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Near-net shape manufacture of B₄C–Co and ZrC–Co composites by slip casting and pressureless sintering

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Abstract

Fabrication of near-net shaped B₄C–Co and ZrC–Co composites by slip casting and pressureless sintering is described. It is shown how B₄C–Co and ZrC–Co concentrated suspensions can be prepared by aqueous colloidal processing, and optimized (in terms of pH, deflocculant contents, and sonication time) to have a shear-thinning rheological behaviour suitable for the near-net shaping of the corresponding cermet compacts by slip casting. It is also demonstrated that the robust, highly-dense compacts so obtained have a uniform green microstructure without macrodefects or gradient density, and which can be fully densified by

pressureless sintering. Specifically, it is shown that B_4C -Co compacts densify by reactive and transient liquid-phase sintering, thus resulting in multi-component ceramics. ZrC -Co compacts densify however by persistent liquid-phase sintering, thus resulting in cermets. An explanation is given for these observations, and general implications are discussed for the near-net shape manufacture of these and similar carbide-metal composites for use in engineering applications.

Keywords: cermets; B_4C ; ZrC ; colloidal processing; pressureless sintering.

1. Introduction

Ceramic-metal composites (i.e., cermets) have great interest as structural materials for a broad variety of engineering applications because they appropriately exploit the benefits of the hard ceramic matrix and of the tough metal binder, reason for which they are known with the collective name of "hardmetals", or "cemented carbides" if the ceramic matrix is specifically a refractory carbide [1]. The most widely used and investigated of the hardmetals is WC -Co/Ni, but other formulations of cemented carbides are equally possible as long as the metal binder allows the liquid-phase sintering of the carbide. In this context, pure B_4C and ZrC are two refractory carbides (a covalent carbide the former, and an interstitial carbide the latter) with great appeal in structural applications [2-4]. Thus for example, owing to their high or ultrahigh hardness, they may find use in the field of contact mechanics and tribology [5-8]. Also, by virtue of its extreme lightness, B_4C is ideal ceramic armour for both personnel and vehicle protection [9], while, due to its great refractoriness, ZrC belongs to the short list of ultra-high-temperature ceramics for extreme environment applications [10]. Unfortunately however, the great potential of B_4C and ZrC as engineering ceramics is limited by their brittleness and poor pressureless sinterability. This latter also conditions the near-net shape manufacture of B_4C and ZrC

ceramics, as they are best densified with the aid of external pressure (for example using hot pressing, hot-isostatic pressing, or especially spark-plasma sintering) [2,5-8,11-18]. They thus appear to be appropriate candidates for the fabrication of cermets [19-23].

Recently, we have demonstrated that it is possible to obtain robust B_4C -Ni compacts by slip casting from B_4C -Ni concentrated suspensions prepared by aqueous colloidal processing [23]. This combination of techniques is certainly interesting since aqueous colloidal processing allows the challenging co-dispersion of the ceramic and metal phases in the form of environmentally-friendly suspensions with optimized rheological properties, whereas slip casting allows the subsequent near-net shaping of compacts demanded by industry. A Co binder offers some benefits over Ni binder, for example, in terms of hardness and corrosion resistance [24,25]. The former is very relevant because high hardness is an important attribute of the covalent and interstitial carbides required in many engineering applications, and especially in those within the field of contact mechanics and tribology. Therefore, it seems timely to extend the previous study on B_4C -Ni to the potentially appealing B_4C -Co composite and, in addition, to expand it to the also potentially interesting ZrC -Co composite to thus cover two representative types of different refractory carbides. Moreover, the earlier study on B_4C -Ni was devoted exclusively to aspects of aqueous colloidal processing and shaping by slip casting, with no information about sintering, reason by which it also seems timely to conduct additional studies on the B_4C -Co and ZrC -Co composites of both pressureless sintering and microstructural characterization. These all were indeed the objectives of the present study.

2. Experimental procedure

The starting materials used were commercially available B_4C (>99%, $d_{50} \sim 0.5 \mu m$, $\rho = 2.51 \text{ g/cm}^3$; Grade HD 20, H.C. Starck, Germany), ZrC (>99%, $d_{50} \sim 3-5 \mu m$, $\rho = 6.7 \text{ g/cm}^3$; Grade B,

H.C. Starck, Germany), and Co (>99.8%, $d_{50} \sim 1.6 \mu\text{m}$, $\rho = 8.92 \text{ g/cm}^3$; Alfa Aesar, Germany) powders. They were characterized independently by scanning electron microscopy (SEM; Quanta 3D FEG, FEI, The Netherlands) to confirm the particle sizes indicated by the manufacturers, and by X-ray diffractometry (XRD; D8 Advance, Bruker AXS, Germany) and X-ray photoemission spectroscopy (XPS; K-Alpha, Thermo Scientific, UK) to qualitatively examine the phase compositions and the bonding environments or statuses, respectively.

