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Posted Date: 6 May 2023

doi: 10.20944/preprints202305.0410.v1

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Article

Influence of Ion Nitriding on Microstructure and Properties of Haynes 282 Nickel Alloy Specimens Produced Using Additive Technique

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Abstract: The paper investigates the influence of ion nitriding on the microstructure, corrosion resistance, and tensile strength at elevated temperatures of Haynes 282 nickel superalloy specimens produced by the Direct Metal Laser Sintering-DMLS technique. Analysis of the specimens' surface morphology revealed that the ion nitriding process significantly affects the physical and chemical phenomena occurring on the surface of the specimens. Phase composition and stress analysis showed that Haynes 282 nickel alloy specimens produced by the DMLS technique and a chromium nitride layer produced in the ion nitriding process have tensile stress. The layer is also characterized by high nanohardness compared to specimens produced using the classical method (wrought) and DMLS technique. The conducted corrosion resistance tests confirmed the positive effect of the nitrided layer on the corrosion resistance of Haynes 282 nickel alloy produced by the DMLS technique. The ion nitriding process did not significantly affect the elevated-temperature tensile strength of Haynes 282 nickel alloy specimens produced by the DMLS technique.

Keywords: Haynes 282 nickel alloy; additive manufacturing; ion nitriding; corrosion resistance; microstructure; static tensile test

1. Introduction

Nickel-based superalloys are widely used in various industries, e.g. energy, aerospace, oil and gas processing [1-5]. Their versatility stems from both their mechanical properties and their resistance to high-temperature oxidation and electrochemical corrosion in many aqueous environments [6-8]. The passive oxide layer which spontaneously forms on the surface, as in the case of titanium alloys or alloy steels, significantly increases corrosion resistance compared to the substrate in the active state [9]. However, an oxide cathode layer makes the passive oxide layer susceptible to local corrosion in the presence of aggressive ions, such as chloride ions (Cl⁻), often found in many industrial environments [10]. In the case of nitrided layers produced on Ni-Cr alloy [11] exposed in chloride-free environments (H₂SO₄), the CrN+Cr₂N layer is not susceptible to pitting corrosion. In addition, it has been shown that the best corrosion resistance is achieved by when phases are combined (CrN + Cr₂N) rather than when a homogeneous phase Cr₂N or phase π (Cr_{12.8}Ni_{7.2}N_{4.0}) are used alone. It is assumed that the differentiated corrosion resistance of chromium nitrides is related to the different crystallographic structures of the CrN (fcc A1) and Cr₂N (hcp A3) phases and the packing density of atoms in the unit cell. In the case of corrosion solutions with a mixture of aggressive ions (Cl⁻ and

SO₄²⁻) [12], the durability of nitrided layers is additionally determined by the quality and degree of the substrate's defectiveness, which is determined, among others, by the value of breakdown potential (Enp) and the intensity of local corrosion (e.g. pitting). In nitride layers formed on alloys rich in chromium atoms, it is also possible to have insignificant amounts of other phases (α -Cr, Cr₂O₃) that further reduce the corrosion resistance of the substrate. One of the newer nickel superalloys developed for use at high temperatures in aggressive environments is Haynes 282 nickel alloy [13,14]. This alloy is strengthened with the γ' phase and is characterized by high strength properties and good weldability [15]. Good weldability is the result of a carefully defined aluminum and titanium content, which affects the volume fraction of the γ' phase. Its high content significantly limits the technological properties of the produced materials, especially those obtained using additive techniques [16]. However, it should be emphasized that materials produced by additive techniques usually demonstrate anisotropic properties. They are mainly caused by crystallographic texture, the structure of smelting pools, the privileged distribution of certain phases and carbides, and the presence of asymmetrical defects [17]. Boswell et al. in their study [18] showed that the anisotropic effect could be effectively reduced by appropriate dissolution and ageing heat treatment, leading to recrystallization and local texture reduction. Applying high temperature to the building platform during the LPBD process has also been shown to be effective in reducing adverse anisotropy effects, such as grain boundary cracking, associated with accumulating γ' and carbide precipitates [19].

Ion nitriding processes belong to modern thermo-chemical treatments enabling the production of protective diffusion layers with controlled phase composition and high-performance properties. Such materials include chromium nitride, characterized by high hardness, corrosion resistance and resistance to abrasive wear or oxidation. This material in diffusion layers or coatings is successfully produced on some steels [20-22]. However, there is much less information on the ion nitriding of nickel alloys, in particular, nickel alloys having the characteristics of a strongly defective anisotropic structure.

The study aims to investigate the influence of the anisotropic structure of Haynes 282 nickel alloy substrate on the microstructure of a chromium nitride layer formed by ion nitriding and its resistance to corrosion, hardness, Young's modulus and mechanical properties at elevated temperatures.

2. Materials and Methods

2.1. Production of Haynes 282 Specimens by Additive Technique

Test specimens measuring 20x20x10 mm were produced from Haynes 282 (HY 282) nickel alloy powder using an EOS M100 printer working in Direct Metal Laser Sintering (DMLS) technology. The specimens were produced in a 5.0 purity argon atmosphere, with the platform heated to 80 °C. Each subsequent layer was scanned in a direction rotated by 67° relative to the previous layer. The specimen production process was carried out with the following parameters: energy density E-100 J/mm³, laser power P – 100 W, laser speed V – 800 mm/s, and distance between successive paths H – 0.05 mm.

