



## Retraction Note to: Revisiting the Evolution of IR Spectra of CO Adsorbed on Au Nanoparticles Supported on Non-reducible Supports

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**Retraction to: Topics in Catalysis (2020)  
63:1596–1605 <https://doi.org/10.1007/s11244-020-01372-2>**

The authors have retracted this Article because, after re-examination, some of the results were found to be no longer reliable.

In their article, the authors related the temporal evolution of IR bands of CO adsorbed on Au to a hypothetical reconstruction process. A recent work [1] pointed out that a similar effect could be induced by the presence of nickel through contamination by Ni(CO)<sub>4</sub>.

The authors re-examined their data and noted that up to 5 ppm of Ni(CO)<sub>4</sub> were unexpectedly observed in the effluent gases of the diffuse reflectance FT-IR (DRIFTS) cell that had not previously been noticed. No Ni(CO)<sub>4</sub> was detected in the high-purity 5% CO/He from Air Liquide S.A., which had been further purified using a liquid N<sub>2</sub> trap. Ni(CO)<sub>4</sub> could also not be detected when CO had been co-fed with O<sub>2</sub> or if the cell reactor had not previously been raised to 300 °C under H<sub>2</sub>. The authors conclude that part of the cell (from Harrick) had been corroded, probably by NO<sub>x</sub> used during the same period, and reacted with CO to form Ni(CO)<sub>4</sub> in situ, only when the nickel present had been reduced to the metallic state.

The authors attempted to reproduce the experiment reported in this Article over the same materials using a new Harrick cell and they could no longer observe the major Au–CO band shift from ca. 2106–2110 cm<sup>-1</sup> down to 2070–2060 cm<sup>-1</sup> with concomitant several-fold intensity increase that occurred in less than 10 min (see Figs. 3.A and 6.A). By contrast, they found that the band position and intensity remain constant over 60 min (Fig. 1).

The authors therefore conclude that the spectral evolutions reported in this Article were likely not the result of any restructuring, but rather Ni(CO)<sub>4</sub> contamination and deposition of Ni over the Au nanoparticles, leading to Ni–CO species being formed and replacing Au sites.

The authors note that a sample of Au encapsulated in a silicalite-1 zeolite that had been exposed to Ni(CO)<sub>4</sub> did not show this band shift, indicating that Ni(CO)<sub>4</sub> could not enter the pores of silicalite-1 at 50 °C.

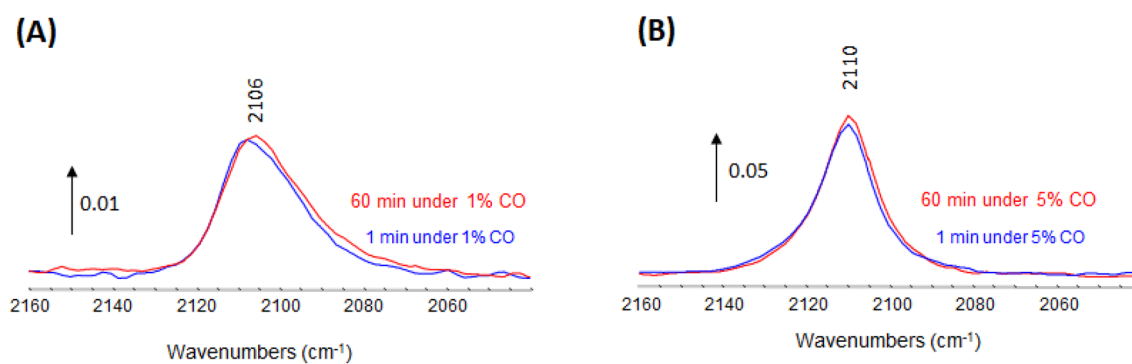
The authors hope that their experience will raise more awareness of the potential pitfalls associated with using CO at low temperatures on corroded steel cells.

All authors agree with this retraction.

The original article can be found online at <https://doi.org/10.1007/s11244-020-01372-2>.

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**Fig. 1** In situ DRIFTS spectra at 50 °C under 1% or 5% CO/He observed over **A** 2.1% Au/alumina and **B** 1% Au/silica. The samples were pre-reduced in situ with H<sub>2</sub> at 300 °C

## Reference

1. Yao Y, Chen L, Mao X, Yang Y, Chen J, Zhou L (2021) J Phys Chem C 125:8606–8619

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