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Description

The present invention relates to a process for producing chromium dioxide which has a high coercive force and is modified with foreign elements.

5 A number of processes are known in which ferro-magnetic chromium dioxide is produced starting from a chromium oxide or mixture of chromium oxides having a mean valency of not less than 4, the process being carried out under superatmospheric pressure and at
10 elevated temperatures in the presence of water. Moreover, it is stated in a number of publications that the magnetic properties of the end product can be improved by modifying such a chromium dioxide with a number of metals. For example, US-A 30 34 988 describes the production of an improved chromium dioxide by converting
15 chromium trioxide in the presence of a metal at atomic number 22 to 28, or of a compound of this metal, and an element of group V of the periodic table and, if required, water, at from 300 to 500°C under a pressure of up to 2940 bar. The chromium dioxides produced in this
20 manner have coercive forces of 34 kA/m at best. In another process as disclosed in NL-A 66 17 476, chromium oxides or mixtures of these in which the chromium has a mean valency of not less than 4 are treated in the presence of alkali metal dichromates and known dopants at
25 from 350 to 500°C and under from 245 to 980 bar. However, the resulting product possesses a high coercive force only when the reaction pressure is 980 bar. Moreover, a disadvantage of this process is that the starting material required is a chromium(III) oxide which is expensive
30 to produce.

Furthermore, DE-C 20 22 820 discloses that a chromium dioxide having a high coercive force can be
35 obtained if the conventional process is carried out not only with the addition of antimony, selenium, tellurium or compounds of these as a first modifier, but also with the addition of iron in the form of acicular crystals

and/or acicular iron oxide particles as a second modifier. The amount of iron used is from 0.1 to 10% by weight, and the total amount of modifier is as high as 25% by weight.

5 Although these and other similar conventional processes permit production of ferromagnetic chromium dioxides having a high coercive force, large amounts of modifiers are generally required. Although these large
10 amounts of added materials do not have an adverse effect on the magnetic properties, they result in a crystallite size which has a marked disadvantageous effect on the electroacoustic properties, particularly where chromium dioxides are used in the production of magnetic recording media.

15 According to EP-A 27 640, a further improvement can be achieved by using the cubic modification of antimony oxide, senarmontite. The magnetic properties are improved in a particularly effective manner when the specific surface area of the senarmontite is from 3 to
20 10 m²/g. However, when the process is carried out it is found that it is difficult to obtain pure senarmontite having this surface area. Another disadvantage is that, on comminuting and milling the senarmontite to give the fine powder required for the synthesis of chromium
25 dioxide, some of the senarmontite is converted to the less effective orthorhombic modification. In addition, special safety measures are necessary for handling antimony oxide in dust form.

30 It is an object of the present invention to provide a process for producing ferromagnetic chromium dioxides having a high coercive force while at the same time reducing the amount of modifiers required for this purpose. It is a particular object of the present invention to provide a modifier which can be introduced into
35 the reaction system of the chromium dioxide synthesis in very homogeneous and extremely finely divided form, and is also easy to handle.

We have found that this object is achieved, and that, surprisingly, ferromagnetic chromium dioxide can be produced by converting oxides of trivalent and hexavalent chromium under superatmospheric pressure and at elevated temperatures in the presence of water and with the addition of one or more antimony compounds such as modifiers and, if required, further additives in an amount of not more than 15% by weight, if the antimony compound used as a modifier is present in the form of a water-soluble salt and is added in an amount of less than 4% by weight, based on the resulting chromium dioxide, with the proviso that the anion of the salt-like antimony compound contains one or more organic molecular moieties which are oxidized in concentrated chromic acid.

For the purpose of the present invention, particularly suitable salt-like antimony compounds are antimony tartrate and potassium antimonyl tartrate, as small an amount as less than 2% by weight, based on the resulting chromium dioxide generally being sufficient. Appropriate amounts of from 0.1 to 1.0% by weight of the antimony compound result in an end product having particularly advantageous properties.