2.2. Ion Nitriding Process

Before the nitriding process, the specimens were washed in an ultrasonic scrubber in ethyl alcohol, then dried and placed in a universal ion nitriding furnace on the cathode potential of PDT (Plasma/Ion Diffusion Treatment) processes. The ion nitriding process on the cathode potential was carried out in a mixture of gases (25% N₂ + 75% H₂) for 12 h at 570 °C and 3.5 hPa.

2.3. Microstructure

Microstructures were studied by means of light microscopy, scanning electron microscopy (SEM) as well as transmission electron microscopy (TEM). To reveal the general microstructure, particularly the melt pools, metallographic deposits of Haynes 282 produced by the additive technique was prepared. For comparison, specimen of Haynes 282 nickel alloy produced using the

classical technology (cast and annealed/wrought) was also prepared. The specimens produced using the optimal process parameter were etched using Kalling's No. 2 reagent. The revealed microstructure was examined and revealed microstructures were examined using a Zeiss Axio Scope light microscope. The microstructure of nitriding Haynes 282 produced by the additive technique was analyzed by light and transmission electron microscopy. The cross-section metallographic deposit was prepared. TEM studies were performed using a FEI Tecnai G2 F20 S-TWIN operated at 200 kV. Cross-sectional TEM specimen was prepared by mechanical polishing and subsequent Ar⁺ ion milling until electron transparency using the PIPS system from Gatan. The surface morphology of nickel superalloy specimens as-prepared and after corrosion resistance tests were also studied by means of the TM-1000 scanning electron microscope using backscattered electrons (BSE).

2.4. Nanohardness of the Manufactured Layers

Nanoindentation tests were conducted on Micromaterials' Vantage Alpha using a Berkovich indenter. The determined load curves as a function of the indenter displacement were analyzed according to the Oliver-Pharr method to determine nanohardness, and included thermal drift analysis. The measurements were made along the cross-section of the produced layers with 1 mN load, 10 s loading time, 10 s unloading time and 5 s dwell period.

2.5. Corrosion Resistance Tests

Corrosion resistance tests of Haynes 282 nickel alloy (produced using classical technology and using 3D printing) (in the initial state and with a nitrided layer produced in the ion nitriding process), were performed using impedance (EIS) and potentiodynamic (LSV) methods by means of the AutoLab PGSTAT100 potentiostat in a non-deoxygenated solution of 0.1 M Na₂SO₄ + 0.1 M NaCl at room temperature. Before electrochemical testing, the specimens were exposed to a corrosion solution under electroless conditions for 5000s needed to stabilize the open system potential. Impedance tests (EIS) were carried out in a three-electrode system (test electrode – reference electrode (saturated calomel electrode SCE) – auxiliary electrode (platinum)) in the frequency range of 105 Hz – 10⁻³Hz, with a sinusoidal signal amplitude of 10 mV in potentiostatic mode at the open circuit potential (EOCP). Electrochemical studies were carried out using a Faraday cage. Impedance spectra were analyzed using Baukamp's EQUIVCRT program. For the classic nickel superalloy, an electrical replacement circuit (EC) with one time constant R(RQ) was used, while the 3D printed material employed a replacement circuit with two-time constants R(RQ)(RQ). Two-time constants in the starting material are conditional on a heterogeneous substrate structure, typical for materials after laser processing, in which the paths of penetration of the substrate with a laser beam and familiar 'tears' are observed. The production of a nitrided layer on a substrate using the 3D printing technique significantly modified the parameters and electrochemical nature of the substrate as a result of numerous nitride precipitates on the surface and some layer discontinuities as observed in microscopy studies. They were analyzed using replacement system R(Q[R(RQ)]), which is commonly used for materials susceptible to local corrosion. The resulting spectra are presented in the form of Bode and Nyquist graphs. Potentiodynamic tests were carried out in an identical three-electrode system up to a potential of 1500 mV. After anodic polarization, a return polarization (inverted curve) was performed to determine the presence of a possible hysteresis loop. Regardless of the polarization direction, the potential change was 0.2 mV/s. Surface topography studies were carried out using the ACCURION optical profilometer HALCYONICS_i4 Sensofar Metrology (SPAIN) using the SensoVIEW program.

2.6. Analysis of Phase Composition and Residual Stresses

The phase composition analysis was carried out using Rigaku's Miniflex II X-ray diffractometer (JPN). Filtered X-rays of the CuK_{α1} λ = 0.154 nm tube in the Bragg-Brentano diffraction geometry were used. An angle value range of 2θ = 20 ÷ 100°, a radiation intensity measurement period of 2θ = 0.02° and a measurement time of t = 3s were assumed. The phase composition was determined using the

diffraction database PDF (Powder Diffraction File), developed by ICDD (The International Centre for Diffraction Data). The identification of phase components consisted of adjusting the profile of the obtained diffractogram, i.e. the calculated distances between hkl planes (d_{hkl}) for individual reflections and their intensity, to the data in the PDF database.

The self-stresses were determined for the marked points on the surface of the test sleeve. Proto and XRD Combo X-ray diffractometer and Proto Manufacturing (CAN) XRD Win 2.0 computer software were used. The test used a lamp with a manganese anode $MnK\alpha_1$ $\lambda = 0.210$ nm, a collimator diameter of 2 mm, an anode current of four mA, and an anode voltage of 20 kV. To calculate the self-stress values at a given measuring point, the $\sin^2\Psi$ method [23] was used. This is a standard method involving the use of symmetrical Bragg-Brentano diffraction. This method uses a goniometer of the Ψ type, which makes it possible to obtain the proper inclinations of the diffraction vector by angles Ψ and in a plane perpendicular to the diffraction plane [24]. Natural stresses were determined for constant values of the angle Ψ in the range from 25 to -25° . Elastic deformations in the tested element were carried out for the diffraction line from the family of {311} planes at an angle of $2\theta = 155.2$. In the self-stress measurements, a Poisson's coefficient value of $\nu = 0.319$ and a Young's modulus value of $E = 217$ GPa were assumed [25].

