

Microstructure and Mechanical Properties of WC-Co Reinforced With NbC

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Cemented carbides such as tungsten carbide-cobalt WC-Co composites have been widely used as cutting tool materials. Several reports have shown the influence of different factors such as grain size, type and amount of binder phase and the addition of hard particles on the properties of WC-Co. The purpose of this work was to investigate the effect of niobium carbide on the microstructure and mechanical properties of WC-Co. Specimens of WC-Co reinforced with NbC were mixed and subsequently hot-pressed in a inert atmosphere. The WC-Co-NbC composite material exhibited high hardness values (18.9 GPa), flexural strength (2100 MPa) and fracture toughness (11.2 MPa.m^{1/2}). TEM analysis has shown a bimodal grain size distribution of WC.

Keywords: tungsten carbide, hardness, fracture toughness, niobium carbide, microstructure

1. Introduction

Cemented carbides such as WC-Co and WC-Co-TiC are the most widely used material for metalworking¹. As a consequence, a considerable amount of research effort has been spent to develop alternatives cemented carbide systems in order to improve the microstructure and mechanical properties of these materials. Figure 1 shows the influence of TaC(NbC) addition on the mechanical performance of WC². The hardness decreases only slightly with increasing TaC(NbC) content, but the decrease in transverse-rupture strength is remarkably high for TaC(NbC) contents exceeding 20 wt. (%). Recent studies have shown an increase in hardness and fracture toughness of WC through the use of different binding phases and the synthesis of nanocomposites^{3,6}. The addition of some hard refractory particles such as TiC, VC and Mo₂C has produced a pinning effect, reducing the grain growth of the WC-Co matrix^{3,7,8}, which contributes to the final properties of the composite material. Literature data about the use of NbC in WC is still scant. Brazil holds the main world niobium reserves, making the study on possible application of niobium very strategic.

On the other hand, WC and NbC have also been reported

as reinforcement in alumina matrix, due its high hardness and elastic modulus^{9,10}. Hot-pressed alumina reinforced with NbC has caused an increase of the hardness and fracture toughness from 16.5 GPa and 2.9 MPa.m^{1/2} to 19.0 GPa and 4.2 MPa.m^{1/2}, respectively.

The objective of this work was to study the influence of NbC on the microstructure and mechanical properties of hot-pressed WC-Co composite material.

2. Experimental Procedure

The starting powders used in this work consisted of WC (Wolfram Bergbau, Austria), with D50 = 1 µm and 10 wt. (%) of cobalt powder as binder phase. Niobium carbide (Herman Starck Berlin) with D50 = 2.3 µm was used as reinforcing element. The WC and Co powders were mixed with 2 wt. (%) of NbC, homogenized for four hours in a planetary ball mill and uniaxially hot-pressed at 1250 °C during 1 h under a pressure of 20 MPa in flowing argon. Microstructural analysis was carried out on sintered samples in a transmission electron microscope (TEM, Philips, CM 30) operated at 300 kV. Energy dispersive X-ray spectrometry (EDS) was

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performed using a SiLi-detector (type 7370, ISIS 30, Link, Oxford) attached to the TEM. Hardness and fracture toughness were determined using a Vickers hardness testing machine by using the method of Shetty *et al.*¹¹. The analysis of crystalline phases was determined by X-ray diffraction.

3. Results and Discussion

The X-ray powder diffraction pattern of pure and reinforced WC-Co solely showed the presence of WC and Co phases (Table 1). No peaks of NbC or other crystalline reaction product phases were identified. The addition of NbC to WC-Co system seems to promote the formation of a carbide phase that contains both W and Nb, possibly (W,Nb)C. The atomic radii of W (140 pm) and Nb (146 pm) are nearly the same. Thus, the NbC may be dissolved within the WC-phase due to substitution of W by Nb. There is no (W-Nb)C phase diagram published in the literature. There are well known phase diagrams that show the mutual solubility of (Nb,gU), (V,Nb), (bTi,Nb), (Ti,V), (Ti,W) and (Cr,W)¹² and Nb-W¹³.

Similar behavior was also observed in alumina reinforced with (Ti,W) carbides¹⁴. The results indicated the formation of a solid solution of (Ti-W)C, which corresponds to the Ti-W binary phase diagram¹⁴. Hot-pressing resulted in dense materials, with fraction densities in the range from 98.5 to 99.0% theoretical density (Table 1). Similar densities have been reported for hot-pressed alumina-NbC composites¹⁵.