The procedure of aqueous colloidal processing followed the standard protocol. In particular, in a first stage the colloidal stability of the ceramic (i.e., B_4C and ZrC) and metal (i.e., Co) powders was studied individually using dilute suspensions (0.1 g/l) with a short equilibrium (i.e., stabilization) time (30 min). To this end, systematic zeta potential measurements (Zetasizer Nano-ZS, Malvern, UK) were first made (in triplicate) as a function of pH (adjusted within the acidic or basic ranges using 10^{-1} M HCl or KOH solution, respectively) on single-phase suspensions prepared without deflocculant using deionized water as suspension medium and KCl 10^{-2} M as inert electrolyte. Next, additional zeta potential measurements were also made (in triplicate) at natural pH as a function of the deflocculant content (in the range 0–2.2 wt.%). In the three cases, deflocculation was done using commercially available polyelectrolytes, specifically, a synthetic polyelectrolyte of unknown composition (PKV; Produkt KV5088, Zschimmer-Schwarz, Germany – it is nonetheless thought to be of type polycarboxylic) for both B_4C and ZrC , and an ammonium salt of polyacrylic acid with a molecular weight of 2400 and a content of active matter of 35% (PAA; DuramaxTM D-3005, Rohm & Haas, USA) for Co.

In a second stage, multi-component concentrated suspensions were prepared in deionized water containing ceramic (either B_4C or ZrC) and metal (Co) in relative concentrations of 80 and 20 vol.%, respectively. These suspensions were prepared to a total solids loading of 30 vol.%

because the earlier study on B₄C–Ni indicated that higher solids loadings (in the range ~30–40 vol.%) increase the suspension viscosity thus complicating the slip casting performance without improvement of the green-body densification [23]. The suspensions were prepared with continuous vigorous mechanical stirring with helices to ensure correct mixing, using the following protocol of sequential addition. First, the pH of the deionized water was adjusted to the desired value by dropwise addition of an aqueous solution of 25 wt.% tetramethylammonium hydroxide (TMAH, Aldrich-Chemie, Germany). Next, the PAA content required to appropriately disperse the Co powder was added, followed by incorporation of the Co powder, stabilization (i.e., equilibration) for 5 min, and sonication for 1 min. Then, the PKV content required to appropriately disperse the ceramic powder was added, followed by addition of the ceramic powder, stabilization for 15 min, and lastly sonication for different times to thus investigate its effect on the rheological behaviour of the suspension. The pH was maintained at the desired value throughout the entire process.

The rheological behaviour of the B₄C–Co and ZrC–Co suspensions was studied using a rheometer (Haake Mars, Thermo Fisher Scientific, Germany) operated in controlled shear rate mode. The measuring system consisted of a double-cone and plate, with cone angle of 2°, provided with a cover plate to avoid evaporation. The measurement cycle of the flow curves involved a linear stretch of shear rate increase from 0 to 1000 s⁻¹ in 300 s, then a plateau at 1000 s⁻¹ for 60 s, and lastly a linear decrease to zero shear rate also in 300 s. The thixotropy/rheopexy was determined from the area of the flow curve's hysteresis loop, and the viscosity by direct reading at 1000 s⁻¹ in the uploading stretch.

For the fabrication of the B₄C–Co and ZrC–Co composites, the corresponding multi-component concentrated suspensions with appropriate rheological behaviour were next slip cast

on plaster moulds to obtain compacts (at least 6 specimens per each composition) with different morphology, whose green-body microstructure was examined by SEM after drying in air at room temperature for 48 h within the moulds. The degree of green-body densification was also evaluated (in triplicate), using to that end Hg intrusion porosimetry (PoreMaster 60, Quantachrome Instruments, UK). Finally, these compacts were pressureless sintered (1000-3560-FP20, Thermal Technology Inc., USA) at 1700 °C for 2 h (heating and cooling ramps of 10 and 20 °C/min, respectively) in a flowing Ar-gas atmosphere of 99.999% purity. The 1700 °C temperature was chosen because it ensures the existence of a low-viscosity liquid phase during pressureless sintering given that Co has a melting point of ~1500 °C. Once polished using conventional ceramographic methods, their microstructure was investigated by SEM together with energy-dispersive X-ray spectrometry (EDS) and XRD. The sintered B₄C–Co and ZrC–Co composites were also broken, and their fracture surface was observed by SEM. The degree of densification of the sintered B₄C–Co and ZrC–Co composites was evaluated by image analysis of multiple SEM images (taken randomly on both polished and fracture surfaces).