2.7. Static Tensile Test

The static tensile test was carried out at 750 °C on axially symmetrical specimens with a circular cross-section and a diameter of 6 mm. The length of the parallel part of the specimen was 36 mm, and the measuring base of the extensometer was 25 mm. The tensile test was performed using an electromechanical testing machine equipped with a force sensor at a measuring range of 50 kN. The control variable of the tensile test was the constant displacement of the testing machine traverse of 0.5 mm/min, which for a given specimen geometry gives an initial tensile speed of 3.3×10^{-4} s $^{-1}$. For each of the materials, 2 specimens were subjected to a tensile test, and then the average values of the analyzed mechanical properties were determined.

3. Research Results and Discussion

3.1. Microstructure

Figure 1 shows the microstructure of Haynes 282 nickel alloy specimens in a cast and annealed state, and produced by DMLS in the XY plane. In the microstructure of the Haynes 282 nickel alloy in the cast and annealed state (Fig. 1a), austenite grains, twins and carbides are visible. The microstructure of the Haynes 282 nickel alloy specimen produced by the additive technique comprises layers/welds arranged alternately corresponding to the laser beam's scanning directions and the melted powder's particle size (Figure 1b).

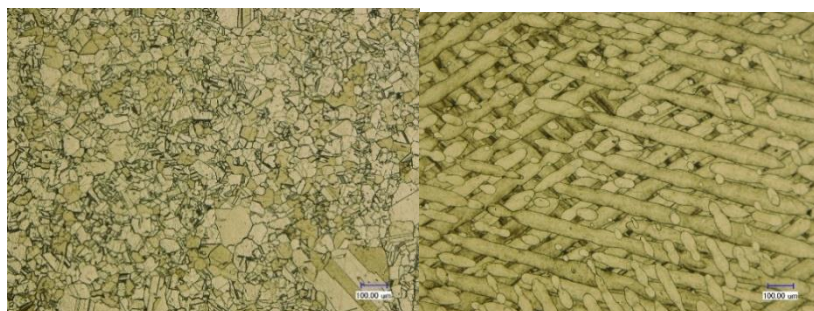


Figure 1. Microstructure of Haynes 282 nickel alloy specimen: wrought –(a), in-built (XY) plane using DMLS technique.

Figure 2 shows TEM micrographs of the Haynes 282 nickel alloy substrate prepared by the additive technique. One can see the high density of dislocations that form the regular cellular structure. The austenite grains consist of slightly misoriented sub-grains about the size of

500 – 1000 nm. The cells are of elongated shape (Fig. 2a). The direction of elongation probably can be attributed to the laser beam's scanning direction. In edge-on view, the cells adopt a hexagonal shape (Fig. 2b). Such kind of dislocation structure was also observed in an additively manufactured austenitic steel [33].

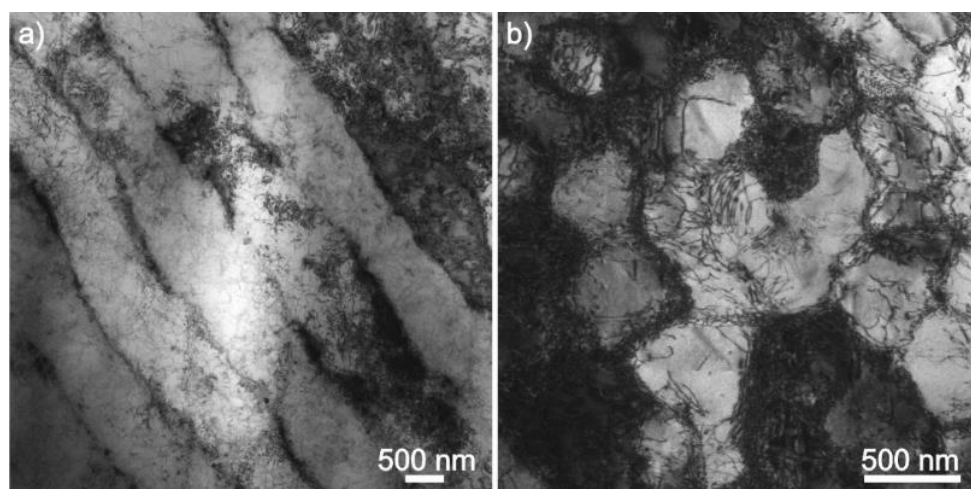


Figure 2. TEM images of Haynes 282 substrate produced by the DMLS technique.

Figure 3 shows the cross-section and morphology of a layer produced in the ion nitriding process at a temperature of 570 °C in 12 h on a Haynes 282 nickel alloy substrate. Microscopic observations show that the thickness of the nitrided layer can be estimated at approx. 7 μm (Fig. 3a). This layer is homogeneous and continuous over the entire specimen's cross-section, adheres well to the substrate, and there are no visible voids at the boundary of the substrate layer. Fine particles are visible on the surface of the layers (Fig. 3b), adhering well to the specimen's surface.

