SiC-based sandwich material for Flow Channel Inserts in DCLL blankets: Manufacturing, characterization, corrosion tests

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highlights

- · A wide range of porous SiC materials with different properties were produced.
- The produced 50% porous SiC has $k \approx 5$ W/m K at 700° C and $\approx 3.10^{-5}$ S/m at 550° C.
- The produced 50% porous SiC has a flexural strength in the range 50-80 MPa.
- A 200 m dense CVD-SiC coating offers reliable protection against static PbLi corrosion.

abstract

Flow Channel Inserts (FCIs) are key elements in a DCLL blanket concept for DEMO, since they provide the required thermal insulation between the He cooled structural steel and the hot liquid PbLi flowing at \approx 700° C, and the necessary electrical insulation to minimize MHD effects. In this work a SiC-based sandwich material is proposed for FCIs, consisting of a porous SiC core covered by a dense CVD-SiC layer. A method to produce the porous SiC core is presented, based on combining a starting mixture of SiC powder with a spherical carbonaceous sacrificial phase, which is removed after sintering by oxidation, in such a way that a microstructure of spherical pores is achieved. Following this technique, a porous SiC material with low thermal and electrical conductivities, but enough mechanical strength was produced. Samples were covered by a 200 m thick CVD-SiC coating to form a SiC-sandwich material. Finally, corrosion tests under static PbLi were performed, showing that such a dense layer offers a reliable protection against static PbLi corrosion.

1. Introduction

The Dual Coolant Lead Lithium (DCLL) blanket is one of the most attractive concepts for its application in a future DEMO reac-tor. In the high temperature DCLL design a eutectic Pb-15.7Li alloy flows through long poloidal channels acting as self-cooled breeder, absorbing the neutron flux energy and reaching tempera-tures around 700 °C. Helium is used to cool the blanket structure, made of reduced activation ferritic-martensitic (RAFM) steel, that has a maximum allowable PbLi interface temperature of around 470 °C due to corrosion issues [1]. Such a blanket concept provides

final thermal efficiencies around ≈45%, considerably higher than those achieved in other blanket designs [2].

Flow channel inserts (FCIs) are key elements in the DCLL design; they consist of hollow channels containing the flowing PbLi and being separated from the RAFM steel wall by a thin gap, also filled with PbLi. The thicknesses typically proposed for the FCI's wall and gap in the high temperature DCLL are 5 and 2 mm, respec-tively [3]. FCIs serve as electrical insulators, to reduce MHD effects, and as thermal insulators, to avoid heat loses from the hot metal to the steel walls. To fulfill these functions, FCI's materials must have low thermal and electrical conductivities, good PbLi corrosion resistance, low activation and degradation by neutrons and excellent high temperature performance. Besides, FCIs will be subject to mechanical stresses during operation due to high thermal gradi-

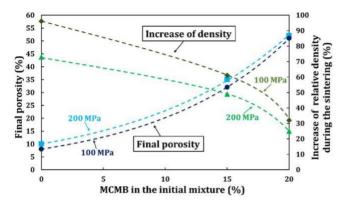


Fig. 1. Total porosity of the samples after sintering and oxidation, and increase of their relative density during sintering, as a function of the initial amount of sacrificial phase.

ent across its walls [4]. For this reason, FCI's materials must exhibit enough mechanical strength to withstand this thermal stresses.

At present, SiC is the main candidate material for the FCIs in the high-temperature DCLL, thanks to its excellent stability at high temperatures, low thermal expansion, high thermal shock resis-tance, low activation and low corrosion by PbLi [5]. However, the need for low electrical and thermal conductivities requires sub-stantial R&D efforts in the development and fabrication of new SiC-based materials, and different possibilities are being consid-ered. One promising option is a SiC-sandwich material formed by a core of porous SiC covered by a dense SiC coating fabricated by chemical vapor deposition (CVD): the porous core provides the insulating properties while the dense layer offers protection against PbLi corrosion and infiltration. However, a reliable method to pro-duce porous SiC with the required characteristics for FCI application has to be developed, and the resulting sandwich material should be characterized and tested under relevant conditions.

