Electrochemical Fabrication of Niobium Silicon Alloys from Oxide Powder Mixtures

Fanke Meng and Huimin Lu
School of Materials Science and Engineering, Beihang University, No.37 Xueyuan Road, Haidian District, Beijing 100191, China

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1. Introduction

Nickel alloys are the mostly used superalloys in aviation industry. Due to the demand of exploring new generation aircraft turbine engines, scientists and engineers are devoted to investigate less expensive superalloys with higher melting points to substitute the Ni-based alloys [1, 2]. Following this idea, NbSi alloys (0.5–37.5 at % Si) are attracting more and more attention because of the high melting points of the two composed phases (melting points: Nb, 2469°C; Nb₅Si₃, 2520°C) in the alloy system [1]. Moreover, NbSi alloys still have three other excellent properties: close coefficient of heat expansion of the two compositions (Nb, 7.3 × 10⁻⁶ K⁻¹; Nb₅Si₃, 6.1 × 10⁻⁶ K⁻¹), relative lower density (7.16 g cm⁻³), and reasonable strength at high temperature. Thus, NbSi alloys are considered to be a potential candidate of the next generation superalloys. Until now, the mainly used techniques for fabricating NbSi alloys are orientation solidification [3], smelting with argon arc [4], physical vapor deposition (PVD), as well as some other methods of powder metallurgy. Generally, all of these techniques have the following procedures: first, extracting pure Nb and Si from minerals; second, melting and mixing molten Nb and Si by using arc or electron beam melting method; third, casting the mixed molten alloys to ingots. However, the notorious weakness of the process is resulted from high energy consumption and complex production procedures. As a typical example, the only commercial process for the production of metallic Nb is reducing K₂NbF₇ or Nb₂O₅ with sodium reduction or carbothermal reduction, respectively [5]. For production of silicon, the conventional technique is reducing SiO₂ with carbothermal reduction technique. The typical reduction temperature is as high as 1700°C, which therefore needs intensive energy and releases carbon oxides as wasted gases in the process [6]. Also, another critical disadvantage should be paid more attention: it is difficult to homogenize Nb and Si in any melting process due to the very high melting point of metallic Nb.

During the recent decade, in order to produce high-melting-point metals and superalloys, many new skills were studied by scientists and engineers. Among them, direct electrochemical reduction of metal oxides or mixed oxides in molten electrolytes was an effective method for production of metals and alloys. The typical examples were FFC Cambridge Process and the derived methods. In this process, metal oxides were pressed to solid pellets as a cathode and the graphite were acted as an anode, which were soaked into the molten CaCl₂ at about high temperatures (700–900°C) for electrolysis simultaneously. Consequently, a potential was loaded between the two electrodes. Generally, this technique was used in production of many metals and nonmetals for...
example, Cr, Si, Nb, Ni, and Zr [7–12]. The whole reaction process can be described with the following equations:

Cathode

\[ \text{AO}_x + 2xe^- = A + xO^{2-} \]  \hspace{1cm} (1)

Anode

\[ 2O^{2-} - 4e^- + C = CO_2 \quad \text{or} \quad O^{2-} - 2e^- + C = CO \]  \hspace{1cm} (2)

In these equations, the A is a symbol of metal or nonmetal elements. If the cathode was mixed oxides, the reduction production might be oxygen-free alloys. Therefore, this electrochemical method can be effectively used for refractory metals, high-melting-point nonmetals, and superalloys.

Herein, we used this electrochemical reduction method for the production of NbSi alloys in the molten CaCl\(_2\) at high temperature (900°C). The processors were mixed Nb\(_2\)O\(_5\) and SiO\(_2\) powders of four various atomic ratios of Nb and Si. After electrolysis, the NbSi alloys were successfully obtained.

After scanning electron microscope (SEM), X-ray diffraction (XRD), and energy-dispersive X-ray spectroscopy (EDX) analysis, the NbSi alloys were composed of different phases and almost free of oxygen. Therefore, this electrochemical reduction method had paved a way for the NbSi alloys production, which is also able to be applied to fabricate other refractory alloys.

