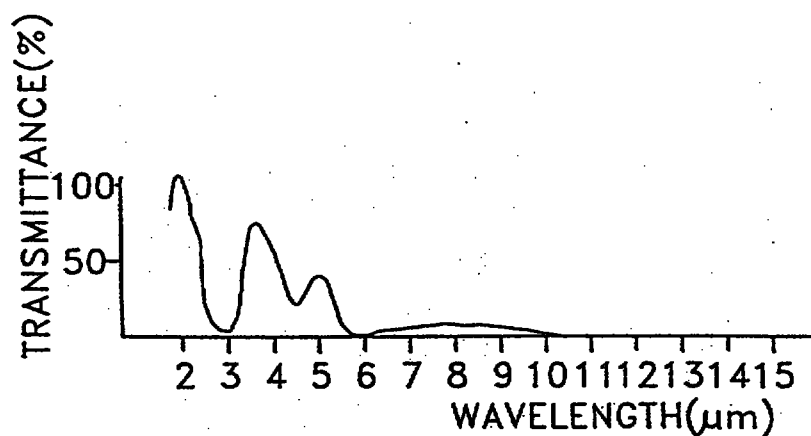
FAR-INFRARED EMITTING CATHODE RAY TUBE

A cathode ray tube includes far-infrared emitting material therein, and thus emits far-infrared rays useful for human body. The far-infrared emitting cathode ray tube is obtained by including far-infrared emitting material (4) in at least one of a phosphor layer (5), a black matrix (2), an outer graphite layer (12), and a shadow mask (7). This far-infrared emitting cathode ray tube provides a user favorable working surroundings.

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FIG.1



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FIG.2A

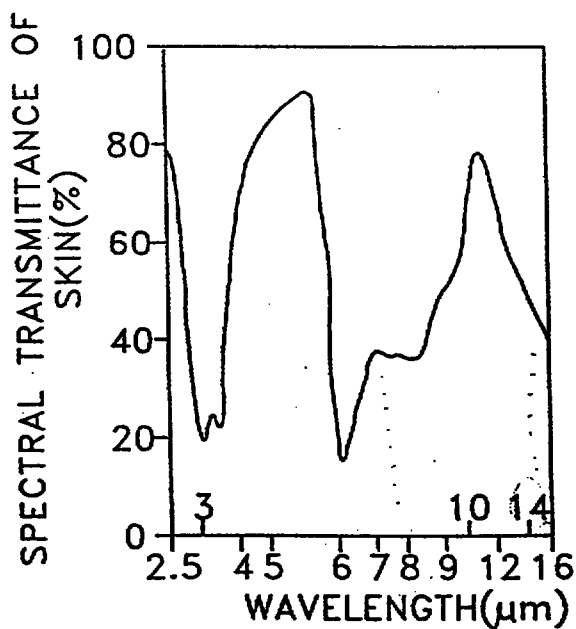
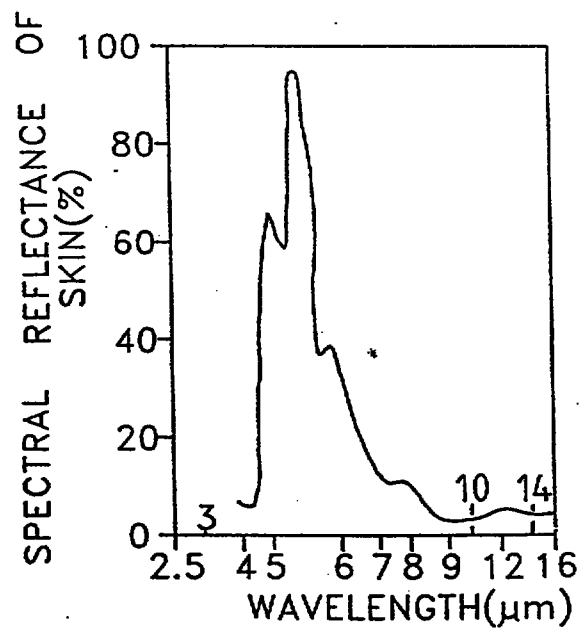


FIG.2B



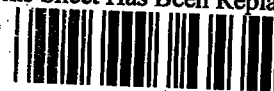


FIG.3A

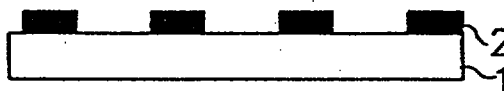


FIG.3B



FIG.3C



FIG.3D

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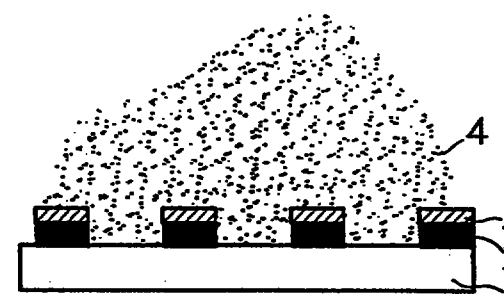


FIG.3E



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FIG. 3F



FIG. 3G

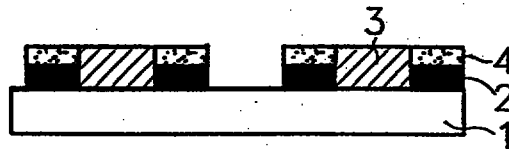
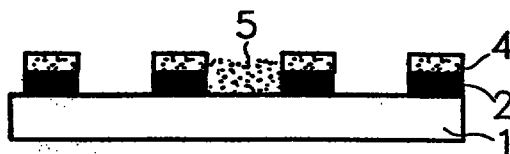


FIG. 3H



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FIG. 3I



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FIG.4A

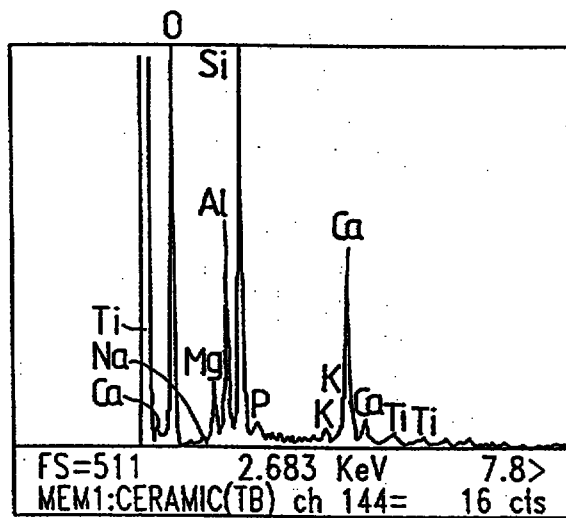
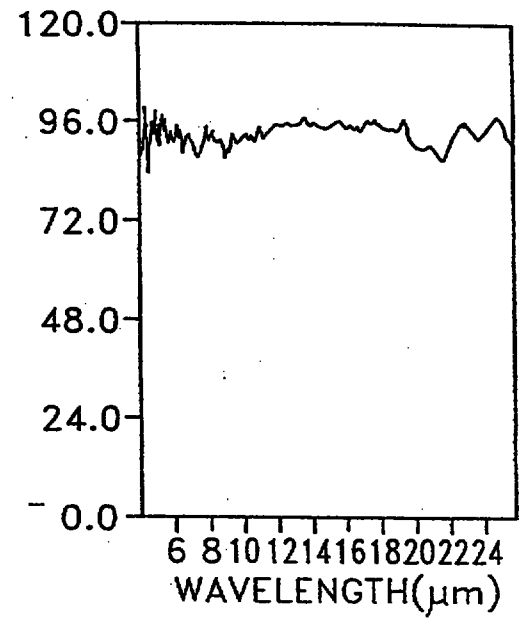
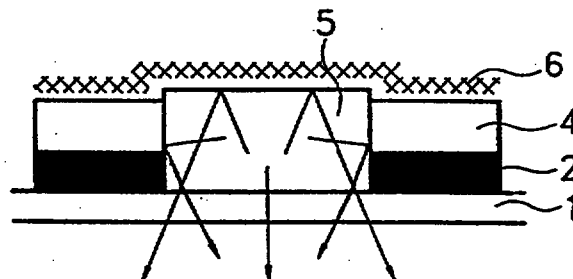


FIG.4B



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FIG.5



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FIG.6A

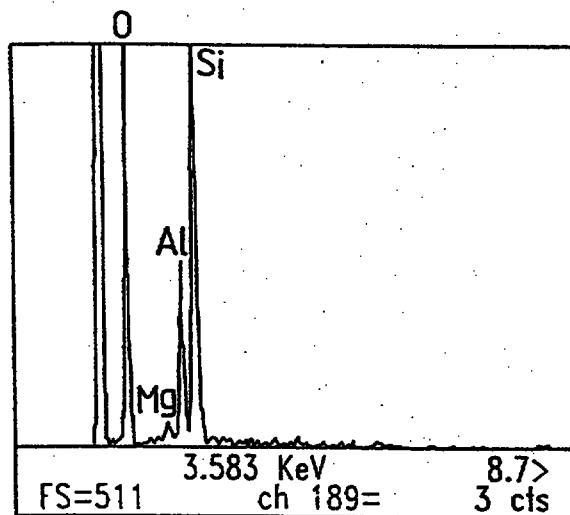


FIG.6B

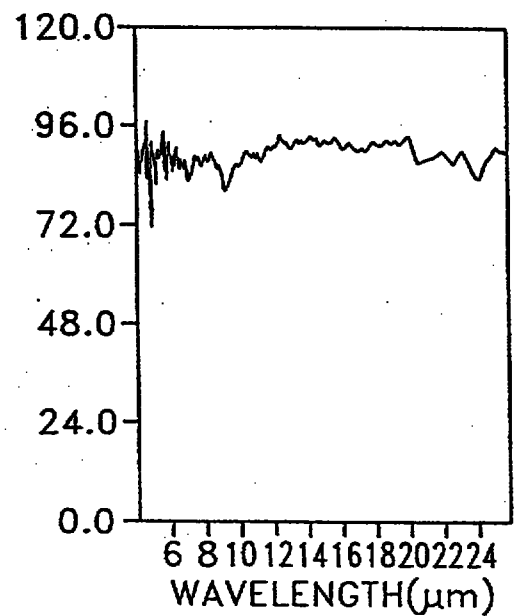


FIG.7

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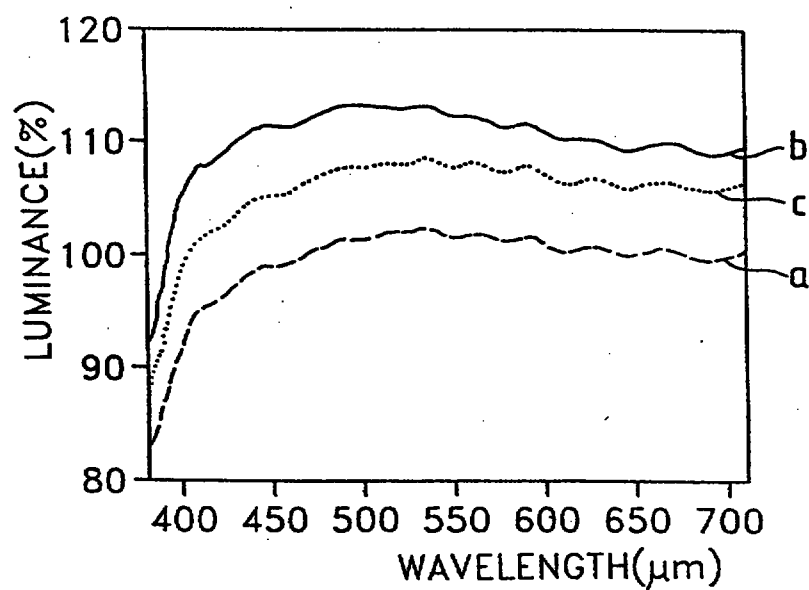




