

Description

The invention relates to a scandate cathode having a cathode body comprising a matrix of at least a high melting-point metal and/or alloy, and a barium compound in contact with the matrix material for supplying barium to the emissive surface by chemical reaction with the matrix material.

The invention also relates to methods of manufacturing such a cathode and to an electron beam tube comprising such a cathode.

Cathodes of the type mentioned above are described in the Article "properties and manufacture of top-layer scandate cathodes", Applied Surface Science 26 (1986), 173-195, J. Hasker, J. v. Esdonk and J.E. Crombeen. In the cathodes described in this Article scandium oxide (Sc_2O_3) grains of several microns or tungsten (W) grains which are partially coated with either scandium (Sc) or scandium hydride (ScH_2) are processed at least in the top layer of the cathode body. The cathode body is manufactured by means of pressing and sintering whereafter the pores are impregnated with barium-calcium-aluminate. By chemical reaction with the tungsten of the matrix during operation of the cathode, the barium-calcium-aluminate supplies barium to the emissive surface in order to maintain the electron emission. To be able to realize a very high cathode load after assembly in, for example, a cathode ray tube and activation of the cathode, it is important that a scandium-containing layer having a thickness of some monolayers has formed on the cathode surface during impregnation by reaction with the impregnant. To this end the impregnation process must be performed very carefully. As compared with an impregnated tungsten cathode, which is either or not coated with, for example osmium, this may be considered a drawback.

As has been proved by experiments described in the abovementioned Article, an ion bombardment which may occur in practice, for example during the manufacture of television tubes, may entirely or partly remove the scandium containing layer, with the attendant detrimental results for the emission. Since Sc_2O_3 is not very mobile (in the cathodes manufactured by means of W partially coated with Sc or ScH_2 whole or partial oxidation of the scandium occurs during impregnation as described for cathodes manufactured with W partially coated with ScH_2 in US-A-4 594 220) the said scandium-containing layer cannot be fully regenerated by reactivating the cathode. According to the experiments described a regeneration sufficient for a complete recovery of the emission was not achieved. As compared with an impregnated tungsten cathode, this may also be considered a drawback.

The object of the invention is to provide scandate cathodes which are improved with respect to the drawbacks mentioned hereinbefore. The invention is based on the recognition that this can be achieved by using scandium-containing materials which segregate scandium to their surface upon heating. Due to the relatively low surface energy of scandium there are scandium metal compounds and scandium alloys which exhibit this scandium segregation. At an elevated temperature in vacuo a monolayer of scandium is deposited on the surface of these compounds and alloys. After removal of this layer - by means of ion bombardment or another process - a new layer of scandium will be deposited on the surface at a sufficiently high temperature. This can of course be repeated until the scandium is depleted.

To this end a scandate cathode according to the invention is characterized in that at least the top layer of the cathode body comprises a scandium metal compound or scandium alloy for exhibiting scandium segregation for depositing a layer of scandium on the surface of these metal compounds or alloys.

The speed at which the scandium is dispensed to the emissive surface may also depend on chemical reactions between the barium compound used and the source supplying scandium.

The metal compound or alloy preferably yields scandium already at the operating temperature of the cathode, but this is not absolutely necessary. If the scandium is dispensed at a higher temperature, the emission may decrease during operation due to evaporation and/or ion bombardment, but then it can in principle be restored by reactivating the cathode at a sufficiently high temperature. The scandium may also segregate if the temperature becomes high enough during the manufacture (for example during impregnation).

Notably metal compounds and/or alloys of scandium comprising one or more of the metals rhenium (Re), ruthenium (Ru), hafnium (Hf), nickel (Ni), cobalt (Co), palladium (Pd), zirconium (Zr) or tungsten (W) were found to be satisfactory.

In US-A-4 626 470 a thin porous film composed of a refractory metal and scandium or scandiumoxide is formed on an electron-emissive surface of a cathode body to prevent damage due to ion-bombardment. Although alloys of the refractory metals may be used no scandium-segregating metal compound or alloy for depositing a layer of scandium on the surface of these metal compounds or alloys is disclosed.

