(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization

International Bureau





(10) International Publication Number WO 2017/017483 A1

(43) International Publication Date 2 February 2017 (02.02.2017)

(51) International Patent Classification: C23C 2/00 (2006.01) C23C 2/12 (2006.01)

(21) International Application Number:

PCT/IB2015/001281

(22) International Filing Date:

30 July 2015 (30.07.2015)

(25) Filing Language:

English

(26) Publication Language:

English

- (71) Applicant: **ARCELORMITTAL** [LU/LU]; 24-26, Boulevard d'Avranches, L-1160 Luxembourg (LU).
- (72) Inventors: ALLELY, Christian; 30, rue de l'Aubépine, F-57000 Metz (FR). MACHADO AMORIM, Tiago; 46 rue du Général De Gaulle, F-57050 Longeville-les-Metz (FR). VAN DEN BERGH, Krista, Godelieve, Oscar; Doornstraat 71 C, B-9170 Sint-Gillis-Waas (BE). DE STRYCK-ER, Joost; Kil 25, BE-9240 Zele (BE).
- (74) Agent: PLAISANT, Sophie; ArcelorMittal France, Research & Development Intellectual Property, 6, rue André Campra - Immeuble le Cézanne, F-93200 Saint-Denis (FR).

- (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.
- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

with international search report (Art. 21(3))



WO 2017/017483 PCT/IB2015/001281

Steel sheet coated with a metallic coating based on aluminum

The present invention relates to a steel sheet coated with a metallic coating. The invention is particularly well suited for the manufacture of automotive vehicles.

5

10

15

20

25

30

Zinc-based coatings are generally used because they allows for a protection against corrosion thanks to barrier protection and cathodic protection. The barrier effect is obtained by the application of a metallic coating on steel surface. Thus, metallic coatings prevent the contact between steel and corrosive atmosphere. The barrier effect is independent from the nature of coating and substrate. On the contrary, sacrificial cathodic protection is based on the fact that zinc is a metal less noble that steel. Thus, if corrosion occurs, zinc is consumed preferentially to steel. Cathodic protection is essential in areas where steel is directly exposed to corrosive atmosphere, like cut edges where surrounding zinc will be consumed before steel.

However, when press hardening process is performed on such zinc coated steel sheets, for example by hot-stamping, microcracks are observed in steel which spread from the coating. Additionally, the step of painting of some hardened parts coated with zinc necessitates sanding operations before phosphatation due to the presence of a weak layer of oxides at the part surface.

Other metallic coatings usually used for the production of automotive vehicle are aluminum and silicon based coatings. There is no microcrack in steel when press hardening process is performed due to the presence of an intermetallic layer Al-Si-Fe. Moreover, they have a good aptitude for painting. They allow for a protection by barrier effect and can be welded. However, they do not allow for a cathodic protection or they have a very low cathodic protection.

The patent application EP1225246 discloses a Zn-Al-Mg-Si alloy-plated material wherein the coating comprises, in terms of weight%, Al: at least 45% and no greater than 70%, Mg: at least 3% and less than 10%, Si: at least 3% and less than 10%, with the remainder Zn and unavoidable impurities, wherein the Al/Zn ratio is 0.89-2.75 and the plating layer contains a bulky Mg₂Si phase. It also discloses a Zn-Al-Mg-Si alloy-plated steel material wherein the coating comprises, in terms of weight%, Al: at least 45% and no greater than 70%, Mg: at least 1% and less than 5%, Si: at least 0.5% and less than 3%, with the remainder Zn and

10

15

20

25

30

unavoidable impurities, wherein the Al/Zn ratio is 0.89-2.75 and the plating layer contains a scaly Mg₂Si phase. These specific coatings show unpainted corrosion resistance and edge creep resistance at cut edge sections after painting.

However, the fabrication of specific Mg_2Si phases, scaly or bulky, is complex. Indeed, it depends on the size and on the ratio of the short diameter mean size with respect to the long diameter of Mg_2Si phases, as observed with a 5° polished cross-section. The size is affected most predominantly by the cooling rate after hot-dip plating. Moreover, the fabrication of Mg_2Si phases also depends on the quantity of Mg and Si.

