Description

Method for the production of magnet cores, magnet core and inductive component with a magnet core

[1] The invention relates to a method for the production of magnetic powder composite cores pressed from a mix of alloy powder and binder. It further relates to a magnet core produced from a mix of alloy powder and binder and to an inductive component with a magnet core.

[2] In powder composite cores of this type, low hysteresis and eddy-current losses and low coercitive field strength are desired. The powder is typically supplied in the form of flakes provided by comminuting a soft magnetic strip produced using melt spinning technology. These flakes may, for example, have the form of platelets and are typically first provided with an electrically insulating coating and then pressed to produce a magnet core. While flakes of pure iron or iron/nickel alloys are so ductile that they are plastically deformed under the influence of the compacting pressure and result in pressed cores of high density and strength, flakes or powders of relatively hard and rigid materials cannot be pressed with just any pressure. Rigid flakes would break in unsuitable conditions, resulting not in the desired compaction, but only in a further reduction of particle size. In addition, the break-up of the flakes releases fresh surfaces without any electrically insulating coating, which lead to a drastic reduction of the resistivity of the magnet core and thus to high eddy-current losses at high frequencies.

[3] As described, for example, in DE 103 48 810 A1, it is possible to use powders with a multi-modal particle size distribution. A multi-modal size distribution permits a relatively dense packing of the particles and thus the production of a relatively dense magnet core.

[4] When using FeAlSi-based materials, the high energy input required for comminution results in structural damage in the production of fine-grain particle fractions, but these are healed virtually completely in the subsequent heat treatment process and hardly affect the magnetic properties of the finished magnet core. In mixes with ductile materials, packing density can be increased by increasing the ductile component, for example the pure iron component. This procedure is, for example, described in JP 2001-196216.

[5] Problems are posed, however, by the production of dense magnet cores from amorphous FeBSi-based materials, which are favoured owing to their good magnetic properties. In the energy-intensive production of the fine-grain particle fractions, FeBSi-based materials form phases of iron borides, which represent permanent structural damage and adversely affect magnetic properties.

[6] The invention is therefore based on the problem of specifying a method for the production of a powder composite core, which allows the production of particularly dense magnet cores from alloys produced in a rapid solidification process. It is further based on the problem of specifying a particularly dense magnet core with low coercitive field strength.

[7] According to the invention, this problem is solved by the subject matter of the independent patent claims. Advantageous further development of the invention form the subject matter of the dependent patent claims.

[8] A method according to the invention for the production of a magnet core comprises the following steps: First, at least one coarse-grain powder fraction is produced from an amorphous strip of a soft magnetic alloy. In addition, at least one fine-grain powder fraction is produced from a nanocrystalline strip, likewise of a soft magnetic alloy. Following comminution, the particle fractions may be sized in order to obtain an optimum particle size distribution. The particle fractions are then mixed to produce a multi-modal powder, the particles of the coarse-grain particle fraction having an amorphous structure, while the particles of the fine-grain particle fraction have a nanocrystalline structure. The multi-modal powder is then pressed to produce a magnet core.

[9] The soft magnetic strip material is typically produced as amorphous strip in a rapid solidification process, the term “strip” in this context including a foil-like form or pieces of strip. To produce a nanocrystalline strip, the amorphous strip can then be subjected to heat treatment to obtain the nanocrystalline structure.

[10] According to a basic concept of the invention, the aim is to minimise energy input in the comminution of the strip material to produce a powder. Energy input can be reduced by converting the strip into a nanocrystalline state prior to comminution, thus making it very brittle. In this brittle state, the fine-grain powder fractions can be produced without increasing energy input sufficiently for the formation of FeB phases. In this way, irreversible structural damage can be avoided. On the other hand, the production of the coarse-grain powder fraction from nanocrystalline strip is not advisable, because the flakes produced from nanocrystalline strip would also be nanocrystalline and therefore so brittle that they would not be compacted under pressure, but rather disintegrate.

