

SURFACE HARDENING OF TUNGSTEN-BASED HEAVY ALLOY VIA PACK BORONIZING

A. R. Ataei¹, M. Jalaly^{2,*} and M. Tamizifar¹

* maisam_jalaly@iust.ac.ir

Received: December 2016

Accepted: February 2017

¹ School of Metallurgy and Materials Engineering, Iran University of Science and Technology, Tehran, Iran.

² Nanotechnology Department, School of New Technologies, Iran University of Science & Technology, Tehran, Iran.

Abstract: The boronizing of a tungsten heavy alloy containing Ni and Fe as the major alloying elements were performed in the present study to increase its surface hardening. Pack cementation method was employed as a well-known, successful solid-state process for boronizing. The coating treatment was accomplished at different temperatures of 1000, 1050 and 1100°C for 6 and 9 hours. The formation of tungsten boride phase was confirmed, although a silicide layer covered the surface of the specimen as the outer layer. The mechanism of the formation of a multilayered surface was explained. The maximum thickness of reaction zone and surface hardness achieved in the current work were 300 μm and 2470 HV, respectively.

Keywords: Tungsten alloy, Hardening, Boronizing, Pack cementation.

1. INTRODUCTION

Due to the significant developments of industries and need for materials of superior quality, ceramic materials have attracted more attention. Although ceramics possess desired features like good thermal stability, high hardness, excellent corrosion resistance and high stiffness, these materials suffer from the disadvantages of high cost of raw materials, low ductility and susceptibility to brittle fracture [1]. Therefore, surface engineering has been considered for a long time to use the benefits of ceramics as the coating layers on the metallic substrates to improve their oxidation, wear, fatigue, and hardness properties [2].

Boronizing is a well-known thermochemical surface treatment that is applicable for surface hardening of a wide range of ferrous and nonferrous metals. In this process, boron atoms are introduced into the metal lattice at the surface of a work-piece to react with substrate atoms, forming a boride phase. This inward movement of the boron atoms takes place through the high temperature diffusion mechanism. Boronizing can be performed in gas state, liquid state and solid state. Because of the problems of toxic nature and expensiveness of initial materials in gaseous boronizing and technical problems in

liquid boronizing, solid boronizing is normally selected in practical treatments [3, 4].

Pack cementation as the solid boronizing process can be used even for irregular surfaces of the pieces or inside the holes. This process involves burying a well-cleaned metallic substrate in a boron-containing solid powder pack and heating it to between 700 to 1100°C for 1 to 12 hours [5]. Formation of surface boride phase onto the substrate results in a substantial increase in hardness, wear resistance, corrosion resistance, and serviceability of the parts.

Literature survey shows numerous research works on the boronizing of the ferrous materials including various types of steels and cast irons to improve their surface properties [6–11]. Some of intermetallic compounds or cermet materials have also been reported to be covered by boronizing method [12–14]. There is a few works on the boronizing of nonferrous metals. The boride phases of transition metals (like WB, VB₂, ZrB₂, CrB, NbB₂, etc) are very hard and can be excellent resistive films against wear and corrosion for their weak substrates of transition metals. Usta et al. [2, 15] studied the boronizing of pure tungsten, niobium and chromium at 940°C for 2-8 hours with a constant pack composition and compared their surface characterizations. Taracki et al. [16] boronized

pure vanadium at 1100°C for 4-16 hours under the argon atmosphere. Boronizing powder pack used by them had a chemical composition of 90% SiC, 5% B₄C and 5% KBF₄. Boronizing of titanium alloys and the effect of rare earth additions on the process results were also performed [17, 18].

Tungsten is one of the hardest metals in the periodic table. Tungsten has few, if any, substitutes available in its key applications due to its unique properties of heat resistance and wear rate. Hence, tungsten is frequently identified as a critical and strategic material. However, tungsten is often employed as tungsten heavy alloys instead of pure metal. Heavy metal alloys are combinations of tungsten with nickel-iron or nickel-copper alloying components. They are usually produced through powder metallurgy and sintering processes. These heavy alloys with a boronized layer will find broader tribological applications. Therefore, the present work focused on the boronizing of a tungsten-based heavy alloy by pack cementation, which has not been reported so far.

2. EXPERIMENTAL

A tungsten heavy alloy with a chemical composition of ~92wt.% W, ~5 wt.% Ni and ~3wt.% Fe was used as the initial substrate. The specimens for boronizing treatment were prepared by wire cutting in a cylindrical shape with diameter of 7 mm and height of 5 mm. The specimens were ground using 180-2500 grit emery paper, polished by 0.3 µm Al₂O₃ particles, and cleaned by acetone in an ultrasonic cleaner for 10 min. Boronizing process was performed in a tube furnace under an argon atmosphere of 99.99% purity at different temperatures of 1000, 1050 and 1100°C for durations of 6 and 9 hours by pack cementation method. The powder pack had a chemical composition of 90wt.% SiC, 5wt.% B₄C (99%, Sigma-Aldrich), and 5wt.% KBF₄ (99%, JamPack Enterprise Ltd). This composition for powder pack is a typical one for surface boronizing [2, 15, 16]. This mixture was sufficiently blended before treatment to make a homogeneous combination. The powder pack was enclosed by a completely sealed vessel and

subjected to the temperatures. The vessel dimensions were 3cm×4.5cm×7cm.

