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Additional Information

1 Microstructure and mechanical properties of plasma spraying
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3 coatings from YSZ feedstocks comprising nano- and submicron-
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5 sized particles
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Abstract

Atmospheric plasma spraying (APS) is an attractive technique to obtain nanostructured coatings due to its versatility, simplicity and relatively low cost. However, nanoparticles can not be fed into the plasma using conventional feeding systems, due to their low mass and poor flowability, and must be adequately reconstituted into sprayable micrometric agglomerates.

In this work, nanostructured and submicron-nanostructured powders of yttria-stabilised zirconia (YSZ) were deposited using APS, with a view to obtaining high-performance thermal barrier coatings (TBC). All powders were reconstituted by spray-drying from different solid loading suspensions, followed by a thermal treatment of the spray-dried granules to reduce agglomerate porosity and enhance powder sinterability. The reconstituted granules were characterised by XRD, SEM, pore sizing and flowability evaluation.

The reconstituted feedstocks were successfully deposited onto metallic substrates by APS. A metallic bond coat was sprayed between the substrate and the ceramic layer. The coating microstructure, characterised by SEM, was formed by partially melted zones, which retained the initial powder microstructure, embedded in a fully melted matrix which acts as a binder. It was shown that feedstock characteristics which in turn are very dependent of starting suspension characteristics, in particular agglomerate density and primary particle size, impact on coating microstructure (porosity and amount of partially-melted areas). For this reason mechanical properties of coatings are also strongly affected by feeding powder characteristics.

Key words: YSZ coatings; atmospheric plasma spraying; reconstituted powders; thermal barrier coatings

Introduction

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3 Thermal barrier coatings (TBC) protect metallic substrates against exhaustive hot gases.
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5 Nowadays these materials are widely studied owing to their application on aeronautical
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7 or power technologies [1]. A top ceramic layer with thermal insulation properties and a
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9 metallic bond coat form these coatings. The bond coat usually comprises a MCrAlY (M
10 = Ni and/or Co) alloy which develops two main functions: reducing the thermal
11
12 contraction mismatch between the ceramic layer and metallic substrate and acting like
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14 oxidation barrier. The most used material for ceramic layer is yttria-stabilised zirconia
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16 (YSZ) due to its low thermal conductivity, relatively high thermal contraction
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18 coefficient and good mechanical properties at high temperature [2,3].
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25 Atmospheric plasma spraying (APS) is an economical and versatile technique to obtain
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27 TBC. It consists on a plasma source where a feed material is melted and accelerated
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29 until it impacts upon substrate. On the substrate, molten (or partially molten) material is
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31 quickly cooled forming the coating [4]. In the last years, coatings from nanoparticles
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33 have been extensively addressed because different architectures as well as enhanced
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35 properties, compared with their convectional counterparts (from microstructured
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37 feedstocks) can be obtained. In the case of TBC, thermal conductivity can significantly
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39 be reduced and the thermal shock resistance can be ~~then~~ improved [5]. However,
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41 nanoparticles cannot be directly injected into the plasma plume due to their low specific
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43 weight and poor flowability. One possible solution for this problem is to reconstitute the
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45 nanoparticles into sprayable micrometric granules. The reconstitution process consists
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47 of spray drying of stable suspensions to obtain spherical free-flowing granules.
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54 Subsequently spray-dried powders are usually thermally treated in order to enhance the
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56 sinterability for the following, extremely fast sintering process in the plasma torch [6,7].
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59 Coatings from reconstituted, nanostructured feedstocks usually display a bimodal
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1 microstructure formed by partially melted agglomerates surrounded by fully melted
2 particles which act as binder [6,7].
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5 Although many research effort has been made to pass from the micro-scale to either the
6 nano- or the submicron-scale using both powders or liquids as feedstock [8,9], few
7 attempts have been made to use feedstocks comprising mixtures of different particle
8 size distributions, e.g. submicron- and nano-sized particles. The use of a bimodal
9 distribution of submicron-nano particles in the precursor suspension of the spray-dried
10 powder can give rise to significant benefits during the suspension processing, i.e higher
11 solids loading and lower viscosity leading to better properties in the resulting feedstock
12 agglomerates, such as higher agglomerate bulk density and improved powder
13 flowability [10,11]. However the relationship between these agglomerate feedstock
14 characteristics and the final coating properties has been scarcely treated in the thermal
15 spray literature despite the fact that this connection has been already proven in many
16 other shaping processes [12-14]. Thus, some previous research was found in which the
17 effect of agglomerate density and microstructure on TiO₂ coatings properties were
18 addressed [15,16]. In addition some recent papers with YSZ systems have shown that
19 agglomerate morphology and density can have an influence on coating microstructure
20 and properties [17,18]. In this respect some of the authors of this paper have recently
21 reported improved photocatalytic activity in APS TiO₂ coatings or enhanced mechanical
22 properties in Al₂O₃-TiO₂ coatings when a mixture of nano- and submicron-sized
23 particles in the spray-dried feedstock was used. Much higher density agglomerates were
24 obtained which led to better properties in the final coatings [10,11]. These works
25 reported that the microstructure and properties of plasma sprayed coatings not only
26 depend on various process parameters but also are markedly influenced by the
27 characteristics of the feedstock. Nevertheless this type of research trying to relate
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1 feedstocks characteristics with coating microstructure and properties when nano- or
2 submicron-structured feedstocks are used has not been addressed for YSZ coatings.
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5 In a previous research the authors focused on the preparation of different YSZ spray-
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7 dried feedstocks starting from nano- and submicron-sized powders [19]. The paper
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9 aimed at obtaining suitable YSZ feedstocks for APS process including the starting
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11 suspension preparation and characterisation, spray-drying of suspensions,
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13 characterisation of spray-dried powders and their thermal treatment to complete the
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15 aforementioned reconstitution process. Finally, some of these powders were used to
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17 demonstrate that this methodology managed to produce feedstocks with adequate
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19 characteristics to be used in APS process. Most importantly the paper showed that the
20
21 main feedstocks properties in terms of APS depositing process (agglomerate size and
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23 density as well as powder flowability) were very dependent of starting suspension
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25 characteristics.
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32 Following this previous research this paper tries to close the loop by demonstrating the
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34 influence of feedstocks characteristics on the resulting APS coatings. Thus powders
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36 obtained from suspensions with different solid loadings and particle size distributions
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38 (nano- and submicron-sized mixtures) were deposited by APS onto stainless steel
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40 substrates. In addition two commercial (micro- and nano-structured) YSZ feedstocks
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42 were also deposited for comparison purposes. Then coatings were microstructurally and
43
44 mechanically characterised. On the one hand, voids and partially melted areas were
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46 quantified by image analysis. On the other hand, microhardness and toughness were
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48 determined.
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3 Experimental procedure
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6 Feedstock preparation
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8 The following commercial YSZ powders were used as raw materials: a nano-sized
9 powder (5932HT, Nanostructured and Amorphous Materials Inc., USA) with a mean
10 particle size of 40 nm and a specific surface area of 25.1 m²/g; and a submicron-sized
11 powder (TZ-3YS, Tosoh Co., Japan) with a mean particle size of 400 nm and a specific
12 surface area of 6.8 m²/g. Figure 1 shows the particle size distribution of these two
13 powders obtained by dynamic light scattering (Zetasizer NanoZS, Malvern, UK) in the
14 case of nanoparticles and laser diffraction (Mastersizer 200G, Malvern, UK) for
15 submicron-sized particles. Both powders were made up of tetragonal phase while the
16 nanopowder contained some minor (not quantified) monoclinic phase. On the other
17 hand a commercial polyacrylic acid-based polyelectrolyte (DURAMAXTM D-3005,
18 Rohm & Haas/Dow Chemicals, USA) was used as dispersant for suspension preparation
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38 In the mentioned previous research the colloidal stability and rheological behaviour
39 were evaluated to obtain stable and well-dispersed suspensions [19]. From this study,
40 four aqueous YSZ suspensions with different solid loadings and different particle size
41 distributions were selected. These suspensions were then reconstituted into sprayable
42 granules by spray-drying (Mobile Minor, Gea Niro, Denmark) followed by thermal
43 treatment. This treatment was carried out in an electric kiln with a soaking time of 60
44 min, at 1000 °C for the powders obtained from nanoparticles (YnD and YnC), at 1200
45 °C for the powder obtained from submicron-sized particles (YsC), and at 1050 °C for
46 the powder obtained from bimodal (nano- and submicron-sized) particle size
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1 distribution (Y_{sn}C). These temperatures were chosen in the previous work with the aim
2 of obtaining denser granules but at the same time of preserving as much as possible the
3 nanostructure of the initial agglomerates [19]. Hence the spray dried powders were
4 thermally treated at temperatures in which the sintering degree was only incipient
5 (contraction value of around 1%).
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11 Additionally two commercial YSZ feedstocks were used. One of them is a hollow-
12 sphere (HOSP) micrometric powder (METCO 204NS, Sulzer-Metco, Germany) and the
13 other one is a micrometric, nanostructured powder (Nanox S4007, Inframat Corp.,
14 USA). Hereafter these powders are referenced as YR_m and YR_n respectively. Further
15 information and micrographs of these two powders can be found elsewhere [21]. Due to
16 the preparation routes followed to obtain these commercial powders YR_m agglomerates
17 were much more sintered than YR_n agglomerates which were highly porous. All the
18 feedstocks used in this work are described in table 1.
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31 Feedstock characterisation

