

5 MAGNESIUM-ALUMINUM-ZINC ALLOY, METHOD FOR THE PRODUCTION THEREOF AND
 USE THEREOF

10 The subject matter of the patent application relates to a magnesium alloy and to a method
 for the production thereof and to the use thereof.

15 It is known that the properties of magnesium alloys are decisively defined by the type
 and quantity of the alloying elements and impurities as well as the production
 conditions. The effects of the alloying elements and impurities on the properties of the
 magnesium alloys have been known for a long time to a person skilled in the art and
 illustrate the complex nature of determining the properties of binary or ternary
 magnesium alloys for the use thereof as implant materials.

20 The alloying element used most frequently for magnesium is aluminum, resulting in
 increased tensile strength due to solid solution and precipitation hardening and fine
 grain formation, but also in microporosity. Moreover, in the melt aluminum shifts the
 iron precipitation boundary toward drastically lower iron contents at which the iron
 particles precipitate or form intermetallic particles together with other elements.

25 Calcium exhibits a pronounced grain refining effect and worsens the castability and
 corrosion resistance.

 Undesirable accompanying elements in magnesium alloys include iron, nickel, cobalt and
 copper, which cause a considerable increase in the corrosion tendency due to the
 electropositive nature thereof.

30 Manganese can be found in all magnesium casting alloys and binds iron in the form of
 AlMnFe precipitations, whereby the formation of local elements is reduced. On the other
 hand, manganese is not able to bind all the iron, and therefore a remainder of iron and a
 remainder of manganese are always left in the melt.

Silicon lowers the castability and viscosity, and as the content of Si rises, a worsened corrosion behavior is to be expected. Iron, manganese and silicon have a very high tendency to form an intermetallic phase.

5 The electrochemical potential of this phase is very high and can thus act as a cathode controlling the corrosion of the alloy matrix.

As a result of solid solution hardening, zinc improves the mechanical properties and results in grain refining, however it also leads to microporosity with a tendency toward hot cracking starting at a content of 1.5 to 2% by weight in binary Mg-Zn and ternary
10 Mg-Al-Zn alloys.

Alloying additions made of zirconium increase the tensile strength without lowering the expansion and lead to grain refining, but also to a strong impairment of dynamic recrystallization, which is manifested in an increase of the recrystallization temperature and therefore requires high energy expenditure. Moreover, zirconium cannot be added to
15 melts containing aluminum and silicon because the grain refining effect is lost.

Rare earths such as Lu, Er, Ho, Th, Sc and In all exhibit a similar chemical behavior and form eutectic systems with partial solubility on the magnesium-rich side of the binary phase diagrams such that precipitation hardening is possible.

The addition of further alloying elements, in conjunction with the impurities, is known to
20 cause the formation of different intermetallic phases in binary magnesium alloys. For example, the intermetallic phase $\text{Mg}_{17}\text{Al}_{12}$ forming at the grain boundaries is brittle and limits the ductility. As compared to the magnesium matrix, this intermetallic phase is more noble and able to form local elements, whereby the corrosion behavior worsens.

25 In addition to these influencing factors, the properties of the magnesium alloys also decisively depend on the metallurgical production conditions. Conventional casting methods automatically introduce impurities when adding, by alloying, the alloying elements. The prior art (US 5,055,254 A) therefore defines tolerance limits for impurities in magnesium casting alloys, which, for example for a magnesium-aluminum-zinc alloy
30 containing approximately 8 to 9.5% by weight Al and 0.45 to 0.9% by weight Zn, mentions tolerance limits of 0.0015 to 0.0024% by weight Fe, 0.0010% by weight Ni, 0.0010 to 0.0024% by weight Cu and no less than 0.15 to 0.5% by weight Mn.

Tolerance limits for impurities in magnesium and the alloys thereof as well as the production conditions are mentioned in many known documents and listed as follows in % by weight:

Alloy	Production	State	Fe	Fe/Mn	Ni	Cu
Pure Mg	no information		0.017		0.005	0.01
AZ 91	Die casting	F		0.032	0.005	0.040
	High-pressure die casting			0.032	0.005	0.040
	Low-pressure die casting			0.032	0.001	0.040
		T4		0.035	0.001	0.010
		T6		0.046	0.001	0.040
	Gravity die casting	F		0.032	0.001	0.040
AM60	Die casting	F		0.021	0.003	0.010
AM50	Die casting	F		0.015	0.003	0.010
AS41	Die casting	F		0.010	0.004	0.020
AE42	Die casting	F		0.020	0.020	0.100

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It has been found that these tolerance definitions are not sufficient to reliably exclude the formation of corrosion-promoting intermetallic phases, which in terms of electrochemistry have a more noble potential than the magnesium matrix.