In this work, a method to produce porous SiC based in the sacrificial template technique is presented. In this method a sac-

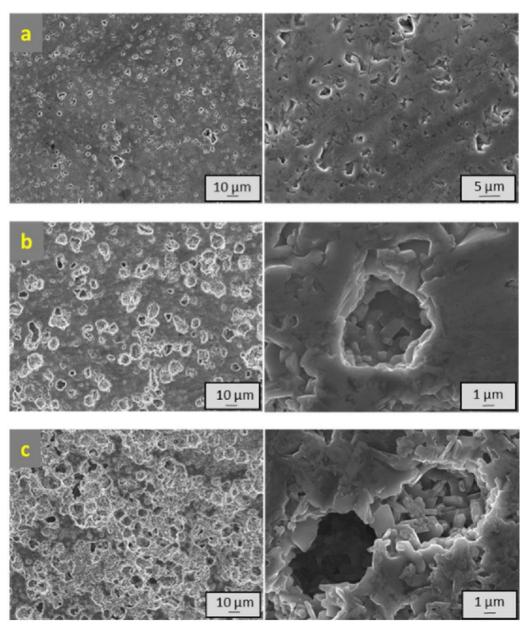


Fig. 2. FESEM micrographs of samples fabricated with: a) 0% MCMB; b) 15% MCMB and c) 20% MCMB in the initial powder mixture, pressed at 100 MPa.

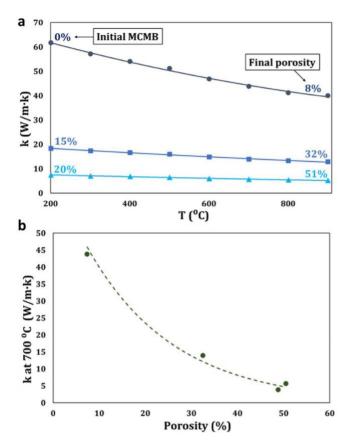


Fig. 3. a) Thermal conductivity as a function of temperature for different initial MCMB contents; b) Thermal conductivity at 700 °C versus porosity.

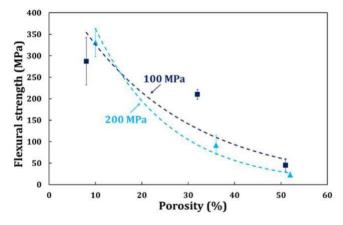


Fig. 4. Flexural strength versus porosity for samples pressed at 100 and 200 MPa.

rificial phase is introduced in the initial powder mixture, being removed after sintering and leaving spherical pores instead. Mate-rials produced by this method and their characterization in terms of microstructure, thermal and electrical conductivities, and flexural strength are presented.

A previous thermomechanical study of the basic heat transfer problem of FCIs described in [4] provides a guideline for the mate-rial's development. The first objective marked by this study was to produce a material with thermal conductivity $\leq 7 \text{ W/(m K)}$ and flex-ural strength well above 25 MPa to offer reliable protection against thermal stresses. The achieved materials properties are compared with the requirements marked by this study, to determine their suitability as porous SiC core of a sandwich FCI.

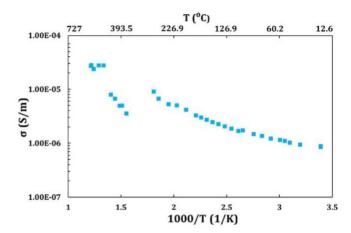


Fig. 5. Electrical conductivity versus the inverse of temperature of a 50% porous SiC sample.

Another important point to determine the viability of the SiC sandwich concept for FCIs is to study the reliability of the dense CVD-SiC coating covering the porous core, in terms of quality (lack of defects) and corrosion resistance. According to the mentioned thermo-mechanical study [4], a coating thickness resulting in rea-sonable mechanical stresses would be 200 m. Thus, porous SiC samples produced in this work were covered by a dense CVD-SiC coating of ≈200 m. In order to test if such a dense coating offers effective protection against PbLi corrosion, laboratory tests have been performed on SiC-sandwich samples under static PbLi at ≈700 °C for 1000 h. Tests under flowing PbLi in presence of a magnetic field will be performed shortly.

