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Organosilane oxidation by water catalysed by large gold nanoparticles in a membrane reactor†

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We show that gold nanoparticles catalyse the oxidation of organosilanes using water as oxidant at ambient conditions. Remarkably, monodispersions of small gold particles (3.5 nm diameter) and large ones (6–18 nm diameter) give equally good conversion rates. This is important because separating large nanoparticles is much easier, and can be done using ultrafiltration instead of nanofiltration. We introduce a simple setup, constructed in-house, where the reaction products are extracted through a ceramic membrane under pressure, leaving the gold nanoparticles intact in the vessel. The nominal substrate/catalyst ratios are ca. 1800 : 1, with typical TONs of 1500–1600, and TOFs around 800 h⁻¹. But the actual activity of the large nanoparticles is much higher, because most of their gold atoms are “inside”, and therefore unavailable. Control experiments confirm that no gold escapes to the membrane permeate. The role of surface oxygen as a possible co-catalyst is discussed. Considering the ease of product separation and the robustness of the ceramic membrane, this approach opens opportunities for actual applications of gold catalysts in water oxidation reactions.

Silicon alcohols (silanols) are widely used in organic synthesis, particularly in cross-coupling reactions, and in industry as synths for silicon-based polymeric materials.1 Traditionally, they are synthesized by hydrolysis of halosilanes, stoichiometric oxidation of organosilanes, or reactions of siloxanes with alkali reagents.2 However, these are not clean processes. In contrast, the catalytic oxidation of silanes with water is environmentally benign. It produces silanols with a high selectivity and hydrogen gas as the only by-product. Catalysing chemical reactions in water at room temperature is one of the challenges facing chemists today.3 Oxidizing silanes to silanols is usually catalysed by noble metals, such as rhenium, ruthenium, iridium, silver and gold.

Although gold is traditionally considered (and prized) as inert, its nanoparticles (AuNPs) are enjoying a newly acquired status as catalysts for many reactions.4–17 Kaneda and co-workers18 showed that hydroxyapatite-supported gold particles (3.0 ± 0.9 nm) catalysed silane oxidation in water at 80 °C. Using a different approach, Asao et al.19 reported the use of nanoporous gold with pore size around 30 nm received from a gold–silver alloy for this type of reactions. Recently, carbon nanotube-supported AuNPs (AuCNT) were reported as highly active for silane oxidation in THF at room temperature.20 Gold nanoparticles also catalyse selective activation of alkynes, giving fast hydrogenation or hydration reactions,21 as well as catalysts for the selective conversion of biomass into chemicals22 and in fuel cell reactions.23 The particles’ size and shape, dictated by the preparation recipes,24 often determine the catalytic performance. Generally, the smaller the particle, the higher the catalytic activity, down to the three-layered Au55 cluster. Indeed, small gold particles (2–5 nm diameter) catalyse a variety of reactions.25–29 Exceptions are few yet notable, such as the unusually large 38 nm AuNPs reported as catalysts for hydrogen peroxide decomposition to hydroxyl radicals, enhancing the chemiluminescence of the luminol–H₂O₂ system.30

The problem is that although small AuNPs are exciting as catalysts, their size hampers practical application. Gold is too expensive a catalyst to throw away, and separating particles smaller than 2 nm is no mean feat. Acceptable Au separation levels often require repeated purification cycles or a combination of methods, increasing operational costs and decreasing throughput. One way to solve this is by using pressure-driven membrane separation, a relatively new technology. State-of-the-art nanofiltration (NF) membranes can retain even single-nm particles, but these membranes are expensive, delicate, and have low flow rates.28 Alternatively, separation using ultrafiltration...
(UF) membranes demands significantly lower transmembrane pressures, and the membranes are cheaper and more robust.\textsuperscript{31} The problem is that UF membranes are only effective for particles 8–10 nm diameter and larger.

In this paper we show that AuNPs as large as 18 nm can catalyze the oxidation of silanes to silanols with water, generating hydrogen as the only by-product (Scheme 1). We demonstrate the effectiveness of these reactions in a dedicated membrane pressure reactor, using a ceramic membrane for separating the product mixture from the catalyst. Using large AuNPs in such a reactor combines the advantages of high conversion rates and good flux values with full retention of the catalysts.

We prepared monodispersed gold aquasols of varying sizes using the citric acid/tannic acid method (Fig. 1 shows TEM images and gold nanoparticle size distribution; more details are given in the ESI\textsuperscript{1}). In a typical reaction (Scheme 1), organosilanes were oxidized in water in presence of gold nanoparticle catalyst under 1 bar N\textsubscript{2} pressure. A production of corresponding silanols was accompanied by a formation of hydrogen gas (this was quantified volumetrically, see ESI\textsuperscript{1}). Reactions were run for up to 24 h in a stainless steel high-pressure reactor, equipped with a membrane, constructed in-house. The gold was retained in the reactor by a porous ceramic membrane placed in the bottom, and the product was isolated and determined using \textsuperscript{1}H NMR and gas chromatography (full details and drawings in the ESI\textsuperscript{1}). The extraction of reaction products through the membrane was possible using only 4–5 bar additional nitrogen pressure.