- 10 Biodegradable implants require a load-bearing function and consequently strength, together with sufficient expandability, during the physiologically necessary support periods thereof. However, especially in this respect, the known magnesium materials cannot come even close to the properties achieved by permanent implants such as titanium, CoCr alloys and titanium alloys. The ultimate tensile strength R_m for permanent
- 15 implants is approximately 500 MPa to >1000 MPa, while that of magnesium materials is <275 MPa so far, and in most cases <250 MPa.

Another drawback of many technical magnesium materials is that the difference thereof between ultimate tensile strength R_m and proof stress R_p is only small. In the case of

20 implants that allow plastic deformation, such as cardiovascular stents, this means that no further resistance exists against deformation after initial deformation of the material, and the regions that have already been deformed are deformed further without any load

increase, whereby overstretching of parts of the component may be caused and fracture may occur.

Many magnesium materials additionally exhibit a clearly pronounced mechanical asymmetry, which is manifested in the difference in the mechanical properties, especially the proof stress R_p with tension load and compression load. Such asymmetries are created, for example, during forming processes such as extrusion, rolling and drawing, which are used to produce suitable semi-finished products. A difference between the proof stress R_p during tension and the proof stress R_p during compression that is too large may result in inhomogeneous deformation of a component, such as a cardiovascular stent, which later undergoes multiaxial deformation, and may cause cracking and fracture.

Because of the low number of crystallographic slip systems, magnesium alloys can generally also form textures during forming processes such as extrusion, rolling and drawing used to produce suitable semifinished products by orienting the grains during the forming process. Specifically, this means that the semifinished product has different properties in different directions in space. For example, high deformability or elongation at fracture occurs in one direction in space after forming, and reduced deformability or elongation at fracture occurs in another direction in space. The formation of such textures should likewise be avoided, because a stent is subjected to high plastic deformation, and reduced elongation at fracture increases the risk of failure of the implant. One method for substantially avoiding such textures during forming is to adjust as fine a grain as possible prior to forming. Because of the hexagonal lattice structure of magnesium materials, the ability of these materials to deform at room temperature is low, which is characterized by slip in the base plane. If the material additionally has a coarse microstructure, which is to say a coarse grain, so-called twinning is forcibly produced upon further deformation, at which shear strain occurs, which transforms a crystal region into a position that is mirror symmetrical to the starting position.

The resulting twin grain boundaries constitute weak points in the material, where incipient cracking starts, especially with plastic deformation, which ultimately leads to the destruction of the component.

If the grain of the implant materials is sufficiently fine, the risk of such implant failure is drastically reduced. Implant materials should therefore have as fine a grain as possible so as to prevent such undesirable shear strain.

5 All available technical magnesium materials for implants are subject to high corrosion in physiological media. Attempts have been made in the prior art to curb the corrosion tendency by providing the implants with a corrosion-inhibiting coating, for example made of polymeric materials (EP 2 085 100 A2, EP 2 384 725 A1), an aqueous or alcoholic conversion solution (DE 10 2006 060 501 A1) or an oxide
10 (DE 10 2010 027 532 A1, EP 0 295 397 A1).

The use of polymeric passivation layers is highly contested, because virtually all appropriate polymers also cause strong inflammations in the tissue at times. Thin structures without such protective measures do not reach the required support periods. The corrosion on thin-walled traumatological implants is often times accompanied by an
15 excessively fast loss of tensile strength, which poses an additional burden by forming excessive amounts of hydrogen per unit of time. The consequences are undesirable gas inclusions in the bones and tissue.

In the case of traumatological implants having larger cross-sections, there is a need to be able to deliberately control the hydrogen problem and the corrosion rate of the implant by
20 way of the structure thereof.

Specifically with biodegradable implants, there is a desire for maximum biocompatibility of the elements, because all the chemical elements that are contained are absorbed by the body after decomposition. In any case, highly toxic elements such as Be, Cd, Pb, Cr and
25 the like should be avoided.

Degradable magnesium alloys are especially suitable for implementing implants which have been employed in a wide variety of forms in modern medical technology. Implants are used, for example, to support vessels, hollow organs and vein systems (endovascular
30 implants, such as stents), for fastening and temporarily fixing tissue implants and tissue transplantations, but also for orthopedic purposes, such as nails, plates or screws. A particularly frequently used form of an implant is the stent.

