Ceria-supported metal nanoparticles for CO abatement in combustion exhaust gases

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Abstract: In order to reduce CO emissions in power and industrial plants, a series of ceria-supported catalysts have been synthesized and characterized. Encouraging preliminary catalytic tests show, at least for one sample, a reactivity comparable to the *state-of-the-art* Pt-based catalyst.

136 m²·g⁻¹

Two synthetic routes for supports were attempted: soft template using CTAB as templating agent (s-ceria; Fig. a)¹ and hard template using SBA-15 as templating agent (*h*-ceria; Fig. b).² Dimension of the particles, aspect ratio and surface area values varied depending on the procedure employed. Catalysts were produced by incipient wetness impregnation with nominal 5%_{mol} of Ni, Fe and Pt.¹



104 m²·g⁻

(a)



ATR-IR (Fig. c) identifies two peaks related to Ce-O v_{symm} at 250 cm⁻¹ (i)³ and v_{anti} at 500 cm⁻¹ (ii).⁴ Ceria template do not affect greatly crystal structure and dimensions (Fig. d), even though aspect ratio changes considerably; Ni and Fe metal active phase is not detectable by XRD prior and after testing (small particles), while Pt aggregates already in the synthetic step (Fig. e).



Fig. f) CO oxidation (CO = $2\% v/v; O_2 = 8\% v/v; N_2$ balance; 50 mg catalyst; $0.12 \cdot 10^6$ cm³ h⁻¹ g_{cat}⁻¹)⁵ performed with Fe@s-55% stops at ceria conversion (far from law requirements). Ni@s-ceria reaches 100% conversion at 400 aggregated °C vs the 300 °C of Pt; however, the exhaust excellent gases temperature is reached by close to 400 °C, making Ni a valid alternative to Pt for CO oxidation. explored in the future.

Conclusions

We optimized the syntheses of different ceria morphologies, characterized with a wide set of techniques. We deposited Fe and Ni active phases, with excellent dispersion level (undetected from XRD). Conversely Pt considerably; however, it retained its conversion, Ni-based catalyst, that will be further

References: ¹ Int. J. Hydrog. Ener., 2016, 41, 6316; ² Microporous Mesoporous Mater., 2016, 226, 466; ³ J. Sol-Gel Sci. Technol., 2009, 52, 356; ⁴ Sci. Tot. Environ., 2011, 409, 2987; ⁵ ACS Catal., 2020, 10, 11356.