32 Feedstock powders were characterised before APS deposition. Field emission scanning
33 electron microscope (FEG-SEM, S-4800, Hitachi, Japan) was used to analyse feedstock
34 microstructure and the crystalline phases were determined by XRD (D8 Advance,
35 Bruker AXS, Germany). Moreover, granule size distribution was measured by sieving.
36 Granule apparent specific mass (ρ_{granule}) was calculated from tapped powder specific
37 mass by assuming a theoretical packing factor of 0.6, which is characteristic of
38 monosized, spherical particles [22]. Powder flowability was evaluated in terms of the
39 Hausner ratio, which is the ratio of the tapped powder specific mass to the powder
40 apparent specific mass. Free-flowing powders display a Hausner ratio below 1.25 [22].
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3 Coating deposition
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5 YSZ coatings were deposited by an atmospheric plasma spray (APS) system. It
6 consisted of a gun (F4-MB, Sulzer Metco, Germany) operated by an industrial robot
7 (IRB 1400, ABB, Switzerland). Before spraying, the substrate was grit blasted with
8 corundum at a pressure of 4.2 bars and cleaned with ethanol to remove any remaining
9 dust or grease from the surface. A NiCrAlY bond coat was used to enhance the
10 adhesion between the substrate and the ceramic layer. Deposition was performed using
11 argon and hydrogen as plasma-forming gases. The same spraying conditions were used
12 for all YSZ depositions: Ar flow=35 slpm, H₂ flow=12 slpm, arc intensity=600 A,
13 spraying distance=0.1 m, spraying velocity=1 m/s (slpm: standard litre per minute). The
14 necessary number of spraying passes was carried out to obtain a ceramic layer thickness
15 of 150 µm.
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36 Coating characterisation
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38 First, coating microstructures were observed by FEG-SEM. Voids and partially melted
39 areas of the coatings were evaluated by image analysis from 10 micrographs at 500x
40 magnification. Averages values were then calculated.
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46 Coatings were also mechanically characterised. The coating microhardness and fracture
47 toughness were determined by microhardness tester (LECO M400, Leco Co., USA)
48 performing 10 indentations for each coating to obtain an average value. 50 g indentation
49 load was used to determine the microhardness while 1000 g indentation load was used
50 for toughness assessment since a minimum load is required for the crack growth to
51 occur. Niihara equation was used for toughness calculation [23].
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Results and discussion

Feedstock characterisation

Morphology and structure of reconstituted powders (spray-dried+thermally treated) in this work are shown in figure 2. As it can be observed the powders contain spherical agglomerates displaying the typical doughnut-shape morphology of spray-dried agglomerates [24,25]. Most of granule sizes range from approximately 10 μm to 200 μm . The presence of some satellite-like granules stuck to bigger ones is also observed. Granules from diluted suspension of nanoparticles (YnD) seem to be smaller and their surface is quite rough due to the low viscosity of the starting suspension as set out in table 1. This finding coincides with that observed for nanoparticle suspensions of TiO_2 or Al_2O_3 - TiO_2 mixtures obtained with different solids content [24,25]. On contrary the powder obtained from the high solids content suspension of nanoparticles (YnC) is made up of coarser granules containing large inner holes. This behaviour is due to the high viscosity of the starting suspension of this sample as indicated in table 1 [26]. On the other hand, higher magnification of the solid areas of the agglomerates of all the powders reveals that the granules are porous and formed by the agglomeration of the individual nano- and submicron-sized particles. On comparing nanostructured agglomerates (YnD and YnC) it can be observed that as the solid loading in the starting suspension rises the resulting agglomerates are denser [25]. Also when a bimodal mixture is used (YsnC versus YnC) i.e when submicron-sized particles are added to a nanoparticle suspension the resulting agglomerates are also denser due to enhanced particle packing effect [11]. Finally particle necking and coarsening is quite evident in all the samples as a consequence of the thermal treatment. As expected samples containing submicron-sized particles gave rise to coarser grains after the thermal treatment.

1 With regard to commercial powders, the characteristics of both samples have been
2 reported elsewhere [21]. YRm powder consists of hollow spherical (HOSP) granules
3 obtained by spray-drying and subsequent plasma-densifying process while YRn powder
4 contains spherical, highly porous agglomerates formed by individual nanoparticles with
5 diameters varying from approximately 30 to 130 nm.
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11 Figure 3 shows the size distribution of the agglomerates comprising all the feedstocks.
12 Although the curves represent sizes of thermal treated agglomerates it must be noticed
13 that these sizes should practically coincide with those of the thermally untreated
14 powders since the agglomerate shrinkage was, in all the samples, lower than 1% as
15 mentioned above. As observed all the distributions display a monomodal pattern,
16 significant differences among them appear though. The curves also confirm the
17 micrometer size range of the spray-dried agglomerates. With regard to commercial
18 samples, YRn nanostructured powder comprises agglomerates coarser than those of the
19 reconstituted powders while granules of conventional YRm powder were slightly finer
20 than those of the reconstituted powders. Among powders reconstituted in this work the
21 graph shows that the higher the viscosity of the starting suspension (see table 1) the
22 coarser the agglomerates confirming the aspect of the agglomerates displayed in the
23 micrographs in figure 1. The correlation between starting suspension viscosity and
24 agglomerate average size calculated from figure 3 data is plotted in figure 4. As it can
25 be seen a good fit between both variables is obtained confirming the findings stated
26 above.
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51 To further characterise the agglomerate structure in all the samples mercury pore sizing
52 was determined (figure 5). Although this technique determines any type of pore volume
53 trapped in a powder bed, i.e. inter and intragranular porosity, in these curves only
54 intragranular porosity is displayed owing to the scarce interest of the porosity trapped
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1 between agglomerates when the performance of the agglomerates inside the plasma
2 torch in an APS process is to be analysed. Focusing on the reconstituted powders, YnD
3 and YnC display a very similar intragranular porosity. However when submicron-sized
4 particles are mixed with nanoparticles (sample YsnC) the resulting agglomerates are
5 slightly less porous but more importantly display larger pore sizes as a consequence of
6 the increased particle size inside the agglomerates. This same trend is observed when
7 the amount of submicron-sized particles is increased up to 100% (YsC sample). These
8 findings show that granule porosity and pore size basically depends on the particle size
9 distribution of the solids comprising the agglomerates. Thus the wider particle size
10 distribution of the submicron-sized powder (see figure 1) when compared with the
11 distribution of the nanosized powder results in denser particle packing inside the
12 agglomerate. These findings agree with theoretical models on particle packing of
13 continuous particle size distributions [27]. YsnC sample porosity lies between YnC and
14 YsC porosity values. Regarding commercial samples, low porosity in YRm powder
15 curve relates to the HOSP nature of this feedstock (the large inner hole of the HOSP
16 granule is not considered in the pore sizing curve) while YRn sample exhibits its high
17 porosity agglomerate characteristic.

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41 Figure 6 shows Hausner ratio and agglomerate apparent specific mass of the feedstocks.
42 All powders present a Hausner ratio lower than 1.25 therefore they display a free-
43 flowing nature. Even though little differences on flowability were observed among the
44 different feedstocks, YnD sample shows the worst flowability (the highest Hausner
45 ratio) as a consequence of a combined effect of small agglomerate size and high surface
46 roughness. Regarding granule apparent specific mass, significant differences were
47 found between the reconstituted powders as expected from the pore sizing assessment.
48 To better visualise these differences, figure 7 plots the variation of agglomerate
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apparent specific mass of the four reconstituted powders versus the amount of
submicron-sized particles in the starting suspension. As it can be observed, a clear,
linear relationship between both variables has been found confirming the findings
observed in the porosimetry analysis. The higher the amount of submicron-sized
particles in the starting suspension the higher the agglomerate density is. Figure 8
confirms the good agreement between the values of agglomerate apparent specific mass
calculated from the powder tapped specific mass as set out above and those obtained
from mercury pore sizing curves (figure 5). In respect of commercial feedstocks as
expected agglomerate apparent specific mass of YRm (HOSP) agglomerate is much
higher than that of the YRn agglomerate. Finally it is important to note that all the
powders met the recommendations to be deposited by APS (Hausner ratio lower than
1.25, granule apparent specific mass higher than 1700 kg/m^3 and mean granule size
higher than $20 \text{ }\mu\text{m}$) [4,7].