In an advantageous embodiment of the process according to the invention, iron, either in the form of acicular iron particles having a BET specific area of from 5 to 40 m²/g or, particularly advantageously, in the form of acicular iron oxide particles, such as α - or γ -iron oxide hydroxides, magnetite or α - or γ -iron(III) oxides, are used as a further modifier, in addition to the antimony compound employed as a modifier. The added amounts are from 0.4 to 10% by weight for the acicular iron particles, and from 0.5 to 15% by weight for the acicular iron oxide particles.

The invention is based on conventional processes for producing ferromagnetic chromium dioxide under superatmospheric pressure and at elevated temperatures in the presence of water. For example, in the present

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process water is initially taken in a reaction vessel, chromic acid is introduced while stirring, and the modifiers according to the novel embodiment of this process are added. When the evolution of gas, which takes place after the addition, has ceased, chromium(III) oxide is then added, with further stirring, in the stoichiometric amount required for the synproportionation reaction. Formation of the chromium dioxide occurs during the subsequent treatment of the mixture in a high pressure reactor under from 100 to 700 bar and at from 200 to 600°C. After from 10 to 50 hours, the chromium dioxide is removed mechanically from the reaction vessel and, if necessary, its surface is stabilized chemically in a conventional manner by the action of a reducing agent. The chromium dioxide thus produced consists of acicular particles having a mean particle length of from 0.1 to 2.0 μm , in particular from 0.2 to 0.9 μm , a length/width ratio of from 15:1 to 5:1, and a BET specific surface area of from 10 to 50 m^2/g .

Surprisingly, the modification according to the invention gives a chromium dioxide having a coercive force greater than 35 kA/m, in spite of the small amount of added antimony compound, ie. less than 4.0, preferably from 0.1 to 1.0% by weight, and a total amount of modifier of less than 15, in particular from 0.4 to 10% by weight, based in each case on the end product. Moreover, the fact that the chromium dioxides obtained by the novel method are more finely divided, and the particularly advantageous narrow particle size distribution, are noteworthy.

The advantageous properties of a chromium dioxide produced according to the invention are particularly evident when it is used as magnetic material for the production of magnetic recording media.

Chromium dioxide produced in this manner is processed by a conventional method. To produce the magnetic layer, from 2 to 5 parts by weight of chromium

dioxide, one part of the binder or binder mixture and suitable dispersants, lubricants and other conventional additives in a total amount of not more than 10% by weight, based on the chromium dioxide, are converted to a dispersion. The dispersion thus obtained is filtered, and applied with a conventional coating apparatus, eg. a knife coater, onto the non-magnetic base in one or more thin layers, or in a thin layer on a magnetic recording medium already provided with another magnetic layer. Before the liquid coating mixture is dried at from 50 to 90°C, the chromium dioxide particles may be oriented magnetically if required. Special surface treatment of the magnetic layer is carried out by passing the coated webs between heated polished rollers under pressure. The thickness of the magnetic layers is then usually from 1.5 to 12 μm .

Suitable binders for the magnetic layers are the conventional polymer binders, such as vinyl chloride copolymers, acrylate copolymers, polyvinylacetals, such as polyvinylformal or polyvinylbutyral, fairly high molecular weight epoxy resins, polyurethanes and mixtures of these and similar binders. Substances which have proven advantageous are the elastomeric, virtually isocyanate-free linear polyester urethanes which are soluble in a volatile organic solvent, as can be prepared by reacting a polyester of an aliphatic dicarboxylic acid of 4 to 6 carbon atoms, such as adipic acid, and one or more aliphatic diols of 3 to 10 carbon atoms, such as 1,2- or 1,3-propylene glycol, butane-1,4-diol, diethylene glycol, neopentylglycol or octane-1,8-diol, with a diisocyanate of 6 to 24, in particular 8 to 20, carbon atoms, such as toluylene diisocyanate or 4,4'-diisocyanatodiphenylmethane, preferably in the presence of a small amount of glycol of 4 to 10 carbon atoms, such as butane-1,4-diol, which acts as a chain extender. Preferred polyester urethanes are those which are obtained from adipic acid, butane-1,4-diol and 4,4'-diisocyanatodi-



