

Surface characterization of Fe-Al-Mn alloys for biomaterial applications

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1 **Surface characterization of Fe–10Al–25Mn alloy for biomaterial applications**

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11

12 **Abstract**

13 The austenitic stainless-steel biomaterial, AISI 316L stainless steel, is one of the most
14 widely used for orthopedic or prosthetic implant devices because it is easy to
15 manufacture at a relatively low cost. However, corrosion is a challenging issue related
16 to major alloying compounds with body fluids and wear. Therefore, this study aimed to
17 develop a biomaterial of Fe–Al–Mn alloys by minimizing chromium (Cr) and nickel (Ni)
18 contents. In the research, an attempt has been made to increase the corrosion
19 resistance of Fe–10Al–25Mn alloy using plasma nitriding, which was considered one
20 of the most cost-effective surface treatment. The processes were carried out at
21 various treatment temperatures between 350 and 550 °C, with a pressure of 1.8 mbar
22 in three hours. Several tests were performed, such as chemical compositions,
23 scanning electron microscopy (SEM) combined with energy dispersive spectroscopy
24 (EDS), hardness, and corrosion. The results indicated that the treatment of plasma
25 nitriding at temperatures up to 500 °C significantly enhances the corrosion resistance
26 and increases the hardness of the alloy. At 500 °C, the percentage of nitrogen atoms
27 reaches a peak and then decreases. It means that the nitride formation process on the
28 alloy surface occurs more massively. The amount of nitrogen deposited on the surface
29 of the Fe–10Al–25Mn alloy, as well as a thin layer of iron nitride, is noticeable in the
30 SEM-EDS test. Furthermore, the formation of phases due to the nitriding temperature
31 significantly impacts the alloy properties.

32

33 **Keywords:** Plasma nitriding, Materials characterization, Fe–10Al–25Mn alloy,
34 Biomaterials, Surface Hardness, Corrosion.

35

36 **Highlights**

- 37 – Plasma nitriding improves the corrosion resistance of stainless steel
- 38 – Plasma nitriding enhances the surface hardness of the Fe–10Al–25Mn alloy
- 39 – The Fe–10Al–25Mn nitride alloy has a great potential for medical applications
- 40 – The increase of manganese contents reduces the mechanical resistance
- 41 – Nickel and chromium are two of the potential hazards in medical stainless steels

42

43 **1. Introduction**

44 In the last decades, the application of metallic biomaterials in the medical sector
45 has **been overgrown**, such as in orthopedic implant devices and dentures. With
46 material engineering and the development of the latest technology, millions of people
47 seem to have a new spirit in improving their quality of life. **The metallic biomaterials**
48 **play a crucial role in reconstructing the human body in orthopedic surgery and dentistry**
49 **[1]. Some literature noted that at least three implant materials are broadly used for**
50 **medical purposes, such as stainless steel [2], titanium (Ti) alloys [3], and cobalt-**
51 **chromium (Co-Cr) based alloys [4]. Because of its high strength, dependable wear**
52 **and corrosion resistance [5], and antibacterial properties [6], the Co-Cr alloy-based**
53 **material is considerably used in medical implants and dental prosthetics. The suitability**
54 **of the biological system of the human body is an essential consideration in developing**
55 **biomaterials. Therefore, materials should not cause harmful effects and not damage**
56 **the tissues of the human body. It must be free from toxic products and not trigger**
57 **allergies, inflammation, or cancer [7]. Furthermore, biomaterials require clinical testing**
58 **to ensure their use [8], [9].**

59 Austenitic stainless steel, AISI 316L, is extensively used for orthopedic and
60 prosthetic implant devices. This material has reliable mechanical characteristics at an
61 affordable price. The manufacturing process of this material is also relatively easy
62 compared to others [10]–[12]. **The main advantages of SS-AISI-316L are inexpensive,**
63 **stable mechanical properties and easy manufacturing process. However, problems**
64 **with medical stainless steel have been discovered in recent decades of clinical use.**
65 **Firstly, the stainless steel used in medical applications commonly has a higher elastic**
66 **modulus than bone. This incompatibility of strength or modulus can result in a stress**
67 **shielding effect, which can impede bone healing. Secondly, stainless steel is**
68 **considered less able to withstand wet conditions in a fluid body environment, which**
69 **has a greater potency in corrosion and wear.** This could lead to premature fracture and
70 massive corrosion of the implanted device, followed by releasing harmful substances
71 to the human body. A study reported that both nickel (Ni) and chromium (Cr) are among
72 the potential hazards in medical stainless steels [13]. Allergic contact dermatitis
73 affected by Ni is the most common type of metal hypersensitivity reaction. Furthermore,
74 high Cr levels in the human body may cause cancer and other diseases [1].