Figure 2 shows the mechanical properties of WC-Co and WC-Co-NbC composites investigated in this work. For comparison, the hardness and fracture toughness of Al₂O₃

and WC-Co reinforced with hard particles are also presented in Fig. 2. The mechanical properties of WC-Co-NbC material are comparable to those of the reinforced materials available in the literature. The introduction of NbC into WC-Co cemented carbide resulted in a slight decrease of fracture toughness from 12 to 11.2 MPa.m^{1/2}. The addition of NbC to the WC-Co composite caused practically no effect on the hardness values as compared to pure WC-Co. The Hv-values of the WC-Co-NbC material remained almost the same as that for the WC-Co composite (19 GPa). The addition of different types of reinforcing elements did not influence considerably the fracture toughness values of WC-Co. K_{IC} values vary from 10.2 to 12 MPa.m^{1/2}. Figure 2 shows that the hardness is improved when WC is incorporated in alumina. The alumina-WC composite material exhibits a hardness value of 21 GPa, slightly higher than WC-Co reinforced material¹⁵.

Introduction of TiC and TiC+Mo₂C into WC-Co cemented carbide results in a drastic decrease in the hardness from 19 to 14 GPa. Substitution of TiC by NbC improves the hardness of WC-Co material. Further increase in hardness is observed when VC is added to the system. The hardness of a WC-composite material depends strongly on the processing method and the grain size of the raw materials^{3,6,7}. This dependence can explain the variation in hardness values of the different WC-Co composites, presented in Fig. 2.

Table 1. Mechanical properties and crystalline phases of WC-Co-NbC sintered specimens.

Properties	WC-Co	WC-Co-NbC
Theoretical Density (%)	99.0	98.5
Flexural Strength (MPa)	2000 ± 220	2100 ± 200
Crystalline Phases	WC, Co	WC, Co

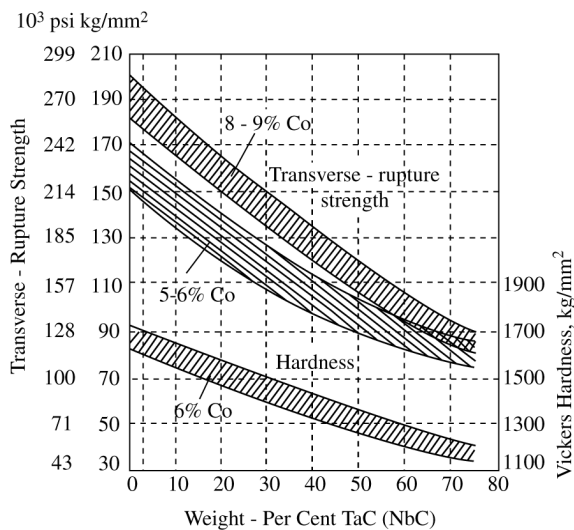


Figure 1. Mechanical properties of WC-Co with TaC-NbC¹¹.

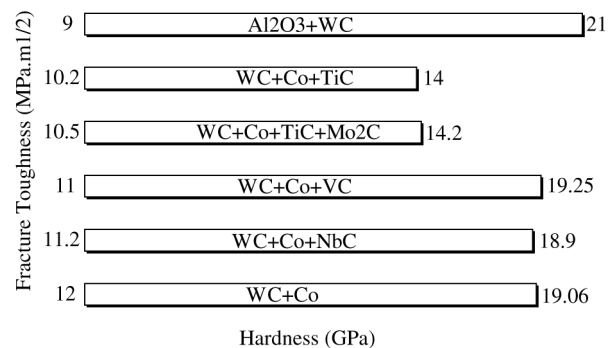
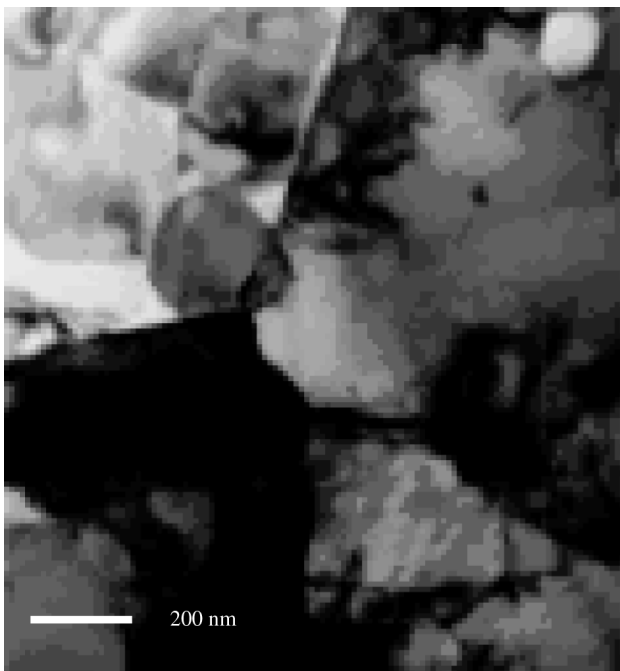


Figure 2. Comparison of hardness and fracture toughness values from various different WC-materials^{2-6,8,14}.