The boronized samples were cleaned to remove physically bonded powder. X-ray diffraction (XRD) analysis were carried out on the surface of the samples to identify the phases formed during the treatment. A JEOL JDX-8030 instrument with Cu-K_α radiation was used for XRD analysis. The specimens were then cut along their diameter to study cross-sectional surfaces. The cut samples were mounted into epoxy resin for better microscopic observations. Microstructures were investigated by Zeiss optical microscope and Tescan Vega II XMU scanning electron microscope (SEM), equipped by Rontec energy dispersive X-ray spectroscopy (EDS) system. Vickers microhardness measurements were carried out on the surfaces of the specimens by a Leitz microhardness tester under a load of 300 g for 10 s.

3. RESULTS AND DISCUSSION

3. 1. Characterization

Fig. 1 (a) shows spheroidized microstructure of the base alloy before heat treatment obtained by an optical microscope. Spherical grains are pure tungsten, which are surrounded by a softer, Ni-Fe binder phase containing some dissolved tungsten. This is typical structure of this alloy after liquid phase sintering (LPS), which can provide high mechanical properties. Fig 1 (b) also shows the XRD patterns of the substrate material before surface treatment. As seen, the major phase is the bcc pure tungsten phase (ICCD PDF #004-0806), and a minor phase of Ni-Fe-W solid solution with fcc crystal structure is observed. In this solid solution, Ni (ICCD PDF #004-0850) is the matrix which Fe and W are introduced within its lattice.

After boronizing treatment, XRD analyses were performed on the surfaces of the specimens boronized at different temperatures and times. Figures 2–4 show the XRD results of boronizing at 1000, 1050 and 1100 °C, respectively, for 6 and 9 hours. All XRD patterns are almost identical and show a silicide phase as a major surface product. At more extreme conditions of

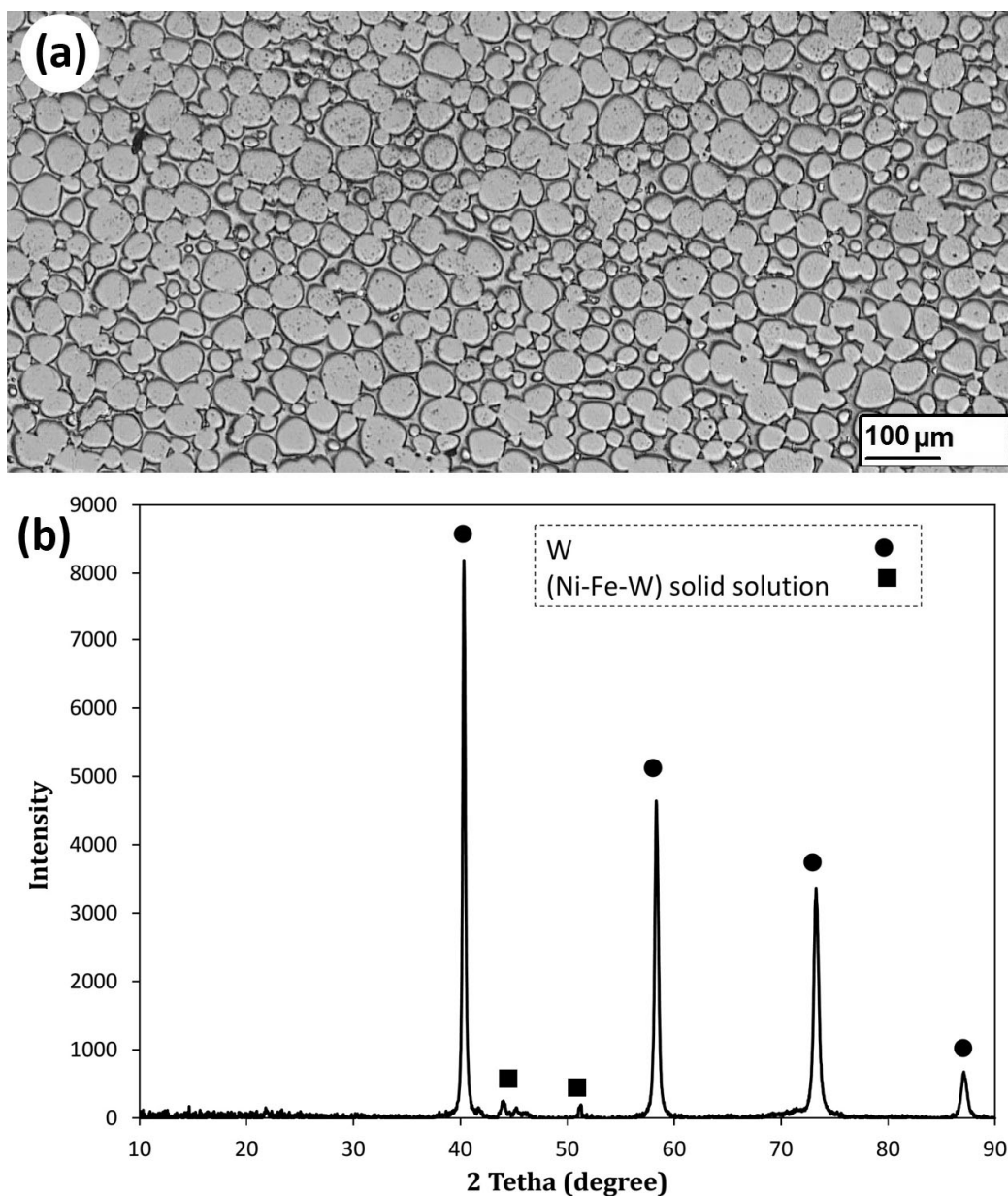


Fig. 1. (a) Optical microstructure and (b) XRD pattern of initial tungsten based heavy alloy.

treatment (1050°C-9h, 1100°C-6 and 9h), the major silicide phase is Ni_5Si_2 (ICCD PDF #003-1073), while at less extreme conditions (1000°C-6 and 9h, 1050°C-6h), Ni_2Si (ICCD PDF #073-2029) is the major silicide phase. A small amount of tungsten boride, WB (ICCD PDF #035-0738), can also be detected along with silicide phases.

