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Physico-chemical investigation of catalytic oxidation sites in 4%Rh/CeO₂ catalysts prepared by impregnation and deposition–precipitation methods



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ABSTRACT

4 wt% Rh/CeO₂ catalysts were prepared by two different methods: deposition–precipitation (DP) and impregnation (Imp). X-ray diffraction (XRD), differential scanning calorimetry and thermogravimetry (DSC-TG), temperature-programmed reduction (TPR), and electron paramagnetic resonance (EPR) were used for physicochemical characterization. The solids were tested in propylene and carbon black oxidation reactions. The 4%Rh/ CeO₂ (DP) showed better catalytic performance in both reactions compared to the catalyst prepared by the impregnation method. The XRD technique evidenced the formation of Rh₂O₃ phase in the DP-solid after calcination of this latter at 400 °C for 4 h. EPR evidenced, only in the DP-solid, the presence of O_2^- species in interaction with the CeO₂ surface whereas Rh⁴⁺ ions in the form of clusters were identified in both solids. The TPR technique showed that the DP-solid was reduced by hydrogen at lower temperature compared to the impregnated one. The higher catalytic performance of the DP-solid was attributed to the presence of O₂⁻ species along with the presence of Rh₂O₃ phase in ceria and to the better reducibility and lower particle size of the rhodium species.

1. Introduction

Ceria (CeO₂) has a great reputation in catalysis and was well investigated due to its high oxygen storage capacity and its use in catalytic oxidation reactions [1-3]. It is also well known that adding transition metals over ceria enhances the activity of the latter [4,5].

Rhodium supported on CeO₂ is extensively used as catalyst in many reforming reactions and it is also involved in a number of oxidation reactions [6–12]. Haneda et al. [6] have shown that the Ir-Rh/CeO₂-ZrO₂ improves three-way catalytic activity by oxidizing CO and C₃H₆ molecules and reducing NO one. Shimizu et al. [7] have prepared Rh/ CeO₂ catalysts by a supercritical CO₂ impregnation technique and they used them for automotive exhaust. The catalysts have been characterized by energy-dispersive X-ray spectroscopy (EDX), X-ray diffraction (XRD), and transmission electron microscopy (TEM) to analyze the Rh loading, Rh structure, and the morphology of the particles, respectively. Li et al. [8] have investigated Rh/CeO₂-SiC catalyst in the partial oxidation of ethanol for hydrogen production. Cuauhtémoc et al. [9] have used Rh/ $\gamma\text{-Al}_2\text{O}_3$. CeO $_2$ as catalyst to oxidize gasoline oxygenates by wet air.

The EPR technique is highly sensitive compared to other spectroscopic techniques (XPS, IR, UV–visible, etc.) making it suitable for the investigation of adsorbed paramagnetic species at solid surfaces. These paramagnetic species can be intermediate compounds implied in catalytic oxidation and reduction reactions. Therefore, their identification could help in accessing reactions mechanisms which depend on the redox properties of the active sites located at the support surface and the support itself.

The aim of the present study is to compare the catalytic performances of two series of 4%Rh/CeO₂ catalysts (prepared by deposition–precipitation and conventional impregnation) towards the total oxidation of propylene and carbon black particles and to use physicochemical techniques such as EPR and TPR to explain the performances obtained. Propylene is an example of the pollutants volatile organic compounds and carbon black is used as soot model.

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2. Experimental part

2.1. Catalysts preparation

Cerium hydroxide was prepared by precipitation of cerium (III) nitrate hexahydrate, Ce(NO₃)₃·6H₂O (ACROS) with sodium hydroxide NaOH (ACROS). The suspension was filtered, washed several times with hot deionized water, and dried at 100 °C for 24 h. CeO₂ was obtained by calcination under air at 400 °C for 4 h.

The rhodium chloride $RhCl_3$:xH₂O (STREM CHEMICALS) was used as rhodium precursor and introduced into ceria according to two methods: wet impregnation and deposition–precipitation.

Wet impregnation: The supported rhodium/CeO₂ catalysts were prepared by the conventional wet impregnation method. CeO₂ was impregnated with rhodium solution. The slurry was stirred 2 h at ambient temperature and then water was slowly evaporated at 75 °C in a rotary evaporator. The as-dried sample was put in an oven set at 100 °C for 20 to 24 h. The obtained powder was calcined at 400 °C (1 °C.min⁻¹, 4 h, 2 L dried air.h⁻¹). Two rhodium loadings were prepared: 0.5 wt% and 4 wt%. The samples will be respectively designated by 0.5% Rh/CeO₂ (Imp) and 4% Rh/CeO₂ (Imp).

