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Fabricating toughened super-hard B₄C composites at lower temperature by transient liquid-phase assisted spark plasma sintering with MoSi₂ additives

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Abstract

Toughened, super-hard B₄C triplex-particulate composites were densified by spark plasma sintering with MoSi₂ additives (5, 10, and 15 vol.%) at temperatures in the range 1750-1850 °C at which the reference monolithic B₄C ceramics are porous. It is proved that MoSi₂ is a reactive sintering additive that promotes densification by transient liquid-phase sintering, thus yielding fully-dense B₄C-MoB₂-SiC composites at relatively lower temperatures. Specifically, the MoSi₂ first reacts at moderate temperatures (<1150 °C) with part of B₄C to form MoB₂, SiC, and Si. This last is a transient component that eventually melts (at ~1400 °C), contributing to densification by liquid-phase sintering, and then (at 1500-1700 °C) reacts with free C present in the B₄C starting powders to form more SiC, after which densification continues by solid-state sintering. It is found that these B₄C-MoB₂-SiC composites are super-hard (~30 GPa), tough (~3-4 MPa·m^{1/2}), and fine-grained, a combination that renders them very appealing for structural applications. Finally, research opportunities are discussed for the future microstructural design of a novel family of toughened, ultra-hard/super-hard multi-particulate composites based on B₄C plus refractory borides and carbides.

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Keywords: B₄C; super-hard ceramic composites; spark-plasma sintering; sintering additives;
mechanical properties.

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1. Introduction

There is growing demand for super-hard monolithic ceramics and ceramic composites for use in a myriad of structural applications requiring great resistance to contact damage, both without and with friction. Examples of such applications are armours for the protection of personnel and vehicles against ballistic projectiles, components for industrial machinery (dies, bearings, seals, valves, nozzles, etc.), and machining/cutting tools (drills, end mills, reamers, blades, wheels, etc.), to name but a few. Unfortunately however, the current list of super-hard ceramics (i.e., with hardness ideally above 30 GPa under Vickers indentation) is very short [1], and is limited basically to diamond [2-4], B₄C [5,6], and a few nitrides (i.e., BN, C₃N₄, and BC₂N) [2-3,7-9], borides (i.e., OsB₂, ReB₂, and WB₂) [10,11], and oxides (i.e., B₆O) [12,13]. Of these, B₄C receives most attention because diamond is prohibitively expensive and the rest are very exotic. In addition, B₄C is ultra-lightweight (i.e., ~2.52 g/cm³ density), which is always very desirable in any engineering application but essential in many of them (armour, rotating/moving tribocomponents, etc.). Moreover, B₄C is ultra-refractory (i.e., ~2450 °C melting point) too, and therefore appealing for high-temperature engineering applications.

The bane of B₄C is, as for the rest of the ultra-refractory ceramics, its poor sinterability [14]. This is due to the strong covalent bonding and low self-diffusion coefficients imposing serious kinetic restrictions, and to the oxide impurities promoting coarsening over densification [15-17]. It is then quite common to circumvent this problem by using densification aids [5,14]. Which one is chosen is not a trivial decision, however, because they all have their own advantages and drawbacks. Overall, solid-state-sintered B₄C ceramics are harder and more refractory, but are brittle and have coarse-grained microstructures undesirable in terms of strength and wear resistance. Liquid-phase-sintered B₄C ceramics have fine-grained

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4 microstructures, and are stronger and tougher, but the residual intergranular phase limits their
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6 hardness and refractoriness. This dichotomy could be avoided by using so-called transient liquid-
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8 phase sintering,^I which exploits the benefits of both liquid-phase sintering and solid-state
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10 sintering. This enticing approach has recently been applied to B₄C, in particular using metallic
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12 and intermetallic sintering additives [18,19].
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16 Given its ultra-high refractoriness, sintering additives suitable for B₄C could be those
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18 already used successfully for densifying ultra-high-temperature ceramics (UHTCs)^{II} [20]. Of
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20 these, there stands out MoSi₂, which is regularly used for carbide-based UHTCs [21-29]. Its
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22 great appeal is that it also strengthens and toughens UHTCs, and provides them with superior
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24 oxidation resistance. Unsurprisingly, MoSi₂ has already been used to fabricate super-coarse-
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26 grained, dense B₄C composites by hot-pressing [30]. Interestingly, the hot-pressing conditions
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28 used were relatively smooth (for B₄C), and resulted in super-hard, tougher B₄C composites. That
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30 study raised various questions of great interest, namely: (i) can MoSi₂ also be used to densify
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32 fine-grained B₄C composites by ultra-fast sintering (spark plasma sintering (SPS))^{III} in
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34 particular?; (ii) if so, what is the mechanism of this sintering?; and (iii) how does the MoSi₂
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36 addition affect the densification kinetics and the mechanical properties of these composites? In
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38 the present study we shall answer these questions, with the expectation that the scientific
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48 ^I In transient liquid-phase sintering, the sintering aids first melt at an early stage of sintering, and then that liquid
49 reacts with the solid, disappearing as sintering proceeds further.

50 ^{II} UHTCs are a family of super-refractory ceramics (ideally with melting points above 3000 °C) based on borides,
51 carbides, and nitrides of the early-transition metals.

52 ^{III} SPS is currently the best exponent of ultra-fast sintering techniques. It uses much shorter densification cycles than
53 either hot-pressing or conventional pressureless sintering (i.e., heating/cooling ramps are one order of magnitude
54 faster (hundreds vs tens of °C/min), and holding times are one order of magnitude shorter (min vs h)). **SPS**
55 essentially lies in the repeated application of high energy, low voltage, pulsed direct electrical current under
56 mechanical pressure, and is then an electric current-assisted sintering technique of low electric field (only a few
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4 knowledge gained may contribute to guiding the future fabrication of novel B₄C composites for
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6 engineering applications.
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10 11 **2. Experimental Procedure** 12 13

14 The starting powders, obtained from a commercial source (H.C. Starck, Germany) and
15 used in their as-received condition, were B₄C (Grade HD20; d_{50} ~0.3-0.6 μ m) and MoSi₂ (Grade
16 B; d_{50} ~3.5-5 μ m). According to the manufacturer's specifications, the B₄C powder has a B:C
17 atom ratio of 3.8, and the following mass fractions (in %): C 22.0, N 0.3, O 1.8, Si 0.09, Al 0.01.
18 The corresponding data for the MoSi₂ are: Si 36.7, C 0.06, O 0.52, and N 0.02. Consequently, the
19 B₄C powder contains free C, and the MoSi₂ powder does not contain residual Si. Three powder
20 batches were prepared having B₄C-MoSi₂ relative compositions of 95-5 vol.%, 90-10 vol.%, and
21 85-15 vol.%. To this end, the B₄C and MoSi₂ powders were first combined in appropriate
22 proportions in wt.%, then mixed/homogenized by wet ball-milling in methanol for 24 h, dried on
23 a hot-plate while stirring, and finally de-agglomerated by crushing.
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38 Next, the as-received B₄C powder and these three B₄C-MoSi₂ powder mixtures were
39 loaded into graphite dies (2-cm diameter) lined with graphite foil and covered by graphite
40 blankets, and were SPS-ed (HP-D-10, FCT Systeme GmbH, Germany) in dynamic vacuum. The
41 common SPS conditions were 100 °C/min heating, 50 MPa pressure, and 5 min soaking, whereas
42 the target temperature was varied in the range 1700-1850 °C. The SPS furnace is equipped with
43 an axial optical pyrometer that measures the temperature on a hole machined across the upper
44 graphite punch, and therefore these temperatures are not directly comparable with those
45 measured by others in SPS furnaces equipped with radial optical pyrometers focused on the
46 graphite die. Except for the ultra-fast sintering cycle, these SPS conditions are similar to those
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4 used in the earlier hot-pressing study (i.e., 1700-1900 °C for 2 h under 30-50 MPa). Table 1 lists
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6 the particular conditions used in each case. Given that the expectation is that MoSi₂ would
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8 facilitate the densification, lower target SPS temperatures were tested for the powder mixtures
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10 with 10 or 15 vol.% MoSi₂. The as-received B₄C powder is the baseline reference for
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12 comparison, and therefore was consolidated at all target SPS temperatures. During each SPS
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14 cycle the shrinkage curve was logged, then corrected for the expansion of the graphite parts, and
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16 converted to a densification curve for subsequent analysis.
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21 The resulting monolithic ceramics and ceramic composites were ground and polished to a
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23 1-μm finish, and also intentionally broken. They were characterized microstructurally by X-ray
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25 diffractometry (XRD; D8 Advance, Bruker AXS, Germany), scanning electron microscopy
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27 (SEM; S-3600N, Hitachi, Japan or Quanta 3D, FEI, The Netherlands), and energy-dispersive X-
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29 ray spectrometry (EDS; Xflash Detector 3001, Röntec GmbH, Germany). The mechanical
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31 characterization (hardness and toughness) was done by Vickers indentation tests (MV-1,
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33 Matsuzawa, Japan) on the polished surface of selected SPS-ed samples.
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38 Model SPS experiments were also conducted to investigate in detail the microstructural
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40 development of the B₄C composites. To this end, the B₄C powder with 15 vol.% MoSi₂ was SPS-
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42 ed at temperatures in the range 1150-1500 °C. Modified SPS cycles (100 °C/min heating, 50
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44 MPa pressure, dynamic-vacuum atmosphere) without soaking time were used, in which just
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46 immediately after reaching the desired target SPS temperature the electrical power of the SPS
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48 furnace was intentionally shut off and the pressure was deliberately released. The resulting B₄C
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50 composites were ground, and were all characterized microstructurally by XRD and some
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52 selectively by SEM.
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3. Results and Discussion

Figures 1-4 show representative SEM images of all the materials fabricated, and Table 1 gives the degrees of densification (qualitative information) inferred from the extensive microstructural observations. It can be seen that the degree of densification increases with both the target SPS temperature and the MoSi₂ proportion. The former was expected (diffusion is a thermally-activated process obeying an Arrhenius-type law [31]), while the latter proves that MoSi₂ is a suitable SPS additive for B₄C. Note that the differences of densification are attributable to effects of temperature and composition, and not to electrical effects because during SPS (i) the voltages applied were very low (~ 6 V) and nearly the same in all cases (difference of less than ~0.22 V), which would exclude field effects, and (ii) the amperages applied were also, for a given temperature, nearly the same regardless of the MoSi₂ content used as sintering additive (difference of less than ~1.4%), which would rule out current effects. Importantly, it can be seen that the monolithic ceramics are all to a greater or lesser extent porous, although the one SPS-ed at 1850 °C is near fully-dense. On the contrary, all the composites eventually densified completely, in particular, at target SPS temperatures of 1850 (5 vol.% MoSi₂), 1800 (10 vol.% MoSi₂), and 1750 °C (15 vol.% MoSi₂). In addition, at ~50 °C below these temperatures they reached near full-densification. It can also be seen in Figs. 2-4 that the composites contain Mo/Si-rich second phases over their entire microstructure. This was further confirmed by EDS compositional maps, an example of which is shown in Fig. 5. There are however pockets of Mo/Si-rich second phases, which indicates that their distribution is not completely uniform. This is attributable to the difference between the particle sizes of the B₄C (~0.3-0.6 μm) and MoSi₂ (~3.5-5 μm) powders, and therefore future processing efforts are needed to gain in microstructural homogeneity (for example introducing a step of ball-

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4 milling/co-ball milling, using colloidal processing, or simply utilizing starting powders with
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6 more similar granulometric features).