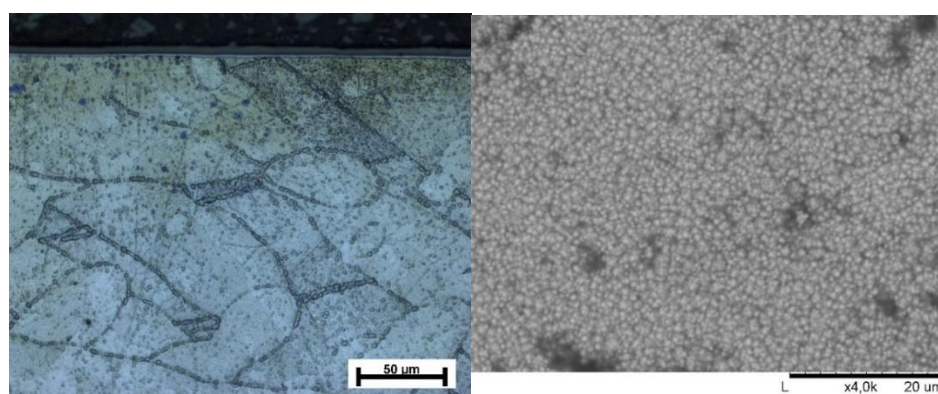


Figure 3. Cross-section-a), and morphology (SEM)-b) of nitride layer on Haynes 282 nickel alloy produced by ion nitriding.

3.2. Phase Analysis and Residual Stress

Analysis of the phase composition of Haynes 282 nickel alloy as delivered and specimens produced by 3D printing showed very similar lines on diffractograms and the occurrence of two phases: gamma matrix $\text{Ni}\gamma$ (PDF Card 01-077-9326) and the strengthening phase γ' (PDF Card 04-004-2742) (Fig. 4).

The ion nitriding process changed the phase composition on the surface of Haynes 282 nickel alloy specimens. Two additional phases were found: CrN (PDF Card 04-007-0676) and Cr_2N (PDF Card 00-035-0803). The CrN phase crystallizes in the orthorhombic structure (Pnmm). The lattice parameters of the unit cell are $a_0 = 0.287$, $b_0 = 0.297$ and $c_0 = 0.4132$ nm. In turn, the Cr_2N phase

crystallizes in the hexagonal structure (P31m), with the unit cell lattice parameters equal to $a_0 = 0.481\text{nm}$ and $c_0 = 0.448\text{ nm}$.

Diffractogram patterns (Fig. 4) obtained from the XRD analysis, containing both phases from the Haynes 282 alloy substrate ($\text{Ni}\gamma$ and γ') and the phases of CrN and Cr_2N after the ion nitriding process, are similar to those analyzed in studies [26-27].

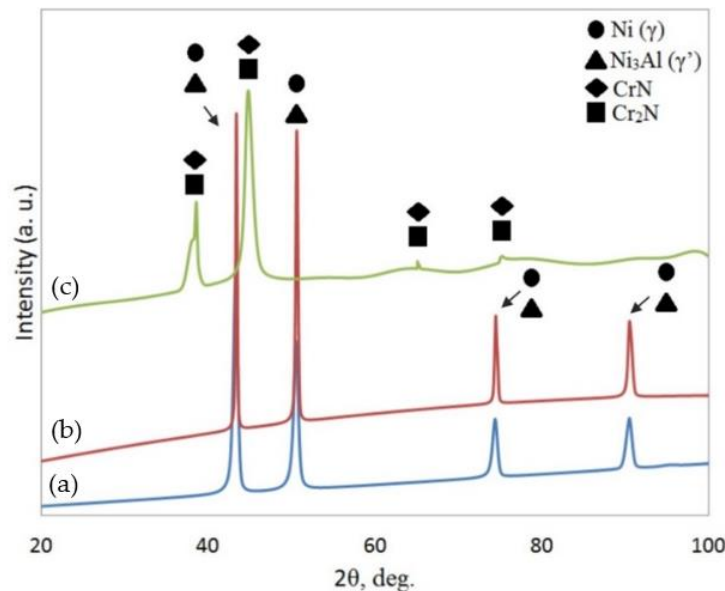


Figure 4. Diffractogram pattern for specimens: Haynes 282 alloy wrought –(a), produced DMLS –(b), and DMLS with nitride layer –(c).

TEM studies of the nitriding alloy (Fig. 5) revealed a thin layer at the surface. The layer has irregular thickness of about 200 – 400 nm and consists of nano-sized grains (several nanometers) in an amorphous matrix with the presence of larger grains (tens to hundreds of nanometers) and voids (Fig. 5a-b). Below the layer, a Moiré pattern is visible (Fig. 5c) that indicates nanograins precipitated in the austenite matrix of the alloy. Interestingly, this region does not exhibit a high density of dislocations. It is extending to a depth of about 8 μm and then the cellular dislocation structure, like for the alloy substrate (Fig. 2), was revealed. It corresponds to the layer thickness determined by light microscopy observations. Electron diffraction analysis (Fig. 5d) indicates the formation of a CrN phase, but the presence of a Cr_2N phase cannot be excluded due to similar interplanar distances.