Due to the high melting point and the fact that rhenium or ruthenium do not evaporate during operation and manufacture, $\text{Re}_{24}\text{Sc}_5$, Re_2Sc and Ru_2Sc are extremely suitable, particularly the rhenium compounds because they exhibit scan-

dium segregation already at the operating temperature.

A first method of manufacturing a scandate cathode according to the invention as claimed in claim 8 is characterized in that a porous body comprising the scandium metal compound or scandium alloy at least in the top layer is obtained by means of mixing, pressing and sintering powders of a high melting-point metal and/or alloy and of a scandium metal compound or scandium alloy which can exhibit scandium segregation, whereafter said body is at least partly provided by means of impregnation with a barium compound which can supply barium to the emissive surface by chemical reaction with the high melting-point metal and/or alloy.

Another method as claimed in claim 9 is characterized in that the cathode body comprising in at least its top layer a scandium metal compound or scandium alloy which can exhibit scandium segregation is obtained by means of mixing, pressing and sintering powders of a high melting-point metal and/or alloy and of the scandium metal compound or scandium alloy combined with the powder of a barium compound which can supply barium to the emissive surface by chemical reaction with the high melting-point metal and/or alloy during operation of the cathode. In this method the sintering temperature is the highest temperature the cathode body ever acquires. This temperature may be substantially lower than the impregnation temperature which is generally used in the previous method. Consequently, the reaction of the barium compound with the scandium metal compound or scandium alloy is reduced. In fact, a too vigorous reaction may give rise to a considerable scandium oxidation so that the supply of scandium is reduced.

The invention will now be described in greater detail, by way of example, with reference to the accompanying drawing in which

Figure 1 shows diagrammatically an experimental set-up for testing scandium metal compounds and alloys,

Figure 2 shows a result of measurements on a scandium metal compound,

Figure 3 is a diagrammatic representation of a cathode according to the invention, and

Figure 4 is a diagrammatic representation of another cathode according to the invention.

Figure 1 is a longitudinal sectional view of an experimental set-up. A pulverulent scandium metal compound or scandium alloy 2 is pressed and sintered in the molybdenum tray 1. Subsequently, it is welded onto the shaft 3 comprising a heating element 4. The assembly is mounted in a Scanning Auger Microscope to measure the scandium concentration on the surface. This concentration

can be reduced by means of ion bombardment and it may increase again after this bombardment due to scandium segregation. In this way various scandium metal compounds and scandium alloys have been tested, such as $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{43}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ and $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.

Figure 2 shows a measuring result for the metal compound $\text{Re}_{24}\text{Sc}_5$. The measurement indicated by curve a is considered first. Prior to the instant $t = 0$ in the Figure the experimental set-up had been at a temperature of 950°C for some time and this temperature was also maintained during the measurement. At the instant $t = 0$ approximately one mono-layer of scandium is then present on the surface - the experimental set-up was exposed to an ion bombardment. Consequently, the scandium concentration on the surface decreased until at $t = t_1$ a balance was achieved between the supply and removal of scandium. After the ion bombardment had been switched off at $t = t_2$ the original concentration was achieved again in a short time by scandium segregation. No scandium depletion was observed when the experiment was repeated several times. Curve b shows a similar result measured on the same experimental set-up at a temperature of 1100°C . The balance during bombardment was set at a higher concentration than at 950°C . Scandium depletion was neither observed when the experiment was repeated several times. Another result of the investigations was that the metal compound Ru_2Sc at the operating temperature (approximately 950°C) or the usual temperature for activating a scandate cathode (approximately 1100°C) did not exhibit any scandium segregation.