[11] This problem can be solved by producing the fine-grain and the coarse-grain powder fractions by different means. By separately producing the fine-grain fractions from nanocrystalline strip and the coarse-grain fractions from amorphous strip, the roles played by the powder fractions in the production of the magnet core and their properties in the pressing process are taken into consideration. The production processes for the different powder fractions are “tailor-made” in a manner of speaking. As

a result, the properties of the powder can be precisely adapted to pressing conditions and to the desired density of the finished magnet core before the pressing process.

[12] In this way, an alloy capable of nanocrystallisation can be used even for the amorphous strip, provided it is still in a amorphous state at the time of pressing. However, the initially amorphous alloy capable of nanocrystallisation can be converted into a nanocrystalline alloy by heat treatment. As a result, a variety of alloy combinations can be used for the coarse- and fine-grain fractions: The fine-grain fraction is produced from an alloy capable of nanocrystallisation, which is already in a nanocrystalline state in the pressing process. The coarse-grain fraction, on the other hand, can be produced either from an alloy not capable of nanocrystallisation or from an alloy capable of nanocrystallisation, and in the latter case, the alloy can be converted into a nanocrystalline state after pressing.

[13] The particles representing the fine-grain powder fraction advantageously have a diameter between 20 and 70 µm, while the particles representing the coarse-grain powder fraction have a diameter between 70 and 200 µm. With particles in this size range, relatively dense packing and therefore dense magnet cores can be obtained.

[14] In one embodiment of the method, the amorphous strip is pre-embrittled by heat treatment at a pre-embrittling temperature $T_{embrittle}$ prior to the production of the coarse-grain powder fraction in order to simplify comminution, the pre-embrittling temperature $T_{embrittle}$ and the crystallisation temperature $T_{crystal}$ of the amorphous strip having the relationship $T_{embrittle} < T_{crystal}$. The pre-embrittling temperature $T_{embrittle}$ is therefore chosen low enough to avoid (nano-) crystallisation. It is further chosen low enough, and the duration of the heat treatment is chosen short enough, to make the particles produced from the strip ductile enough to avoid break-up in the pressing process. The pre-embrittling temperature $T_{embrittle}$ advantageously is $100°C <= T_{embrittle} <= 400°C$, preferably $200°C <= T_{embrittle} <= 400°C$. The duration of the heat treatment may be 0.5 to 8 hours.

[15] In an alternative embodiment of the method, the amorphous strip is comminuted to produce the coarse-grain powder fraction without any preceding heat treatment for pre-embrittling in the "as cast" state, i.e. in the state in which it is following the rapid solidification process. The amorphous strip is advantageously comminuted to produce the coarse-grain powder fraction at a grinding temperature T_{mill} of $-196°C <= T_{mill} <= 20°C$.

[16] The nanocrystalline strip used to produce the fine-grain powder fraction is, for example, comminuted in a cutting mill. Using a cutting mill instead of, for example, a ball mill reduces energy input to a minimum and avoids irreversible structural damage.

[17] In one embodiment of the method, the same alloy is used for the amorphous strip and for the nanocrystalline strip. In this case, the strip used to produce the fine-grain

powder fraction is nanocrystallised by heat treatment following the rapid solidification process, while the strip used to produce the coarse-grain powder fraction is left in its amorphous state.

[18] It is, however, alternatively possible to use different alloys. The first soft magnetic alloy for the amorphous strip may, for instance, be an alloy which is particularly suitable for processing in the amorphous state and which is sufficiently ductile, while the second soft magnetic alloy for the nanocrystalline strip may be an alloy which can be nanocrystallised particularly easily.

[19] In view of these considerations, suitable soft magnetic alloys for both the amorphous and the nanocrystalline strip are soft magnetic iron-based alloys.

[20] In one embodiment, the amorphous particles have the alloy composition $M_{alpha}Y_{beta}Z_{gamma}$, wherein M is at least one element from the group including Fe, Ni and Co, wherein Y is at least one element from the group including B, C and P, wherein Z is at least one element from the group including Si, Al and Ge, and wherein a, b, and g are specified in atomic percent and meet the following conditions: 70 <= alpha <= 85; 5 <= beta <= 20; 0 <= gamma <= 20, wherein up to 10 atomic percent of the M component may be replaced by at least one element from the group including Ti, V, Cr, Mn, Cu, Zr, Nb, Mo, Ta and W and up to 10 atomic percent of the (Y+Z) component may be replaced by at least one element from the group including In, Sn, Sb und Pb.

