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Synthesis of Hard Layer by Titanium Addition During Welding Process and Quenched Directly

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ABSTRACT

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Keywords: Hardfacing Ti Welding Hardness Quenching Oil Enhance the surface hardness of materials usually conducted through a hardfacing technique. Hardfacing is popular, whereby materials with better properties are deposited over cheaper bulk material. This work fabricated hard layers by adding titanium (Ti) wire during the welding process. This research used low-carbon steel as the base material, wire optime Ti grade 1 for Ti addition, and an HV 600 electrode with a diameter of 3.2 mm for filler metal. A single-layer weld was conducted with SMAW (positive polarity and 90 A). The samples were directly quenched in a different solution after welding. The properties of the weld layer were examined phase, structure, microstructure, macrostructure, and hardness using optical emission spectroscopy (OES), x-ray diffraction (XRD), an optical microscope, a digital camera, and a hardness device, respectively. Adding titanium (Ti) to the weld layer and quenching the samples after welding in the solution enhances the hardness. This phenomenon is attributed to different phase compositions, oxides, and microstructures. A fine dispersion of small particles and oxide amount is important in increasing hardness. There is no cracking in the weld and base metal. In conclusion, samples BNTiO and BNTiM are recommended for lathe-cutting tools.

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1. INTRODUCTION

High Speed Steel (HSS) is a commonly used cutting tool in the lathe. For dry-cutting operational conditions, four factors need consideration: pressure, temperature, sliding speed, and the interface. Thus, factors could affect high wear rate, heat, and short tool life [1]. These conditions could be reduced by increasing the hardness. HSS commercial lathe-cutting tool hardness is around 765.68 HV [2].

Enhance the surface hardness of materials usually conducted through a hardfacing technique. Hardfacing is popular, whereby materials with better properties are deposited over cheaper bulk material [3]. Furthermore, hardfacing is one of the most economical techniques to mitigate the wear of tools and components of a machine [4]. By increased the hardness on the surface of the metal with less hardness is the cheapest than using a bulk metal with high hardness. Hardfacing applies a hard surface layer on the base material to resist abrasion, impact, and erosion.

Welding is one technique for fusion between two materials [5]. The welding process also could be conducted to reach the hard layer. Therefore, a hard layer could result from welding processes such as shielded metal arc welding (SMAW) [6, 7], Flux core arc welding (FCAW) [6], plasma transferred arc welding (PTA) [8], electro-slag welding (ESW) [9], and gas tungsten arc welding (GTAW) [10]. The SMAW technique is more flexible and easy to operate than others.

Commonly filler metals used for hardfacing are iron (Fe)-based alloys, cobalt (Co)-based alloys, and nickel (Ni)-based alloys. Singh [11] has reported hardfacing single layers through various hard alloy filler metals (iron (Fe)-based alloys), resulting in hardness between 375 and 543 HV. A different electrode composition has resulted in different hardness. A filler metal with a high carbon (C), Manganese (Mn), and chromium (Cr) has a resulting

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high hardness. Deng et al. [12] reported that hardfacing by cobalt (Co)-based alloys resulted in different hardness for different thicknesses. Less thickness is resulting high hardness. This phenomena is caused by the absence of the fine grain when the thickness is decreased. Kesavan and Kamaraj [8] have found that aging treatment of a nickelbased hardfacing has reduced the hardness when treatment time increased.

Titanium (Ti) and Ti-alloys are widely applied in marine, petrochemical, chemical, aerospace, and biomedical industries because promising high specific strength and high corrosion resistance [13, 14]. Furthermore, Ti also promises good hardness and erosion resistance [15]. Therefore adding a Ti into the weld layer could increase the hardness. Zhou et al. [16] have added 0.263 wt% Ti in the flux of the SMAW filler metal (Fe-Cr13-C-Nb). Exhibit Ti leads to a decreasing hardness from 46.2 to 45.2 HRC. Ti could be aglomerated perfectly between filler and based metal because the flux is changed into gasses as shielding for molten metal when the SMAW process is performed. Zhou et al. [16] found that the hardness of the layer increased from 56 to 61 HRC with titanium content from 0 to 0.63 wt%. Increased Ti content to 1.17 wt% could form TiC carbide because too much carbon is consumed by titanium. It led to a change in the microstructure to a hypereutectoid one and caused a decrease in the hardness. Moreover, the exhibit Ti content in the layer also increases the hardness than Zr addition by approximately 603 and 584 HV, respectively [17].