Nickel, which is present in the binder areas, reacts to form surface silicide compound. The formation of different types silicide phase as a separate layer on the boride layer was also reported for boronizing of pure nickel [19, 20]. Briefly, the presence of SiC as filler material in powder pack gives rise to the formation of

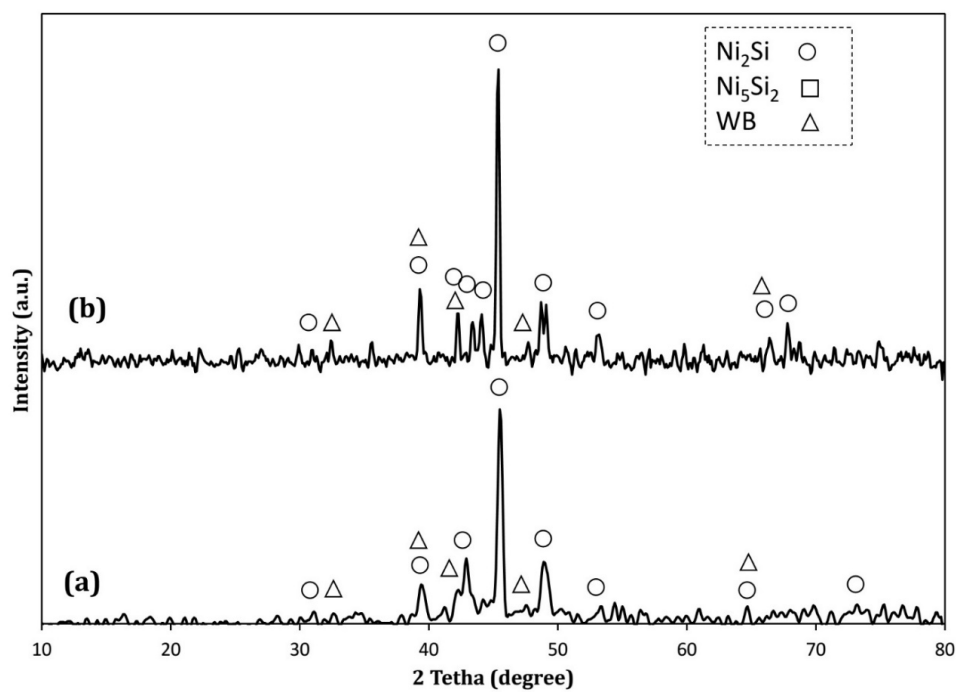


Fig. 2. XRD pattern of the sample boronized at 1000 °C for (a) 6 and (b) 9 hours.

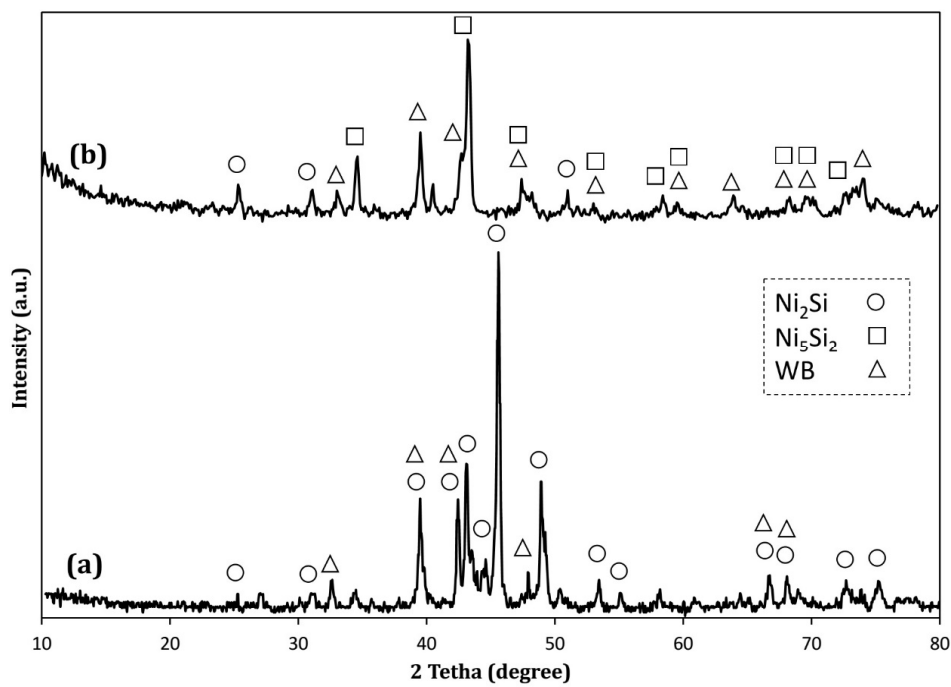


Fig. 3. XRD pattern of the sample boronized at 1050 °C for (a) 6 and (b) 9 hours.

