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Metal–organic framework composites with luminescent gold(III) complexes. Strongly emissive and long-lived excited states in open air and photo-catalysis†

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The encapsulation of luminescent gold(III) complexes by metal–organic frameworks (MOFs) lays the groundwork for new phosphorescent materials with activities that are not readily achieved by the host MOF materials or gold(III) complexes alone. In this work, strong phosphorescence with lifetimes of up to ~50 μs in open air at room temperature has been achieved by incorporation of cationic cyclometalated gold(III) complexes into MOFs with anionic frameworks to form $\text{Au}^{\text{III}}@$ MOFs. The $\text{Au}^{\text{III}}@$ MOFs display solid state two-photon-induced phosphorescence. Photo-reduction of methyl viologen to the reduced radical was achieved inside $\text{Au}^{\text{III}}@$ MOFs and in the presence of Et_3N upon excitation at $\lambda > 370$ nm under ambient conditions. These $\text{Au}^{\text{III}}@$ MOFs comprise a class of reusable and size-selective heterogeneous photo-catalysts for the aerobic oxidation of secondary amines to imines as well as five other reactions, including oxidative C–H functionalization under aerobic conditions.

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Introduction

Metal–organic frameworks (MOFs) have received burgeoning interest over the past decades, with applications in gas storage/separation, proton/electrical conductivity, biomedicines, sensing, and catalysis.^{1,2} The tuning of both the metal nodes and the organic linkers in MOFs, by either pre- or post-synthetic^{1,3} methods involving covalent/dative bond formation, leads to extraordinary versatility in framework structure, porosity, and function. Encapsulation of a functional cargo into the pores of MOFs *via* non-covalent and/or dative interactions,⁴ such as the use of MOFs to encapsulate metal catalysts or luminescent metal ions/complexes by the cation exchange method,^{4a} is an alternative approach to MOF modification. Pores with defined environments inside MOFs constitute a unique platform to confine guest species, and as a result, new

functions/performance of the encapsulated guest species may emerge.

We are attracted to using MOFs for housing metal complexes with long-lived emissive excited states for heterogeneous photochemical catalysis and luminescence applications. In the literature, other porous materials besides MOFs, such as mesoporous silica, have also been employed to encapsulate photoluminescent and/or photochemically active compounds,^{5,6} including confinement of a photoluminescent trinuclear gold(I) pyrazolate complex in the channels of mesoporous silica.^{6b} Our attention has been directed towards recently reported luminescent gold(III) complexes which display strongly emissive excited states with long lifetimes of up to 506 μs (emission quantum yields: up to 11.4%) in degassed solutions at room temperature.⁷ These complexes may have useful applications in photo-catalysis and emission sensing, particularly if they are located in an aerobic environment in which their photo-physical characteristics can be retained, and if the photochemical reactions they catalyse can be endowed with specificity. Here we report our findings on the encapsulation of strongly luminescent gold(III) complexes with various dimensions ($\text{Au}1\text{--Au}4$,^{7,8} Scheme 1; Table S1 in the ESI†) by two MOFs with differing porous structures. The encapsulation was performed using a cation exchange method.^{4a,9} The $\text{Au}^{\text{III}}\text{--MOF}$ composites ($\text{Au}^{\text{III}}@$ MOFs) display long emission lifetimes of up to 48.8 μs at room temperature and in open air, exhibit solid state two-photon-induced phosphorescence, and can act as selective and recyclable catalysts for light-induced electron transfer reactions

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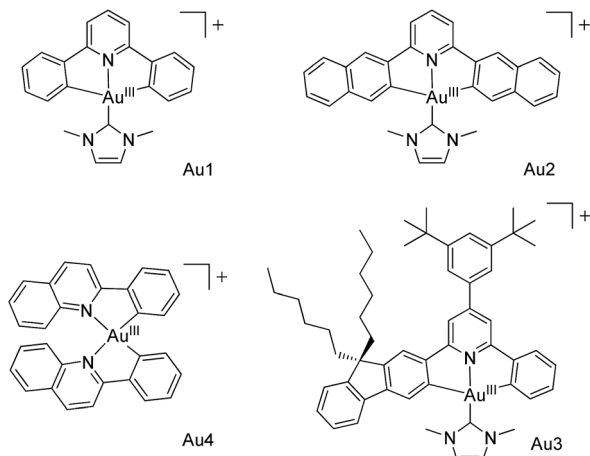
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Scheme 1 Au^{III} complexes Au1–Au4 used in this work.

and aerobic C–N/C–C/C–O bond forming reactions. Of note, Corma and co-workers¹⁰ have described a recyclable Au^{III}-functionalized MOF catalyst for domino coupling/cyclization and alkene hydrogenation, prepared by post-synthetic dative bond formation.