[21] The particles capable of nanocrystallisation may have the alloy composition $(Fe_{1-a}M_a)_{100-x-y-z-alpha-beta-gamma}Cu_xSi_yB_zM'_{alpha}M''_{beta}X_{gamma}$, wherein M is Co and/or Ni, wherein M' is at least one element from the group including Nb, W, Ta, Zr, Hf, Ti and Mo, wherein M" is at least one element from the group including V, Cr, Mn, Al, elements of the platinum group, Sc, Y, rare earths, Au, Zn, Sn and Re, wherein X is at least one element from the group including C, Ge, P, Ga, Sb, In, Be und As, and wherein a, x, y, z, alpha, beta and gamma are specified in atomic percent and meet the following conditions: 0 <= a <= 0.5; 0.1 <= x <= 3; 0 <= y <= 30; 0 <= z <= 25; 0 <= y+z <= 35; 0.1 <= alpha <= 30; 0 <= beta <= 10; 0 <= gamma <= 10.

[22] As an alternative, the particles capable of nanocrystallisation may have the alloy composition $(Fe_{1-a-b}Co_aNi_b)_{100-x-y-z}M_xB_yT_z$, wherein M is at least one element from the group including Nb, Ta, Zr, Hf, Ti, V and Mo, wherein T is at least one element from the group including Cr, W, Ru, Rh, Pd, Os, Ir, Pt, Al, Si, Ge, C and P, and wherein a, b, x, y and z are specified in atomic percent and meet the following conditions: 0 <= a <= 0.29; 0 <= b <= 0.43; 4 <= x <= 10; 3 <= y <= 15; 0 <= z <= 5.

[23] For the strip capable of nanocrystallisation, at least one of the alloys $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, $Fe_{73.5}Nb_3Cu_1Si_{13.5}B_9$, $Fe_{86}Cu_1Zr_7B_6$, $Fe_{91}Zr_7B_3$ and $Fe_{84}Nb_7B_9$ can be used.

[24] The multi-modal powder obtained by mixing coarse- and fine-grain powder fractions

is advantageously pressed at a pressing temperature T_{press} of $T_{press} > T_{embrittle}$ to produce a magnet core. This ensures that the coarse-grain particles, in particular, behave in a very ductile manner and that there is no further mechanical comminution during the pressing process.

[25] After pressing, the magnet core is advantageously subjected to heat treatment at a heat treatment temperature T_{anneal} in order to relieve mechanical stresses introduced into the magnet core by pressing and to obtain good magnetic properties, in particular a low coercitive field strength. The heat treatment temperature T_{anneal} is expediently chosen such that the heat treatment temperature T_{anneal} and the crystallisation temperature $T_{crystal}$ of the first soft magnetic alloy have the relationship $T_{anneal} >= T_{crystal}$. This results in the nanocrystallisation of the coarse-grain particles which still have an amorphous structure at this point. For this purpose, the heat treatment temperature is typically set above 500°C.

[26] As an alternative, the heat treatment temperature T_{anneal} may be chosen such that the heat treatment temperature T_{anneal} and the crystallisation temperature $T_{crystal}$ of the first soft magnetic alloy have the relationship $T_{anneal} <= T_{crystal}$. In this case, the nano-crystallisation of the amorphous particle fraction is avoided. The only purpose of the heat treatment in this case is the relief of mechanical stresses, and typically 400°C <= T_{anneal} <= 450°C.

[27] All heat treatment processes are advantageously conducted in a controlled atmosphere to prevent corrosion and thus the premature ageing of the magnet core combined with a deterioration of its magnetic properties.

[28] Prior to pressing, processing aids such as binders and/or lubricants are advantageously added to the multi-modal powder. The particles representing the coarse- and/or fine-grain powder fractions may be pickled in an aqueous or alcohol solution prior to pressing to apply an electrically insulating coating and then dried. An electrically insulating coating may also be applied by different means. It is used to reduce the resistivity of the magnet core and to reduce eddy-current losses.