75 It is well known that the liquid contains about 0.9% salt in the human body, a pH
76 of 7.4, and a temperature of about 37 °C [14]. Therefore, biomaterials naturally interact
77 continuously with body fluids. This interaction certainly affects the metals and alloys
78 attached to the human body. Naturally, the complexity of the human body with a wet
79 environment triggers corrosion of almost all metal-based biomaterials, which is
80 followed by chemical and electrochemical degradation. Even the most corrosion-
81 resistant materials are difficult to escape from this natural process. Therefore,
82 biomaterials must be biocompatible and non-stimulating by the body systems, non-
83 toxic, and withstand repeated loads in an aggressive body environment [15]. Besides,
84 biomaterials must have physical and mechanical properties that are reliable to replace
85 body systems, be easily formed and produced at a relatively low cost [4].

86 **Recently, high manganese (Mn) austenitic steels with nitrogen alloys have**
87 **improved strength, toughness, corrosion resistance, and nonmagnetic properties,**
88 **making them a viable replacement for traditional Cr-Ni stainless steels in orthopedic**
89 **implants and other medical applications.** The development of manganese-based alloys
90 has been reported for bio-absorbable implants [16]. A metal vapor vacuum is used to
91 implant the biocompatible manganese (Mn) compound into the biomedical Mg surface.

92 This is subjected to evaluate the impact of Mn ion cultivation on the corrosion
93 phenomenon of biomedical Mg. This study found that the surface roughness could be
94 reduced by Mn ion implantation. In a recent study using Fe–Mn–C alloys, the increase
95 of Mn contents reduced the mechanical resistance [17]. Another study found that the
96 addition of Ca to Fe–Mn–Si alloys could improve the osteoinduction and
97 osteoconduction processes better than Fe–Mn–Si alloys or standard AISI 316L
98 stainless steel. The ability to degrade at higher corrosion rates appears to be more
99 optimal [18]. In addition, the suitability of Fe–35Mn–5Si as a biodegradable implant has
100 been improved by considering its mechanical and corrosion properties [19].

101 Metallic biomaterials appear to be required for patients to support diseased
102 tissue for as long as necessary. However, the capability of implants to degrade
103 uniformly under various conditions in the human body to avoid cytotoxic effects and
104 inappropriate tissue responses remains a significant concern. It means that the new
105 alloys being developed must be high strength, wear-resistant, corrosion-resistant,
106 antibacterial and non-toxic. One of the most widely used industrial processes is plasma
107 nitriding, including nitrogen absorption by diffusion into the structure of a material. This
108 approach mainly applies to tools and low alloy steels, considering their most
109 inadequate surface treatments costs. The main benefit of plasma nitriding is to
110 increase the mass transfer of molecules and high-energy nitrogen ions to the surface
111 of the material and to improve the control of process parameters. In this regard, this
112 study focuses on developing a biomaterial by eliminating both Cr and Ni contents. New
113 biomaterials are directed towards stainless steel without a nickel to reduce the toxic
114 properties of AISI 316L stainless steel. The biomaterial compounds made are also
115 expected to be more robust and corrosion-resistant compared to pure metal. In the
116 study, several tests such as microstructure, surface hardness, and corrosion have
117 been realized to evaluate the surface characteristics of Fe–10Al–25Mn alloy after
118 plasma nitriding. The nitride alloys were characterized using a spectrometer, scanning
119 electron microscopy (SEM) combined with energy dispersion spectroscopy (EDS),
120 micro-Vickers hardness and corrosion testing.

