

Cite this: *J. Mater. Chem. A*, 2017, 5, 20024

The water gas shift reaction over Pt–CeO₂ nanoparticles confined within mesoporous SBA-16†

D. Carta,^{ab} T. Montini,^c M. F. Casula,^{bd} M. Monai,^c S. Bullita,^b P. Fornasiero^{cd} and A. Corrias^{bd}

Novel nanocomposite catalysts for the single step Water Gas Shift Reaction (WGSR) were prepared by deposition–precipitation and impregnation of Pt–CeO₂ nanophases onto an ordered mesoporous silica support featuring a cubic arrangement of mesopores (SBA-16 type). The highly interconnected porosity of the SBA-16 developed in three dimensions (3D) provides a scaffold which is easily accessible to reactants and products by diffusion. The textural and morphological properties of the final catalyst were affected by the procedure utilized for dispersion of the nanophases onto SBA-16. The catalysts prepared by deposition–precipitation present highly dispersed nanocrystalline CeO₂ on the surface of SBA-16 and retain a high surface area, high thermal stability and high Pt accessibility. The catalysts prepared by impregnation show improved Pt–CeO₂ interaction but a more significant decrease of surface area compared to pure SBA-16, due to the confinement of the CeO₂ crystallites within the mesoporous matrix. As a result, the catalysts prepared by deposition–precipitation are effective for the WGSR under working conditions in the high temperature range (around 300–350 °C), whereas the catalysts prepared by impregnation are suitable for the process operating at low temperature. Our results point out that these catalyst preparation procedures can be used to optimise the performance of heterogenous catalysts, by controlling the CeO₂ crystallite size and optimizing the Pt–CeO₂ contact by embedding. Improved thermal and chemical stabilities were achieved using a mesoporous scaffold.

Received 27th April 2017
Accepted 24th August 2017

DOI: 10.1039/c7ta03640j

rsc.li/materials-a

1. Introduction

The water gas shift reaction (WGSR):



is a key process in the modern chemical industry, since it allows controlling the H₂/CO_x ratio and the CO_x composition starting from syn-gas produced by reforming processes. A low H₂/CO ratio is required for the Fischer–Tropsch synthesis of liquid fuels and for hydroformylation reactions, while H₂/CO/CO₂ mixtures are required for methanol synthesis. On the other hand, applications that use large amounts of pure H₂, such as

NH₃ synthesis, oil refining and hydrogenation processes involved in fine chemical production, require very high starting H₂/CO₂ ratios in order to effectively remove CO₂ by absorption into basic solutions. In the last few decades, H₂ production attracted great attention for the development of upgrading processes in biorefineries (*e.g.* hydrodeoxygenation) and in view of the possible use of large amounts of H₂ as the energy vector, in combination with the fuel cell technology. In the last case, the diffuse energy production employing Polymer Electrolyte Membrane Fuel Cells (PEMFCs) will require an extremely low CO content (usually below 10 ppm) to avoid the poisoning of Pt-based anode electrocatalysts.¹ It must be underlined that, the WGSR being an equilibrium reaction, the CO content at the exit of the WGSR reactors is still too high and a further purification process is required. This is the so-called PROX (PReferential OXidation of CO in the presence of H₂).

The WGSR is an exothermic reaction and therefore thermodynamically limited at high temperatures and kinetically limited at low temperatures. Because of this, the production of H₂ in industrial large-scale plants involves a two-step process: a high temperature (HT-WGSR) step (350–450 °C) in the presence of Fe₂O₃–Cr₂O₃ based catalysts and a low-temperature (LT-WGSR) step (180–250 °C) in the presence of Cu–ZnO catalysts. The conventional Cu–ZnO-based LT-WGSR catalysts

^aDepartment of Chemistry, University of Surrey, Guildford GU2 7XH, UK

^bDepartment of Chemical and Geological Sciences, INSTM, University of Cagliari, S.P. Monserrato-Sestu, Km 0.700, I-09042 Monserrato, Cagliari, Italy

^cDepartment of Chemical and Pharmaceutical Sciences, INSTM, ICCOM-CNR, University of Trieste, Via L. Giorgieri 1, 34127 Trieste, Italy

^dSchool of Physical Sciences, University of Kent, Ingram Building, Canterbury, CT2 7NH, UK. E-mail: a.corrias@kent.ac.uk

† Electronic supplementary information (ESI) available: Details of the synthesis of Pt and CeO₂ precursors by the impregnation method, the typical SAED pattern and additional SEM characterization, additional catalytic tests and characterization of the catalyst after the WGSR. See DOI: 10.1039/c7ta03640j



present severe limitations, such as low activity in the relevant fuel cell temperature range, pyrophoric nature and weak stability under cyclic operation.² Recently, research has focused on noble metals (Pt, Ph, Ru, Au, and Pd) supported on reducible oxides (CeO₂ and TiO₂) as highly active WGS catalysts.³ Among all the systems studied, Pt–CeO₂ is regarded as the most promising catalyst due to the ability of CeO₂ to undergo rapid (Ce(IV)/Ce(III)) reduction and oxidation cycles, further promoted by the addition of Pt on the CeO₂ surface, with consequent enhancement of the WGS activity, in particular, in the low temperature process.⁴ The redox and catalytic properties of ceria are also strongly dependent on the particle size, a decrease in particle size being associated with an increase of density of oxygen defects and hence an enhancement of the number of active sites for gas–solid catalysis.^{5,6} However, pure ceria has a poor thermal stability and undergoes sintering at high temperatures.^{7–9} To overcome this problem, CeO₂ is usually doped with another metal oxide (generally ZrO₂ or lanthanide oxides)⁵ or is dispersed on a high surface area thermally stable support (such as Al₂O₃).^{5,10,11} Recently, dispersion in a silica matrix with an ordered mesoporous structure and a hexagonal symmetry such as the MCM-41 type¹² and SBA-15 type^{6,12,13} has been proposed as a strategy to improve the thermal stability of nanocrystalline ceria. Silica-supported nanocrystalline ceria materials are of particular interest in catalysis, since the mesoporous silica supports provide thermal stability as well as a high surface area. So far, only ordered mesoporous silica supports with a hexagonal porous geometry have been used to stabilize CeO₂. In this work, we address the design of novel catalysts for the LT-WGS by dispersing Pt–CeO₂ nanophases onto an ordered mesoporous silica support with a cubic arrangement of mesopores (SBA-16 type) developed in three dimensions (3D). The highly interconnected pores provide the scaffold easy accessibility from all directions to reactants and products by diffusion.¹⁴ Additionally, the SBA-16 scaffold presents high thermal stability,¹⁵ thick walls and enhanced resistance to local pore blockage,¹⁶ making it very attractive for nanoparticle dispersion.¹⁷ It should be pointed out that to successfully produce the cubic arrangement of pores typical of SBA-16 type silicas, a careful adjustment of the synthetic conditions needs to be achieved.¹⁸ For this reason the literature on SBA-16 is more limited than that on SBA-15. However, very recently the potential of SBA-16 as a support for copper-based catalysts for the WGS has been investigated.¹⁹

In this work, SBA-16 type silica is used as a support for CeO₂ and Pt nanophases with the aim of designing innovative catalysts for the LT-WGS. The catalyst synthesis was designed to confine both CeO₂ and Pt nanoparticles within the mesoporous silica scaffold. This allows us to maximize the contact between the two nanophases and to hinder their sintering, finally leading to high catalytic activity. Different wet chemistry routes were tested in order to achieve an efficient dispersion of the CeO₂ and Pt nanophases in the SBA-16. A 1 wt% loading of Pt was selected based on recent findings on Pt-based catalysts for the WGS process,⁷ whereas two different CeO₂ loadings were investigated (10 and 20 wt%). The samples were characterized by Transmission Electron Microscopy (TEM), powder X-ray

diffraction (XRD), N₂ physisorption and H₂ chemisorption. The catalytic activity results indicate that the confinement within the SBA-16 pores is effective in stabilizing CeO₂ and maximizing Pt–CeO₂ interactions.