Figure 3 is a longitudinal sectional view of a scandate cathode according to the invention. The cathode body 13 has a top layer 23 and an emissive surface 33. This body, having a diameter of 1.8 mm, is obtained by pressing a matrix of W powder with a top layer on it comprising a mixture of W powder and a powder of a scandium metal compound or scandium alloy according to the invention. After pressing a sintering operation is carried out at 1500°C in a hydrogen atmosphere. The thickness of the matrix is then approximately 0.5 mm and that of the top layer is approximately 0.1 mm. The pressure during pressing of the cathode body is such that the increase in weight is substantially 4.5% after impregnation with $4\text{BaO} - 1\text{CaO} - 1\text{Al}_2\text{O}_3$ in a hydrogen atmosphere. The impregnated cathode body, either or not provided with an envelope 43, is welded onto the cathode shaft 53. A coiled cathode filament 63 which may consist of a helically wound metal core 73 with an aluminium oxide insulation layer 83 is present in the shaft 53. The emission of such a cathode, after

assembly and activation, is measured in a diode arrangement with a cathode-anode gap of 0.3 mm at a 1000 Volt pulse load. Cathodes were manufactured as examples with top layers consisting of W with 25 and 50% by weight of Re_2Sc and with top layers consisting of W with 10 and 25% by weight of $\text{Re}_{24}\text{Sc}_5$. In all cases the measured emission was substantially 100 A/cm² at an operating temperature of approximately 950°C. In another example the top layer consisted of W with 10 and 25% by weight of Ru_2Sc . The emission was again substantially 100 A/cm² but, unlike the previous examples, it exhibited a decrease of approximately 30% after 8000 hours of a continuous load of 1.5 A/cm². In yet another example the top layer consisted of W with 5, 10 and 20% by weight of $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$. The measured emission varied between approximately 70 and 90 A/cm². The above examples show that the high emissions characteristic of scandate cathodes can be realized by using scandium metal compounds or scandium alloys according to the invention.

Figure 4 is a longitudinal sectional view of another scandate cathode according to the invention. The cathode body 14 has an emissive surface 24. This body, with a diameter of 1.8 mm and a thickness of approximately 0.5 mm is obtained by pressing a mixture of W powder and 10% by weight of $\text{Re}_{24}\text{Sc}_5$ powder and 7% by weight of barium-calcium-aluminate powder ($4\text{BaO}-1\text{CaO}-1\text{Al}_2\text{O}_3$) and by subsequently sintering at 1500°C in a hydrogen atmosphere. The cathode body, either or not provided with a molybdenum envelope 34, is then welded onto the cathode shaft 44. The shaft 44 accommodates a coiled filament 54 which may consist of a helically wound metal core 64 having an aluminium oxide insulation layer

1. The measured emission after activation was approximately 100 A/cm² at a cathode temperature of 950°C. An advantage of this cathode is the simple method of its manufacture: impregnation and subsequent cleaning is not necessary. Auger measurements have proved that the scandium concentration on the surface is very low before activation. During activation, as described in the Article mentioned in the opening paragraph, the scandium concentration required for the measured emission is formed on the surface.

The invention is of course not limited to the examples shown, but several variations within the scope of the invention are possible to those skilled in the art. The emissive material may be present in a storage space under the 6 actual matrix (L-cathode), whilst many design variations are also possible. Moreover, the barium supply to the emissive surface is not necessarily confined to the mechanism described herein but can also be imagined to originate e.g. from segregation from

barium compounds or alloys because the surface energy of barium is lower than that of scandium.

Claims

1. A scandate cathode having a cathode body comprising a matrix of at least a high melting-point metal and/or alloy, and a barium compound in contact with the matrix material for supplying barium to the emissive surface by chemical reaction with the matrix material, characterized in that at least the top layer of the cathode body comprises a scandium metal compound or scandium alloy for exhibiting scandium segregation for depositing a layer of scandium on the surface of these metal compounds or alloys.
2. A scandate cathode as claimed in Claim 1, characterized in that the scandium metal compound or scandium alloy is adapted for exhibiting scandium segregation at the operating temperature of the cathode.
3. A scandate cathode as claimed in Claim 1, characterized in that the scandium metal compound or scandium alloy is adapted for exhibiting scandium segregation at an activation temperature which is higher than the operating temperature of the cathode.
4. A scandate cathode as claimed in Claim 1, characterized in that the scandium metal compound or scandium alloy is adapted for exhibiting scandium segregation at a temperature to which the cathode is subjected during one of its manufacturing steps.
5. A scandate cathode as claimed in any one of the preceding Claims, characterized in that the scandium metal compound or scandium alloy is a compound of scandium with one or more of the metals rhenium (Re), ruthenium (Ru), hafnium (Hf), nickel (Ni), cobalt (Co), palladium (Pd), zirconium (Zr) or tungsten (W).
6. A scandate cathode as claimed in Claim 5, characterized in that the scandium metal compound or scandium alloy is $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{43}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ or $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.
7. A scandate cathode as claimed in Claim 5, characterized in that at least the top layer of the cathode body comprises 5 to 50% by weight of Re_2Sc or $\text{Re}_{24}\text{Sc}_5$.