[29] A magnet core according to the invention comprises a soft magnetic powder made from particles, the particle size distribution being multi-modal. It further comprises processing aids such as binders. The powder comprises at least one coarse-grain powder fraction with particles with an amorphous structure and at least one fine-grain powder fraction with particles with a nanocrystalline structure.

[30] A magnet core of this type may combine an exceptionally high density with a low coercitive field strength, because the multi-modal particle size distribution permits a particularly dense packing of the particles, while the particle surfaces suffer only minor deformation and structural damage.

[31] The magnet core according to the invention can be used in inductive components

such as storage chokes, PFC chokes (chokes for power factor correction), switching power supplies, filter chokes or smoothing chokes.

[32] Embodiments of the invention are explained in greater detail below.

[33] Example 1

[34] From a strip with the nominal composition $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, particle fractions with the following particle diameters were produced: The nanocrystalline particles of the first fraction had diameters between 28 and 50 µm, the amorphous particles of the second fraction had diameters between 80 and 106 µm, and the likewise amorphous particles of the third fraction had diameters between 106 and 160 µm. The press-ready powder mix consisted of 29% flakes of the first fraction, 58% flakes of the second fraction and 10% flakes of the third fraction in addition to 2.8% binder mix and 0.2% lubricant. The mix was pressed at a pressure of 8 t/cm2 and a temperature of 180°C to produce a magnet core. After pressing, the core had a density of 67 percent by volume. After pressing, the magnet core was subjected to a heat treatment lasting one hour in a controlled atmosphere at 560°C. The finished magnet core had a static coercitive field strength of 51.6 A/m.

[35] Example 2

[36] From a strip with the nominal composition $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, particle fractions with the following particle diameters were produced: The nanocrystalline particles of the first fraction had diameters between 40 and 63 µm, and the amorphous particles of the second fraction had diameters between 80 and 106 µm. The press-ready powder mix consisted of 48.5% flakes of the first fraction, 48.5% flakes of the second fraction and 2.8% binder mix and 0.2% lubricant. The mix was pressed at a pressure of 8 t/cm^2 and a temperature of 180°C to produce a magnet core. After pressing, the core had a density of 68.3 percent by volume. After pressing, the magnet core was subjected to a heat treatment lasting one hour in a controlled atmosphere at 560°C. The finished magnet core had a static coercitive field strength of 55.4 A/m.

[37] For comparison, magnet cores were produced in the conventional way from purely amorphous powders.

[38] Comparative example 1

[39] From a strip with the nominal composition $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, purely amorphous particles with particle diameters between 80 and 106 µm were produced. The press-ready powder mix consisted of 97% of these amorphous particles and 2.8% binder mix and 0.2% lubricant. The mix was pressed at a pressure of 8 t/cm^2 and a temperature of 180°C to produce a magnet core. After pressing, the core had a density of 61.7 percent by volume. After pressing, the magnet core was subjected to a heat treatment lasting one hour in a controlled atmosphere at 560°C. The finished magnet core had a static coercitive field strength of 71.0 A/m.

[40] Comparative example 2

[41] From a strip with the nominal composition $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, purely amorphous particle fractions with the following particle diameters were produced: The particles of the first fraction had diameters between 40 and 63 µm, and the particles of the second fraction had diameters between 80 and 106 µm. The press-ready powder mix consisted of 48.5% flakes of the first fraction, 48.5% flakes of the second fraction and 2.8% binder mix and 0.2% lubricant. The mix was pressed at a pressure of 8 t/cm^2 and a temperature of 180°C to produce a magnet core. After pressing, the core had a density of 63.2 percent by volume. After pressing, the magnet core was subjected to a heat treatment lasting one hour in a controlled atmosphere at 560°C. The finished magnet core had a static coercitive field strength of 100.5 A/m.

[42] These examples show that high densities can be combined with low coercitive field strengths of the magnet cores when using the method according to the invention. The low coercitive field strength in the magnet cores from examples 1 and 2 is due to the fact that the fine-grain particles, as a result of their production from a nanocrystalline material, do not suffer any significant irreversible structural damage caused by the formation of FeB phases.