Deposition–precipitation: Rhodium (4 wt%) was loaded on CeO_2 preliminary suspended in water. The precipitation was carried under control of the following parameters: pH was maintained constant at 8 by adding drops of 0.1 mol.L⁻¹ NaOH solution, the temperature was fixed at 80 °C, and the stirring speed was constant. The slurry was stirred during four hours, filtered, washed with hot water, dried overnight in an oven 100 °C, and then calcined at 400 °C (1 °C.min⁻¹, 4 h, 2 L dried air.h⁻¹). The sample will be designated by 4% Rh/CeO₂ (DP).

2.2. Catalysts characterization

XRD measurements were performed with an automatic powder diffractometer BRÜKER AXS D8 Advance, using CuK α radiation and a scintillation counter LynxEye. The diffraction patterns were recorded in a step-scan mode with a step of 0.02° (2 θ), counting time 2 s, in the angular interval 20–80° at room temperature. The Eva program was used for diffraction data processing. Identification of the crystalline phases was made with the Joint Committee on Powder Diffraction Standards (JCPDS) files from the International Center for Diffraction Data (ICDD). Differential scanning calorimetry and gravimetric analysis (DSC-TG) was conducted in air flow (75 mL.min⁻¹) at a heating rate of 5 °C.min⁻¹ from room temperature to 400 °C followed by an isotherm at the same temperature for four hours with about 30 mg of sample with a Netzsch STA 409 apparatus.

EPR measurements were performed at two different temperatures: – 196 °C and room temperature, on an EMX BRUKER spectrometer with a cavity operating at a frequency of 9.5 GHz (X band). The measurements were performed on 50–55 mg of the calcined solids previously treated under vacuum (1.2×10^{-5} bar). The magnetic field was modulated at 100 kHz and the power supply was sufficiently low to avoid saturation effects. The g values were determined from precise frequency and magnetic field values.

The temperature-programmed reduction (TPR) measurements were performed on a device AMI-200 from ZETON ALTAMIRA. A trap for removing water during reduction was mounted in the gas line prior to the thermal conductivity detector. A hydrogen-argon mixture (5% H₂/Ar) was used to reduce the samples at a flow rate of 30 mL.min⁻¹. The temperature was linearly raised at a rate of 5 °C.min⁻¹ from room temperature till 900 °C and then the sample was kept one hour at 900 °C. The current of the thermal conductivity detector was 75 mA. Before the reducing step, the sample was flushed by an argon flow of 30 mL.min⁻¹ during one hour at 150 °C.min⁻¹ (5 °C.min⁻¹). The sample mass used was about 50 mg.



Fig. 1. Determination of $T_{\rm i}$ (beginning of CB combustion under air), $T_{\rm max}$ (maximum combustion rate), and $T_{\rm f}$ (end of CB combustion) on the DSC and TG curves.

2.3. Catalytic test

The catalysts were tested in the total oxidation of propylene. The experiments were carried out in a conventional fixed bed micro reactor between 25 °C and 500 °C (1 °C.min⁻¹). The reactive flow (100 mL.min⁻¹) consisted of a mixture of air and 6000 ppm of propylene. Before the catalytic test, the solid (100 mg) was activated under a flow of air (2 L.h⁻¹) at 400 °C (1 °C.min⁻¹). The reactants and products were identified and quantified by a Varian CP-4900 micro gas chromatography equipped with thermal conductivity detectors (TCD).

The carbon black (CB)–catalyst mixture was prepared in tight contact conditions. 2 wt% of CB and 98 wt% of catalyst were manually grinded for 15 min. About 30 mg of CB–catalyst mixture were loaded in an alumina crucible and heated from room temperature to 1000 °C (heating rate: 5 °C min⁻¹) in air flow (75 mL·min⁻¹). Three characteristic temperature values are determined from DSC–TG curves (Fig. 1): T_i where the CB oxidation begins, T_{max} where the rate of oxidation is at its maximum, and T_f where the combustion ends. T_{max} values were used to compare the catalytic performances and CB oxidation rates.