assembly and activation, is measured in a diode arrangement with a cathode-anode gap of 0.3 mm at a 1000 Volt pulse load. Cathodes were manufactured as examples with top layers consisting of W with 25 and 50% by weight of Re_2Sc and with top layers consisting of W with 10 and 25% by weight of $\text{Re}_{24}\text{Sc}_5$. In all cases the measured emission was substantially 100 A/cm² at an operating temperature of approximately 950°C. In another example the top layer consisted of W with 10 and 25% by weight of Ru_2Sc . The emission was again substantially 100 A/cm² but, unlike the previous examples, it exhibited a decrease of approximately 30% after 8000 hours of a continuous load of 1.5 A/cm². In yet another example the top layer consisted of W with 5, 10 and 20% by weight of $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$. The measured emission varied between approximately 70 and 90 A/cm². The above examples show that the high emissions characteristic of scandate cathodes can be realized by using scandium metal compounds or scandium alloys according to the invention.

Figure 4 is a longitudinal sectional view of another scandate cathode according to the invention. The cathode body 14 has an emissive surface 24. This body, with a diameter of 1.8 mm and a thickness of approximately 0.5 mm is obtained by pressing a mixture of W powder and 10% by weight of $\text{Re}_{24}\text{Sc}_5$ powder and 7% by weight of barium-calcium-aluminate powder ($4\text{BaO}-1\text{CaO}-1\text{Al}_2\text{O}_3$) and by subsequently sintering at 1500°C in a hydrogen atmosphere. The cathode body, either or not provided with a molybdenum envelope 34, is then welded onto the cathode shaft 44. The shaft 44 accommodates a coiled filament 54 which may consist of a helically wound metal core 64 having an aluminium oxide insulation layer

1. The measured emission after activation was approximately 100 A/cm² at a cathode temperature of 950°C. An advantage of this cathode is the simple method of its manufacture: impregnation and subsequent cleaning is not necessary. Auger measurements have proved that the scandium concentration on the surface is very low before activation. During activation, as described in the Article mentioned in the opening paragraph, the scandium concentration required for the measured emission is formed on the surface.

The invention is of course not limited to the examples shown, but several variations within the scope of the invention are possible to those skilled in the art. The emissive material may be present in a storage space under the 6 actual matrix (L-cathode), whilst many design variations are also possible. Moreover, the barium supply to the emissive surface is not necessarily confined to the mechanism described herein but can also be imagined to originate e.g. from segregation from

barium compounds or alloys because the surface energy of barium is lower than that of scandium.