Results and discussion

Syntheses and structures of MOF1 and MOF2

Two MOFs, MOF1 and MOF2, built from similar ligands but possessing different types of pores with suitable window sizes, were chosen in this work to allow investigation of the effect of porous structure on the luminescence of Au(III) complexes.

MOF1, formulated as $[\text{Me}_2\text{NH}_2]_2[\text{Cd}_2(\text{TATMB})_2] \cdot 10\text{DMF} \cdot 3\text{H}_2\text{O}$ ($\text{H}_3\text{TATMB} = 3,3',3''\text{-}((1,3,5\text{-triazine-2,4,6-triyl})\text{tris(azanediy)})\text{tribenzoic acid}$) based on single-crystal X-ray diffraction, elemental analysis, thermogravimetric analysis (TGA), and charge-balance consideration, was obtained as colourless rod-like crystals by the reaction of H_3TATMB with $\text{Cd}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ in DMF at 120 °C for 2 days (the $[\text{Me}_2\text{NH}_2]^+$ cations came from decarbonylation of DMF¹¹). The phase purity of MOF1 was examined by powder X-ray diffraction (PXRD) analysis.

Single-crystal X-ray structure determination of MOF1 revealed a 3D framework (Fig. 1a) that can be described as a 4-connected uninodal net with hxg-d-4-C2/m topology¹² (Fig. S1d, ESI†). The asymmetric unit contains one crystallographically independent Cd^{2+} ion and one TATMB ligand (the other cations and guest molecules are disordered and have not been precisely defined). The Cd^{2+} ion is seven-coordinated by six carboxylate oxygen atoms of three TATMB ligands and one N atom from the triazine ring of another TATMB ligand; each pair of nearest-neighbour Cd^{2+} ions are bridged by two TATMB ligands to form an 18-membered ring (Fig. S1a, ESI†). There are large nano-scale tubular channels (window size: $17 \times 24 \text{ \AA}^2$, Fig. 1b) running along the $[023]$ plane, with solvent molecules and $[\text{Me}_2\text{NH}_2]^+$ ions residing in the channels. The effective free volume of MOF1 is 70.7% of the crystal volume (calculated by PLATON analysis¹³). The TGA data (Fig. S2, ESI†) revealed a decrease in weight of 38.4%, attributed to a loss of solvent and

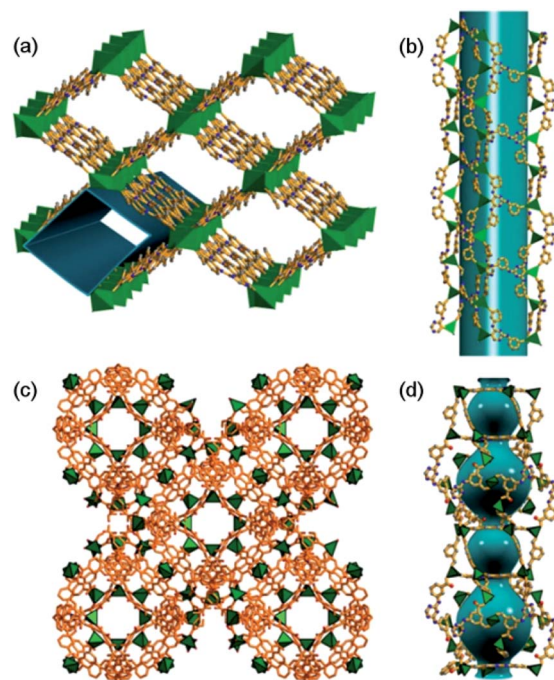


Fig. 1 (a) 3D polyhedral structure of MOF1, viewed along the c axis; (b) 1D mesoporous nanotube component of MOF1; (c) 3D polyhedral structure of MOF2, viewed along the a axis; (d) trigonal and hexagonal prism-shaped nanocages in MOF2.

water molecules; the de-solvated framework remained stable until $\sim 290 \text{ }^\circ\text{C}$ (above this temperature the framework began to decompose and the final residue was CdO [6.32% observed, 6.20% calculated]). N_2 adsorption measurements for MOF1 revealed an isotherm (Fig. S3, ESI†) which corresponds to a Type 1 adsorption isotherm for porous materials.¹⁴

MOF2, $[\text{Me}_2\text{NH}_2]_2[\text{Zn}(\text{TATAT})_{2/3}] \cdot 3\text{DMF} \cdot \text{H}_2\text{O}$ ($\text{TATAT} = 5,5',5''\text{-}((1,3,5\text{-triazine-2,4,6-triyl})\text{tris(azanediy)})\text{trisophthalate}$), was prepared according to the literature.¹⁵ MOF2 has a 3D meso-cage structure (Fig. 1c) featuring alternately arranged hexagonal prismatic cages (window size: $6.3 \times 10.5 \text{ \AA}^2$) and trigonal prismatic cages (window size: $14.3 \times 11.5 \text{ \AA}^2$) (Fig. 1d), with a potential solvent volume that accounts for 60% of the empty volume.¹⁵ The $[\text{Me}_2\text{NH}_2]^+$ cations and DMF/ H_2O molecules reside in these nano-scale cages.