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9 Figure 6 shows the SPS densification curves as a function of temperature for the B₄C
10 powder with different MoSi₂ contents (i.e., 0, 5, 10, and 15 vol.%). Overall, these curves have
11 the three-stretch shape observed for other refractory carbides and borides [32-39]. Specifically,
12 there are two stretches during the non-isothermal heating, the first of moderate densification
13 attributable to mechanical powder compaction and the second of accelerated densification
14 attributable to diffusion inducing mass transport. Lastly, there is a third stretch of gradual
15 densification during the isothermal heating. Table 2 lists the ultimate densifications reached,
16 which are not necessarily 100%.^{IV} The SPS densification curves thus confirm that B₄C
17 sinterability increases with the MoSi₂ proportion because the densification kinetics speeds up.
18 Thus for example, the intermediate sintering regime [40]^V was reached at ever lower
19 temperatures during the non-isothermal heating, in particular at ~1770 (0 vol.% MoSi₂), 1685 (5
20 vol.% MoSi₂), 1600 (10 vol.% MoSi₂), and 1550 °C (15 vol.% MoSi₂). The final sintering
21 regime^V was also reached earlier with increasing MoSi₂ proportion, in particular after ~130 s at
22 1800 °C or 80 s at 1850 °C for 0 vol.% MoSi₂, after ~50 s at 1800 °C or 10 s at 1850 °C for 5
23 vol.% MoSi₂, at ~1790 °C for 10 vol.% MoSi₂, and at ~1700 °C for 15 vol.% MoSi₂.

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26 The SPS densification curves are also a reflection of the MoSi₂ addition altering the
27 sintering mechanism of B₄C. This is clearer however in the shrinkage rate curves, an example of
28 which is shown in Fig. 7 for the B₄C powder without and with 15 vol.% MoSi₂. The curve of the
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54 ^{IV} When this happens, prolonging the SPS cycle is of little or no effectiveness because densification tends towards a
55 certain limiting value. This is more evident when the SPS densification curves are plotted as a function of time.

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^V The intermediate sintering regime is that during which the pore distribution transforms from open porosity to
closed porosity, and is typically taken to begin at ~70% relative density. The final sintering regime is that during
which the closed porosity is eliminated, and is normally taken to start at ~90% relative density.

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4 former reflects a partial densification by solid-state sintering because the shrinkage rate first
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6 remains invariably low up to the sintering onset temperature (~ 1420 °C), and then increases
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8 gradually with temperature. But the curve of the latter exhibits a distinctive peak of accelerated
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10 shrinkage rate at ~ 1390 °C, and sintering started long before (at ~ 1300 °C). This peak is
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12 indicative of densification by liquid-phase sintering, and in particular is associated with the
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14 sudden formation of a liquid that spreads rapidly between grains, filling the pores. The rest of the
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16 curve follows the expected temperature dependence, with the shrinkage rate increasing gradually
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18 up to the temperature of most active sintering, and then decreasing [40].
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24 Figure 8 compares, by way of example, representative SEM images of the microstructure
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26 of both the monolithic ceramic and the composite with 15 vol.% MoSi₂ SPS-ed at 1800 °C. It
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28 can be seen that while the former is porous and fine-grained, the latter is fully dense and also has
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30 a more complex triplex-particulate microstructure. According to EDS analyses shown in Fig. 9,
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32 the major phase is rich in B and C, whereas the other grains are rich in either Si and C or Mo and
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34 B. Clusters of these grains were also observed in the lower-magnification SEM observations,
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36 which is consistent with the earlier observations (Figs. 2-4).
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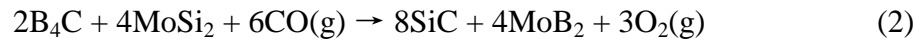
41 Figure 10 shows the XRD patterns of all the materials fabricated. It can be seen that the
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43 monolithic ceramics contain only B₄C and some residual graphite (an impurity already present in
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45 the B₄C starting powder [19,41,42]). The composites, however, contain B₄C plus various
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47 additional compounds, none of which is MoSi₂. These are SiC (cubic polytype (β -SiC)) and
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49 MoB₂ (trigonal and hexagonal polymorphs (β -MoB₂ and α -MoB₂, respectively)). This then is
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51 consistent with the EDS analyses. Of the two MoB₂ polymorphs, β -MoB₂ predominates for SPS
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53 temperatures up to ~ 1750 °C, whereas α -MoB₂ does so thenceforward. Previous work has also
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55 shown that β -MoB₂ and α -MoB₂ are, respectively, low- and high-temperature polymorphs [43].
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4 There is also some, but less, residual graphite in the composites fabricated with 5 vol.% MoSi₂,
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6 but not in those made with 10 and 15 vol.% MoSi₂. The present composites fabricated by SPS
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8 and their counterparts fabricated earlier by hot-pressing thus have the same phase composition,
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10 because the trigonal-Mo₂B₅ compound that has been reported for the latter [30] is actually β-
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12 MoB₂ [44,45].^{VI}
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16 Thus, the above results reveal that the whole of the MoSi₂ reacted with part of the B₄C
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18 during SPS. A possible global reaction path would be:
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22 where the C reactant comes from the graphite impurities in the B₄C starting powders and/or is
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24 taken up during SPS (diffusion from the graphite tools and/or foils). This reaction has already
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26 been proposed for the hot-pressed B₄C composites [30].^{VII} Another possible global reaction path
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28 would be:
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33 where the CO(g) reactant would be generated in-situ during SPS because the graphite tools/foils
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35 are heated under low oxygen partial pressure (due to the dynamic vacuum during SPS). There
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37 were however two experimental observations in favour of the occurrence of Reaction (1). Firstly,
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39 the B₄C-MoSi₂ powder mixtures did not emit more gas (attributable to O₂ release) than the B₄C
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41 powder. By way of example, Fig. 11 compares the vacuum levels registered for the B₄C powder
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43 without and with 15 vol.% MoSi₂ SPS-ed at 1700 °C, illustrating this observation. This is indeed
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54 ^{VI} It is now well established that what was formerly described as trigonal-Mo₂B₅ is actually trigonal-Mo₂B₄ (i.e., β-
55 MoB₂) because crystallographic studies have demonstrated that Mo₂B₅ is actually a non-stoichiometric compound
56 without the centring boron atoms in the puckered sheets of condensed cyclohexane-like boron chairs.