2. Experimental Details

2.1. Materials Preparation. In this research work, the electrolyte was CaCl\(_2\). In this work, CaCl\(_2\) (≥99.0%), Nb\(_2\)O\(_5\) (≥99.9%), and SiO\(_2\) (>99.8%) powders were purchased from Chengde Optical and Electrical Materials Ltd., Hebei Province, China. The pellets preparation was following the below steps. First, CaCl\(_2\) powders were dried at 80°C overnight and then pressed to small cylindrical blocks with a pressure of 30 MPa. Second, the cylindrical blocks were dried again at 80°C for several hours and then pulverized to small cylindrical pellets with an average sized less than 1 mm. For the preparation of the Nb\(_2\)O\(_5\)/SiO\(_2\) pellets, the process was identical as that used for CaCl\(_2\) pellets preparation. The Nb\(_2\)O\(_5\) and SiO\(_2\) powders were mixed with the atomic ratios of Nb-10Si (Nb, 90 at%; Si, 10 at%), Nb-20Si (Nb, 80 at%; Si, 20 at%), Nb-30Si (Nb, 70 at%; Si, 30 at%), and Nb-37.5Si (Nb, 62.5 at%; Si, 37.5 at%) between Nb and Si. The total mixed powder was 200 g. The graphite crucibles (inner diameter: ~9.0 cm, height: ~18.0 cm, wall thickness: ~1.0 cm) were cleaned with ethanol (>95%, Beijing Keshi Co. Ltd., Beijing, China) for at least 5 times until the inner walls of the graphite crucible were almost free of graphite powders. At last, the mixed Nb\(_2\)O\(_5\)/SiO\(_2\) pellets were placed in the cleaned crucible and then the CaCl\(_2\) pellets (~ 2000 g) were placed on the top of the Nb\(_2\)O\(_5\)/SiO\(_2\) pellets.

2.2. Potentiostatic Electrolysis. The process of electrolysis was conducted in a programmable temperature-controlled furnace produced by China Iron & Steel Research Institute Group. The design of the instrument was shown in the schematic of Figure 1. The graphite crucible filled with Nb\(_2\)O\(_5\)/SiO\(_2\) and CaCl\(_2\) was placed in a graphite stage in the furnace. As an anode, a graphite rod (diameter: 6.0 cm) was just hanged above the crucible vertically by a mechanic instrument, which can push the graphite rod move up and down to ensure that the rod can contact with the molten electrolyte during the whole electrolysis process. The direct current for the electrolysis was supplied by a KGHF-1000A/6V rectifier, made by Beijing Chunshu Electric rectifier Ltd., Beijing, China. The temperature-controlled furnace was evacuated and then filled with argon (≥99.9%, O\(_2\) ≤10 ppm, H\(_2\)O ≤15 ppm, N\(_2\) ≤5 ppm) as soon as possible. It should be noted that the argon flow was maintained with a rate of 1.0–1.5 L min\(^{-1}\) and the outlet was adjusted with a valve to ensure that the gas pressure inside the furnace was about 0.01 MPa larger than the ambient pressure, which was helpful for avoiding air to ingress into the furnace in the electrolysis process. The heating process was of three steps: first, the temperature was heated to 780°C with a ramping rate of 10°C min\(^{-1}\); second, kept at 780°C for 30 min, the CaCl\(_2\) electrolyte was completely melted in the crucible; third, the temperature continued to rise up to 900°C with a heating rate of 10°C min\(^{-1}\). Consequently, the potential was loaded and then stabilized at 3.0 V for electrolysis between the two electrodes. The electrolysis would be lasted for about 4–5 hours, and the graphite crucible with the products was removed out the furnace as the temperature in the furnace was decreased to room temperature after electrolysis. The product was obtained by smashing the crucible.

2.3. Characterizations of Morphology and Composition. Micrograph and composition analyses were conducted with a VEGA\(\backslash\)LMU SEM system equipped with a CCD camera, from TESCAN. EDX equipment was purchased from OXFORD and analysis software was INCA. The phase composition of the NbSi alloys was identified by a Rigaku D/Max 2200 PC X-ray diffractometer.

3. Results and Discussion

After electrolysis at 900°C in molten CaCl\(_2\), the samples were obtained. Figure 2 showed the pictures of the obtained
product after electrolysis. As shown in Figure 2(a), the CaCl$_2$ electrolyte and the sample were molten together after electrolysis. By smashing the crucible, a block of ingot was obtained in Figure 2(b). Clearly, the ingot had two layers of different materials. A metal-like layer with a color of grayish was rich of alloys and other layer was mainly composed of electrolyte.