[43] As a result of the separate production of coarse-grain amorphous and fine-grain nano-crystalline powder fractions, the resulting powder mix meets all requirements: It is multi-modal and allows, even when using FeBSi-based alloys capable of nano-crystallisation, a very dense packing of the particles, resulting in a high density of the magnet core. Owing to their amorphous structure, the coarse-grain particles are ductile enough not to break up in the pressing process. And finally, being produced from a nanocrystalline starting material, the fine-grain particles are not irreversibly damaged by the formation of iron boride phases which would adversely affect the magnetic properties of the core.

Claims

[1] Method for the production of a magnet core, comprising the following steps:
- production of at least one coarse-grain powder fraction from an amorphous soft magnetic strip;
- production of at least one fine-grain powder fraction from a nanocrystalline soft magnetic strip made of an alloy capable of nanocrystallisation;
- mixing of the coarse- and fine-grain powder fractions to produce a powder with a multi-modal particle size distribution, wherein the particles of the coarse-grain particle fraction have an amorphous structure and the particles of the fine-grain powder fraction have a nanocrystalline structure;
- pressing of the multi-modal powder to produce a magnet core.

[2] Method according to claim 1,
characterised in that
the particles representing the fine-grain powder fractions have particle diameters between 20 and 70 µm.

[3] Method according to claim 1 or 2,
characterised in that
the particles representing the coarse-grain powder fractions have particle diameters between 70 and 200 µm.

[4] Method according to any of claims 1 to 3,
characterised in that
the amorphous strip is pre-embrittled by heat treatment at a pre-embrittling temperature $T_{embrittle}$ prior to the production of the coarse-grain powder fractions, the pre-embrittling temperature $T_{embrittle}$ and the crystallisation temperature $T_{crystal}$ of the amorphous strip having the relationship $T_{embrittle} < T_{crystal}$.

[5] Method according to claim 4
characterised in that
$100°C <= T_{embrittle} <= 400°C$.

[6] Method according to claim 4 or 5
characterised in that
$200°C <= T_{embrittle} <= 400°C$.

[7] Method according to any of claims 1 to 3,
characterised in that
the amorphous strip is comminuted to produce the coarse-grain powder fraction without any preceding heat treatment for pre-embrittling in the “as cast” state.

[8] Method according to any of claims 1 to 7,
characterised in that

the amorphous strip is comminuted to produce the coarse-grain powder fraction at a grinding temperature T_{mill} of -196°C <= T_{mill} <= 20°C.

[9] Method according to any of claims 1 to 8,
characterised in that
the nanocrystalline strip used to produce the fine-grain powder fraction is comminuted in a cutting mill.

[10] Method according to any of claims 1 to 9,
characterised in that
an alloy not capable of nanocrystallisation is used for the amorphous strip.

[11] Method according to claim 10,
characterised in that
an iron-based alloy is used as an alloy for the amorphous strip.

[12] Method according to claim 10,
characterised in that
an alloy of the composition $M_{alpha}Y_{beta}Z_{gamma}$, is used as an alloy for the amorphous strip, wherein M is at least one element from the group including Fe, Ni and Co, wherein Y is at least one element from the group including B, C and P, wherein Z is at least one element from the group including Si, Al and Ge, and wherein alpha, beta, and gamma are specified in atomic percent and meet the following conditions: 70 <= alpha <= 85; 5 <= beta <= 20; 0 <= gamma <= 20, wherein up to 10 atomic percent of the M component may be replaced by at least one element from the group including Ti, V, Cr, Mn, Cu, Zr, Nb, Mo, Ta and W and up to 10 atomic percent of the (Y+Z) component may be replaced by at least one element from the group including In, Sn, Sb und Pb.

[13] Method according to any of claims 1 to 9,
characterised in that
the same alloy capable of nanocrystallisation is used both for the amorphous strip and for the nanocrystalline strip.

[14] Method according to any of claims 1 to 9,
characterised in that
different alloys are used for the amorphous strip and for the nanocrystalline strip, both alloys being capable of nanocrystallisation.