From an industrial point of view, Mg₂Si phases can be difficult to obtain because of these specifics criteria. Therefore, there is a risk that the desired Mg₂Si phase is not obtained.

The purpose of the invention is to provide a coated steel sheet easy to form having a reinforced protection against corrosion, i.e. a sacrificial cathodic protection in addition to barrier protection, before and after the forming.

In terms of sacrificial protective corrosion, electrochemical potential has to be at least 50mV more negative than the potential of steel, i.e. a maximum potential of -0.78V with respect to a saturated calomel electrode (SCE). It is preferable not to decrease the potential at a value of -1.4V/SCE, even -1.25V/SCE which would involve a fast consumption and would finally decrease the period of protection of steel.

This is achieved by providing a steel sheet coated with a metallic coating according to claim 1. The coated steel sheet can also comprise any characteristics of claims 2 to 19 taken alone or in combination.

The invention also covers parts coated with a metallic coating having a sacrificial cathodic protection according to claim 20.

The invention also covers the use of a coated part for the manufacture of an automotive vehicle according to claim 21.

To illustrate the invention, various embodiments and trials of non-limiting examples will be described, particularly with reference to the following Figure:

Figure 1 illustrates one corrosion cycle corresponding to 168 hours of the norm VDA 233-102.

10

15

20

25

30

Other characteristics and advantages of the invention will become apparent from the following detailed description of the invention.

Any steel can be advantageously used in the frame of the invention. However, in case steel having high mechanical strength is needed, in particular for parts of structure of automotive vehicle, steel having a tensile resistance superior to 500MPa, advantageously between 500 and 2000MPa before or after heat-treatment, can be used. The weight composition of steel sheet is preferably as follows: $0.03\% \le C \le 0.50\%$; $0.3\% \le Mn \le 3.0\%$; $0.05\% \le Si \le 0.8\%$; $0.015\% \le Ti \le 0.2\%$; $0.005\% \le Al \le 0.1\%$; $0\% \le Cr \le 2.50\%$; $0\% \le S \le 0.05\%$; $0\% \le P \le 0.1\%$; $0\% \le B \le 0.010\%$; $0\% \le Ni \le 2.5\%$; $0\% \le Mo \le 0.7\%$; $0\% \le Nb \le 0.15\%$; $0\% \le N \le 0.015\%$; $0\% \le Ca \le 0.01\%$; $0\% \le W \le 0.35\%$, the balance being iron and unavoidable impurities from the manufacture of steel.

For example, the steel sheet is 22MnB5 with the following composition: $0.20\% \le C \le 0.25\%$; $0.15\% \le Si \le 0.35\%$; $1.10\% \le Mn \le 1.40\%$; $0\% \le Cr \le 0.30\%$; $0\% \le Mo \le 0.35\%$; $0\% \le P \le 0.025\%$; $0\% \le S \le 0.005\%$; $0.020\% \le Ti \le 0.060\%$; $0.020\% \le Al \le 0.060\%$; $0.002\% \le B \le 0.004\%$, the balance being iron and unavoidable impurities from the manufacture of steel.

The steel sheet can be Usibor®2000 with the following composition: 0.24% \leq C \leq 0.38%; 0.40% \leq Mn \leq 3%; 0.10% \leq Si \leq 0.70%; 0.015% \leq Al \leq 0.070%; 0 % \leq Cr \leq 2%; 0.25% \leq Ni \leq 2%; 0.020% \leq Ti \leq 0.10%; 0% \leq Nb \leq 0.060%; 0.0005% \leq B \leq 0.0040%; 0.003% \leq N \leq 0.010%; 0.0001% \leq S \leq 0.005%; 0.0001% \leq P \leq 0.025%; it being understood that the contents of titanium and nitrogen satisfy Ti/N \geq 3.42; and that the contents of carbon, manganese, chromium and silicon satisfy:

$$2,6C + \frac{Mn}{5.3} + \frac{Cr}{13} + \frac{Si}{15} \ge 1,1\%$$

the composition optionally comprising one or more of the following: $0.05\% \le Mo \le 0.65\%$; $0.001\% \le W \le 0.30\%$; $0.0005\% \le Ca \le 0.005\%$, the balance being iron and unavoidable impurities from the manufacture of steel.