31 32 33 Coating microstructure

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Commercial and reconstituted feedstocks were deposited by APS. Figure 9 shows the
cross-section microstructure of the resultant coatings. As observed all the coatings were
porous, exhibiting the typical lamellar structure of plasma sprayed coatings. This
microstructure comprises smaller pores located within individual lamellae and larger
pores trapped along the interlamellar boundaries. Figure 9 also reveals that coatings
microstructure is influenced by the feedstock characteristics. Hence layers deposited
from the nanostructured feedstocks (YnD and YnC) exhibit a bimodal microstructure
formed by partially melted agglomerates (marked PM in the micrographs) that retain
some of the initial nanostructure, surrounded by a fully melted matrix (marked M in the
micrographs). Such microstructure has been extensively reported in literature [6-8] in

1 which details on the features of these partially-melted areas can be found. Thus despite
2 the nanostructured character of the primary particles comprising the feedstock, the
3 relatively high porosity associated to these agglomerates results in coatings containing
4 partially melted areas which preserve in some extent the starting nanostructure. This
5 statement is particularly evident in the commercial, nanostructured powder which
6 showed the highest agglomerate porosity (figure 5) giving rise to much higher amount
7 of partially-melted areas as observed in the corresponding micrograph of figure 9.
8
9 Respecting coatings obtained from submicron-sized particles ($Y_{sn}C$ and Y_sC
10 feedstocks) higher amount of partially-melted areas when compared with coatings
11 obtained from nanostructured feedstocks (Y_nD and Y_nC) are observed. These findings
12 seem to indicate that for similar values of agglomerate porosity feedstocks containing
13 coarser particles (submicron-sized) lead to higher amount of partially-melted areas in
14 the coatings provided that energy conditions during the plasma spray are also similar.
15
16 These findings have been previously observed during deposition by APS of nano- and
17 submicron-sized titania particles [11]. The different features of these partially-melted
18 zones can be observed in figure 10. As expected nanoparticles were retained in the
19 partially melted zones of Y_nD and Y_nC coatings but some of them were partially
20 sintered so that micrometric voids and coarser grains were also found. On the other
21 hand partially melted zones of Y_sC coatings exhibit the initial agglomerate
22 microstructure. Finally, coating obtained from the commercial, microstructured powder
23 (YR_m sample) presents a quite homogeneous microstructure containing low amount of
24 partially-melted areas what agrees with the highly sintered microstructure of the HOSP
25 agglomerates.
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28 In view of the above, the total porosity of the coatings as well as the amount of
29 partially-melted areas were estimated by image analysis at 500 magnifications from
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SEM pictures following a procedure set out elsewhere [11,25]. Figure 11 shows the porosity and partially-melted areas values obtained for all the coatings. Porosity data agree with the typical values observed in these types of coatings. Besides, in all the coatings reconstituted in this work the porosity values were very similar. This is because, at this low range of coating porosities the contribution of the agglomerate density to the final coating porosity is negligible as compared with the contribution coming from the melting and subsequent deformation of the agglomerates during the deposition process. On contrary larger differences among the coatings were observed for the values of the partially-melted areas content. Coatings obtained from feedstocks containing submicron-sized particles (YsC and YnsC) display higher amount of partially-melted areas in comparison with the feedstocks exclusively formed by nanoparticles (YnD and YnC). Thus for a given plasma spray conditions (plasma energy) the amount and the size of the particles inside these partially-melted zones obviously depend on the size of the particles comprising the feedstock agglomerates (nanoparticles or submicron- sized particles) but also on other feedstock characteristics such as agglomerate size and density. As the agglomerate size and porosity of the reconstituted feedstocks are not so different the findings obtained highlight the role of the particle size in reconstituted, spray-drying agglomerated powders with regard to the appearance of the always existing partially-melted areas. Finally, the quantified amount of partially-melted areas for the coatings obtained from the commercial feedstocks (YRm and YRn) agrees well with the appreciation deduced from the corresponding micrographs in figure 9.

Coating properties

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3 Figure 12 displays microhardness and toughness of the coatings obtained from the
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5 different feedstocks. Coatings from commercial powders exhibit lower values of
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7 microhardness, in particular YRn coating which presented the highest value of porosity
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9 as set out above. Thus the negative effect of porosity on microhardness can easily be
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11 noted in this coating in which the porosity value almost doubles those of the rest of
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13 coatings. This effect has been extensively reported in APS literature [28,29]. In less
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15 extent, the effect of porosity can be also observed in the other commercial coating
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17 (YRm) which also shows lower microhardness due to its relatively high porosity. As
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19 regards the coatings obtained from the reconstituted feedstocks, higher values of
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21 microhardness were found on comparing with the coatings obtained from the
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23 commercial feedstocks owing to their lower porosity values. Nevertheless no clear
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25 relationship between porosity and microhardness were observed for this group of
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27 coatings. This is because differences in porosity were too small for the correlation to be
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29 established (see figure 11). On the other hand literature on coatings obtained from
30
31 nanostructured feedstocks reports that the presence of mechanically weak partially-
32
33 molten zones featuring these coatings can result in mechanical properties decrease [30].
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35 Hence, although this effect can contribute to the low hardness of YRn coating, it has not
36
37 been observed in the case of YsC and YsnC coatings despite the fact that these coatings
38
39 contained higher amount of partially-melted areas than YnD and YnC coatings. This
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41 discrepancy with the reported data can be caused by the higher density of the partially-
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43 melted areas in coatings which have been obtained from feedstocks containing
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45 submicron-sized particles (YsC and YsnC coatings) as observed in figure 10.
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57 Finally, according to literature, partially melted zones present in coatings obtained from
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59 nanostructured feedstocks can behave as barriers for crack propagation [5,7]. However
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1 few changes in terms of toughness were found between conventional coating (YRm)
2 and coatings from nanostructured feedstocks (YnC, YnD and ~~YRe~~). It is probably
3 because of the poor cohesion presented by the partially melted zones of these coatings.
4
5 On contrary, the higher toughness value for YsC coating can be attributed to its larger
6
7 amount of denser partially molten zones as set out above.
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10 11 12 13 14 15 **Conclusions**

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18 In this work, YSZ coatings were obtained from different feedstocks by APS. On the one
19
20 hand four reconstituted powders from nano- and submicron-sized particles were
21
22 deposited. The reconstitution process consisted in spray-drying of the starting
23
24 suspension followed by a thermal treatment of the spray-dried agglomerates. In
25
26 addition, two commercial powders (one nanostructured and the other one conventional)
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28 were also deposited to compare differences between the final coatings. It was proven
29
30 that suspension characteristics strongly affect the spray-dried agglomerate properties.
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32 Thus the higher the suspension viscosity the coarser the spray-dry agglomerate size
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34 whereas the agglomerate density is directly related to the amount of submicron-sized
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36 particles in the starting suspension. All the reconstituted feedstocks showed adequate
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38 properties to be used in an APS process.
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46 The coatings obtained displayed a bimodal microstructure formed by partially melted
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48 zones surrounded by a fully melted matrix. Coating porosity and the amount of partially
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50 melted areas depend on the feedstock characteristics. As the coating microstructure
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52 (porosity and amount of partially melted areas) strongly defines the mechanical
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54 properties of the final coatings this work has proven the connection between the
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56 mechanical properties of the coatings and the characteristics of the feedstock
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1 agglomerates. On the one hand, coatings obtained from the reconstituted feedstocks
2 gave rise to higher values of microhardness when compared with the coatings obtained
3 from the commercial feedstocks owing to the lower porosity of the reconstituted
4 powders. On the other hand, the effect of the characteristics of the partially melted areas
5 on microhardness has been also put forward. Nevertheless the effect of these partially-
6 melted areas on microhardness is not so impacting as that of coating porosity. Finally
7 the reinforcing role of the amount of partially-melted areas on the coating toughness as
8 reported in the literature has not been observed in this work. In fact no differences in
9 toughness were found between the conventional coating which did not contain any
10 partially-melted areas and the coatings obtained from reconstituted feedstocks which
11 presented different amounts of these areas.
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Figure captions

1
2
3 Figure 1. Particle size distribution of the commercial nano- (n) and submicron-sized (s)
4
5 particles

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7
8 Figure 2. FEG-SEM micrographs at different magnifications of reconstituted feedstocks
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10 (thermally treated spray-dried powders): a) YnD; b) YnC; c) YsC; d) YsnC

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12
13 Figure 3. Agglomerate size distributions of the feedstocks