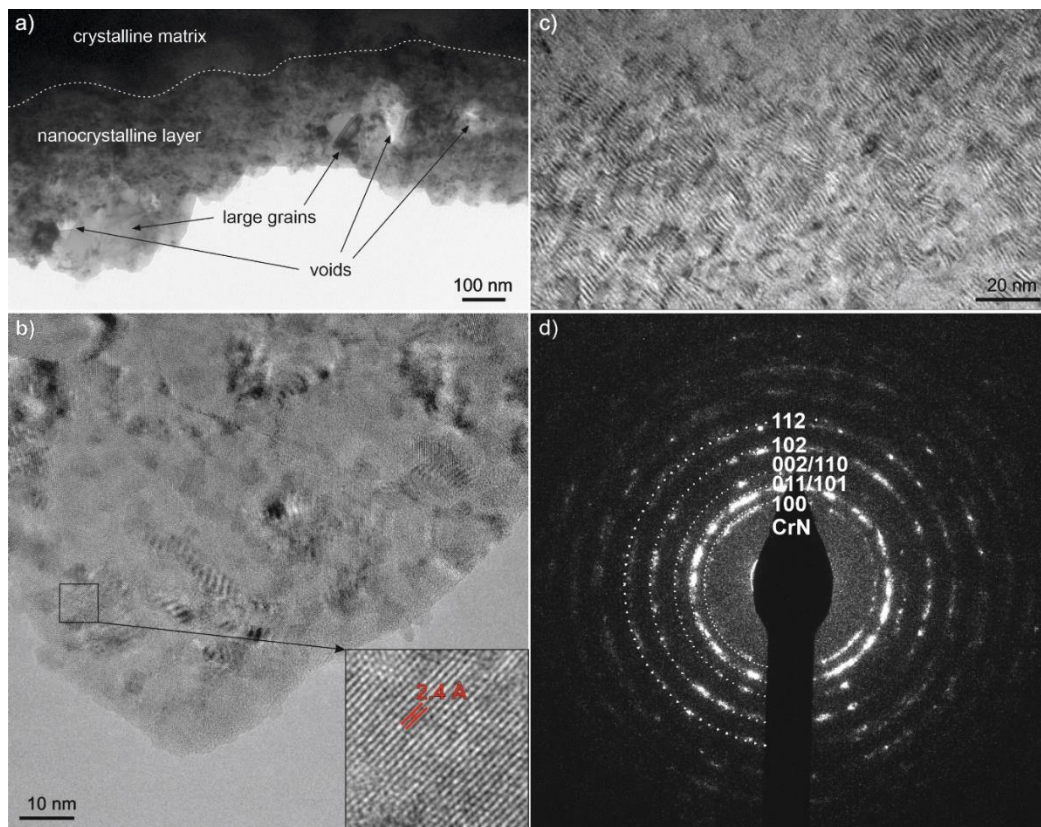


Figure 5. TEM images of the nitriding Haynes 282 produced by the additive technique. a)-b) The nanocrystalline layer is present at the surface. c) The Moiré pattern is visible in the matrix below the layer. d) Electron diffraction pattern obtained in b) region.

Residual stress values were measured in the initial state, after 3D printing and after ion nitriding. In each case, a tensile stress state was found. The highest tensile stress value of $+384 \pm 40$ MPa was found in the alloy in its initial state, while the 3D printed specimen showed a slightly lower value of $+152 \pm 24$ MPa. The lowest tensile stress value was found in the Haynes 282 alloy after ion nitriding and amounted to $+64 \pm 45$ MPa.

In the analyzed studies, the stresses found in the surface layer after fluorescent nitriding are either mostly compressive [28] or on the border of tensile and compressive [29]. However, B. Bouaouina et al. also obtained eigen-tensile stresses [30].

3.3. Nanohardness

Nanohardness distribution analysis along the cross-section in the near-surface zone showed a strengthening of the material after ion nitriding to a depth of about $6 \mu\text{m}$ (Fig. 6), which corresponds to the layer thickness determined based on microscopic observations (Fig. 3). The hardness of the core is 13.5 ± 1.9 GPa. As it approaches the surface, it smoothly increases to a maximum value of about 27 GPa at a depth of about $1.4 \mu\text{m}$, followed by a slight decrease associated with the morphology of the surface zone of the nitrided layer.

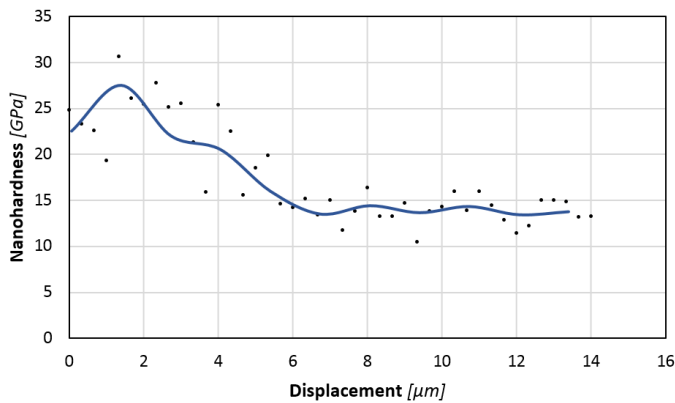


Figure 6. Nanohardness distribution in the cross-section of the layer.

3.4. Corrosion Resistance

Figure 7 shows the impedance spectra obtained for specimens of Haynes 282 nickel alloy wrought, produced by DMLS, and produced by DMLS and coated with a CrN+Cr₂N layer. The analysis of the spectra and data presented in Table 2 confirmed the positive effect of the nitrided layer on the corrosion resistance of Haynes 282 nickel alloy.