For example, the steel sheet is Ductibor®500 with the following composition: $0.040\% \le C \le 0.100\%$; $0.80\% \le Mn \le 2.00\%$; $0\% \le Si \le 0.30\%$; 0%

 $0\% \le Cr \le 0.100\%$; $0\% \le Mo \le 0.100\%$; $0\% \le Ca \le 0.006\%$, the balance being iron and unavoidable impurities from the manufacture of steel.

Steel sheet can be obtained by hot rolling and optionally cold rolling depending on the desired thickness, which can be for example between 0.7 and 3.0mm.

5

10

15

20

25

30

The invention relates to a steel sheet coated with a metallic coating comprising from 2.0 to 24.0% by weight of zinc, from 7.1 to 12.0% by weight of silicon, optionally from 1.1 to 8.0% by weight of magnesium, and optionally additional elements chosen from Pb, Ni, Zr, or Hf, the content by weight of each additional element being inferior to 0.3% by weight, the balance being aluminum and optionally unavoidable impurities and residuals elements, wherein the ratio Al/Zn is above 2.9. Metallic coatings according to the invention have a high sacrificial protection.

Preferably, the metallic coating does not comprise elements selected among Cr, Mn, Ti, Ce, La, Nd, Pr, Ca, Bi, In, Sn and Sb or their combinations. In another preferred embodiment, the metallic coating does not comprise any of the following compounds: Cr, Mn, Ti, Ce, La, Nd, Pr, Ca, Bi, In, Sn and Sb. Indeed, without willing to be bound by any theory, it seems that when these compounds are present in the coating, there is a risk that the properties of the coating, such as electrochemical potential, are altered, because of their possible interactions with the essential elements of the coatings.

Preferably, the ratio Al/Zn is below or equal to 8.5. Preferably, the ratio Al/Zn is between 3.0 and 7.5, advantageously between 4.0 and 6.0. Without willing to be bound by any theory, it seems that if these conditions are not met, there is a risk that the sacrificial protection decreases because zinc rich phases are not in sufficient amount in the coating.

In a preferred embodiment, the coating layer further comprises an Al-Zn phase.

Advantageously, the metallic coating comprises from 10.0 to 20.0%, preferably from 10.0 to 15.0%, by weight of zinc.

Preferably, the metallic coating comprises from 8.1 to 10.0% by weight of silicon.

10

15

20

25

30

PCT/IB2015/001281

Advantageously, the coating comprises from 3.0 to 8.0% by weight of magnesium, preferably, from 3.0 to 5.0% by weight of magnesium. Without willing to be bound by any theory, it has been found that the addition of magnesium in the above range further improve the anti-corrosion properties.

Preferably, the microstructure of said coating comprising a Mg₂Si phase. In another preferred embodiment, the microstructure of said coating further comprises a MgZn₂ phase.

Advantageously, the amount of aluminum is above 71%, preferably above 76%, by weight.

The coating can be deposited by any methods known to the man skilled in the art, for example hot-dip galvanization process, electrogalvanization process, physical vapour deposition such as jet vapor deposition or sputtering magnetron. Preferably, the coating is deposited by hot-dip galvanization process. In this process, the steel sheet obtained by rolling is dipped in a molten metal bath.

The bath comprises zinc, silicon, aluminum and optionally magnesium. It can comprise additional elements chosen from Pb, Ni, Zr, or Hf, the content by weight of each additional element being less than 0.3% by weight. These additional elements can improve among others ductibility, coating adhesion on the steel sheet.

The bath can also contain unavoidable impurities and residuals elements from feeding ingots or from the passage of the steel sheet in the molten bath. Residual element can be iron with a content up to 3.0% by weight.

The thickness of the coating is usually between 5 and 50µm, preferably between 10 and 35µm, advantageously between 12 and 18µm or between 26 to 31µm. The bath temperature is usually between 580 and 660°C.

