Amorphous mesostructured zirconia with high (hydro)thermal stability

Bénédicte Lebeau^{1,2}, Issam Naboulsi³, Laure Michelin^{1,2}, Claire Marichal^{1,2}, Séverinne Rigolet^{1,2}, Cédric Carteret⁴, Sylvette Brunet⁵, Magali Bonne^{1,2}, Jean-Luc Blin^{3*},

¹: Université de Haute Alsace (UHA), CNRS, IS2M UMR 7361, F-68100 Mulhouse, France

²: Université de Strasbourg, 67000 Strasbourg, France

³: Université de Lorraine, CNRS, L2CM, F-54000 Nancy, France

⁴: Université de Lorraine, CNRS, LCPME, F-54000 Nancy, France

⁵: Université de Poitiers, CNRS, IC2MP, UMR 7285, 86073 Poitiers Cedex 9 France.

* Corresponding authors :
Pr. Jean-Luc Blin
Université de Lorraine
L2CM 7053
Faculté des Sciences et Technologies
BP 70239
F-54506 Vandoeuvre-lès-Nancy cedex, France
Tel. +33 3 83 68 43 70
E-mail: Jean-Luc.Blin@univ-lorraine.fr

Supporting information S1: TG and heat-flow curves of the hybrid mesophase and EtOH-extracted mesostructured ZrO_2



Supporting information S2: ${}^{1}H{}^{-13}C$ CPMAS NMR spectra of as-synthesized (A) and dehydrated at 70 °C (B) ZrO₂ hybrid mesophase



Supporting information S3: Raman spectra of Pluronic P123 (A) and ZrO_2 hybrid mesophase (B)



Supporting information S4: ¹H MAS NMR spectra of as-synthesized (A) and dehydrated at 70 $^{\circ}$ C (B) ZrO₂ hybrid mesophase



Supporting information S5: Raman spectrum of amorphous mesostructured ZrO₂ recovered after surfactant extraction.



Supporting information S6: SAXS pattern (A), nitrogen adsorption-desorption isotherm (B) and mesopores size distribution (C) of ZrO_2 after calcination at 480 °C under air atmosphere in a furnace.



Supporting information S7: Evolution as a function of the immersion time of the specific surface area (\blacksquare) and pore volume (\bigcirc) of amorphous ZrO₂ after calcination at 440 °C under air atmosphere in a furnace. Lines are just a guide for the eyes.