2. Experimental

2.1 Catalyst preparation

Pt–CeO₂/SBA-16 catalysts were prepared either by deposition–precipitation (DP) or by co-impregnation (IMP) of dodecanethiol-protected Pt nanoparticles (Pt-DT) and of cerium(IV) tetrakis(decyloxy) (Ce(OR)₄) on the SBA-16 support. The details of the preparation of Pt-DT nanoparticles and of Ce(OR)₄ are reported in the ESI.†

2.1.1 Synthesis of pure SBA-16 silica. The synthesis of pure SBA-16 was performed according to a procedure¹⁸ which makes use of Pluronic F127 (F127, Aldrich), a high molecular weight ($M_{av} = 12\,600$) block copolymer formed by a sequence of poly(ethylene oxide)–poly(propylene oxide)–poly(ethylene oxide) units with a high EO/PO ratio (EO₁₀₆–PO₇₀–EO₁₀₆), as a structure directing agent. 4.0 g of P127 were added to a mixture of 30 g of water and 120 g HCl 2 M under stirring at room temperature. Tetraethoxysilane (8.5 g, TEOS, Aldrich 98%) was then added to the solution, which became opalescent due to the precipitation of the SBA-16 after about 30 minutes. The solution was left under stirring at room temperature for 20 hours and then aged at 80 °C for 2 days. The precipitate of SBA-16 was separated from the mother solution by centrifugation, washed with distilled water and dried at room temperature. The organic template (F127) was removed by calcination of the dried powder in static air at 500 °C with a heating rate of 1 °C min^{−1}, and held at the final temperature for 6 hours.

2.1.2 Synthesis of Pt–CeO₂/SBA-16 by the DP route. Two Pt–CeO₂/SBA-16 samples were prepared by the deposition–precipitation method according to ref. 13 modified for SBA-16. CeO₂ was deposited into the previously prepared SBA-16 by precipitation of cerium hydroxide. To this end, a solution of cerium(IV) ammonium nitrate (either 0.63 g or 1.26 g in 100 mL) and urea (either 1.02 g or 2.04 g) was added to 1.6 g of SBA-16 in a flask thermostated at 90 °C. The suspension was magnetically stirred for 150 minutes, cooled to room temperature and then filtered. The sample was then washed three times with deionized water, dried at 120 °C for 24 h and calcined at 500 °C for 4 h. Pt was deposited by the addition of 10 mL of an aqueous solution of H₂PtCl₆ to 1 g of the previously prepared CeO₂/SBA-16 to obtain a final Pt content of 1 wt% with respect to the total mass of the catalyst. The solution was stirred for 1 hour at room temperature and 4 hours at 80 °C. The final sample was filtered and dried at 120 °C overnight and calcined at 500 °C for 4 hours. The two samples obtained varying in the CeO₂ content (10 and 20 wt%) will be hereafter labeled DP_Pt/CeO₂(10%)/SBA-16 and DP_Pt/CeO₂(20%)/SBA-16.

2.1.3 Synthesis of Pt–CeO₂/SBA-16 by the IMP route. Two Pt–CeO₂/SBA-16 samples were obtained by impregnation of SBA-16 with a solution containing Pt-DT nanoparticles and Ce(OR)₄ in toluene, whose synthesis is reported in the ESI (Sections S1.1 and S1.2†). Briefly, 1.00 g of SBA-16 was degassed



overnight at 250 °C to remove adsorbed water and then suspended into 10 mL of toluene. Pt-DT and Ce(OR)₄ were mixed together in toluene and added dropwise to the SBA-16 suspension. After stirring for 5 h, the solvent was slowly removed by evaporation at reduced pressure. After drying at 120 °C overnight, the powders were calcined in static air at 500 °C for 5 h. These two catalysts will be hereafter labeled IMP_Pt@CeO₂(10%)/SBA-16 and IMP_Pt@CeO₂(20%)/SBA-16.

To improve the diffusion of Pt-DT and Ce(OR)₄ within the mesopores of SBA-16, the surface of SBA-16 was treated with triethoxyoctyl silane (TEOOS, Sigma-Aldrich, 97.5%) to obtain a hydrophobic support. After degassing the synthesized SBA-16, the support was suspended in 25 mL of toluene and 523 μL of TEOOS was added, and the suspension was refluxed overnight. The powder was recovered by centrifugation, washed 3 times with 10 mL of toluene and vacuum dried at room temperature. Impregnation with Pt-DT and Ce(OR)₄ was performed using the same procedure adopted for the IMP_Pt@CeO₂(XX%)/SBA-16 materials described above. These two catalysts will hereafter be labeled IMP_Pt@CeO₂(10%)/H-SBA-16 and IMP_Pt@CeO₂(20%)/H-SBA-16, where the symbol H indicates that the SBA-16 was made hydrophobic before impregnation.

2.2 Catalyst characterization

XRD patterns were recorded in the range of 15–85° (2θ) on a Panalytical Empyrean diffractometer equipped with a Cu Kα source, a graphite monochromator on the diffracted beam, and a X'Celerator linear detector. The average size of the CeO₂ nanoparticles was estimated using the Scherrer formula, $t = 0.91\lambda / (B \cos \theta)$, where t is the crystallite size, λ is the incident radiation wavelength, θ is the Bragg angle and B is the full-width at half-maximum of the diffraction peak (corrected for instrumental broadening).

Textural characterization was performed by N₂ adsorption-desorption measurements at liquid nitrogen temperature recorded on a Micromeritics ASAP2020. Prior to the analysis, the samples were outgassed under vacuum at 200 °C. Surface areas were estimated using the Brunauer-Emmett-Teller (BET) model,^{20,21} and pore size and pore volumes were estimated using the Non-Local Density Functional Theory (NLDFT) using the Tarazona model for cylindrical pores.²²

H₂ chemisorption experiments were carried out using a Micromeritics ASAP 2020C. Typically, 150 mg of the fresh material were loaded in a U-shaped reactor and reduced by flowing H₂(5%)/Ar (40 mL min⁻¹) at 200 °C for 1 h. The adsorbed hydrogen is then removed by evacuation at 300 °C for 6 h. Chemisorption analysis was performed firstly at -90 °C (liquid/solid acetone bath) dosing H₂ (2–400 torr). Then, the sample was again evacuated at 300 °C for 6 h and the chemisorption analysis was repeated at 35 °C. In both the cases, the contribution of physisorption was subtracted by extrapolating the linear part of the isotherms to zero. In the case of the aged samples, the sample was speedily transferred into the chemisorption reactor after the catalytic tests and only evacuated at 350 °C for 6 h. In this case, the chemisorption analysis was performed only at -90 °C.

Scanning Electron Microscopy (SEM) coupled with Energy Dispersive X-ray spectroscopy (EDX) was performed on a Hitachi S-3400N microscope equipped with a W gun, a Secondary Electron detector and an Oxford Instruments "X-max 80" EDX spectrometer. Samples were deposited on sticky carbon tape mounted on a 15 mm aluminium stub.

Transmission Electron Microscopy (TEM) images were recorded on a Hitachi H-7000 microscope equipped with a W thermoionic filament operating at 125 kV. The catalysts were ground in an agate mortar and deposited on a carbon-coated copper grid for observations. The images were acquired using an AMT DVC CCD camera in bright field (bf) and dark field (df) modes; selected area electron diffraction (SAED) patterns were acquired with a camera length of 20 cm.