FIG. 8

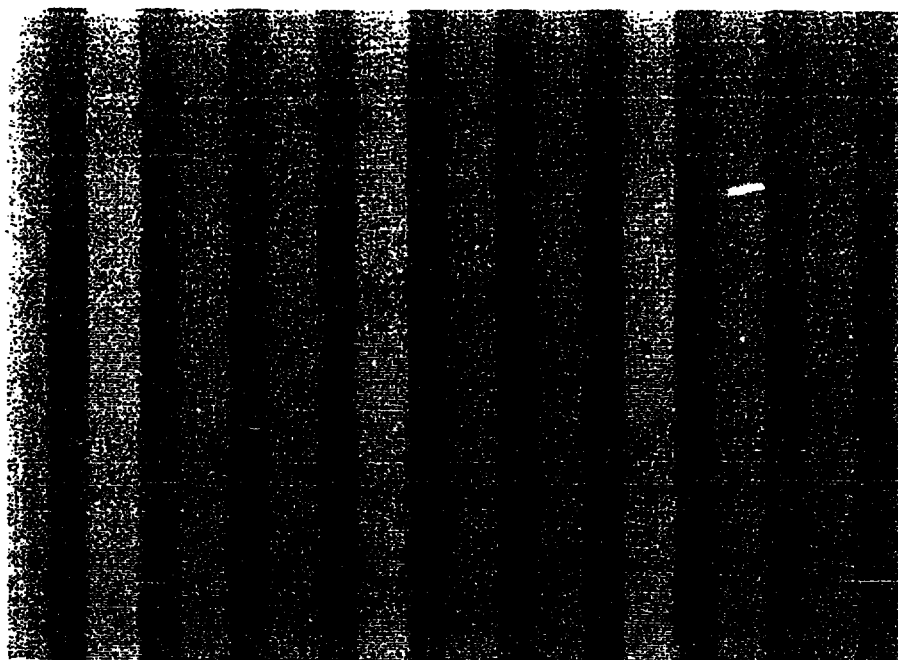


FIG. 9A FIG. 9B FIG. 9C FIG. 9D

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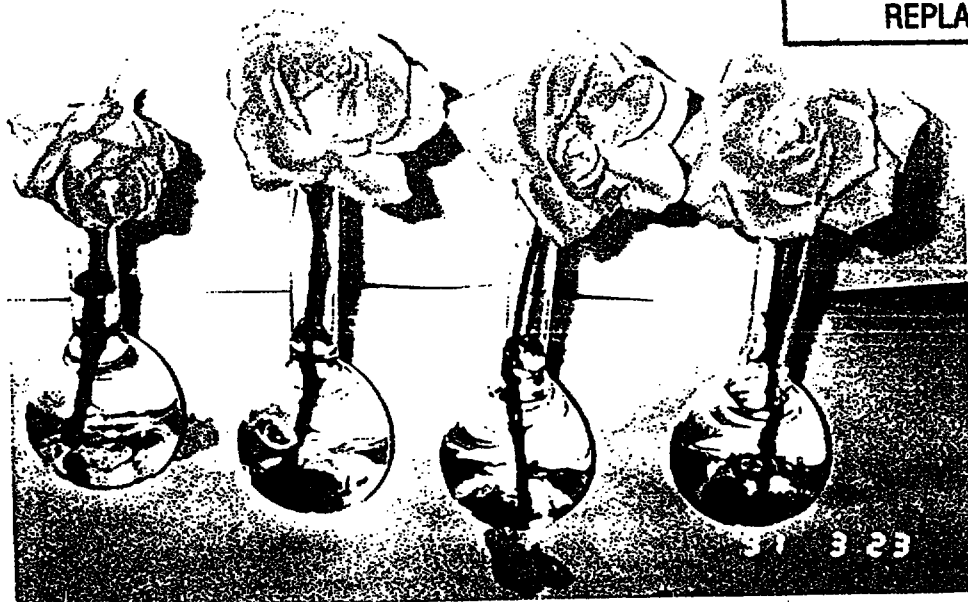




FIG.10A

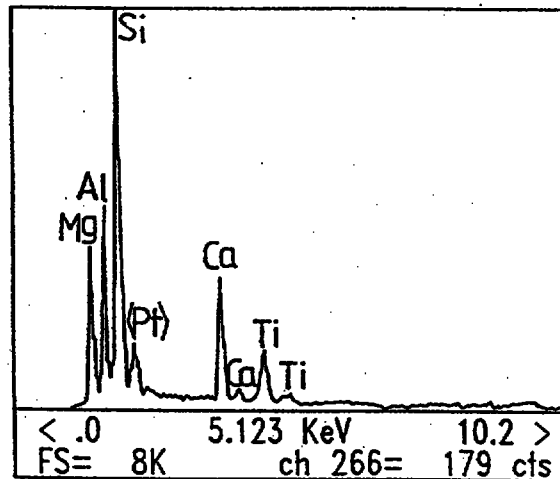
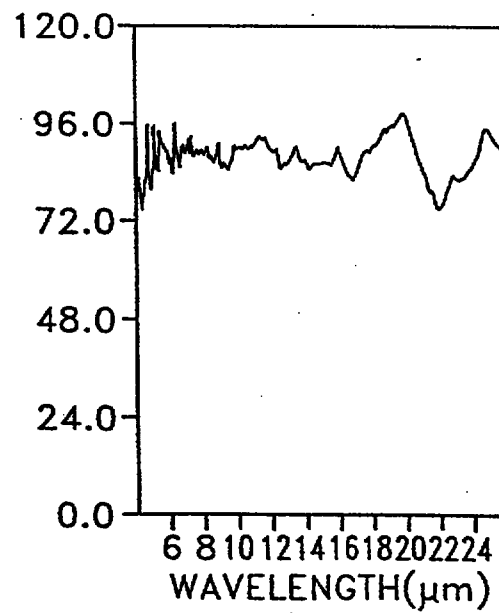


FIG.10B

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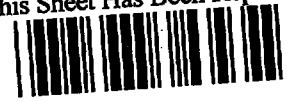


FIG. 11

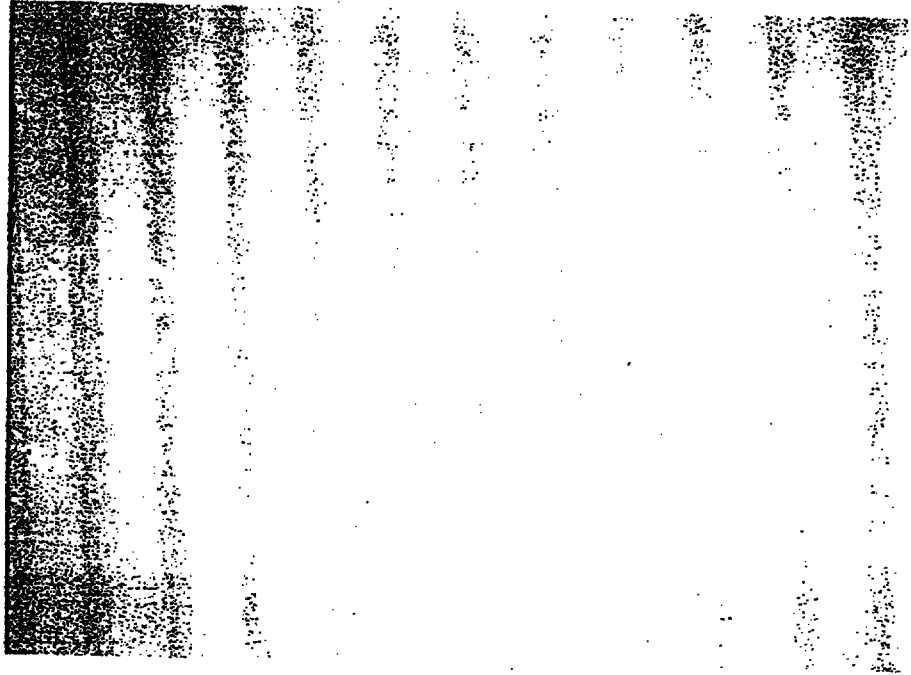
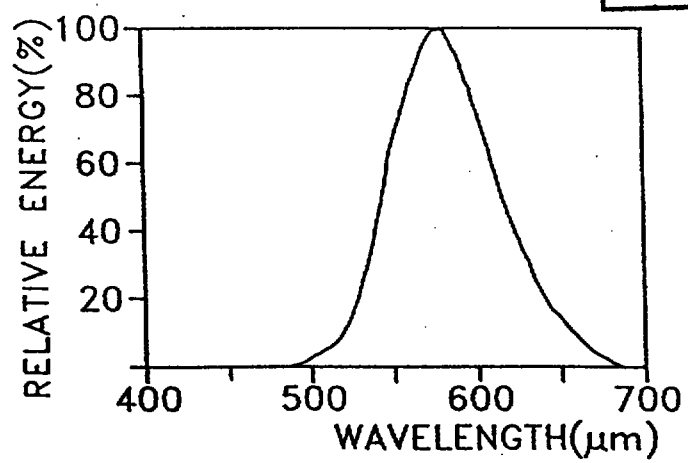


FIG. 12

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FIG.13

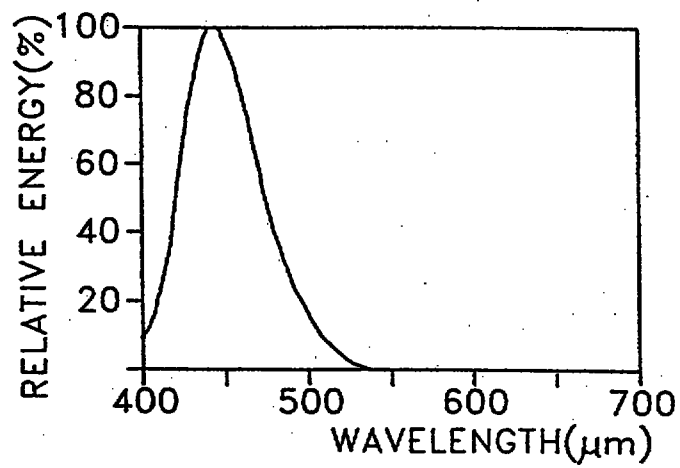
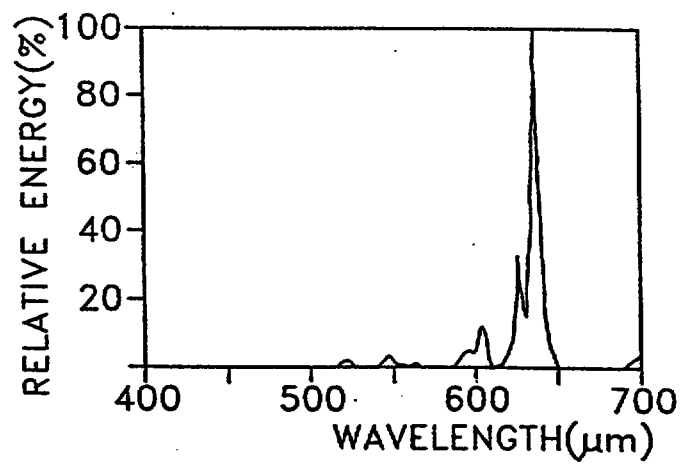