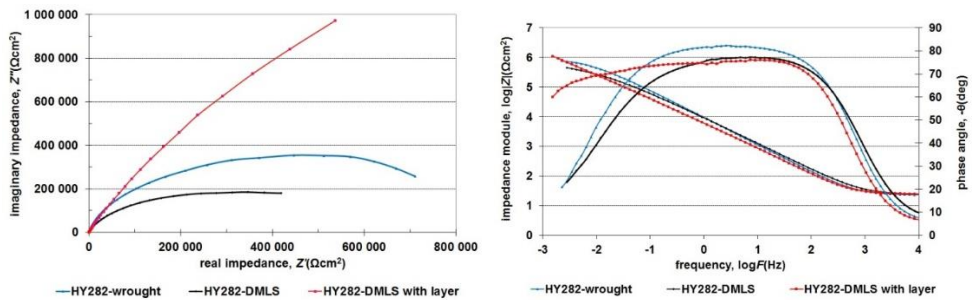


Figure 7. Impedance spectra obtained for Haynes 282 alloy specimens wrought, produced by DMLS, and by DMLS containing a nitride layer.

Table 2. Characteristic electrochemical values of tested specimens (impedance tests).

	Substitute arrangement		Dielectric layer	double layer
Haynes 282-wrought	R(RQ)	$R(\Omega\text{cm}^2)$		8.07×10^5
		$Y_0\text{CPE}(F\text{cm}^{-2}\text{s}^{n-1})$		1.99×10^{-5}
		n		0.91
Haynes 282-DMLS	R(RQ)(RQ)	$R(\Omega\text{cm}^2)$	6.26×10^4	4.36×10^5
		$Y_0\text{CPE}(F\text{cm}^{-2}\text{s}^{n-1})$	4.99×10^{-5}	3.87×10^{-5}
		n	0.86	0.88
Haynes 282 DMLS with layer	R(Q[R(RQ)])	$R(\Omega\text{cm}^2)$	8.61×10^3	7.43×10^6
		$Y_0\text{CPE}(F\text{cm}^{-2}\text{s}^{n-1})$	2.67×10^{-5}	1.03×10^{-5}
		n	0.89	0.66

R – resistance, Y0 CPE – capacity of constant phase element, n – coefficient of imperfections of constant phase element (CPE); an empirical constant ranging from 0 to 1. It is worth noting that when n = 1, the CPE behaves as a pure capacitor, while when n = 0, the CPE behaves as a pure resistor.

Increasing the value of charge transfer resistance through the double layer (resistance R_t) from $4.4 \times 10^5 \Omega\text{cm}^2$ (Haynes 282-DMLS) to $7.4 \times 10^6 \Omega\text{cm}^2$ for Haynes 282 alloy with CrN+Cr₂N layer led to significant increases in the corrosion resistance. However, the presence of a dielectric layer with

reduced resistance ($8.6 \times 10^3 \Omega\text{cm}^2$), despite the relatively high value of the parameter $n = 0.89$ (indicating its capacitive character), may imply local discontinuities facilitating degradation and initiation of pitting corrosion observed in potentiodynamic studies. It can be assumed that the structure of the specimens produced using the DMLS technique, in which few pores are observed, promotes the initiation of pitting corrosion in chloride solutions.

The data analysis in Tables 2 and 4 indicates a significant development of the nitrided layer surface, thus increasing the active surface several times. This is indicated by the values of the parameter 'n' of the double layer ($n=0.66$) and the parameter S_a ($S_a=118\text{nm}$) of the roughness profile. Both of these value indicate a significant share of diffusion factors determining the material's corrosion resistance.

The potentiodynamic tests were correlated in Fig. 8 and summarized in Table 3. It confirmed the positive effect of the ion nitriding process on the corrosion resistance of the Haynes 282 alloy specimens produced using the DMLS technique. This is reflected in the decrease of the density values of corrosion currents from $4.09 \times 10^{-2} \mu\text{A}/\text{cm}^2$ (Haynes-DMLS) to $8.89 \times 10^{-3} \mu\text{A}/\text{cm}^2$ and an increase in the value of corrosion potentials. A leading shift in the value of corrosion potentials ($\Delta E=200 \text{ mV}$) towards anodic indicates an increase in the nobility of the substrate and the presence of a cathodic surface layer. Its presence increases the corrosion resistance of Haynes 282 alloy but also sensitizes the specimens to pitting corrosion in chloride environments. In the case of a chromium nitride layer produced on a substrate produced using the DMLS technique, the observed constant increase in current density that accompanies the polarization intensity increase (up to approx. 400 mV) is typical for nitride layers produced on metallic substrates. This result is associated with the presence of nitrogen in the top layer. The presence of nitrogen and its susceptibility to oxidation facilitates the oxidation of the alloy during anodic polarization. Taguchi and Kurihara's research has additionally shown that chromium loses its self-passivation capacity during nitriding [31]. An increase in the intensity of polarization results in the initiation of pitting corrosion (E , e.g. = 430 mV, E_{np} – puncture potential initiating pitting corrosion) and flaking of the nitrided layer (Fig. 9A). A further increase in the intentionality of polarization (surf. 1200 mV) allows the start of another electrochemical process associated with releasing oxygen from the solution and the decomposition of water. The reverse polarization process showed the existence of a large hysteresis loop and the absence of regasification potential. This result indicates that the damaged nitrided layer does not undergo spontaneous repassivation and is completely degraded by its exfoliation. This result shows that the possible application of nitrided layers can be proposed for conditions that do not damage mechanical nitrided layers or for chloride-free environments.