3. Results and discussion

Figs. 1 and 2 show representative low- and high-magnification SEM images, respectively, of the B₄C, ZrC, and Co starting powders. It can be seen that the particles in the B₄C powder are equiaxed and have an average size in the submicrometre range (Figs. 1A and 2A). The particles in the ZrC powder are also equiaxed (although less faceted), but have greater size dispersion. Indeed, there are particles with sizes from the ultrafine range to the micrometre scale (Figs. 1B and 2B). Finally, it can also be seen that the particles in the Co powders have a worn-like morphology, with average length and thickness in the micrometre and submicrometre ranges, respectively (Figs. 1C and 2C). This is because they are actually intergrowths of finer

particles (Fig. 2C). There is also a rough mat on the surface of the Co particles (Fig. 2C), which is likely a passivating oxide layer. The XRD patterns shown in Fig. 3 indicate that the B₄C powder contains B₄C together with H₃BO₃ and graphite impurities (very minor phases), that the ZrC powder contains only ZrC, and that the Co powder contains a combination of α -Co (fcc) and β -Co (hcp). Lastly, the XPS spectra shown in Fig. 4 confirm the presence of H₃BO₃ in the B₄C powders, and also indicate that the particles of ZrC and Co in the corresponding powders are notably passivated (surface ZrO₂ in the former case, and both surface CoO and Co(OH)₂ in the latter case). Unlike what occurs for the solid-state sintering of these refractory carbides, fortunately however these oxidic impurities are not likely to play a dominant role in their densification by liquid-phase sintering.

Fig. 5 shows the variation of the zeta potential against pH for the individual dilute suspensions of B₄C, ZrC, and Co. It can be seen that the B₄C used here has its surface negatively charged within the entire pH range investigated, always with high absolute values of the zeta potential (approximately in the range -35 mV to -60 mV) [23]. This is attributable to the formation of surface boron oxides and interfacial boron oxycarbides [26,27]. Consequently, this B₄C will be very stable colloiddally in its current state under the vast majority of possible aqueous processing conditions. Note however that other B₄C powders, even from the same supplier, exhibited zeta potential curves in aqueous media with an isoelectric point at pH in the range 5–6 [28], which is likely due to their different grades making them to behave differently. It can also be seen that ZrC has its isoelectric point at pH ~ 3.4 , above which the ZrC particles are already negatively charged. This reflects the surface passivation of the ZrC particles [29]. This ZrC reaches high absolute zeta potential values (i.e., -30 mV) for pHs greater than ~ 6 , above which its colloidal stability in water seems to be ensured. Finally, it can be seen that Co has its

isoelectric point at pH \sim 8.3, exhibiting an abrupt change from positive to negative zeta potentials. This is attributable to the surface passivation of the Co particles [29]. Above pH \sim 8.5, and particularly at pH \sim 10, Co exhibits zeta potentials high enough (i.e., -30 mV) to ensure its colloidal stability in water. Considering this set of results, it seems clear that pHs above 8.5 are required to prepare the multi-component B₄C–Co and ZrC–Co suspensions in the absence of deflocculants, with pH \sim 10 being optimum. However, as will be described next, anionic polyelectrolytes were also used to improve the dispersion at pH 10 by shifting the isoelectric points towards more acidic pHs.

Fig. 6 shows the variation of the zeta potential against deflocculant content for the individual dilute suspensions of B₄C, ZrC, and Co, at their natural pH. Deflocculation of both B₄C and ZrC was done using PKV because it is suitable for carbides [23,30-32], whereas Co was deflocculated using PAA because it has been found to be more appropriate for metals [23,33]. It can be seen that both B₄C and ZrC always exhibit negative zeta potentials. This was to be expected for B₄C (which has negative zeta potentials over the entire pH range), and in the case of ZrC is understandable considering that the natural pH is higher than the isoelectric point (i.e., \sim 6.1 vs. 3.4). In both cases the zeta potential increases in magnitude with the PKV addition, but the increase is more marked for ZrC (from -33 mV to -48 mV, and therefore \sim 45.5% increase) than for B₄C (from -47 mV to -55 mV, and therefore 17% increase). Moreover, the zeta potentials of both B₄C and ZrC are already stabilized (at -55 and -48 mV, respectively) for a PKV addition of 1 wt.%, an amount that will be used to deflocculate the ceramic particles in the multi-component concentrated suspensions. The effect of the deflocculant addition is much more pronounced in the case of Co. Its zeta potential changes from positive to negative values with increasing PAA content up to \sim 0.3 wt.%, attributable to the natural pH being lower than the

isoelectric point (i.e., ~ 7.7 vs. 8.3), and then continues to increase in absolute value until it stabilizes (at -32 mV) for a PAA addition of 1 wt.%. The metal particles in the multi-component concentrated suspensions will therefore be deflocculated using 1 wt.% PAA.