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16 Figure 4. Correlation between starting suspension viscosity displayed in table 1 and
17
18 agglomerate average size worked out from figure 3

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21 Figure 5. Pore size distribution of the feedstocks obtained by mercury intrusion
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23 porosimetry. Only intragranular porosity is taken into account

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26 Figure 6. Hausner ratio and agglomerate apparent specific mass ($\rho_{\text{agglomerate}}$) of the
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28 feedstocks

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31 Figure 7. Variation of agglomerate apparent specific mass ($\rho_{\text{agglomerate}}$) versus the
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33 amount of submicron-sized particles in the starting suspension

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36 Figure 8. Relation between agglomerate apparent specific mass of the six powder
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38 feedstocks as obtained from the powder tapped specific mass and mercury pore sizing
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40 procedures

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43 Figure 9. Coating cross section micrographs at 500x magnification. Melted and
44
45 partially-melted areas are referenced as M and PM respectively

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48 Figure 10. Partially melted zone detail of different coatings

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51 Figure 11. Porosity and partially melted areas of the coatings as determined by SEM

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54 Figure 12. Microhardness and toughness of the different coatings

Table caption

Table 1. Reference and description of all the YSZ feedstocks used through this work

Reference	Starting suspension			$T_{\text{treatment}}^3$ (°C)
	Solids	SL ¹ (vol.%)	μ^2 (mPa·s)	
YnD	100% nanoparticle	10	2.8	1000
YnC	100% nanoparticle	30	46.5	1000
YsC	100% submicron-sized particle	30	13.3	1200
YsnC	50% nano- + 50% submicron-sized particle	30	17.9	1150
YRm	Commercial (reference) microstructured powder			
YRn	Commercial (reference) nanostructured powder			

¹ SL: solid loading

² μ : suspension viscosity measured from downward flow curves at shear rate of 1000 s⁻¹ (reference 19).

³ $T_{\text{treatment}}$: temperature of the powder thermal treatment (reference 19).

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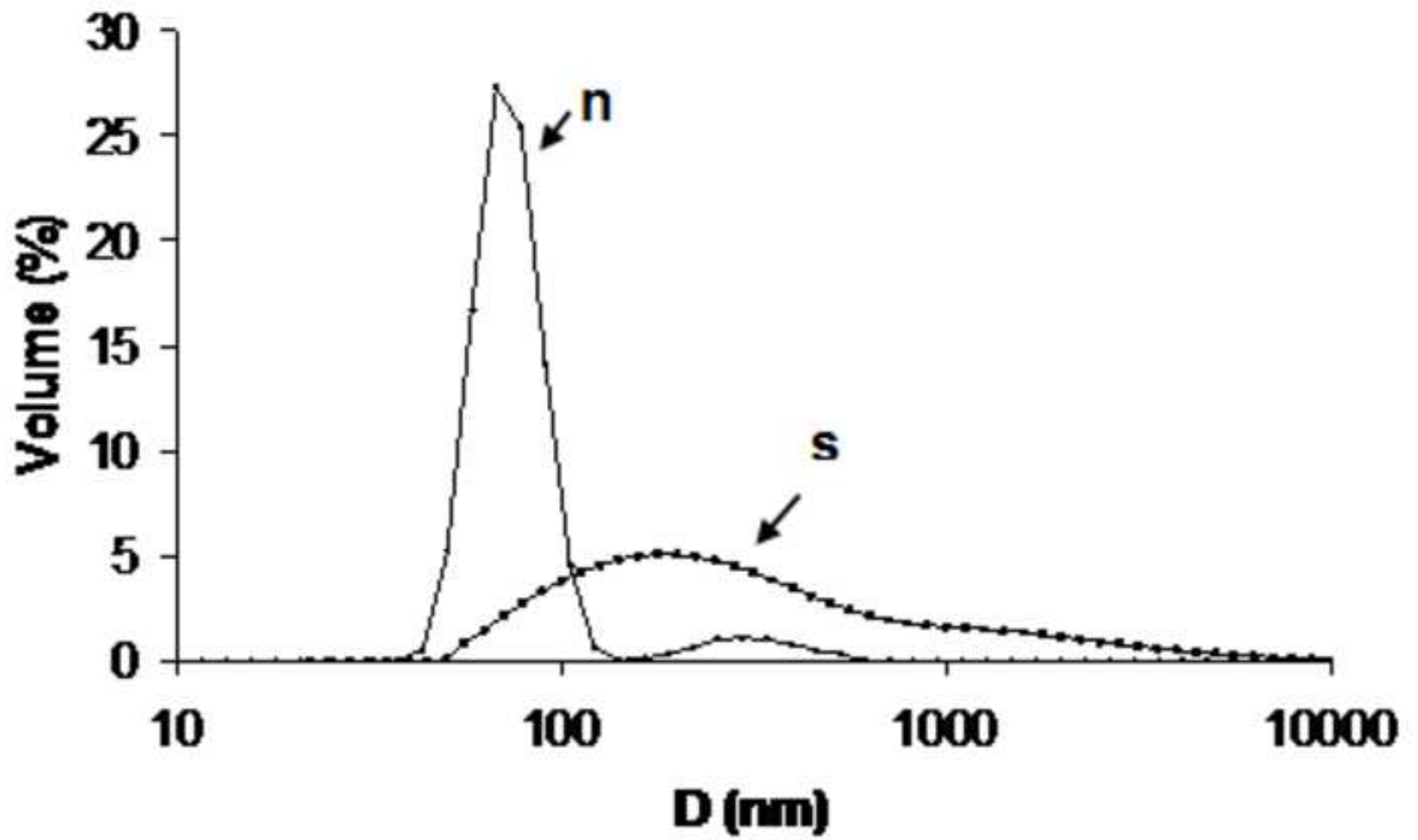


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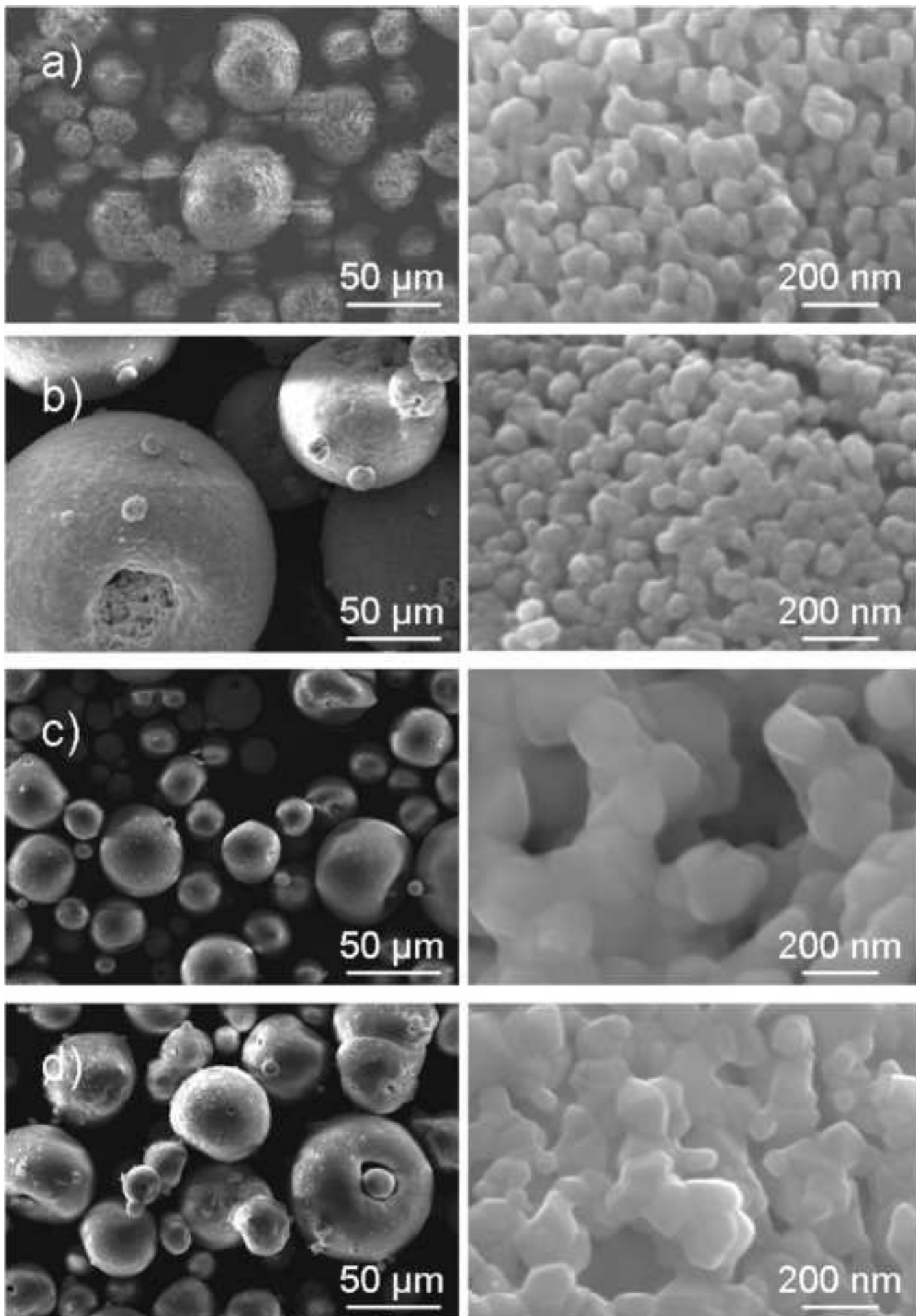