After the deposition of the coating, the steel sheet is usually wiped with nozzles ejecting gas on both sides of the coated steel sheet. The coated steel sheet is then cooled. Preferably, the cooling rate is above or equal to 15°C.s⁻¹ between the beginning of the solidification and the end of the solidification. Advantageously, the cooling rate between the beginning and the end of the solidification is superior or equal to 20°C.s⁻¹.

Then, a skin-pass can be realized and allows work hardening the coated steel sheet and giving it a roughness facilitating the subsequent shaping. A

10

15

30

degreasing and a surface treatment can be applied in order to improve for example adhesive bonding or corrosion resistance.

Then, the coated steel sheet according to the invention can be shaped by any method known to the man skilled in the art, for example cold-stamping and/or hot-forming.

In a preferred embodiment, the part is obtained by cold-stamping. In this case, the coated steel sheet is cut to obtain a blank and then cold-stamped in order to obtain a part.

In another preferred embodiment, the part coated is obtained by a press hardening process including the hot-forming. In this case, this method comprises the following steps:

- A) the provision of a steel sheet pre-coated with a metallic coating comprising from 2.0 to 24.0% by weight of zinc, from 7.1 to 12.0% by weight of silicon, optionally from 1.1 to 8.0% by weight of magnesium, and optionally additional elements chosen from Pb, Ni, Zr, or Hf, the content by weight of each additional element being less than 0.3% by weight, the balance being aluminum and unavoidable impurities and residuals elements, wherein the ratio Al/Zn is above 2.9,
- B) the cutting of the coated steel sheet to obtain a blank,
- 20 C) the thermal treatment of the blank at a temperature between 840 and 950°C to obtain a fully austenitic microstructure in the steel,
 - D) the transfer of the blank into a press tool,
 - E) the hot-forming of the blank to obtain a part,
- F) the cooling of the part obtained at step E) in order to obtain a microstructure in steel being martensitic or martensito-bainitic or made of at least 75% of equiaxed ferrite, from 5 to 20% of martensite and bainite in amount less than or equal to 10%.

Indeed, after, the provision of steel sheet pre-coated with the metallic coating according to the present invention the cutting to obtain a blank. A thermal treatment is applied to the blank in a furnace under non protective atmosphere at an austenitization temperature Tm usually between 840 and 950°C, preferably 880 to 930°C. Advantageously, said blank is maintained during a dwell time tm

10

15

20

25

30

between 1 to 12 minutes, preferably between 3 to 9 minutes. During the thermal treatment before the hot-forming, the coating forms an alloy layer having a high resistance to corrosion, abrasion, wear and fatigue.

After the thermal treatment, the blank is then transferred to a hot-forming tool and hot-formed at a temperature between 600 and 830°C. The hot-forming comprises the hot-stamping and the roll-forming. Preferably, the blank is hot-stamped. The part is then cooled in the hot-forming tool or after the transfer to a specific cooling tool.

The cooling rate is controlled depending on the steel composition, in such a way that the final microstructure after the hot-forming comprises mostly martensite, preferably contains martensite, or martensite and bainite, or is made of at least 75% of equiaxed ferrite, from 5 to 20% of martensite and bainite in amount less than or equal to 10%.

A coated part according to the invention can thus obtained by cold or hot forming but also by any suitable combination of cold-stamping and hot-forming.

For automotive application, after phosphating step, the part is dipped in an e-coating bath. Usually, the thickness of the phosphate layer is between 1 and 2 μ m and the thickness of the e-coating layer is between 15 and 25 μ m, preferably inferior or equal to 20 μ m. The cataphoresis layer ensures an additional protection against corrosion.

After the e-coating step, other paint layers can be deposited, for example, a primer coat of paint, a basecoat layer and a top coat layer.

Before applying the e-coating on the part, the part is previously degreased and phosphated so as to ensure the adhesion of the cataphoresis.

The invention will now be explained in trials carried out for information only. They are not limiting.

Examples

For all samples, steel sheets used are 22MnB5. The composition of the steel is as follows: C = 0.2252%; Mn = 1.1735%; P = 0.0126%, S = 0.0009%; N = 0.0037%; Si = 0.2534%; Cu = 0.0187%; Si = 0.0197%; Si = 0.180%; Si = 0.004%; Si = 0.00371%; Si = 0.00371%

All coatings were deposited by hot-dip galvanization process. All coatings have a thickness of 15µm.