[15] Method according to any of claims 1 to 14,
characterised in that
at least one of the alloys capable of nanocrystallisation has the composition $(Fe_{1-a}M_a)_{100-x-y-z-alpha-beta-gamma}Cu_xSi_yB_zM'_{alpha}M''_{beta}X_{gamma}$, wherein M is Co and/or Ni, wherein M' is at least one element from the group including Nb, W, Ta, Zr, Hf, Ti and Mo, wherein M'' is at least one element from the group including V, Cr,

Mn, Al, elements of the platinum group, Sc, Y, rare earths, Au, Zn, Sn and Re, wherein X is at least one element from the group including C, Ge, P, Ga, Sb, In, Be und As, and wherein a, x, y, z, alpha, beta, and gamma are specified in atomic percent and meet the following conditions: 0 <= a <= 0.5; 0.1 <= x <= 3; 0 <= y <= 30; 0 <= z <= 25; 0 <= y+z <= 35; 0.1 <= alpha <= 30; 0 <= beta <= 10; 0 <= gamma <= 10.

[16] Method according to any of claims 1 to 15,
characterised in that
at least one of the alloys capable of nanocrystallisation has the composition $(Fe_{1-a-b}Co_aNi_b)_{100-x-y-z}M_xB_yT_z$, wherein M is at least one element from the group including Nb, Ta, Zr, Hf, Ti, V and Mo, wherein T is at least one element from the group including Cr, W, Ru, Rh, Pd, Os, Ir, Pt, Al, Si, Ge, C and P, and wherein a, b, x, y and z are specified in atomic percent and meet the following conditions: 0 <= a <= 0.29; 0 <= b <= 0.43; 4 <= x <= 10; 3 <= y <= 15; 0 <= z <= 5.

[17] Method according to any of claims 1 to 16,
characterised in that
at least one of the alloys capable of nanocrystallisation has the composition $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, $Fe_{73.5}Nb_3Cu_1Si_{13.5}B_9$, $Fe_{86}Cu_1Zr_7B_6$, $Fe_{91}Zr_7B_3$ or $Fe_{84}Nb_7B_9$.

[18] Method according to any of claims 1 to 17,
characterised in that
the multi-modal powder is pressed at a pressing temperature T_{press} of $T_{press} > T_{embrittle}$ to produce a magnet core.

[19] Method according to any of claims 1 to 18,
characterised in that
the magnet core is subjected to heat treatment at a heat treatment temperature T_{anneal} after pressing.

[20] Method according to claim 19,
characterised in that
the heat treatment temperature T_{anneal} and the crystallisation temperature $T_{crystal}$ of the first soft magnetic alloy have the relationship $T_{anneal} >= T_{crystal}$.

[21] Method according to claim 19 or 20,
characterised in that
Tanneal > 500°C.

[22] Method according to claim 19,
characterised in that
the heat treatment temperature T_{anneal} and the crystallisation temperature $T_{crystal}$ of the amorphous strip have the relationship $T_{anneal} <= T_{crystal}$.

[23] Method according to claim 19 or 22,
characterised in that
$400°C <= T_{anneal} <= 450°C$.

[24] Method according to any of claims 1 to 23,
characterised in that
processing aids such as binders and/or lubricants are added to the multi-modal powder prior to pressing.

[25] Method according to any of claims 4 to 24,
characterised in that
the heat treatments are conducted in a controlled atmosphere.

[26] Method according to any of claims 1 to 25,
characterised in that
the particles representing the coarse- and/or fine-grain powder fractions are pickled in an aqueous or alcohol solution prior to pressing to apply an electrically insulating coating and then dried.

[27] Magnet core comprising a soft magnetic powder with a multi-modal particle size distribution and processing aids, wherein the powder comprises at least one coarse-grain powder fraction of particles with an amorphous structure and at least one fine-grain powder fraction of particles with a nanocrystalline structure.

[28] Magnet core according to claim 27,
characterised in that
the amorphous particles and the nanocrystalline particles have the same alloy composition capable of nanocrystallisation.