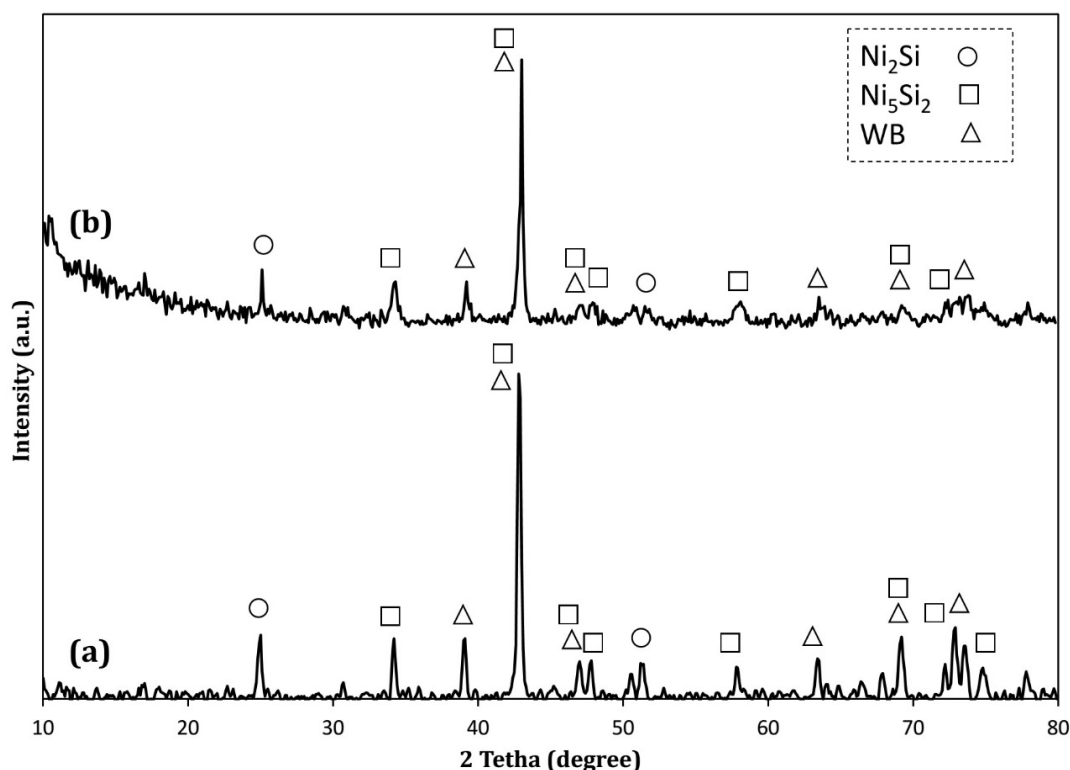


Fig. 4. XRD pattern of the sample boronized at 1100 °C for (a) 6 and (b) 9 hours.

silicide phase on the surface of specimens. In the following, the process mechanism will be explained in detail. The reason for smallness of WB peaks in XRD patterns is that the silicide layer is thick and prevents X-ray from penetration to the deeper layers.

After boronizing treatment, surface and cross section of the samples were examined. As a typical microstructure obtained after treatment, optical cross-sectional micrographs of the samples coated at 1100°C for 6 and 9 h are shown in Fig. 5 (a, b). Three regions can obviously be observed in the surfaces of the samples. These include: (a) outer layer which is mainly composed of silicide compounds (according to XRD patterns) (A region), (b) reaction zone (B region), and (c) non-affected base metal (C region). The thicknesses of reaction zones for different samples are shown in Fig. 6. As seen in this figure, increasing treatment temperature resulted in an increase in reaction zone depth. At

any constant temperature, the thickness of reaction zone is also observed to increase by increasing the duration of boronizing process. In fact, the diffusion depth of boron increases with treatment time and temperature. The thicknesses of the reaction zones obtained from boronizing at various temperatures ranged between 115–207 μm and 150–300 μm for 6 and 9 hours of boronizing, respectively.

SEM micrographs in Fig. 7 show the microstructural evolutions within the reaction zone created after boronizing treatment at different processing times and temperatures. Similar to optical micrographs, the three-layered natures of the surfaces are obviously seen for all samples. The formation of such a configuration can be explained as follows.

3. 2. Governing Mechanisms

The formation of the outer silicide layer

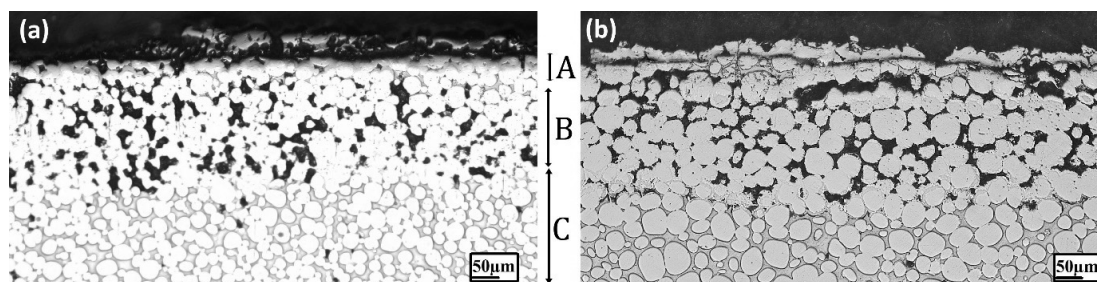


Fig. 5. Typical optical microstructure of the sample boronized at 1050 °C for (a) 6 and (b) 9 hours.