Example 1: Cut edge potential test:

Trials 1 to 4 were prepared and subjected to an electrochemical potential test.

A test consisting in measuring the cut edges potential of coated steel sheet was realized. To this end, each steel sheet was dipped in a solution comprising 2.43% by weight of sodium sulfate and 0.1% by weight of sodium chloride. A saturated calomel electrode (SCE) was also immersed into the solution. The coupling potential of cut edges was measured. Results are shown in the following Table 1:

Trials		Coatir	ng		Thickness	Coupling	
IIIais	Al Si		Zn	Mg	(µm)	potential(V/SCE)	
1*	81	9	10	-	15	- 0.84	
2*	77	9	10	4	15	- 0.84	
3*	73	9.	10	8	15	- 0.84	
4	91	9	_	-	15	- 0.625	

^{*:} examples according to the invention.

15

25

5

10

Trials according to the invention (Trials 1 to 3) have a lower coupling potential than a coating comprising aluminum and 9% by weight of silicon. Coupling potentials of Trials 1 to 3 are under -0.78V/SCE as required.

20 Example 2: Cut edge corrosion test:

Trials 5 to 12 were prepared and subjected to a corrosion test to evaluate the cut edge protection of the coated steel sheets.

All trials were dipped in a solution comprising 2.43% by weight of sodium sulfate and 0.1% by weight of sodium chloride during 50 hours. The presence of corrosion on cut edges of coated steel sheet was observed with the naked eye: 0 means excellent, in other words, there is little or no corrosion and 5 means very

bad, in other words, there are is a lot of corrosion on the cut edges. Results are shown in the following Table 2:

Trials		Coati	ng	Thickness	Corrosion	
IIIais	Al	Si	Zn	Mg	(µm)	0011031011
5*	86	9	5	-	15	2
6*	81	9	10	_	15	1.5
7*	71	9	20	-	15	1
8*	77	9	10	4	15	0
9*	73	9	10	8	15	0
10*	67	9	20	4	15	0
11*	63	9	20	8	15	0
12	91	9	_	-	15	5

^{*:} examples according to the invention.

5

10

15

Trials 5 to 11 have very good protection against corrosion on the cut edges of coated steel sheet. By contrast, Trial 12 does not show enough corrosion resistance on the cut edges.

Example 3: Electrochemical behavior test:

Trials 13 to 16 were prepared and subjected to an electrochemical potential test.

A test consisting in measuring the electrochemical potential of the coated steel surface sheet was realized. Steel sheets and coatings were separated and dipped in a solution comprising 5% by weight of sodium chloride at pH 7. A saturated calomel electrode (SCE) was also immersed into the solution. The coupling potential of the surface was measured over time. Results are shown in the following Table 3:

		Coating			Thickness	Coupling
Trials	Al	Si	Zn	Mg	(µm)	potential
	ΛI	Oi	Z11	ivig		(V/SCE)
13*	81	9	10		15	- 0.98
14*	.77	9	10	4	15	- 0.98
15*	73	9	10	8	15	- 0.99
16	0.2	-	99.8	1	7	- 1.00

^{*:} examples according to the invention.

10

15

20

Trials 13 to 15 are sacrificial such as zinc coating. Coupling potential are under -0.78V/SCE as required.

Example 4: Corrosion test:

Trials 17 to 20 were prepared and subjected to a corrosion test to evaluate the protection of the coated steel sheets.

A test, consisting in submitting coated steel sheet to corrosion cycles according to the norm VDA 233-102, was realized. At this end, trials were put in a chamber wherein an aqueous solution of sodium chloride of 1% by weight was vaporized on trials with a rate of flow of 3mL.h⁻¹. The temperature varied from 50 to -15°C and the humidity rate varied from 50 to 100%. Figures 1 illustrates one cycle corresponding to 168 hours, i.e. one week.