2. Production of porous SiC

2.1. Experimental procedure

SiC powder (Superior Graphite, 0.5 m) was mixed with 2.5 wt.% Al₂ O₃ (0.4 m) and Y₂ O₃ (1 m) powders as sintering additives, added in a 3/2 relation. The sacrificial phase proposed in this work is a graphitized powder of spherical mesocarbon microbeads (MCMB, 15 m). 3 wt.% of an aqueous polymer dispersion was used as binder. Additives, SiC and binder were mixed in ethanol for 16–18 h, being the MCMB subsequently added to the solution and mixed for 30 min. The resulting blend was dried and uniaxially pressed to the required geometries at 100 and 200 MPa. The green compacts were sintered at 1850 $^{\circ}$ C for 1 h. The sintered samples were heated at 700 $^{\circ}$ C in air for 5 h to burn out the carbonaceous sacrificial phase.

Density of the samples was determined, in green state and before and after the oxidation treatment, by geometrical mea-surements, and porosity was calculated from geometrical and theoretical densities; the latter was obtained by the rule of mix-ture. Microstructure was studied by field emission gun scanning electron microscopy (FESEM) and energy dispersive X-ray spectroscopy (EDS). Thermal conductivity as a function of temperature was determined from the heat capacity (obtained from [6]), density of the samples and their thermal diffusivity, measured by the Laser Flash method. Flexural strength was determined at room temper-ature by three point bending tests (3PBT) using 4 samples for each condition. The electrical conductivity was measured in high vac-uum between 15 and 550 $^{\circ}$ C. The corrosion test was carried out under static eutectic PbLi at ≈ 700 $^{\circ}$ C, immersing the samples for 1000 h in a niobium container placed in a vacuum chamber.

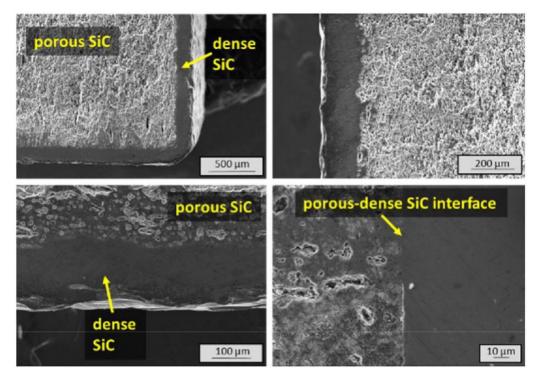


Fig. 6. Cross-sections of SiC-sandwich samples formed by a ≈50% porous SiC and a dense CVD-SiC coating of ≈200

m thickness (secondary electrons).

2.2. Results

In Fig. 1 the final porosity of the samples pressed at 100 and 200 MPa as well as the increase of relative density during sinter-ing are shown as a function of three different initial amounts of MCMB (sacrificial phase): 0, 15 and 20 wt.%. It can be observed that the porosity increases in an exponential way with increasing MCMB content, while the variation of relative density during sin-tering decreases. The sintering process is less effective in presence of the MCMB phase; this is probably due to the fact that the MCMB particles are external to the SiC-Al₂ O₃ -Y₂ O₃ sintering system, and they do not suffer any change in composition or volume during sin-tering, being rather an obstacle in the shrinkage of the SiC matrix. Besides, the sintering process seems to be less effective on samples pressed at 200 MPa; this may indicate the presence of cracks inside the green samples caused by internal stresses, usually appearing in ceramics during uniaxial pressing [7].

Fracture surfaces of the materials produced are shown in Fig. 2. The final porosity of the samples with initial MCMB is formed by smaller pores (≤ 1 m) present in the SiC matrix caused by par-tial sintering, and by spherical pores ($\approx 10-15$ m) formed by burn out of the sacrificial phase, as can be seen in b) and c). The higher sintering grade of b), the sample with less initial MCMB, can be appreciated. Micrographs in a) show the pores present at the sur-face of a sample sintered without any sacrificial phase, where pores due to partial sintering are more abundant than inside, as has been reported in other SiC materials sintered with Al₂ O₃ -Y₂ O₃ as addi-tives [8].