In addition, increasing the hardness of a material can also be conducted by quenching techniques in various solutions such as oil [18, 19]. Vegetables and engine oil were used for quenching operations to avoid undesirable microstructure [18, 20]. Dauda et al. [20] quenched carbon steel in engine oil, and Totten et al. [21] also quenched steel in vegetable oil which could increase the hardness. Various oils for quenching solution would be resulting different microstructures due to different cooling rates. Engine oil and vegetable oil at 40 °C have a cooling rate of 64.4 and 71.9 K/s, respectively [21]. The cooling rate decreases as the viscosity of the solution increases [22].

Before quenching, the sample was heated in a muffle furnace. Commonly, the muffle furnace temperature is set to > 800°C, and the specimen was heated 0.5 until 1 hour [18]. For cost efficiency, the sample quench after the welding process is similar to when the sample heating with a muffle furnace and then rapidly quenching because the temperature during the welding process is around 1200 °C, which is similar to heating with a muffle furnace [23].

In this research, the hardfacing by the commercial electrode and adding Ti during welding on the lowcarbon steel was conducted. Moreover, after welding, each sample was directly quenched in vegetable oil and engine oil. It is worth to be mentioned and to highlight the major concern about properties and hardness result by adding Ti in the weld layer and directly quenching method after weld. This method is more cost-effective than heating the sample with a muffle furnace. The application of this research is for lathe cutting tools application.

2. MATERIALS AND METHODS

This research used low-carbon steel $(150 \times 10 \times 10 \text{ mm})$ as the base material. The composition of the base material is 0.128 wt% C, 0.359 wt% Si, 0.996 wt% Mn, 0.225 wt% Cr, 0.167 wt% Cu, and Fe balance (see Figure 1). Wire optime Ti grade 1 with a diameter of 0.65 mm (from the USA) was used for Ti addition. The weight of each Ti wire used is 0.225 g. HV 600 electrode (typical composition: 0.6 wt% C, 1 wt% Mn, 0.4 wt% Si, 4wt% Cr, and Fe balance) with a diameter of 3.2 mm (from Nikko Steel Manufacturer) was used for filler metal.

Prior welding process, the based material surface was cleaned by hand grinder. HV 600 electrode was dried for 1 hour (150 °C). Three pieces of Ti wire ($150 \times \emptyset$ 0.65mm) were placed on the surface of low carbon steel and welded single layer with SMAW (Fro BF 443, positive polarity, and 90 A).

This research was resulting three different samples that were welded using an HV 600 electrode without Ti addition and directly cooled in the natural air, quenched in 1L of engine oil (SAE 20-50), and 1 L of palm oil, namely BNU, BNO, BNM, respectively. Afterward, three samples were welded using an HV 600 with Ti addition. They directly cooled in the natural air, quenched in 1 L of palm oil, and 1L of engine oil (SAE 20-50) were BNTiU, BNTiO, and BNTiU, respectively. The sample after the weld is shown in Figure 2.

The samples were checked with Was Foundry Master-Optical Emission Spectroscopy (OES) according to ASTM E415. X-ray diffraction (XRD) PANalytical Aeris with CoK=1.789 Å and step size 0.0217° was used to analyze the phase composition and crystal structure. The Rietveld method was used to calculate the lattice parameters [24]. Furthermore, the sample was mounted in the epoxy resin, grinding, polishing, etching in 3%



Figure 1. Base material



Figure 2. Weld samples

Nital, and captured by a digital camera for macrostructure. It was also checked by an inverted optical metallurgy microscope Olympus BX41M - LED for microstructure investigation (ASTM E407). Afterward, samples were hardness measured by Vickers hardness tester FV-300e (5kg_f of load). Five readings were collected to have average hardness.