Syntheses, characterization and photo-physical properties of Au^{III}@MOF composites

Au^{III}@MOFs, including Au^{III}@MOF1 (Au^{III} = Au1–Au4) and Au^{III}@MOF2 (Au^{III} = Au1, Au2), were obtained as yellow or pale yellow solids (Fig. S4, ESI†) by immersing MOF1 or MOF2 in a DMF solution of the CF_3SO_3^- or BF_4^- salt of the corresponding Au^{III} complex for several days. The incorporated Au^{III} complexes, comprising 0.91–8.26 wt% of the Au^{III}@MOFs (determined by inductively coupled plasma (ICP) spectroscopy, Table S2, ESI†), are likely to reside in the inner pores of the MOF host materials based on the following: (i) the MOF hosts of these Au^{III}@MOFs contain nanochannels/nanocages with window sizes larger than the sizes of the incorporated Au^{III}



complexes (Table S1, ESI[†]); (ii) complex Au₃, which is larger than the window sizes of the nano-cages in MOF2, did not give Au₃@MOF2 (under similar conditions the MOF2 in the reaction mixture remained colourless, and ICP analysis revealed no Au₃ incorporation into MOF2), suggesting that adsorption of the Au^{III} complex outside the pores (or in gaps/cracks between crystal grains) of the MOF is negligible; (iii) the PXRD data of the Au^{III}@MOFs were nearly identical to those of the corresponding MOF matrices (Fig. S5, ESI[†]); (iv) examination of the cross section of a crystal of Au₂@MOF2 (1.43 wt% Au₂) by optical microscopy under UV light irradiation at 365 nm showed the characteristic yellowish green emission of Au₂ (ref. 7), congruent with incorporation of Au₂ inside the crystal (Fig. S8, ESI[†]); (v) further analysis of the cross section of Au₂@MOF2 by scanning electron microscopy (SEM) and electron dispersive X-ray spectroscopy (EDX) revealed that Au₂ was disorderly dispersed at the cross section (Fig. S9, ESI[†]) and that no F was detected in Au₂@MOF2 by EDX, even when the content of incorporated Au₂ increased from 1.43 to 8.26 wt% (Fig. S10, ESI[†]); (vi) N₂ adsorption experiments (Fig. S11, ESI[†]) showed that, from free MOF2 to Au₂@MOF2 (1.43 wt% Au₂), the pore volume decreased from 0.303 to 0.276 cm³ g⁻¹, with a decrease in BET surface area from 1112 to 1011 m² g⁻¹. Increasing the Au₂ content of Au₂@MOF2 to 8.26 wt% led to a reduction in the pore volume and BET surface area to 0.183 cm³ g⁻¹ and 656 m² g⁻¹, respectively. The decreases in the pore volume and BET surface area are attributed to the encapsulation of Au₂ in the pores of MOF2. Au^{III}@MOFs display strong emission under air at room temperature. As depicted in their emission spectra (Fig. 2), two emission bands are observed. The one at ~450 nm

corresponds to the MOF host. The other is a vibronic structured emission with peak λ_{max} at 478–550, 520–610, 540–630 and 510–580 nm for Au^{III} = Au₁, Au₂, Au₃ and Au₄, respectively. These bands are all similar to those observed in the emission spectra of the corresponding Au^{III} complexes in degassed solution.^{7,8} The measured emission lifetimes (τ₀), quantum yields (φ_{em}), and estimated radiative (k_r) and non-radiative (k_{nr}) decay rate constants of Au^{III}@MOFs and Au₁–Au₄ under different conditions are compiled in Table 1.