phenylmethane and have a Shore hardness A of from 70 to 100, a strength of from 400 to 420 kp/cm² and an elongation of about 440-560%. Other polymer binders which have proven satisfactory are those based on a copolymer of from 70 to 95, in particular from 75 to 90, % by weight of vinyl chloride and from 5 to 30, in particular from 10 to 25, % by weight of an alkyl ester of an olefinically unsaturated carboxylic acid of 3 to 5 carbon atoms, such as acrylic acid, methacrylic acid or maleic acid, where the alkyl radical is preferably of 1 to 3 carbon atoms. The corresponding vinyl chloride copolymers with one or more C₁-C₃-dialkyl maleates, such as copolymers of from 70 to 90% by weight of vinyl chloride, from 5 to 15% by weight of dimethyl maleate and from 5 to 15% by weight of diethyl maleate, are noteworthy. The K value according to H. Fikentscher is from 40 to 60 for the particularly suitable polymer binders.

The magnetic recording media produced using the chromium dioxide produced according to the invention possess the familiar good electroacoustic properties and, where relevant, video characteristics of conventional chromium dioxide magnetic tapes. Furthermore, they have a particularly high signal-to-noise ratio as well as low print-through properties. The high signal-to-noise ratio results in the tapes having a particularly wide dynamic range at both low and high frequencies, ie. the tapes have an optimum maximum output level. The ratio of noise at rest to signal-to-print-through ratio is also surprisingly good.

The Examples which follow illustrate the invention in comparison with a prior art Comparative Experiment. In the Examples and Comparative Experiment, parts and percentages are by weight, unless stated otherwise. Parts by volume bear the same relation to parts by weight as that of the liter to the kilogram.

EXAMPLE 1

2.19 l of water are initially taken in a reaction

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vessel having a capacity of 4 l, and 5.68 kg of chromic anhydride (CrO_3) are first added while stirring. After 10 minutes, 33 g (0.38% by weight, based on chromium dioxide) of potassium antimonyl tartrate ($\text{KSbO}(\text{C}_4\text{H}_4\text{O}_6)$).
5 0.5 H_2O , molecular weight 333.93) and 104 g (1.3% by weight, based on chromium dioxide) of acicular $\gamma\text{-Fe}_2\text{O}_3$ are added. When the evolution of gas which occurs on addition of the antimony-containing modifier has ceased, 2.84 kg of chromium(III) oxide are introduced while stirring
10 constantly, and stirring is continued for a further 20 minutes. The reaction slurry is then heated at from 300 to 350°C in an autoclave, chromium dioxide being formed during this process. The resulting oxygen causes an increase in pressure, and the pressure is kept at from
15 300 to 400 bar by means of a regulating valve. When the reaction is complete, the reactor is let down and cooled in such a way that the resulting chromium dioxide has a residual moisture content of from 1 to 5%. It is removed mechanically from the reaction vessel, milled, and
20 suspended in an aqueous sodium sulfite solution, and the agglomerates are broken up by wet-milling. During this procedure, 10% of the chromium dioxide is reduced at the surface of the crystals and converted to chromium(III) oxide hydroxide. The product is filtered, washed and then
25 dried at 50°C under reduced pressure.

The BET specific surface area in [m^2/g] of the resulting chromium dioxide was measured and its magnetic properties were determined by means of a vibrating sample magnetometer in a magnetic field of 160 kA/m at a mean
30 tap density ρ [g/cm^3], the particular magnetic properties determined being the coercive force H_c in (kA/m), the specific remanence M_r/ρ and the specific magnetization M_s/ρ in (nTm^3/g).

The measured values obtained are shown in the
35 Table.

EXAMPLE 2

The procedure described in Example 1 is followed,

except that 42 g of potassium antimonyl tartrate are used. The results of the measurements are shown in the Table.

EXAMPLE 3

5 The procedure described in Example 1 is followed, except that 55 g of potassium antimonyl tartrate are used. The results of the measurements are shown in the Table.

EXAMPLE 4

10 The procedure described in Example 1 is followed, except that 39 g of potassium antimonyl tartrate [$\text{Sb}_2(\text{C}_4\text{H}_4\text{O}_6)_3 \cdot 6 \text{H}_2\text{O}$, molecular weight 795.81] are used. The results of the measurements are shown in the Table.

