Influence of Niobium or Molybdenum Addition on Microstructure and Tensile Properties of Nickel-Chromium Alloys

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Abstract: This work discusses on influence of niobium or molybdenum addition on microstructure and tensile properties of NiCr-based dental alloys. In this regard, the Ni-24Cr-8Nb, Ni-22Cr-10Nb and Ni-20Cr-12Nb (wt. %) alloys produced by arc melting process. To compare the typical Ni-22Cr-10Mo dental alloy was also produced. These ternary alloys were analyzed by chemical analyses, X-ray diffraction (XRD), scanning electron microscopy (SEM), electron dispersive spectrometry (EDS), thermogravimetric analysis (TG), Vickers hardness and tensile tests. Although the mass losses of the samples during arc melting, the optical emission spectrometry showed that the initial compositions were kept. The Ni-22Cr-10Mo alloy produced a matrix of Ni_{ss} (ss—solid solution), whereas Ni_{3}Nb disperse in a Ni_{ss} matrix was also identified in Ni-Cr-Nb alloys. Excepting for the Ni-22Cr-10Nb alloy with mass gain of 0.23%, the as-cast Ni-Cr alloys presented mass gains close to 0.4% after heating up to 1000 °C under synthetic airflow. The hardness values, the modulus of elasticity, yield strength and ultimate tensile strength have enhanced, whereas the ductility was reduced with increasing niobium addition of up to 12 wt.-%. The Ni-22Cr-10Mo alloy presented an intergranular fracture mechanism containing deep dimples and quasi-cleavage planes, whereas the shallow dimples were identified on fracture surface of the as-cast Nb-richer Ni-Cr alloys due to the presence of higher Ni_{3}Nb amounts.

Keywords: microstructure; tensile properties; intermetallics; casting; phase transformation

1. Introduction

As-cast nickel-chromium (Ni-Cr) and nickel-chromium-molybdenum (Ni-Cr-Mo) based alloys are applied in dentistry to partial removable prosthesis since 1930 because of their excellent characteristics, such as corrosion resistance and mechanical properties, compared with the more expensive gold-based alloys. Thus, the less expensive and lighter Ni-Cr alloys with high elastic module (170–190 GPa) have been developed for producing metal-ceramic restorations, intra-radicular pins, crowns and removable partial prosthesis [1–3]. In these alloys, the chromium is responsible for the formation of an outer protective oxide layer [4]. Further, the control on the quantities released from dental materials has also been considered for these alloy designs.
Typical chemical compositions of different commercial Ni-Cr alloys used for metal-ceramic restorations presented the Cr and Mo contents varying between 18.5–20.4 wt. % and 4.2–9.5 wt. %, respectively [5]. Other alloys and traces such as cobalt, iron, zinc and manganese containing up to 2.5, 2.5, 0.65 and 0.1 wt. %, respectively, can be found in these alloys. Studies with Be-free Ni-Cr alloys have also been considered for dental applications [6]. In accordance with the phase diagrams of the nickel-chromium [7] and nickel-molybdenum [8] systems, the adopted chromium and molybdenum amounts in these commercial Ni alloys are located in regions of Ni solid solutions, respectively. For comparison, the commercial Ni-20Cr-5Mo (wt. %) alloy with a microstructure based on Ni<sub>33</sub> (ss—solid solution) was adopted, and its mechanical properties are based mainly on the strengthening solid solution and grain refining mechanisms. However, the use of metal matrix-intermetallic composites can also improve the need flexural strength for dentistry devices [9].

It is well known on the use of niobium as alloying in the new Ti alloys for orthopedic and dentistry devices, since their superior biocompatibility and osseointegration features compared to commercial Ti-6Al-4V alloy [10–12]. In accordance with the phase diagram of the nickel-niobium system [13], the nickel can dissolve up to 4 wt. % (11 at. %) Nb whereas the eutectic reaction \( L \leftrightarrow Ni + Ni_3Nb \) at 1216 °C can happen during solidification of these richer-Ni alloys. The liquidus projection of the Ni-Cr-Nb system indicates the presence of a ternary eutectic reaction \( [L \leftrightarrow NbCr_2\text{ (high-temperature)} + Nb_6Ni_3 + Nb_3Ni_7] \) at 1170 °C in the rich-Ni region [14].

In this sense, this work reports on the influence of niobium or molybdenum addition in microstructure and tensile properties of nickel-chromium alloys.