The presence of corrosion on coated steel sheet was observed by naked eyes: 0 means excellent, in other words, there is little or no corrosion and 5 means very bad, in other words, there is a lot of corrosion. Results are shown in the following Table 4:

	Coating		Thickness	Number of cycles							
Trials	Al	Si	Zn	Mg	(µm)	1	5	7	10	15	20
17*	81	9	10	-	15	0	0	0.5	1	3	4
18*	77	9	10	4	15	0	0	0	0	0	0
19*	73	9	10	8	15	0	0	0	0	0	0
20	0.2	-	99.8	-	7	0	2	4	ND	ND	ND

^{*:} examples according to the invention, ND: not done.

Trials 17 to 19 show excellent protection against corrosion, in particular when the coating comprises magnesium (Trials 18 and 19).

Example 5: Corrosion test on scratched trials:

Trials 21 to 24 were prepared and subjected to a corrosion test to evaluate the protection of the coated steel sheets.

Firstly, all trials were scratched on a width of 0.5, 1 and 2mm. then, all trials were submitted to corrosion cycles according to the norm VDA 233-102 represented in Figure 1.

The presence of corrosion on coated steel sheet around scratches was observed by naked eyes: 0 means excellent, in other words, there is little or no corrosion around scratch and 5 means very bad, in other words, there is a lot of corrosion around scratch. Results are shown in the following Table 5:

	Coating			Thickness (µm)		Nur	mber	of c	ycle	es	
Trials	Al	Si	Zn	Mg	15	1	2	3	4	5	6
21*	81	9	10	_	15	0	0	0.5	1	2	3
22*	77	9	10	4	15	0	0	0	0	0	0
23*	73	9	10	8	15	0	0	0	0	0	0.5
24	0.2	-	99.8		10	0	0	0	1	2	3

^{*:} examples according to the invention.

10

15

O 2017/017483 PCT/IB2015/001281

Trials according to the invention (Trials 21 to 23) have an excellent protection against corrosion, in particular when the coating comprises magnesium (Trial 22 and 23).

Example 6: Corrosion test on heat treated and scratched trials:

Trials 25 to 28 were prepared and subjected to a corrosion test to evaluate the protection of the coated steel sheets after austenitization treatment.

All trials were cut in order to obtain a blank. Blanks were then heated at a temperature of 900°C during a dwell time varying between 5 and 10 minutes. Blanks were transferred into a press tool and hot-stamped in order to obtain parts. Then, parts were cooled to obtain a hardening by martensitic transformation. All trials were submitted to 6 corrosion cycles according to the norm VDA 233-102 represented in Figure 1.

The presence of corrosion on coated steel sheet around scratches was observed by naked eyes: 0 means excellent, in other words, there is little or no corrosion around scratch and 5 means very bad, in other words, there is a lot of corrosion around scratch. Results are shown in the following Table 6:

		Coa	Thickness (µm)	Dwell time (min)			
Trials	Al	Si	Zn	Mg	(μπ)	5	10
25*	71	9	20	_	15	1	1
26*	77	9	10	4	15	0.5	0.5
27*	73	9	10	8	15	2	3
28	91	9	_	-	15	5	5

^{*:} examples according to the invention.

20

5

10

15

Trials 25 to 27 show good protection against corrosion compared to the coating comprising aluminum and silicon (Trial 28).

10

15

Example 7: Electrochemical behavior test:

Trials 29 to 40 were prepared and subjected to an electrochemical potential test after austenitization treatment.

All trials were cut in order to obtain a blank. Blanks were then heated at a temperature of 900°C during a dwell time of 5 minutes. Blanks were transferred into a press tool and hot-stamped in order to obtain parts. Then, parts were cooled to obtain a hardening by martensitic transformation.