57 ^{VII} An analogous reaction (i.e., 5B₄C + 8MoSi₂ + 11C → 16SiC + 4Mo₂B₅) was also formulated to explain the (now
58 known to be wrong) formation of Mo₂B₅ (and concurrently of more SiC as well), but of course this reaction can
59 never take place because Mo₂B₅ is actually β-MoB₂.
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4 the result expected from Reaction (1), whereas Reaction (2) predicts the opposite result. And
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6 secondly, C is necessarily one of the reactants because the graphite impurities in the B₄C starting
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8 powders were consumed only when there was MoSi₂. This is the case of Reaction (1), but not of
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10 Reaction (2). Also note that Reaction (1) has been proved to be thermodynamically enabled in
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12 the range of SPS temperatures used [30]. Having identified that the overall reaction path is
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14 Reaction (1), one can then calculate that the composites fabricated with 5, 10, and 15 vol.%
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16 MoSi₂ have a B₄C-MoB₂-SiC composition in wt.% of 85.9-8.4-5.7, 73.7-15.6-10.7, and 63.2-
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18 21.9-14.9, respectively.
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24 The conclusion is that MoSi₂ acted as a reactive sintering additive, not as a permanent
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26 sintering additive. MoSi₂ is not the first transient sintering additive used to densify B₄C with in-
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28 situ formation of refractory carbides and borides [18,19], but it can offer certain advantages over
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30 the others. In particular, it will also notably increase the oxidation resistance of B₄C at high
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32 temperatures, which otherwise would undergo non-protective passive or active oxidations due to
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34 the formation of a liquid B₂O₃ scale or gaseous B₂O₃ [46]. Thus, the numerous oxidation studies
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36 carried out on boride-based UHTCs indicate that the SiC grains in the microstructure of the
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38 present composites will oxidize to SiO₂, which will react with the B₂O₃ resulting from the
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40 oxidation of the B₄C grains, thus developing protective borosilicate-glass scales [47-49]. This
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42 might allow the use of these composites at high temperatures in oxidizing atmospheres, which
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44 are conditions that the monolithic B₄C ceramics normally do not withstand. Indeed, a recent
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46 study [50] has confirmed experimentally that the B₄C-ZrB₂-SiC triplex-particulate composites
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48 have greater oxidation resistance than the monolithic B₄C ceramics.
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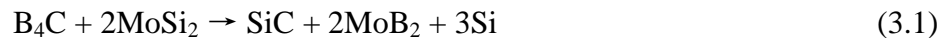
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54 To identify the sintering mechanism of these B₄C-MoB₂-SiC composites, additional
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56 model SPS experiments were conducted on the B₄C powder with 15 vol.% MoSi₂. They
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4 consisted in interrupting deliberately the SPS cycle at preselected temperatures in the range
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6 1150-1500 °C by shutting off the electrical power of the furnace and releasing the pressure.
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8 Owing to the ultra-fast cooling to room-temperature of the SPS furnace, these samples provide
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10 “frozen snapshots” of the microstructural development of the composites. **Figure 12** shows the
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12 XRD patterns of some selected composites resulting from these interrupted SPS experiments. It
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14 can be seen that the composite SPS-ed at 1150 °C contains B₄C, H₃BO₃ (typical impurity in this
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16 and in other B₄C starting powders [19,41,42,51,52]),^{VIII} C (impurity in the B₄C starting powder
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18 [19,41,42]), and MoSi₂, plus β-MoB₂, SiC, and Si in minor amounts. The reaction of in-situ
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20 formation of refractory carbides and borides thus began slightly below 1150 °C, with Si being
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22 another reaction product. The composites SPS-ed between 1150 and 1350 °C had the same phase
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24 inventory, but with less H₃BO₃ and MoSi₂ and more β-MoB₂, SiC, and Si as the SPS temperature
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26 increased. This was also the trend observed for the composites SPS-ed at 1400 and 1450 °C, but
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28 with the difference that they no longer contain crystalline H₃BO₃, but nevertheless some α-
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30 MoB₂. Lastly, the composite SPS-ed at 1500 °C is formed by B₄C, C, SiC, β-MoB₂, α-MoB₂,
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32 and Si, but not MoSi₂. Consequently, B₄C finished consuming all the MoSi₂ at a temperature
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34 between 1450 and 1500 °C. According to their XRD patterns (Figs. 10C-D), the B₄C composites
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36 SPS-ed at higher temperatures (i.e., ≥1700 °C) have only B₄C, SiC, and MoB₂, reflecting that Si
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38 and C eventually consume each other at a temperature between 1500 and 1700 °C.
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40 Concomitantly, the composites become richer in α-MoB₂ at the expense of β-MoB₂. **Figure 13**
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42 shows representative SEM images of the composites resulting from the model SPS experiments
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44 interrupted at 1250 and 1400 °C. As expected, the two composites are still extremely porous. The
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46 relevant microstructural observation is that, whereas the former exhibits only particles in a very
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58 ^{VIII} H₃BO₃ will dehydrate into B₂O₃ during heating (2H₃BO₃→B₂O₃+3H₂O(g)), and will re-hydrate again during
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4 early adhesion state (incipient sinter bond), the latter already exhibits dense regions with a
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6 molten phase embedding the particles. According to the EDS analyses (not shown), that phase is
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11 It is then clear that the formation of refractory carbides (i.e., SiC) and borides (i.e.,
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13 MoB₂) in-situ during SPS actually came accompanied by the formation of Si as an intermediate
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15 component, and consequently that Reaction (1) really occurred in the following two sub-steps:
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24 Thus, the MoSi₂ sintering additive reacts with B₄C to form MoB₂ (mostly β-MoB₂) and SiC, plus
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26 free Si (Reaction (3.1)). This first reaction starts slightly below 1150 °C, and is complete at
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28 ~1450-1500 °C (Fig. 12). Importantly, this reaction does not itself play a role in the densification
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30 because the SPS densification curves (Figs. 6B-D) and the shrinkage rate curves (Fig. 7) show no
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32 signs of enhanced densification below ~1300 °C. Eventually during the heating ramp the SPS
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34 temperature reaches ~1410 °C or more, and therefore free Si melts. This molten Si quickly
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36 spreads and wets efficiently the neighbouring particles (Figs. 13B-C), thus increasing the
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38 densification by both pore filling and particle rearrangement. This explains the accelerated
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40 shrinkage rate observed in the B₄C-MoSi₂ powder mixtures at ~1390 °C (Fig. 7).^{IX} Nonetheless,
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42 as seen in both the SPS densification curves (Figs. 6B-D) and the SEM observations (Figs. 13B-
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44 C), the amount of molten Si formed is insufficient to reach complete densification only by pore
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46 filling and particle rearrangement. As the SPS temperature increases further during the heating
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48 ramp, molten Si reacts with the C impurities of the B₄C starting powder producing more SiC
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50 (Reaction (3.2)). This second reaction occurs at a temperature between 1500 and 1700 °C (Figs.
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59 ^{IX} Note that the actual temperature within the graphite die is slightly greater than that measured by the optical
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61 pyrometer focused on the interior of the upper graphite punch.
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4 **10 and 12**). Si is therefore a transient component that does not jeopardize the refractoriness of
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6 these composites. Finally, once all the molten Si has been consumed forming SiC, during the rest
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8 of the SPS cycle the composites continue to densify gradually by conventional solid-state
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10 sintering at a lower rate until their full densification. Concomitantly, β -MoB₂ progressively
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12 transforms polymorphically to α -MoB₂. Consequently, despite its high melting point of
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14 ~1930 °C, MoSi₂ then promotes the lower-temperature densification of B₄C by transient liquid-
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16 phase sintering.
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22 Table 2 also lists the hardness of all the materials fabricated. It can be seen that it always
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24 increases to or above 25 GPa with increasing target SPS temperature, as expected for B₄C-based
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26 materials. Importantly, for the same given target SPS temperature, the composites are always
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28 harder than their reference monolithic ceramic. In addition, the higher the MoSi₂ content used as
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30 sintering additive, the greater the difference in hardness between the composite and the
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32 monolithic ceramic. To put these differences in perspective, note for example that the monolithic
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34 ceramic SPS-ed at 1750 °C is very soft (~14 GPa hardness), whereas the corresponding
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36 composite fabricated with 15 vol.% MoSi₂ is super-hard (~30 GPa hardness). The hardest of the
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38 monolithic ceramics (i.e., the one SPS-ed at 1850 °C) also reached an ultra-high hardness of ~25
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40 GPa, but this is far less than expected for super-hard B₄C (i.e., ~35 GPa). Besides the residual
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42 graphite that softens them, this essentially reflects that the hardness of monolithic B₄C ceramics
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44 depends largely on their degree of densification, decreasing exponentially with a rate constant of
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46 ~7 as the residual porosity increases [53]. Indeed, near fully-dense monolithic B₄C ceramics
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48 have a hardness in the range ~25-29 GPa [54,55]. The same is the case for the composites, whose
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50 hardness also increases gradually up to ~30 GPa as their residual porosity decreases. It is a little
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52 surprising, however, that the hardness of the fully-dense composites is independent of the MoSi₂
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4 content used as sintering additive (in the range 5-15 vol.%). This is attributable to the differences
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6 in their phase compositions. Firstly, the fully-dense composite fabricated with 5 vol.% MoSi₂
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8 contains some residual graphite (Fig. 10B; SPS-ed at 1850 °C), and is therefore softer. And
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10 secondly, the fully-dense composites fabricated with 5, 10, and 15 vol.% MoSi₂ contain,
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12 respectively, α-MoB₂ but no β-MoB₂ (Fig. 10B; SPS-ed at 1850 °C), much α-MoB₂ and a little
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14 β-MoB₂ (Fig. 10C; SPS-ed at 1800 °C), and a little α-MoB₂ and much β-MoB₂ (Fig. 10D; SPS-
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16 ed at 1750 °C), and α-MoB₂ is much softer than β-MoB₂ (~13 vs 19 GPa at 9.8 N) [43]. The net
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18 result is that increasing the MoSi₂ content in the powder mixtures enhances B₄C sinterability
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20 without any deterioration in the hardness of the resulting composites – at least up to 15 vol.%
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22 MoSi₂. Finally, one also observes in Table 2 that, for the same degree of densification, the
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24 monolithic ceramics are harder than the composites, which is because B₄C is harder than SiC, α-
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26 MoB₂, and β-MoB₂.
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34 There are more triplex-particulate composites based on B₄C that are ultra-hard/super-
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36 hard, namely (relative composition in volume fractions): 1B₄C-1TiB₂-1SiC (~33 GPa) [56],
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38 15B₄C-70TiB₂-15SiC (~30.9 GPa) [56], 1B₄C-1ZrB₂-1SiC (~28.9 GPa) [57], 90B₄C-8ZrB₂-2SiC
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40 (~29 GPa) [50], 80B₄C-16ZrB₂-4SiC (~30 GPa) [50], and 70B₄C-24ZrB₂-6SiC (~27 GPa) [50].
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42 Interestingly, they were all fabricated by conventional solid-state sintering using hot-pressing
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44 [50,56,57], not by transient liquid-phase assisted SPS as done here. This suggests that triplex-
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46 particulate composites based on B₄C, SiC and early-transition metal diborides are a novel family
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48 of potentially ultra-hard/super-hard materials, as could also be the case for other possible
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50 combinations of three or more carbides and borides.
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56 Table 2 also gives the fracture toughness of the fully-dense materials. Vickers testing is
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58 not applicable to measure the fracture toughness of porous brittle materials, and therefore the
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4 reference toughness will be the $\sim 1.9\text{-}2.3 \text{ MPa}\cdot\text{m}^{1/2}$ measured earlier by single edge V-notched
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6 beam tests for fine-grained fully-dense monolithic B_4C ceramics [53]. It can then be seen that the
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8 composites are increasingly tougher than the monolithic ceramic with increasing MoSi_2 content
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10 used as sintering additive. This attribute thus adds to the enhanced sinterability without hardness
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12 penalization. As with other B_4C composites [58], this greater fracture toughness is because the
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14 increasing residual stresses [59,60]^X induced by the thermoelastic mismatches between the B_4C ,
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16 MoB_2 , and SiC grains result in microstructures with increasingly weak interfaces that favour
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18 crack deflection and bridging.
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24 Let us now discuss some final considerations that arise from the present study that are of
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26 interest for the ceramics community in general, and for that of hard materials in particular.
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28 Firstly, while dense monolithic B_4C ceramics are very challenging to fabricate, on the contrary, it
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30 is relatively easy to fabricate, maybe even by conventional pressureless sintering, dense multi-
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32 particulate composites based on B_4C plus SiC and one or various refractory diborides (i.e.,
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34 MeB_2 , $\text{Me} = \text{Ti, Zr, Hf, V, Nb, Ta, Cr, Mo, W, and Re}$) from powder mixtures of B_4C with the
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36 corresponding early-transition metal disilicide(s) (i.e., MeSi_2). These composites are expected to
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38 possess a unique generic combination of ultra-high/super-high hardness, enhanced toughness,
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40 and superior oxidation resistance, not to mention other properties such as strength, wear
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42 resistance, etc., and therefore to hold great promise for structural applications. This could also be
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44 the case for the alternative multi-particulate composites fabricated (foreseeably at even lower
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46 temperatures) from powder mixtures of B_4C with aluminide(s) (i.e., ZrAl_3 , TiAl_3 , TaAl_3 , etc.)
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48 instead of silicide(s). Future research needs to be encouraged that is targeted at investigating the
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50 relationships between the processing, microstructure, and properties of such families of B_4C
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58 ^X Thermoelastic residual stress exhibits a parabolic functional dependence on the volume fraction of second phases.
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4 composites. Secondly, the fact that the molten phase formed eventually reacts with C presents a
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6 unique opportunity for further toughening these B_4C composites via innovative microstructural
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8 designs. Specifically, in principle it seems potentially possible to fabricate B_4C composites
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10 reinforced in-situ with an ample variety of hard carbide phases of high aspect ratio, and therefore
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12 superior toughening effect, simply by introducing intentionally in the starting powder mixtures
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14 carbon nanotubes, graphene nanoplatelets, reduced graphene oxide nanosheets, carbon fibres, or
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16 other high aspect ratio carbonaceous phases as the carbon source for Reaction (3.2). There is thus
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18 a requirement for the stimulation of new processing efforts to make these appealing B_4C
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20 composites a reality. And thirdly, there remain important scientific aspects to be clarified. For
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22 example, it is unclear whether, in the absence of a solid carbon source in the powder mixtures,
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24 the $CO(g)$ generated in the graphite furnaces would react with the molten phase to form more
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26 refractory carbide **if this reaction were thermodynamically possible**. If this reaction does not take
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28 place, it would then be important to determine whether the molten phase (L) remains in the
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30 microstructure as a residual secondary phase or, on the contrary, disappears either by entering in
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32 substitutional solid solution within the B_4C host to form $B_4(C,L)$ or by reacting with B_4C to form
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34 a C-deficient B_4C plus a refractory carbide. Indeed, new studies aimed at answering these
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36 questions are expected. In sum, the multi-particulate composites based on B_4C plus refractory
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38 borides and carbides fabricated in-situ at fairly smooth sintering conditions constitute a fertile
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40 ground for future research framed within one of the eight grand challenges (i.e., that of ceramics
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42 for extreme environments) with societal import identified recently by the ceramics community
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44 [61].
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4. Summary

Super-hard monolithic B_4C ceramics are very appealing for many engineering applications, but brittle and difficult to densify. To palliate these two weaknesses, here $MoSi_2$ additives were used to fabricate B_4C multi-particulate composites with superior toughness ($\sim 3-4$ $MPa \cdot m^{1/2}$) and super-high hardness (~ 30 GPa) by SPS at lower temperatures. It was demonstrated that increasing $MoSi_2$ addition (in the range 5-15 vol.%) benefits B_4C sinterability and toughness, without penalizing its hardness. It was also demonstrated that the B_4C - $MoSi_2$ powder mixtures lead to B_4C - MoB_2 - SiC triplex-particulate composites with fine-grained microstructures, and that $MoSi_2$ acts a reactive sintering additive that promotes the lower-temperature densification by transient liquid-phase sintering. Finally, research opportunities were presented for the future microstructural design of novel structural ceramic composites based on B_4C plus refractory borides and carbides.

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Figure Captions

Figure 1. Representative SEM micrographs of the fracture surface of the monolithic B₄C ceramics fabricated by SPS using target temperatures of (A) 1700, (B) 1750, (C) 1800, and (D) 1850 °C.

Figure 2. Representative SEM micrographs of the fracture surface of the B₄C composites fabricated by SPS using 5 vol.% MoSi₂ sintering additive and target temperatures of (A) 1800 and (B) 1850 °C.

Figure 3. Representative SEM micrographs of the fracture surface of the B₄C composites fabricated by SPS using 10 vol.% MoSi₂ sintering additive and target temperatures of (A) 1700, (B) 1750, and (C) 1800 °C.

Figure 4. Representative SEM micrographs of the fracture surface of the B₄C composites fabricated by SPS using 15 vol.% MoSi₂ sintering additive and target temperatures of (A) 1700, (B) 1750, and (C) 1800 °C.

Figure 5. (A) Representative SEM micrograph of the fracture surface of the B₄C composite fabricated by SPS using 15 vol.% MoSi₂ sintering additive and a target temperature of 1750 °C, and the corresponding elemental compositional maps of (B) Mo and (C) Si obtained by EDS.