2. Materials and Methods

In this work, high-purity starting powders of Ni, Cr and Nb were used to prepare the Ni-22Cr-10Mo, Ni-24Cr-8Nb, Ni-22Cr-10Nb and Ni-20Cr-12Nb alloys (wt. %) by arc melting process. For comparison, the typical Ni-20Cr-4Mo dental alloy (wt. %) was also evaluated. In this way, the elemental Ni-Cr-Nb and Ni-Cr-Mo powder mixtures were mixed and then compacted square-section blocks with a length of 55 mm and a 5 mm thickness were obtained by axial compaction using 7 ton (approximately 25 MPa). The Ni-Cr-Mo and Ni-Cr-Nb ingots weighing close to 50 g were prepared in an arc melting furnace under argon atmosphere using a water-cooled copper hearth (own USP project, Lorena, Brazil), non-consumable tungsten electrode, and gettered by titanium. Five melting steps were carried out for each alloy in order to obtain homogeneous ingots. The mass losses occurred during the melting process were lower than 0.6%, and the nominal compositions were then adopted.

The as-cast Ni-24Cr-8Nb, Ni-22Cr-10Nb, Ni-20Cr-12Nb and Ni-22Cr-10Mo (wt. %) alloys were characterized by chemical analysis, scanning electron microscopy (SEM, Carl Zeiss Microscopy GmbH, Jena, Germany), X-ray diffraction (XRD, Shimadzu Corporation, Kyoto, Japan), electron dispersive spectrometry (EDS, INCA Energy Oxford, Oxfordshire, UK), Vickers hardness and tensile tests (Buehler Global Head Quarters, Lake Bluff, IL, USA) at room temperature.

Chemical analysis of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys was conducted in an ARL 3460 Optical Emission Spectrometer (OES, Thermo Scientific, Waltham, MA, USA). The total C, S, N and O contents of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys were measured by combustion analyses in CS-125 and TC-436 LECO analyzers (Leco Corporation, St. Joseph, MI, USA). Conventional metallographic techniques, including the use of silica colloidal suspension for polishing and chemical attack with aqueous acid solution (20 vol. % HF:HNO<sub>3</sub> (2:1)), were adopted in this work.

XRD experiments at room temperature of the monolithic Ni-Cr-Nb and Ni-Cr-Mo alloys were performed in XRD-6000 Shimadzu equipment (Shimadzu Corporation, Kyoto, Japan) using Ni-filtered Cu-Kα radiation, voltage of 40 kV, current of 30 mA, angular range (2θ) from 10° to 80°, angular pass of 0.05° and counting time per pass of 1 s. Based on the JCPDS database [15] and Powdercell computer program [16], the phases formed in microstructure of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys were indexed.
Electron images of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys were obtained in a Zeiss (LEO) 1450-VP SEM (Carl Zeiss Microscopy GmbH, Jena, Germany) using the secondary electron and back-scattered electron (BSE) detectors. EDS analysis was adopted to measure the amounts of Ni, Cr, N and Mo presents in phases of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys. At least three measurements per phase were obtained for each sample. The amounts of precipitates in as-cast Ni-Cr-Nb alloys were determined using the ImageJ computer program [17] from their SEM images.

To determine the oxidation resistance of the Ni-Cr alloys evaluated in this work, the thermogravimetric analysis of these as-cast samples was conducted after heating up to 1000 °C in alumina crucible under synthetic air flow with heating rate of 10 °C per minute.

Vickers hardness tests of as-cast Ni-Cr-Nb and Ni-Cr-Mo samples were performed in a Wolpert durometer using 9.81 N load and loading time of 15 s, according to ASTM E-92 standard [18]. At least eight measurements were obtained for each specimen.

Tensile tests of as-cast Ni-Cr-Nb and Ni-Cr-Mo samples were conducted according to the ASTM E-8M [19] standard in an INSTRON model 8801 servo hydraulic testing machine (Instron Company, Norwood, MA, USA) with load cell of 100 kN capacity. Cylinder specimens with 20 mm gage length and diameter of 4 mm were carefully machined. At least three specimens per alloy were tested. Details on their fractured surfaces were obtained by SEM analysis.

3. Results

Table 1 shows the Ni, Cr, Nb and Mo contents (wt. %) of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys measured by ICP-AES analysis.

