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Correction: Improving time-resolution and sensitivity of *in situ* X-ray photoelectron spectroscopy of a powder catalyst by modulated excitation

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Correction for 'Improving time-resolution and sensitivity of *in situ* X-ray photoelectron spectroscopy of a powder catalyst by modulated excitation' by M. Roger *et al.*, *Chem. Sci.*, 2023, 14, 7482–7491, <https://doi.org/10.1039/D3SC01274C>.

The originally published version of this article contained an incorrect figure and figure caption for Fig. 5, in which the labels 'O₂ env.' and 'CO env.' in Fig. 5b and c, respectively, were inverted. The correct Fig. 5 and corresponding caption are displayed below. Two paragraphs that refer to Fig. 5 on page 7488 in the Results and discussion section under the sub-heading 'Identification of species on 5 wt% Pd/Al₂O₃' have also been updated to replace the original text.

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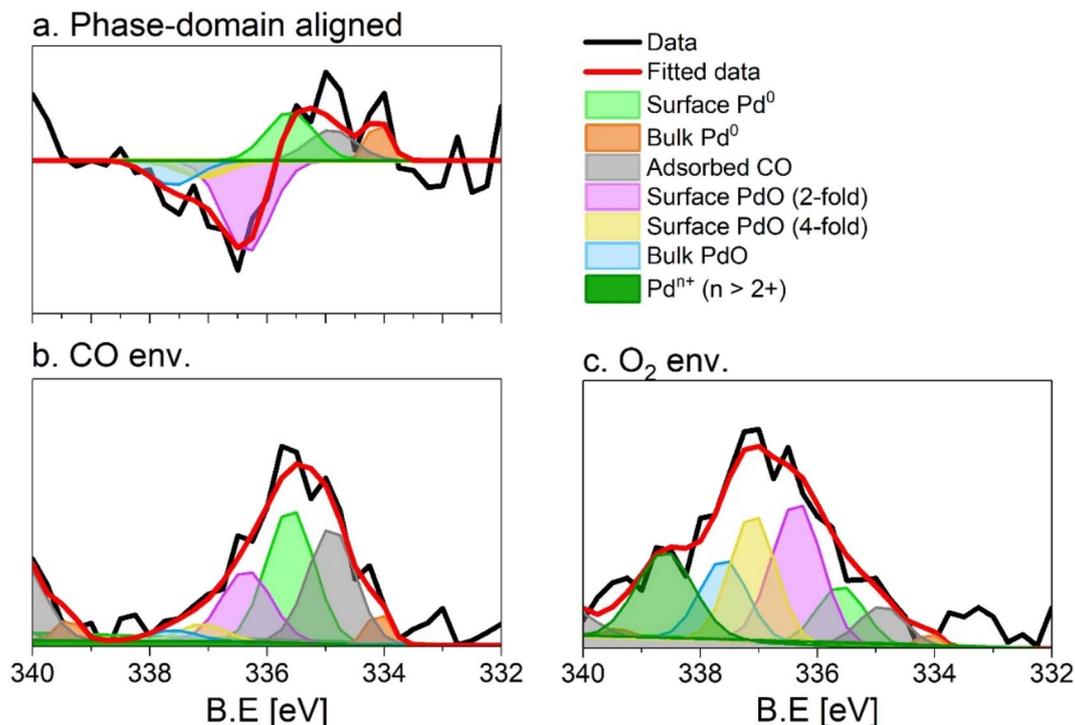


Fig. 5 Fit of the (a) phase (0°) and (b and c) time domain spectra ($t = 150$ s), X-ray photoelectron spectra of 5 wt% Pd/ Al_2O_3 acquired at the Pd 3d core level. (b) Averaged spectrum in reducing conditions, (c) averaged spectrum in oxidising conditions. Peaks are defined in Table 1. Only the Pd 3d_{5/2} core level is shown.

The fit of the averaged spectrum obtained at 150 s under oxidising conditions is dominated by cationic Pd species (Fig. 5c and Table 2). The contribution of Pd⁰ is low (7%), while surface Pd⁰ is absent. The fraction of Pd⁰ with adsorbed CO (11%) is not negligible and is attributed to the Pd surface poisoning by CO due to the strong Pd–CO bond. The poisoning effect of CO on Pd sites is supported by the very low levels of CO₂ all along the O₂ half-period in the MS data (Fig. S3), after a less pronounced sharp peak than that detected in the CO half-period. The presence of the signals of surface and bulk Pd⁰ under oxidizing conditions suggests that the thickness of the oxide layer formed is smaller than the mean escape depth of photoelectrons and thus that metallic Pd is coated by a thin skin of oxide.

Under reducing conditions, the fit of the averaged spectrum at 150 s (Fig. 5b and Table 2) shows predominantly peaks corresponding to metallic Pd species as well as Pd⁰ with adsorbed CO together with minor contributions from bulk and surface PdO but no signal of Pdⁿ⁺. The poor contribution from the surface Pd⁰ (4%) is justified by the presence of adsorbed CO molecules.

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