3. Results and discussion

3.1. Physico-chemical characterization

3.1.1. X-ray diffraction

Fig. 2 shows the XRD patterns obtained for both 4% Rh/CeO₂ (DP) and (Imp). The lines attributed to CeO₂ cerianite phase (JCPDS-ICDD 34-0394) located at 20: 28.55°; 33.07°; 47.48°; 56.34°; 59.44°; 69.86°; 77.08°; and 79.83°, were observed for both solids. In addition, a line with a weak intensity, located at 35.2°, was observed and assigned to Rh₂O₃ hexagonal phase (JCPDS-ICDD 41-0541). This line was absent in the XRD pattern of 4% Rh/CeO₂ (Imp). Mulukutla et al. [13] were also able to detect Rh₂O₃ by XRD in rhodium incorporated into MCM-41.

3.1.2. Thermal analysis

Fig. 3a and b show the DSC and TG curves obtained respectively for 4% Rh/CeO₂ (DP) and 4% Rh/CeO₂ (Imp), before calcination, heated under dry air at a rate of 5 °C·min⁻¹ from room temperature up to 400 °C followed by a 4 h dwell at the final temperature.

For the DP solid, the percentage of weight loss from room temperature up to 400 °C and before the 4 h-dwell at this latter temperature is of 3.1%. This loss occurs in two steps: the first one is due to the loss of physisorbed water on the solid (2.1%) and the second one to the desorption of water present in RhCl₃·xH₂O (1%). Indeed, when this latter precursor was brought from room temperature up to 400 °C and kept at this temperature for 4 h (Fig. 3c), the total weight loss recorded is about



Fig. 2. XRD patterns of 4% Rh/CeO₂ (DP) and 4% Rh/CeO₂ (Imp).

24%. Since, in the prepared catalyst, the rhodium weight corresponds to 4% of the catalyst total weight, the water loss resulting from the precursor dehydration must be equal to about $24 \times \frac{4}{100} = 0.96\%$, value close to 1%, the value obtained for the second loss step. In addition, the very slight gain of weight (0.1–0.2%) after 150 min of calcination at 400 °C can be due to the oxidation of RhCl₃ into Rh₂O₃ -detected by



Fig. 4. EPR spectra recorded at -196 °C of: a) 4% Rh/CeO₂ (DP) evacuated at room temperature for 1 h, b) 4% Rh/CeO₂ (DP) evacuated at 400 °C for 1 h, c) 1% HCl/CeO₂ evacuated at 400 °C for 1 h.

XRD- and to the formation of CeOCl or Ce-Cl species on ceria surface as reported by Force et al. [10]. These formations are illustrated by the following reaction:

 $2RhCl_3/CeO_2 + 4CeO_2 + \frac{3}{2}O_2 \rightarrow Rh_2O_3/6ClCeO_2$

For the Imp solid, the weight loss also takes place in two steps, with a total percentage of 4.7%. The first step, 3.4%, corresponds to the loss of physisorbed water and the second step, 1.3%, to dehydration of the precursor RhCl₃·xH₂O. No trace of gain was detected during the 4 h-dwell at 400 °C. This result can explain the absence of Rh₂O₃ formation in the impregnated solid.

3.1.3. Electron paramagnetic resonance

3.1.3.1. 4% Rh/CeO₂ (DP). The EPR spectrum, recorded at -196 °C, of 4% Rh/CeO₂ (DP) subjected to vacuum at room temperature during 1 h, is given in Fig. 4a. It is a complex spectrum. It is apparently formed by the superimposition of three signals. The first one, Ce³⁺, has an axial symmetry with $g_{\perp} = 1.965$ and $g_{//} = 1.945$. This signal was extensively studied by numerous authors [1,3,14–16] and was attributed to Ce³⁺



Fig. 3. DSC-TG analysis under air flow (75 mL.min⁻¹) at a heating rate of 5 °C.min⁻¹ from room temperature to 400 °C followed by an isotherm at 400 °C for four hours: a) 4% Rh/CeO₂ (DP), b) 4% Rh/CeO₂ (Imp), and c) RhCl₃xH₂O.