The implantation of stents has become established as one of the most effective therapeutic measures for the treatment of vascular diseases. Stents have the purpose of assuming a supporting function in hollow organs of a patient. For this purpose, stents
5 featuring conventional designs have a filigree supporting structure comprising metal struts, which is initially present in compressed form for introduction into the body and is expanded at the site of the application. One of the main application areas of such stents is to permanently or temporarily widen and hold open vascular constrictions, particularly constrictions (stenosis) of coronary blood vessels. In addition, aneurysm stents are
10 known, which are used primarily to seal the aneurysm. The support function is additionally provided.

The implant, notably the stent, has a base body made of an implant material. An implant material is a non-living material, which is employed for applications in medicine and
15 interacts with biological systems. A basic prerequisite for the use of a material as an implant material, which is in contact with the body environment when used as intended, is the body friendliness thereof (biocompatibility). For the purpose of the present application, biocompatibility shall be understood to mean the ability of a material to induce an appropriate tissue reaction in a specific application. This includes an adaptation
20 of the chemical, physical, biological, and morphological surface properties of an implant to the recipient's tissue with the aim of a clinically desired interaction. The biocompatibility of the implant material is also dependent on the temporal process of the reaction of the biosystem in which it is implanted. For example, irritations and inflammations occur in a relatively short time, which can lead to tissue changes.
25 Depending on the properties of the implant material, biological systems thus react in different ways. According to the reaction of the biosystem, the implant materials can be divided into bioactive, bioinert and degradable or resorbable materials.

Implant materials comprise polymers, metallic materials, and ceramic materials (as
30 coatings, for example). Biocompatible metals and metal alloys for permanent implants comprise, for example, stainless steels (such as 316L), cobalt-based alloys (such as CoCrMo cast alloys, CoCrMo forge alloys, CoCrWNi forge alloys and CoCrNiMo forge

alloys), technical pure titanium and titanium alloys (such as cp titanium, TiAl6V4 or TiAl6Nb7) and gold alloys. In the field of biocorrosible stents, the use of magnesium or technical pure iron as well as biocorrosible base alloys of the elements magnesium, iron, zinc, molybdenum, and tungsten are proposed.

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The use of biocorrosible magnesium alloys for temporary implants having filigree structures is made difficult in particular in that the degradation of the implant progresses very quickly in vivo. So as to reduce the corrosion rate, this being the degradation speed, different approaches are being discussed. For one, it is attempted to slow the degradation on the part of the implant material by developing appropriate alloys. In addition, coatings are to bring about a temporary inhibition of the degradation. While the existing approaches are promising, none of them has so far led to a commercially available product. Regardless of the efforts made so far, there is rather a continuing need for solutions that make it possible to at least temporarily reduce the corrosion of magnesium alloys in vivo, while optimizing the mechanical properties thereof at the same time.

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In light of this prior art, the techniques disclosed herein may allow for provision of a biodegradable magnesium alloy, a method for the production thereof and a use for implants, which may allow the magnesium matrix of the implant to remain in an electrochemically stable state over the required support period with fine grain and high corrosion resistance without protective layers, and to utilize the formation of intermetallic phases, which electrochemically are more noble than the magnesium matrix, while also improving the mechanical properties, such as increasing the tensile strength and proof stress, as well as reducing the mechanical asymmetry so as to adjust the degradation rate of the implants.

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These may be achieved by a magnesium alloy, a method, and the use according to the independent claims.

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The characteristics listed in the dependent claims allow advantageous refinements of the magnesium alloy, of the method for the production thereof, and of the use thereof.

The solution may allow for the corrosion resistance and deformability of the magnesium matrix of the implant to be assured over the support period such that the implant is able to absorb multiaxial permanent load without fracture or cracking, and to also utilize the magnesium matrix as a means for the decomposition triggered by the physiological liquids.

This is achieved by the magnesium alloy comprising:
less or equal 4.0% by weight Zn, 2.0 to 10.0% by weight Al, wherein the alloy content of Al in % by weight is greater than or equal to the alloy content of Zn in % by weight,
the remainder being magnesium which contains impurities, which promote electrochemical potential differences and/or the formation of precipitations and/or intermetallic phases, in a total amount of no more than 0.0063% by weight of Fe, Si, Mn, Co, Ni, Cu, Zr, Y, Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to 103, Be, Cd, In, Sn and/or Pb as well as P, wherein the matrix of the alloy is solid solution hardening due to Al and Zn and is also particle hardening due to the intermetallic phases formed of Mg and Al.