121

122 2. Materials and Methods

123 2.1 Materials and samples preparation

124 The Fe–10Al–25Mn alloy smelting process was carried out using a high-
125 frequency induction chamber with a capacity of 50 kg. The raw materials used in this
126 study were mild steel scrap, Fe–Mn med C, pure Al, Fe–C. The ingot-shaped Fe–10Al–
127 25Mn alloy casting has a size of 30 m x 30 m x 200 m. An inductively coupled plasma
128 optical spectrometer was used to examine the chemical compositions of the alloys.
129 Table 1 shows the chemical compositions of the tested specimen.

130

131 **Table 1.** The chemical compositions of Fe–10Al–25Mn alloy.

132

133 2.2 Surface characterization and experiments

134 For nitriding, specimens measuring 5 mm x 10 mm x 10 mm were prepared.
135 The reference material was a 2 mm thick AISI 316L stainless steel plate cut into 10
136 mm x 10 mm squares. Furthermore, the specimen surfaces were smoothed by
137 sandpaper up to 2000 mesh and cleaned by an ultrasonic cleaner using a polishing

138 machine. Plasma nitriding equipment consists of a metal vacuum vessel with an
139 emptiness system, a nitrogen gas input, a 300 1200-volt DC high voltage system, and
140 a temperature regulator. The nitriding process was completed in three hours at 350,
141 400, 450, 500, and 550 °C with a pressure of 1.8 mbar.

142 The tests carried out included the composition, microstructures, hardness, and
143 corrosion. The composition test was carried out using Baird FSQ Foundry Spectrovac
144 Spectrometer based on ASTM E2209 standard test. A JEOL type JSM.6360-LA-EDX
145 (JED 2200 series) Scanning Electron Microscope-Energy Dispersive X-Ray System
146 (SEM-EDS) were used to examine the microstructures. The mechanical testing was
147 performed by Schmierplan/Lubrication plan LA-H-250 RC 16-02/Hardness Tester DIA
148 Testory micro-Vickers method. The Vickers hardness test procedure was based on
149 ASTM E384. Finally, a corrosion polarization test was performed using a CMS 100
150 Gamry Instrument to quantify the corrosion rate. The ASTM G5 standard was used to
151 determine the polarization potential.

152 2.3 Data analyses

153 The results of microstructure tests were qualitatively evaluated based on
154 various temperatures of plasma nitriding. The comparison of shapes, patterns, sizes,
155 and types of microstructures were interpreted in the analyses. Quantitatively, graphs
156 were created to present the effects of hardness and corrosion. The current research
157 data were also analyzed considering the viewpoint of the findings data by other
158 researchers for more advanced analyses.

159

160 3. Results and Discussion

161 The Fe–10Al–25Mn alloy has been experimentally investigated to evaluate its
162 potency for biomaterials. The plasma nitriding was realized to give treatments in the
163 surface material. The microstructure change of the surface alloy plays an important
164 role to impact its hardness and corrosion rate. The main results and the advanced
165 discussion are presented as follow.

166 3.1 Microstructures

167 Figure 1 gives the SEM micrograph and EDS spot analysis of Fe–10Al–25Mn
168 alloy before plasma nitriding. It is found in Figure 1(a) that the Fe–10Al–25Mn as-cast
169 alloy has austenite, ferrite, and kappa structure. The structure of austenite tends to be
170 dominant because of the element Mn as an austenite stabilizer. The ferrite structure is
171 related to the Al element as a ferrite stabilizer, while the kappa phase is associated
172 with the relatively high C content, as found by another researcher [20]. As addressed
173 by Chen et al. [21], Mn is dissolved in the Fe system as a solid solution with a
174 disordered FCC structure. The presence of the Al atom in the system changes the
175 disordered FCC structure to an ordered FCC, and the C atom causes the formation of
176 the κ (Fe, Mn)₃AlC phase [22]. The κ phase is seen around the α/γ duplex system (see
177 Figure 2).

178 Based on the magnification of the SEM micrograph in Figure 2, both the austenite
179 structure and the κ form lamellas. It is similar to the findings of another research [21].
180 Thus, the aluminum content of 7.5% is a ferrite phase stabilizer, and 20% manganese
181 is an austenite stabilizer, and a high enough C content encourages the kappa phase
182 formation.