Fig. 5. EPR spectra recorded at room temperature of: a) $4\% \text{ Rh/CeO}_2$ (DP) evacuated at room temperature for 1 h, b) $4\% \text{ Rh/CeO}_2$ (DP) evacuated at 400 °C for 1 h.

ions. The second one, nominated O_D , possessing an axial symmetry with $g_{//}=2.030,\ g_{\perp}=2.010,\ and\ g_{iso}=2.017$ was unambiguously attributed to O_2^- species located on Ce^{4+} ions of ceria. Indeed, a similar signal with a slight difference in the $g_{//}$ value was obtained after adsorption of molecular O_2 on pure ceria [17–20]. The O_D signal disappeared when the spectrum is recorded at room temperature (Fig. 5a). This means that the O_2^- species is weakly adsorbed on ceria. The third signal, R_D , is symmetric and centred at g=2.17 with peak-to-peak width $\Delta H=96$ G. This latter can be attributed to Rh^{4+} (Z=45; 4d^5 with S=5/2; 3/2; 1/2) of RhO_2 small particles [21–23]. This latter oxide can be formed beside of the Rh_2O_3 one but in an amount not detectable by XRD. In addition, since the fine structure is not observed, it could be said that S=1/2.

The EPR spectrum, recorded at -196 °C, of 4% Rh/CeO₂ (DP) subjected to vacuum at 400 °C during 1 h, is illustrated in Fig. 4b. This kind of treatment can be considered as a smooth reduction of the solid. Apparently, beside the Ce³⁺ and R_D signals, the spectrum is formed of four signals: i) the first one O_P, apparently similar to O_D, with a little difference in the $g_{//}$ value (2.032), but, on the contrary, the O_2^- species corresponding to O_P remained stable when the spectrum is recorded at room temperature (Fig. 5b); ii) an isotropic signal, called S_D, centred at g = 2.0015 with a peak-to-peak width $\Delta H = 10$ G, it was attributed to a trapped electron in the ceria matrix [24,25]; iii) a signal, nominated R_{P} , with an axial symmetry is characterized by $g_{//} = 2.16$, $A_{//} = 115$ G, $g_{\perp} = 1.984$, $A_{\perp} = 19$ G, and $g_{iso} = 2.043$ [26–29]. Since the signal possesses a hyperfine structure of two lines, it was unambiguously attributed to Rh species (I = 1/2). It is well known that the metallic rhodium (Rh⁰; Z = 45; $5 s^{1} 4d^{8}$) cannot be detected by EPR except at low temperatures (less than -196 °C) [23,27,30] due to the presence of the unpaired electron in the s orbital. In addition, Rh^+ (4d⁸) and Rh^{3+} $(4d^6 \text{ with high spin})$ possess respectively a total electron spin S = 1 and 2. Since the fine structure is absent in the spectrum, therefore R_P signal could not be due to such species. Rh^{3+} (4d⁶ with low spin) has a S = 0 and is silent in EPR. Finally, the Rh^{2+} ions (4d⁷) has S = 3/2 in high spin and S = 1/2 in low spin. Since the fine structure is absent in the spectrum, hence R_P signal could be ascribed to Rh²⁺ low spin; iv) a symmetric signal, S_P , with $\Delta H = 15$ G and centred at g = 2.043. This latter value is equal to giso of the RP signal, consequently it can be attributed to Rh²⁺ species in the form of clusters. Since the R_D signal, attributed to Rh⁴⁺ ions, remained stable after this smooth reduction, this means that the $Rh^{2\,+}$ species responsible for the $R_{\rm P}$ signal can be obtained by reduction the Rh³⁺ ions of Rh₂O₃ oxide and not by the reduction of Rh4+ ions.



Fig. 6. EPR spectra recorded at -196 °C of 4% Rh/CeO₂ (Imp): a) evacuated at room temperature for 1 h, b) evacuated at 400 °C for 1 h.

The formation of O_D and S_D signals is due to the presence of Cl⁻ ions, coming from the RhCl₃ precursor, in ceria [24,31]. Indeed, 1 wt% Cl was impregnated on CeO₂ already calcined at 400 °C using a solution of HCl. The obtained product was calcined at 400 °C for 4 h. The EPR spectrum of this sample, after being subjected to O₂ adsorption, exhibited two signals similar to O_D and S_D (Fig. 4c). The small difference in the g values is due to the absence of rhodium on the latter solid.

When the EPR spectrum of 4% Rh/CeO₂ (DP) subjected to vacuum at 400 °C during 1 h is recorded at room temperature (Fig. 5b), the R_D, R_P, and S_P disappeared whereas, the Ce³⁺, O_P, and S_D remained observable but their intensities decreased, compared to those obtained at

 $-\,196\,^\circ\text{C},$ following the Curie law. The stability of O_2^- species responsible for the O_P signal is an indication of the solid oxidizing property.