Multi-component concentrated suspensions of B_4C-Co and of $ZrC-Co$ were then prepared at pH 10 to a total solids loading of 30 vol.%, whose formulation contains 80 vol.% ceramic (deflocculated with 1 wt.% PKV) and 20 vol.% metal (deflocculated with 1 wt.% PAA). Fig. 7 shows the flow curves of those B_4C-Co and $ZrC-Co$ suspensions as a function of the sonication time. It can be seen in Fig. 7A that the flow curves of the B_4C-Co suspensions are very similar, and indicative of shear-thinning rheological behaviour. This is the desirable behaviour because the suspension viscosity under rest conditions is high enough to avoid settling and therefore to ensure proper storage, but low enough under manufacturing conditions to flow appropriately and thus facilitate the shaping of the piece [34]. The suspension viscosity is in the range $\sim 32-35$ mPa·s at a shear rate of 1000 s $^{-1}$, and the slight difference between the suspensions prepared is simply that those without sonication and sonicated for 1 min exhibit some thixotropy (~ 1160 and 660 Pa·s $^{-1}$, respectively) while the one sonicated for 2 min exhibits very little rheopexy (210 Pa·s $^{-1}$). In terms of processing this latter is very desirable because these low thixotropy or rheopexy values indicate that there is no formation or destruction of structures in the suspension. The rheological behaviour of the $ZrC-Co$ suspensions is somewhat different to that of the B_4C-Co suspensions. Thus, it can be seen in Fig. 7B that, like the B_4C-Co suspensions, they all exhibit the desirable shear-thinning rheological behaviour, but in all cases with rheopexy (in the range $\sim 2450-2990$ Pa·s $^{-1}$). Also, there is now a greater difference of viscosity between the suspensions prepared without (~ 40 mPa·s) and with sonication (for 1, 2, or 3 min), these latter three being almost equally viscous (in the range $\sim 29-31$ mPa·s). Lastly, the

comparison between the B₄C–Co and ZrC–Co suspensions indicates that they have similar viscosity for the same preparation condition (difference less than 6 mPa·s), although the former have flow curves with less hysteresis loop. This is most likely attributable to the larger size of the ZrC particles (B₄C with submicrometre size, but ZrC with micrometre size) favouring the formation of structures in the suspension. Another interesting difference between the B₄C–Co and ZrC–Co suspensions is that the flow curves of the former shows evidence of a yield point because the shear stress increases first very abruptly and later gradually with increasing shear rate, which is something not observed in the flow curves of the latter in which the shear stress always increases progressively. Therefore, the B₄C–Co suspensions exhibit very shear-thinning rheological behaviour (i.e., plastic behaviour), whereas the ZrC–Co suspensions exhibit only shear-thinning rheological behaviour (i.e., pseudo-plastic behaviour).

The B₄C–Co and of ZrC–Co suspensions prepared here all have a rheological behaviour suitable for the near-net shaping of the corresponding compacts by slip casting. Their 30 vol.% solids loading is also entirely adequate because in slip casting both shaping and consolidation occur simultaneously since the driving force for the wall formation is the filtration through a permeable mould, so that the solids loading is not as critical as in other direct consolidation methods in which the drying stage is performed after the consolidation stage. Thus for example, the B₄C–Co and ZrC–Co suspensions sonicated for 2 and 3 min, respectively, were slip cast on plaster moulds with different shapes. Fig. 8 shows photographs of some resulting green compacts once dried in air at room temperature for 48 h. Apparently the green compacts did not shrink laterally during drying, which is because water elimination occurs by filtration (i.e., drainage assisted by capillary forces). They are robust and have good handling and storage characteristics. Also, these green compacts of B₄C–Co and ZrC–Co have high relative densities of ~58 and 52%,

respectively, achieved without the application of external pressure. The B_4C -Co compacts have higher green-body densification, which is due to the smaller size of the B_4C particles favouring tighter packaging. Fig. 9 shows representative low- and high-magnification SEM images of the fracture surface of the B_4C -Co and ZrC-Co compacts in their as-cast condition. It can be seen that the green compacts exhibit a uniform microstructure, with no evidence of macrodefects or density gradients (Figs. 9A and 9C). Moreover, the particles are well packed, and the aggregates of metal particles are homogeneously distributed among the ceramic particles (Figs. 9B and 9D), especially in the case of the ZrC-Co compacts due to the larger size of the ZrC particles. The metal aggregates in the B_4C -Co compacts can have B_4C particles entrapped, which is due to the smaller size of the B_4C particles.

Fig. 10 shows representative low- and high-magnification SEM images, and Fig. 11 the XRD patterns, of the B_4C -Co and ZrC-Co composites obtained by pressureless sintering at 1700 °C for 2 h in Ar atmosphere. It can be seen in Figs. 10A and 10B that the B_4C -Co composite is fully dense, that it has shrunk preserving the shape (~18% lateral shrinkage), and that its microstructure consists of submicrometre grains embedded in an intergranular phase plus large pockets of an additional secondary phase. Clearly, densification occurs by liquid-phase sintering. Also, relative to the un-sintered composite, the XRD pattern in Fig. 11A indicates that the H_3BO_3 and C peaks disappeared, that the B_4C peaks modified their relative intensities and shifted toward larger diffraction angles, and that CoB and Co_2B peaks appeared (with some shifting), this latter compound with marked crystallographic texture. The EDS compositional analyses revealed that the Co:B ratio in the pockets is close to 2, with the presence of C and O atoms. With all this information, it is then reasonable to infer that basically the grains are C-richer B_4C , that the intergranular phase is CoB, and that the secondary-phase pockets are textured single-