183

184

Fig. 1. The microstructure of Fe–10Al–25Mn alloy.

185

186

Austenite remains stable at low Al and high C compositions, while κ -carbide remains stable at high C and high Al compositions. Thus, in austenite, κ -carbide precipitation is part of the dispersion of both C and Al. It is in line with the research findings by Kim et al. [23]. Based on the measurement of EDS composition, the Fe–10Al–25Mn alloy contains no nitrogen, as shown in Figure 1(b).

191

192

Fig. 2. The SEM micrograph Fe–10Al–25Mn alloy high magnification.

193

194

Figure 3(a) depicts the SEM test results on the nitride cross-section of the Fe–10Al–25Mn by plasma nitriding process. As previously found by Manfredini et al. [24], the nitride layer consists of γ -Fe(N), Fe₄N and AlN compounds. Fe₄N tends to be dark, whereas AlN is bright. The dominant austenite phase in the Fe–10Al–25Mn alloy encourages the formation of the AlN nitride. This result is similar to that of a study carried out by Chen [25]. The nitride layer on the transverse surface of the Fe–10Al–25Mn alloy produced by plasma nitriding at a temperature of 350 °C is unclear. The higher the nitriding temperature, the thicker the nitride layer. This finding denotes that the higher the plasma nitriding temperature, the more nitrogen diffuses the Fe–10Al–25Mn alloy surface, forming a nitride compound.

204

Figure 3(b) provides the EDS test results on the surface of the Fe–10Al–25Mn alloy after the nitriding process. There is a thin layer of iron nitride on the Fe–10Al–25Mn alloy surface and a percentage of nitrogen deposited. The nitrogen content on the surface of the Fe–10Al–25Mn alloy is sensitive to the plasma nitriding temperature. The distance between the atoms of the Fe–10Al–25Mn alloy specimen becomes increasingly tenuous with the rise of the plasma nitriding temperature. It is due to nitrogen atoms diffuse more easily into the Fe crystal system. The increased nitriding temperature also causes the atoms to vibrate in a position of instability. This causes it more straightforward for nitrogen atoms to enter and diffuse between the atoms making up the Fe–10Al–25Mn alloy. The nitrogen atom then binds with Fe to form the intermetallic compound Fe₃N and Fe₄N, as Chen found [1]. When the nitrogen atom meets Al, it creates the intermetallic AlN compound.

216

217

Fig. 3. The microstructure of Fe–10Al–25Mn nitride alloy.

218

219

Figure 4 displays N content on the surface of the Fe–10Al–25Mn alloy after nitriding treatment at various temperatures between 350 and 550 °C. The percentage of nitrogen atoms increases from 350 to 500 °C and declines after 550 °C. This decrease is related to the nitriding temperature, proportional to the depth of nitrogen atoms in the specimen. The distance between the particles in the sample stretches as the nitriding temperature increases at 500 °C. Thus, it is easier for nitrogen atoms to diffuse onto the surface of the specimen to form a layer of iron and aluminum nitride. The distance between the atoms would be even greater if the nitriding temperature is increased to 550 °C. The percentage of nitrogen atoms on the specimen surface decreases as the specimen surface diffuses deeper below the cut surface. The

228

229 nitriding process at 350 to 550 °C yields in α' -Fe(N) with a percentage of nitrogen
230 atoms up to 19 %, whereas at 19-21 %, nitrogen levels cause the formation of Fe₄N
231 iron nitride phase. The amount of nitrogen atoms deposited on the specimen surface
232 significantly impacts the percentage of the Fe₄N phase formed in its region. This
233 finding agrees well with another research [24].

234

235 **Fig. 4.** The N content on the surface of the Fe-10Al-25Mn Mn alloy.