FIG.14

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FIG.15A

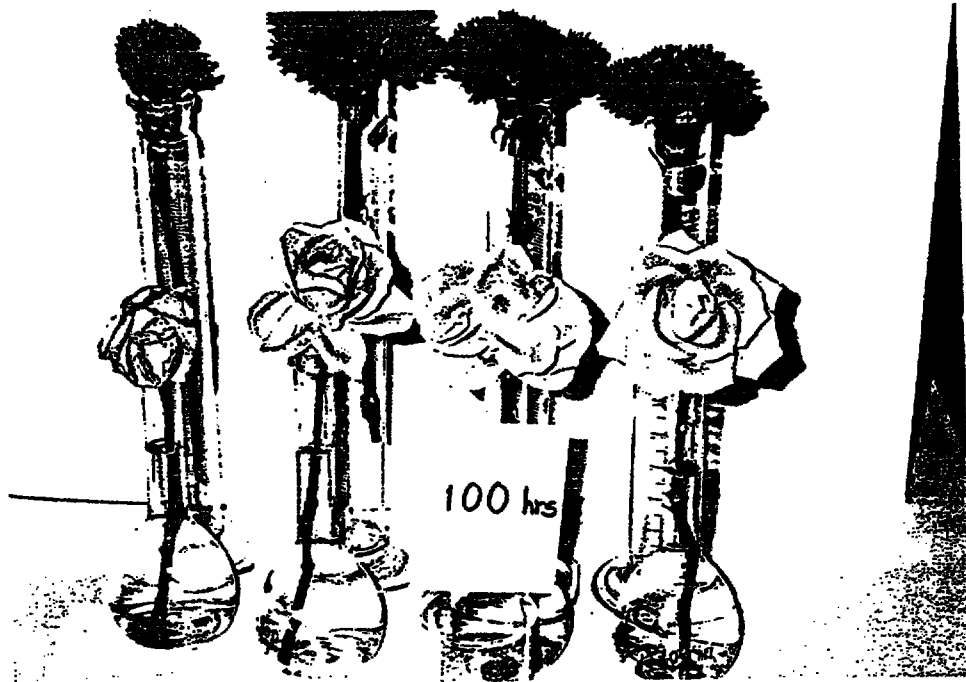
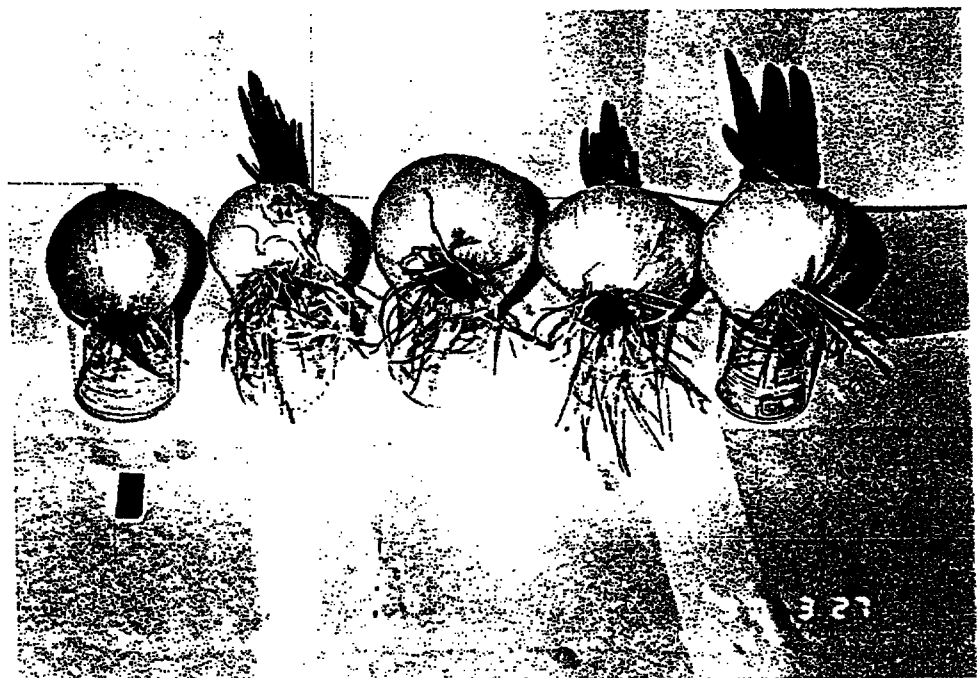


FIG.15B

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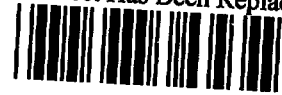


FIG. 16

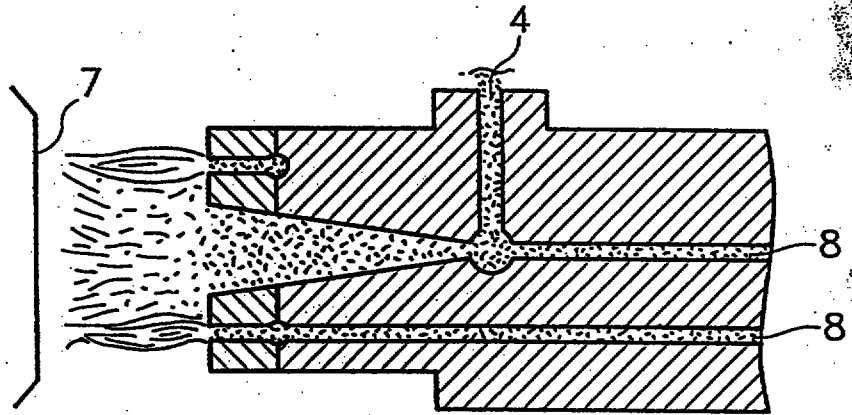
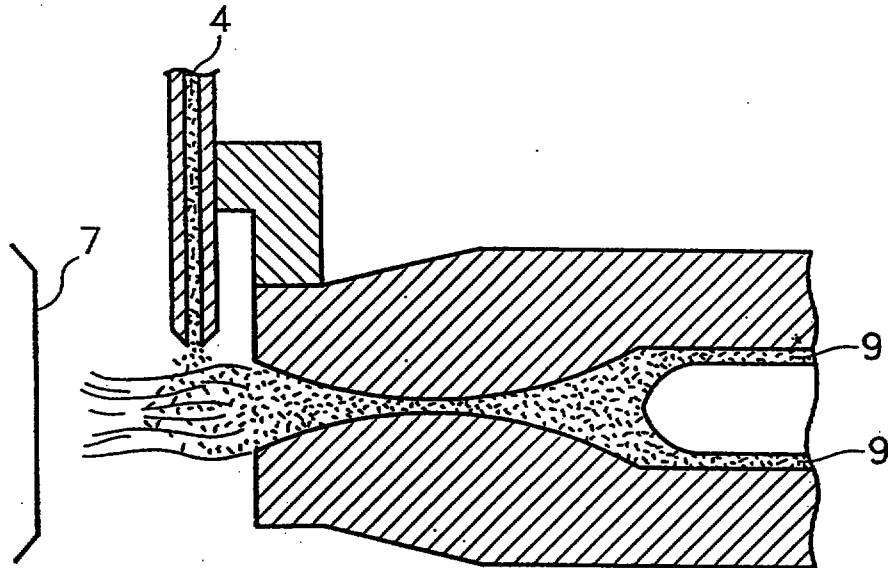


FIG. 17

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FIG.18

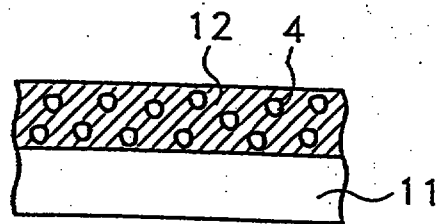


FIG.19

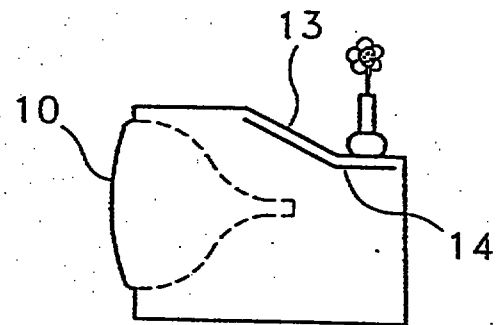
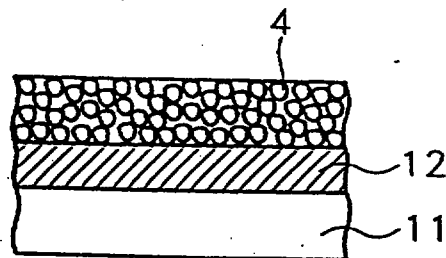


FIG.20



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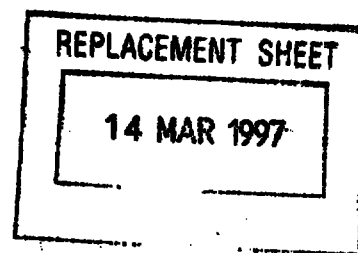
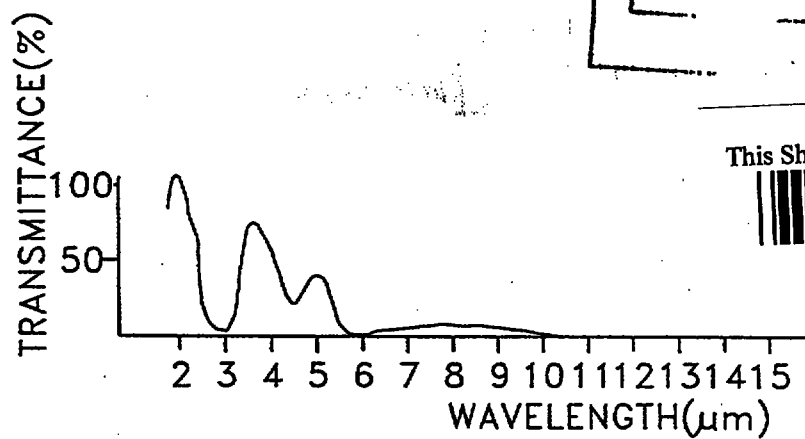


FIG. 1



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FIG. 2A

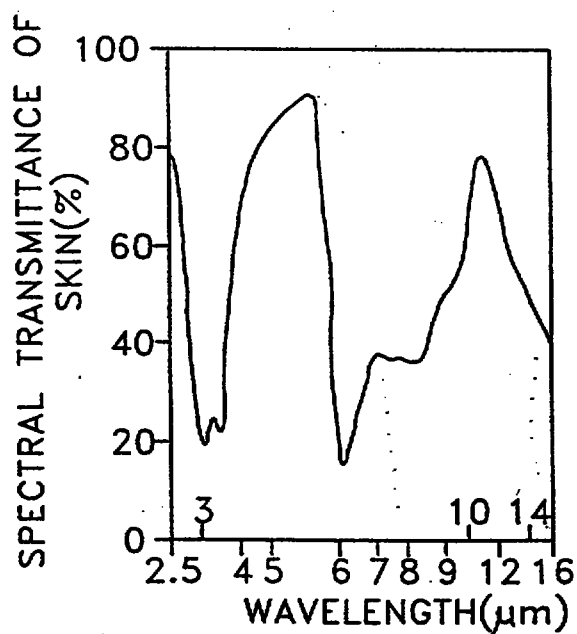
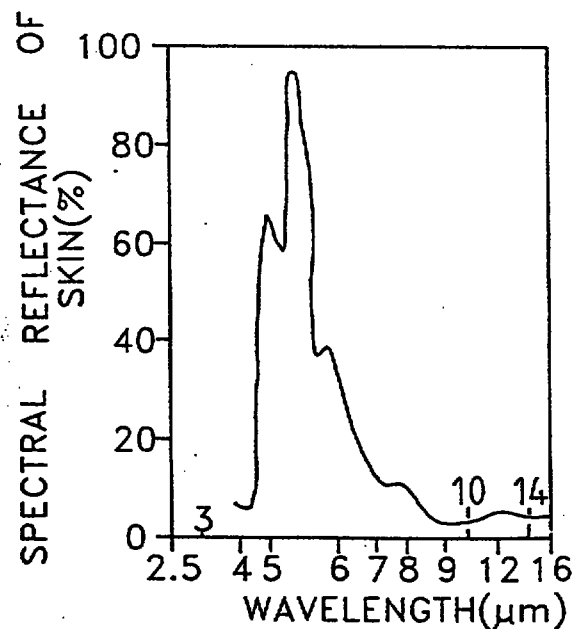