3.1.3.2. 4% Rh/CeO₂ (Imp). Fig. 6a shows the EPR spectrum, recorded at -196 °C, of 4% Rh/CeO₂ (Imp) subjected to vacuum at room temperature during 1 h. The spectrum is composed of three signals: the first one was already attributed to Ce³⁺ ions, the second one, nominated R_I, is a broad symmetric signal centered at g = 2.16 with Δ H = 225 G, and the last one (D_I) is not well-resolved. When the spectrum is recorded at room temperature, the intensities of the above signals decreased following the Curie law. The R_I signal can be attributed to Rh⁴⁺ in the form of clusters. The oxidation of RhCl₃, which is in the form of large particles loaded on ceria surface due to the preparation method (impregnation), is responsible for the formation of Rh⁴⁺ clusters.

When the solid 4% Rh/CeO2 (Imp) was treated under vacuum at 400 °C for 1 h (Fig. 6b), the Ce³⁺ signal and R_I one disappeared whereas the D_I signal became well resolved with an axial symmetry and a hyperfine structure of two lines (g_// = 2.176, A_// = 53 G, g_{\perp} = 2.034, $A_{\perp} = 25$ G, and $g_{iso} = 2.081$) but its intensity is very weak (Fig. 6b). Such a signal, attributed to Rh²⁺ species, was obtained in rhodium doped lithium dithionate (Li₂S₂O₆·2H₂O) [28]. In our sample, these species can be obtained by a soft reduction of some RhCl₃ species which could resist to the oxidation during the calcination because of their localisation in the bulk of large particles. The disappearance of the R_I signal can be due to the reduction of Rh⁴⁺ into Rh³⁺ since this latter species is in general silent toward the EPR technique. The symmetric signal with a weak intensity, centred at g = 2.0015 (S_D), can unambiguously be attributed to a trapped electron in ceria matrix. Finally, the isotropic signal S_I centred at g = 2.29 with $\Delta H = 65$ G, can be due to traces of Rh²⁺ clusters [29,32,33] resulting from the reduction of some Rh³⁺ species obtained after reduction of Rh⁴⁺ clusters.



Fig. 7. TPR profiles: a) CeO₂, b) 4% Rh/CeO₂ (Imp), c) 4% Rh/CeO₂ (DP), d) 4% Rh₂O₃/CeO₂ (DP).

3.1.4. Temperature-programmed reduction

Fig. 7 shows the TPR profiles of CeO₂, 4% Rh/CeO₂ (Imp), 4% Rh/CeO₂ (DP), and 4% Rh₂O₃/CeO₂ (DP). The latter is a sample prepared by using Rh₂O₃ as precursor instead of RhCl₃. In the case of ceria (Fig. 7a), two broad reduction peaks were observed at 472 °C and 776 °C and were ascribed, respectively, to the reduction of ceria surface with the formation of Ce³⁺ species and an oxygen lacuna, and to the removal of bulk oxygen from the ceria structure along with the reduction of Ce⁴⁺ into Ce³⁺ ions [17,34–36].

Adding rhodium to ceria, following both DP and Imp methods, leads in one side to the shift of the two reduction peaks obtained for the pure ceria to lower temperatures and in the other side to the appearance of new reduction peaks at a lower range of temperatures (< 220 °C). The shift of ceria reduction peaks to lower temperatures suggests that rhodium makes ceria reduction easier. Furthermore, the amount of Ce⁴⁺ reduced into Ce³⁺ either on the surface or in the bulk of ceria, decreases after rhodium addition. The latter fact suggests a modification in ceria reduction behavior due to rhodium.

The peaks obtained at low temperatures (60 and 141 °C) for the 4% Rh/CeO₂ (DP) (Fig. 7c) can be due to Rh₂O₃ and O₂⁻ ions reduction. Indeed, the TPR profile of Rh₂O₃/CeO₂ (Fig. 7d) shows two peaks, respectively at 53 and 148 °C. These two temperatures are close to those obtained during 4% Rh/CeO₂ (DP) reduction. The small differences which exist could be due to the fact that in the latter, Rh₂O₃ is formed upon calcination of RhCl₃. The reduction peak at 60 °C was attributed to the reduction of Rh₂O₃ and O₂⁻ ions present on the CeO₂ surface whereas the peak at 141 °C could be due to the reduction of Rh₂O₃ and O₂⁻ ions rather located in the bulk (not on the surface) [37]. It was stated that the existence of two reduction peaks is due to the reduction



Fig. 8. Conversion of propylene during its total oxidation versus reaction temperature over CeO₂, 4% Rh/CeO₂ (DP), 4% Rh/CeO₂ (Imp), and 0.5% Rh/CeO₂ (Imp).

of Rh_2O_3 in different environments and/or with different particle size [13,38]. Rh_2O_3 is reduced into Rh^0 [36,39,40]. Conversely, Boutros et al. [38] stated that the first reduction peak (at lower temperature) is due to the reduction of Rh_2O_3 into Rh_2O and RhO and the second one to the reduction of Rh_2O and RhO.

