Au-Modified Pd catalyst exhibits improved activity and stability for NO direct decomposition

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The direct decomposition of NO, at moderate temperatures is a challenge because of the poisoning of the surface of noble metals by oxygen, yet the modification of Pd with Au enables obtaining a highly stable catalyst under an O₂-free feed at 450 °C. Care was taken to ensure that the reaction was not due to the reduction of NOx with carbonaceous residues by monitoring CO₂ evolution. Pd–Au alloyed domains with a wide range of compositions were evidenced by in situ and ex situ XRD, the Pd content of which decreased with the increasing oxidising nature of the feed due to phase segregation giving rise to PdO and an Au-rich alloy. Under air, the limiting alloy composition was Pd₀.₂Au₀.₈. The catalyst was yet inactive in the presence of 5% O₂.

The proportion of vehicles using internal combustion engines will remain significant in the foreseeable future and the drive to reduce CO₂ emissions will favor the use of diesel and lean-burn engines. Yet, these engines exhibit significant drawbacks due to the emissions of NOₓ, particulate matters and organic pollutants (e.g. unburnt hydrocarbons and formaldehyde, notably present when using natural gas as fuel). The corresponding emission control technologies are complex, e.g. NOₓ-storage-reduction, NH₃-selective catalytic reduction (SCR) of NOₓ with hydrocarbons in which oxygen atoms strongly inhibit the reaction through competitive adsorption. In addition, noble metals can become poisoned through extensive oxidation leading to the formation of a less active oxidic phase. This led to the interest in selective catalytic reduction (SCR) of NOₓ with hydrocarbons in which oxygen atoms strongly bound to the noble metal could be removed by forming CO₂ and water, though this method only led to limited practical applications in the case of heavy-duty engines. Another approach was proposed by Toyota researchers consisting in Au- and Ir-modifications of Pt/alumina to

10% O₂, yet the rates are too low to be of practical interest and it is strongly poisoned by SO₂. NO decomposition can be carried out on perovskites, but only at reaction temperatures higher than 600 °C.

Many metals and oxides catalyse NO decomposition, but O₂ inhibits the reaction through competitive adsorption. In addition, noble metals can become poisoned through extensive oxidation leading to the formation of a less active oxidic phase. This led to the interest in selective catalytic reduction (SCR) of NOₓ with hydrocarbons in which oxygen atoms strongly bound to the noble metal could be removed by forming CO₂ and water, though this method only led to limited practical applications in the case of heavy-duty engines.

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Fig. 1 Thermodynamic equilibrium concentrations of NO, NO₂ and N₂O in a system initially containing 200 ppm of NO and 10% O₂ in N₂ at a total pressure of 1 bar. The much larger equilibrium concentrations of N₂ (ca. 90%) and O₂ (ca. 10%) are not shown. The data were calculated using the HSC Chemistry 6.2 software.
facilitate oxygen desorption from Pt.\textsuperscript{18} The improved NO decomposition activity obtained was assigned to the better resistance against oxidation brought about by Au. However, the role of individual promoters was not determined since only trimetallics Pt–Au–Ir were studied. Yet, the Au-rich sample appeared totally inactive, pointing out to the inability of Au alone to decompose NO. Bimetallics Ir–Au supported on alumina and dendrimers were reported later,\textsuperscript{19} though the activity was only measured in transient modes, making difficult differentiating true catalytic activity and adsorption phenomena, or even the potential reduction of NO with residues from organic dendrimers.

Au oxide is among the least stable of metal oxides, as shown by the plots giving the Gibbs energy of decomposition of various oxides of noble metals (active for NO decomposition) over the temperature range of interest and in the presence of 10\% O\textsubscript{2} (Fig. 2).