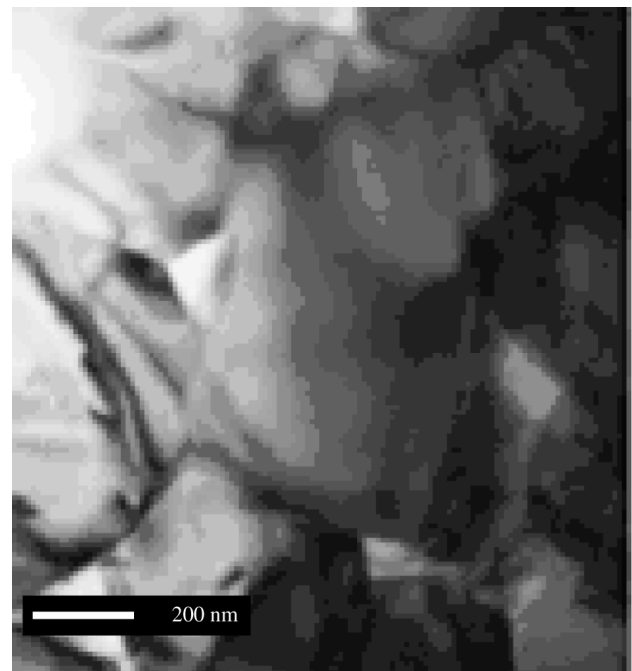
TEM investigation was used to observe the morphological aspects and the microstructure of WC-Co-NbC composites. The composite ceramic material exhibited a heterogeneous microstructure with a bimodal WC-grain size distribution (Figs. 3a and 3b). The microstructure consisted of large (1 μ m) and fine (100-200 nm) grains of WC. The WC-particles were analyzed by EDS (Fig. 3c). The main elements found in the spectra were W, Co and Fe, whereas Nb was only hardly detectable (see inset in Fig. 3c). The results obtained by EDS indicate that the

particles formed were principally composed of WC. A typical STEM image and a EDS line scan with element distribution of W, Co, Nb, C, and Fe is given in Fig. 4. The results suggest that the dark phase consisted primarily of tungsten and carbon (WC) and the bright phase of Co and Fe, respectively. The presence of niobium was not detected, possibly due the small amount of this element in the analyzed region.

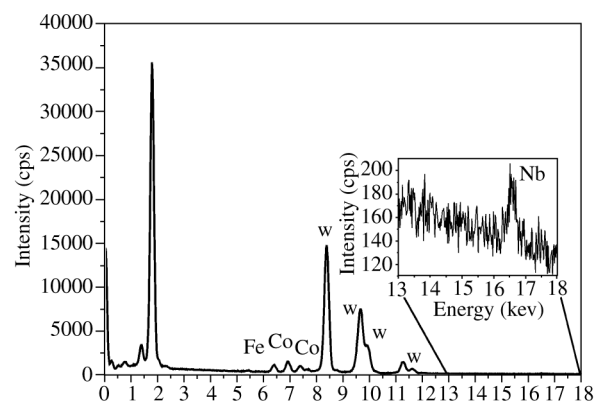
Figure 5 shows EDS maps of the spatial distribution of the elements within the analyzed region. The region in-



(a)



(b)



(c)

Figure 3. a) and b) TEM micrographs of WC-Co-NbC composites; c) EDS analysis of the WC-Co-NbC composites in the TEM.

investigated was the same of Fig. 4. The observed data confirm that the dark grains correspond to W and very small amounts of Nb and the bright region basically to Co and

Fe. The Nb appears to be equally distributed over the analyzed region suggesting the formation of (W,Nb)C. The results obtained in this work indicated that the WC-Co-