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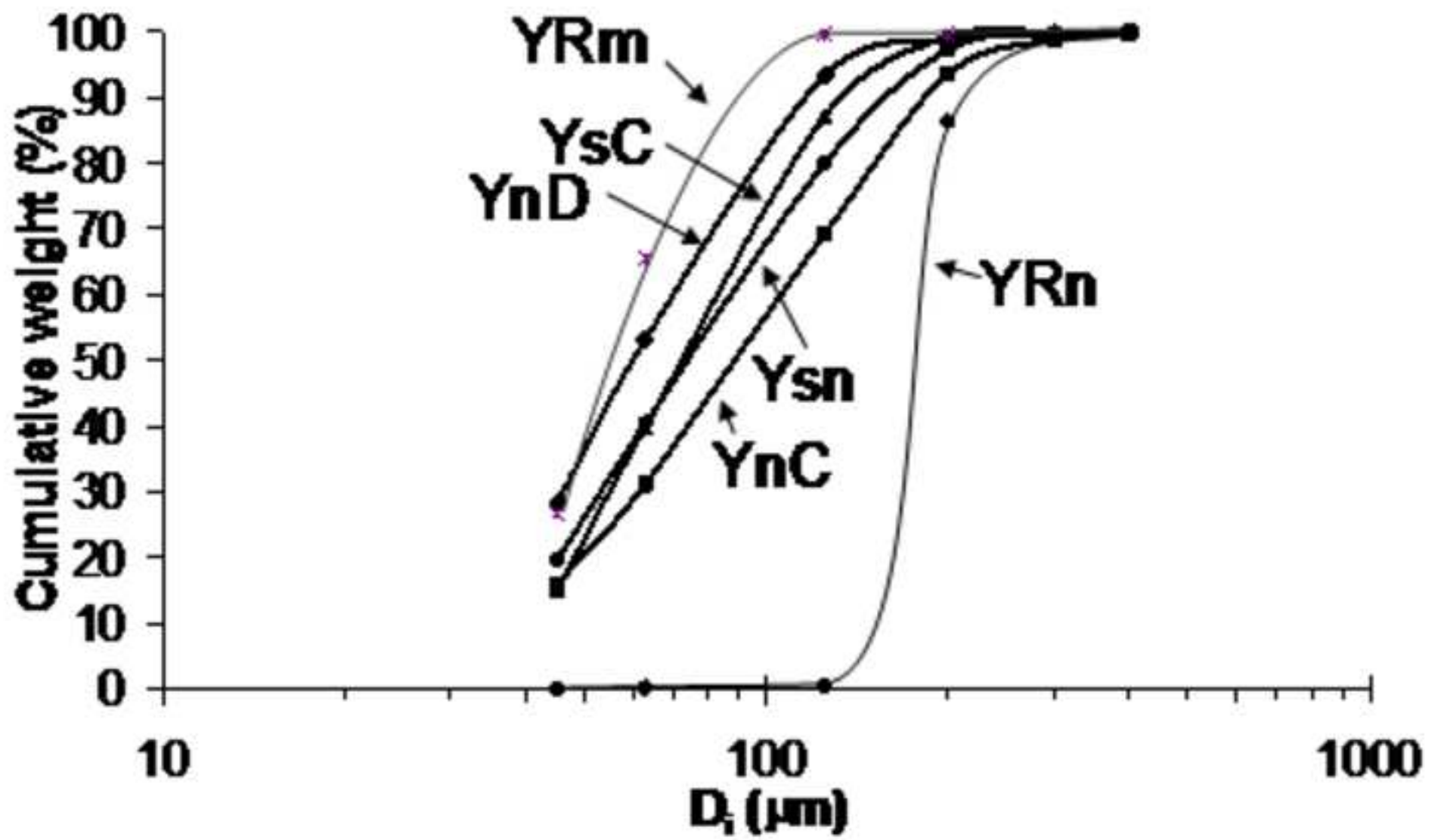


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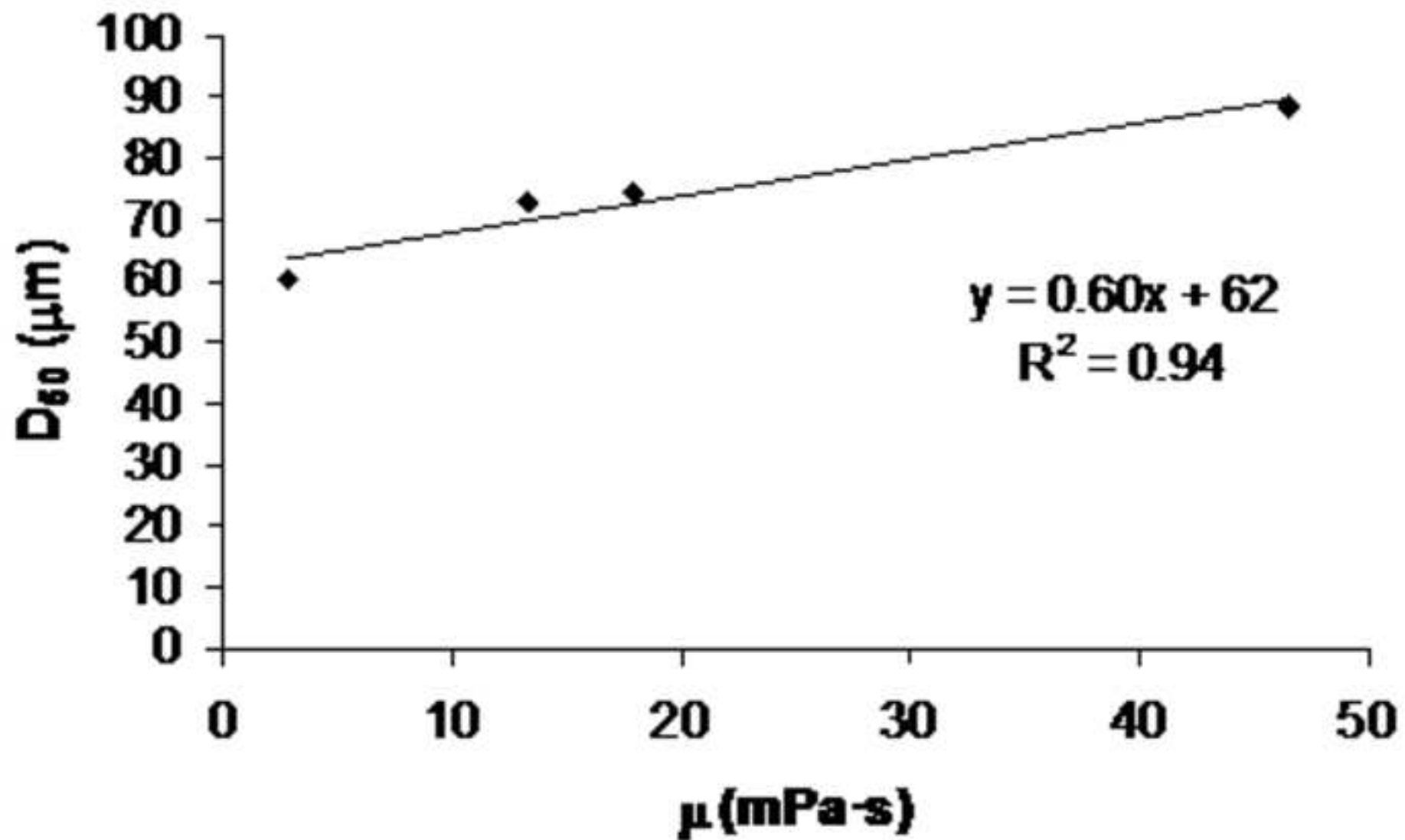