EXAMPLE V

15 The procedure described in Example 1 is followed, except that, instead of potassium antimonyl tartrate, the corresponding amount, based on antimony, of antimony(III) oxide [Sb_2O_3 , molecular weight 291.5], ie. 14.4 g, is used. The results of the measurements are shown in the
20 Table.

EXAMPLE B1

115 parts of a chromium dioxide produced as described in Example 1, 2 parts of zinc stearate, 1.5 parts of soya lecithin, 3 parts of a mixture of
25 liquid fatty acids having a melting point of from 50 to 59°C, 110 parts of a mixture of equal amounts of tetrahydrofuran and 1,4-dioxane and 200 parts of 15% strength binder solution prepared by dissolving 19.5 parts of an elastomeric thermoplastic polyurethane (obtained as
30 described in German Published Application DAS 1,106,959 from adipic acid, butane-1,4-diol and 4,4'-diphenylisocyanatodiphenylmethane) and 10.5 parts of a vinyl chloride polymer of 80 parts of vinyl chloride and 10 parts of dimethyl maleate in 170 parts of a mixture of
35 equal amounts of tetrahydrofuran and 1,4-dioxane are introduced into a cylindrical steel mill which has a capacity of 1000 parts by volume and contains 1000 parts

of steel balls having a diameter of from 4 to 7 mm. The mixture is dispersed for 5 days, and the resulting dispersion is filtered under pressure through a glass/paper filter layer and is applied, on a conventional coating apparatus by means of a knife coater, onto a 12 μm thick polyethylene terephthalate film in a thickness such that the layer obtained after drying and calendering is 5.5 μm thick. Immediately after casting the liquid dispersion, the acicular chromium dioxide particles are oriented along the recording direction by means of a magnetic field. The surface of the magnetic layer has an average peak-to-valley height R_z , measured according to DIN 4756, sheet 1, vpm [sic] 0.15 μm . The coated film is slit into 3.81 mm wide magnetic tapes.

The magnetic tapes are tested as follows:

1. Magnetic properties

The magnetic properties of the resulting magnetic tapes are determined using a vibrating sample magnetometer in a magnetic field of 160 kA/m. The coercive force H_c in [kA/m], the residual induction M_r and the maximum magnetization M_m in [mT] are measured, and the orientation ratio R_f is calculated as the quotient of the residual induction along the preferred magnetic direction and that in a crosswise direction.

2. Electroacoustic properties

The electroacoustic tape data are measured according to DIN 45,512, sheet II, against the standard chromium dioxide reference tape S 4592 A, using a HF biasing current of 20 mA. All electroacoustic tape data, maximum output level at long wavelengths A_L and sensitivity E_L at 1 kHz, maximum output level at short wavelengths A_H and sensitivity E_H at 10 kHz, signal-to-bias noise ratio at rest RG_0 and signal-to-print-through ratio K_0 , are based on reference tape S 4592 A, the latter being set to 0 db for all parameters measured.

The corresponding measured parameters are shown in the Table.

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EXAMPLES B2, B3 and BV

The procedure described in Example B1 is followed, except that the corresponding chromium dioxide samples of Examples 2 and 3 and of the Comparative Experiment are used instead of that of Example 1. The results of the measurements are shown in the Table.

TABLE

		Examples				
		1/B1	2/B2	3/B3	4/B4	V/BV
<u>Powder</u>						
	BET	34.2	39.4	45.4	35	25.4
	ρ	1.15	1.12	1.13	1.19	1.35
	H_c	51.5	51.1	47.4	50.1	49.5
15	M_T/ρ	39.3	39.1	36.8	40.3	41.7
	M_s/ρ	68.9	68.7	64.8	71.0	76.0
<u>Tape</u>						
	H_c	52.9	49.6	45.2	50.7	50.9
	M_T	152	125	112	148	186
20	M_m	171	148	141	170	204
	R_f	2.85	2.36	1.96	2.88	3.92
	d	5.5	6.0	6.3	5.3	5.0
	E_T	-2.2	-2.5	-3	-2.0	-0.5
	E_H	+3.5	+3	+1.5	+3.2	+2.5
25	A_T	-2.5	-4	-1	-2.1	+0.5
	A_H	+5	+4.5	+2	+4.5	+4
	RG_o	+4.8	+5.3	+6.5	+4.5	+1.2
	Ko	56.5	53.0	42.5	55.0	57.0