This observation led us to consider alloys based on Au and Pd, since (i) these metals are fully miscible (in contrast to Au and Pt), (ii) Pd is active for NO decomposition\textsuperscript{10} and (iii) Au should facilitate oxygen desorption as suggested earlier.\textsuperscript{18} Au was already proposed to protect Pd against oxidation by oxygen,\textsuperscript{20} sulfur\textsuperscript{20} and chloride\textsuperscript{21} through geometric (e.g. Pd site dilution) and electronic (e.g. electronic transfer from Au to Pd) effects, when used for hydrodechlorination or hydrodesulfurization.

Facilitating oxygen release from Pd at moderate temperature is a challenge, as the heat of adsorption of O atoms on metallic Pd is high (\textit{i.e.} 364 kJ mol\textsuperscript{-1}).\textsuperscript{22} Reddy et al.\textsuperscript{23} reported on the nature and activity of Pd phases during NO decomposition over silica-supported catalysts. NO decomposition was found to be mostly stoichiometric (\textit{i.e.} non-catalytic) at temperatures lower than 500 °C, by which metallic Pd was converted into inert PdO. PdO was shown to decompose significantly to metallic Pd and O\textsubscript{2} only above ca. 500 °C, temperature above which truly catalytic NO decomposition could occur.

The support used here was a commercial silica (surface area of 275 m\textsuperscript{2} g\textsuperscript{-1}), which was first functionalized by grafting APTMS (3-amino-propyltrimethoxysilane) according to a method reported elsewhere.\textsuperscript{24} The two monometallic catalysts were prepared using wet impregnation.\textsuperscript{25} The metal precursors were H\textsubscript{3}AuCl\textsubscript{4}.3H\textsubscript{2}O and Pd(NO\textsubscript{3})\textsubscript{2}.xH\textsubscript{2}O, both from Aldrich. The metals were reduced with NaBH\textsubscript{4} at room temperature. The bimetallic catalyst was prepared using the same batch of the monometallic Pd catalyst. Au was deposited on Pd using a surface redox method\textsuperscript{26} by which Au was reduced over the Pd/silica suspension in water in which H\textsubscript{2} was bubbled at room temperature. The final suspension was filtered and washed several times with distilled water and then kept in an oven overnight at 90 °C. After the preparation, the catalysts were kept in a refrigerator in the dark to avoid any sintering or ageing effects on the Au-based particles.

The metal loadings were determined by ICP analyses. The parent Pd/SiO\textsubscript{2} comprised 4.3 ± 0.3 wt\% of Pd and the bimetallic had a composition of 4.7 ± 0.3 wt\% Pd + 1.0 ± 0.2 wt\% Au/SiO\textsubscript{2}. The loading of the reference Au/SiO\textsubscript{2} was 1.0 ± 0.2 wt\%. The TEM analysis (using a Jeol 2010 LaB\textsubscript{6}) of the Au/SiO\textsubscript{2} revealed particle sizes comprised between 2 and 9 nm and a surface-weighted diameter $d_{sw} = (\sum n_i d_i^3/\sum n_i d_i^2)$ of 4.7 nm (Fig. 3, top). The parent Pd/SiO\textsubscript{2} showed particle sizes ranging mostly from 2 to 5 nm and a few larger particles up to 14 nm (Fig. 3, middle) and the $d_{sw}$ was 9.1 nm. The bimetallic catalyst exhibited a similar particle size.

![Fig. 2. Gibbs energy of decomposition $M_xO_y \rightarrow x M + y/2 O_2$ for several bulk metal oxides (in equilibrium with 10 kPa of $O_2$). Data calculated using the HSC Chemistry 6.2 software.](image1)

![Fig. 3. TEM and particle size distributions of (top) Au/SiO\textsubscript{2}, (middle) Pd/SiO\textsubscript{2} and (bottom) Pd-Au/SiO\textsubscript{2}. The samples were calcined in flowing air at 500 °C and then reduced with $H_2$ at 450 °C. More than 350 particles were counted for each sample.](image2)
distribution, with a few larger particles up to 24 nm, and $d_{\text{sw}} = 10.4$ nm (Fig. 3, bottom).