FIG. 2B



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FIG.3A



This Sheet Has Been Replaced



FIG.3B



FIG.3C



FIG.3D

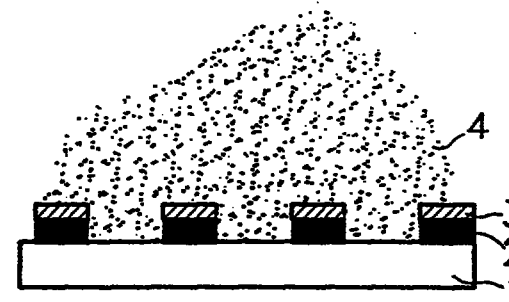


FIG.3E



14 MAR 1997

FIG.3F



This Sheet Has Been Replaced



FIG.3G

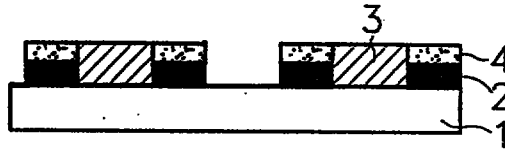


FIG.3H

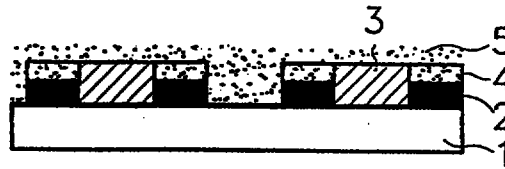
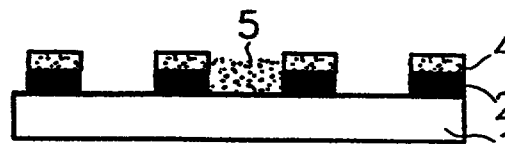


FIG.3I



This Sheet Has Been Replaced



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REPLACEMENT SHEET

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FIG.4A

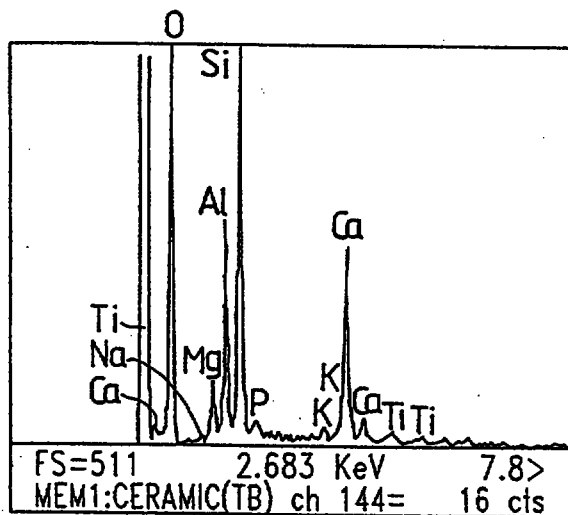
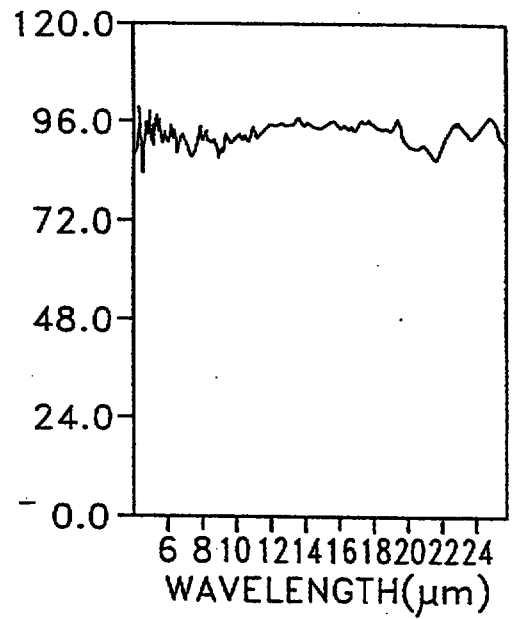
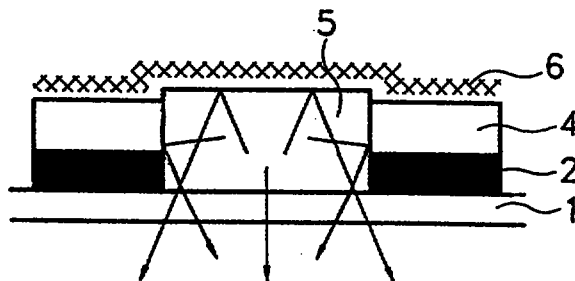


FIG.4B



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FIG.5



This Sheet Has Been Replaced



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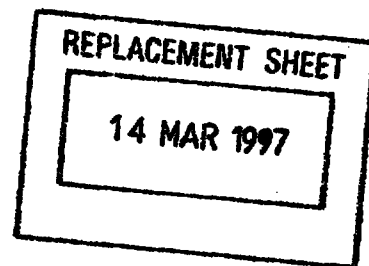


FIG.6A

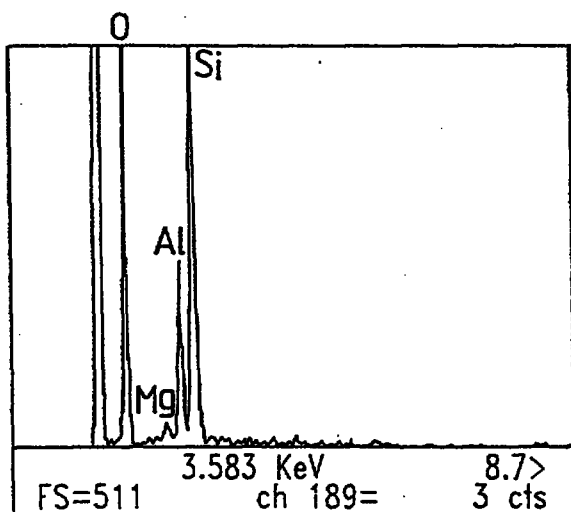


FIG.6B

THIS SHEET HAS BEEN REPLACED

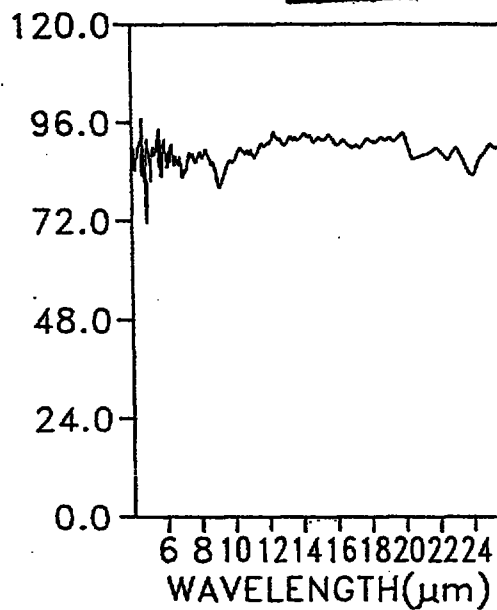
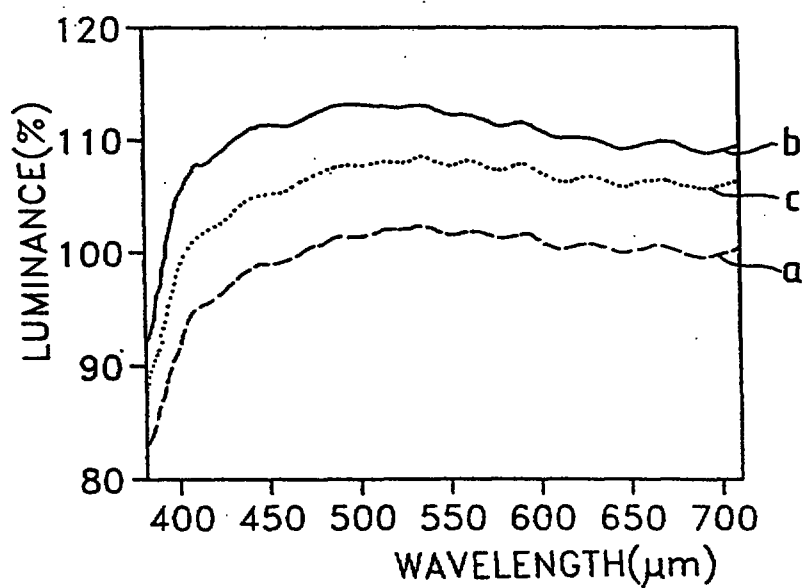


FIG.7



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FIG.8

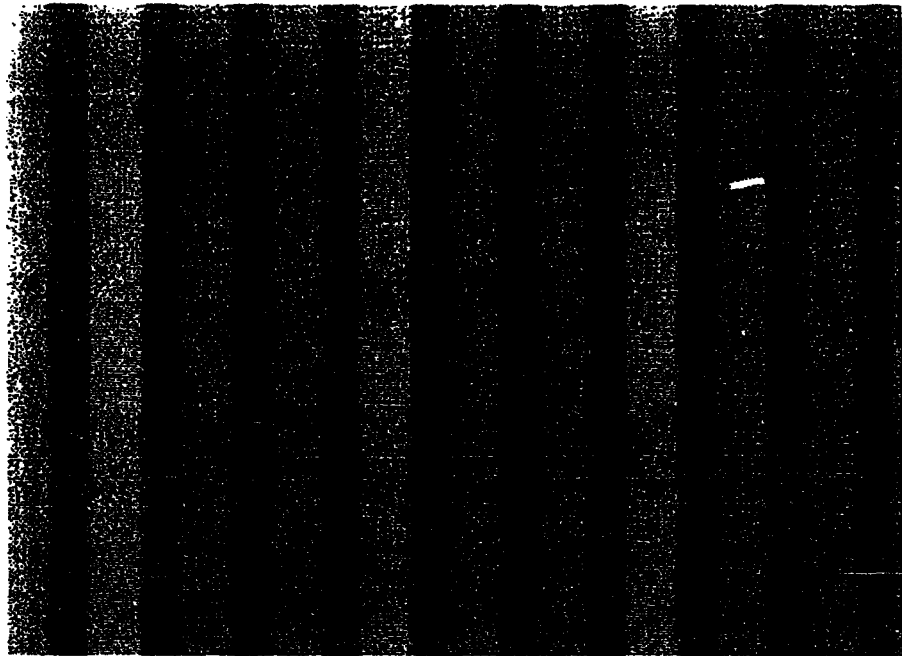
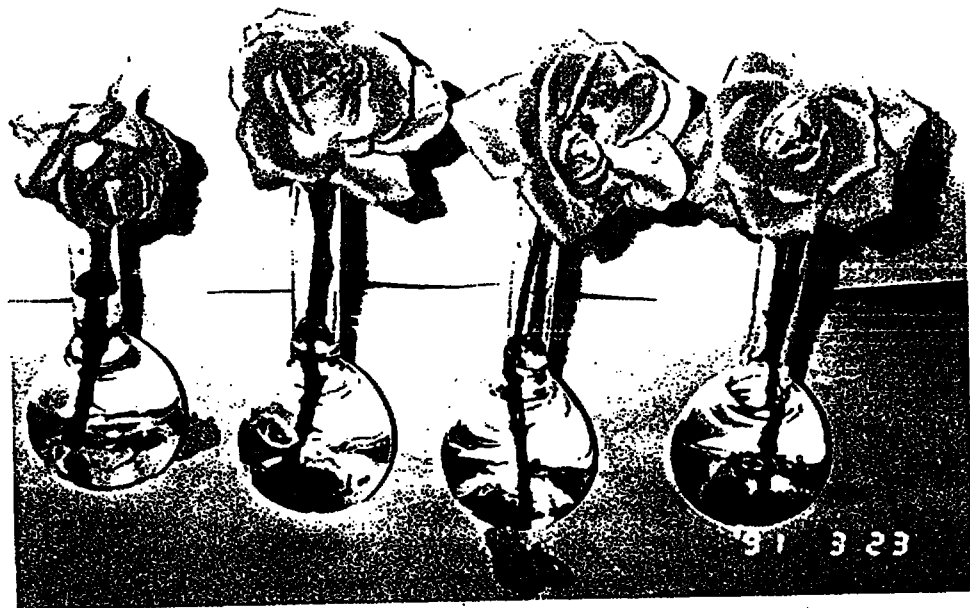