In the case of starting materials (Haynes 282 alloy wrought and produced using DMLS), regardless of the structure, this alloy remains passive in a wide range of tested potentials. The data analysis in Table 2 indicates a slight decrease (by approximately 5%) in the corrosion resistance of the specimens produced by the DMLS technique compared to the wrought Haynes 282 alloy. In both the analyzed cases, the durability of passive layers and the current densities in the passive area have been compared. The observed increase in current density at approx. 500 mV is caused, by the hydration of nickel oxide (at approx. 500 mV) and change in the oxidation stage of chromium (at approx. 650 mV), i.e. the main alloying components of Haynes 282 nickel alloy. In the case of nickel, hydration of nickel oxide to hydroxide (from NiO to $\text{Ni}(\text{OH})_2$) occurs at a potential of about 500 mV [32] while a further increase in current density is caused by corrosion of passive chromium oxide Cr_2O_3 to the soluble form of CrO_4^{2-} . The formation of a thick layer of corrosion products and the achievement of a diffusion limit current by the system stabilizes the intensity of electrochemical processes. The return polarization shows the absence of a typical hysteresis loop, which shows that the observed increase in current density above 500 mV is associated with the trans-passive region, in which the intensity of processes is related to oxidation of the substrate and release of oxygen from the solution. The tight formed layer of oxide corrosion products is shown in Fig. 9B. The created scale slows down the availability of electrolytes to the substrate and the resulting cathode reactions determine the intensity of corrosion processes. The results of the EIS test of Haynes 282 alloy with a formed corrosion layer

confirm its capacitive character ($n=0.93$) with a high resistance ($6.34 \times 10^6 \Omega\text{cm}^2$) typical for oxide barrier layers.

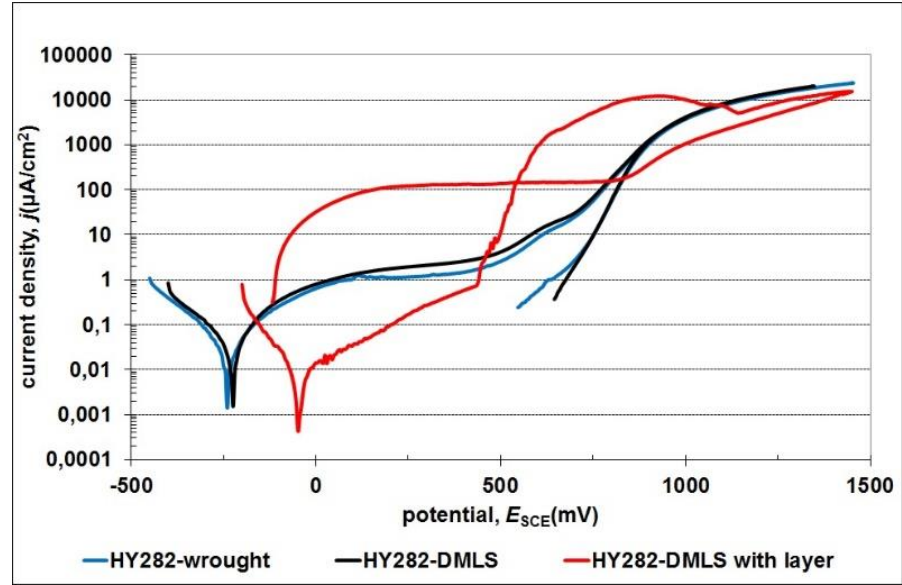


Figure 8. Potentiodynamic curves obtained for the specimens: Haynes 282 alloy wrought, produced by DMLS, and by DMLS with a nitride layer.

Table 3. Characteristic electrochemical values of the tested specimens.

	Haynes 282 wrought	Haynes 282 DMLS	Haynes 282 DMLS with layer
R_p (kΩcm²)	809	637	2763
I_{corr} (μA/cm²)	3.88×10^{-2}	4.09×10^{-2}	0.89×10^{-2}
E_{corr} (mV)	-240	-225	-45

R_p – polarization resistance; I_{corr} . – corrosion current density; E_{corr} . – corrosion potential.

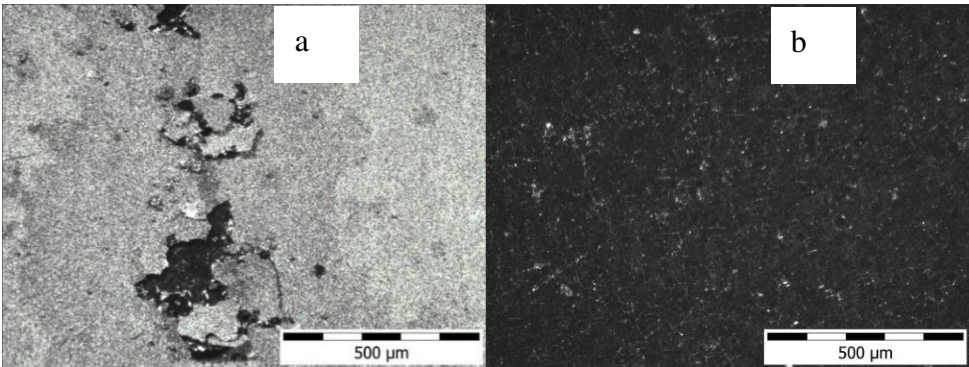


Figure 9. Surface topography of Haynes 282 alloy obtained by DMLS and coated with a chromium nitride layer (after potentiodynamic tests) - a), and oxide barrier layer -b).