Encapsulation of Au₁–Au₄ by MOF1 or MOF2 was observed to cause significant increases in the intensity and lifetime of the emission of these Au^{III} complexes (Fig. 2 and Table 1; Fig. S6 and S7, ESI[†]). As an example, at room temperature, Au₂ is non-emissive in the solid state, and the intensity of its emission at 520 nm in degassed dichloromethane solution decreased more than 100-fold upon exposure of the solution to air.⁷ Remarkably, Au₂@MOF2 is luminescent under aerobic conditions,¹⁶ and the quantum yield decreased only ~5-fold compared with that measured under argon (Table 1). For Au₂@MOF1 and Au₄@MOF1, the decrease in emission quantum yield on changing the atmosphere from argon to air is ~10- and ~20-fold, respectively. Moreover, while Au₂ is virtually non-emissive in DMF solution in air, Au₂@MOF2 (1.43 wt% Au₂), Au₂@MOF2 (8.26 wt% Au₂), and Au₂@MOF1 are luminescent in air, with emission lifetimes of 14.5, 15.8, and 7.9 μs, respectively. Notably, the emission lifetime of Au₁@MOF2 in air is 19.4 μs, which is >60-fold longer than that of Au₁ in the solid state (0.20 μs) and in DMF solution (0.32 μs) under aerobic conditions (Table 1).¹⁷ The emission lifetime of Au₃@MOF1 in open air is 48.8 μs, the longest emission lifetime reported for luminescent Au^{III} complexes under aerobic conditions. Compared with Au^{III}@MOF2, the Au^{III}@MOF1 composites are more susceptible to luminescence quenching under aerobic conditions (Fig. 2 and Table 1), due probably to the larger window size of the pores in MOF1 than in MOF2. The intrinsic triplet excited state properties and molecular sizes of the Au^{III} complexes could also affect the luminescence quenching behaviour of Au^{III}@MOFs upon exposure to air. The elevated intensities and lifetimes of the emissions of Au^{III}@MOFs in the solid state under aerobic conditions are attributed to the following: (i) the emission lifetimes of Au^{III}@MOFs under argon are longer, and their non-radiative decay rate constants (k_{nr}) are smaller, than those of the corresponding Au^{III} complexes in degassed solution (Table 1). Electrostatic binding of the Au^{III} complexes on the surfaces of the inner pores of the MOFs would restrict the molecular motion of the Au^{III} complexes, thereby slowing down non-radiative decay and self-quenching (by diffusion) of the emissive excited state.¹⁸ The radiative decay rate constants (k_r) of Au^{III}@MOFs and Au₁–Au₄ are in the order of 10² to 10³ s⁻¹ (Table 1), typical of ³IL excited states of luminescent gold(III) complexes.^{8a} (ii) The steady state concentration of oxygen in the pores of Au^{III}@MOFs may be lower than that of the oxygen dissolved in the solvent and in free atmosphere. This may reduce oxygen quenching of the emission of Au^{III}@MOFs in air.¹⁹ Given that the activity of O₂ in the matrix would equal that in the external environment if the matrix and the ambient environment are at equilibrium, and considering the use of O₂

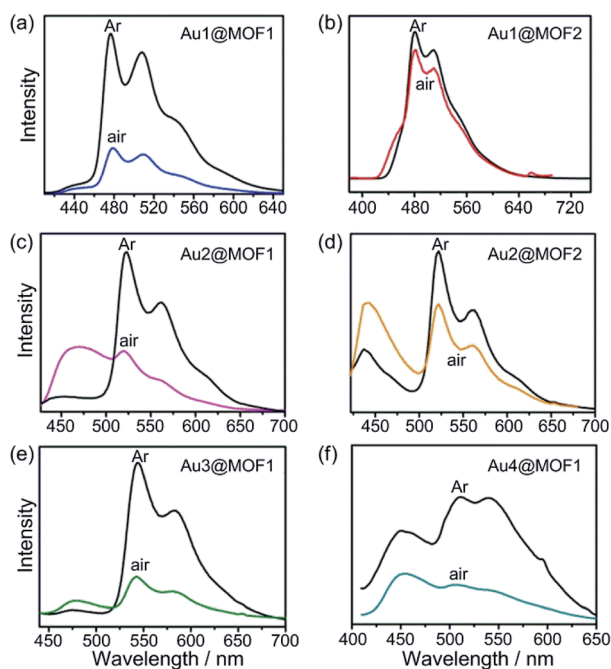


Fig. 2 Emission spectra of Au^{III}@MOFs in Ar (black lines) and air (color lines) upon excitation at λ = 365 nm at room temperature. (a) Au₁@MOF1; (b) Au₁@MOF2; (c) Au₂@MOF1; (d) Au₂@MOF2; (e) Au₃@MOF1; (f) Au₄@MOF1.