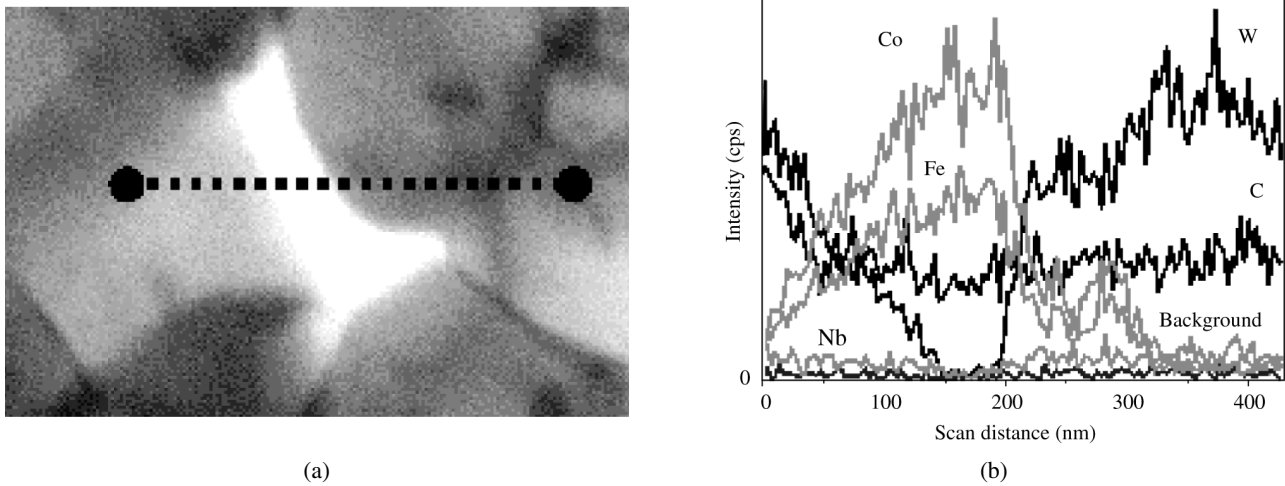


Figure 4. STEM image and corresponding EDS line scan.