Table 4. Surface roughness parameters of specimens [nm].

	S_a	SD	R_a	SD	R_z	SD
Haynes 282 wrought	12.78 ± 0.15		7.35 ± 0.62		76.14 ± 13.74	
Haynes 282 DMLS	16.45 ± 0.24		6.39 ± 0.40		71.82 ± 10.63	
Haynes 282 DMLS with CrN+Cr ₂ N layer	118.93 ± 1.37		84.25 ± 5.26		1186.65 ± 263.02	

S_a - average arithmetic deviation of the roughness surface from the median line; R_a - average arithmetic deviation of the roughness profile from the median line along the elementary length (2D tests); R_z – distance from the

highest point of the roughness profile to its lowest point measured along the elementary length (2D tests); SD - standard deviation.

3.7. Static Tensile Test

Based on the strength tests carried out in the form of a static tensile test (Fig. 10), it can be concluded that specimens produced by the DMLS technique have comparable yield strength (YS_{0.2}) as well as strength (UTS) with respect specimens covered with a chromium nitride layer. Simultaneously, the specimens produced by DMLS and specimens with a chromium nitride layer showed significantly higher values than those in the wrought state (Table 5). At a temperature 750 °C, the specimens produced using DMLS show a decrease in strength with small deformations after reaching (YS_{0.2}).

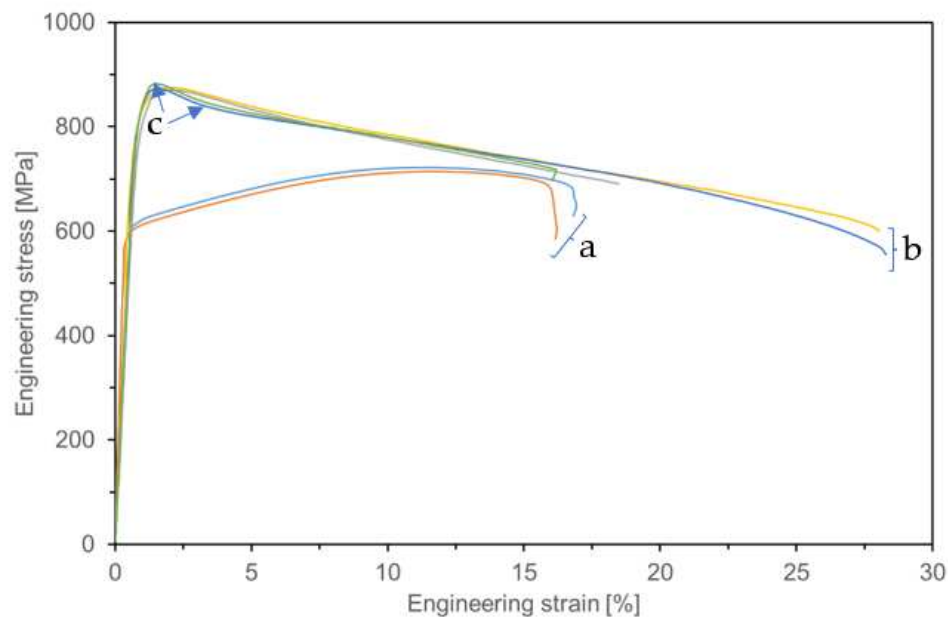


Figure 10. Examples of stress-strain curves for specimens: Haynes 282 nickel alloy (wrought) (a), HY282 DMLS (b) and HY282 DMLS with a CrN+Cr₂N layer (c).

Table 5. Average mechanical properties of tested specimens.

Specimens	YS _{0.2} [MPa]	UTS [MPa]	A [%]
Haynes 282 alloy (wrought)	600±4	719±4	15±1.5
Haynes 282 DMLS	808±3	873±4	22±6.4
Haynes 282 DMLS with a CrN+Cr ₂ N layer	834±5	878±6	21±8.5

4. Conclusions

Based on the research carried out, the following conclusions can be drawn:

1. Use of the DMLS technique on Haynes 282 nickel alloy powder makes it possible to produce specimens characterized by low porosity and a fine-crystalline anisotropic structure.
2. Ion nitriding conducted at 570 °C leads to the formation of a 7µm-thick continuous layer containing CrN+Cr₂N phases on a substrate of Haynes 282 nickel alloy.
3. Tensile stress is found to occur in specimens produced by the DMLS technique and in specimens containing the CrN+Cr₂N layer produced via ion nitriding.
4. The produced CrN+Cr₂N layer demonstrates high corrosion resistance and comparable tensile strength to the substrate material (produced DMLS technique).

Author Contributions: Conceptualization, R.S. J.K., and J.K.; methodology, R.S. K.P.; formal analysis, A.K.; investigation, R.S., J.K., K.G., K.P., J.M., and K.K.; resources, K.K.; data curation, K.K., K.P. and R.S.; writing—original draft preparation, R.S. and J.K.; writing—review and editing, J.K., A.K., J.M., and K.G.; visualization, A.K., K.K., J.M., and K.G.; supervision, R.S. and J.K.; project administration, R.S.; funding acquisition, R.S.

Acknowledgments: The research was funded by POB Technologie Materiałowe of Warsaw University of Technology in the framework of the Excellence Initiative: Research University (IDUB) programme.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: Data are available with the first author and can be shared with anyone upon reasonable request.

Conflicts of Interest: The authors declare no conflicts of interest.

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