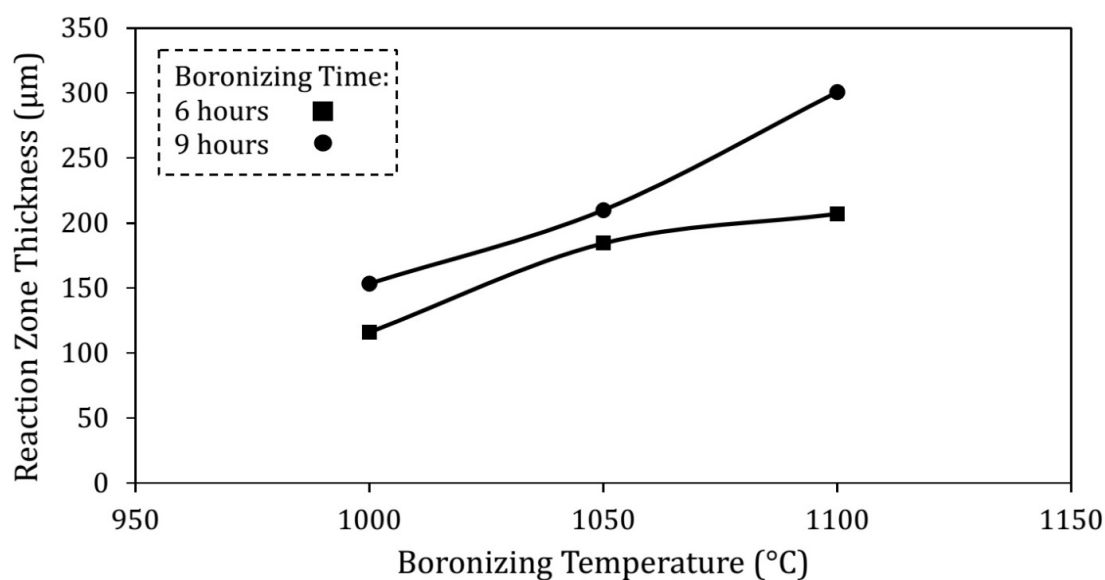


Fig. 6. Reaction zone thickness versus boronizing temperature for different boronizing time.

together with inner boride layer can be explained here. It is believed that the mechanism of the coating formation is directly related to the diffusion phenomena occurred in the material. This system is a multi-component alloy, and is different from a simple metal treatment. In an imaginary A-B alloy, it is well-known that if adding B to A lowers the melting point of A (or the liquidus line), it will increase the chemical diffusion coefficient at any given temperature [21]. Moreover, boron and silicon as the alloying

constituents have been recognized to be melting point depressing elements in high-temperature conditions [22]. Boron and silicon can be introduced into the surface as nascent atoms or gaseous species which is explained later. Therefore, the addition of boron and silicon into the surface of the base metal by diffusion causes the reduction of nickel and iron melting point and increasing their diffusion coefficients, leading to the redistribution phenomena [23]. Hence, nickel and iron that exist in the binder phase within the