Figure 6. Densification curves as a function of temperature for (A) the B₄C powder, and for the B₄C-MoSi₂ powder mixtures with (B) 5, (C) 10, and (D) 15 vol.% MoSi₂ logged during their SPS cycles with different target temperatures. The degrees of densification corresponding to the

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4 onset of the intermediate (70%) and final (90%) stages of sintering are marked by dotted and
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6 dashed lines, respectively.
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10 **Figure 7.** Shrinkage-rate curve as a function of temperature for the B₄C powder and for the B₄C-
11 MoSi₂ powder mixture with 15 vol.% MoSi₂ logged during their SPS cycle with a target
12 temperature of 1700 °C.
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19 **Figure 8.** Representative SEM micrographs of the electrochemically-etched, polished surface of
20 (A) the monolithic B₄C ceramic and (B) the B₄C composite with 15 vol.% MoSi₂ sintering
21 additive fabricated by SPS using a target temperature of 1800 °C.
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29 **Figure 9.** EDS spectra corresponding to the locations indicated by the numbers in Fig. 8B. Peak
30 assignments are included. K and O come from the electrochemical etching with KOH solution.
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32 Si detected in 1, B and Mo detected in 2, and Si and C detected in 3 come from neighbouring
33 grains because the effective interaction volume is larger than grain size.
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40 **Figure 10.** XRD patterns of (A) the monolithic B₄C ceramics and of the B₄C composites with
41 (B) 5, (C) 10, and (D) 15 vol.% MoSi₂ sintering additives fabricated by SPS using different
42 target temperatures. Peak assignments are included.
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49 **Figure 11.** Evolution of the vacuum level within the chamber of the SPS furnace as a function of
50 temperature for the B₄C powder and for the B₄C-MoSi₂ powder mixture with 15 vol.% MoSi₂
51 logged during their SPS cycle with a target temperature of 1700 °C.
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4 **Figure 12.** XRD patterns of the B₄C composites fabricated by SPS using 15 vol.% MoSi₂
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6 sintering additive and cycles interrupted at 1150, 1400, 1450, and 1500 °C. Peak assignments are
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8 included. Logarithmic scales have been used to facilitate the observation of the weaker XRD
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10 peaks.
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15 **Figure 13.** Representative SEM micrographs of the fracture surface of the B₄C composite with
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17 15 vol.% MoSi₂ sintering additive fabricated by SPS using a cycle interrupted at (A) 1250 and
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19 (B) 1400 °C. (C) SEM micrograph at higher magnification of a region in (B), showing more
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21 detail.
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Tables

Table 1. Fabrication conditions of all monolithic ceramics and ceramic composites, and their corresponding degrees of densification.

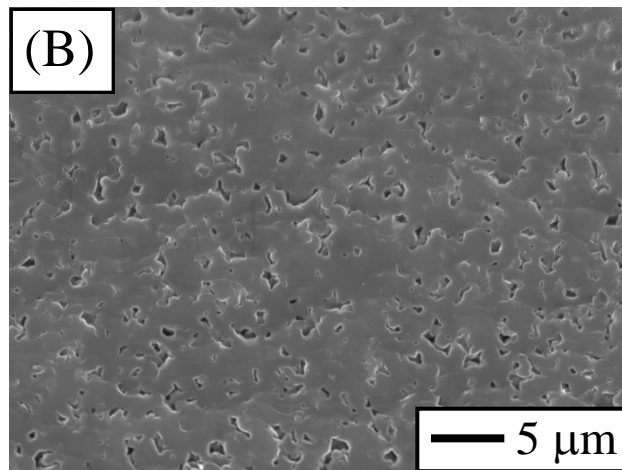
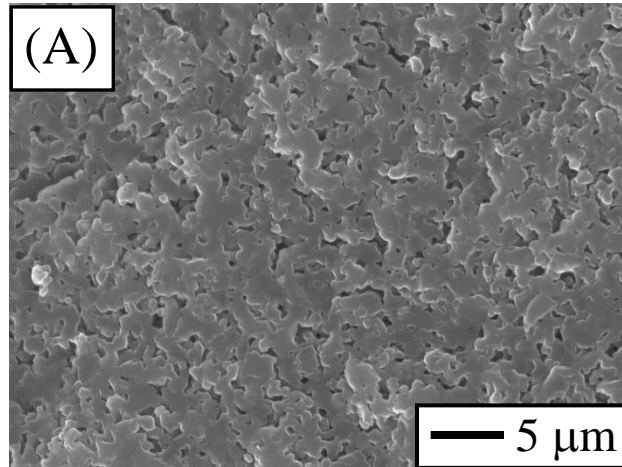
Type &	Target SPS temperature (°C) #			
	1700	1750	1800	1850
B ₄ C	✓ extremely porous	✓ very porous	✓ porous	✓ near-fully dense
B ₄ C-5vol.%MoSi ₂	—	—	✓ near-fully dense	✓ fully dense
B ₄ C-10vol.%MoSi ₂	✓ porous	✓ near-fully dense	✓ fully dense	—
B ₄ C-15vol.%MoSi ₂	✓ near-fully dense	✓ fully dense	✓ fully dense	—

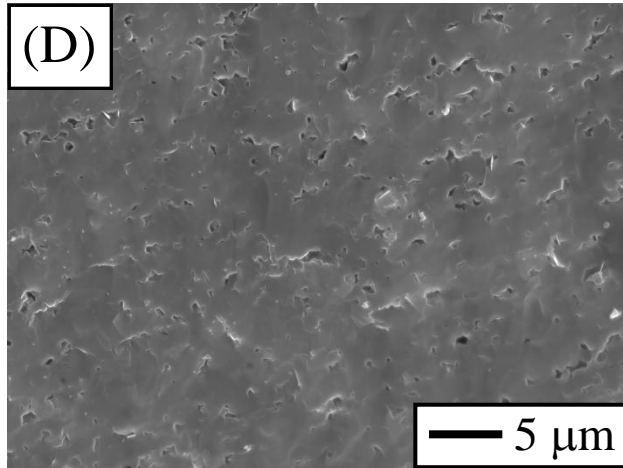
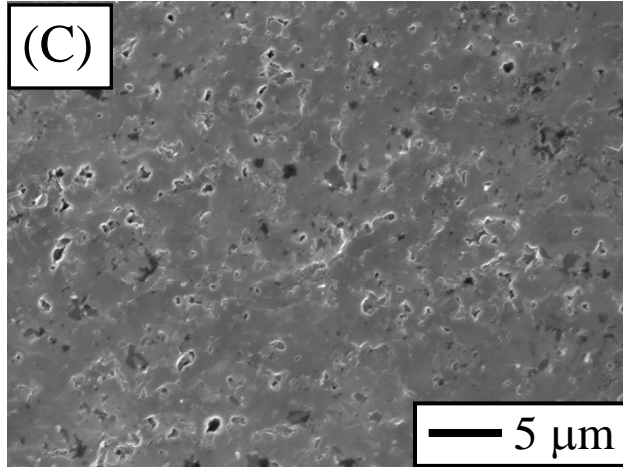
[&] Designations refer to the composition of the powder batches. [#] Other SPS conditions were: heating ramp of 100 °C/min and soaking time of 5 min at the target temperature. The lowest target SPS temperatures required to achieve near-full and full densification are in blue and red, respectively. The following five ranges of densification (d) have been defined: Full dense ($d=100\%$), near-full dense ($95\leq d<100\%$), porous ($90\leq d<95\%$), very porous ($80\leq d<90\%$), and extremely porous ($d<80\%$).

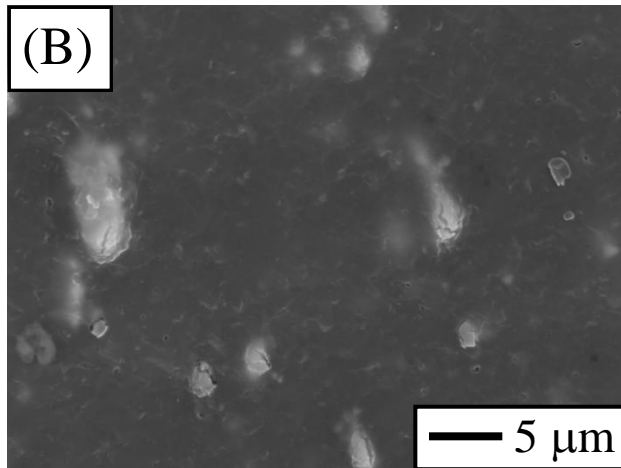
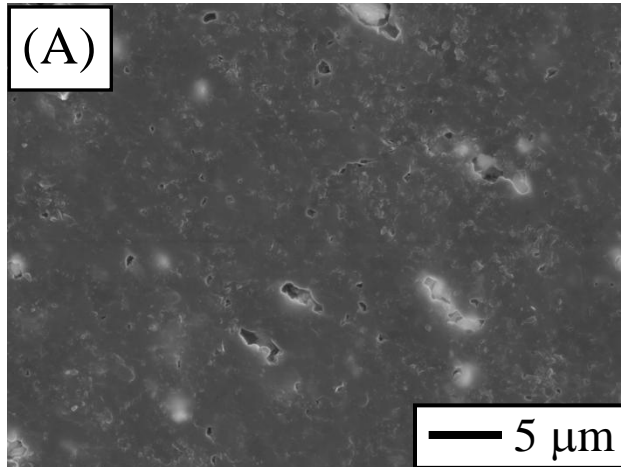
Table 2. Fabrication conditions, degrees of densification achieved, and mechanical properties of the monolithic ceramics and ceramic composites.

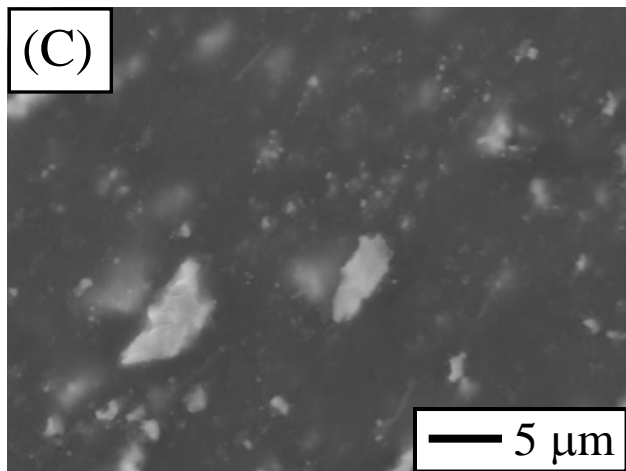
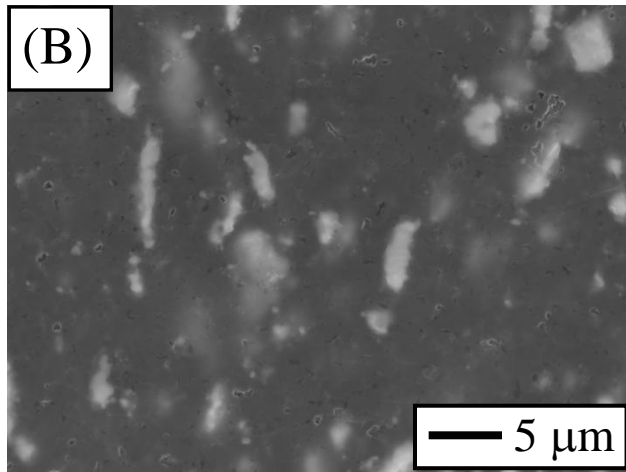
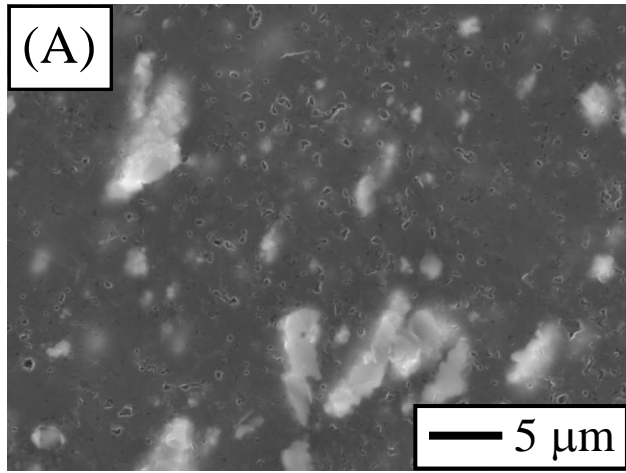
Type &	Target SPS temperature (°C) #	Densification (%)	Hardness (GPa)	Fracture toughness (MPa·m ^{1/2})
B ₄ C	1700	78	13±1	—
	1750	82	14±1	—
	1800	93	19±1	—
	1850	95	25.0±0.7	—
B ₄ C-5vol.%MoSi ₂	1800	96	19.8±0.8	—
	1850	100	29.5±0.9	3.1±0.2
B ₄ C-10vol.%MoSi ₂	1700	93	15.0±0.5	—
	1750	96	20±1	—
	1800	100	30.1±0.9	3.7±0.3
B ₄ C-15vol.%MoSi ₂	1700	97.5	23±1	—
	1750	100	30.1±0.7	4.1±0.3
	1800	100	30.1±0.6	4.1±0.3