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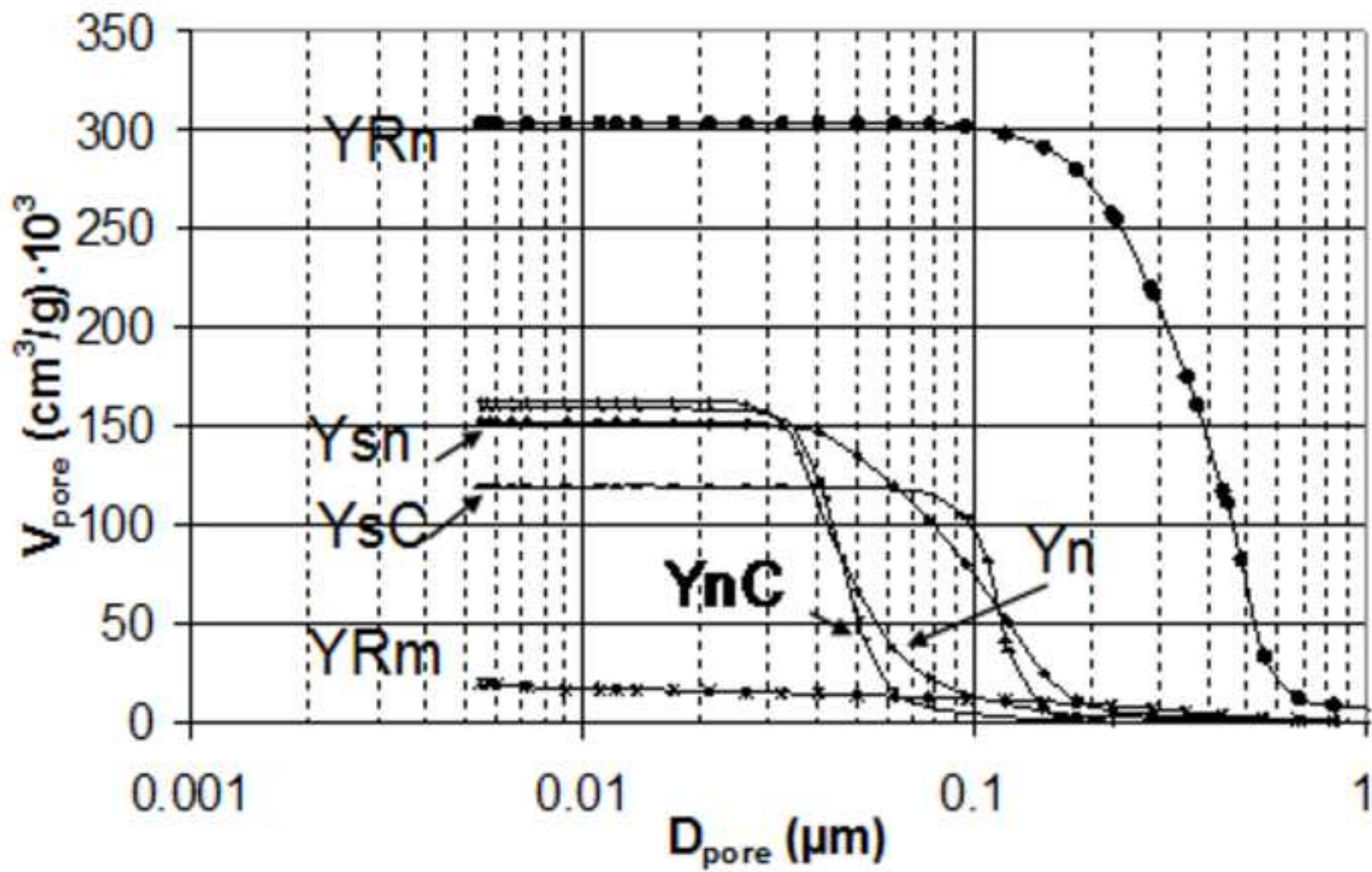


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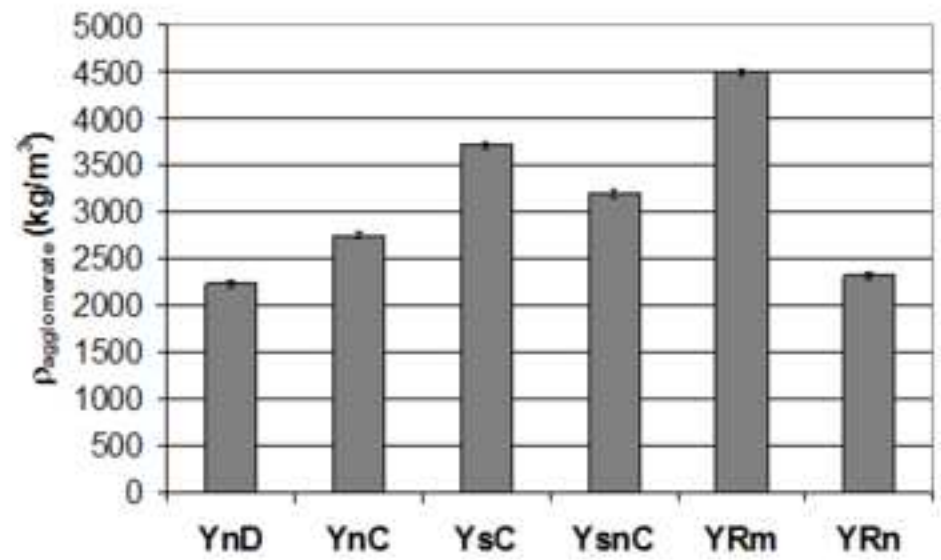
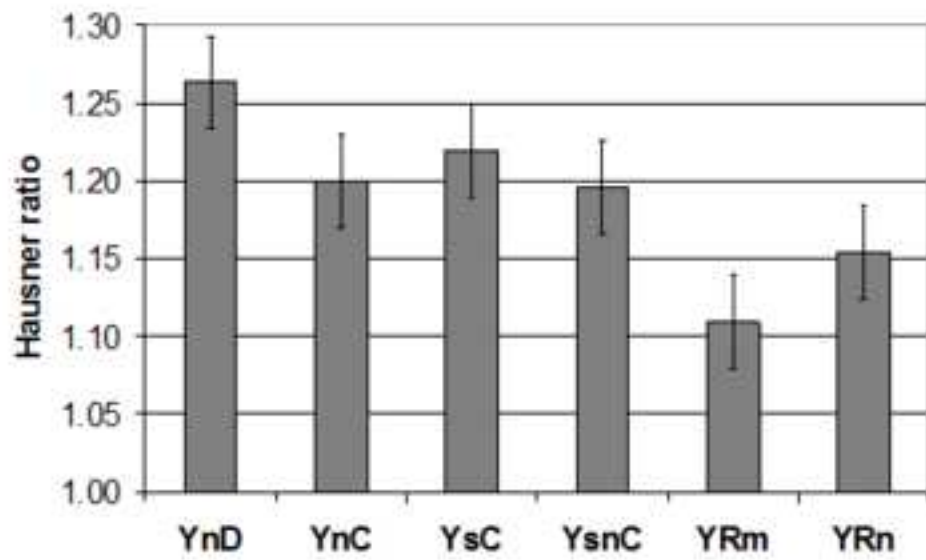