Table 1 Emission lifetimes, quantum yields, and radiative and non-radiative decay rate constants of the Au^{III} complexes and Au^{III}@MOFs

Compound	Conditions	λ_{\max} (nm)	τ_0 (μs)	ϕ_{em} (%)	k_r^a (10^3 s^{-1})	k_{nr}^b (10^5 s^{-1})
Au1 solid	Air	478, 511	0.20			
Au1 (DMF solution)	Air	478, 511	0.32	0.06	1.88	31.23
	Degassed	478, 511	0.59	0.09	1.53	16.93
Au1@MOF1	Air	478, 511	10.1	1.3	1.29	0.98
	Ar	478, 511	33.6	3.5	1.04	0.29
Au1@MOF2	Air	478, 511	19.4	2.0	1.03	0.51
	Ar	478, 511	21.2	2.7	1.27	0.46
Au2 (DMF solution)	Air	Non-emissive				
	Degassed	520, 558	21.4	0.64	0.30	0.46
Au2@MOF1	Air	520, 558	7.9	1.1	1.39	1.25
	Ar	520, 558	146.3	11.4	0.78	0.06
Au2@MOF2 ^c	Air	520, 558	14.5	1.5	1.03	0.68
	Ar	520, 558	98.6	8.1	0.82	0.09
	O ₂ ^d	520, 558	6.0			
Au2@MOF2 ^e	Air	522, 560	15.8			
	Ar	522, 560	104.2			
Au3 (DMF solution)	Air	540, 580	0.88	0.09	1.02	11.35
	Degassed	540, 580	117.7	7.0	0.60	0.08
Au3@MOF1	Air	540, 580	48.8	11.9	2.44	0.18
	Ar	540, 580	130.7	34.8	2.66	0.05
Au4 (DMF solution)	Air	510, 540	0.52	0.12	2.31	19.21
	Degassed	510, 540	8.0	0.95	1.19	1.24
Au4@MOF1	Air	510, 540	0.97	0.20	2.06	10.29
	Ar	510, 540	21.7	3.9	1.80	0.44

^a Radiative decay rate constant estimated by $k_r = \phi/\tau$. ^b Non-radiative decay rate constant estimated by $k_{\text{nr}} = (1 - \phi)/\tau$. ^c Au2 content: 1.43 wt%.

^d Au2@MOF2 immersed in acetonitrile saturated with O₂. ^e Au2 content: 8.26 wt%.

as a terminal oxidant in photo-catalytic reactions within the matrix (see the section below on the photo-catalytic properties of the Au^{III}@MOF composites) as well as the partial luminescence quenching under aerobic conditions, O₂ diffusion in the matrix of Au^{III}@MOFs is likely to occur at a finite rate, with the mass transport of O₂ through the matrix being sufficiently slow that the above-mentioned equilibrium did not fully occur.

Light-induced electron transfer reactivity of Au2@MOF2

Au^{III}@MOFs were observed to display light-induced electron transfer reactivity. As an example, Au2@MOF2 (1.43 wt% Au2) was first immersed into a solution of methyl viologen dication (MV²⁺) in DMF at room temperature for 12 h to allow encapsulation of MV²⁺ inside the pores of the MOF composite, followed by removal of the solution, washing with DMF and drying. Upon xenon lamp ($\lambda > 370 \text{ nm}$) irradiation of a mixture of the resulting Au^{III}@MOF-MV²⁺ and a drop of Et₃N liquid in

air, the characteristic blue colour of the MV^{•+} radical appeared within a few seconds (the colour intensity increased with irradiation time), as depicted in Fig. 3. Heating the mixture (after removing the xenon lamp) to remove Et₃N was observed to revert MV^{•+} back to MV²⁺, and the photolysis was recycled three times. For comparison, irradiation of a mixture of Au2, MV²⁺ and Et₃N in MeCN under the same aerobic conditions did not give a similar net change in colour of the solution mixture. In fact, a recently reported photochemical reaction of a Au^{III} complex with MV²⁺ was conducted under degassed conditions.^{8a} We measured the N₂ adsorption of Au2@MOF2-MV²⁺ without sample pretreatment, and found that very little N₂ could be adsorbed by the sample ($\sim 3.7 \text{ cm}^3 \text{ g}^{-1}$). The corresponding BET surface area is $6.3 \text{ m}^2 \text{ g}^{-1}$ (Fig. S12, ESI[†]), which is quite small compared with the BET surface area of the pre-treated Au2@MOF2 ($1011 \text{ m}^2 \text{ g}^{-1}$). Thus, for Au2@MOF2, the observed photochemical reaction with MV²⁺ and Et₃N in air can be attributed to the occurrence of the reaction inside the pores of the MOF composite, which could be occupied by solvent and Et₃N molecules, thus decreasing the quenching of MV^{•+} by O₂.

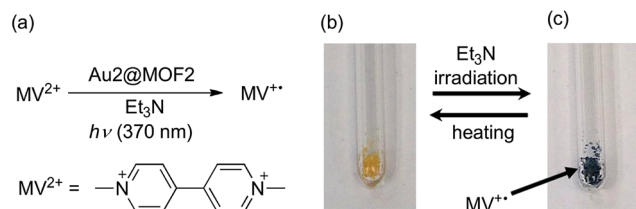


Fig. 3 (a) Photochemical reaction (light source: xenon lamp) between Au2@MOF2 and MV²⁺, and photographs of the reaction mixture before (b) and after (c) irradiation with light for 1 min.