3. RESULT AND DISCUSSION

3.1. OES Based on the OES result, the base material was welded with HV 600 electrodes, resulting welded layer in the composition: 0.342 wt%C, 0.182 wt%Si, 0.608 wt%Mn, 2.09 wt%Cr, 0.139 wt%Cu, and Fe balance. The 0.342 wt%C in the steel is close to medium carbon steel [25].

The base material was welded with HV 600 electrode and added a Ti wire resulting welded layer in the composition: 0.443 wt%C, 0.366 wt%Si, 0.923 wt%Mn, 2.19 wt%Cr, 0.144 wt%Cu, 0.147 wt% Ti, and Fe balance. Ti was perfect adding in the weld layer, as proved by the OES result.

3. 2. XRD Figure 3 shows the diffraction pattern of the various samples. The plane peaks (110) and (211) are reflections of the α -iron (ferrite) phase. These α -iron phase plane peaks are in good agreement with the results of other studies [26-28]. The peak magnification of 110 plane diffraction patterns in the inset of Figure 3 ($2\theta = 50-53^{\circ}$) shows an asymmetric peak shape. According to Han et al. [28], this asymmetrical peak denotes the presence of an α '-iron (martensite) phase. The peak at an angle of $2\theta = 41.3^{\circ}$, identified as the Fe₃O₄ phase, illustrates how the welding process used results in the oxidation of the steel.

Different intensity values can be seen in the diffraction patterns of the four samples. This intensity value can be connected to the full width at half maximum (FWHM) value of each sample. The BNTiO sample has the smallest crystallite size, according to the qualitative analysis of the FWHM values of the α -iron and α '-iron phases in the (110) and (211) planes, as summarized in Table 1.



TABLE 1. Quantitative analysis of diffraction patterns for various samples

Source	BNU	BNTiU	BNTiO	BNTiM
Ferrite phase				
Crystal system: Body Center Cubic (BCC)				
Space group: Im-3m				
Lattice a (nm)	0.2878	0.2867	0.2881	0.2875
Martensite phase				
Crystal system: Body Center Tetragonal (BCT)				
Space group: I4/mmm				
Lattice a (nm)	0.2855	0.2857	0.2859	0.2821
Lattice c (nm)	0.2935	0.2958	0.2931	0.2876
α-Fe (wt.%)	81.24	60.35	72.75	85.41
α'-Fe (wt.%)	15.99	32.30	23.86	13.02
Fe ₃ O ₄ (wt.%)	2.77	7.35	3.39	1.57
FWHM (110) (°)	0.695	0.858	0.548	0.680
FWHM (211) (°)	0.980	1.090	1.030	1.120
Rwp (%)	3.30	5.31	3.46	3.52
χ^2	1.360	1.369	1.662	1.268

The composition of the phase formed and lattice constants were determined through the quantitative analysis of the diffraction patterns. Figure 4 compares the results of the calculated and observed diffraction patterns for the BNTiM sample. This study uses the Rietveld method to analyze diffraction patterns quantitatively. The overlapping peak (110) of the ferrite and martensite phase fractions were calculated using the Rietveld method in order to determine its composition. The ferrite and martensite phase fractions for the BNTiM sample were 85.41 wt.% and 13.02 wt.%, respectively (see Table 1). The sample with the greatest α -iron phase content out of the other three is the BNTiM sample.



Figure 4. Diffraction patterns for the BNTiM sample calculated with the Rietveld method

The BNTiM sample, which has a body center cubic (BCC) structure with the space group Im-3m, has a lattice constant value (a=b=c) of 0.2875 nm. Table 1 shows the lattice constants for the four samples. These constant lattice values are identical to those previously reported [29-31].

3. 3. Microstructure Microstructure observation for the samples with an optical microscope was conducted as shown in Figure 5.