FIG.9A FIG.9B FIG.9C FIG.9D



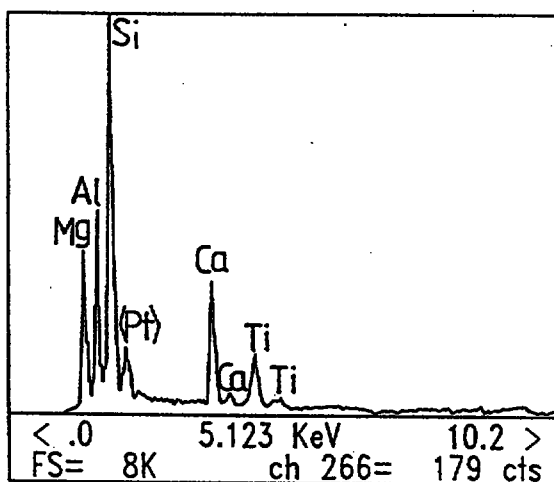
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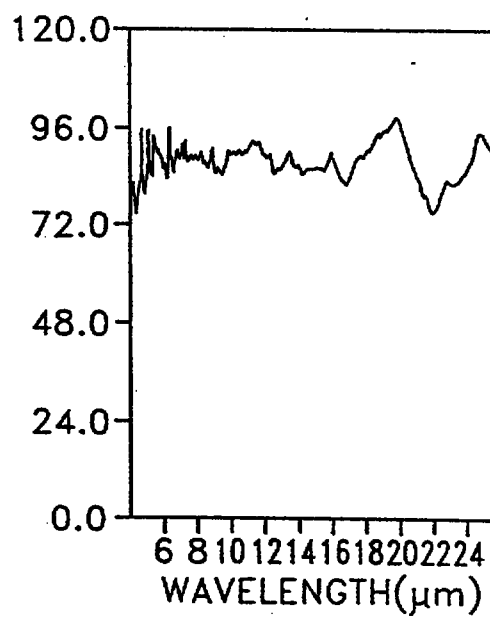
FIG.10A



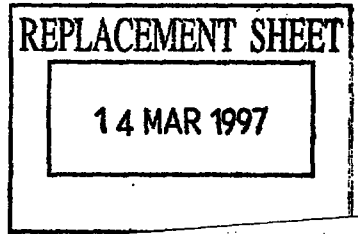
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FIG.10B



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This Sheet Has Been Replaced



FIG. 11

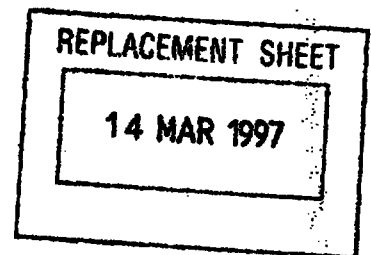
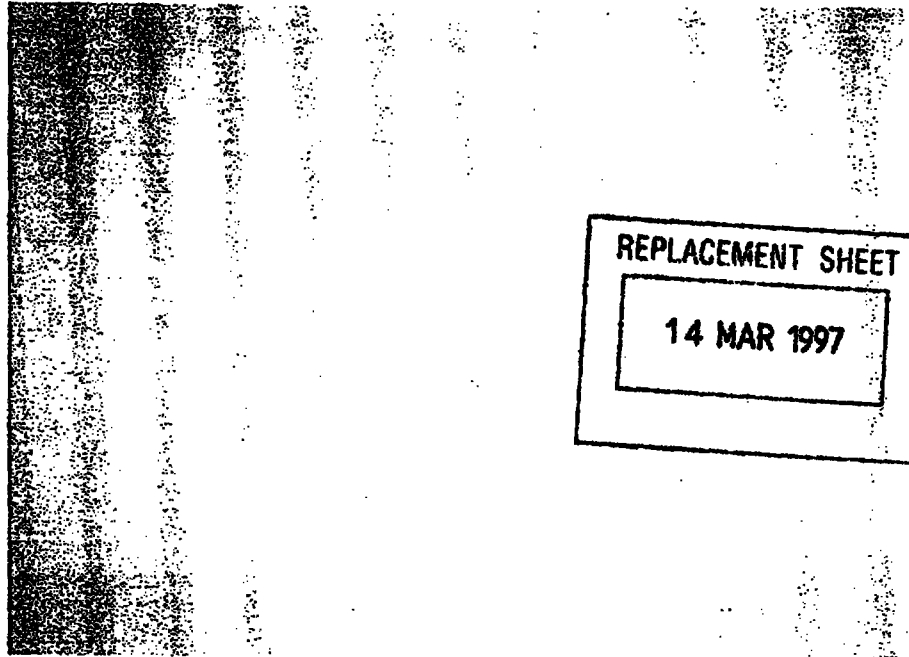
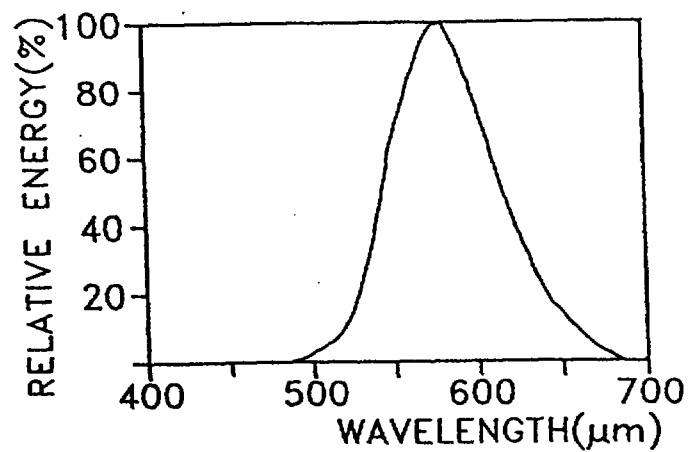


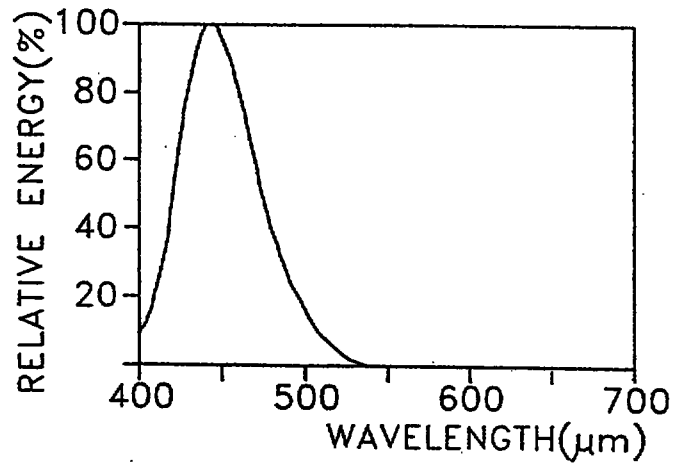
FIG. 12



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FIG.13

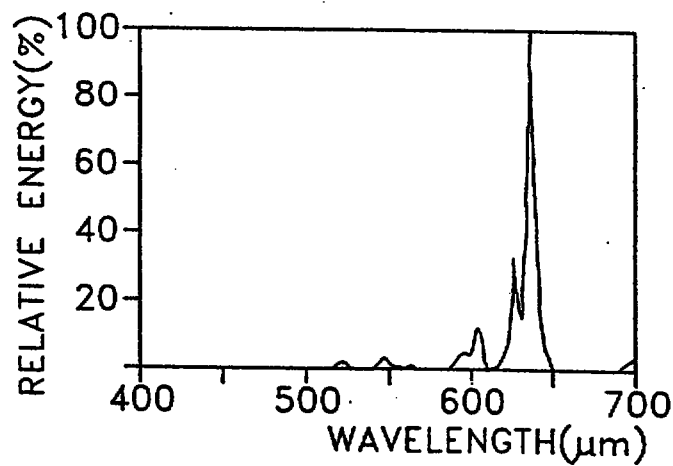
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REPLACEMENT SHEET