Two-photon absorption of Au3@MOF1 in the solid state

Two-photon absorption has useful applications in chemistry and biological science;^{20,21} the two-photon-excited emission of a phosphorescent Au^{III} complex in solution has recently been reported.^{8a} However, there are few reported examples of this property in the solid state, owing to aggregation-induced quenching.²¹ In the present study, we observed that Au^{III}@MOFs display two-photon-induced phosphorescence²² in the



solid state. For Au₃@MOF1, excitation at 756 nm with a focused laser beam gave bright yellow emission (Fig. 4a) with peak maxima at 545 and 585 nm (characteristic of the emission of Au₃) and an intensity showing a quadratic dependence ($y = x^2$) on the laser power density (Fig. 4b and c).

Photo-catalytic properties of Au^{III}@MOF composites

Cyclometalated Au^{III} complexes have recently been reported to sensitize the formation of ¹O₂ upon light excitation for the oxidation of secondary amines to imines and for oxidative cyanation of tertiary amines.^{7,8a,23} Using the former as a paradigm, we examined the photo-catalytic activities of Au^{III}@MOFs. The photochemical reactions were performed in MeCN. The solution mixture was under light irradiation ($\lambda > 400$ nm) with constant bubbling of oxygen. Control experiments showed that both MOF1 and MOF2 are stable in MeCN for at least 24 h, as revealed by PXRD measurements (see Fig. S13 of ESI†). The following was observed: (i) for the photo-reaction with dibenzylamine (Fig. 5), Au₁@MOF1 gave the imine product with a turnover number of 692, exceeding the turnover number of 390 obtained with free Au₁. The photo-activity of Au₁@MOF1 showed little variation over a period of 10 h, whereas free Au₁ exhibited a significant decrease in photo-activity after 2 h, with almost no imine product obtained after 6 h (Fig. 5b). (ii) The Au₁@MOF1 catalyst could be recycled by washing with MeCN. After five cycles, the substrate conversion retained a value of ~70% (Fig. 5c). (iii) The photo-catalytic activity is attributed to Au₁ encapsulated by MOF1 rather than Au₁ leached into solution. This is confirmed by the finding that the solution phase of the Au₁@MOF1-catalysed reaction mixture remained colourless during the course of photolysis. This is in contrast to the reaction catalysed by free Au₁, in which there was a colour change attributed to the decomposition of the Au₁ catalyst after 6 h of photolysis. ICP measurements revealed nearly the same content of gold in the Au₁@MOF1 sample before and after photo-catalysis; control experiments with MOF1 (free of Au) as the catalyst gave few product turnovers (Fig. 5b). (iv) Substrate size

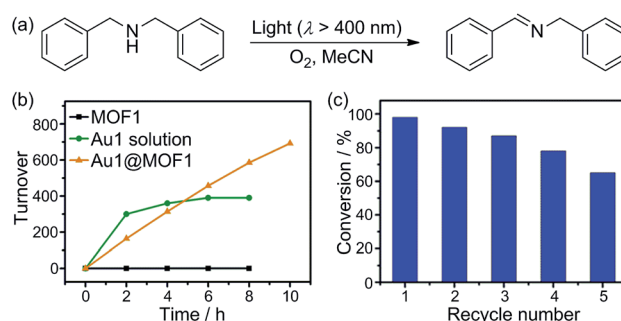


Fig. 5 (a) Photo-catalytic oxidation of dibenzylamine to imine; (b) time course of the oxidation by Au₁, MOF1, and Au₁@MOF1; (c) substrate conversion in recycling experiments using the Au₁@MOF1 catalyst.

selectivity^{1b,2d} was observed for the Au₁@MOF1-catalysed competitive photochemical oxidation of dibenzylamine (S1) in the presence of another secondary amine of larger size (S2, Scheme 2). The photochemical reaction consumed S1 rapidly without significant consumption of S2, and the imine products P1 and P2 were formed in a yield ratio of ~11 : 1 for a reaction time of 1.5 h. Under the same conditions, catalyst Au₁ showed no selectivity in solution, with a P1/P2 yield ratio of ~1 : 1 (Scheme 2). The drastic difference between the yields of P1 and P2 in the Au₁@MOF1-catalysed reaction can be attributed to the reaction mainly occurring inside the channels of the MOF framework. The smaller substrate, S1, could enter the channels of MOF1, but the larger one, S2, could not; the formation of a small amount of P2 in the reaction mixture is attributed to the external diffusion of singlet oxygen (produced inside the MOF) and its subsequent reaction with S2. Au₂@MOF1 and Au₃@MOF1 were also found to catalyse the light-induced aerobic oxidation of dibenzylamine (Fig. S14 and S15, ESI†). The total turnover numbers of the imine product furnished by Au₂@MOF1 (0.91 wt% Au₂) and Au₃@MOF1 (2.34 wt% Au₃) were 557 and 920, respectively, and both were higher than the values of 510 and 760 obtained with catalysts Au₂ and Au₃, respectively. The Au₂@MOF2 (1.43 wt% Au₂) catalyst gave the imine product with turnover number of 610; changing the catalyst to Au₂@MOF2 (8.26 wt% Au₂) led to a ~2-fold increase in conversion rate, albeit with a decrease in product turnover