236

237 3.2 Surface Hardness

238 Figure 5 indicates the surface hardness test results of the Fe-10Al-25Mn alloy
239 after plasma nitriding. The surface hardness of the Fe-10Al-25Mn alloy after plasma
240 nitriding at 350 °C is 445.6 VHN. The higher the plasma nitriding temperature, the more
241 hardness increases until it reaches a maximum of 680.3 VHN at 500 °C. The hardness
242 decreases up to 30% after attaining a peak point at a temperature of 550 °C. It is relevant
243 to the EDS test results, where the percentage of nitrogen atoms reaches a maximum at
244 500 °C and then decreases significantly. The nitride phase formed on the surface also
245 has a powerful effect on the surface hardness of the Fe-10Al-25Mn alloy after plasma
246 nitriding. Treatment temperature up to 450 °C causes the material surface of the Fe-
247 10Al-25Mn alloy nitride phase to be α' -Fe(N). The surface changes to Fe₄N when the
248 temperature is 500 °C. Unsurprisingly, Meka et al. found something similar from the
249 results of their study [26].

250

251 **Fig. 5.** The surface hardness of Fe-10Al-25Mn alloy at various nitriding
252 temperatures.

253

254 Figure 6 shows the hardness distribution test results of the Fe-10Al-25Mn alloy
255 after the plasma nitriding process. At all plasma nitriding temperatures, the deeper the
256 hardness decreases. At a distance of 10 μ m from the surface, the decrease in hardness
257 is not significant, only around 2.5 %. This finding reveals that nitrogen atoms diffuse
258 quickly up to a distance of 50 μ m and form nitride compounds. However, there is a
259 significant decrease in hardness at a distance of 100 and 150 μ m. It means that the
260 nitrogen atom experiences a substantial energy reduction. It can be related to the
261 collisions with particles on the surface to reduce its penetration depth. At a distance of
262 250 μ m, the hardness is equivalent to a Fe-10Al-25Mn alloy that does not undergo
263 nitriding. Nitrogen atoms show no more diffusion in the penetration of N in Fe-N up to
264 70 μ m, as found by other studies [24][26].

265

266 **Fig. 6.** The cross-surface hardness distribution of Fe-10Al-25Mn alloy after plasma
267 nitriding.

268

269 3.3 Corrosion rate

270 Figure 7 represents the corrosion rate of the Fe-10Al-25Mn alloy. The calculation
271 of the corrosion rates adopts the formula used by Li et al. [27]. The corrosion resistance
272 increases and reaches the lowest value at 500 °C when plasma nitriding temperature is

273 higher. It means that the corrosion resistance of Fe–10Al–25Mn alloy increases and gets
274 a maximum value at 500 °C. The corrosion resistance of this alloy is due to the formation
275 of nitride on its surface after plasma nitriding, which becomes more apparent, more
276 massive, and thicker as the temperature of the nitride layer rises. Fe₄N, Fe₂N, and AlN
277 compounds make up this nitride layer. Thus, the nitride compound on the alloy surface
278 increases the superficial properties, especially hardness and corrosion resistance.
279 According to Chen et al. [21], corrosion in this nitride layer appears to cross-grain
280 boundaries and take the form of pitting. Another study found that alloying Fe with Mg
281 reduces its corrosion resistance [28]. Mg corrosion is affected by the production of
282 hydroxide (OH⁻) and an increase in pH. On the other side, the increase in Al content
283 leads to higher noble corrosion.

284

285 **Fig. 7.** The corrosion rate of Fe–10Al–25Mn alloy at various nitriding temperatures.

286

287 **4. Conclusion**

288 Investigation of Fe–10Al–25Mn alloy has been carried out experimentally. It can
289 be summarized that plasma nitriding can increase the corrosion resistance of Fe–
290 10Al–25Mn alloy. Higher plasma nitriding temperature increases corrosion resistance
291 and reaches a maximum of 500 °C with γ-Fe(N), Fe₄N and AlN structures on the
292 surface. Plasma nitriding also increases the surface hardness of the alloy. Plasma
293 nitriding at a temperature of 500 °C produces the highest hardness. It means that the
294 newly developed material will be a reliable replacement for traditional medical
295 stainless steel. This material combines the advantages of a stable austenitic structure
296 and corrosion resistance. Therefore, the Fe–10Al–25Mn nitride alloy can be developed
297 as a prospective biomaterial.

298

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303

304 **Conflicts of interest**

305 The authors declare no conflicts of interest.

306

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