A test consisting in measuring the electrochemical potential of the coated steel surface sheet was realized. Steel sheets and coatings were separated and dipped in a solution comprising 5% by weight of sodium chloride at pH 7. A saturated calomel electrode (SCE) was also immersed into the solution. The power of sacrificial protection, also called galvanic coupling, was measured over time. In other words, it has been assessed how long the coating remains sacrificial in these conditions. Results are shown in the following Table 7:

		Coating			Thickness	Galvanic
Trials	Al	Si	Zn	Mg	(µm)	coupling
	ΛI) Oi	211	ivig		(hours)
29	88	2	10	-	15	0
30	83	2	15	_	15	0
31	80	5	15	-	15	0
32*	81	9	10	_	15	16
33*	77	9	10	4	15	45
34*	73	9	10	8	15	7
35*	76	9	15	-	15	26
36*	83	9	15	2	15	84
37*	71	9	20	-	15	140
38*	67	9	20	4	15	91
39*	63	9	20	8	15	14
40	91	. 9	-	-	15	0

^{*:} examples according to the invention.

Trials 32 to 39 according to the present invention are and remain sacrificial protection over time.

CLAIMS

1. Steel sheet coated with a metallic coating comprising from 2.0 to 24.0% by weight of zinc, from 7.1 to 12.0% by weight of silicon, optionally from 1.1 to 8.0% by weight of magnesium, and optionally additional elements chosen from Pb, Ni, Zr, or Hf, the content by weight of each additional element being inferior to 0.3% by weight, the balance being aluminum and optionally unavoidable impurities and residuals elements, wherein the ratio Al/Zn is above 2.9.

10

5

- 2. Steel sheet according to claim 1, wherein the ratio Al/Zn is below or equal to 8.5.
- 3. Steel sheet according to claim 2, wherein the ration Al/Zn is between 3.0 and 7.5.
 - 4. Steel sheet according to claim 3, wherein the ratio Al/Zn is between 4.0 and 6.0.
- 5. Steel sheet according to anyone of claims 1 to 4, wherein the metallic coating microstructure comprises an Al-Zn phase.
 - 6. Steel sheet according to anyone of claims 1 to 5, wherein the coating comprises from 10.0 to 20.0% by weight of zinc.

25

- 7. Steel sheet according to claim 6, wherein the coating comprises from 10.0 to 15.0% by weight of zinc.
- 8. Steel sheet according to anyone of claims 1 to 7, wherein the coating comprises from 8.1 to 10.0% by weight of silicon.

- 9. Steel sheet according to anyone of claims 1 to 8, wherein the coating comprises from 3.0 to 8.0% by weight of magnesium.
- 10. Steel sheet according to claim 9, wherein the coating comprises from 3.0 to 5.0% by weight of magnesium.

15

30

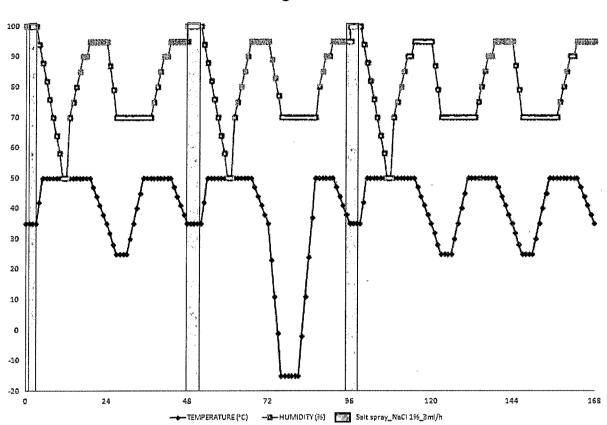
- 11. Steel sheet according to anyone of claims 1 to 10, wherein the microstructure of said coating comprising a Mg₂Si phase.
- 10 12. Steel sheet according to anyone of claims 1 to 11, wherein the microstructure of said coating comprising a MgZn₂ phase.
 - 13. Steel sheet according to anyone of claims 1 to 12, wherein the amount of aluminum is above 71% by weight.
 - 14. Steel sheet according to anyone of claims 1 to 13, wherein the amount of aluminum is above 76% by weight.
- 15. Steel sheet according to anyone of claims 1 to 14, wherein the thickness of the coating is between 5 and 50µm.
 - 16. Steel sheet according to claim 15, wherein the thickness is between 10 and 35µm.
- 17. Steel sheet according to claim 16, wherein the thickness between 12 and 18μm.
 - 18. Steel sheet according to claim 16, wherein the thickness between 26 and 31µm.