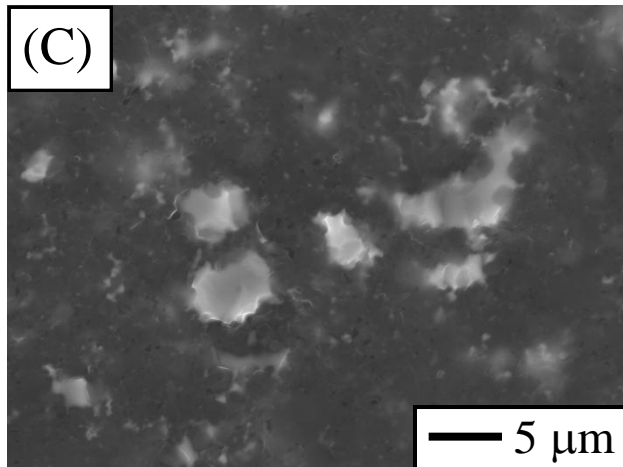
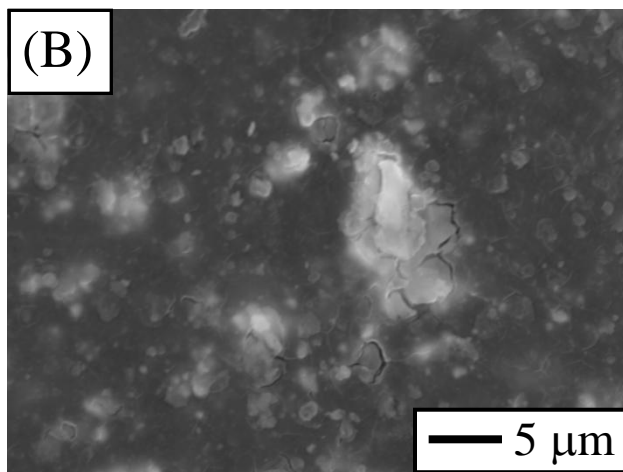
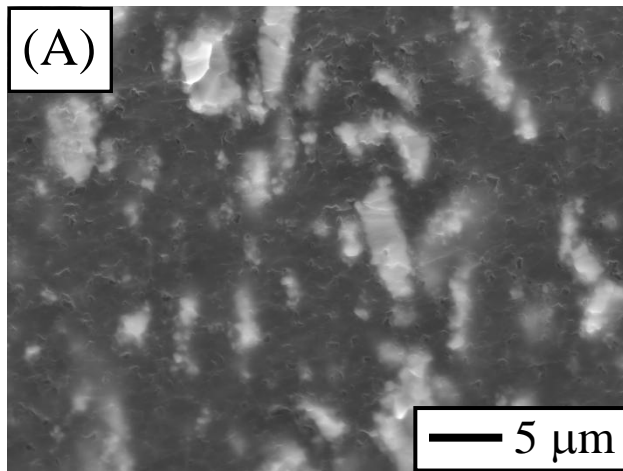
[&] Designations refer to the composition of the powder batches. [#] Other SPS conditions were: heating ramp of 100 °C/min and soaking time of 5 min at the target temperature.

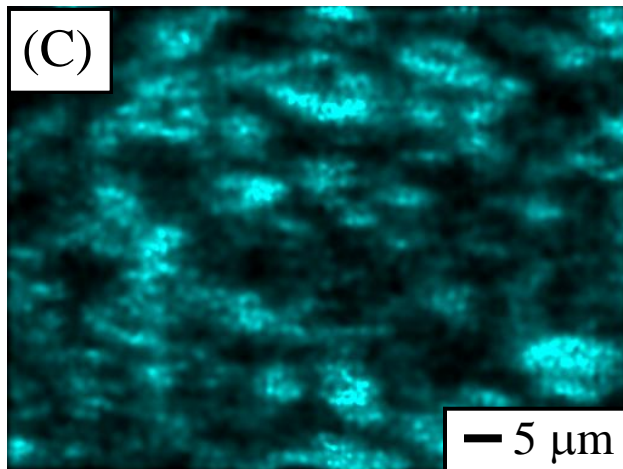
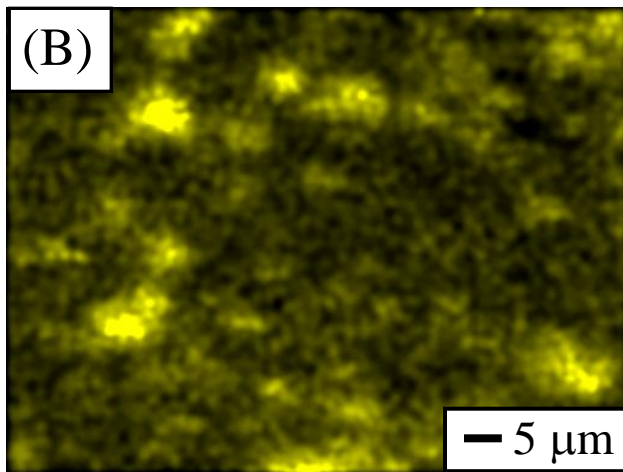
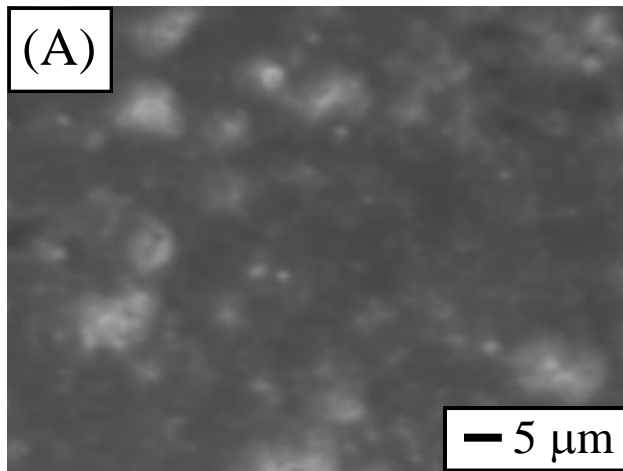


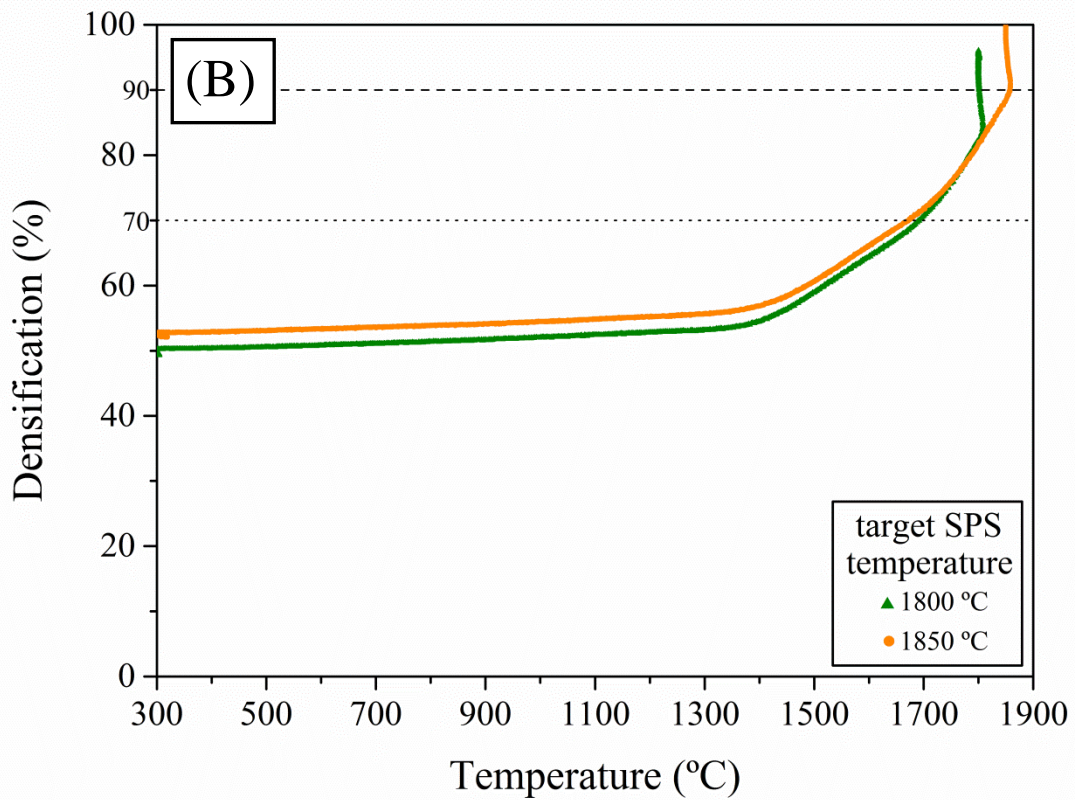
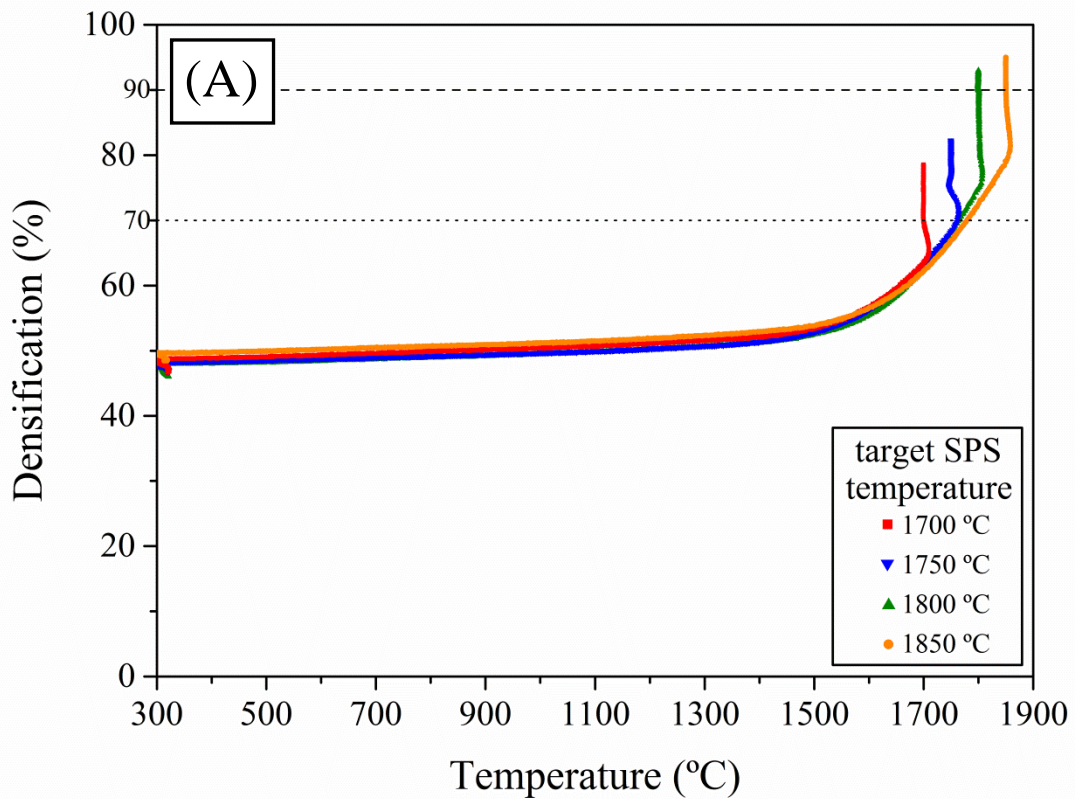