2.3 Catalytic activity measurements

WGS activity tests were performed in a U-shaped tubular reactor at atmospheric pressure, using a reaction mixture containing CO (2.0%) + H₂O (10.0%) in Ar. The gas hourly space velocity (GHSV) was set at 50 000 mL g⁻¹ h⁻¹ using a total flow rate of 40.7 mL min⁻¹ and 48.9 mg of the powdered catalyst. A granular quartz bed was put in the reactor to support the powdered catalyst while a 3 mm layer of granular quartz was put above the catalytic bed to pre-heat the gas flow. The flows of CO and Ar were adjusted using mass flow controllers while 3 μL min⁻¹ of H₂O was introduced in the gas flow by means of a gas-tight syringe, controlled by an infusion pump. Vaporization of H₂O was ensured by heating the reactivity line at 120 °C by using adequate heating tapes. The composition of the effluent from the reactor was monitored on-line using a Hiden HPR20 mass spectrometer.

Before each catalytic test, the catalysts were cleaned by treatment in O₂ (5.0%)/Ar (40 mL min⁻¹) at 400 °C for 30 min. After cooling to 150 °C, the gas flow was switched to pure Ar for 1 h to remove O₂ from the system. Light-off experiments were performed by introducing the reaction mixture into the reactor and stabilizing the signals for 1 h, before increasing the temperature up to the desired value at the rate of 2 °C min⁻¹. Aging of the catalysts was performed by maintaining the reactor at the desired temperature for the desired time, before cooling down to 150 °C at the rate of 2 °C min⁻¹.

3. Results and discussion

The XRD patterns of the pure SBA-16 silica support and of the synthesized catalysts are shown in Fig. 1. The pattern of pure SBA-16 (Fig. 1a) is typical of amorphous silica and only shows a broad halo at 2θ ~ 20–30°. The XRD patterns of the catalysts are similar and show broad peaks detectable in the region 2θ ~ 28–80° ascribed to the cubic fluorite-type structure of CeO₂ (PDF 034-0394), on top of the SBA-16 background (Fig. 1b–g). As expected, the intensity of the peaks is higher for the catalysts containing 20 wt% of CeO₂ compared to those containing 10 wt%. The broad XRD reflections are indicative of nanosized CeO₂, having average nanocrystal sizes in the range of 2–4 nm, as calculated by the Scherrer equation (Table 1). These results



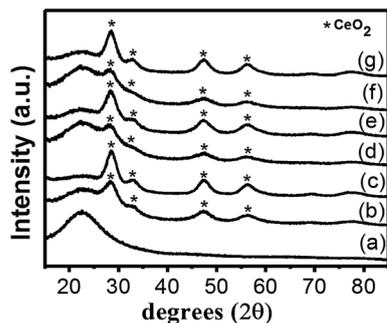


Fig. 1 Wide angle XRD patterns of SBA-16 (a), DP_Pt/CeO₂(10%)/SBA-16 (b), DP_Pt/CeO₂(20%)/SBA-16 (c), IMP_Pt@CeO₂(10%)/SBA-16 (d), IMP_Pt@CeO₂(20%)/SBA-16 (e), IMP_Pt@CeO₂(10%)/H-SBA-16 (f) and IMP_Pt@CeO₂(20%)/H-SBA-16 (g). The positions corresponding to the main reflections of the fluorite structure of CeO₂ are also shown.

indicate that nanocrystalline ceria has been effectively stabilized into SBA-16 with 10 and 20 wt% loadings. In all the materials investigated, no reflections ascribed to Pt-related phases can be detected, as expected for the low metal loading and the small size of the Pt nanoparticles,²³ as already reported for various Pt–CeO₂ systems.^{2,3,24–27}

The texture of the pristine SBA-16 and of the prepared materials was studied by N₂ physisorption at liquid nitrogen temperature. According to the IUPAC classification, the N₂ physisorption isotherms of pure SBA-16 (Fig. 2A) can be classified as type IV isotherms with a H2 type hysteresis, typical of materials containing ink-bottle mesopores and indicative of a cage-like cubic porous structure. The N₂ physisorption isotherms of the materials prepared by DP (Fig. 2B and C), indicate that the mesoscopic cubic order is preserved after the calcination treatments needed to stabilize the CeO₂ and Pt components, and that the pores of SBA-16 are not significantly occluded by the deposition of the active phase. The isotherms of the samples prepared by co-impregnation on the bare SBA-16 (Fig. 2D and E) and on the hydrophobic H-SBA-16 (Fig. 2F and G) show a significant decrease in the overall adsorbed amount of N₂ with respect to the pristine SBA-16 support.

The textural parameters calculated by analyzing the physisorption data are summarized in Table 1. SBA-16 has a very high surface area and pore volume, with the pore size distribution centered at 6.5 nm, in accordance with the literature.^{28,29} After loading of the active phase, the differences in textural

properties are strongly dependent on the preparation route adopted and generally more evident for samples having a 20 wt% CeO₂ loading. For almost all samples, a decrease in the surface area and pore volume is observed, although the pore size distribution is only marginally affected. The deposition of the active phase by DP causes the smallest change in surface area and pore volume while the IMP route results in a significant decrease of the accessible surface area and pore volume. The decrease in pore volume is more evident in the samples prepared using H-SBA-16 and can be associated with the occlusion of a significant fraction of pores. Notably, the process to make the support hydrophobic has been introduced in the preparation procedure in order to maximize the diffusion of hydrophobic Pt-DT and Ce(OR)₄ precursors within the SBA-16 texture. It should be pointed out that plugging of the pores with the catalysts remaining on the SBA-16 surface can be excluded since this would be accompanied by a drastic decrease of the surface area which was not observed for any of the samples. It should also be noted that DP_Pt/CeO₂(20%)/SBA-16 has a surface area and pore volume very similar (within the experimental error) to the pure SBA-16 sample, while DP_Pt/CeO₂(10%)/SBA-16 has a slightly lower surface area.

The mesoporous structure of the catalysts and the dispersion of the nanostructured active phases in the SBA-16 matrix were investigated by TEM. The representative TEM images are shown in Fig. 3 for DP_Pt/CeO₂(10%)/SBA-16 and DP_Pt/CeO₂(20%)/SBA-16, in Fig. 4 for IMP_Pt@CeO₂(10%)/SBA-16 and IMP_Pt@CeO₂(20%)/SBA-16 and in Fig. 5 for IMP_Pt@CeO₂(10%)/H-SBA-16 and IMP_Pt@CeO₂(20%)/H-SBA-16. A highly ordered arrangement of mesopores can be seen in all the samples, confirming that the SBA-16 scaffold is preserved in the final material, independent of the preparation route employed, in agreement with the physisorption analysis results. The samples are composed of spherical nanoparticles homogeneously dispersed within the SBA-16 mesoporous matrix. CeO₂ nanoparticles are particularly evident in the dark field images, in which they appear as bright spots. The SAED patterns of CeO₂ in all the samples show rings corresponding to the fluorite CeO₂ structure, confirming that the nanoparticles are crystalline (ESI, Fig. S1†). The TEM images show that the CeO₂ nanoparticles are well dispersed and confined in the SBA-16 matrix. In particular, in DP_Pt/CeO₂(20%)/SBA-16 (Fig. 3C and D and S2†) CeO₂ nanocrystals smaller than the size of the SBA-16 pores are clearly visible both in the bright field images as

Table 1 Textural properties of SBA-16 and nanocomposite materials obtained from N₂ physisorption data^a and the average crystallite size obtained from XRD using the Scherrer equation^b