[29] Magnet core according to claim 27,
characterised in that
the amorphous particles and the nanocrystalline particles have different alloy compositions capable of nanocrystallisation.

[30] Magnet core according to claim 27,
characterised in that
the amorphous particles consist of an amorphous iron-based alloy.

[31] Magnet core according to any of claims 27 to 30,
characterised in that
the particles representing the fine-grain powder fraction have particle diameters between 20 and 70 μm.

[32] Magnet core according to any of claims 27 to 31,
characterised in that
the particles representing the coarse-grain powder fraction have particle diameters between 70 and 200 μm.

[33] Magnet core according to any of claims 30 to 32,
characterised in that
the amorphous particles have the alloy composition $M_{alpha}Y_{beta}Z_{gamma}$, wherein M is at least one element from the group including Fe, Ni and Co, wherein Y is at least one element from the group including B, C and P, wherein Z is at least one element from the group including Si, Al and Ge, and wherein alpha, beta, and gamma are specified in atomic percent and meet the following conditions: 70 <= alpha <= 85; 5 <= beta <= 20; 0 <= gamma <= 20, wherein up to 10 atomic percent of the M component may be replaced by at least one element from the group including Ti, V, Cr, Mn, Cu, Zr, Nb, Mo, Ta and W and up to 10 atomic percent of the (Y+Z) component may be replaced by at least one element from the group including In, Sn, Sb und Pb.

[34] Magnet core according to any of claims 27 to 33,
characterised in that
the nanocrystalline particles have the alloy composition $(Fe_{1-a}M_a)_{100-x-y-z-alpha-beta-gamma}Cu_xSi_yB_zM'_{alpha}M''_{beta}X_{gamma}$, wherein M is Co and/or Ni, wherein M' is at least one element from the group including Nb, W, Ta, Zr, Hf, Ti and Mo, wherein M" is at least one element from the group including V, Cr, Mn, Al, elements of the platinum group, Sc, Y, rare earths, Au, Zn, Sn and Re, wherein X is at least one element from the group including C, Ge, P, Ga, Sb, In, Be und As, and wherein a, x, y, z, alpha, beta, and gamma are specified in atomic percent and meet the following conditions: 0 <= a <= 0.5; 0.1 <= x <= 3; 0 <= y <= 30; 0 <= z <= 25; 0 <= y+z <= 35; 0.1 <= alpha <= 30; 0 <= beta <= 10; 0 <= gamma <= 10.

[35] Magnet core according to any of claims 27 to 33,
characterised in that
the nanocrystalline particles have the alloy composition $(Fe_{1-a-b}Co_aNi_b)_{100-x-y-z}M_xB_yT_z$, wherein M is at least one element from the group including Nb, Ta, Zr, Hf, Ti, V and Mo, wherein T is at least one element from the group including Cr, W, Ru, Rh, Pd, Os, Ir, Pt, Al, Si, Ge, C and P, and wherein a, b, x, y and z are specified in atomic percent and meet the following conditions: 0 <= a <= 0.29; 0 <= b <= 0.43; 4 <= x <= 10; 3 <= y <= 15; 0 <= z <= 5.

[36] Magnet core according to any of claims 27 to 35,
characterised in that
the nanocrystalline particles have at least one of the alloy compositions $Fe_{73.5}Nb_3Cu_1Si_{15.5}B_7$, $Fe_{73.5}Nb_3Cu_1Si_{13.5}B_9$, $Fe_{86}Cu_1Zr_7B_6$, $Fe_{91}Zr_7B_3$ and $Fe_{84}Nb_7B_9$.

[37] Magnet core according to any of claims 27 to 36,
characterised in that

the magnet core includes processing aids such as binders and/or lubricants.

[38] Inductive component with a magnet core according to any of claims 1 to 37.

[39] Inductive component according to claim 38,
characterised in that
the inductive component is a choke for power factor correction.

[40] Inductive component according to claim 38,
characterised in that
the inductive component is a storage choke.

[41] Inductive component according to claim 38,
characterised in that
the inductive component is a filter choke.

[42] Inductive component according to claim 38,
characterised in that
the inductive component is a smoothing choke.