Fig. 2 shows the conversion profiles of dimethylphenylsilane 1a (top) and the yield profiles for dimethylphenylsilanol 2a (bottom) vs. time in the presence of AuNPs of different sizes. The profiles are divided into small particles and large ones (left and right, respectively) for clarity. For comparison, we also performed a control reaction in the absence of any catalyst. These graphs show two important things. First, there is no significant difference in the conversion when using AuNPs up to 18 nm in diameter. This is important because the filtering of larger particles (>10 nm) is much easier, and can be done using standard ultrafiltration technology. Second, there is a marked difference between substrate conversion and product yield in the absence of AuNPs. Practically, this means that the formation of the disiloxane by-product is suppressed in the presence of the catalyst, as the silane is quickly consumed in the oxidation reaction.

![Scheme 1](image)

**Scheme 1** Oxidation of silanes to silanols with water using Au nanoparticles.

![Fig. 1](image)

**Fig. 1** Transmission electron micrographs of gold nanoparticle suspensions with different average diameters: 3.1 ± 0.5 nm (I); 4.8 ± 0.5 nm (II); 9.2 ± 0.8 nm (III). Size distribution of gold nanoparticle suspensions A–G, determined using dynamic light scattering (IV). The distribution function analysis is displayed as scattered intensity per particle size: A (3.1 nm), B (4.8 nm), C (9.2 nm), D (18.3 nm), E (27.5 nm), F (37.4 nm) and G (45 nm).

![Fig. 2](image)

**Fig. 2** Conversion profiles of dimethylphenylsilane 1a (top graphs) and yield profiles for dimethylphenylsilanol 2a (bottom graphs) in the presence of AuNPs of different diameters (3.1 nm, 4.8 nm, 18.3 nm, 37 nm and 45 nm). Reaction conditions: 2.93 mmol 1a, 103.3 mM acetone, 222 mM water, 0.31 mg catalyst (1.57 × 10\textsuperscript{–6} mol gold; 0.05% Au relative to substrate) and 296 K. All experiments were performed in duplicates.

Our dead-end membrane reactor enables quick and efficient separation of the AuNPs. This is easily monitored using UV-visible spectroscopy.\textsuperscript{32,33} Fig. 3 shows the spectra of the reaction mixture before and after the membrane.
Fig. 3 UV-visible spectra of the reaction mixture at 0, 1, 2 and 4 h for 18 nm AuNPs, compared with the spectrum of the membrane permeate. Note that the increase in the absorbance of the plasmon peak at 520 nm is due to the removal of solvent during the sampling, which increases the concentration of the NPs.34

We then examined the scope of the reaction, testing various aromatic and aliphatic organosilanes in the membrane reactor (structure 1a–e, Scheme 1). Table 1 shows the conversion, yield, and nominal TON values. Aliphatic silanes (entries 2, 5) were more active than the aromatic ones. In all cases, we observed complete retention of the nanoparticles within the membrane reactor. Moreover, the membrane could be reused several times.

We may anticipate that the reaction begins with the insertion of the Si–H bond on the Au nanoparticle to generate a silyl-metal hydride intermediate. The silyl-metal hydride intermediate is subsequently attacked by a nucleophile derived from water to generate the silanol (this attack is most likely an $S_N^2$ type, and therefore an inversion would be expected, since it would be easier for the oxygen to attack trans to the NP surface). However, this is only a suggestion. In the final step of the cycle, Au–H$^+$ and O$_4$⋯H$^+$ species should react to produce H$_2$, regenerating the catalytic surface (Scheme 2).

The critical step of the reaction is most likely the dissociative activation of water. Our observations show that this step does not depend on the particle size alone below a certain size. Recent surface science studies proposed that an oxygen atom on Au(111) surface (O$_4$) acts as a basic co-catalyst that promotes various organic reactions at low temperatures.35–38

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<th>Table 1 Water oxidation of various organosilanes to the corresponding silanols$^a$</th>
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$^a$ Reaction conditions: 0.2 g organosilane, 7.6 ml acetone, 10 ml deionized water, 0.31 mg AuNPs (6 nm) catalyst, reaction time 2 h. Total reaction volume 17.8 ml, magnetic stirring at 400 rpm, 296 K.

$^b$ GC yield, corrected for the presence of an external standard.

$^c$ Isolated yield.
presence of hydroxyl groups on the surface.\textsuperscript{43,44} Carbon-supported Au catalysts, which were inactive for the vapor-phase CO oxidation up to 373 K, showed excellent CO oxidation rates in the aqueous phase at 300 K.\textsuperscript{44} Moreover, a 50-fold increase in rate was observed on going from acidic to basic pH, though Au particles used in aqueous-phase studies were larger than 4 nm.\textsuperscript{43} All these facts show that the adsorbed species play a major role in enhancing the activity even for larger particles. A similar effect may be operating in our case, making larger particles as active as smaller ones.

Conclusions

Water oxidation of silanes to silanols is efficiently catalysed by “large” gold nanoparticles (6–18 nm in diameter). This catalytic oxidation has several advantages, namely using water as a clean oxidant, high activity and selectivity for silanols. Moreover, it works well with a number of silanes with different types of substituents. Combining this reaction with ultrafiltration membrane reactor technology solves the problem of difficult separation of nanoparticles from the reaction mixture, and opens opportunities for bona fide applications.

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Notes and references