Sample	Surface area ^a (m ² g ⁻¹)	Pore width ^a (nm)	Pore volume ^a (cm ³ g ⁻¹)	CeO ₂ crystallite size ^b (nm)
SBA-16	747	6.5	0.49	—
DP_Pt/CeO ₂ (10%)/SBA-16	613	6.5	0.44	3 ± 1
DP_Pt/CeO ₂ (20%)/SBA-16	773	6.5	0.49	3 ± 1
IMP_Pt@CeO ₂ (10%)/SBA-16	434	7.4	0.54	2 ± 1
IMP_Pt@CeO ₂ (20%)/SBA-16	306	7.5	0.43	3 ± 1
IMP_Pt@CeO ₂ (10%)/H-SBA-16	549	6.5	0.35	4 ± 1
IMP_Pt@CeO ₂ (20%)/H-SBA-16	304	6.4–7.1	0.28	3 ± 1



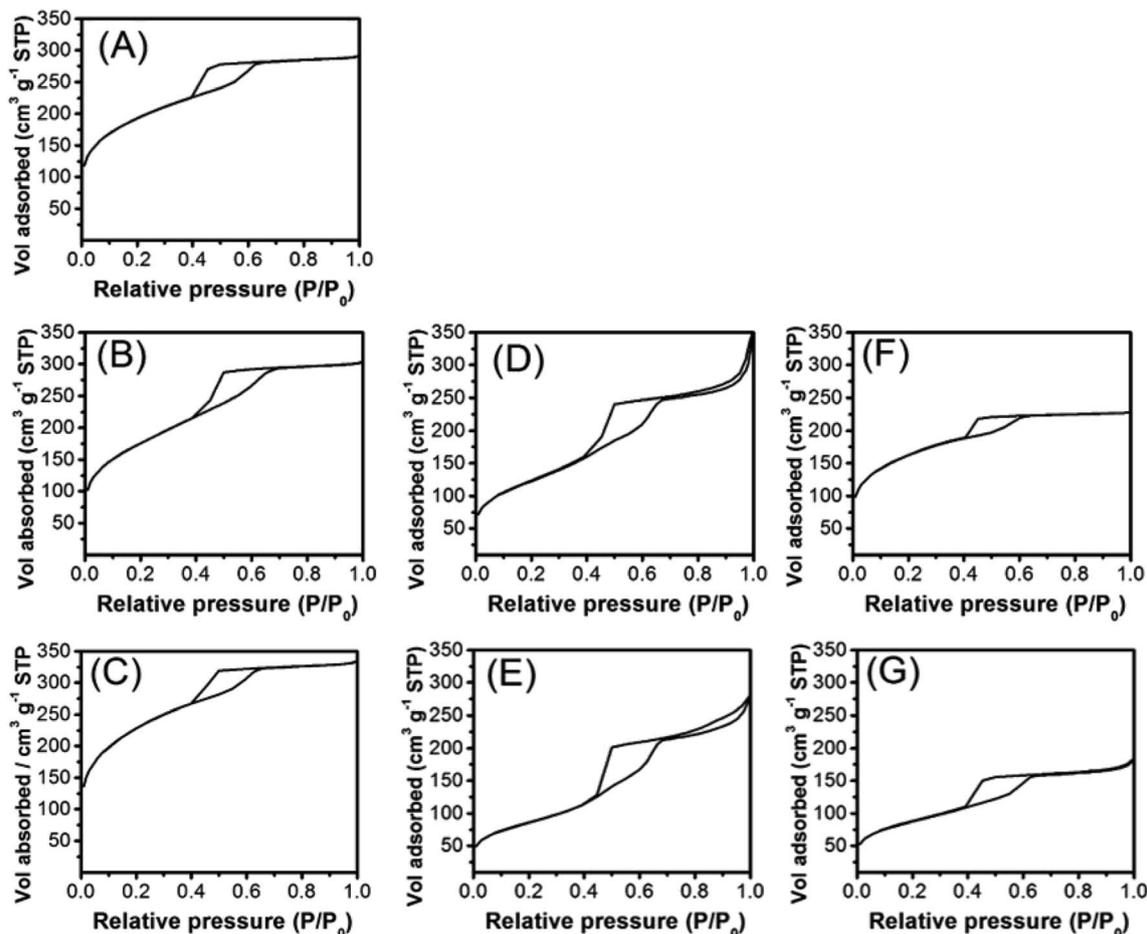


Fig. 2 N_2 physisorption isotherms collected at liquid nitrogen temperature for SBA-16 (A), DP_Pt/CeO₂(10%)/SBA-16 (B), DP_Pt/CeO₂(20%)/SBA-16 (C), IMP_Pt@CeO₂(10%)/SBA-16 (D), IMP_Pt@CeO₂(20%)/SBA-16 (E), IMP_Pt@CeO₂(10%)/H-SBA-16 (F) and IMP_Pt@CeO₂(20%)/H-SBA-16 (G).

dark spots and in the dark field images as bright spots. On the other hand, the CeO₂ nanoparticles in the samples prepared by co-impregnation seem to be less evident compared to the corresponding DP samples. This could be justified by either a less homogeneous dispersion of the nanoparticles in the matrix or by their confinement within the SBA-16 scaffold which would make them more difficult to be detected because of the thickness of the sample under investigation. The presence of Pt nanoparticles cannot be detected in any of the materials, because of the low loading and high dispersion of Pt in the CeO₂-SBA-16 matrix.²³ Further information on the morphology of the catalysts and on the dispersion of the nanostructured active phases in the SBA-16 matrix was obtained by SEM coupled with EDS, as reported in the ESI (Fig S3[†]). As can be observed from SEM images, no CeO₂ aggregates have been observed outside the SBA-16 scaffold, indicating that the active phase is deposited within the mesoporous structure of the SiO₂ scaffold.

H₂ chemisorption analysis was employed to assess the accessibility of the Pt phase and the interaction between the Pt nanoparticles and the CeO₂ promoter. It was performed at -90 °C and 35 °C, with the results summarized in Table 2. In

agreement with previous H₂ chemisorption studies on noble metal nanoparticles supported on CeO₂-based materials,^{30,31} activated H atoms are spilled over the reducible oxide inducing the so-called “reversible reduction” of the promoter.³² The reversible reduction of CeO₂ results in the conversion of the more superficial Ce(IV) to Ce(III) and the formation of superficial OH groups, which allow the H hopping and diffusion on the surface.³² The transfer of activated H atoms is strongly inhibited lowering the analysis temperature.^{30,31} In this study, the H/Pt value measured at -90 °C is used to determine the fraction of accessible Pt atoms. The results obtained indicate that the Pt accessibility depends both on the CeO₂ loading and on the preparation method. A slightly lower accessibility of surface Pt is observed for the materials with the higher CeO₂ content; however, the preparation method has the most prominent effect on accessibility, which is significantly enhanced in the samples prepared by the DP route, independent of the CeO₂ loading. These results can be rationalized considering that the platinum precursor is impregnated after the deposition of CeO₂ by deposition-precipitation on the SBA-16 support. On the other hand, the co-impregnation route could lead to a partial occlusion of Pt nanoparticles within the CeO₂ nanocrystallites, in