Preferably the magnesium alloy according to the this application has a content of Zn less or equal 2.0% by weight, in particular preferably less or equal 1.0% by weight and/or a content of Al in the range of 2.0 to 8.0% by weight, preferably 3.0 to 8.0% by weight and still more preferably 3.0 to 6.0% by weight.

The magnesium alloy according to the subject matter of the patent application may have very high corrosion resistance, which may be achieved by drastically reducing the content of impurities and the combinations thereof in the magnesium matrix, and by having only those alloying elements present which raise the electrochemical potential of the matrix such that the corrosion resistance of the alloy is considerably increased.

The previously known tolerance limits for impurities do not take into account that wrought magnesium alloys often times are subjected to a thermomechanical treatment, and more particularly to an extended annealing process, which creates the near-

equilibrium structures. The metallic elements bond by way of diffusion and form what are known as intermetallic phases, which have a different electrochemical potential, notably a considerably higher potential, than the magnesium matrix, and therefore these intermetallic phases act as cathodes and can trigger galvanic corrosion processes.

5

The applicant has found that a corrosion-stable alloy matrix can be achieved when complying with the following tolerance limits of individual impurities in % by weight:

Fe, Si, Mn, Co, Ni, Cu each with <0.0005; Zr, Y each with <0.0003; and P <0.0002.

Further preferred in this embodiment, the alloy has additional individual impurities in the
10 following tolerance limits (% by weight): Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to 103 in total <0.001; Be, Cd, In, Sn and/or Pb each with <0.0003;.

Preferably the corrosion-stable alloy matrix contains impurities in a total amount of no more than 0.0053 Gew.%, which can be achieved when complying with the following
15 tolerance limits of individual impurities in % by weight:

Fe, Si, Mn each with <0.0005; Co, Ni, Cu each with <0.0002; Zr, Y each with <0.0003; Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to 103 in total <0.001; Be, Cd, In, Sn and/or Pb each with <0.0003; and P <0.0001.

20 In particular the preferred corrosion-stable alloy matrix contains impurities in a total amount of no more than 0.0022 Gew.%, which can be achieved when complying with the following tolerance limits of individual impurities in % by weight:n:

Fe, Si, Mn each with <0.0002; Co, Ni, Cu, Zr, Y each with <0.0001; Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to 103 in total <0.0005; Be, Cd, In, Sn
25 and/or Pb each with <0.0001, and P <0.0001.

It is surprising that the addition of manganese as an alloying element, which is customary in the prior art, can be dispensed with when these tolerance limits are adhered. The formation of the ternary intermetallic phase FeMnSi is suppressed, whereby the corrosion
30 resistance of the alloy is improved.

When the impurity elements are combined, this tolerance limit of the sum of impurities of Fe, Si, Mn, Co, Ni and Cu may be no more than 0.003 % by weight, preferably no more than 0.0021% by weight and in particular preferred no more than 0.0009% by weight.

5 The magnesium alloy according to the subject matter of the patent application may achieve a tensile strength of >275 MPa, and preferably >300 MPa, a yield point of >200 MPa, and preferably >225 MPa, and a yield ratio of <0.8, and preferably <0.75, wherein the difference between the tensile strength and yield point may be >50 MPa, and preferably >100 MPa, and the mechanical asymmetry may be <1.25.

10 These significantly improved mechanical properties can be attributed to the formation of solid solutions between aluminum and zinc in the alloy matrix as well as to the intermetallic phases of magnesium and aluminum occurring in the matrix. The basis for the increased tensile strength is the interaction of the dislocations with the particles, whereby the dislocation movement is adversely impacted and additional tension is
15 required in order to generate the same plastic deformation as in an undisturbed matrix.

The improved mechanical properties of the magnesium alloys may allow the implants, for example cardiovascular stents, to be able to withstand the multiaxial permanent load in the implanted state over the entire support period, despite onsetting degradation of the magnesium matrix due to corrosion.

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For the mechanical asymmetry, it is particularly important for the magnesium alloy to have a particularly fine microstructure having a grain size of no more than 7.5 μm , preferably < 5 μm , and still more preferably < 2.5 μm .