It should be noted that Figure 2 only showed typical pictures after electrolysis of the processor powder for the alloy of Nb-37.5Si. The other three samples were also of a similar picture, which were not shown here. The morphologies of the alloys were shown in the SEM images in Figure 3. The white small particles with an average size about 10–20 nm had Nb concentration of 98.2 at % in Table 1. On the contrary, the concentrations of the Si and the O were only 1.6 at % and 0.2 at %, respectively. Therefore, it was clearly to show that the white small particles were mainly composed of Nb. In the grey areas, the concentrations of NB and Si were 89.4 and 10.1 at % respectively, which was almost equated to the ratio of Nb-10Si.
alloy. It should be noted that only a very small part of O was still remained in both the white small particles and the grey areas. Although the O was not completely electrochemical reduced, it can be removed by other metallurgical methods in the consequent process. Typically, the mixed \( \text{Nb}_2\text{O}_5/\text{SiO}_2 \) powder was almost completely electrodeoxidized in molten \( \text{CaCl}_2 \) at 900°C. For the processor powder mixture of Nb-20Si alloy, the product after electrolysis the SEM image was shown in Figure 3(b), which was similar to the image in Figure 3(a). The EDX results in the Table 1 also showed that the white particles were Nb and the grey areas were NbSi alloy (Nb-20Si). The similar SEM image and EDX results were shown for the sample Nb-30Si alloy. Compare the three alloys (Nb-10Si, Nb-20Si, and Nb-30Si), two conclusions were obtained. First, the grey areas in all the three images had Nb and Si concentrations close to 62.5 and 37.5 at%, respectively, which was the atomic ratio in the Nb\(_5\)Si\(_3\) alloy. Second, the white small particles were almost mainly composed of metallic Nb.

Third, although O was remained in a little part in all the three samples, most of the O were still removed electrochemically. For the powder mixture of Nb-37.5Si alloy, the product after electrolysis, the SEM image, and EDX result were shown in Figure 3(d) and Table 1, respectively. Clearly, the product in the SEM image showed a homogenous phase with a Nb/Si ratio of 62.2/37.0, which was also close to the atomic ratio of Nb\(_5\)Si\(_3\).

To address the phase composition of the products in the four figures, the XRD patterns were obtained in Figure 4. All of the samples have a phase of \( \text{Nb}_5\text{Si}_3 \) (JCPDS: 30-0874) as indicated with solid circle. Only the alloys of the Nb-10Si, Nb-20Si, and Nb-30Si had a phase of Nb (JCPDS: 35-0789) shown in Figures 4(a), 4(b), and 4(c). Based on the data of the XRD, SEM, and EDX, three critical conclusions can be drawn. First, all the samples were composed of two phases: Nb and Nb\(_5\)Si\(_3\), which were the two phases in NbSi alloys. Second, in
Table 1: EDX results of the four samples in Figure 3.

| Positions | Nb (at%) | Si (at%) | O (at%) |
|-----------|----------|----------|---------|
| a-white   | 98.2     | 1.6      | 0.2     |
| a-grey    | 62.1     | 37.6     | 0.3     |
| b-white   | 99.1     | 0.8      | 0.1     |
| b-grey    | 62.6     | 37.4     | —       |
| c-white   | 98.7     | 1.1      | 0.2     |
| c-grey    | 62.7     | 37.0     | 0.3     |
| d         | 62.2     | 37.5     | 0.4     |

a-white, b-white, and c-white showed the concentrations of Nb, Si, and O in the positions of white small particles in Figures 3(a), 3(b), and 3(c), respectively. Similarly, a-grey, b-grey, and c-grey showed the concentrations of Nb, Si, and O in the positions of grey areas in Figures 3(a), 3(b), and 3(c), respectively. The concentrations of Nb, Si, and O in the sample Nb-37.5Si alloy in Figure 3(d) were shown in position d.

Figures 3(a), 3(b) and 3(c), the small white particles were Nb phase, and grey phase was Nb$_5$Si$_3$. Third, the Nb-37.5Si sample was completely composed of Nb$_5$Si$_3$ phase after electrolysis.

4. Conclusion

We used four mixed powders of different Nb-Si compositions (Nb-10Si, Nb-20Si, Nb-30Si, and Nb-37.5Si) for NbSi alloys preparation. NbSi alloys can be prepared by direct electrochemical reduction method in molten CaCl$_2$ at 900°C, which is an easy and one-pot procedure. After electrolysis, Nb phase showed white scattered particles in the grey Nb$_5$Si$_3$ phase under SEM. XRD, and EDX results have proved that the NbSi alloys were successfully obtained, which had paved a creative way for production of alloys with direct electrochemical reduction method.

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