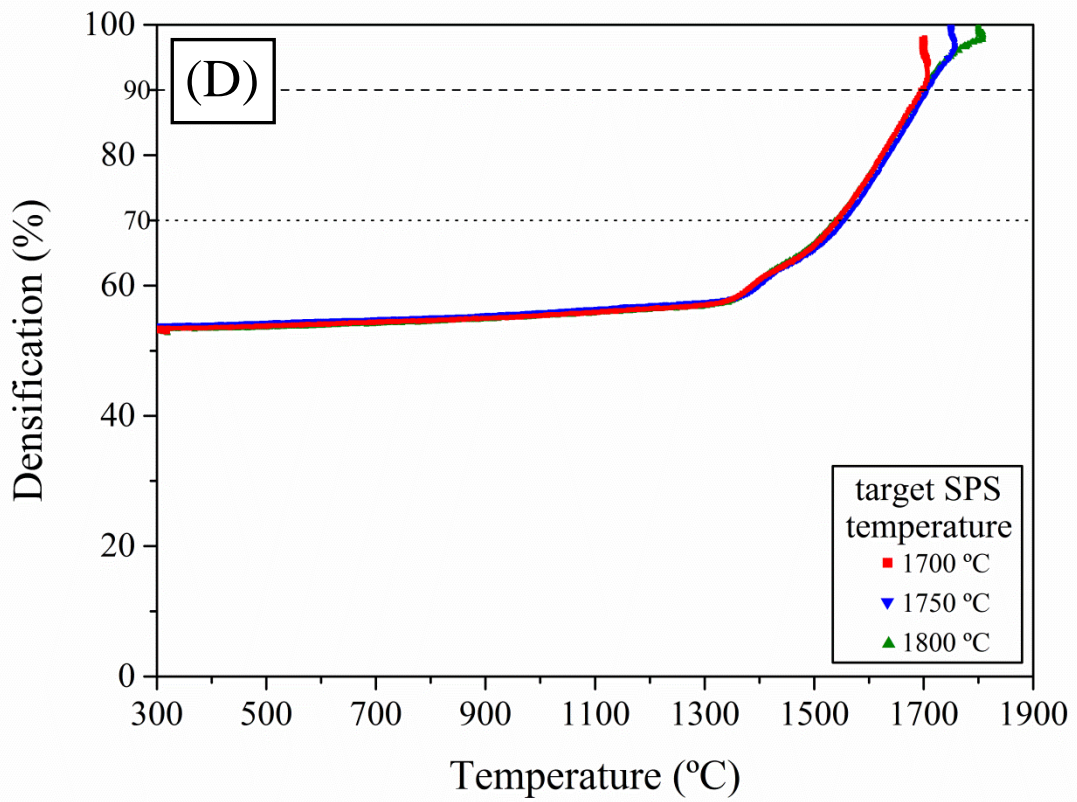
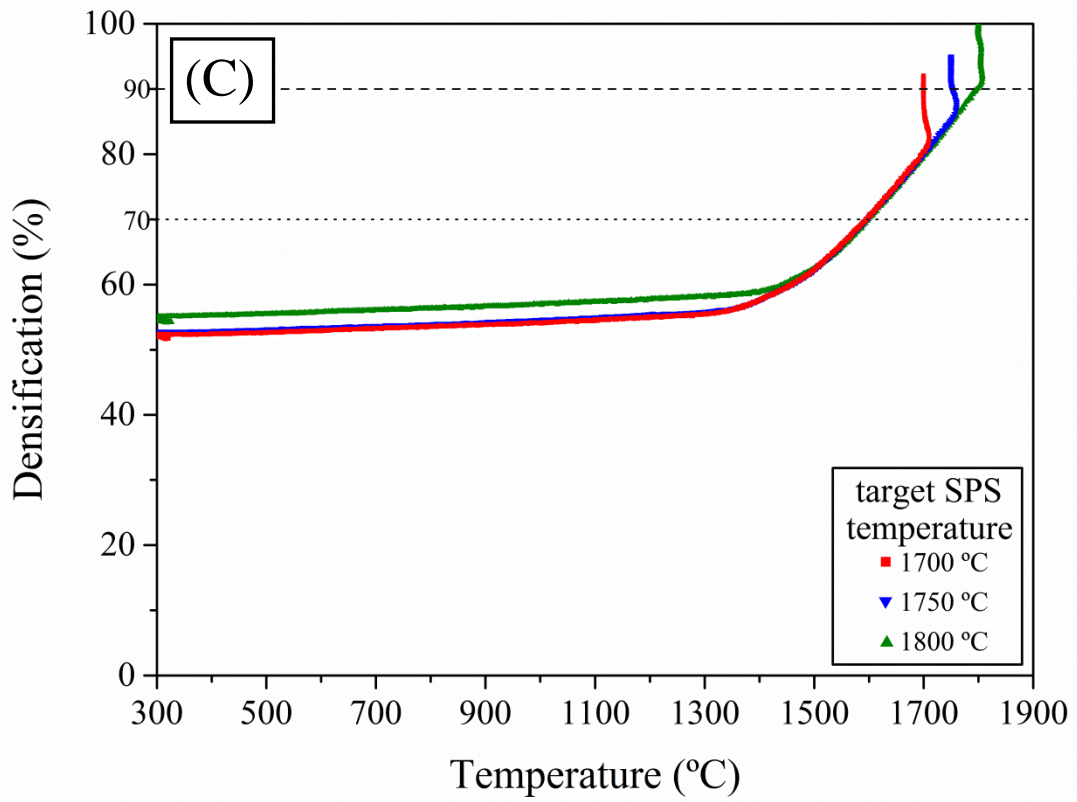


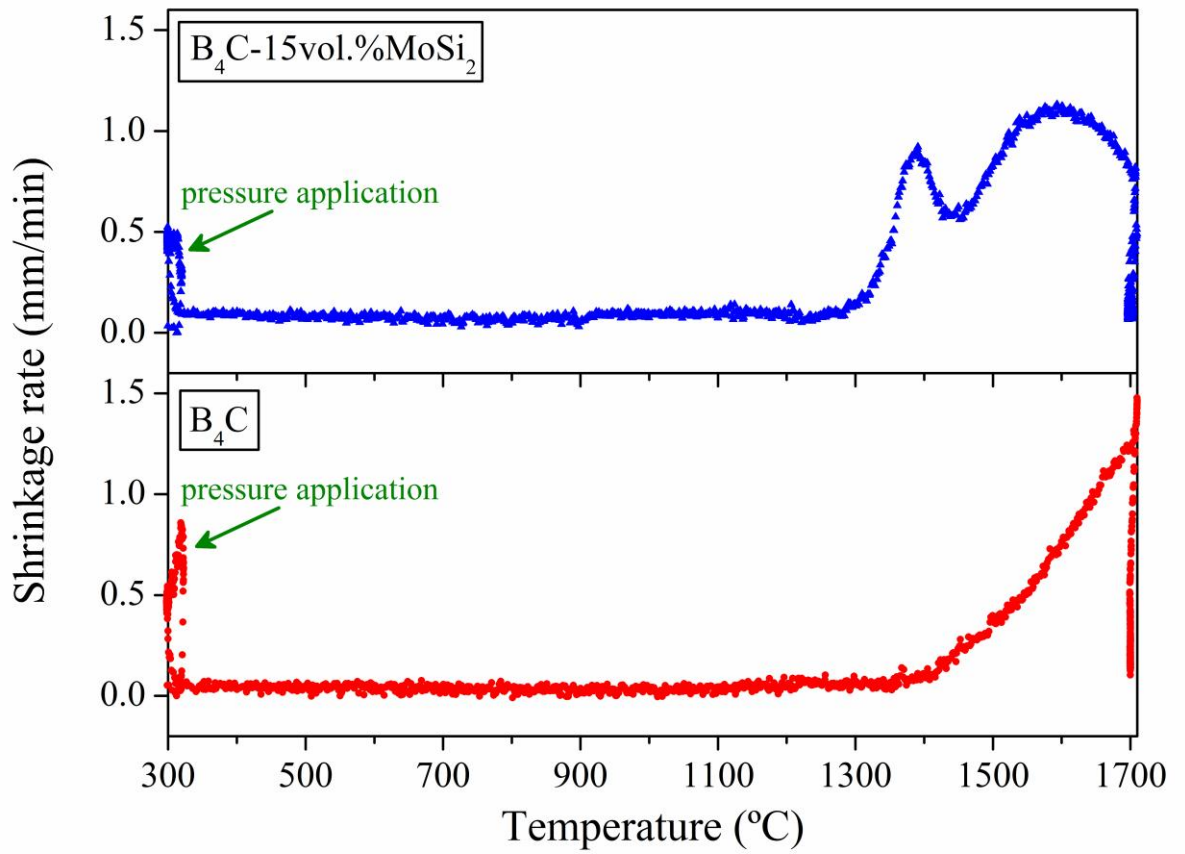


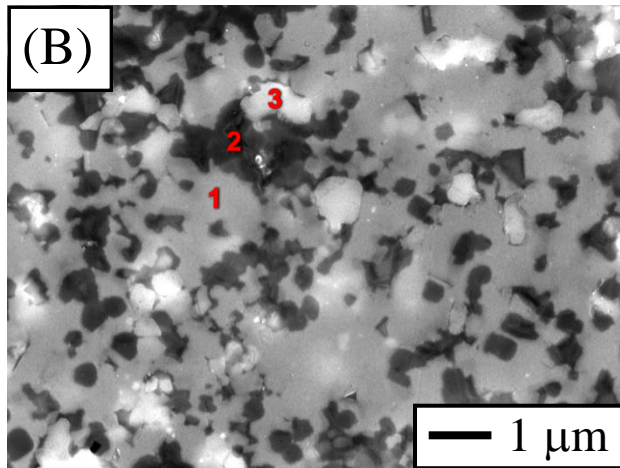
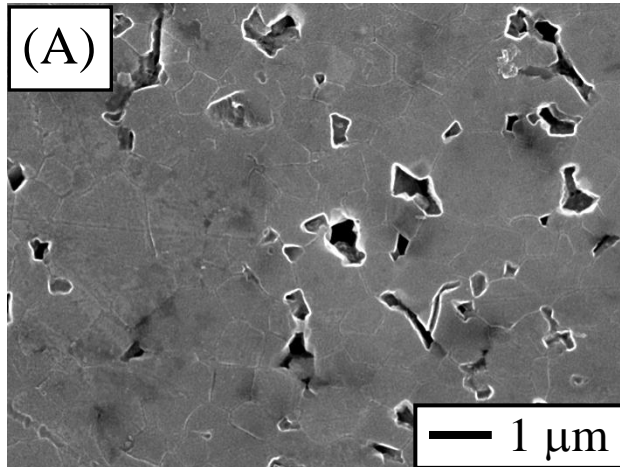