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Claims

1. A process for producing ferromagnetic chromium dioxide by converting oxides of trivalent and hexavalent chromium under superatmospheric pressure and at elevated temperatures in the presence of water and with the addition of an antimony compound as a modifier and, if required, further additives in an amount of not more than 15% by weight, wherein the antimony compound used as a modifier is present in the form of a water-soluble salt and is added in an amount of less than 4% by weight, based on the resulting chromium dioxide, with the proviso that the anion of the salt-like antimony compound contains one or more organic molecular moieties which are oxidized in concentrated chromic acid.
2. A process as claimed in claim 1, wherein the antimony compound used as a modifier is antimony tart-rate.
3. A process as claimed in claim 1, wherein the antimony compound used as a modifier is potassium antimonyl tartrate, and this is added in an amount of less than 2% by weight, based on the resulting chromium dioxide.
4. A process as claimed in claim 3, wherein the potassium antimonyl tartrate used as a modifier is employed in an amount of from 0.1 to 1% by weight, based on the resulting chromium dioxide.
5. A process as claimed in claim 1, wherein acicular iron oxide particles in the form of α - or γ -FeOOH, Fe_3O_4 , or γ - or α - Fe_2O_3 , in an amount of from 0.5 to 15% by weight, based on the chromium dioxide, are used as a further modifier, in addition to the salt-like antimony compound.
6. A process as claimed in claim 1, wherein acicular iron particles having a BET specific surface area of from 5 to 40 m^2/g are added in an amount of from 0.4 to 10% by weight, based on the resulting chromium dioxide, as a further modifier in addition to the salt-like antimony



* C L N S *

compound.

7. A process as claimed in claim 3, wherein acicular iron oxide particles in the form of α - or γ -FeOOH, Fe_3O_4 , or γ - or α - Fe_2O_3 , in an amount of from 0.5 to 15% by weight, based on the chromium dioxide, are used as a further modifier, in addition to the potassium antimonyl tartrate.

8. A process as claimed in claim 3, wherein acicular iron particles having a BET specific surface area of from 5 to 40 m^2/g are added, in an amount of from 0.4 to 10% by weight, based on the resulting chromium dioxide, as a further modifier in addition to the potassium antimonyl tartrate.

9. Use of the ferromagnetic chromium dioxide produced as claimed in claim 1 or 2 or 3 or 4 or 5 or 6 or 7 or 8 for the production of a magnetic recording medium.



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as printed

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Claims

1. A process for the preparation of a ferromagnetic chromium dioxide by converting oxides of trivalent and hexavalent chromium under superatmospheric pressure and at elevated temperatures in the presence of water and with the addition of an antimony compound as a modifier and, if required, further additives in an amount of not more than 15 % by weight, wherein the antimony compound used as a modifier is present in the form of a water-soluble salt and is added in an amount of less than 4 % by weight, based on the resulting chromium dioxide, with the proviso that the anion of the salt-like antimony compound contains one or more organic molecular moieties which are oxidized in concentrated chromic acid.
2. A process as claimed in claim 1, wherein the antimony compound used as a modifier is antimony tartrate.
3. A process as claimed in claim 1, wherein the antimony compound used as a modifier is potassium antimonyl tartrate, and this is added in an amount of less than 2 % by weight, based on the resulting chromium dioxide.
4. A process as claimed in claim 3, wherein the potassium antimonyl tartrate used as a modifier is employed in an amount of from 0.1 to 1 % by weight, based on the resulting chromium dioxide.
5. A process as claimed in claim 1, wherein acicular iron oxide particles in the form of α - or γ -FeOOH, Fe_3O_4 or γ - or α - Fe_2O_3 , in an amount of from 0.5 to 15 % by weight, based on the chromium dioxide, are used as a further modifier, in addition to the salt-like antimony compound.
6. A process as claimed in claim 1, wherein acicular iron particles having a BET specific surface area of from 5 to 40 m^2/g are added in an amount of from 0.4 to 10 % by weight, based on the resulting chromium dioxide, as a further modifier in addition to the salt-like antimony compound.
7. A process as claimed in claim 3, wherein acicular iron oxide particles in the form of α - or γ -FeOOH, Fe_3O_4 or γ - or α - Fe_2O_3 , in an amount of from 0.5 to 15 % by weight, based on the chromium dioxide, are used as a further modifier, in addition to the potassium antimonyl tartrate.
8. A process as claimed in claim 3, wherein acicular iron particles having a BET specific surface area of from 5 to 40 m^2/g are added, in an amount of from 0.4 to 10 % by weight, based on the resulting chromium dioxide, as a further modifier in addition to the potassium antimonyl tartrate.
9. Use of a ferromagnetic chromium dioxide prepared as claimed in claim 1 or 2 or 3 or 4 or 5 or 6 or 7 or 8 for the preparation of a magnetic recording medium.