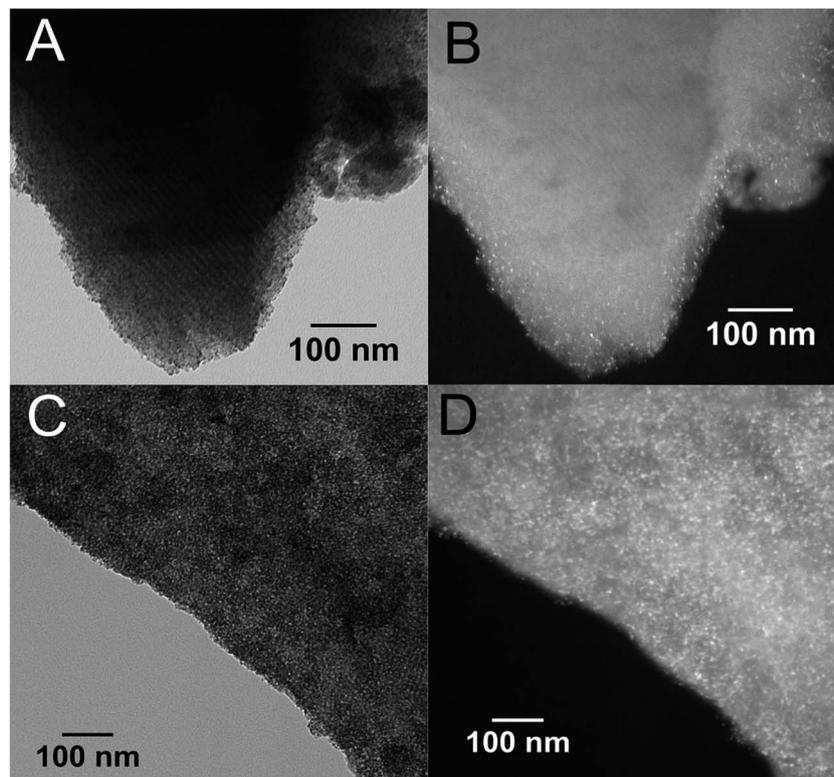


Fig. 3 Representative TEM images of DP_Pt/CeO₂(10%)/SBA-16 (A and B) and DP_Pt/CeO₂(20%)/SBA-16 (C and D). Bright field (A and C) and dark field (B and D).

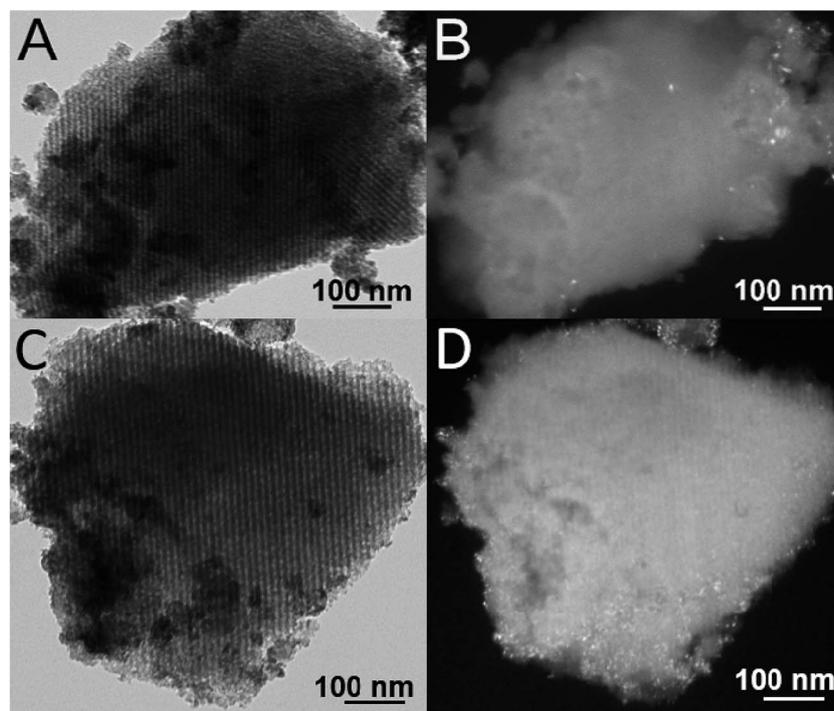


Fig. 4 Representative TEM images of IMP_Pt@CeO₂(10%)/SBA-16 (A and B) and IMP_Pt@CeO₂(20%)/SBA-16 (C and D). Bright field (A and C) and dark field (B and D).



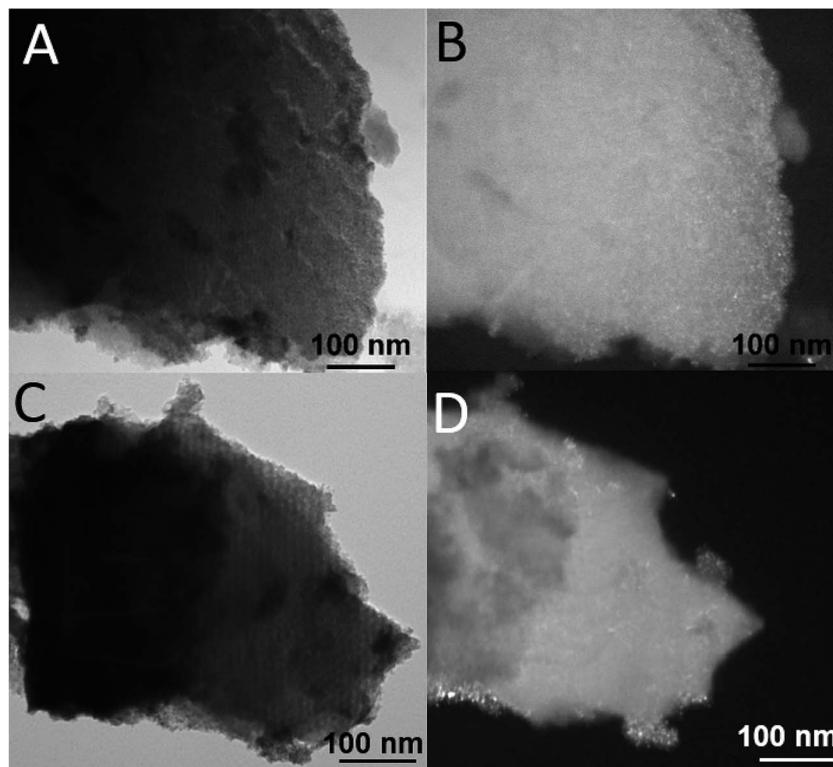


Fig. 5 Representative TEM images of IMP_Pt@CeO₂(10%)/H-SBA-16 (A and B) and IMP_Pt@CeO₂(20%)/H-SBA-16 (C and D). Bright field (A and C) and dark field (B and D).

a similar manner to what was observed for Pd@CeO₂/Si-Al₂O₃ prepared from self-assembled core-shell units.³³

The H/Pt value measured at 35 °C includes the amount of H spilt over reducible CeO₂. Therefore, the ratio between the H/Pt value measured at 35 °C and the one measured at 90 °C (*R* in Table 2) can be considered as an indication of the interaction between Pt and CeO₂ nanoparticles. All the H/Pt values measured at 35 °C are significantly higher than the values measured at low temperature and, notably, higher than 1, indicating that H is effectively spilt over the CeO₂ surface. Higher H/Pt values have been obtained, for each preparation method, with the materials with the higher CeO₂ content, in agreement with the more extended reducible surface available to accommodate the OH groups. On the other hand, for the same CeO₂ loading, the preparation method has a strong

influence on the hydrogen chemisorption capacity of the samples. The materials prepared by DP show the lowest H₂ chemisorption capacity although the Pt accessibility is the highest. These results indicate that the Pt-CeO₂ interaction is not optimal in the DP samples, with a significant fraction of Pt nanoparticles deposited onto the free SBA-16 surface being not in direct contact with CeO₂ nanoparticles. The samples prepared by impregnation of SBA-16 with Pt-DT nanoparticles and Ce(OR)₄ show higher H₂ chemisorption capacities and better abilities to spill activated H onto the CeO₂ surface, despite the lower Pt accessibility and surface area. The impregnation methodology was designed in order to obtain Pt nanoparticles embedded into a matrix of CeO₂ crystallites, with the aim of maximizing the Pt-CeO₂ interaction. The spillover is further enhanced in the samples impregnated on the hydrophobic H-SBA-16 surface, indicating that this treatment allows the best distribution of the Pt and CeO₂ components.