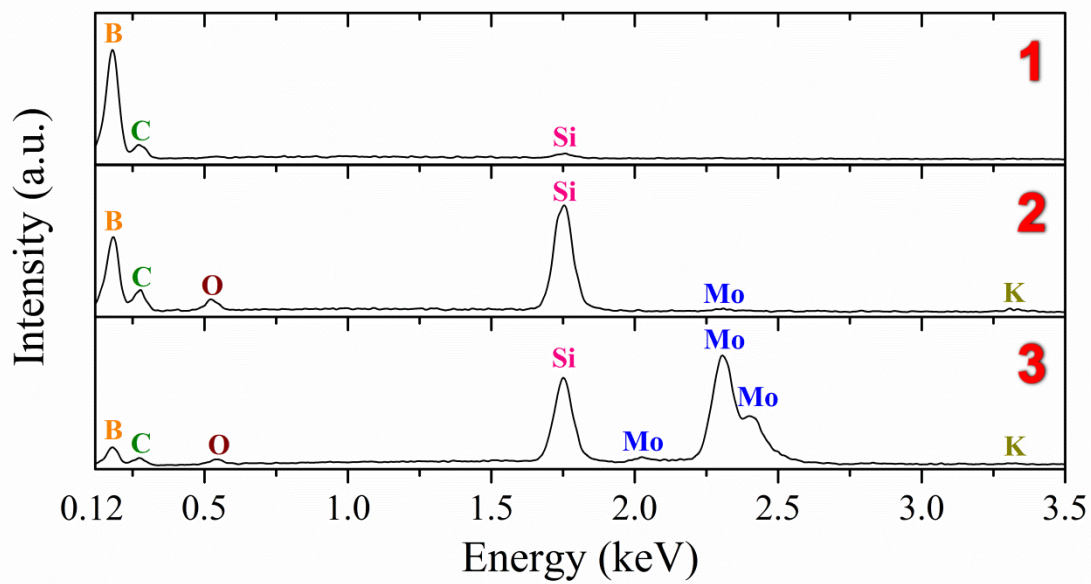


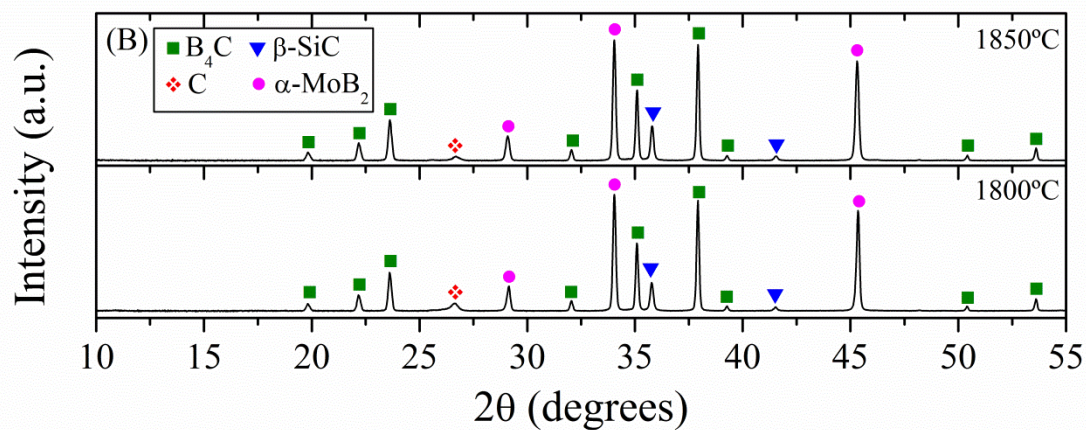
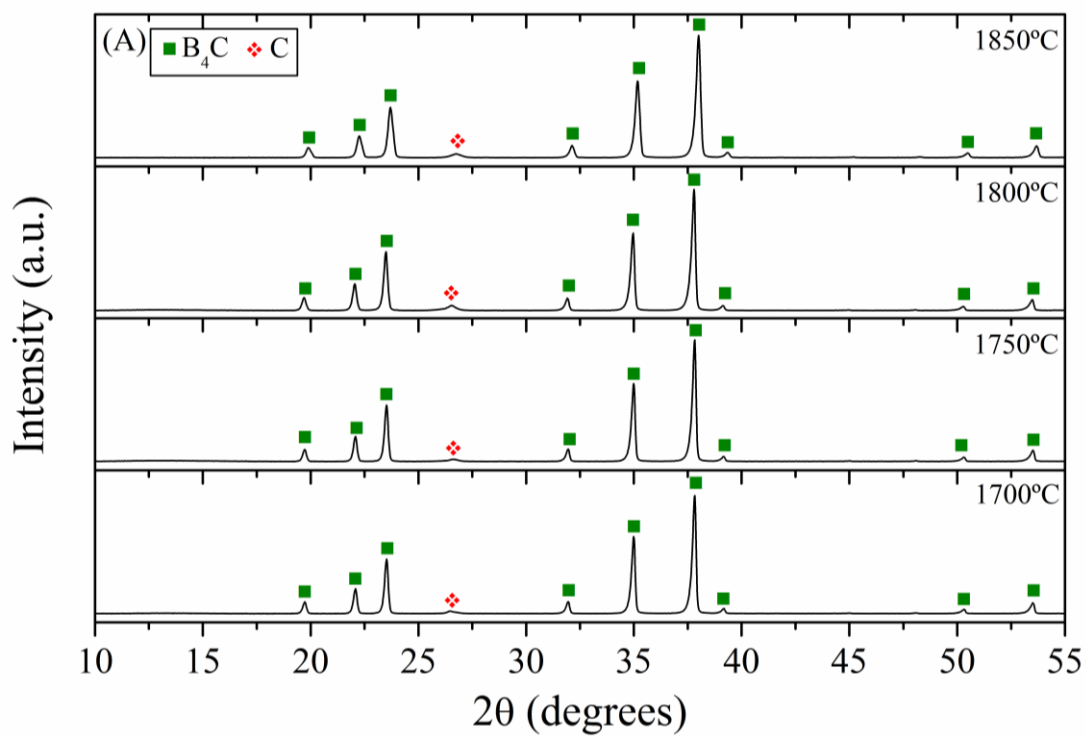


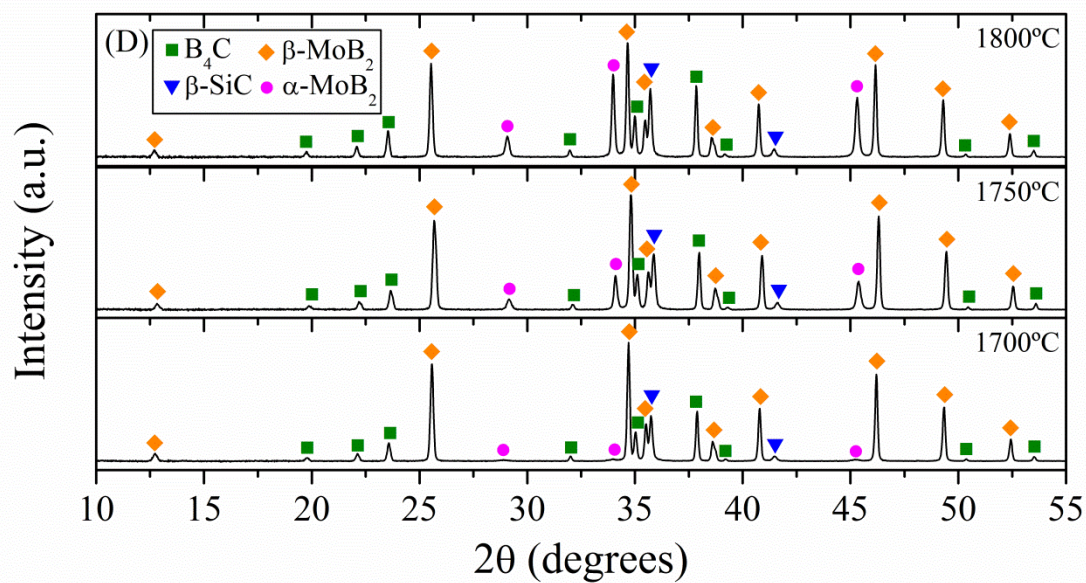
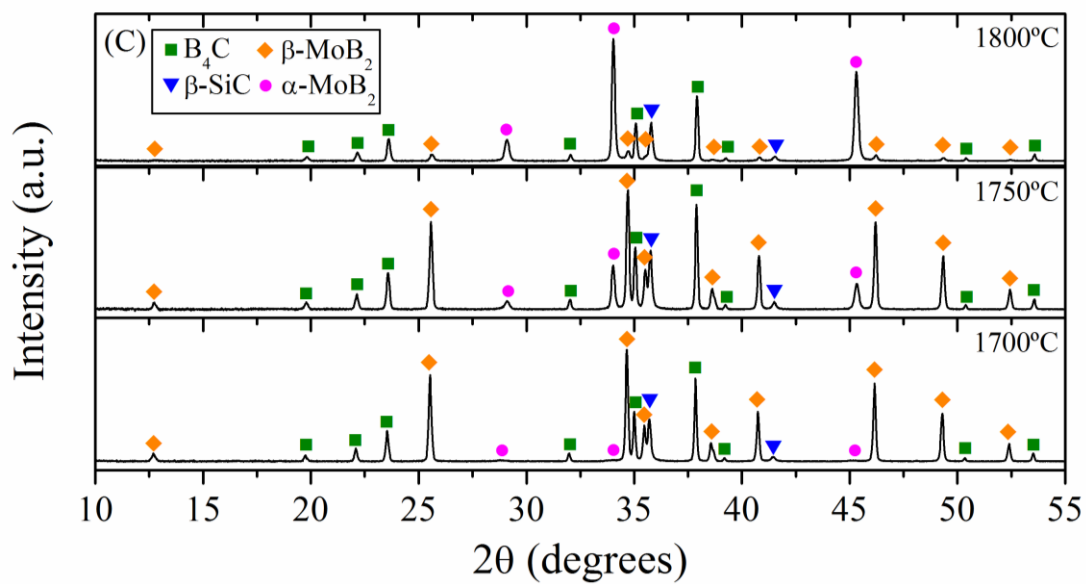