25 In another exemplary technique, there is provided a method for producing a magnesium alloy having improved mechanical and electrochemical properties. The method comprises the following steps:

- a) generating high-purity magnesium by way of vacuum distillation;
- b) generating a billet of the alloy by synthesis of the magnesium according to step a)
30 with less or equal 4.0% by weight Zn, 2.0 to 10.0% by weight Al, wherein the alloy content of Al in % by weight is greater than or equal to the alloy content of Zn in % by weight, the remainder being magnesium containing impurities, which promote

5 electrochemical potential differences and/or the formation of precipitations and/or intermetallic phases, in a total amount of no more than 0.0063% by weight of Fe, Si, Mn, Co, Ni, Cu, Zr, Y, Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to 103, Be, Cd, In, Sn and/or Pb as well as P, wherein the matrix of the alloy is solid solution hardening due to Al and Zn and is also particle hardening due to the intermetallic phases formed of Mg and Al;

- c) homogenizing the alloy by annealing at a temperature between 150°C and 450°C with a holding period of 4 to 40 hours; and
 - d) at least single forming of the homogenized alloy in the temperature range between 200°C and 400°C.
- 10

Preferably steps c) and d) can be repeated at least once.

After step c) and before step d) an ageing treatment step can be performed. Depending on the alloy composition and/or the amount and/or type of the grains and/or the grain size ageing treatment step can be performed at a temperature between 20°C and 300°C with a holding period of 1h to 168h. Preferably the ageing treatment can be performed at a temperature between 20°C and 275°C, still more preferably at a temperature of 150°C with a holding period of 120 hours.

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20 In a preferred value range steps c) is performed at a temperature between 250°C and 450°C and/or step d) is performed at a temperature between 225°C and 400°C.

Preferably the magnesium alloy generated by step a) has a content of Zn less or equal 2.0% by weight, in particular preferably less or equal 1.0% by weight and/or a content of Al in the range of 2.0 to 8.0% by weight, preferably 3.0 to 8.0% by weight and still more preferably 3.0 to 6.0% by weight.

25

Vacuum distillation is preferably used to produce a starting material for a high-purity magnesium-aluminum-zinc alloy having the required threshold values.

The sum of impurities can be selectively adjusted and in % by weight are:

a) for the individual impurities:

5 Fe, Si, Mn, Co, Ni, Cu each with <0.0005 ;
Zr, Y each with <0.0003 ; and
P <0.0002 .

Preferably in this embodiment additional individual impurities in the following tolerance limits (% by weight):

10 Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to 103
in total <0.001 ; Be, Cd, In, Sn and/or Pb each with <0.0003 .

aa) for the individual impurities in a preferred total amount of impurities of no more than 0.0053% by weight,:

15 Fe, Si, Mn each with <0.0005 ;
Co, Ni, Cu each with <0.0002 ;
Zr, Y each with <0.0003 ;
Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to
103 in total <0.001 ;
20 Be, Cd, In, Sn and/or Pb each with <0.0003 ; and
P <0.0001 .

ab) for the individual impurities in a particularly preferred total amount of impurities of no more than 0.0022 % by weight:

25 Fe, Si, Mn each with <0.0002 ;
Co, Ni, Cu, Zr, Y each with <0.0001 ;
Sc or rare earths having the ordinal numbers 21, 57 to 71 and 89 to
103 in total <0.0005 ;
Be, Cd, In, Sn and/or Pb each with <0.0001 ; and
30 P <0.0001 .

b) for the combination of individual impurities in total:

Fe, Si, Mn, Co, Ni, Cu no more than 0.003, preferably no more than 0.0021% by weight and in particular preferred no more than 0.0009% by weight.

5 It is particularly advantageous that the method according to the patent application only requires a small number of forming steps. Extrusion, equal channel angular extrusion and/or multiple forging can thus preferably be employed, which assure that a substantially homogeneous fine grain of $<10\text{ }\mu\text{m}$ is achieved.

10 In another exemplary technique, there is provided a use of the magnesium alloy produced according to the method disclosed herein, which may have the aforescribed advantageous composition and structure, in medical technology, notably for the production of implants, for example endovascular implants such as stents, for fastening and temporarily fixing tissue implants and tissue transplantations, orthopedic and dental
15 implants, and neuroimplants.

All implants in the sense of this patent application are in the Cardiovascular field, osteosynthesis field or other areas.