TEM-EDX microanalyses were carried out over 20 nanoparticles of two aliquots of the 4.7 wt% Pd-1.0 wt% Au bimetallic catalyst, one aliquot being calcined and then reduced by NaBH$_4$ at room temperature and the second aliquot being calcined and reduced by H$_2$ at 450 °C. These TEM-EDX analyses cannot be taken as being fully representative of the bimetallic catalyst due to the limited number of particles probed, yet those showed that the composition varied significantly from one particle to another in terms of Au and Pd proportions (Fig. 4). This indicates that the two metals were not homogeneously distributed over the sample, though most nanoparticles (about 90%) were bimetallic.

An in situ XRD analysis was carried out using an AntonPaar XRK900 reaction chamber fitted in a Bruker D8Advance. The thermal treatment consisted first of a calcination at 500 °C in a synthetic air flow, followed by a reduction in a H$_2$ stream at 450 °C, before returning to 25 °C under H$_2$ and finally air. The pattern of the SiO$_2$ support was subtracted.

The XRD patterns measured over the Au/SiO$_2$ (Fig. 5A) were essentially featureless, in agreement with the small Au particle sizes measured by TEM. A broad peak at ca. 38° corresponding to metallic Au was observed that hardly evolved with the thermal treatment (Fig. 5A.a to f).

In contrast, the Pd/SiO$_2$ patterns (Fig. 5B) exhibited strong peaks that evolved depending on temperature and the composition of the feed. The sample was poorly crystallised just after synthesis (Fig. 5B.a), but then exhibited a well-crystallised PdO pattern after calcination at 500 °C (Fig. 5B.b). The mere contact with H$_2$ at 25 °C led to the reduction of the PdO and the formation of a bulk hydride Pd$_2$H phase (Fig. 5B.c). The Pd$_2$H was decomposed by raising the temperature to 450 °C in H$_2$ and metallic Pd was formed (Fig. 5B.d, the offset from the Pd PDF file line is

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**Fig. 4** Proportion (in wt%) of Pd and Au measured by TEM-EDX over the SiO$_2$-supported bimetallic sample on various nanoparticles (top) calcined at 500 °C and reduced by NaBH$_4$ at room temperature and (bottom) calcined at 500 °C and then reduced with H$_2$ at 450 °C.

**Fig. 5** In situ X-ray diffraction patterns of the SiO$_2$-supported (A) Au, (B) Pd and (C) Pd-Au. The patterns were measured successively at (a) 25 °C in air after synthesis, (b) 25 °C in air after oxidation at 500 °C, (c) 25 °C in H$_2$, (d) 450 °C in H$_2$, (e) 25 °C in H$_2$ and (f) 25 °C in air. The patterns were offset for the sake of clarity. The dotted lines indicate the positions associated with the corresponding ICDD powder diffraction files.
exactly due to the thermal expansion of the lattice parameter from 25 °C to 450 °C. The Pd$_3$H phase reformed when going back to 25 °C (Fig. 5B.e), but the peak width was narrower, indicating that the thermal treatment had increased the size of the crystalline domains. The hydride phase decomposed upon H$_2$ being replaced with N$_2$ then air at 25 °C leading to metallic Pd peak (Fig. 5B.f), probably covered with chemisorbed oxygen.

The XRD results obtained on the Pd/SiO$_2$ stress that the reduction of PdO is fast, even at 25 °C, once the sample is exposed to H$_2$. In contrast, the oxidation of Pd to PdO is slow and requires a high temperature calcination. The latter observation is consistent with earlier reports that Pd oxidation to bulk PdO is slow and limited by O diffusion, while O$_2$ is easily activated at the surface of Pd particles forming chemisorbed O species.$^{27,28}$

The size of the Pd crystalline domains over the Pd/SiO$_2$ after reduction at 450 °C were determined using Scherrer equation and found to be around 13 nm. This represents the higher range of the particles measured by TEM (Fig. 3, middle), showing that the XRD signal is mostly representative of the (few) larger particles present in the sample.