crystal Co_2B with C and O solutes. This reflects the occurrence of reactions between B_4C (and H_3BO_3 as well) and Co during pressureless sintering, which has also been observed during the spark-plasma sintering of B_4C with Ti-Al intermetallics [35]. Moreover, the formation of the Co_2B pockets is consistent with the existence in the compacts of Co micrometre aggregates with B_4C entrapped. Lastly, another interesting conclusion is that the sintered composite is not a cermet, but a multi-component ceramic because under the present sintering conditions Co indeed acted as a reactive and transient liquid phase.

Additionally, it can be seen in Figs. 10C and 10D that the ZrC–Co composite is fully dense as well, and that it has shrunk preserving the shape (~15% lateral shrinkage). According to its XRD pattern in Fig. 11B, this composite only contains ZrC and Co. Therefore, it is a cermet with the typical microstructure of the cemented carbides, in particular with ZrC grains homogeneously embedded in a Co intergranular phase which acted as a persistent liquid phase. This is different from what has been observed earlier in ZrC–Mo composites, in which Mo acted as a reactive and transient liquid phase, thus with extensive formation of MoC_2 occurring [21,22]. Also, as expected for densification by conventional liquid-phase sintering, the microstructure in Figs. 10C and 10D shows evident signs of grain shape accommodation and grain growth by solution-precipitation.

Fig. 12 shows representative SEM images of the fracture surface of the B_4C –Co and ZrC–Co composites obtained by pressureless sintering at 1700 °C for 2 h in Ar atmosphere. It is evident that there is a vast difference in the fracture mode of the two composites. Specifically, the ZrC–Co cermet exhibits many cleavage steps, which is not the case for the B_4C –Co composite. Thus, while the ZrC–Co cermet has broken in transgranular fracture mode, the B_4C –Co composite did so in intergranular fracture mode. This difference is mostly attributable to the

coarseness of the microstructure promoting transgranular fracture in the ZrC–Co cermet. Indeed, earlier observations support this explanation, showing that the ZrC [6,11] ceramics and the ZrC–MoSi₂ [8] and ZrC–SiC [13] composites, as well as other ceramics [36], also break transgranularly when their grain size is coarse, but intergranularly, or intergranularly plus transgranularly, when their grain size is sufficiently fine. Nonetheless, composition also plays a role in the fracture mode because the B₄C [12,16,18] ceramics and the B₄C–SiC [14] and B₄C–SiC–C [14] composites essentially break transgranularly, regardless of the grain size.

To conclude, it is convenient to discuss some final considerations for the near-net shape manufacture of these, and similar, carbide-metal composites by slip casting and pressureless sintering. Specifically, the pressureless sintering cycle (i.e., heating/cooling ramps, temperature, soaking time, and atmosphere) could be customized to thus control the particle sizes in the final microstructure. Customization of the sintering cycle might also force the densification to occur with a persistent liquid phase or with a reactive and transient liquid phase, thus enabling the controlled fabrication of either ceramic composites or cermets. The ceramic:metal ratio in the powder mixtures could also be customized with a view to controlling the volume fraction of intergranular phase (persistent liquid-phase sintering) or secondary phases (reactive and transient liquid-phase sintering), and perhaps the type of microstructure as well (i.e., disconnected-liquid microstructure or connected-liquid microstructure). Beyond compositional aspects (exact type of carbide and metal used), it seems clear that it is potentially possible to fabricate an ample variety of different carbide-metal composites with microstructures, and therefore mechanical properties, tailored for engineering applications. Exploration of this variety of possibilities remains pending, however, for future work. Future research efforts should also be aimed at avoiding the undesirable formation of secondary-phase pockets in the final microstructure when densification

takes places by reactive and transient liquid-phase sintering. This requires, in general, a greater optimization of the ceramic–metal dispersion, using colloidal processing techniques or other possible alternatives. Nonetheless, it also seems that the secondary-phase pockets will be unlikely to form if the ceramic starting powder is coarser than the metal starting powder, or if the volume fraction of metal in the powder mixture is judiciously reduced. Investigation of these aspects also merits future work.