20 Cardiovascular field in the sense of this application means

- the field of diagnostic, prevention and treatment of all diseases of the cardiovascular system, i.e. heart and blood vessel system,
- by mean of active and non-active implants used to support vessels, and vein systems
- including coronary, cerebral and peripheral vascular implants like stents, valves,
25 closure devices, occluders, clips, coils, staples, implantable regional drug delivery devices,
- implantable electrostimulators (like pacemakers and defibrillators), implantable monitoring devices, implantable electrodes,
- system for fastening and temporarily fixing tissue implants and tissue
30 transplantations
- field also includes any type of stent as mechanical fix or temporary scaffold to support hollow organs (or bodies?) including bones, intervertebral disks

Osteosynthesis in the sense of this application means

- the field of treatment of fractured bones for internal fixation and stabilization by mechanical devices such as metal plates, pins, rods, wires, screws, clips, nails, staples excluding stent technology

Examples of areas out of the osteosynthesis field or the cardiovascular field are:

- Devices for the treatment of diseases of the sinews, joints, muscles, cartilages,
- oral (including dental) and maxillo facial implants (excl. osteosynthesis means),
- esthetic implants,
- supporting tools out of the body, (examples?)
- tissue engineering,
- soft tissue implants,
- devices for wound care,
- suture material and clamps,
- neurosurgery
- local drug delivery (excl. cardiovascular, i.e. lever)
- renal

Exemplary Embodiment 1

A magnesium alloy is to be generated which is composed of 2.0% by weight Zn and 6.0% by weight Al, the remainder being Mg, and contains the following individual impurities in % by weight:

Fe: <0.0005; Si: <0.0005; Mn: <0.0005; Co: <0.0002; Ni: <0.0002; Cu <0.0002, wherein the sum of impurities of Fe, Si, Mn, Co, Ni, Cu is no more than 0.0021% by weight and that of Zr is no more than 0.0003% by weight.

The magnesium produced with aid of vacuum distillation is melted with high-purity Al and Zn in a graphite crucible, and the alloy is subjected to homogenizing annealing at a temperature of 360°C for a duration of 24 hours, and subsequently to multiple extrusion

processes at a temperature of 300°C, so as to produce a precision tube for a cardiovascular stent.

The grain size of the microstructure was $< 5.5 \mu\text{m}$, and the particle size of the intermetallic phases dispersely distributed in the alloy matrix was $0.5 \mu\text{m}$.

- 5 The magnesium alloy reached a tensile strength of 310 to 320 MPa and proof stress of approximately 250 MPa [sic]. The yield ratio was 0.79 and the mechanical asymmetry was 1.2.

Exemplary Embodiment 2

10

A magnesium alloy is to be generated which is composed of 0.25% by weight Zn and 2.50% by weight Al, the remainder being Mg, and contains the following individual impurities in % by weight:

- Fe: <0.0005 ; Si: <0.0005 ; Mn: <0.0005 ; Co: <0.0002 ; Ni: <0.0002 ; Cu <0.0002 , wherein
15 the sum of impurities of Fe, Si, Mn, Co, Ni, Cu is no more than 0.0021% by weight and that of Zr is no more than 0.0003% by weight.

- The magnesium produced with aid of vacuum distillation is melted with high-purity Al and Zn in a graphite crucible, and the alloy is subjected to homogenizing annealing at a
20 temperature of 360°C for a duration of 24 hours, and subsequently to multiple extrusion processes at a temperature of 300°C, so as to produce a precision tube for a cardiovascular stent.

The grain size of the microstructure was $< 5.5 \mu\text{m}$, and the particle size of the intermetallic phases dispersely distributed in the alloy matrix was $0.5 \mu\text{m}$.

- 25 The magnesium alloy reached a tensile strength of 310 to 320 MPa and proof stress of approximately 250 MPa [sic]. The yield ratio was 0.79 and the mechanical asymmetry was 1.2.

Exemplary Embodiment 3

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A magnesium alloy is to be generated which is composed of 5.0% by weight Al, the remainder being Mg, and contains the following individual impurities in % by weight:

Fe: <0.0005; Si: <0.0005; Mn: <0.0005; Co: <0.0002; Ni: <0.0002; Cu <0.0002, wherein the sum of impurities of Fe, Si, Mn, Co, Ni, Cu is no more than 0.0021% by weight and that of Zr is no more than 0.0003% by weight.

5 The magnesium produced with aid of vacuum distillation is melted with high-purity Al in a graphite crucible, and the alloy is subjected to homogenizing annealing at a temperature of 360°C for a duration of 24 hours, and subsequently to multiple extrusion processes at a temperature of 300°C, so as to produce a precision tube for a cardiovascular stent.

The grain size of the microstructure was < 5.5 μm, and the particle size of the
10 intermetallic phases dispersely distributed in the alloy matrix was 0.5 μm.

The magnesium alloy reached a tensile strength of 310 to 320 MPa and proof stress of approximately 250 MPa [sic]. The yield ratio was 0.79 and the mechanical asymmetry was 1.2.