Claims

1. A scandate cathode having a cathode body comprising a matrix of at least a high melting-point metal and/or alloy, and a barium compound in contact with the matrix material for supplying barium to the emissive surface by chemical reaction with the matrix material, characterized in that at least the top layer of the cathode body comprises a scandium metal compound or scandium alloy for exhibiting scandium segregation for depositing a layer of scandium on the surface of these metal compounds or alloys.
2. A scandate cathode as claimed in Claim 1, characterized in that the scandium metal compound or scandium alloy is adapted for exhibiting scandium segregation at the operating temperature of the cathode.
3. A scandate cathode as claimed in Claim 1, characterized in that the scandium metal compound or scandium alloy is adapted for exhibiting scandium segregation at an activation temperature which is higher than the operating temperature of the cathode.
4. A scandate cathode as claimed in Claim 1, characterized in that the scandium metal compound or scandium alloy is adapted for exhibiting scandium segregation at a temperature to which the cathode is subjected during one of its manufacturing steps.
5. A scandate cathode as claimed in any one of the preceding Claims, characterized in that the scandium metal compound or scandium alloy is a compound of scandium with one or more of the metals rhenium (Re), ruthenium (Ru), hafnium (Hf), nickel (Ni), cobalt (Co), palladium (Pd), zirconium (Zr) or tungsten (W).
6. A scandate cathode as claimed in Claim 5, characterized in that the scandium metal compound or scandium alloy is $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{43}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ or $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.
7. A scandate cathode as claimed in Claim 5, characterized in that at least the top layer of the cathode body comprises 5 to 50% by weight of Re_2Sc or $\text{Re}_{24}\text{Sc}_5$.



8. A method of manufacturing a scandate cathode, characterized in that a porous body comprising a scandium metal compound or scandium alloy at least in the top layer is obtained by means of mixing, pressing and sintering powders of a high melting-point metal and/or alloy and of a scandium metal compound or scandium alloy which can exhibit scandium segregation for depositing a layer of scandium on the surface of these metal compounds or alloys, whereafter said body is at least partly provided by means of impregnation with a barium compound which can supply barium to the emissive surface by chemical reaction with the high melting-point metal and/or alloy.

9. A method of manufacturing a scandate cathode characterized in that the cathode body comprising in at least its top layer a scandium metal compound or scandium alloy which can exhibit scandium segregation for depositing a layer of scandium on the surface of these metal compounds or alloys is obtained by means of mixing, pressing and sintering powders of a high melting-point metal and/or alloy and of the scandium metal compound or scandium alloy combined with the powder of a barium compound which can supply barium to the emissive surface by chemical reaction with the high melting-point metal and/or alloy during operation of the cathode.

10. A method as claimed in Claim 8 or 9, characterized in that the scandium metal compound or scandium alloy is a compound or alloy comprising one or more of the metals rhenium (Re), ruthenium (Ru), hafnium (Hf), nickel (Ni), cobalt (Co), palladium (Pd), zirconium (Zr) or tungsten (W).

11. A method as claimed in Claim 10, characterized in that the scandium metal compound or scandium alloy is $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{49}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ or $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.

12. A method as claimed in Claim 10, characterized in that at least the top layer of the cathode body comprises 5 to 50% by weight of Re_2Sc or $\text{Re}_{24}\text{Sc}_5$.

13. An electron beam tube provided with a cathode as claimed in any one of Claims 1 to 7.

Patentansprüche

1. Scandatkathode mit einem Kathodenkörper mit einer Matrix aus wenigstens einem hochschmelzenden Metall und/oder einer hochschmelzenden Legierung und mit einer Bariumverbindung im Kontakt mit dem Matrixmaterial zum Liefern von Barium nach der emittierenden Oberfläche durch chemische Reaktion mit dem Matrixmaterial, dadurch gekennzeichnet, daß wenigstens die Oberschicht des Kathodenkörpers eine Scandiummetallverbindung oder eine Scandiumlegierung zum Aufweisen von Scandiumsegregation zum Ablagern einer Scandiumschicht auf der Oberfläche dieser Metallverbindungen oder Legierungen enthält.

2. Scandatkathode nach Anspruch 1, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder die Scandiumlegierung zum Aufweisen von Scandiumsegregation bei der Betriebstemperatur der Kathode ausgelegt ist.

3. Scandatkathode nach Anspruch 1, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder die Scandiumlegierung zum Aufweisen von Scandiumsegregation bei einer Aktivierungstemperatur ausgelegt ist, die höher ist als die Betriebstemperatur der Kathode.

4. Scandatkathode nach Anspruch 1, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder die Scandiumlegierung zum Aufweisen von Scandiumsegregation bei einer Temperatur ausgelegt ist, der die Kathode bei einem seiner Herstellungsschritte unterworfen ist.