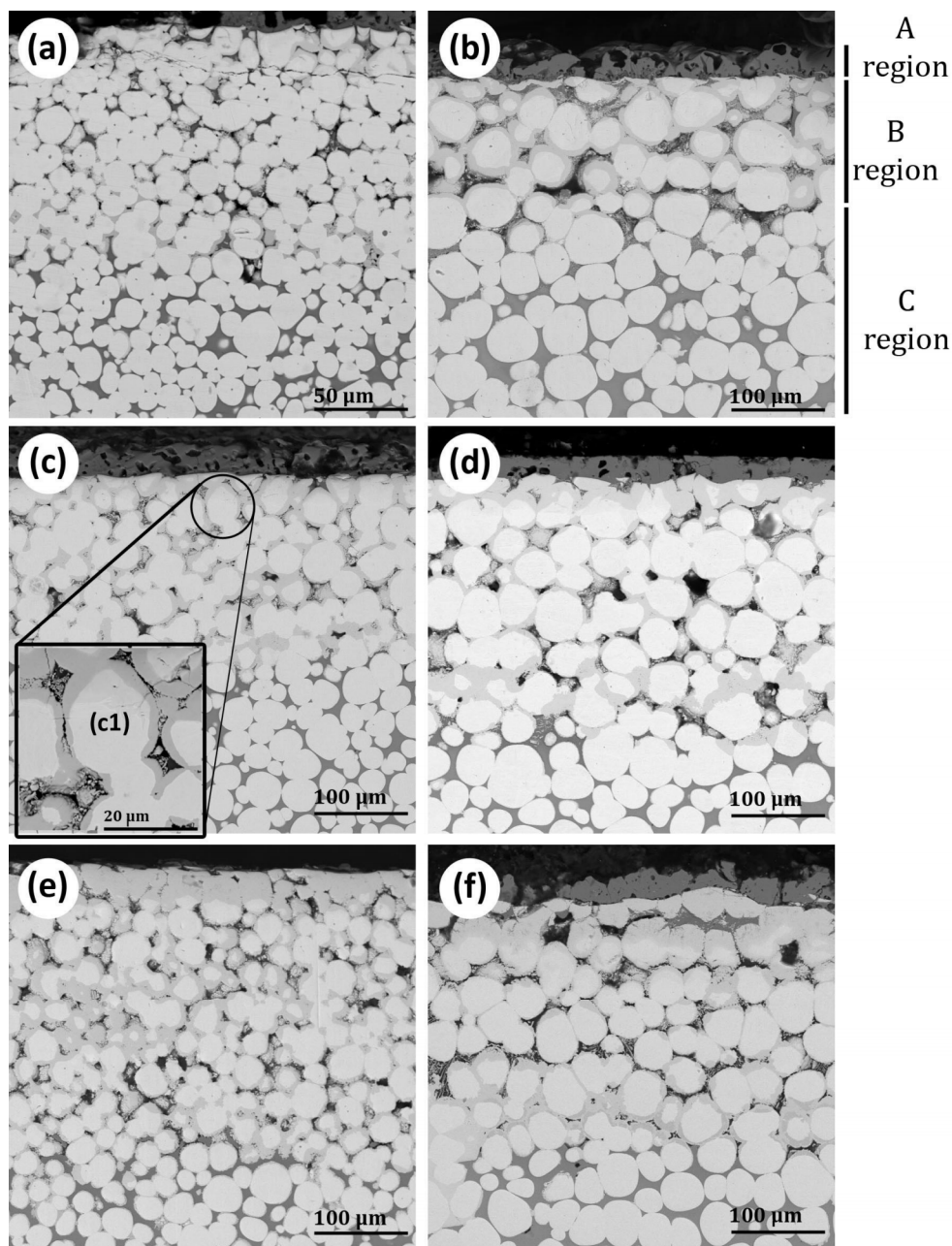


Fig. 7. SEM micrograph of the samples boronized at different conditions including (a) 1000 °C–6 h, (b) 1000 °C–9 h, (c) 1050 °C–6 h, (d) 1050 °C–9 h, (e) 1100 °C–6 h, and (f) 1100 °C–9 h. The inset labeled as C1 is the same micrograph shown in (c) in a higher magnification.

reaction zone tend to change their positions because of their diffusions. They concentrate in the top sections of the surface through an uphill diffusion as a consequence of redistribution phenomena. As a result of this migration to the

top areas, silicon which has a high concentration and activity on the surface due to its presence in powder pack, reacts with nickel and iron to form silicide compounds (like Ni_2Si and Ni_5Si_2 shown in XRD patterns) as an outer layer. No

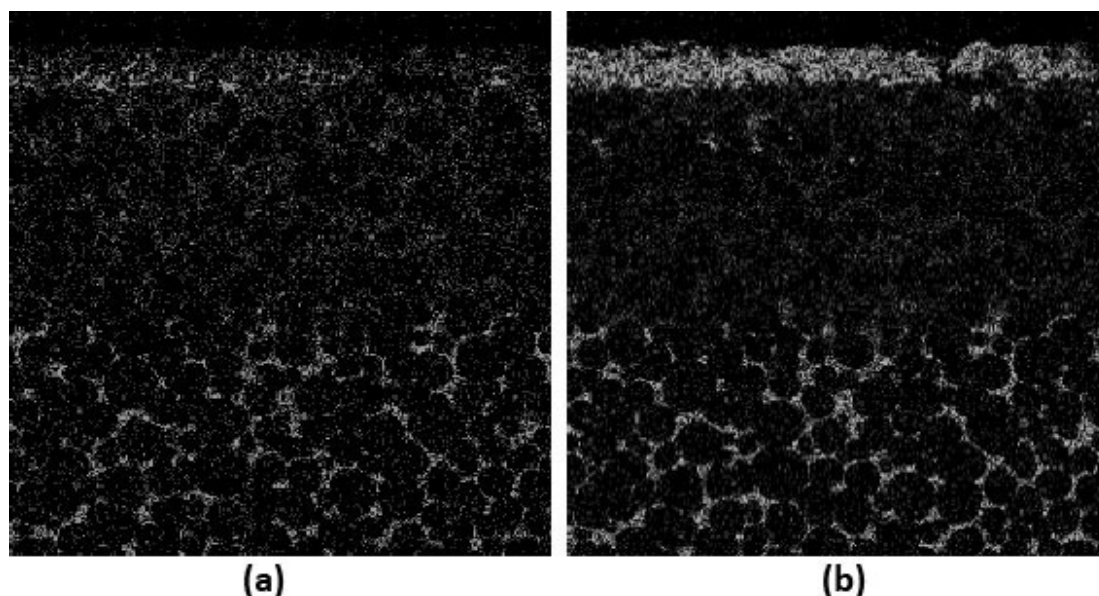
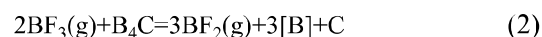


Fig. 8. Elemental maps showing the distributions of (a) Fe and (b) Ni in the surface of the sample boronized at 1050 °C for 6 h.

iron silicide was detected by XRD mostly due to the small amount of iron in this alloy. The elemental mapping taken by EDAX in Fig. 8 shows the concentration of Ni and Fe on the top surface in the outer silicide layer, and the reaction zone can be seen to be almost free of these elements. The formation of outer silicide layer has also been reported by others in boronizing of pure nickel [19, 20]. As can be seen in Figs. 5 and 7, microvoids were formed inside the reaction zone due to the outward transport of Ni and Fe from binder phase areas. Boron has also diffused downhill in an interstitial regime to react with tungsten and create tungsten boride (WB), which is also shown in XRD results. Actually, reaction zone contains spherical unreacted tungsten grains which are surrounded by shell-like features with a darker color (see Fig. 7 (c1)). These affected external shells seem to be formed due to the reaction of diffused B with W on the surface of tungsten grains. Indeed, the boride phase has formed beneath the outer silicide layer. Such a core-shell configuration in reaction zone has already been reported where carburization of tungsten heavy alloy was performed to create a

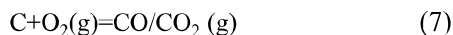
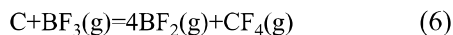
tungsten carbide coating [24].