The analysis of the XRD patterns obtained in the case of the bimetallic Pd–Au sample (Fig. 5C) was complicated by the fact that Au and Pd$_3$H exhibit a major peak roughly at the same position (ca. 38.3°). To make things worse, alloyed phases Pd$_x$Au$_{1-x}$ are expected to show peaks in between those of Au and Pd. As a consequence, the presence of pure Au can only be assessed whenever no Pd hydride is present, hence at 450 °C or in air. The corresponding Pd–Au/SiO$_2$ patterns (Fig. 5C.b, d and f) did not show any significant contribution at the position corresponding to pure Au, indicating that most gold was poorly crystallised or alloyed with Pd (note that the global molar ratio Pd/Au is about 9).

A large peak at the position of pure Pd (ca. 40°) indicated the presence of Pd domains (Fig. 5C.d and f), in agreement with the TEM-EDX data showing the presence of a few pure or Pd-rich particles (Fig. 4). In fact, the nature and evolution of the pattern observed over the bimetallic sample was similar to those observed over the pure Pd/SiO$_2$, with the exception of broader bands being present at positions in between those of Au and Pd that can be assigned to alloyed phases (compare signals in the dotted boxes in Fig. 5B and C for patterns b, d and f).

Three patterns of the bimetallic sample are shown in Fig. 6 to better highlight the peaks associated with alloyed phases. The pattern obtained after the first calcination at 500 °C exhibited PdO peaks (34° and 42°) and a peak at ca. 38.5° (Fig. 6a). The peak at 38.5° correspond to an alloy of composition Pd$_{0.2}$Au$_{0.8}$ based on Vegard’s law. It is interesting to note that this composition corresponds roughly to the limit (i.e. Pd$_{0.15}$Au$_{0.85}$) below which the Pd content of the alloy phase cannot be lowered when oxidising Pd–Au alloy films as those segregate into PdO and an Au-rich alloy phase.$^{29}$

Fig. 6b corresponds to the pattern of the bimetallic Pd–Au sample at 25 °C in air after further reduction at 450 °C in H$_2$ (Fig. 5C.f) to which the weighted Pd pattern collected in the same conditions (Fig. 5B.f) was subtracted. The obtained pattern revealed the superposition of several peaks comprised between those of pure Au and pure Pd that can be assigned to Pd–Au alloyed phases with various compositions (Fig. 6b). In view of the positions of these peaks, their Pd content appears to be higher than that of the Pd$_{0.2}$Au$_{0.8}$ phase observed after calcination (20 mol% Pd). This result is again consistent with the TEM-EDX data (Fig. 4) that show a wide range of Pd/Au compositions with many particles being rich in Pd.

A further oxidation of the bimetallic sample led to a pattern (Fig. 6c) that was similar to that collected initially (Fig. 6a) with PdO and an alloyed phase Pd$_{0.2}$Au$_{0.8}$. This shows the reversibility of the alloying-dealloying process upon reduction–oxidation cycles.

Using the area of the XRD peaks of the Pd and Pd–Au phases for a rough quantitative analysis, the bimetallic sample after reduction contained about 60 wt% of a phase corresponding to pure or almost pure Pd and about 40 wt% of alloyed Pd–Au phases. The quantity of free Au was negligible. It should be reminded at this point that the XRD technique cannot determine whether or not the surface of the “pure” Pd particles would be partly covered with Au atoms. The presence of some Au atoms segregated at the surface can be expected because of the significantly lower surface tension of Au (1145 mJ m$^{-2}$) as compared to that of Pd (1482 mJ m$^{-2}$).$^{30}$ In addition, we stressed above that the XRD data are mostly representative of the larger nanoparticles present, hence the corresponding compositions and reactivities do not encompass those of the smaller nanoparticles. Nonetheless, these XRD data indicate that a fraction of Pd atoms was made oxidation-resistant by forming an alloy (i.e. Pd$_{0.2}$Au$_{0.8}$) with Au.
The catalytic activity for the decomposition of NO was studied using a setup described in details elsewhere. In brief 100 mg of as-prepared catalyst was placed in a U-shaped quartz reactor on a quartz wool plug. The catalyst was reduced at 200 °C under 50% H₂ in He, before being flushed with He. A reaction flow of 50 mL min⁻¹ comprising 500 ppm of NO + 5% O₂ in He was initially used. Reactor effluents were analysed by a combination of μ-gas chromatography (CP490, Agilent), mass spectrometry (Omnistar, Pfeiffer Vacuum) and IR gas cell analyses (MKS MultiGas 2030, MKS). The temperature was raised step-wise to 450 °C.