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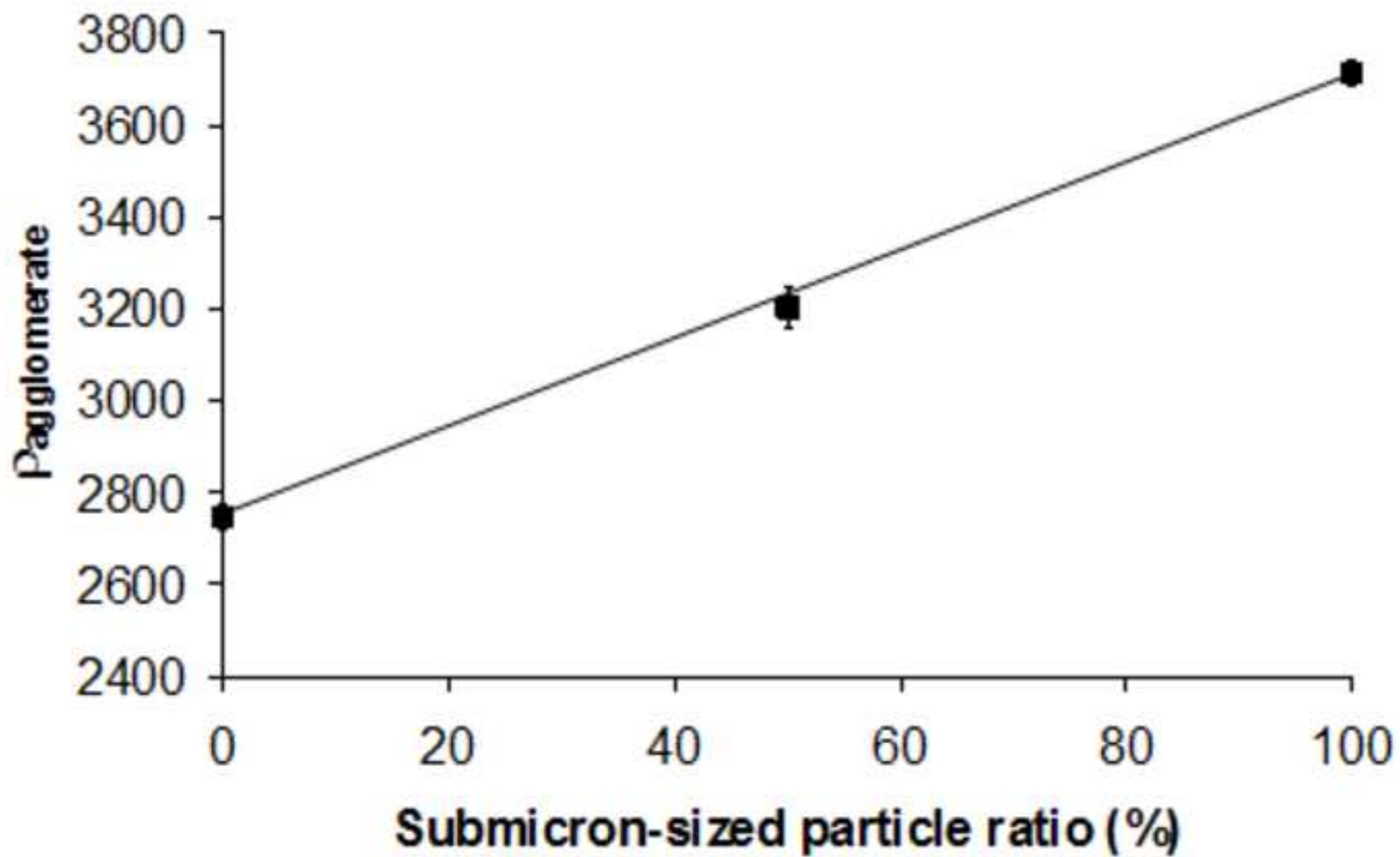


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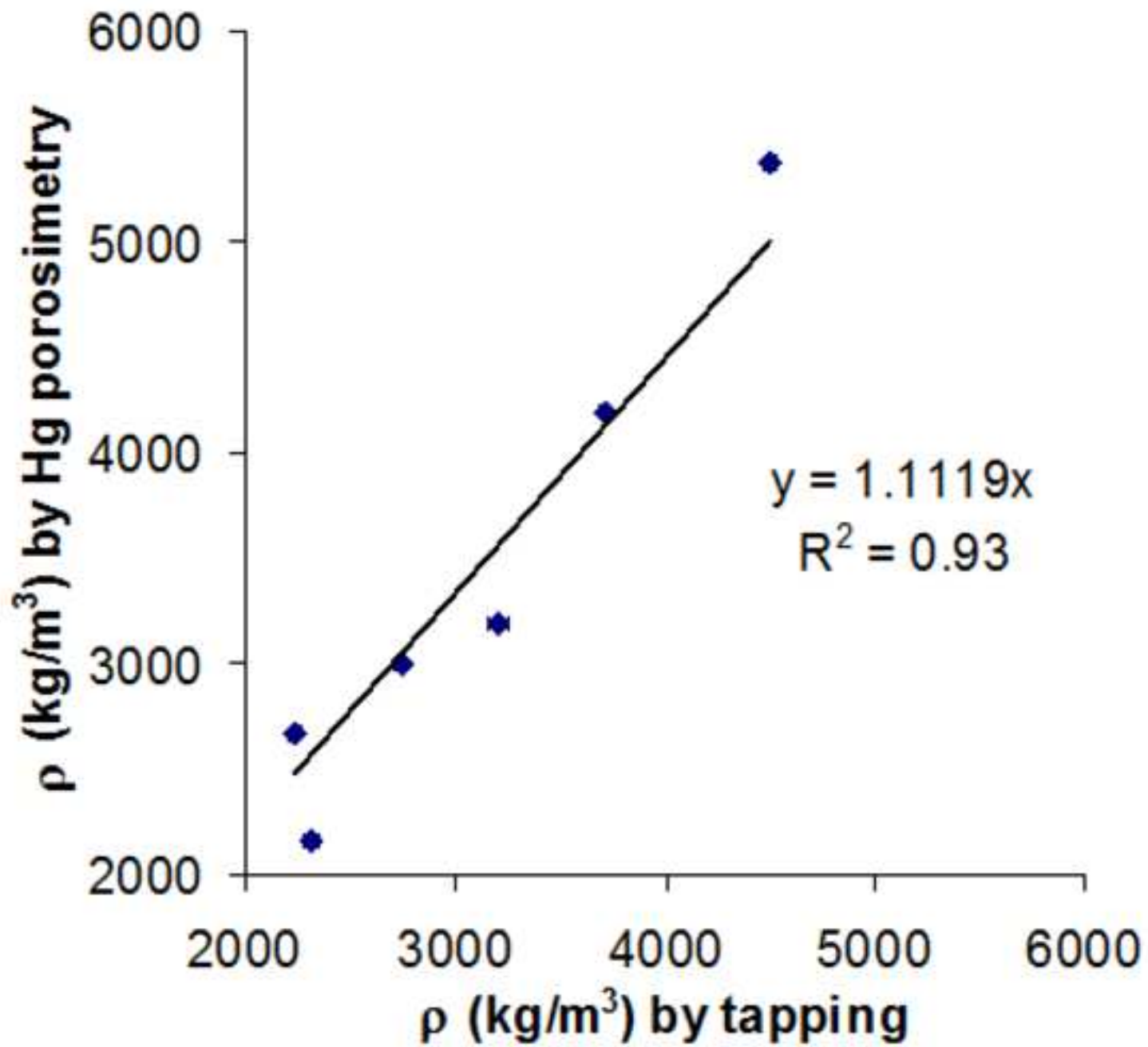


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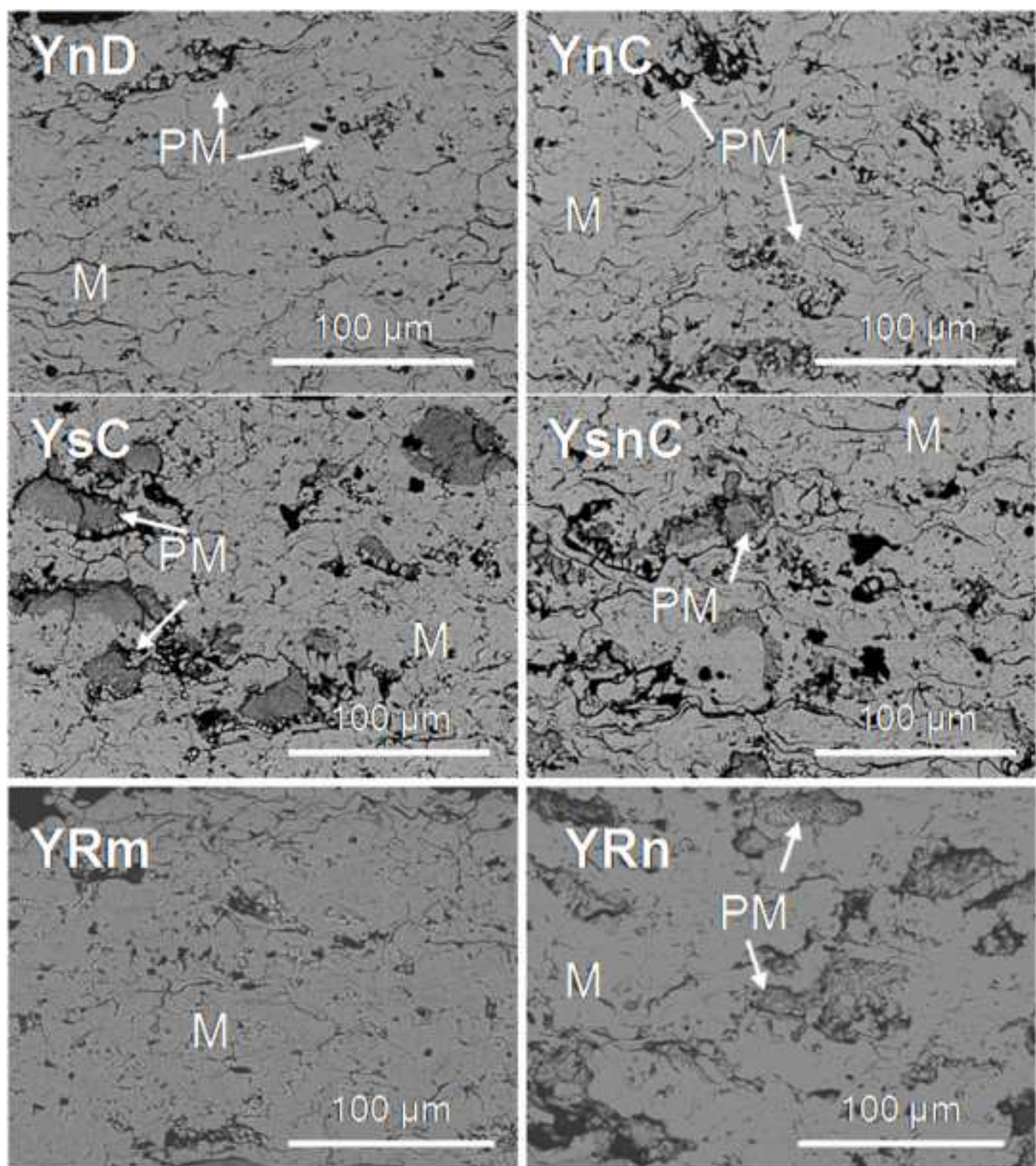