15 Exemplary Embodiment 4

A magnesium alloy is to be generated which is composed of 3% by weight Al, the remainder being Mg, and contains the following individual impurities in % by weight:

Fe: <0.0005; Si: <0.0005; Mn: <0.0005; Co: <0.0002; Ni: <0.0002; Cu <0.0002, wherein
20 the sum of impurities of Fe, Si, Mn, Co, Ni, Cu is no more than 0.0021% by weight and that of Zr is no more than 0.0003% by weight.

The magnesium produced with aid of vacuum distillation is melted with high-purity Al in a graphite crucible, and the alloy is subjected to homogenizing annealing at a temperature
25 of 360°C for a duration of 24 hours, and subsequently to multiple extrusion processes at a temperature of 300°C, so as to produce a precision tube for a cardiovascular stent.

The grain size of the microstructure was < 5.5 μm, and the particle size of the intermetallic phases dispersely distributed in the alloy matrix was 0.5 μm.

The magnesium alloy reached a tensile strength of 310 to 320 MPa and proof stress of
30 approximately 250 MPa [sic]. The yield ratio was 0.79 and the mechanical asymmetry was 1.2.

Exemplary Embodiment 5

A magnesium alloy is to be generated which is composed of 0.25% by weight Zn and 2.0% by weight Al, the remainder being Mg, and contains the following individual
5 impurities in % by weight:

Fe: <0.0005; Si: <0.0005; Mn: <0.0005; Co: <0.0002; Ni: <0.0002; Cu <0.0002, wherein the sum of impurities of Fe, Si, Mn, Co, Ni, Cu is no more than 0.0021% by weight and that of Zr is no more than 0.0003% by weight.

10 The magnesium produced with aid of vacuum distillation is melted with high-purity Al and Zn in a graphite crucible, and the alloy is subjected to homogenizing annealing at a temperature of 360°C for a duration of 24 hours and thereafter to an ageing treatment at 125°C for 120 hours.

Subsequently the material is subjected to multiple extrusion processes at a temperature of
15 200°C, so as to produce a precision tube for a cardiovascular stent.

Before the final extrusion step is applied another annealing process is performed at 150°C for 3 hours.

The grain size of the microstructure was < 5.5 µm, and the particle size of the intermetallic phases dispersely distributed in the alloy matrix was 0.5 µm.

20 The magnesium alloy reached a tensile strength of 320 to 350 MPa and proof stress of approximately 235 MPa [sic]. The yield ratio was 0.70 and the mechanical asymmetry was 1.2.

Exemplary Embodiment 6

25 A magnesium alloy is to be generated which is composed of 1.5% by weight Zn and 3.0% by weight Al, the remainder being Mg, and contains the following individual impurities in % by weight:

Fe: <0.0005; Si: <0.0005; Mn: <0.0005; Co: <0.0002; Ni: <0.0002; Cu <0.0002, wherein
30 the sum of impurities of Fe, Si, Mn, Co, Ni, Cu is no more than 0.0021% by weight and that of Zr is no more than 0.0003% by weight.

The magnesium produced with aid of vacuum distillation is melted with high-purity Al and Zn in a graphite crucible, and the alloy is subjected to homogenizing annealing at a temperature of 360°C for a duration of 24 hours and thereafter to an ageing treatment at 150°C for 120 hours.

- 5 Subsequently the material is subjected to an extrusion process at a temperature of 200°C, so as to produce a rod with 8 mm diameter to produce screws for craniofacial fixations.

The grain size of the microstructure was $< 3.0 \mu\text{m}$, and the particle size of the intermetallic phases dispersely distributed in the alloy matrix was $0.5 \mu\text{m}$.

- 10 The magnesium alloy reached a tensile strength of 340 to 360 MPa and proof stress of approximately 250 MPa [sic]. The yield ratio was 0.71 and the mechanical asymmetry was 1.2.