One crucial property of the porous SiC material for application as core of FCIs is its thermal conductivity. In Fig. 3. a) the thermal conductivity as a function of temperature can be seen for samples of different initial MCMB content and thus of different final porosity. As expected, the thermal conductivity decreases with tempera-ture and with porosity. The thermal conductivity of the samples at 700 °C (the expected operation temperature of a FCI in the high temperature DCLL) versus its porosity is shown in b). The data can be well fitted to an exponential law, as has been noticed in a pre-

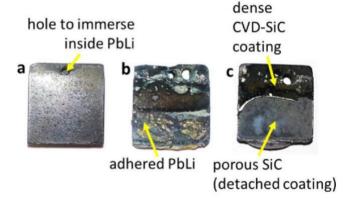


Fig. 7. a) Sample before the corrosion experiment; b) Undamaged sample after the experiment, conserving its whole CVD-SiC coating; the region exposed to liquid PbLi during the experiments, exhibiting metal adhered to the surface, can be clearly appreciated; c) Damaged sample after the corrosion experiment with part of the coating detached during manipulation after the tests.

vious work [9], although this behavior should be confirmed with further measurements.

In Fig. 4, the flexural strength versus the porosity of the samples is shown for the two used compaction pressures: 100 and 200 MPa. Different results are obtained depending on the compaction pres-sure; this behavior, together with the important dispersions of the measured values, seems to confirm the hypothesis that cracks or other defects were present in the samples. For this reason, the flex-ural strength is possibly underestimated in some of these results. In ceramic materials, the relationship between the flexural strength and porosity has been reported as exponential [10,11]; however, the dispersion of the obtained results prevents clear conclusions, and in any case, more work should be done to produce samples free of defects.

Finally, the electrical conductivity of a 50% porous sample has been measured as a function of temperature until 550 $^{\circ}$ C. Results

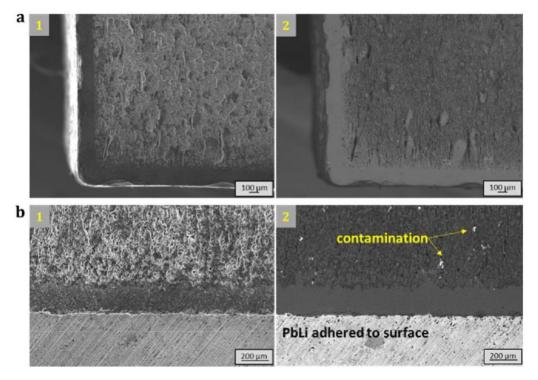


Fig. 8. SEM micrographs of: a) Corner of a sample after the corrosion experiment; b) Detail of the CVD-SiC layer of another sample after the experiment, with PbLi adhered to the outer surface (1-secondary electrons; 2- backscattered electrons).

are shown in Fig. 5, being similar to those obtained in previous work [9].

3. Corrosion test of coated samples

In order to study if a dense \approx 200 m thick CVD-SiC layer offers a reliable protection against PbLi corrosion at 700 °C, $14 \times 14 \times 5$ mm³ porous SiC samples with \approx 50% porosity were cov-ered by a dense SiC coating produced by Archer Technicoat Ltd, UK. Cross-sections of the produced sandwich samples are shown in Fig. 6. The thickness of the dense layer is near 180–200 m in all areas and no cracks or other damages were found.

The covered samples were immersed in static PbLi at ≈700 °C during 1000 h. After the experiment the samples seemed undam-aged, although pieces of the dense layer were detached from some samples during their manipulation after the test, as shown in the photographs of Fig. 7. No metal infiltration or deposition was detected on the surface of the damaged samples, confirming that detachments did not occur during the experiment.

SEM and EDS analysis were performed inside the samples; micrographs of samples after the experiment can be seen in Fig. 8. No lead was detected inside them and no cracks or damages in the dense CVD-SiC coating were found, being the thickness of the coating the same as before the experiment in all samples. Contam-ination observed by backscattered electrons in the interior of the sample (Fig. 8) is superficial and is assumed to originate from the cutting procedure.