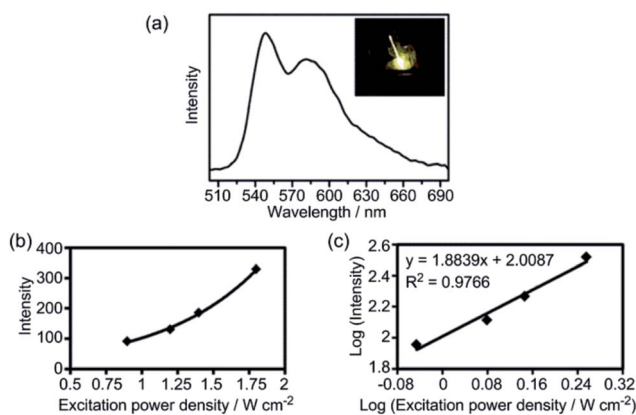
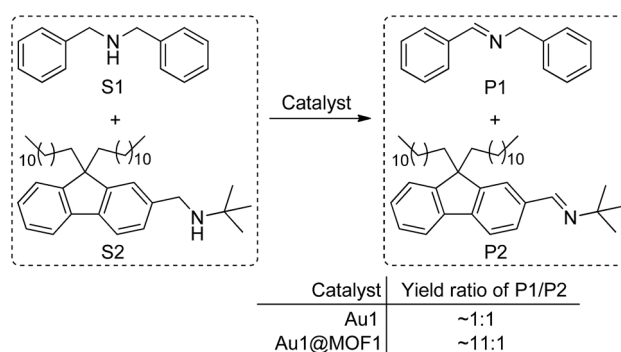


Fig. 4 (a) Emission spectrum of Au₃@MOF1 excited with 756 nm laser light (inset: photo of the emission of the solid); (b) plot of emission intensity against excitation power density (W cm^{-2}); (c) double-log plot of emission intensity against excitation power density.



Scheme 2 Competitive photo-catalytic oxidation of secondary amines.

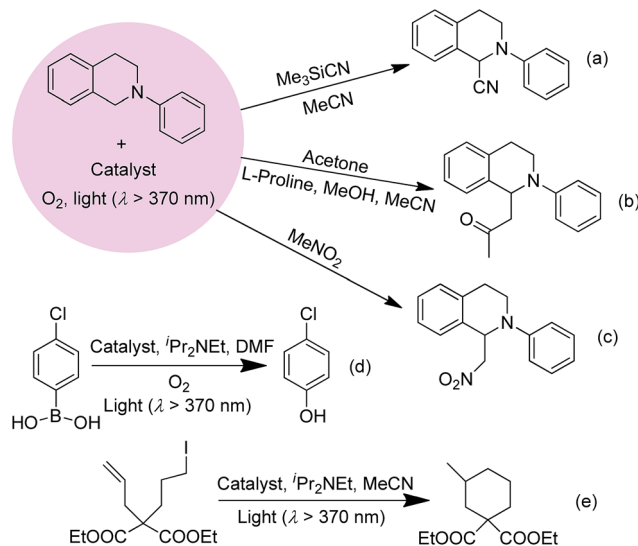


number (calculated based on % wt of Au₂ in Au₂@MOF₂; Fig. S16, ESI†).

The photochemical oxidation of dibenzylamine was also catalysed by Au₄ and Au₄@MOF₁ (Au₄: 6×10^{-7} mol in both cases). Au₄@MOF₁ was a more active catalyst with a turnover number of >143 attained within 2 h (Fig. 6a). Notably, the initial reaction rate of the Au₄@MOF₁ system was two-fold higher than that of Au₄ alone. After recycling five times, the photo-activity of the Au₄@MOF₁ catalyst maintained ~80% of its initial value (Fig. 6b). Increasing the content of Au₄ in solution from 6×10^{-7} to 6×10^{-6} mol decreased the product turnover number from 90 to 18 (Fig. 6a).