Table 1. Elemental contents (wt. %) measured by an ICP-AES analysis of the Ni-Cr-Nb and Ni-Cr-Mo alloys evaluated in this work.

| Element | Ni-20Cr-10Mo | Ni-24Cr-8Nb | Ni-22Cr-10Nb | Ni-20Cr-12Nb | Commercial Alloy |
|---------|--------------|-------------|--------------|--------------|-----------------|
| C       | 0.014        | 0.032       | 0.036        | 0.027        | ND              |
| S       | 0.011        | 0.012       | 0.015        | 0.012        | ND              |
| N       | 0.016        | 0.019       | 0.021        | 0.021        | ND              |
| Al      | 0.002        | 0.000       | 0.001        | 0.001        | ND              |
| Si      | 0.170        | 0.220       | 0.210        | 0.170        | ND              |
| P       | 0.004        | 0.003       | 0.003        | 0.003        | ND              |
| Ti      | 0.011        | 0.113       | 0.136        | 0.162        | ND              |
| V       | 0.003        | 0.006       | 0.006        | 0.007        | ND              |
| Cr      | 21.92        | 24.29       | 22.04        | 20.08        | 19.1            |
| Mn      | 0.010        | 0.030       | 0.010        | 0.010        | 0.65            |
| Ni      | 69.00        | 66.30       | 66.20        | 65.30        | 70.0            |
| Cu      | 0.003        | 0.018       | 0.018        | 0.023        | ND              |
| Nb      | 0.010        | 8.090       | 9.840        | 11.52        | ND              |
| Mo      | 9.110        | 0.045       | 0.011        | 0.006        | 4.2             |

ND—not determined.

Table 2 shows the C, S, O and N contents (wt. %) of the as-cast NiCr alloys evaluated in this work, which were determined utilizing either the gas fusion method (nitrogen and oxygen) or the combustion method (carbon and sulfur).
Table 2. Interstitial contents (wt. %) measured by LECO combustion analyses of the Ni-Cr-Nb and Ni-Cr-Mo alloys.

| Alloy                  | C (wt. %) | S (wt. %) | N (wt. %) | O (wt. %) |
|------------------------|-----------|-----------|-----------|-----------|
| Ni-22Cr-10Mo           | 0.020     | 0.020     | 0.016     | 0.099     |
| Ni-20Cr-12Nb           | 0.010     | 0.007     | 0.022     | 0.039     |
| Ni-22Cr-10Nb           | 0.010     | 0.010     | 0.025     | 0.059     |
| Ni-24Cr-8Nb            | 0.020     | 0.099     | 0.019     | 0.054     |
| Commercial Ni-Cr Alloy | 0.260     | 0.002     | 0.032     | 0.027     |

XRD patterns of the Ni-22Cr-10Mo and Ni-20Cr-12Nb alloys as well as the commercial Ni-Cr alloy are presented in Figure 1.

![XRD patterns](image)

**Figure 1.** XRD patterns of Ni-Cr alloys evaluated in this work: (a) Commercial Ni-Cr alloy, (b) Ni-22Cr-10Mo alloy and (c) Ni-20Cr-12Nb alloy.

Figure 2 displays the SEM images of the Ni-22Cr-10Mo alloy as well as the commercial Ni-Cr alloy evaluated in this work.

![SEM images](image)

**Figure 2.** SEM images of the (a) commercial Ni-Cr alloy and (b) as-cast Ni-22Cr-10Mo alloy without a chemical attack.

SEM images of as-cast Ni-Cr-Nb alloys evaluated in this work are presented in Figure 3.
Figure 3. SEM images of the as-cast (a,d) Ni-24Cr-8Nb, (b,e) Ni-22Cr-10Nb and (c,f) Ni-20Cr-12Nb alloys: (a,c) Without chemical attack and (b,d,f) after chemical attack.

The contents (at. %) of Ni, Cr, Mo, Nb and other elements measured by EDS analysis of phases formed in the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys evaluated in this work are presented in Table 3.
Representative thermogravimetric curves of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys evaluated in this work are illustrated in Figure 4.