5. Scandatkathode nach einem oder mehreren der vorangehenden Ansprüche, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder die Scandiumlegierung eine Verbindung aus Scandium mit einem oder mehreren der Metalle Rhenium (Re), Ruthenium (Ru), Hafnium (Hf), Nickel (Ni), Kobalt (Co), Palladium (Pd), Zirkon (Zr) oder Wolfram (W) ist.

6. Scandatkathode nach Anspruch 5, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder Scandiumlegierung wie folgt aussieht $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{49}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ oder $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.



7. Scandatkathode nach Anspruch 5, dadurch gekennzeichnet, daß wenigstens die Ober-
schicht des Kathodenkörpers 5 bis 50 Gew. %
von Re_2Sc oder $\text{Re}_{24}\text{Sc}_5$ enthält.
8. Verfahren zum Herstellen einer Scandatkathode, dadurch gekennzeichnet, daß ein poroser Körper mit wenigstens in der Oberschicht einer Scandiummetallverbindung oder einer Scandiumlegierung erhalten wird durch Mischen, Pressen und Sintern von Pulvern eines hochschmelzenden Metalls und/oder einer hochschmelzenden Legierung und einer Scandiummetallverbindung oder Scandiumlegierung, die Scandiumsegregierung zum Ablagern einer Scandiumschicht auf der Oberfläche dieser Metallverbindungen oder Legierungen aufweisen kann, wonach der Körper wenigstens teilweise mittels Imprägnieren mit einer Bariumverbindung versehen wird, die durch chemische Reaktion mit dem hochschmelzenden Metall und/oder der hochschmelzenden Legierung Barium nach der emittierenden Oberfläche liefern kann.
9. Verfahren zum Herstellen einer Scandatkathode, dadurch gekennzeichnet, daß der Kathodenkörper, der wenigstens in seinem Oberschicht eine Scandiummetallverbindung oder Scandiumlegierung enthält, die Scandiumsegregierung zum Ablagern einer Scandiumschicht auf der Oberfläche dieser Metallverbindungen oder Legierungen aufweisen kann, erhalten wird durch Mischen, Pressen und Sintern von Pulvern eines hochschmelzenden Metalls und/oder einer hochschmelzenden Legierung und der Scandiummetallverbindung oder Scandiumlegierung in Verbindung mit dem Pulver einer Bariumverbindung, die durch chemische Reaktion mit dem hochschmelzenden Metall und/oder der hochschmelzenden Legierung im Betrieb der Kathode Barium nach der emittierenden Oberfläche liefern kann.
10. Verfahren nach Anspruch 8 oder 9, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder Scandiumlegierung eine Verbindung oder Legierung mit einem oder mehreren der Metalle Rhenium (Re), Ruthenium (Ru), Hafnium (Hf), Nickel (Ni), Kobalt (Co), Palladium (Pd), Zirkon (Zr) oder Wolfram (W) ist.
11. Verfahren nach Anspruch 10, dadurch gekennzeichnet, daß die Scandiummetallverbindung oder die Scandiumlegierung die folgende ist $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc ,

Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{43}\text{W}_7$, $\text{Sc}_{58}\text{Hf}_{24}\text{W}_8$ oder $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.

12. Verfahren nach Anspruch 10, dadurch gekennzeichnet, daß wenigstens die Oberschicht des Kathodenkörpers 5 bis 50 Gewichtsprozent an Re_2Sc oder $\text{Re}_{24}\text{Sc}_5$ enthält.