The TPR profile of 4% Rh/CeO₂ (Imp) shows an intense peak at 82 °C and a small peak at 204 °C. The first peak could be ascribed to the reduction of the Rh⁴⁺ clusters –evidenced by EPR– into Rh³⁺ and Rh²⁺. Indeed, EPR showed the reduction of Rh⁴⁺ clusters after one hour under vacuum at 400 °C. The peak at 204 °C could be due to the reduction of some Rh³⁺ ions, located in the bulk, into Rh²⁺.

3.2. Catalytic tests

3.2.1. Oxidation of propylene

Fig. 8 illustrates the conversion of propylene versus the reaction temperature in the presence of the pure ceria support and the 4% Rh/ CeO_2 catalysts. The catalyst 0.5% Rh/ CeO_2 (Imp) was also evaluated. It is observed that addition of rhodium shifts the temperature of propylene conversion to lower values, whatever is the rhodium content or the preparation method. Furthermore, it is worthy to note that in all cases, propylene was exclusively converted into CO_2 . No other by-product was formed.

Furthermore, 4% Rh/CeO₂ (DP) is more reactive than 4% Rh/CeO₂ (Imp), with a ($T_{50(Imp)} - T_{50(DP)}$) of about 35 °C, T_{50} being the temperature of 50% propylene conversion. This result is in agreement with those reported in the literature [3,17,34,35] where the deposition–precipitation method gave better activities in oxidation reactions than the impregnation one. Indeed, the deposition–precipitation leads to the formation of nanosized particles dispersed on the surface of the solid, which are highly efficient in catalysis, whereas the impregnation method leads to the formation of large particles. The size of these particles depends in general on the active sites concentration added on the support. It decreases with this latter parameter and can become the same dimensions as those of nanoparticles. Indeed, it was demonstrated that when catalysts are prepared by the impregnation method with low rhodium contents, 0.5% Rh/CeO₂ (Imp), its catalytic activity is similar to that obtained for the 4% Rh/CeO₂ (DP).

Other factors could be also in the origin of the higher performance of the 4% Rh/CeO_2 (DP). This catalyst exhibits also the presence of O_2^- species along with the presence of Rh_2O_3 phase and also lower reduction temperatures in TPR than 4% Rh/CeO_2 (Imp). All these factors





could favor higher activities in oxidation reactions.

3.2.2. Combustion of carbon black

Fig. 9 shows the three temperatures, T_i , T_{max} , and T_f , obtained for the different catalysts during the total oxidation of the carbon black. T_{max} was used to compare the catalytic activity. The performance of 4% Rh/CeO₂ (DP) is better than that of CeO₂ particularly in T_i and T_f and much better than 4% Rh/CeO₂ (Imp) with a noticeable difference in temperatures. The better performance of the DP-solid could be due, as in the case of propylene, to the presence of Rh₂O₃ and O₂⁻ as well as to the better reducibility and lower particle size on the solid surface. Indeed, when the size of large particles decreases by decreasing the rhodium content in the Imp solid (0.5% Rh/CeO₂ (Imp)), the combustion activity increases and became similar to that obtained for the DPsolid.

4. Conclusion

The impregnation (Imp) and the deposition–precipitation (DP) methods were used to introduce 4 wt% of Rh, using RhCl₃ as precursor, on CeO₂. A calcination at 400 °C for 4 h followed this step. DP method results in the formation of weakly attached O_2^- species on ceria surface, identified by EPR, and in the formation of the oxide Rh₂O₃. Both species were not obtained by Imp method. These species are responsible

for the better catalytic performance of 4%Rh/CeO₂ (DP) solid in the total oxidation of propylene and carbon black reactions. The easier reducibility of rhodium oxides species and the lower size of rhodium particles are also in the origin of the higher catalytic activities of the solids prepared by DP method.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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