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FIG.14



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REPLACED

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REPLACEMENT SHEET

14 MAR 1997

FIG.15A

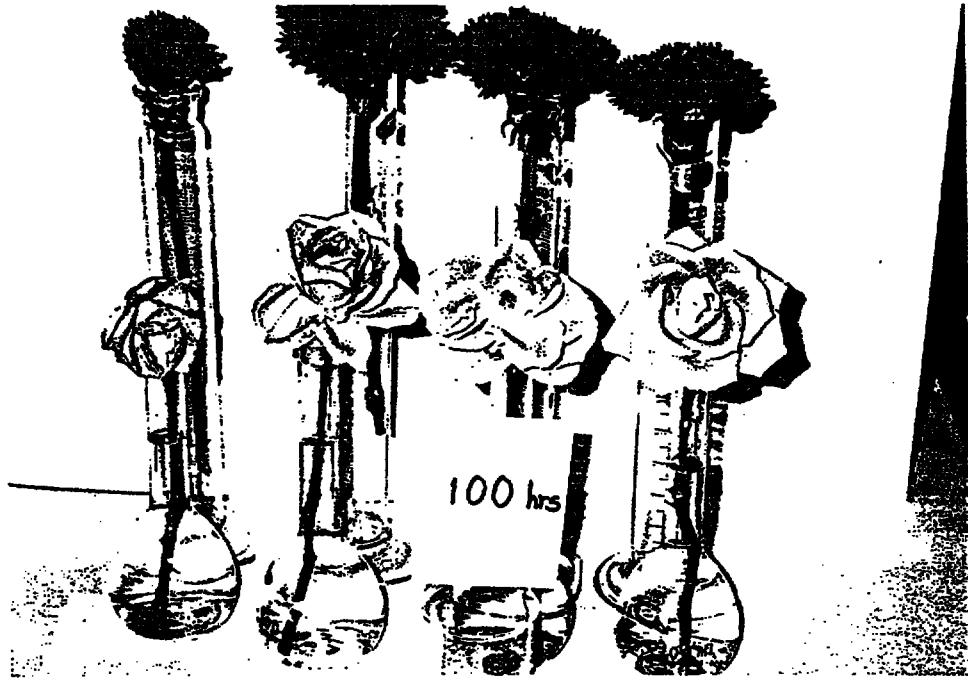
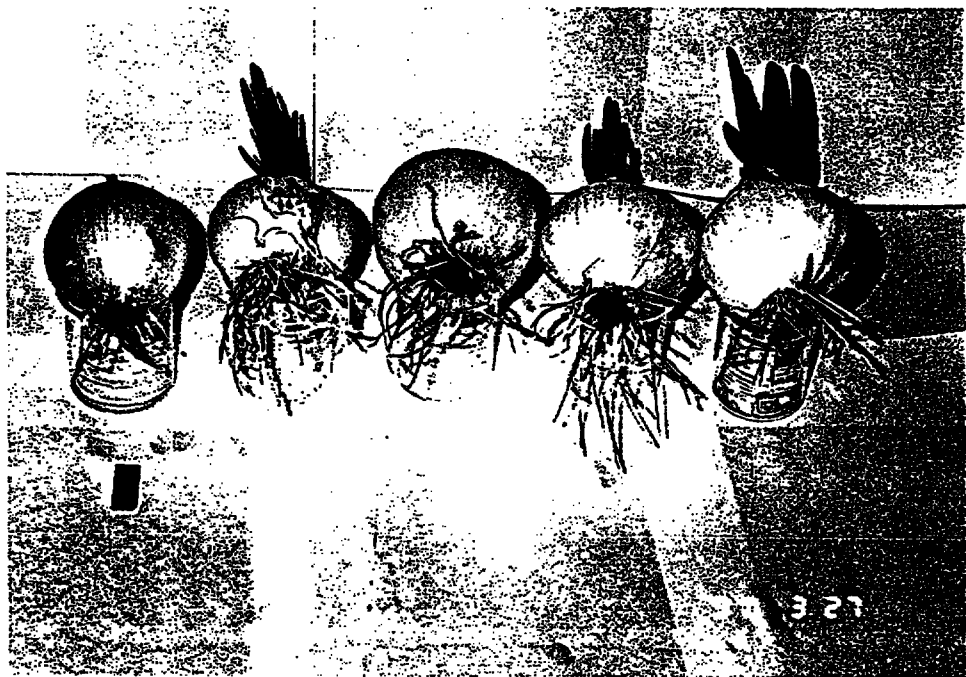


FIG.15B



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FIG. 16

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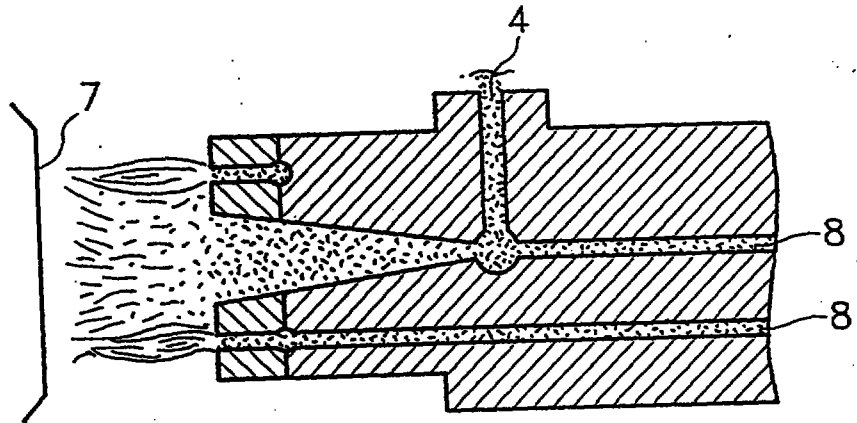
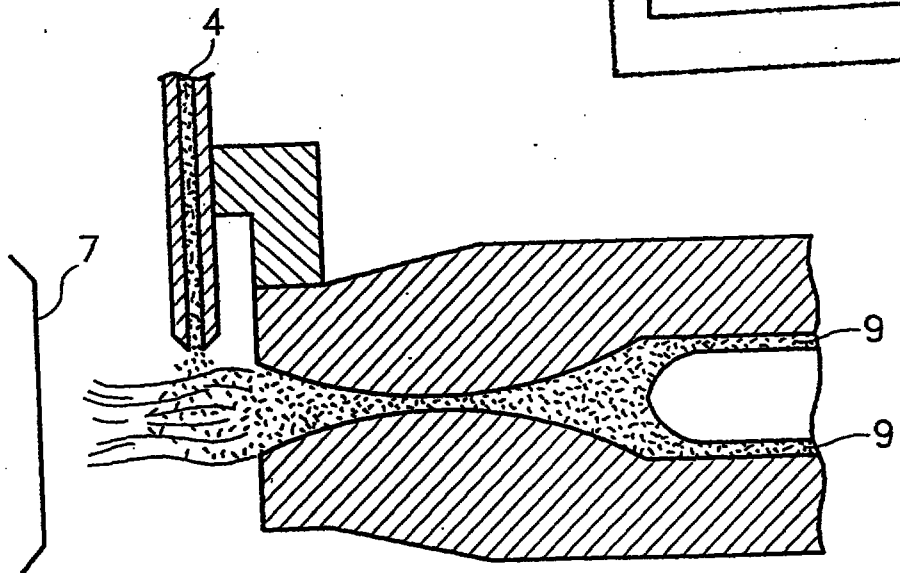



FIG. 17

REPLACEMENT SHEET
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REPLACEMENT SHEET

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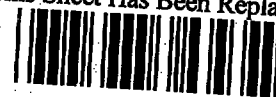


FIG.18

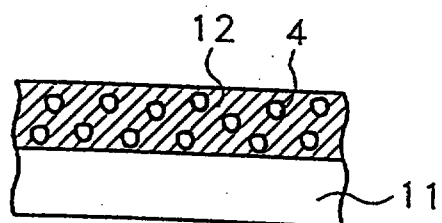


FIG.19

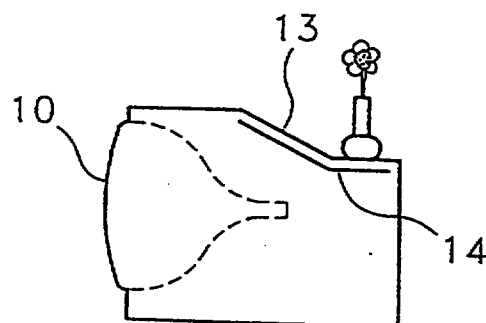
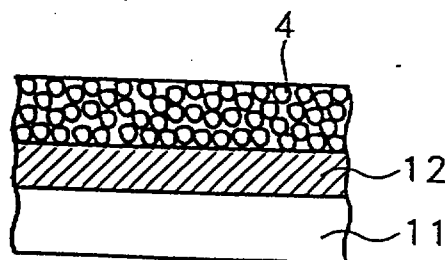


FIG.20



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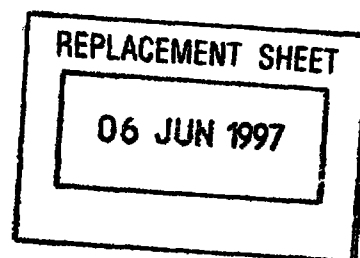


FIG.1

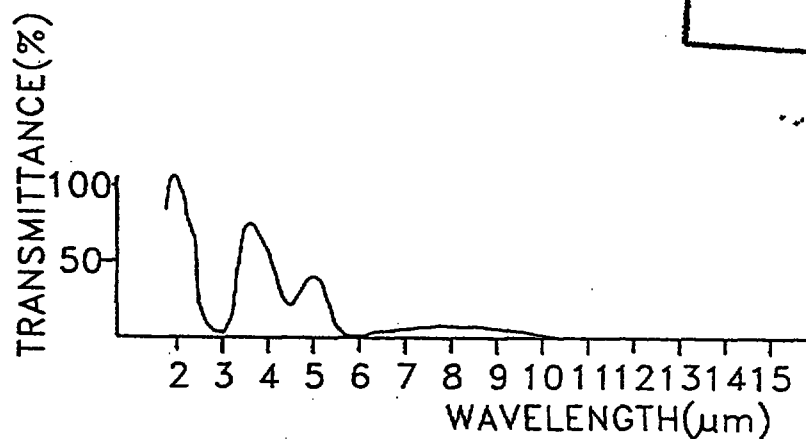


FIG.2A

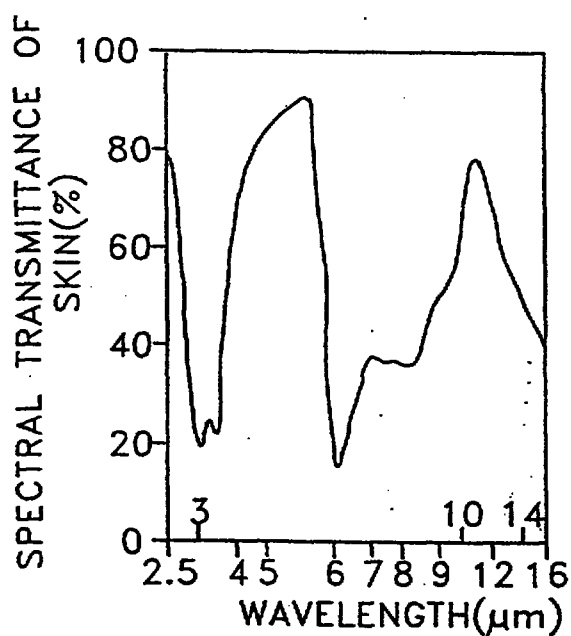
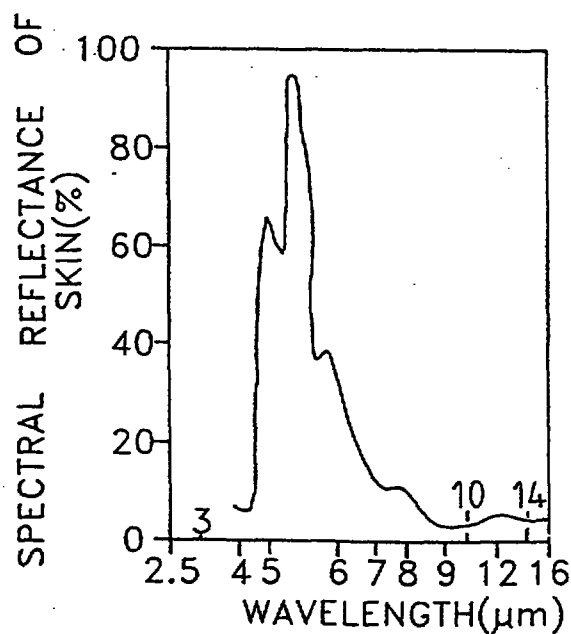


FIG.2B



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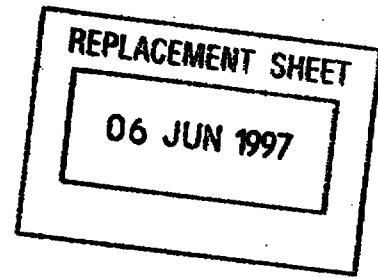