Patent Claims

1. A magnesium alloy having improved mechanical and electrochemical properties, comprising: less or equal 4.0% by weight Zn, 2.0 to 10.0% by weight Al, the alloy
5 content of Al in % by weight being greater than or equal to the alloy content of Zn in % by weight, the remainder being magnesium containing impurities, which promote electrochemical potential differences and/or the formation of precipitations and/or intermetallic phases, in a total amount of no more than 0.0063% by weight of Fe, Si, Mn, Co, Ni, Cu, Zr, Y, Sc or rare earths having the ordinal numbers 21,
10 57 to 71 and 89 to 103, Be, Cd, In, Sn and/or Pb as well as P, wherein the matrix of the alloy is solid solution hardening due to Al and Zn and is also particle hardening due to the intermetallic phases formed of Mg and Al.
2. The magnesium alloy according to claim 1, characterized in that the content of Zn
15 less or equal 2.0% by weight, in particular preferably less or equal 1.0% by weight and/or the content of Al is 2.0 to 8.0% by weight, preferably 3.0 to 8.0% by weight and still more preferably 3.0 to 6.0% by weight.
3. The magnesium alloy according to claim 1, characterized in that the individual
20 impurities in the total sum of impurities amount to the following in % by weight: Fe, Si, Mn, Ni, Co, Cu each with <0.0005; Zr;Y each with <0.0003; and P <0.0002.
4. The magnesium alloy according to claim 1, characterized in that when the impurity
25 elements Fe, Si, Mn, Co, Ni, and Cu are combined, the sum of these impurities is no more than 0.003% by weight.
5. The magnesium alloy according to claim 1 to 4, characterized in that the alloy has a
fine-grained microstructure having a grain size of no more than 7.5 μm , preferably
30 < 5 μm , and still more preferably < 2.5 μm .
6. The magnesium alloy according to claim 1 to 5, characterized in that it has a tensile
strength of >275 MPa, and preferably >300 MPa, a yield point of >200 MPa, and

preferably >225 MPa, and a yield ratio of <0.8, and preferably <0.75, wherein the difference between the tensile strength and yield point is >50 MPa, and preferably >100 MPa, and the mechanical asymmetry is <1.25.

- 5 7. A method for producing a magnesium alloy having improved mechanical and electrochemical properties, comprising the following steps:
- a) generating a high-purity magnesium by way of vacuum distillation;
 - b) generating a billet of the alloy by synthesis of the magnesium according to step
 - a) with less or equal 4.0% by weight Zn, 2.0 to 10.0% by weight Al, wherein
- 10 the alloy content of Al in % by weight is greater than or equal to the alloy content of Zn in % by weight, the remainder being magnesium containing impurities, which promote electrochemical potential differences and/or the formation of precipitations and/or intermetallic phases, in a total amount of no more than 0.0063% by weight of Fe, Si, Mn, Co, Ni, Cu, Zr, Y, Sc or rare
- 15 earths having the ordinal numbers 21, 57 to 71 and 89 to 103, Be, Cd, In, Sn and/or Pb as well as P, wherein the matrix of the alloy is solid solution hardening due to Al and Zn and is also particle hardening due to the intermetallic phases formed of Mg and Al;- c) homogenizing the alloy by annealing at a temperature between 150°C and

20 450°C with a holding period of 4 to 40 hours;- d) at least single forming of the homogenized alloy in the temperature range between 200°C and 400°C.

25 8. The method according to claim 7, characterized in that in the generated billet the content of Zn is less or equal 2.0% by weight, in particular preferably less or equal 1.0% by weight and/or the content of Al is 2.0 to 8.0% by weight, preferably 3.0 to 8.0% by weight and still more preferably 3.0 to 6.0% by weight.

30 9. The method according to claim 7, characterized in that the individual impurities in the total sum of impurities amount to the following in % by weight: Fe, Si, Mn, Ni, Co, Cu each with <0.0005; Zr, Y each with <0.0003; and P <0.0002.

10. The method according to claim 7, characterized in that when the impurity elements Fe, Si, Mn, Co, Ni, and Cu are combined, the sum of these impurities is no more than 0.003% by weight.
- 5 11. The method according to claim 7, characterized in that the forming process is an extrusion, equal channel angular extrusion (EACE) and/or multiple forging process.
12. The method according to claim 7, characterized in that the steps c) and d) are repeated at least once.
- 10 13. The method according to claim 7 or 12, characterized in that the steps c) is performed at a temperature between 250°C and 450°C and/or step d) is performed at a temperature between 225°C and 400°C.
- 15 14. Use of a magnesium alloy according to any one of the claims 1 to 6 for producing a biodegradable implant.
15. Use of a magnesium alloy according to claim 14, characterized in that the implant is elected from the group of endovascular implants such as stents, implants for fastening and temporarily fixing tissue implants and tissue transplantations, orthopedic and dental implants, and neuroimplants.
- 20 16. Use of a magnesium alloy produced by the method according to any one of the claims 7 to 13 for producing a biodegradable implant.
- 25 17. Use of a magnesium alloy according to claim 16, characterized in that the implant is elected from the group of endovascular implants such as stents, implants for fastening and temporarily fixing tissue implants and tissue transplantations, orthopedic and dental implants, and neuroimplants.