Based on Figure 5, we can observe different microstructures were formed. Samples with air cooling have similar characteristics for Ti addition and without Ti addition. It seems 0.147 wt% Ti addition does not influence the microstructure. This phenomenon is due to



(b)

(d) (e)

(f) **Figure 5.** Microstructure result (a) BNU, (b) BNO, (c) BNM, (d) BNTiU, (e) BNTiO, and (f) BNTiM

only 0.147 wt% Ti being formed in the welded layer based on the OES result. Zhou et al. [16] have reported 0.28 wt%Ti could change the microstructure. When the

Ti composition of Ti raised to 1.17 wt%, the microstructure significantly changed.

From Figures 5(a) and 5(d), we can see a un uniform distribution of ferrite (white region) and martensite (dark area). This behavior was confirmed with the XRD result, where 81.24 wt % ferrite and 15.99 wt % martensite were in the BNU. Moreover, 60.35 wt % ferrite and 32.30 wt % martensite were realized in the BNTiU. The distribution and compositions formed between martensite and ferrite could avoid cracking [32]. It has been no cracking seen in the samples based on microstructure observation.

Quenching in palm oil and engine oil significantly influences the microstructure, similar to other cases reported in literature [20]. Figure 5 shows the martensitic phase and perfect agreement with different research results [33]. Quenching the samples in the solution led to uniformly changing ferrite (α -Fe) distribution between martensite (α '-Fe). Randomly martensite structure is distributed in a ferrite matrix after quenching [34]. This phenomenon is due to the oil's rapid cooling; hence, conducted ferrite becomes finer and does not grow more [35]. The quench heat treatment would be resulting smooth distribution of ferrite and martensite [36].

3. 4. Macrostructure The samples were compared to evaluate the effect of the quenching solution and Ti addition to the macrostructure. Macrostructure observation with a digital camera is shown in Figure 6.

Based on Figure 6, we can see different macrostructures were formed. Fusion between the base metal and weld metal is shown for various samples.



Figure 6. Macrostructure result (a) BNU, (b) BNO, (c) BNM, (d) BNTiU, (e) BNTiO, and (f) BNTiM

There are no visible cracks on the macrostructure. Commonly, this occurs influenced by welding conditions, mechanical restraint, and hydrogen trapping [37, 38]. It means the welding condition in this research could avoid this, and no hydrogen trapping appears in the weld metal.

The penetration of the weld layer would increase by raising the current of welding. Shukla et al. [39] showed 1.2 mm deep penetration when welding using 90 A and positive polarity. Rising to 120 A, reach a deep penetration of around 1.75 mm. The negative polarity and 90° of the electrode angle with a current at 120 A revealed full penetration because of more heat to the weld region [35, 39]. Moreover, a higher current for positive polarity is needed to get full penetration than negative polarity [40]. Plate thickness also influences penetration. Shifting to more thicknesses could increase penetration [35]. In this research, a single layer of weld penetrates approximately 3 mm for various samples. It shows perfect agglomeration between the base material and weld metal. Adding Ti in the weld layer is not influence the penetration. This result differs from others because welding parameters such as an electrode, current, and electrode are different. Commonly penetration of the weld layer is affected by the polarity, current, and angle of the electrode for the SMAW process [39].

3.5. Hardness Distributions A comparison of the sample hardness was carried out in Figure 7.

Many factors influence hardness were formed. Khamari et al. [35] found that increasing the current would increase the hardness. Dauda et al. [20] found various oil for quenching media resulting in differences in hardness. An exhibit of a fine dispersion of small particles of ferrite and martensite also contributed to higher hardness [20].

Based on Figure 7, we can see that quenching the sample increased the hardness. BNU sample was resulting an average hardness of 503.48 HV. This hardness agrees with the manufacturer of a diverse range of advanced welding consumable HV 600 for single-layer welding, resulting in hardness between 440 - 540 HV [41]. Quench the sample into oils significantly increase the hardness [20].



Figure 7. Hardness Vickers of the different specimens

Its seem samples where quenching in oils has higher hardness than air cooling. Air and oil cooling could increase the hardness by around 4.5% and 40%, respectively [42]. Sample where quenching in palm oil has a higher hardness than engine oil. This value is attributed to palm oil having less viscosity than engine oil and thus affecting the higher cooling rate. As we know, engine oil has 64.4 and vegetable oil has 71.9 K/s of viscosity at 40° C [20, 21]. A shift to a higher cooling rate led to higher hardness [43]. For this study, we found that palm oil is the best quenching solution to increase the hardness.