The governing chemical mechanism for the formation of boride phase can be described by a series of reactions. Nascent atomic boron must be formed and diffused inward to produce boride phase. KBF_4 cannot produce nascent boron alone and B_4C plays a major role in production of atomic boron. During high temperature process, KBF_4 decomposes into KF and gaseous BF_3 . The following series of reactions may be reasonable for the mechanism [12, 14, 25, 26]:



Gaseous SiF_4 formed in the reaction (3) can be the active agent on the surface to react with Ni and Fe to produce silicide compounds (e.g. $2\text{Ni} +$

$\text{SiF}_4(\text{g}) = \text{Ni}_2\text{Si} + 2\text{F}_2(\text{g})$). The carbon produced in reactions (2) and (3) can be consumed by the following possible reactions:



where the oxygen in the reaction (7) may result from argon atmosphere as a minor impurity, or from remaining air in boronizing media.

3. 3. Hardness Evaluation

The Vickers microhardness of uncoated tungsten substrate was measured to be about 380 HV. The silicide phase which was formed as the outer layer, is a ceramic compound with an intermediate hardness of 800-850 HV [15, 16]. To evaluate the hardness of boride layer below the silicide layer, the surfaces of samples were carefully ground to remove the silicide layer from the top. The new surfaces of the samples were subjected to microhardness test. Fig. 9 (a) shows a typical indentation mark on a grounded surface after microhardness test. Fig. 9 (b) shows variations of surface hardness values for samples boronized at different temperatures and times after removing silicide layer. The dashed line in the diagram specifies the level for hardness of

the substrate. The hardness of borided surfaces was greater than 2000 HV, which is typical of tungsten boride hardness [2]. The maximum hardness (2470 HV) was achieved by the sample coated at 1100°C for 9 hours. The tremendous difference between the hardness of bare and boronized tungsten heavy alloy is an obvious indication for the formation of boride phase on the surface (below the silicide layer). A slight decrease in the hardness of the samples coated at 1050°C may be attributed to the human error resulted from inaccurate grinding the surface to remove the silicide layer. As can be seen, the hardness of the samples increased by increasing the time of boronizing process. This is in agreement with the result that the thickness of reaction zone was shown to increase with increasing treatment time (Fig. 6).

4. CONCLUSIONS

A simple and effective boronizing process was successfully carried out to create a very hard surface on the tungsten-based heavy alloy. Pack cementation containing SiC (90 wt.%), B_4C (5 wt.%) and KBF_4 (5 wt.%) was used. By boronizing at temperatures of 1000, 1050, and 1100 °C for 6 and 9 hours, three distinct regions were identified on the surface of specimens. These include (i) outer silicide layer, (ii) reaction

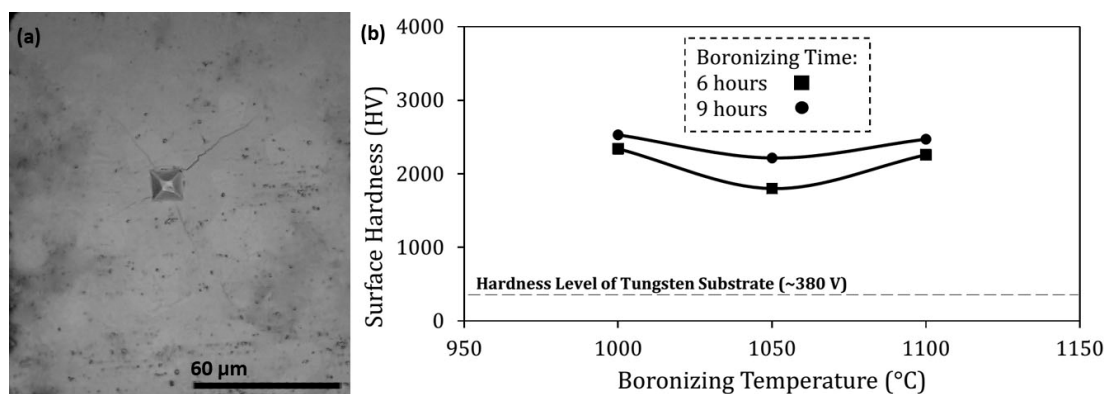


Fig. 9. The microhardness test was accomplished after removing outer silicide layer. (a) Typical indentation mark created after hardness test on the surface. (b) The surface hardness of the samples boronized at different temperatures for different times.

zone, and (iii) base metal. Reaction zone is composed of tungsten grains with a shell of WB phase resulted from downhill diffusion of boron within tungsten lattice. The outer silicide layer creates through a reaction between outwardly diffused nickel with silicon originated from silicon-rich powder pack. The thickness of reaction zone in different samples ranges between 115–300 μm , which increases by increasing boronizing time and temperature. To reach the very hard boride layer, silicide layer was readily grounded. The microhardness of tungsten alloy substrate was approximately 380 HV, which significantly increased to about 2000–2500 HV for various boronizing times and temperatures.