Table 2 H₂ chemisorption results of the synthesized Pt-CeO₂/SBA-16 materials: H/Pt values measured at 35 °C and -90 °C and their ratio (*R*)

Sample	H/Pt at -90 °C	H/Pt at 35 °C	<i>R</i>
DP_Pt/CeO ₂ (10%)/SBA-16	0.743	1.084	1.46
DP_Pt/CeO ₂ (20%)/SBA-16	0.701	1.312	1.87
IMP_Pt@CeO ₂ (10%)/SBA-16	0.401	1.451	3.62
IMP_Pt@CeO ₂ (20%)/SBA-16	0.374	1.896	5.07
IMP_Pt@CeO ₂ (10%)/H-SBA-16	0.384	3.176	8.27
IMP_Pt@CeO ₂ (20%)/H-SBA-16	0.330	4.492	13.60

3.1 WGS activity

The CO conversion in consecutive reaction experiments over the synthesized catalysts is presented in Fig. 6–8. CO conversion increases with increasing temperature, until the thermodynamic equilibrium is reached. The stability of the present catalysts was evaluated by simulated fast aging of the catalysts by prolonged treatment under reaction conditions at 350 °C. The light-off temperatures (*T*₅₀ – corresponding to 50% of CO conversion) and the lower temperature at which the chemical



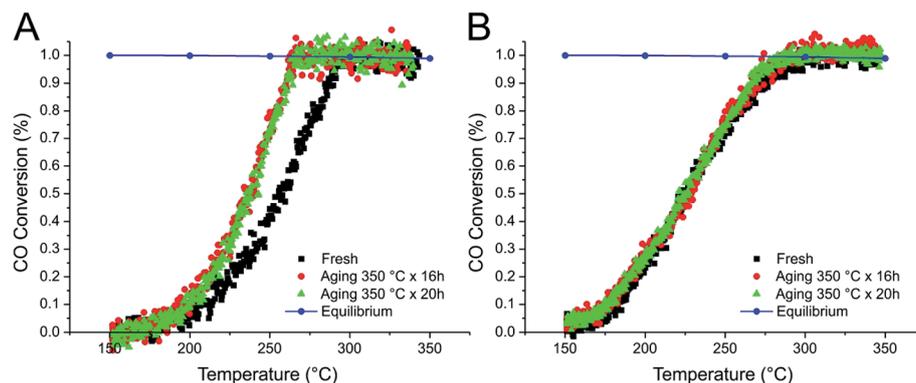


Fig. 6 CO conversion during the WGS over DP_Pt/CeO₂(10%)/SBA-16 (A) and DP_Pt/CeO₂(20%)/SBA-16 (B) catalysts.

equilibrium is reached (T_{EQ}) are summarized in Table 3. Notably, no CO conversion was observed during blank experiments (the reactor containing only the quartz bed and the thermocouple), while a Pt/SBA-16 sample, not containing CeO₂, showed only very poor catalytic activity, leading to CO conversion of 15% at 350 °C (ESI Fig. S4†).

The fresh DP samples (Fig. 6) showed no activity at 150 °C while by increasing the temperature the chemical equilibrium is reached around 290 °C, irrespective of the CeO₂ content. The DP_Pt/CeO₂(10%)/SBA-16 sample revealed a slight increase of activity in light-off experiments after aging for 16 h, being unmodified by further 20 hours of aging. On the other hand, the DP_Pt/CeO₂(20%)/SBA-16 sample showed comparable and stable activities during the consecutive aging treatments.

Significantly higher catalytic activities were observed for the IMP materials (Fig. 7 and 8), with the best performances demonstrated by the H-SBA-16 supported catalysts. The fresh IMP_Pt@CeO₂(10%)/H-SBA-16 and IMP_Pt@CeO₂(20%)/H-SBA-16 catalysts showed remarkable CO conversion at 150 °C (18 and 32%, respectively – Fig. 8). Despite the lower surface area and lower accessibility of Pt nanoparticles with respect to the DP catalysts, the enhanced performances of the materials prepared by impregnation can be related to the more efficient interaction between the Pt nanoparticles and the CeO₂ promoter, as highlighted by H₂ chemisorption experiments. In agreement with

this, the oxidation of CO has been demonstrated to be strongly dependent on the extent of the Pt–CeO₂ interface.³⁴

Apparent activation energy (E_{att}) values ranging from 50 to 80 kJ mol⁻¹ were calculated from CO conversion results during the WGS tests performed on the fresh material for all the investigated catalysts. These values are in good agreement with the results previously reported for Pt supported on reducible (CeO₂ and CeO₂-ZrO₂)^{35,36} and non-reducible oxides (ZrO₂ and Al₂O₃).^{35,37}

After simulated aging of the impregnated catalysts at 350 °C for 16 and 36 h, some deactivation occurred, as evidenced by the shift of the light-off temperatures to slightly higher values (~30 °C). Considering that the operating temperature of a WGS catalyst should be quite close to the temperature at which the chemical equilibrium is reached, to further investigate the stability of the impregnated catalysts, IMP_Pt@CeO₂(10%)/SBA-16 and IMP_Pt@CeO₂(20%)/H-SBA-16 were aged under reaction conditions at the T_{EQ} observed during the first light-off experiment (230 and 200 °C, respectively). Even after a total aging time of 60 hours, the aged IMP_Pt@CeO₂(20%)/H-SBA-16 showed comparable light-off curves to the fresh catalyst, while the IMP_Pt@CeO₂(10%)/H-SBA-16 showed only a slight deactivation (ESI Fig. S5†).

The deactivation of Pt/CeO₂ catalysts is usually related to sintering of Pt nanoparticles³⁸ or with the adsorption of

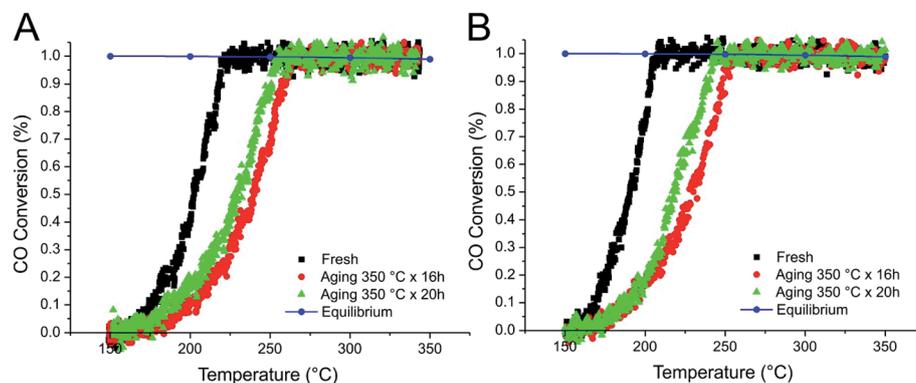


Fig. 7 CO conversion during the WGS over IMP_Pt@CeO₂(10%)/SBA-16 (A) and IMP_Pt@CeO₂(20%)/SBA-16 (B) catalysts.