13. Elektronenstrahlröhre mit einer Kathode nach einem oder mehreren der Ansprüche 1 bis 7.

Revendications

1. Cathode à scandate comportant un corps de cathode comprenant une matrice d'au moins un métal et/ou alliage à haut point de fusion, et un composé du baryum en contact avec la matière de matrice pour fournir du baryum à la surface émissive par réaction chimique avec la matière de matrice, caractérisée en ce qu'au moins la couche supérieure du corps de cathode comprend un composé métallique de scandium ou alliage de scandium pour manifester la ségrégation du scandium en vue de déposer une couche de scandium sur la surface de ces composés métalliques ou alliages.
2. Cathode à scandate suivant la revendication 1, caractérisée en ce que le composé métallique de scandium ou alliage de scandium est à même de manifester la ségrégation du scandium à la température de fonctionnement de la cathode.
3. Cathode à scandate suivant la revendication 1, caractérisée en ce que le composé métallique de scandium ou alliage de scandium est à même de manifester la ségrégation du scandium à une température d'activation qui est supérieure à la température de fonctionnement de la cathode.
4. Cathode à scandate suivant la revendication 1, caractérisée en ce que le composé métallique de scandium ou alliage de scandium est à même de manifester la ségrégation du scandium à une température à laquelle la cathode est soumise pendant l'un de ses stades de fabrication.
5. Cathode à scandate suivant l'une quelconque des revendications précédentes, caractérisée en ce que le composé métallique de scandium ou alliage de scandium est un composé du scandium avec l'un des métaux rhénium (Re), ruthénium (Ru), hafnium (Hf), nickel (Ni), cobalt (Co), palladium (Pd), zirconium (Zr) et tungstène (W).



6. Cathode à scandate suivant la revendication 5, caractérisée en ce que le composé métallique du scandium ou alliage de scandium est $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{43}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ ou $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.
7. Cathode à scandate suivant la revendication 5, caractérisée en ce qu'au moins la couche supérieure du corps de cathode comprend 5 à 50% en poids de Re_2Sc ou de $\text{Re}_{24}\text{Sc}_5$.
8. Procédé de fabrication d'une cathode à scandate, caractérisé en ce qu'un corps poreux comprenant un composé métallique de scandium ou alliage de scandium au moins dans la couche supérieure est obtenu par mélange, pressage et frittage de poudres d'un métal et/ou alliage à haut point de fusion et d'un composé métallique de scandium ou alliage de scandium qui peut manifester la ségrégation du scandium en vue de déposer une couche de scandium sur la surface de ces composés métalliques ou alliages, après quoi le corps est au moins partiellement pourvu, par imprégnation, d'un composé de baryum qui peut fournir du baryum à la surface émissive par réaction chimique avec le métal et/ou alliage à haut point de fusion.
9. Procédé de fabrication d'une cathode à scandate, caractérisé en ce que le corps de cathode comprenant, au moins dans sa couche supérieure, un composé métallique de scandium ou alliage de scandium qui peut manifester la ségrégation du scandium en vue de déposer une couche de scandium sur la surface de ces composés métalliques ou alliages, est obtenu par mélange, pressage et frittage de poudres d'un métal et/ou alliage à haut point de fusion et du composé métallique de scandium ou alliage de scandium combiné avec la poudre d'un composé de baryum qui peut fournir du baryum à la surface émissive par réaction chimique avec le métal et/ou alliage à haut point de fusion, pendant le fonctionnement de la cathode.
10. Procédé suivant la revendication 8 ou 9, caractérisé en ce que le composé métallique du scandium ou alliage de scandium est un composé ou alliage comprenant un ou plusieurs des métaux rhénium (Re), ruthénium (Ru), hafnium (Hf), nickel (Ni), cobalt (Co), palladium (Pd), zirconium (Zr) ou tungstène (W).
11. Procédé suivant la revendication 10, caractérisé en ce que le composé métallique de scandium ou alliage de scandium est $\text{Re}_{24}\text{Sc}_5$, Re_2Sc , Ru_2Sc , Co_2Sc , Pd_2Sc , Ni_2Sc , $\text{Sc}_{50}\text{Zr}_{43}\text{W}_7$, $\text{Sc}_{68}\text{Hf}_{24}\text{W}_8$ ou $\text{Sc}_{47}\text{Hf}_{41}\text{W}_{12}$.
12. Procédé suivant la revendication 10, caractérisé en ce qu'au moins la couche supérieure du corps de cathode comprend 5 à 50% en poids de Re_2Sc ou $\text{Re}_{24}\text{Sc}_5$.
13. Tube à faisceau d'électrons muni d'une cathode suivant l'une quelconque des revendications 1 à 7.