Adding Ti also increases the hardness of the sample when quenching in a different solution. This result perfectly agrees with other research where adding Ti improves the hardness, whereas adding 0.63 wt% Ti could increase the hardness from 56 to 61 HRC [16]. Increasing the Ti from 0.63 to 1.17 wt% could decrease the hardness because too much carbon is consumed when titanium content is 1.17 wt% [16]. Therefore, there is a limitation contained in the layer when Ti is added. Moreover, 0.147 wt% Ti in the weld layer increases hardness by around 0.2% for a sample with and without the quenching process.

Based on Table 1, we can realize that the quenched sample with Ti addition in the engine oil and palm oil led to a decrease in the oxide (Fe₃O₄). This phenomenon may be attributed to the solution cooling rate that increased sample hardness. Iron oxide could affect the strength of the material [44].

Compared to Table 1, an increase in the hardness linearly within increase in the ferrite for a sample with Ti addition. Adding Ti could increase in the hardness of the ferrite matrix [45]. The ferrite phase is the main factor in increasing the hardness in this research. A shift to higher ferrite composition (see Table 1) and fine dispersion of small ferrite particles could enhance the hardness.

Based on the hardness test result, BNTiO, and BNTiM have more hardness than the HSS commercial for a lathe-cutting tool [2]. This value means the samples are recommended for the cutting tool. But this needs deep further investigation.

4. CONCLUSIONS

The effect of Ti addition and quenching of the samples is summarized as follows:

- 1. Adding titanium (Ti) to the weld layer and quenching the samples after welding in the solution enhances the hardness. This phenomenon is attributed to different phase compositions, oxides, and microstructures.
- 2. A fine dispersion of small ferrite particles and oxide amount is important in increasing the hardness.

- 3. Adding Ti in the weld layer did not influence the penetration. Therefore, there is no cracking in the weld and base metal.
- BNTiO, and BNTiM samples are recommended for cutting tools.

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Persian Abstract

چکیدہ

سختی سطح مواد را که معمولاً از طریق تکنیک هارد فاسینگ انجام می شود، افزایش دهید. Hard facing رایچ است، به طوری که مواد با خواص بهتر بر روی مواد فله ارزان تر رسوب می کنند. این کار با افزودن سیم تیتانیوم (Ti) در طول فرآیند جوشکاری، لایه های سختی را ساخت. این تحقیق از فولاد کم کربن به عنوان ماده پایه، سیم بهینه Ti با درجه ۱ برای افزودن Ti، و یک الکترود HV 600 با قطر ۳.۳ میلی متر برای فلز پرکننده استفاده کرد. یک جوش تک لایه با SMAW (قطب مثبت و ۹ A) انجام شد. نمونه ها به طور مستقیم پس از جوشکاری در محلول دیگری خاموش شدند. خواص لایه جوش به ترتیب فاز، ساختار، ریز ساختار، درشت ساختار و سختی با استفاده از طیف سنجی انتشار نوری (OES)، پراش اشعه ایکس (XRD)، میکروسکوپ نوری، دوربین دیجیتال و دستگاه سختی مورد بررسی قرار گرفت. افزودن تیتانیوم (Ti) به لایه جوش و کوئنچ کردن نمونه ها پس از جوشکاری در محلول باعث افزایش سختی می شود. این پدیده به ترتیب فازی، اکترار می و را گرفت. افزودن تیتانیوم (Ti) به لایه پرای و کوئنچ کردن نمونه ها پس از جوشکاری در محلول باعث افزایش سختی می شود. این پدیده به ترکیبات فازی، اکسیدها و ریز ساختارهای مختلف نسبت داده می شود. پراکندگی ریز ذرات کوچک و مقدار اکسید در افزایش سختی مهم است. هیچ ترک خوردگی در جوش و فلز پایه وجود ندارد. در نتیجه، نمونههای می قلود. پراک گر در از می روس می شوند.