REFERENCES

- Callister, W. D. and Rethwisch, D. G., "Materials science and engineering: an introduction", 9th Edition, John Wiley & Sons Inc, New Jersey, USA, 2014, 468–500.
- Usta, M., Ozbek, I., Ipek, M., Bindal, C. and Ucisik, A. H., "The characterization of borided pure tungsten". *Surf. Coat. Technol.*, 2005, 194, 330–334.
- Fichtl, W., "Boronizing and its practical applications". *Mater. Eng.* 1981, 2, 276–286.
- Anthymidis, K. G., Maragoudakis, N., Stergioudis, G., Haidar, O. and Tspas, D. N., "A comparative study of boride coatings obtained by pack cementation method and by fluidized bed technology". *Mater. Lett.*, 2003, 57, 2399–2403.
- Sinha, A. K., *Boriding (Boronizing)*, ASM Handbook, vol. 4, ASM International, Materials Park, OH, USA, 1991, 437–447.
- Campos-Silva, I., Flores-Jiménez, M., Rodríguez-Castro, G., Hernández-Sánchez, E., Martínez-Trinidad, J. and Tadeo-Rosas, R., "Improved fracture toughness of boride coating developed with a diffusion annealing process". *Surf. Coat. Technol.*, 2013, 237, 429–439.
- Calik, A., Simsek, M., Karakas, M. S., Ucar, N., "Effect of boronizing on microhardness and wear resistance of steel AISI 1050 and chilled cast iron". *Met. Sci. Heat Treat.*, 2014, 56, 89–92.
- Bektes, M., Calik, A., Ucar, N. and Keddari, M., "Pack-boriding of Fe–Mn binary alloys: Characterization and kinetics of the boride layers". *Mater. Charact.*, 2010, 61, 233–239.
- Balusamy, T., Narayanan, T. S. N. S., Ravichandran, K., Park, I. S., Lee, M. H., "Pack boronizing of AISI H11 tool steel: Role of surface mechanical attrition treatment". *Vac.* 2013, 97, 36–43.
- Anthymidis, K. G., Stergioudis, G., Roussos, D., Zinoviadis, P., Tspas, D. N., "Boriding of ferrous and non-ferrous metals and alloys in fluidized bed reactor". *Surf. Eng.*, 2002, 18, 255–259.
- Gencer, Y., "Influence of manganese on pack boriding behavior of pure iron". *Surf. Eng.*, 2011, 27, 634–638.
- Guobiao, L., Zhongjian, Z., Zhihai, Q., Xiang, L., Jiahua, W. and Feifei, Z., "Boronizing mechanism of cemented carbides and their wear resistance". *Int. J. Refract. Met. Hard Mater.*, 2013, 41, 351–355.
- Ekmekçiler, E., Polat, A. and Usta, M., "Hard boride coating on iron aluminide (FeAl)". *Surf. Coat. Technol.*, 2008, 202, 6011–6015.
- Li, C., Li, M. S., Zhou, Y. C., "Improving the surface hardness and wear resistance of Ti_3SiC_2 by boronizing treatment". *Surf. Coat. Technol.*, 2007, 201, 6005–6011.
- Usta, M., Ozbek, I., Bindal, C., Ucisik, A. H., Ingole, S. and Liang, H., "A comparative study of borided pure niobium, tungsten and chromium". *Vac.* 2006, 80, 1321–1325.
- Tarakci, M., Gencer, Y., Calik, A., "The pack-boronizing of pure vanadium under a controlled atmosphere". *Appl. Surf. Sci.*, 2010, 256, 7612–7618.
- Zhu, Y. S., Lu, W. Z., Zuo, D. W., Sun, Y. L., Wang, H. and Xu, J., "Effects of rare earth additions during surface boriding on microstructure and properties of titanium alloy TC21". *Surf. Eng.*, 2013, 30, 1765–1769.
- Wang, H., Lu, W. Z., Xu, J., Zhu, Y. S., Huang, Q. C., Zhang, L. F., Zuo, D. W., "Effect of rare earth (RE) on pack boronizing process of titanium alloy". *Surf. Eng.*, 2013, 30, 123–128.
- Mu, D., Shen, B., Yang, C., Zhao, X., "Microstructure analysis of boronized pure nickel using boronizing powders with SiC as

- diluent". Vac.2009, 83, 1481–1484.
20. Ozbek, I., Akbulut, H., Zeytin, S., Bindal, C., Ucisik, A. H., "The characterization of borided 99.5% purity nickel". Surf. Coat. Technol., 2000, 126, 166–170.
21. Shewmon, P., "Diffusion in Solids", 2nd Edition, John Willey & Sons Inc, New Jersey, USA, 1991, 147–148.
22. Dinkel, M. K., Heinz, P., Pyczak, F., Volek, A., Ott, M., Affeldt, E., Vossberg, A., Göken, M., Singer, R. F., "New boron and silicon free single crystal diffusion brazing alloys", 11th International Symposium on Superalloys, Ed.R. C. Reed, Pennsylvania, USA, 2008, 211–220.
23. Palombarini, G., Carbucicchio, M., "Influence of carbon on the chromium redistribution when boriding iron alloys". J. Mater. Sci. Lett.,1993, 12, 797–798.
24. Eroglu, S., Ekren, H., Baykara, T., Surface hardening of tungsten heavy alloys. Scr. Mater.,1998, 38, 131–136.
25. Lu, X. X., Liang, C., Gao, X. X., An, J. and Yang, X. H., "Catalysis of Rare Earth Element Nd on Boriding of AISI 1045 Steel". ISIJ Int.,2011, 51, 799–804.
26. Mareels, S., Wettinck, E., Van Peteghem, A. P., "Thermodynamics of carbide-fluoride systems in pack cementation deposition", Thermochemistry of Alloys, Recent Developments of Experimental Methods, Ed. H. Brodowsky and H.J. Schaller, Springer, Berlin, Germany, 1989, 513–526.