4. Conclusions

Near-net shaped B_4C –Co and ZrC –Co composites have been manufactured by slip casting from aqueous colloidal suspensions, and subsequent pressureless sintering. Based on the experimental results and analyses, the following conclusions can be drawn:

1. Concentrated suspensions of both B_4C –Co and ZrC –Co can be prepared by aqueous colloidal processing having a shear-thinning rheological behaviour suitable for slip casting. Advisable preparation conditions for such suspensions are high pHs (i.e., pH ~ 10), moderate deflocculant contents (i.e., ~ 1 wt.% for both the ceramic and the metal), and short sonication times (i.e., 1–2 min). The most critical preparation aspect is the correct dispersion of the metal particles.
2. Robust, highly-dense B_4C –Co and ZrC –Co compacts can be shaped by slip casting. These green bodies have good handling and storage characteristics, and possess a uniform microstructure without macrodefects or density gradients.
3. The B_4C –Co and ZrC –Co compacts can be fully densified by pressureless liquid-phase sintering, preserving their shape. Under the present heat-treatment conditions (i.e., 1700 °C for 2 h in Ar atmosphere) the former densify by reactive and transient liquid-phase

sintering resulting in multi-component ceramics, and the latter by persistent liquid-phase sintering resulting in cermets.

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Figure Captions

Figure 1. Representative low-magnification SEM images of the (A) B₄C, (B) ZrC, and (C) Co particles in the starting powders.

Figure 2. Representative high-magnification SEM images of the (A) B₄C, (B) ZrC, and (C) Co particles in the starting powders.

Figure 3. XRD patterns of the B₄C, ZrC, and Co starting powders. Peak assignments are included.

Figure 4. High-resolution XPS spectra of the B₄C, ZrC, and Co starting powders. Core-levels monitored in each case are indicated. Peak assignments are included too.

Figure 5. Evolution of the zeta potential of the dilute suspensions of the B₄C, ZrC, and Co starting powders as a function of pH.

Figure 6. Evolution of the zeta potential of the dilute suspensions of the B₄C, ZrC, and Co starting powders as a function of the deflocculant content (PKV for both B₄C and ZrC, and PAA for Co) at the natural pHs.

Figure 7. Flow curves of the concentrated suspensions of (A) 80 vol.% B₄C+20 vol.% Co and (B) 80 vol.% ZrC+20 vol.% Co prepared to a total solids loading of 30 vol.%, without and with

different sonication times. Sonication was not prolonged further once the flow curve worsened. The arrows indicate the uploading and downloading stretches of the flow curves.

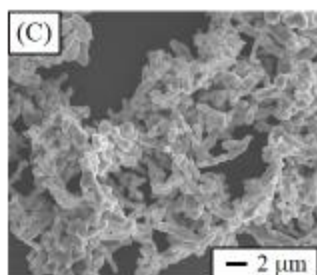
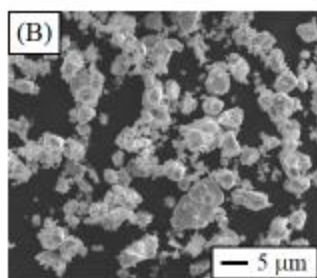
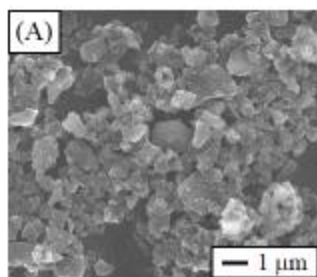
Figure 8. Photographs of various B_4C-Co and $ZrC-Co$ compacts obtained by slip casting. Photographs were taken after drying in air at room temperature for 48 h.

Figure 9. Representative SEM images of the fracture surface of the B_4C-Co compact at (A) low- and (B) high-magnification as well as of the $ZrC-Co$ compact at (C) low- and (D) high-magnification. The insets in (B) and (D) are SEM images taken with backscattered electrons.

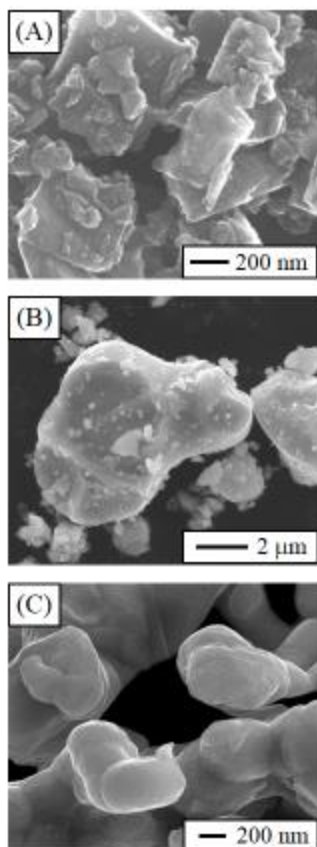
Figure 10. Representative SEM images of the polished surface of the B_4C-Co composite at (A) low- (A) and (B) high-magnification as well as of the $ZrC-Co$ composite at (C) low- and (D) high-magnification. The two composites were obtained by pressureless sintering at $1700\text{ }^\circ\text{C}$ for 2 h in Ar atmosphere. The insets in (A) and (C) compare the as-cast and sintered bodies.

Figure 11. XRD patterns of the (A) B_4C-Co and (B) $ZrC-Co$ composites obtained by pressureless sintering at $1700\text{ }^\circ\text{C}$ for 2 h in Ar atmosphere. Peak assignments are included, in (A) only for relevant peaks up to $\sim 52.5\text{ }^\circ 2\theta$. The intensity scale in (A) is dominated by the atomic scattering factor of Co, and the texture of Co_2B .