FIG.3A

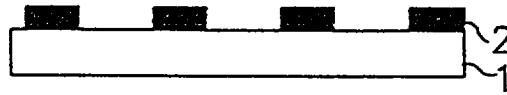


FIG.3B



FIG.3C



FIG.3D

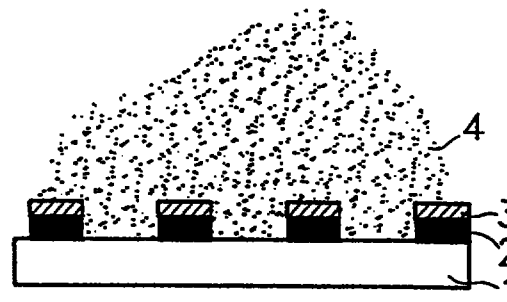


FIG.3E



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FIG.3F

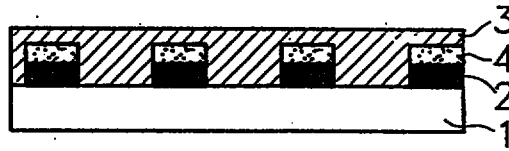


FIG.3G



FIG.3H

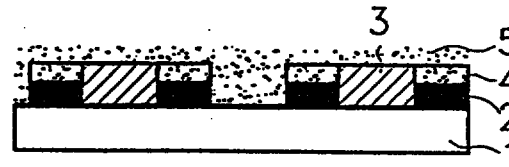
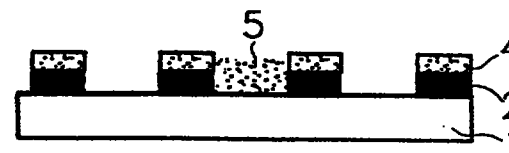


FIG.3I



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FIG.4A

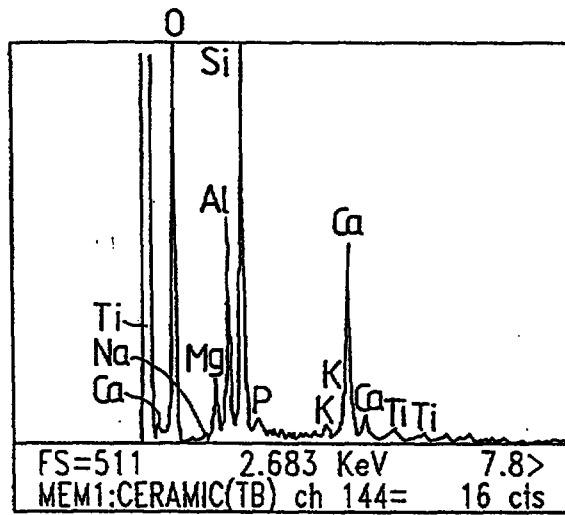


FIG.4B

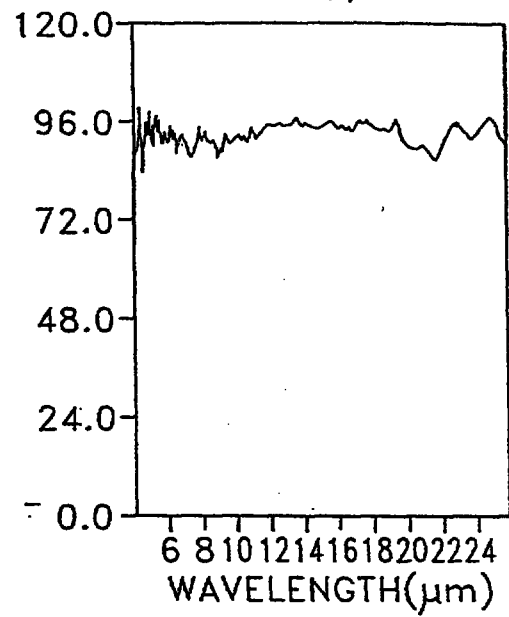
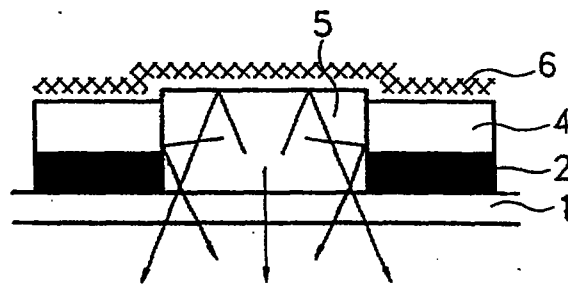


FIG.5



06 JUN 1997

FIG.6A

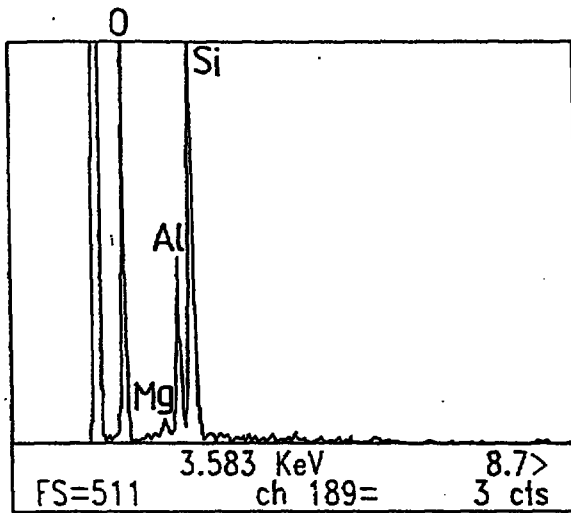


FIG.6B

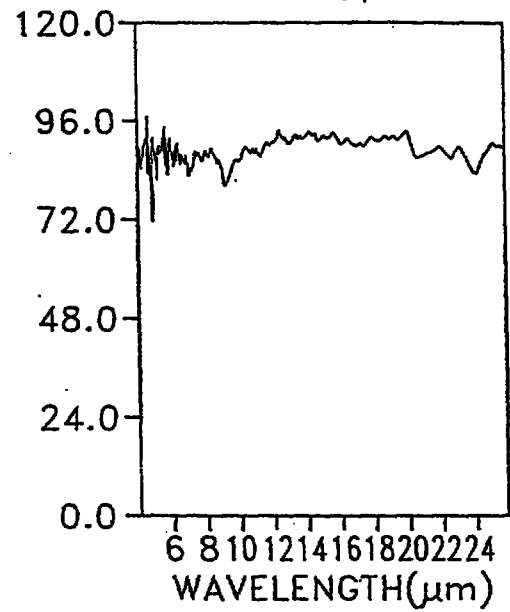
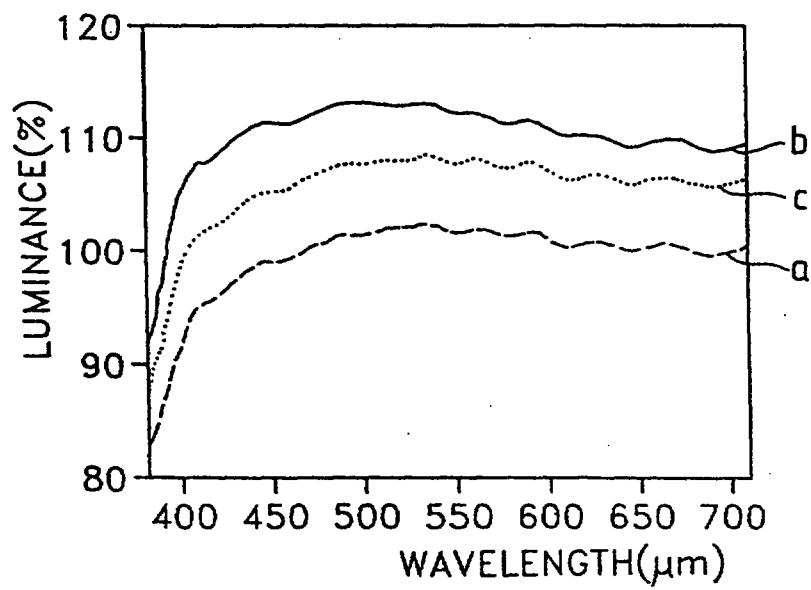


FIG.7



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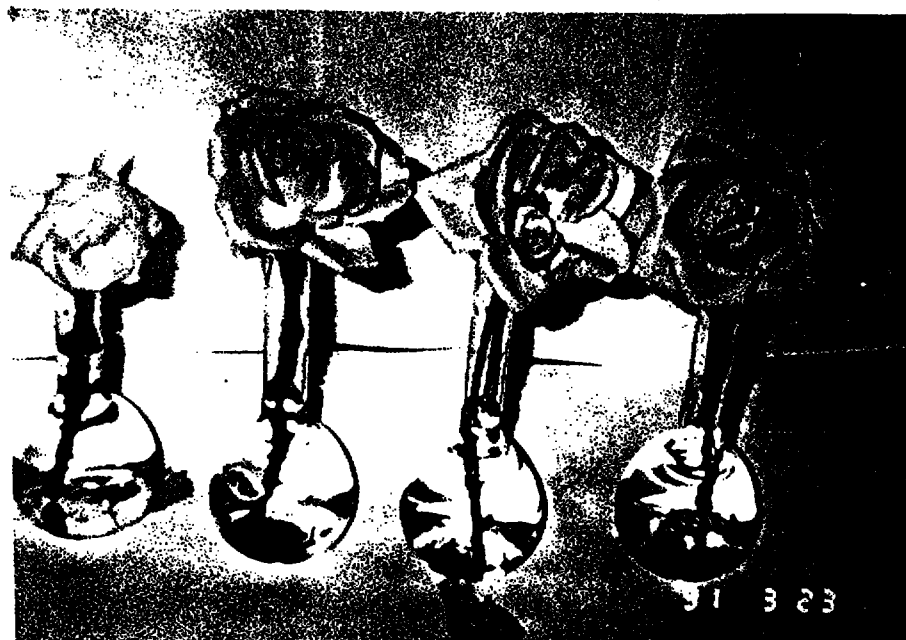
REPLACEMENT SHEET

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FIG. 8



FIG. 9A FIG. 9B FIG. 9C FIG. 9D



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FIG.10A

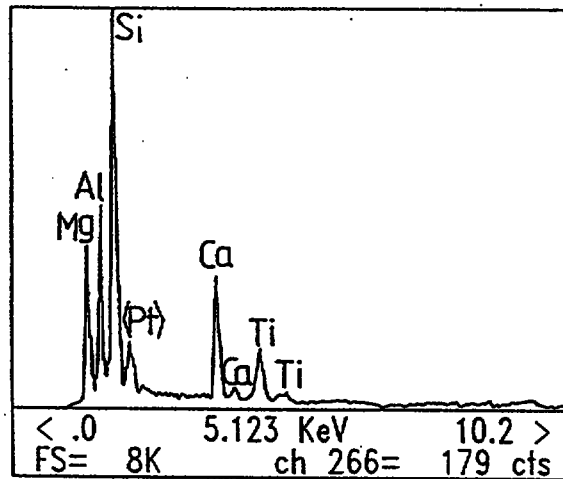
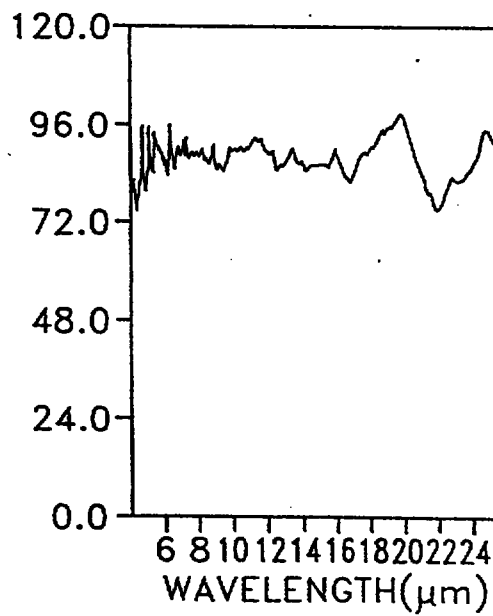