Table 3. The elemental contents (at. %) measured by electron dispersive spectrometry (EDS) analysis of phases formed in the as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys.

| Alloy          | Phase | Ni  | Cr  | Mo  | Nb  | Others                  |
|----------------|-------|-----|-----|-----|-----|-------------------------|
| Ni-22Cr-10Mo   | Ni$_{ss}$ | 68.5–70.5 | 24.0–25.5 | 5.1–6.0 | - | -                       |
| Ni-24Cr-8Nb    | Ni$_{ss}$ | 68.9–69.4 | 25.4–27.8 | - | 2.8–5.5 | -                       |
| Ni-20Cr-12Nb   | Ni$_{ss}$ | 70.2–71.1 | 22.5–25.0 | - | 3.9–7.1 | -                       |
| Commercial Ni-Cr Alloy | Ni$_{ss}$ | 72.3 | 9.6–10.5 | - | 17.2–18.1 | -                       |
| Ni-24Cr-8Nb    | Ni$_{ss}$ | 59.5–70.7 | 22.0–26.7 | 2.0–4.5 | - | Si (3.6–8.6); Mn (0.6–0.7) |

The elemental contents measured by electron dispersive spectrometry (EDS) analysis of phases formed in the as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys are indicated in Table 4.

Table 4. Vickers hardness of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys investigated in this work.

| Alloy          | Vickers Hardness (HV) |
|----------------|-----------------------|
| Ni-22Cr-10Mo   | 150.0 ± 5.3           |
| Ni-24Cr-8Nb    | 243.0 ± 19.8          |
| Ni-22Cr-10Nb   | 309.0 ± 6.7           |
| Ni-20Cr-12Nb   | 374.0 ± 21.1          |
| Commercial Ni-Cr Alloy | 231.0 ± 14.6 |

The Vickers hardness results of as-cast Ni-Cr-Nb and Ni-Cr-Mo alloys investigated in this work are indicated in Table 4.

The representative tensile σ–ε curves of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys are illustrated in Figure 5 whereas their compression mechanical properties are presented in Table 5.
Tensile mechanical properties of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys evaluated in this work.

Table 5. Tensile mechanical properties of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys evaluated in this work.

| Alloy            | Yield Strength (MPa) | Ultimate Tensile Strength (MPa) | Total Strain (%) | Elastic Module (GPa) |
|------------------|----------------------|---------------------------------|------------------|----------------------|
| Ni-22Cr-10Mo     | 212.4 ± 46.2         | 380.6 ± 36.5                    | 4.2 ± 0.2        | 28.9 ± 16.4          |
| Ni-24Cr-8Nb      | 334.2 ± 22.3         | 505.3 ± 22.2                    | 6.5 ± 0.5        | 41.4 ± 0.6           |
| Ni-22Cr-10Nb     | 448.4 ± 48.4         | 551.3 ± 51.7                    | 4.5 ± 0.7        | 61.0 ± 5.8           |
| Ni-20Cr-12Nb     | 510.5 ± 81.5         | 576.5 ± 133.4                   | 1.8 ± 0.5        | 69.5 ± 18.3          |

Details on the fracture surface of the Ni-22Cr-10Mo and Ni-20Cr-12Nb alloys after tensile tests are illustrated in Figure 6a,b.

Figure 5. Representative tensile $\sigma$-$\epsilon$ curves of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys investigated in this work.

Figure 6. SEM fractographies of the as-cast (a,b) Ni-22Cr-10Mo and (c,d) Ni-20Cr-12Nb alloys after tensile tests.
4. Discussion

Chemical analysis by ICP method of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys has indicated that the Cr, Nb and Mo amounts were kept close to their nominal compositions (see Table 1). The commercial Ni-Cr alloy presented the Ni, Cr and Mo contents of 70, 19.1 and 4.2 wt. %, respectively. Trace of manganese (0.65 wt. %) was also detected in this commercial alloy.

Similar values of C, S and N were measured from the combustion gases within the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys (see Table 2). Despite the route adopted in this work the amount of total oxygen of up to 0.099 wt. % were picked up during arc melting from the compacted elemental powder mixtures only. On contrary, this route has produced alloys with lower C and N interstitial amounts than the commercial alloy. Nitrogen contents higher than 0.1 wt. % can contribute to a reduction on the ductility of Ni-based alloys [20]. Similar results were found for a silver-palladium alloy [21]. The commercial alloy had also amounts of Co, Fe, Mn and Zn of 0.25, 0.68, 0.65 and 0.1 wt. %, respectively.