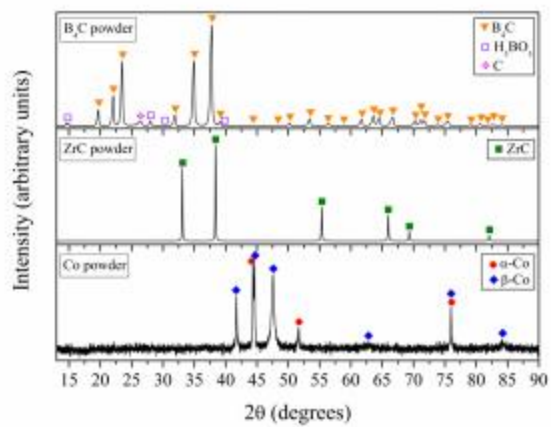
Figure 12. Representative SEM images of the fracture surface of the broken (A) B_4C-Co and (B) $ZrC-Co$ compacts obtained by pressureless sintering at $1700\text{ }^\circ\text{C}$ for 2 h in Ar atmosphere.



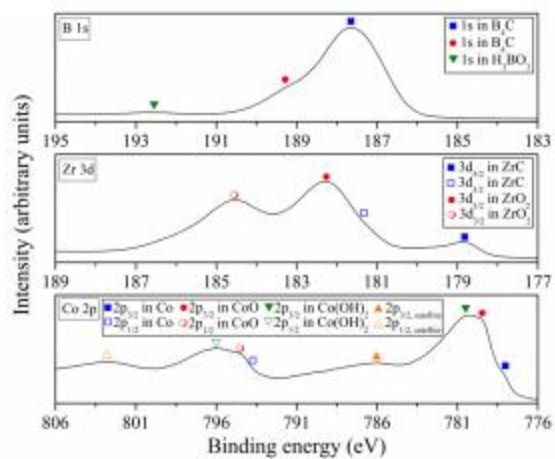
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Figure 1



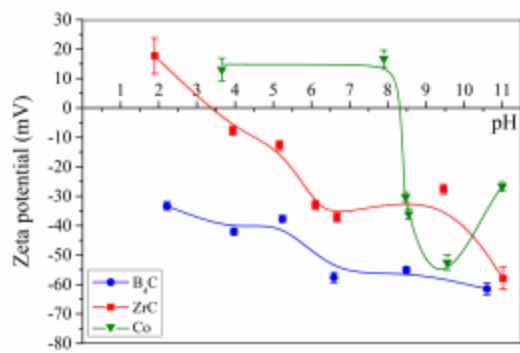
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Figure 2



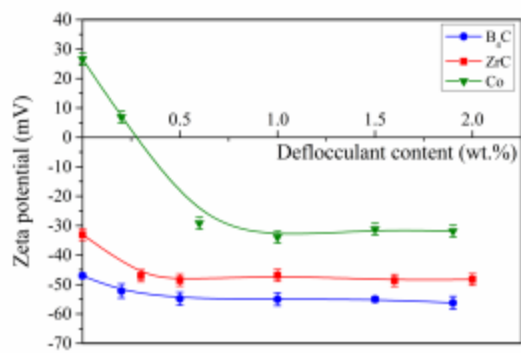
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Figure 3



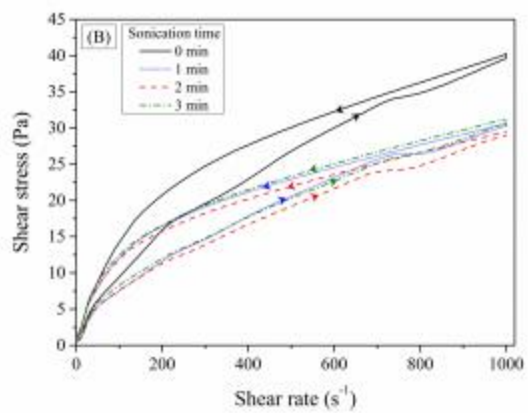
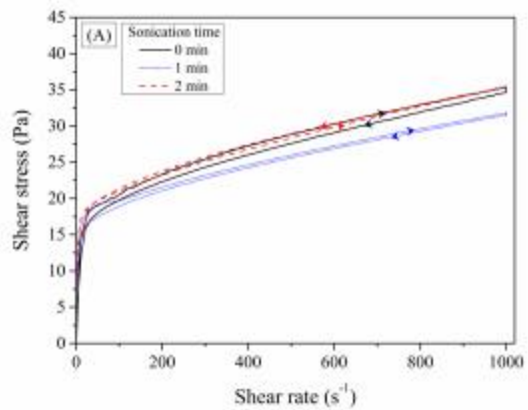
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Figure 4