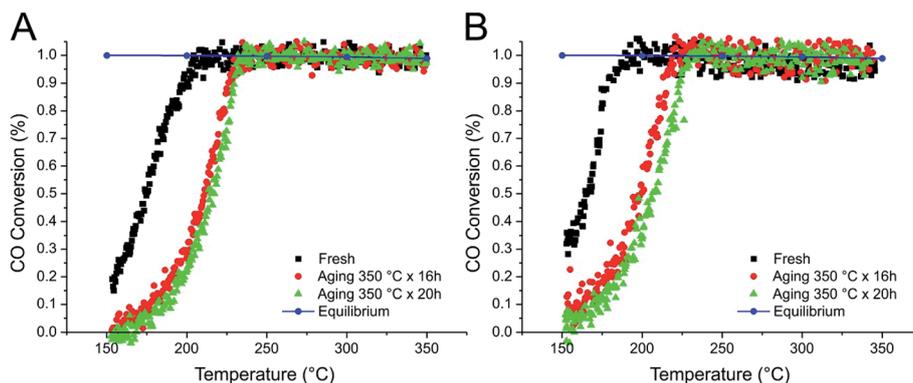


Fig. 8 CO conversion during the WGS over IMP_Pt@CeO₂(10%)/H-SBA-16 (A) and IMP_Pt@CeO₂(20%)/H-SBA-16 (B) catalysts.

intermediates, mainly formates, on the active sites.³⁹ The latter is usually recognized as a major cause of deactivation at lower temperatures (below 300 °C), when formates are spectators of the reaction and accumulate on the catalyst surface.³⁹ At higher temperature, formates are further converted and become intermediates in the catalytic process,³⁹ while sintering of Pt is considered the main reason for catalyst deactivation.³⁸ In order to hinder Pt sintering, hierarchical Pt@CeO₂/Si-Al₂O₃ samples were prepared from self-assembled core-shell nanoparticles,⁴⁰ obtaining stable catalytic activity after pre-treatment of the catalyst at 500 °C. Differently from Pt, Pd@CeO₂/Si-Al₂O₃ demonstrates a fast decrease of the catalytic activity,^{40,41} mainly because of an electronic influence due to reduced CeO_{2-x} and the partial occlusion of Pd nanoparticles within CeO_{2-x} crystallites.⁴¹ To overcome these problems, Pd@CeO₂/MWCNT hybrid materials have been synthesized, observing that the balance between the organic and inorganic components favors their electronic interaction, finally leading to stable catalytic performances in the WGS.⁴²

To investigate the origin of deactivation observed in the Pt/CeO₂ catalysts supported on SBA-16, the materials after catalytic tests have been characterized in detail with a view to highlight differences induced by prolonged aging under WGS conditions. Also in this case, SEM coupled with EDX does not evidence the segregation of the Pt/CeO₂ active phase outside the SBA-16 (see ESI Fig. S6†). The surface area of the materials subjected to severe simulated aging at 350 °C under WGS

conditions significantly decreases with respect to the fresh materials (see ESI Table S1 and Fig. S7†). Notably, the extent of this sintering phenomenon is lower in the case of the catalysts tested under prolonged aging under realistic LT-WGS conditions (200–230 °C) (see ESI Table S1†). The most important modification highlighted by XRD analysis (Fig. S8†) is the presence of a broad (but detectable) peak at $2\theta \sim 40^\circ$, corresponding to the main reflection of Pt. This reflection is present in all the aged samples, to a different extent depending on the material composition and preparation, being the most intense in the IMP_Pt@CeO₂(XX%)/H-SBA-16 materials. This result suggests that the Pt nanoparticles undergo a partial sintering and/or crystallization process during aging under WGS conditions. Notably, this phenomenon does not depend on the treatment temperature, being comparable for samples after severe simulated aging at 350 °C and prolonged aging under LT-WGS conditions (200–230 °C). In agreement with this, the H/Pt values measured by H₂ chemisorption analysis at –90 °C (see ESI Table S2†) are lower than those measured for the fresh materials. The decreased ability/accessibility of Pt to H₂ can be due to a combination of different phenomena, including partial sintering, occlusion of Pt nanoparticles by the sintered support and some extent of electronic strong metal support interaction (SMSI). Also in this case, the decrease of H/Pt values is less important in the case of the catalysts aged under LT-WGS conditions. On the basis of these considerations, the deactivation of the present impregnated catalysts is expected to take place following different

Table 3 Light-off temperature (T_{50}) and the temperature at which the chemical equilibrium for the WGS is achieved (T_{EQ}) during light-off experiments after different aging treatments

Sample	T_{50}			T_{EQ}		
	Fresh	Aging 350 °C × 16 h	Aging 350 °C × 20 h	Fresh	Aging 350 °C × 16 h	Aging 350 °C × 20 h
DP_Pt/CeO ₂ (10%)/SBA-16	258	237	239	294	262	265
DP_Pt/CeO ₂ (20%)/SBA-16	224	229	228	294	291	292
IMP_Pt@CeO ₂ (10%)/SBA-16	203	239	230	228	257	265
IMP_Pt@CeO ₂ (20%)/SBA-16	190	229	218	210	254	245
IMP_Pt@CeO ₂ (10%)/H-SBA-16	175	210	216	210	234	238
IMP_Pt@CeO ₂ (20%)/H-SBA-16	165	212	207	200	221	231



mechanisms depending on the aging temperature. The very limited deactivation observed during aging under LT-WGSR conditions (see ESI Fig. S5†) can be related essentially to deposition of intermediates on the active sites of the catalysts. In agreement with this, deactivation is moderate and is less evident in the materials designed for maximizing the Pt–CeO₂ interaction by confinement within the nanometric pores of the SBA-16 scaffold. In this case, deactivation by electronic influence of reduced CeO_{2–x} is less important, in agreement with the characterization results of the aged catalysts and previous studies on H₂ and CO chemisorption on Pt/CeO₂⁴³ and Pt/CeO₂–ZrO₂³⁰ reduced at various temperatures. In any case, the IMP_Pt@CeO₂(20%)/H-SBA-16 material showed promising performance for application as an active and stable catalyst in the LT-WGSR.

During aging at higher temperature (350 °C), a more severe deactivation occurred, in agreement with the fact that an SMSI state is obtained in the materials. The loss of catalytic activity can be related to the electronic deactivation induced by reduced CeO_{2–x} and by encapsulation of Pt nanoparticles resulting from partial sintering of the material, in a similar manner to Pd@CeO₂/Si–Al₂O₃ catalysts.⁴¹

The catalysts prepared by deposition–precipitation are not active enough to allow their application in the LT-WGSR, although they are not negatively affected by severe aging under WGSR conditions. The lower activity derives from the less extent of Pt–CeO₂ interaction while the higher stability is obtained, thanks to the higher dispersion/accessibility of the Pt component.

4. Conclusions

Nanocomposite materials with a Pt–CeO₂ active phase dispersed within highly porous silica with a well-defined cubic symmetry of a porous scaffold (SBA-16) have been prepared and tested as catalysts for the LT-WGSR, an important industrial process involved in large scale production of pure H₂. Different synthetic methods have been compared with the final aim to maximize the interactions of Pt and CeO₂ nanocrystallites, yielding active and stable catalysts. The preparation methodology greatly affected the textural and morphological properties of the synthesized materials. The key to achieving more active and stable catalysts is the confinement of the active phase within the SBA-16 scaffold, and the enhancement of the interaction between Pt nanoparticles and CeO₂ nanocrystallites, as evidenced by H₂ chemisorption and catalytic tests. The catalytic WGSR performance was optimized in IMP materials by exploitation of the interactions of functionalized Pt-DT nanoparticles with cerium decyloxyde and by the hydrophobization of the SBA-16 support.

Conflicts of interest

The authors declare that there are no conflicts of interest.