Both the commercial alloy (with 19 wt. % Cr and 4 wt. % Mo) and the experimental Ni-22Cr-10Mo alloy presented similar XRD results, as are shown in Figure 1. In these alloys, the intense Ni peaks can be observed only, which could be related for X-ray diffraction experiments with monolithic samples. However, these Ni peaks have shifted toward the direction of smaller diffraction angles for the Mo-richer alloy, suggesting that the larger molybdenum amount in Ni-22Cr-10Mo alloy is preferentially dissolved into the Ni lattice. Traces of others peaks besides the intense Ni peaks were also identified in XRD patterns of the Ni-Cr-Nb alloys evaluated in this work. Nonetheless, it is very important to note that there are two overlapping peaks between the major diffraction lines of Ni, Ni$_8$Nb, Ni$_3$Nb and Ni$_6$Nb$_7$ [22].

The commercial Ni-Cr alloy evaluated in this work presents primary Ni$_{ss}$ grains and a eutectic region located in their grain boundary as is indicated in the SEM micrograph of Figure 2a. It can be also observed the presence of regions with different contrasts into the Ni$_{ss}$ grains, which had the contents of Ni, Cr, Mo, Mn and Si measured by EDS analysis varying between 59.5–70.0, 26.7–22.0, 4.5–2.0, 0.7–0.6 and 8.6–3.8 at. %, respectively (see Table 3). In this sense, the brighter regions presented the higher Nb contents. Excepting to the manganese content with 0.9 at. %, similar elemental values were found in eutectic regions.

Agreeing with the phase diagram of the Ni-Cr-Mo system [23], the microstructure of the as-cast Ni-22Cr-10Mo alloy presented in Figure 2b has revealed the presence of a homogeneous structure formed by coarse Ni$_{ss}$ (ss—solid solution) grains, and the different contrasts observed are relative to their different crystallographic orientations only. No segregation tendency was detected by EDS analysis in these regions in which the Cr and Mo contents varied between 24.0–25.5 and 5.1–6.0 at. %, respectively (see Table 3). In this sense, the brighter regions presented the higher Nb contents. Some diffusion pores can be also noted in microstructure of this ternary alloy.

The microstructures of the as-cast Ni-Cr-Nb alloys presented in Figure 3 have indicated the same solidification path. Agreeing with the phase diagram of the Ni-Cr-Nb system [14], the solidification is started with the formation of primary Ni$_{ss}$ grains followed by simultaneous precipitation of Ni$_{ss}$ and Ni$_3$Nb for Ni-rich alloys. Comparing with the Ni-24Cr-8Nb alloy, it can be noted the presence of larger Ni$_3$Nb amounts in microstructure of the Ni-20Cr-12Nb alloy. Coherently with the liquidus projection of this ternary system, this alloy containing more eutectic regions is located closer to the L + Ni + Ni$_3$Nb monovariant line [14]. For the as-cast Ni-24Cr-8Nb alloy, the Ni$_{ss}$ grains containing different contrasts have presented 25.4–27.8 at. % Cr and 2.8–5.5 at. % Nb, whereas the Nb-rich precipitates containing up to 15 at. % Nb were also measured by EDS analysis from very fine regions. In addition, it was observed that the brighter Ni$_3$Nb precipitates were darker after chemical attack with aqueous acid solution. EDS analysis also indicated that the Nb contents dissolved into the Ni$_{ss}$ lattice were increased in the as-cast Ni-20Cr-12Nb alloy, which have varied from 3.9 at. % to 7.1 at. % Nb. In this ternary alloy, the coarser Nb-rich precipitates presented 17.2–18.2 at. % Nb, indicating to be the Ni$_3$Nb phase with higher enthalpy of formation (31 kJ·mol$^{-1}$) instead to the Ni$_8$Nb phase (with 15 kJ·mol$^{-1}$) containing contents near the 11.1 at. % Nb [21]. In this way, these EDS results have confirmed that the traces identified in XRD patterns of as-cast Ni-20Cr-12Nb alloy could be associated with the Ni$_3$Nb
According with the ImageJ computer program [17], the SEM images of the as-cast Ni-24Cr-8Nb, Ni-22Cr-10Nb and Ni-20Cr-12Nb alloys have indicated the presence of precipitates for 6.3%, 6.9% and 14.4% in area, respectively.

Remelting operations and thermal cycles are commons for Ni-Cr alloys used in dental restorations and high-temperature applications. Similar behavior was noted during heating up to 1000 °C of Ni-based ternary alloys investigated in this work (see Figure 4). Conversely, the commercial Ni-Cr alloy presented a mass gain of 0.3% after heating up to 400 °C followed by a continuous mass loss under heating in synthetic air between 400 °C and 1000 °C. These facts can be associated with initial oxidation and subsequent reactive evaporation of chromium oxide at high temperatures in the presence of oxygen and/or water vapor, respectively [24]. Thermogravimetric curves of the as-cast Ni-22Cr-10Mo, Ni-24Cr-8Nb, Ni-22Cr-10Nb and Ni-20Cr-12Nb alloys are illustrated in Figure 4, which have indicated mass gains of 0.4, 0.43, 0.23 and 0.49%, respectively. Excepting to the Ni-22Cr-10Nb alloy, similar oxidation resistance was noted for experimental Ni-Cr alloys evaluated in this work.