Revendications

1. Procédé de préparation de dioxyde de chrome ferromagnétique par réaction d'oxydes de chrome trivalent et hexavalent sous pression élevée et à température élevée en présence d'eau avec addition d'un composé de l'antimoine en tant qu'agent modificateur ainsi éventuellement d'autres additifs en quantité ne dépassant pas 15 % en poids, caractérisé en ce que le composé d'antimoine introduit en tant qu'agent modificateur est sous forme d'un sel soluble dans l'eau et en quantité non inférieure à 4 % en poids, par rapport au dioxyde de chrome qui se forme pourvu que l'anion du composé de l'antimoine sous forme de sel contienne au moins une partie de molécule organique qui s'oxyde dans l'acide chromique concentré.
2. Procédé selon la revendication 1, caractérisé en ce que le composé de l'antimoine introduit en tant qu'agent modificateur est un tartrate d'antimoine.
3. Procédé selon la revendication 1, caractérisé en ce que le composé de l'antimoine introduit en tant qu'agent modificateur est le tartrate de potassium et d'antimonyle et est utilisé en quantité non inférieure à 2 % en poids, par rapport au dioxyde de chrome qui se forme.
4. Procédé selon la revendication 3, caractérisé en ce que le tartrate de potassium introduit en tant qu'agent modificateur est utilisé en quantité de 0,1 à 1 % en poids par rapport au dioxyde de chrome résultant.
5. Procédé selon la revendication 1, caractérisé en ce que, en plus du composé d'antimoine sous forme de sel, on introduit comme autre agent modificateur des particules d'oxyde de fer en aiguilles sous forme de FeOOH α ou γ , Fe_3O_4 ou Fe_2O_3 γ ou α en quantité comprise entre 0,5 et 15 % en poids par rapport au dioxyde de chrome.
6. Procédé selon la revendication 1, caractérisé en ce que, en plus du composé d'antimoine sous forme de sel, on introduit comme autre agent modificateur des particules de fer sous forme d'aiguilles ayant une surface spécifique BET comprise entre 5 et 40 m^2/g en quantité comprise entre 0,4 et 10 % en poids par rapport au dioxyde de chrome résultant.
7. Procédé selon la revendication 3, caractérisé en ce que, en plus du tartrate de potassium et d'antimonyle, on introduit comme autre agent modificateur des particules d'oxyde de fer en aiguilles sous forme de FeOOH α ou γ , Fe_3O_4 ou Fe_2O_3 γ ou α en quantité comprise entre 0,5 et 15 % en poids par rapport au dioxyde de chrome.
8. Procédé selon la revendication 3, caractérisé en ce que, en plus du tartrate de potassium et d'antimonyle, on introduit comme autre agent modificateur des particules de fer sous forme d'aiguilles ayant une surface spécifique BET comprise entre 5 et 40 m^2/g en quantité comprise entre 0,4 et 10 % en poids par rapport au dioxyde de chrome résultant.
9. Utilisation du dioxyde de chrome ferromagnétique obtenu selon les revendications 1 à 8, pour la préparation de panneaux d'affichage magnétiques.