- 19. Steel sheet according to anyone of claims 1 to 18, wherein the coating does not comprise elements selected among Cr, Mn, Ti, Ce, La, Nd, Pr, Ca, Bi, In, Sn and Sb or their combinations.
- 5 20.Part coated with a metallic coating obtainable from the coated steel sheet according to anyone of claims 1 to 19 by hot-forming or cold-stamping or both.
 - 21. Use of a part according to claim 20, for the manufacture of automotive vehicle.

WO 2017/017483 PCT/IB2015/001281

1/1





INTERNATIONAL SEARCH REPORT

International application No PCT/IB2015/001281

A. CLASSIFICATION OF SUBJECT MATTER INV. C23C2/00 C23C2/12 ADD.									
According to International Patent Classification (IPC) or to both national classification and IPC									
B. FIELDS	SEARCHED								
Minimum do C23C	Minimum documentation searched (classification system followed by classification symbols)								
Documentat	Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched								
Electronic da	ata base consulted during the international search (name of data bas	e and, where practicable, search terms use	d)						
EPO-In	ternal, WPI Data								
C. DOCUME	ENTS CONSIDERED TO BE RELEVANT								
Category*	Citation of document, with indication, where appropriate, of the rele	vant passages	Relevant to claim No.						
А	EP 1 225 246 A1 (NIPPON STEEL CORP [JP]; 1-21 DAIDO STEEL SHEET CORP [JP]) 24 July 2002 (2002-07-24) cited in the application the whole document								
А	EP 2 644 736 A1 (JFE STEEL CORP 2 October 2013 (2013-10-02) the whole document	1-21							
А	WO 2013/156688 A1 (ARCELORMITTAL INVESTIGACION Y DESARROLLO SL [ES CHRISTIAN [FR) 24 October 2013 (2013-10-24) the whole document	1-21							
Furth	ner documents are listed in the continuation of Box C.	X See patent family annex.							
* Special ca	ategories of cited documents :	"T" loter degument wishlighted -ft the 'steem	notional filing data as suitable						
"A" dogumo	nt defining the general state of the art which is not considered	"T" later document published after the intended date and not in conflict with the application."	ition but cited to understand						
	f particular relevance	the principle or theory underlying the in	nvention						
	pplication or patent but published on or after the international ate	"X" document of particular relevance; the cl							
"L" docume	document which may throw doubts on priority claim(s) or which is step when the document is taken alone								
specia	cited to establish the publication date of another citation or other special reason (as specified) "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is								
"O" document referring to an oral disclosure, use, exhibition or other means combined with one or more other such documents, such combination being obvious to a person skilled in the art									
"P" document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family									
Date of the actual completion of the international search Date of mailing of the international search report									
1	5 December 2015	15/01/2016							
Name and mailing address of the ISA/ Authorized officer Authorized officer									
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016 Brown, Andrew									

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/IB2015/001281

		1 317 1523	710,001201
Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 1225246 A1	L 24-07-2002	AT 508212 T AU 763740 B2 AU 6473000 A CN 1369020 A EP 1225246 A1 EP 2108712 A2 ES 2483969 T3 JP 4136286 B2 JP 2001115247 A US 6635359 B1 WO 0111100 A1	15-05-2011 31-07-2003 05-03-2001 11-09-2002 24-07-2002 14-10-2009 08-08-2014 20-08-2008 24-04-2001 21-10-2003 15-02-2001
EP 2644736 A1	L 02-10-2013	CA 2818296 A1 CN 103228812 A EP 2644736 A1 JP 2012126993 A KR 20130112052 A RU 2013128958 A TW 201231717 A US 2013236739 A1 WO 2012070694 A1	31-05-2012 31-07-2013 02-10-2013 05-07-2012 11-10-2013 10-01-2015 01-08-2012 12-09-2013 31-05-2012
WO 2013156688 A1	L 24-10-2013	AU 2012377741 A1 CA 2870532 A1 CN 104302802 A EA 201401136 A1 EP 2839049 A1 JP 2015520797 A KR 20150008114 A US 2015284861 A1 WO 2013156688 A1	04-12-2014 24-10-2013 21-01-2015 31-03-2015 25-02-2015 23-07-2015 21-01-2015 08-10-2013