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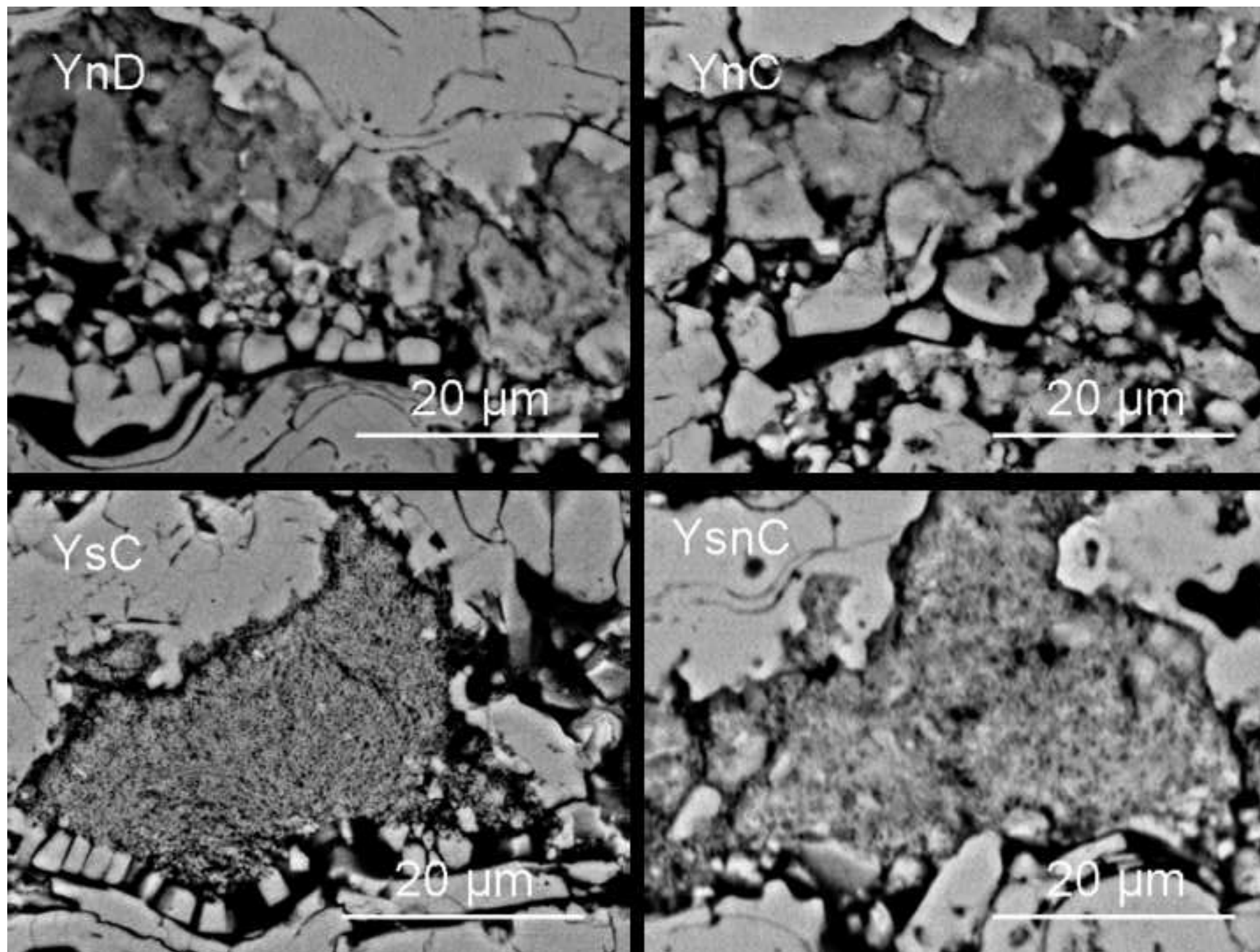


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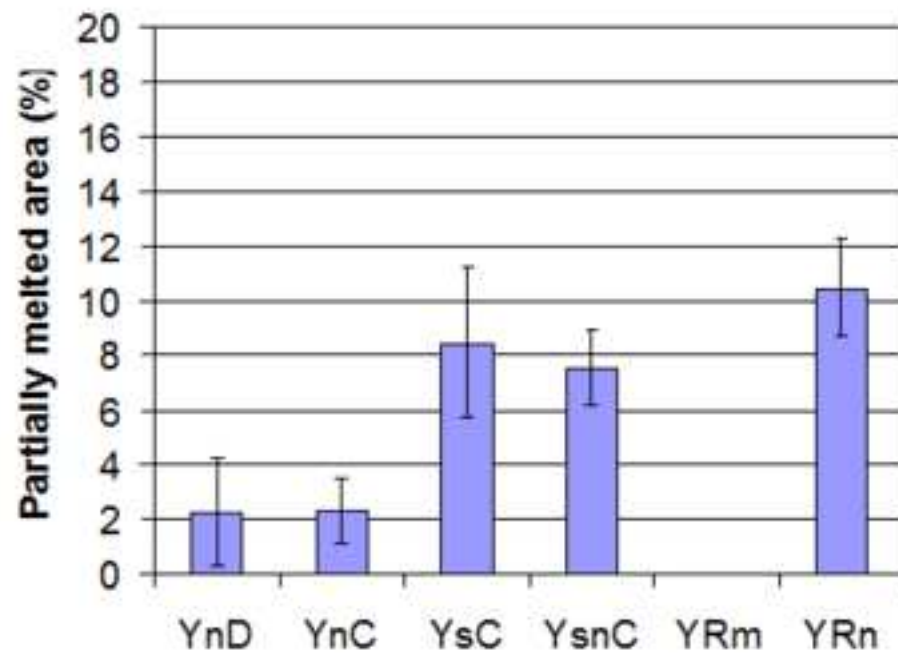
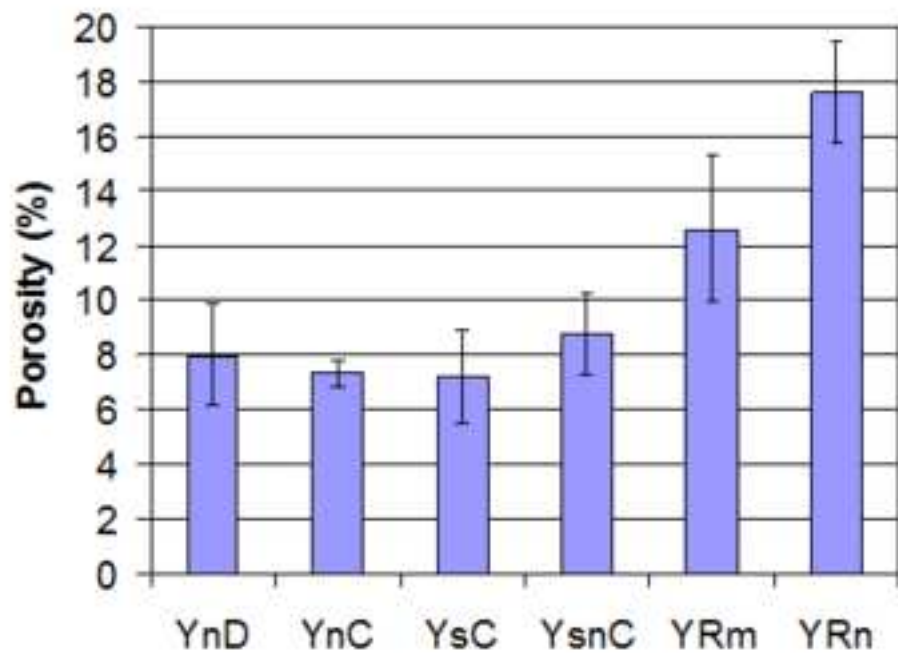


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