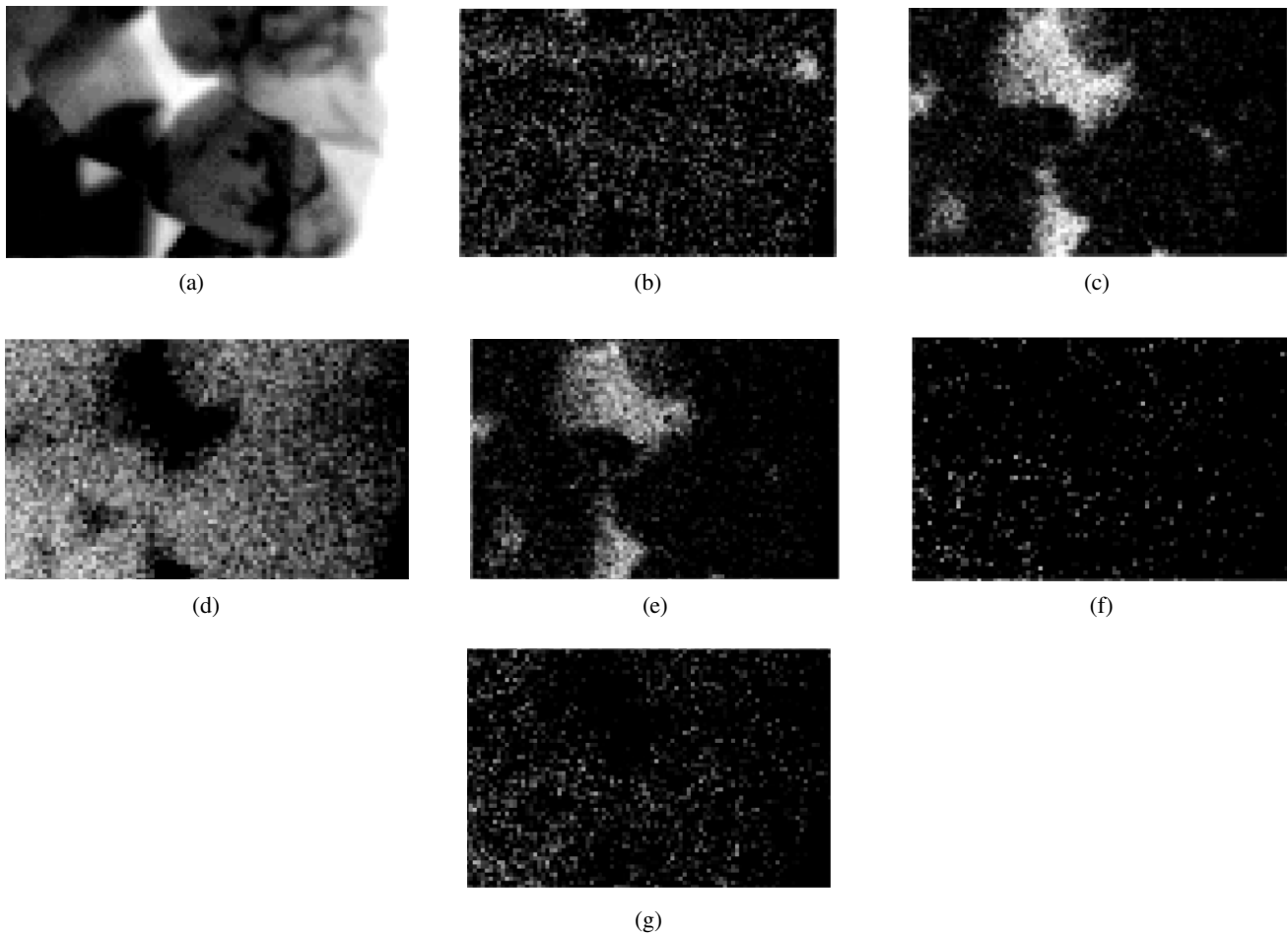


Figure 5. STEM image and EDS maps of W, Co, Nb, C and Fe.

NbC ceramic composites show a good potential to be developed. Further investigation is still under way to study the dependence of higher niobium carbide content on the microstructure and mechanical performance of NbC-reinforced WC-Co.

4. Conclusions

Dense specimens of WC-Co-NbC were fabricated by hot-pressing. No NbC or other new Nb-phases were detected by X-ray diffraction. The addition of NbC to the WC-Co system seems to promote the formation of a carbide phase that consisted of tungsten and niobium (W,Nb)C.

Incorporation of 2 wt. (%) NbC into WC-Co cemented carbide produced a composite material with mechanical properties comparable to the other cemented carbides systems, suggesting that the composite can be a good candidate to metalworking. TEM micrographs have shown a microstructure consisting of large and fine grains of tungsten carbide.

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References

1. Bradt, G. *Mater. Technology* v. 14, n. 1, p. 17-24, 1999.
2. Kopf, P.S.; Kieffer, R.; *Cemented Carbides*, The Macmillan Company, p. 188-198, 1960.
3. Schubert, W.D.; Bock, A.; Lux, B. *Int. J. of Refractory & Hard Mater* v. 13, p. 281-296, 1995.
4. Bhaumik, S.K.; Balasubramaniam, R.; Upadhyaya, G.S.; Vaidya, M.L.; *J. Mater. Sci. Lett.* v. 11, p. 1457-59, 1992.
5. Zhu, Y.T.; Manthiran, A. *J. Amer. Ceram. Soc.*, v. 77, n. 10, p. 2777-78, 1994
6. Leiderman, M.; Botstein, O. A Rosen, Proceeding of the 4th European Conference on *Advanced Materials* and Process, Italy, September 1995.
7. Bhaumik, S.K.; Upadhyaya, G.S.; Vaidya, M.L. *Mater. Sci. Tech.* v. 7, p. 723-27, 1991.
8. Sepulveda, R.; Arenas, F. *Inter. J. Refract. & Hard Mater.* v. 19, p. 389-396, 2001.
9. Acchar, W.; Martinelli, A.E.; Vieira, F.A.; Cairo, C.A. *Mat. Sci. Eng.* v. 284, p. 84-87, 2000.
10. Pasotti, M.R.; Bressiani, A.H.; Bressiani, J.C. *Inter. J. Refrac. & Hard Mater.* v. 16, p. 423-27, 1998.
11. Shetty, D.K.; Wright, I.G.; Mincer, P.N.; Clauer, A.H. *J. Mater. Sci.* v. 20, p. 1873-82, 1985.
12. Murray, J.L. *Bull. Alloy Phase Diagram* v. 2 n. 2, 1981.
13. *Metals Handbook*, ed. 5, v. 8, p. 284, 1973.
14. Acchar, W.; Fonseca, J.L. *Mater. Sci. Eng A.*, v. 371, p. 382-387, 2004.
15. Acchar, W.; Greil, P.; Martinelli, A.E.; Cairo, C.A.; Bressiani, A.H.; Bressiani, J.C. *J. Europ. Ceram. Soc.* v. 20, p. 1765-69, 2000.
16. Weon-Pil Tai, Watanab, T. *J. Am. Ceram. Soc.* v. 81, n. 6, p. 1673-76, 1998.