Acknowledgements

The authors would like to thank the Fondazione Banco di Sardegna, IIT, INSTM and the University of Trieste through the FRA2015 project for support.

References

- 1 F. de Bruijn, *Green Chem.*, 2005, **7**, 132.
- 2 L. Torrente-Murciano and F. R. Garcia-Garcia, *Catal. Commun.*, 2015, **71**, 1–6.
- 3 R. Jain and R. Maric, *Appl. Catal., A*, 2014, **475**, 461–468.
- 4 J. H. Lin, P. Biswas, V. V. Gulians and S. Misture, *Appl. Catal., A*, 2010, **387**, 87–94.
- 5 T. Montini, M. Melchionna, M. Monai and P. Fornasiero, *Chem. Rev.*, 2016, **116**, 5987–6041.
- 6 J. M. de Souza e Silva, M. Strauss, C. M. Maroneze, E. R. Souza, Y. Gushikem, F. A. Sigoli and I. O. Mazali, *J. Mater. Chem.*, 2011, **21**, 15678.
- 7 R. Tiwari, B. Sarkar, R. Tiwari, C. Pendem, T. Sasaki, S. Saran and R. Bal, *J. Mol. Catal. A: Chem.*, 2014, **395**, 117–123.
- 8 J. Káspár and P. Fornasiero, *Catalysis by Ceria and Related Materials*, Imperial College Press, London, 2002.
- 9 A. Trovarelli, *Catal. Rev.*, 1996, **38**, 439–520.
- 10 T. M. Onn, S. Zhang, L. Arroyo-Ramirez, Y. Xia, C. Wang, X. Pan, G. W. Graham and R. J. Gorte, *Appl. Catal., B*, 2017, **201**, 430–437.
- 11 R. Di Monte, P. Fornasiero, J. Kašpar, M. Graziani, J. M. Gatica, S. Bernal and A. Gómez-Herrero, *Chem. Commun.*, 2000, 2167–2168.
- 12 J. Strunk, W. C. Vining and A. T. Bell, *J. Phys. Chem. C*, 2011, **115**, 4114–4126.
- 13 Y. D. Bi, W. Zhang, H. Y. Xu and W. Z. Li, *Catal. Lett.*, 2007, **119**, 126–133.
- 14 A. Salis, D. Meloni, S. Ligas, M. F. Casula, M. Monduzzi, V. Solinas and E. Dumitriu, *Langmuir*, 2005, **21**, 5511–5516.
- 15 R. M. Grudzien, B. E. Grabicka and M. Jaroniec, *Appl. Surf. Sci.*, 2007, **253**, 5660–5665.
- 16 L. Sierra, S. Valange, J. Barrault and J. L. Guth, *Microporous Mesoporous Mater.*, 2008, **113**, 352–361.
- 17 D. Carta, S. Bullita, M. F. Casula, A. Casu, A. Falqui and A. Corrias, *ChemPlusChem*, 2013, 364–374.
- 18 D. Zhao, Q. Huo, J. Feng, B. F. Chmelka and G. D. Stucky, *J. Am. Chem. Soc.*, 1998, **120**, 6024–6036.
- 19 C. Marras, D. Loche, D. Carta, M. F. Casula, M. Schirru, M. G. Cutrufello and A. Corrias, *ChemPlusChem*, 2016, **81**, 421–432.
- 20 S. Brunauer, P. H. Emmett and E. Teller, *J. Am. Chem. Soc.*, 1938, **60**, 309–319.
- 21 F. Rouquerol, J. Rouquerol and K. Sing, *Adsorption by Powders and Porous Solids, Principles, Methodology and Applications*, Academic Press, London, 1999.
- 22 P. Tarazona, U. Marini Bettolo Marconi and R. Evans, *Mol. Phys.*, 1987, **60**, 573–595.
- 23 T. Hyde, *Platinum Met. Rev.*, 2008, **52**, 129–130.
- 24 G. Cavusoglu, D. Miao, H. Lichtenberg, H. W. P. Carvalho, H. Xu, A. Goldbach and J. D. Grunwaldt, *Appl. Catal., A*, 2015, **504**, 381–390.
- 25 H.-S. Roh, H. S. Potdar, D.-W. Jeong, K.-S. Kim, J.-O. Shim, W.-J. Jang, K. Y. Koo and W. L. Yoon, *Catal. Today*, 2012, **185**, 113–118.



- 26 A. Luengnaruemitchai, S. Osuwan and E. Gulari, *Catal. Commun.*, 2003, **4**, 215–221.
- 27 P. Lu, B. Qiao, N. Lu, D. C. Hyun, J. Wang, M. J. Kim, J. Liu and Y. Xia, *Adv. Funct. Mater.*, 2015, **25**, 4153–4162.
- 28 D. Carta, M. F. Casula, S. Bullita, A. Falqui, A. Casu, C. M. Carbonaro and A. Corrias, *Microporous Mesoporous Mater.*, 2014, **194**, 157–166.
- 29 D. Carta, G. Mountjoy, R. Apps and A. Corrias, *J. Phys. Chem. C*, 2012, **116**, 12353–12365.
- 30 J. Gatica, R. Baker, P. Fornasiero, S. Bernal and J. Kaspar, *J. Phys. Chem. B*, 2001, **105**, 1191–1199.
- 31 J. M. Gatica, R. T. Baker, P. Fornasiero, S. Bernal, G. Blanco and J. Kaspar, *J. Phys. Chem. B*, 2000, **104**, 4667–4672.
- 32 A. Norman and V. Perrichon, *Phys. Chem. Chem. Phys.*, 2003, **5**, 3557–3564.
- 33 M. Cargnello, J. Delgado Jaén, J. Hernández Garrido, K. Bakhmutsky, T. Montini, J. Calvino Gámez, R. Gorte and P. Fornasiero, *Science*, 2012, **337**, 713–717.
- 34 M. Cargnello, V. V. Doan-Nguyen, T. R. Gordon, R. E. Diaz, E. A. Stach, R. J. Gorte, P. Fornasiero and C. B. Murray, *Science*, 2013, **341**, 771–773.
- 35 D. W. Jeong, H. S. Potdar, J. O. Shim, W. J. Jang and H. S. Roh, *Int. J. Hydrogen Energy*, 2013, **38**, 4502–4507.
- 36 D. W. Jeong, W. J. Jang, J. O. Shim, W. B. Han, H. M. Kim, Y. L. Lee, J. W. Bae and H. S. Roh, *Renewable Energy*, 2015, **79**, 78–84.
- 37 P. Panagiotopoulou and D. I. Kondarides, *Catal. Today*, 2006, **112**, 49–52.
- 38 W. Ruettinger, X. Liu and R. J. Farrauto, *Appl. Catal., B*, 2006, **65**, 135–141.
- 39 F. C. Meunier, D. Tibiletti, A. Goguet, S. Shekhtman, C. Hardacre and R. Burch, *Catal. Today*, 2007, **126**, 143–147.
- 40 L. Arroyo-Ramírez, C. Chen, M. Cargnello, C. B. Murray and R. J. Gorte, *Catal. Today*, 2015, **253**, 137–141.
- 41 N. L. Wieder, M. Cargnello, K. Bakhmutsky, T. Montini, P. Fornasiero and R. J. Gorte, *J. Phys. Chem. C*, 2011, **115**, 915–919.
- 42 A. Beltram, M. Melchionna, T. Montini, L. Nasi, R. J. Gorte, M. Prato and P. Fornasiero, *Catal. Today*, 2015, **253**, 142–148.
- 43 D. W. Daniel, *J. Phys. Chem.*, 1988, **92**, 3891–3899.