FIG.10B



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REPLACEMENT SHEET

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FIG. 11

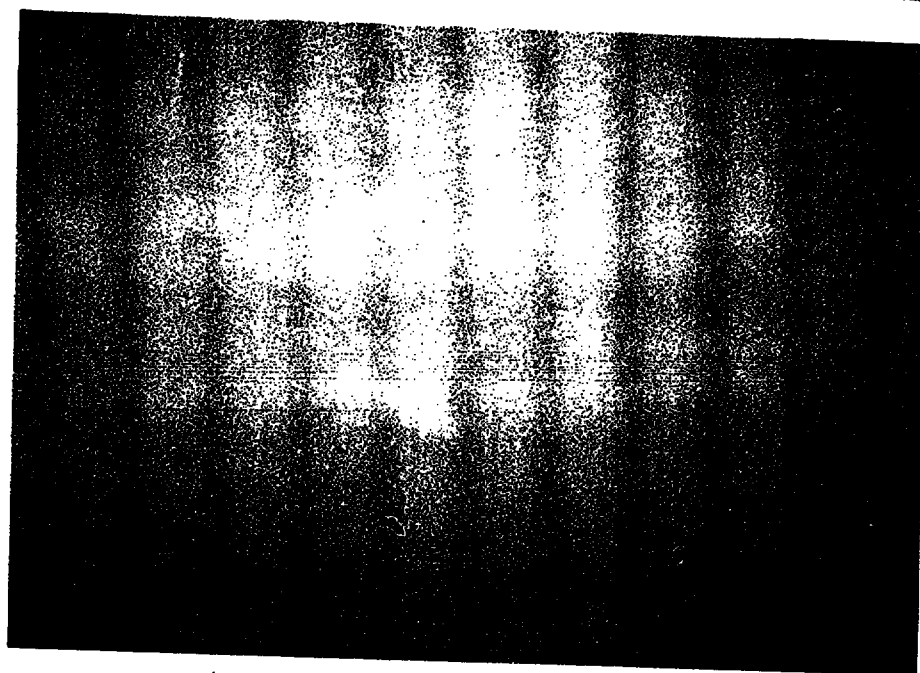
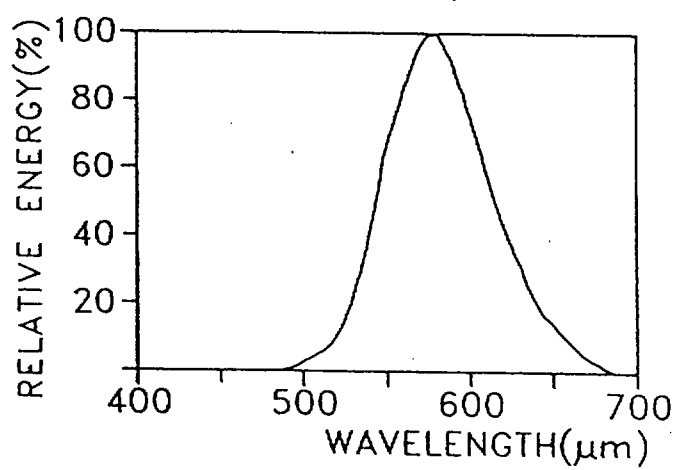


FIG. 12



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FIG.13

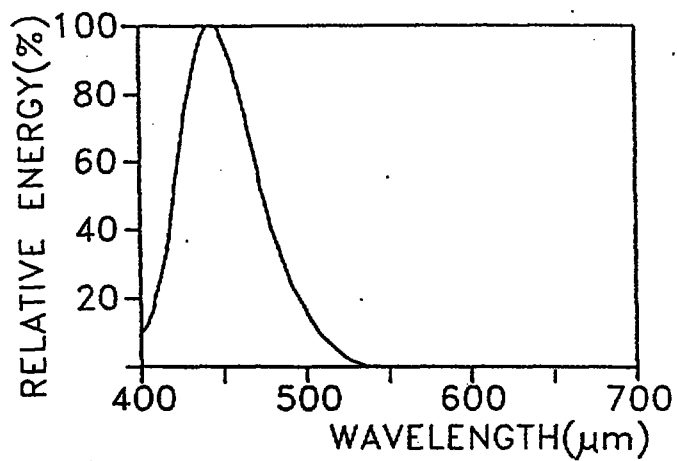
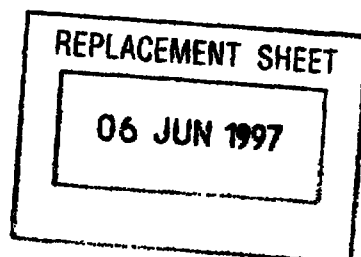
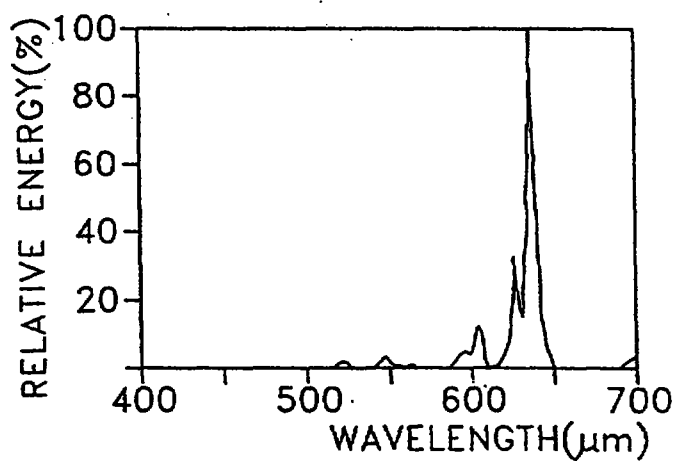


FIG.14



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REPLACEMENT SHEET

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FIG.15A

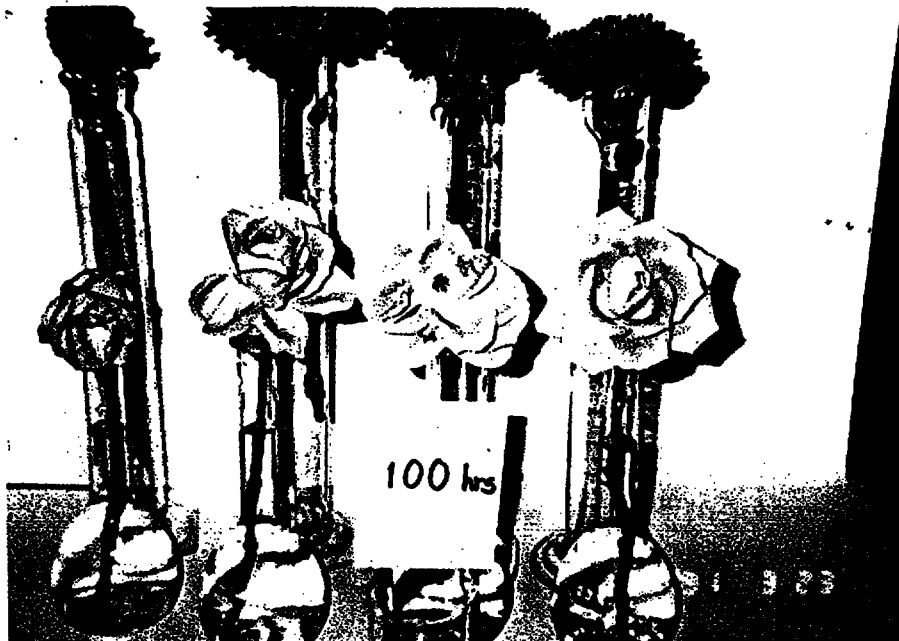


FIG.15B



FIG. 16

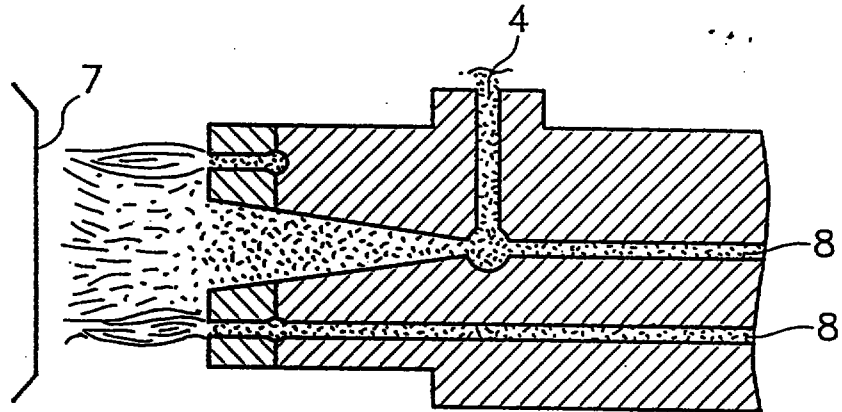
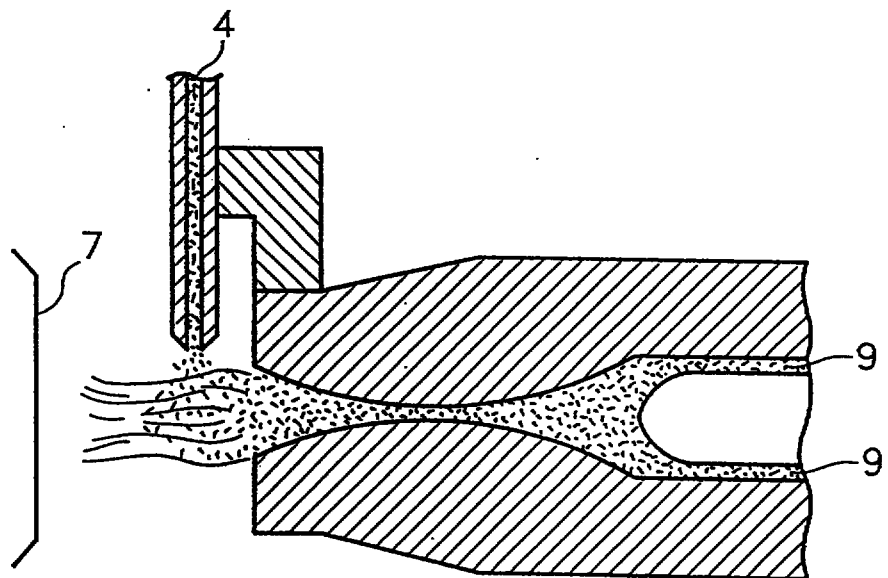


FIG. 17



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FIG.18

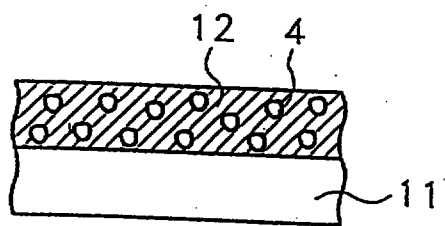


FIG.19...

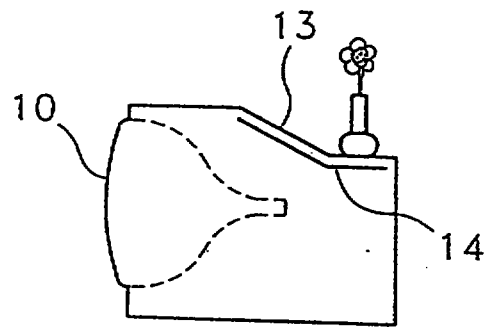
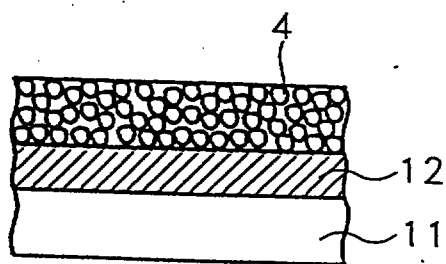


FIG.20



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