The evolution of products measured over the non-calcined Pd–Au/SiO₂ revealed significant N₂ and N₂O production, which correlated with the evolution of CO₂ (Fig. 7). This indicates that APTMS residues present at the surface of the silica support were able to react with NOₓ and lead to NOₓ reduction. This stresses that catalysts should be thoroughly calcined before carrying out NOₓ direct decomposition to prevent the risk of unwanted reduction.

The samples were left under 500 ppm NO + 5% O₂ for 10 h at 450 °C to burn off all carbonaceous residues, which was ascertained by the absence of CO₂ in the reactor effluent (Fig. 7). Unfortunately, the activity of the Pd–Au/SiO₂ dropped to zero once CO₂ production ceased, stressing that this catalyst was not active for NO decomposition in the presence of 5% O₂.

The catalysts were then reduced in situ under 50% H₂ in He at 450 °C for 1 h to reduce the PdO and ensure the realloying of Pd and Au. The NO decomposition activity of the catalysts was then measured isothermally at 450 °C over 1000 min. O₂ was left out of the feed this time to limit potential fast deactivation due to O-poisoning, as suggested by the initial deactivation observed at 200 °C that was not correlated to the CO₂ signal (Fig. 7).

It is important to note that Au/SiO₂ was essentially not active for NO decomposition reaction under these conditions, as the level of N₂ + N₂O formed (ca. 3 ppm) was close to the precision of the measure (Fig. 8, top). Interestingly, the Au-promoted Pd sample appeared significantly more active than the parent Pd/SiO₂ sample, indicating that a synergy existed between these two metals (Fig. 8, top).

In addition, the bimetallic sample activity was stable over the duration of the experiment (following a fast but limited initial deactivation), while that of the monometallic parent sample was continuously decaying. The concentrations of O₂ and NO₂ released during the reaction showed that the release of oxygen was significantly greater in the case of the bimetallic sample (Fig. 9). These observations support the conclusions derived from the XRD data that Au induced a greater resistance against oxidation of at least a fraction of the Pd atoms, which may explain the stability of the Au-promoted Pd catalyst.

The selectivity to N₂ measured over these three catalysts is reported in Fig. 8, bottom. (Note: the complement to 100% represents the selectivity to N₂O, NO₂ being not considered as a reduction product). The high selectivity to N₂ measured over Au/SiO₂ should be taken with caution in view of the low
activity of this sample (part of the N\textsubscript{2} measured may be the results of minute leaks). The Pd/SiO\textsubscript{2} sample showed a rising selectivity to N\textsubscript{2} nearing 90\%, while that of the Pd-Au/SiO\textsubscript{2} remained essentially constant at 35\%. This significant selectivity difference suggests that the Pd active sites were modified by Au, likely through the dilution of the Pd centres that may have induced both geometric and electronic effects.

As reported in the literature\textsuperscript{29,32-34} and shown in Fig. 6, heating Pd-Au alloys in O\textsubscript{2} above 450 °C results in the formation of partially oxidised Pd (or even PdO) at the alloy surface and segregation. Similarly, a fraction of the Pd in the Pd-Au alloyed nanoparticles may have eventually got segregated under NO, a strong oxidizer. The XRD of the bimetallic sample was collected after the NO decomposition reaction shown in Fig. 8 and 9. Interestingly, the pattern obtained after NO decomposition at 450 °C exhibited peaks of PdO, alloyed phases Pd\textsubscript{x}Au\textsubscript{1-x} and Pd (Fig. 10c). This pattern is therefore intermediate between that obtained after reduction at 450 °C showing Pd\textsubscript{0.2}Au\textsubscript{0.8} and Pd (Fig. 10a) and that obtained after calcination at 450 °C showing PdO and Pd\textsubscript{0.2}Au\textsubscript{0.8} (Fig. 10b). This can be simply rationalised by the fact that 500 ppm of NO has a lower oxidising potential than air.