4. Discussion

Assuming a porous sandwich core of \approx 5 mm thick and follow-ing the guidelines of the thermomechanical study mentioned in the introduction [4], a porous SiC with a porosity around \approx 43% would have the required thermal conductivity of \approx 7 W/m K, as can be deduced from the relationship showed in Fig. 3 b). Such a material can be produced with the proposed method by adding

≈18 wt.% sacrificial phase to the initial powder mixture, as can be inferred from the relationship between the initial MCMB and the final porosity of the samples showed in Fig. 1. Even though these relationships should be taken as guidelines and not as exact val-ues, it seems possible to produce a porous SiC material with the required properties to be used in FCIs. Observing the values of flexural strength obtained (in the range 50–80 MPa), it seems that such a porous material would have enough strength to support the expected thermal stresses. However, the dispersion obtained in the flexural strength results indicates that future work is required to reduce the defects present in the material and to assure its integrity in service. Regarding the electrical properties, the measured electri-cal conductivity is well below the most restrictive requirements for FCIs found in literature so far (≈ 1 S/m for inboard blankets [2]), and although the values showed in Fig. 5 correspond to the porous material, an important increase in the electrical conductivity due to the dense CVD-SiC layers is not expected.

5. Conclusions

The following conclusions can be drawn from this work:

- With the proposed method, a porous material with required ther-mal conductivity and enough mechanical strength to form the core of a SiCsandwich FCI can be produced. The properties of the material can be varied in a wide range by tailoring its porosity.
- The electrical conductivity of the produced porous SiC is suffi-ciently low to assure the FCI insulation.
- A 200 m thick free-of-defects dense CVD-SiC coating offers a reliable protection against static PbLi corrosion.

Acknowledgments

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References

- [1] C.P.C. Wong, M. Abdou, M. Dagher, Y. Katoh, R.J. Kurtz, S. Malang, E.P. Marriott, B.J. Merrill, K. Messadek, N.B. Morley, M.E. Sawan, S. Sharafat, S. Smolentsev, D.K. Sze, S. Willms, A. Ying, M.Z. Youssef, An overview of the US DCLL ITER-TBM program, Fusion Eng. Des. 85 (7–9) (2010) 1129–1132.
- [2] S. Smolentsev, N.B. Morley, M.A. Abdou, S. Malang, Dual-coolant lead-lithium (DCLL) blanket status and R&D needs, Fusion Eng. Des. (2015) 1–11.
- [3] S. Smolentsev, N.B. Morley, C. Wong, M. Abdou, MHD and heat transfer considerations for the US DCLL blanket for DEMO and ITER TBM, Fusion Eng. Des. 83 (10–12) (2008) 1788–1791.
- [4] C. Soto, J.M. Martínez-Esnaola, C. García-Rosales, Thermomechanical analysis of a Flow Channel Insert based on a SiC-sandwich material concept, Nucl. Mater. Energy 7 (2016) 5–11.

- [5] B. Riccardi, L. Giancarli, A. Hasegawa, Y. Katoh, A. Kohyama, R.H. Jones, L.L. Snead, Issues and advances in SiCf/SiC composites development for fusion reactors, J. Nucl. Mater. 329-333 (1–3) (2004) 56–65 (PART A).
- [6] K.H.O. Knacke, O. Kubaschewski, Thermochemical Properties of Inorganic Substances, 2nd ed., Springer, Düseldorf, 1991.
- [7] A. Singh, N.F. Complex, End-capping and Other Defects in Pressed Ceramic Compacts, 1996, pp. 196–200 (no. October).
- [8] T. Grande, H. Sommerset, E. Hagen, K. Wiik, M. a Einarsrud, Effect of weight loss on liquid-Phase-Sintered silicon carbide, J. Am. Ceram. Soc. 80 (4) (1997) 1047–1052.
- [9] N. Ordás, A. Bereciartu, C. García-Rosales, A. Morono, M. Malo, E.R. Hodgson, J. Abellà, S. Colominas, L. Sedano, Testing of porous SiC with dense coating under relevant conditions for Flow Channel Insert application, Fusion Eng. Des. 89 (7–8) (2014) 1274–1270
- [10] R.W. Rice, Evaluation and extension of physical property-porosity models based on minimum solid area, J. Mater. Sci. 31 (1) (1996) 102–118.
- [11] J. Magdeski, The porosity dependence of mechanical properties of sintered alumina, J. Univ. Chem. Technol. Metall. 45 (2) (2010) 143–148.