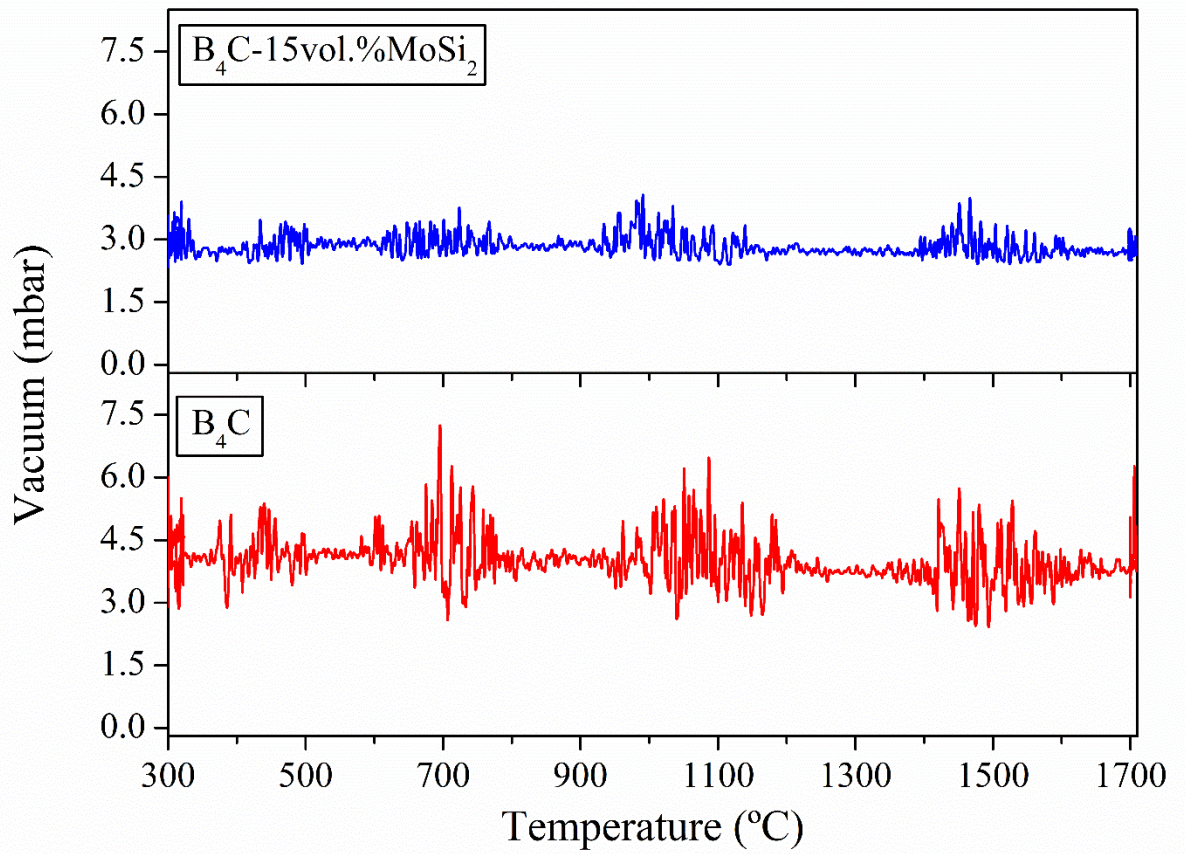


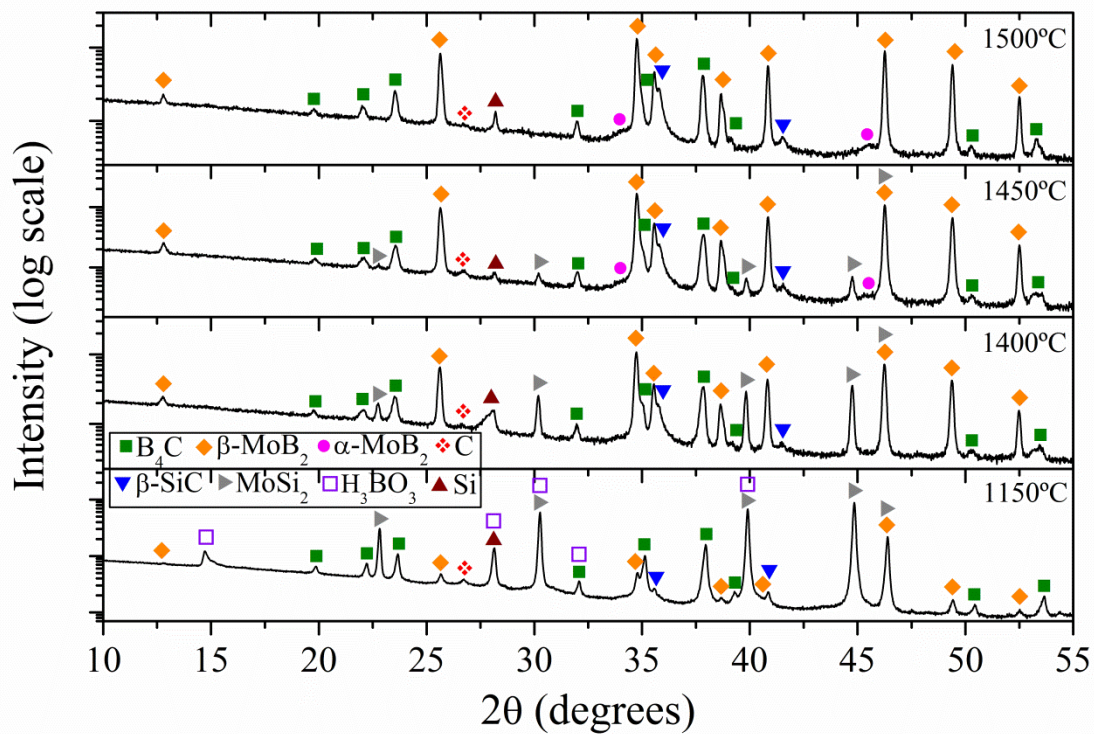


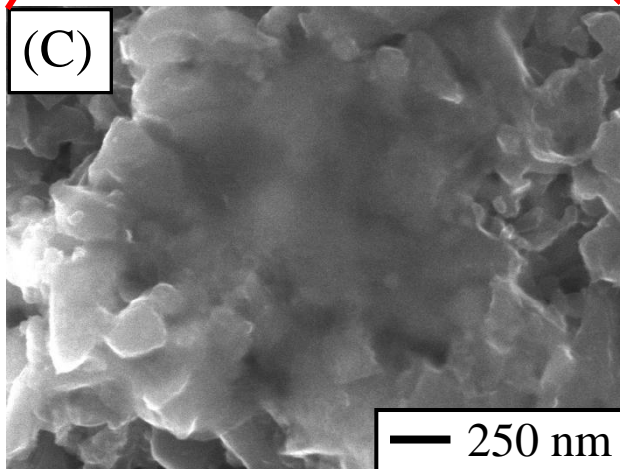
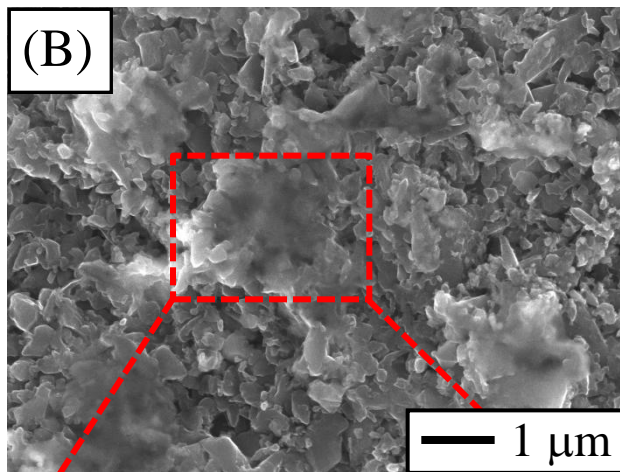
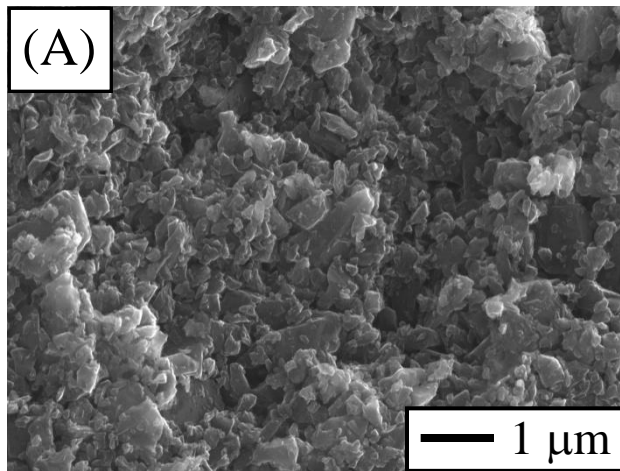












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Figure Captions

Figure 1. Representative SEM micrographs of the fracture surface of the monolithic B₄C ceramics fabricated by SPS using target temperatures of (A) 1700, (B) 1750, (C) 1800, and (D) 1850 °C.

Figure 2. Representative SEM micrographs of the fracture surface of the B₄C composites fabricated by SPS using 5 vol.% MoSi₂ sintering additive and target temperatures of (A) 1800 and (B) 1850 °C.

Figure 3. Representative SEM micrographs of the fracture surface of the B₄C composites fabricated by SPS using 10 vol.% MoSi₂ sintering additive and target temperatures of (A) 1700, (B) 1750, and (C) 1800 °C.

Figure 4. Representative SEM micrographs of the fracture surface of the B₄C composites fabricated by SPS using 15 vol.% MoSi₂ sintering additive and target temperatures of (A) 1700, (B) 1750, and (C) 1800 °C.

Figure 5. (A) Representative SEM micrograph of the fracture surface of the B₄C composite fabricated by SPS using 15 vol.% MoSi₂ sintering additive and a target temperature of 1750 °C, and the corresponding elemental compositional maps of (B) Mo and (C) Si obtained by EDS.

Figure 6. Densification curves as a function of temperature for (A) the B₄C powder, and for the B₄C-MoSi₂ powder mixtures with (B) 5, (C) 10, and (D) 15 vol.% MoSi₂ logged during their SPS cycles with different target temperatures. The degrees of densification corresponding to the

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4 onset of the intermediate (70%) and final (90%) stages of sintering are marked by dotted and
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6 dashed lines, respectively.
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10 **Figure 7.** Shrinkage-rate curve as a function of temperature for the B₄C powder and for the B₄C-
11 MoSi₂ powder mixture with 15 vol.% MoSi₂ logged during their SPS cycle with a target
12 temperature of 1700 °C.
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19 **Figure 8.** Representative SEM micrographs of the electrochemically-etched, polished surface of
20 (A) the monolithic B₄C ceramic and (B) the B₄C composite with 15 vol.% MoSi₂ sintering
21 additive fabricated by SPS using a target temperature of 1800 °C.
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29 **Figure 9.** EDS spectra corresponding to the locations indicated by the numbers in Fig. 8B. Peak
30 assignments are included. K and O come from the electrochemical etching with KOH solution.
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50 temperature for the B₄C powder and for the B₄C-MoSi₂ powder mixture with 15 vol.% MoSi₂
51 logged during their SPS cycle with a target temperature of 1700 °C.
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4 **Figure 12.** XRD patterns of the B₄C composites fabricated by SPS using 15 vol.% MoSi₂
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6 sintering additive and cycles interrupted at 1150, 1400, 1450, and 1500 °C. Peak assignments are
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Tables

Table 1. Fabrication conditions of all monolithic ceramics and ceramic composites, and their corresponding degrees of densification.

Type &	Target SPS temperature (°C) #			
	1700	1750	1800	1850
B ₄ C	✓ extremely porous	✓ very porous	✓ porous	✓ near-fully dense
B ₄ C-5vol.%MoSi ₂	—	—	✓ near-fully dense	✓ fully dense
B ₄ C-10vol.%MoSi ₂	✓ porous	✓ near-fully dense	✓ fully dense	—
B ₄ C-15vol.%MoSi ₂	✓ near-fully dense	✓ fully dense	✓ fully dense	—

[&] Designations refer to the composition of the powder batches. [#] Other SPS conditions were: heating ramp of 100 °C/min and soaking time of 5 min at the target temperature. The lowest target SPS temperatures required to achieve near-full and full densification are in blue and red, respectively. The following five ranges of densification (d) have been defined: Full dense ($d=100\%$), near-full dense ($95\leq d<100\%$), porous ($90\leq d<95\%$), very porous ($80\leq d<90\%$), and extremely porous ($d<80\%$).

Table 2. Fabrication conditions, degrees of densification achieved, and mechanical properties of the monolithic ceramics and ceramic composites.

Type &	Target SPS temperature (°C) #	Densification (%)	Hardness (GPa)	Fracture toughness (MPa·m ^{1/2})
B ₄ C	1700	78	13±1	—
	1750	82	14±1	—
	1800	93	19±1	—
	1850	95	25.0±0.7	—
B ₄ C-5vol.%MoSi ₂	1800	96	19.8±0.8	—
	1850	100	29.5±0.9	3.1±0.2
B ₄ C-10vol.%MoSi ₂	1700	93	15.0±0.5	—
	1750	96	20±1	—
	1800	100	30.1±0.9	3.7±0.3
B ₄ C-15vol.%MoSi ₂	1700	97.5	23±1	—
	1750	100	30.1±0.7	4.1±0.3
	1800	100	30.1±0.6	4.1±0.3

[&] Designations refer to the composition of the powder batches. [#] Other SPS conditions were: heating ramp of 100 °C/min and soaking time of 5 min at the target temperature.

- Toughened, super-hard B_4C triplex-particulate composites have been fabricated at lower than usual temperatures by spark plasma sintering with $MoSi_2$ additives (5, 10, and 15 vol.%).
- It is proved that $MoSi_2$ is a reactive sintering additive that promotes densification by transient liquid-phase sintering, thus yielding fully-dense B_4C-MoB_2-SiC composites at relatively lower temperatures.
- It is demonstrated that the $MoSi_2$ first reacts at moderate temperatures with part of B_4C to form MoB_2 , SiC , and Si . This last is a transient component that eventually melts, contributing to densification by liquid-phase sintering, and then reacts with free C present in the B_4C starting powders to form more SiC , after which densification continues by solid-state sintering.
- It is shown that that the B_4C-MoB_2-SiC composites are super-hard (~ 30 GPa), tough ($\sim 3-4$ $MPa \cdot m^{1/2}$), and fine-grained, a combination that renders them very appealing for structural applications.
- Research opportunities are discussed for the future microstructural design of a novel family of toughened, ultra-hard/super-hard multi-particulate composites based on B_4C plus refractory borides and carbides.