The presence of PdO, Pd and Pd\textsubscript{x}Au\textsubscript{1-x} post-reaction stresses the complex speciation of Pd existing over the bimetallic catalyst, due to the wide range of particle sizes and composition existing in the as-prepared sample (Fig. 3 and 4). It is therefore difficult to infer on the nature of the most active and stable phase. Surprisingly, the Pd/SiO\textsubscript{2} post-reaction exhibited an almost negligible fraction of PdO (Fig. 10d), in contrast to the bimetallic sample (Fig. 10c). This observation can possibly be explained by an improved diffusion of O in the alloyed domains (presenting more defects), as O diffusion has been proposed as the limiting step during Pd oxidation.\textsuperscript{27,28}

The stability of the bimetallic sample during NO decomposition (Fig. 8, top) indicates yet that under our reactions conditions most bimetallic nanoparticles exhibited a sufficient Au fraction at the surface to prevent deactivation, in agreement with the TEM-EDX analysis (Fig. 4). The XRD analysis was yet unable to capture this nanoscale insight by pointing out a large fraction of seemingly pure Pd domains, which may have actually been superficially covered with Au. Again, the XRD analysis is likely to have overemphasized the signal of larger particles that are less important for catalysis.

In situ characterization methods would need to be employed to determine the nature of the active surface, since surface segregation is likely to occur when altering the atmosphere composition, even at room temperature, as reported in the case of CO adsorption over Au–Pd and other nanoalloys.\textsuperscript{35,36} We have carried out operando DRIFTS analyses to try to monitor NO\textsubscript{2} species adsorbed on the catalyst under reaction conditions at 450 °C, but no meaningful signal could be obtained (data not shown).

In conclusion, catalysts should be free of carbonaceous deposits when investigating NO decomposition, as the unexpected presence of potential reductants (or using carbonaceous supports) could be a cause of data misinterpretation. Monitoring CO\textsubscript{2} production can help in assessing this point, as demonstrated here (Fig. 7). The addition of Au does promote the activity of Pd/SiO\textsubscript{2} leading to a more active catalyst that exhibited a stable activity under our reaction conditions over more than 16 h. This raises the prospect of developing better oxidation-resistant catalysts based on noble metals and Au for the direct decomposition of NO. The effect of water will be important to investigate because it is known to deactivate Pd-based catalysts used for methane oxidation by forming strongly bound surface hydroxyls.\textsuperscript{37} Water can also facilitate sintering.
Since the activity for NO decomposition in the presence of 5% O2 was essentially nil (Fig. 7), other formulations (possibly based on metals other than Pd) will have to be considered for practical applications in the presence of large O2 concentrations. Formulations based on Pt or Ag alloyed to Au are currently being investigated, following the insights given by the thermodynamics of oxide decomposition reported in Fig. 2. Another challenge will be to retain activity in the presence of SOx.

**Author contributions**

FCM designed the project, carried out the thermodynamic analysis, monitored the data and drafted the manuscript. TE carried out the catalyst synthesis and participated to the literature review and data analysis. LB carried out the TEM-EDX analysis. FB carried out the XRD analysis. AT and YS contributed to data analysis and manuscript writing. JS carried out the NO decomposition experiments, data analysis and reviewed the manuscript. LD provided support in the synthesis of the catalysts and reviewed the manuscript. CT built and validated the catalytic set-up, monitored the catalytic results and reviewed the manuscript.

**Conflicts of interest**

There are no conflicts to declare.

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