The Ni-22Cr-10Mo presented lower hardness values than the commercial alloy containing lesser Mo contents, which could be related with its lower interstitial amounts (see Table 4). Similar hardness values were noted between the Ni-24Cr-8Nb alloy and commercial Ni-Cr alloy. In contrast, the as-cast Ni-Cr alloys with 10 wt. % and 12 wt. % Nb presented higher hardness values than the commercial Ni-Cr alloy probably due to the higher Ni$_3$Nb amounts in microstructure of these as-cast ternary alloys.

Depending on the composition and processing route, the Ni-Cr alloys can present the yield strength and ultimate tensile strength of 260–830 MPa and 440–1200 MPa, respectively [25–28]. The commercial Ni-Cr alloy evaluated in this work presented the yield strength (offset 0.2%) and ultimate tensile strength of 417 MPa and 550 MPa, respectively. Table 5 displays the tensile mechanical properties at room temperature of the as-cast Ni-Cr-Mo and Ni-Cr-Nb alloys investigated in this work. Coherently to the hardness results, the Ni-22Cr-10Mo alloy exhibited the lowest values of yield stress (offset 0.2%) and ultimate tensile strength besides the elastic module of 29 GPa close to that related to bone tissues [25–28]. Similarly, the as-cast Ni-24Cr-8Nb alloy containing small amounts of Ni$_3$Nb also exhibited lower yield strength and ultimate tensile strength. Nevertheless, these tensile properties of as-cast Ni-20Cr-12Nb alloy were slightly increased and kept close to those of the commercial Ni-Cr alloy. Consequently, the total strain of the Ni-24Cr-8Nb alloy was reduced from 6.5% to 1.8% in samples of as-cast Ni-20Cr-12Nb alloy whereas their elastic module was increased between 41 GPa and 70 GPa, respectively. It is well known that the Ni$_3$Nb platelets has a detrimental effect on ductility and other mechanical properties that are related to precipitate within the inter-dendritic regions of the as-cast microstructures [29]. In this regard, the development of multicomponent alloys containing the Mo and Nb additions could produce both the high ductility and mechanical strength for structural applications.

The fracture surface of the as-cast Ni-22Cr-10Mo alloy presented an intergranular fracture mechanism and two different regions constituted with variable dimple sizes, which were associated mainly to crystallography orientation of Ni$_{52}$ grains formed during solidification. Moreover, it was also noted the existence of a small amount of quasi-cleavage planes. Shallow dimples were identified on the fracture surfaces of the as-cast Ni-Cr-Nb alloys, which were less deep with increasing Ni$_3$Nb amount as shown in Figure 3c,d, denoting that the Ni$_3$Nb precipitates favored the occurrence of brittle fracture.

5. Conclusions

Arc melting successfully produced the low-interstitial Ni-Cr-Nb and Ni-Cr-Mo alloys from elemental powder mixtures.

The values of Vickers hardness, elastic module, yield strength and ultimate tensile strength were enhanced from 243 HV, 41 GPa, 334 MPa and 505 MPa to 374 HV, 70 GPa, 510 MPa and 576 MPa, respectively, with increasing Nb addition up to 12% in the alloy composition owing the Ni$_3$Nb formation during solidification of Ni-Cr-Nb alloys.
The Ni-22Cr-10Mo alloy with deep dimples and quasi-cleavage planes on its fracture surface has exhibited intergranular fracture mechanism whereas the shallow dimples were identified on fracture surface of the as-cast Nb-richer Ni-Cr alloys due to the presence of higher Ni$_3$Nb amounts.

Author Contributions: M.A.d.S. produced the Ni-Nb alloys and participated in all stages; B.d.O.F. performed the XRD experiments; T.M.H. participated in all stages; A.P. conducted the thermal analysis; C.A.N. obtained SEM images and EDS analysis; C.A.R.P.B. conducted the compression mechanical tests; A.S.R. conceived, participated and coordinate the present work in all stages as well as wrote the paper.

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