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Figure 5



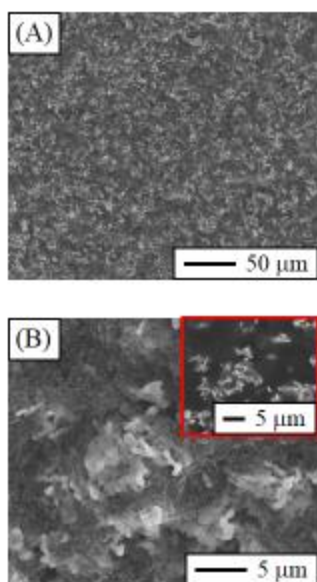
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Figure 6



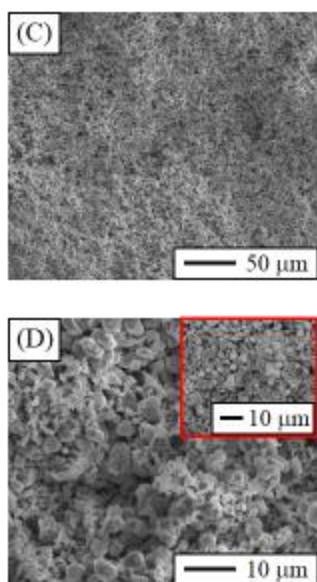
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Figure 7



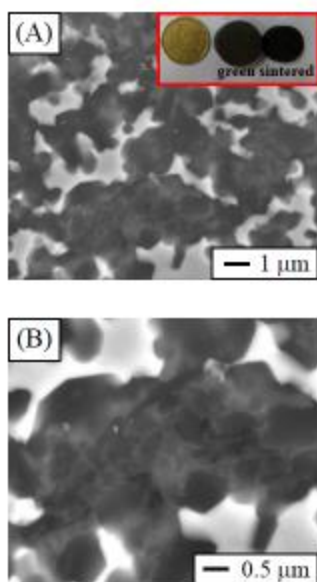
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Figure 8



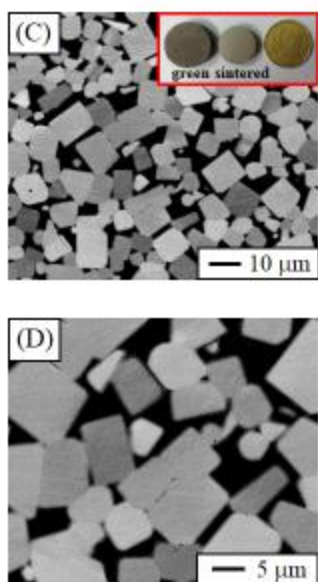
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Figure 9



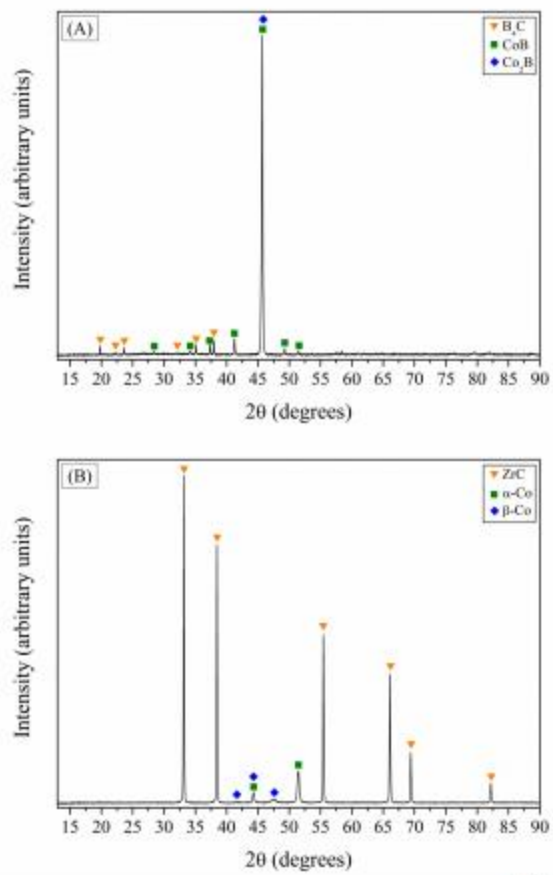
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Figure 9



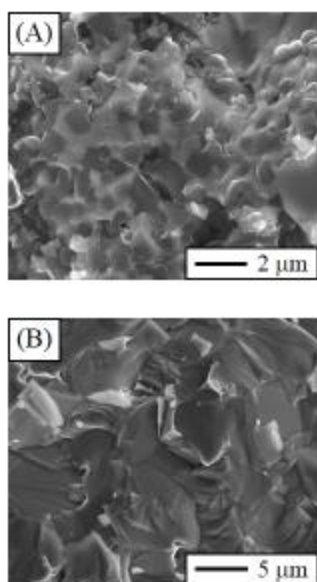
Ortiz *et al.*
Figure 10



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Figure 10



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Figure 11



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Figure 12