The improvement in photo-catalytic activity of Au^{III}@MOFs relative to the free Au^{III} complexes could be extended to the oxidative cyanation of a tertiary amine, a Mannich-type reaction, the aza-Henry reaction, the hydroxylation of 4-chlorophenylboronic acid, and the reductive cyclization of alkyl iodide.²⁴ These findings are depicted in Scheme 3 and in Fig. S17 and Tables S3–5 of the ESI.† For example, the steady formation of product was observed over 8 h in the Au₁@MOF₁-catalysed oxidative cyanation of *N*-phenyl-1,2,3,4-tetrahydroisoquinoline, in contrast to a significant decrease in the activity of the free Au₁ catalyst after 2.5 h of photolysis (Fig. S17, ESI†). The product yield obtained with the Au₁@MOF₁ catalyst was ~81% (turnover number of 923). For the other photochemical reactions with Au₂@MOF₂ (1.43 wt% Au₂) as catalyst, the products were obtained in 63–86% yields with turnover numbers of 137–549. These values were higher than the corresponding product yields and turnover numbers obtained with Au₂ alone (Tables S3–S5, ESI†).

To gain further insight into the photochemical oxidation reactions catalysed by Au^{III}@MOFs under aerobic conditions, which are likely to involve the formation of ¹O₂,^{7,23b} we measured the reduction potentials, $E(\text{Au}^{0/1-})$, of Au₁–Au₄ (Table S6, ESI†) and estimated the excited state reduction potentials, $E(\text{Au}^*/\text{Au}^-)$, from the electrochemical and emission data. The estimated $E(\text{Au}^*/\text{Au}^-)$ values of Au₁–Au₄ range from +0.74 to +1.38 V vs. Cp₂Fe⁺⁰ (Table S6, ESI†), indicating that they are strong oxidants in the excited state. The potentials of dibenzylamine and *N*-phenyl-1,2,3,4-tetrahydroisoquinoline, $E(\text{amine}^{+/0})$, were measured to be +0.50 V and +0.25 V vs. Cp₂Fe⁺⁰, respectively. Thus, the thermodynamic driving force for the reaction of the excited Au₁–Au₄ with dibenzylamine was estimated to be in the range of +0.24 to +0.88 V, whereas that of the excited Au₂



Scheme 3 Photo-catalytic oxidative cyanation of a tertiary amine (a), Mannich-type reaction (b), aza-Henry reaction (c), hydroxylation of 4-chlorophenylboronic acid (d), and reductive cyclization of alkyl iodide (e) catalysed by Au^{III}@MOFs.

complex with *N*-phenyl-1,2,3,4-tetrahydroisoquinoline is +0.49 V. Nanosecond time-resolved transient absorption (TA) measurements (Table S7, ESI†) revealed that, in the presence of dibenzylamine (0.1 M) in degassed DMF, the TA of Au₂ and Au₃ was quickly quenched (τ_{TA} : 1.4 μs for Au₂, 2.2 μs for Au₃) and returned to the baseline, whereas the TA of Au₁ and Au₄ evolved to give long-lived species (τ_{TA} : 145 μs for Au₁, 24 μs for Au₄), attributable to the formation of Au₁⁻ and Au₄⁻.²⁵ The TA spectrum of Au₂ in the presence of *N*-phenyl-1,2,3,4-tetrahydroisoquinoline (0.01 M) shows the formation of a long-lived species ($\tau = 65 \mu\text{s}$) assignable to Au₂⁻ (Table S8, ESI†). Thus, in the photochemical oxidation of dibenzylamine catalysed by Au₁ and Au₄, and that of *N*-phenyl-1,2,3,4-tetrahydroisoquinoline catalysed by Au₂, the excited gold(III) complexes may undergo electron transfer with the amines (to give amine radical cations) besides producing singlet oxygen for oxidation.

Conclusions

A series of luminescent Au^{III}-encapsulated MOF composites (Au^{III}@MOFs) have been synthesized. These Au^{III}@MOF solids exhibit long emission lifetimes in air, display solid state two-photon-induced phosphorescence, and function as reusable and size-selective heterogeneous photo-catalysts. The simple approach of incorporating phosphorescent metal complexes with long-lived emissive excited states into MOFs provides a means of developing new classes of heterogeneous photo-functional materials/photo-catalysts with useful applications.

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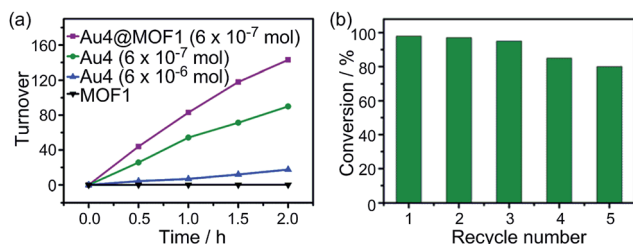


Fig. 6 (a) Product turnovers in the photo-catalytic oxidation of dibenzylamine to imine by MOF₁, Au₄@MOF₁, and Au₄ solution; (b) substrate conversion in recycling experiments using the Au₄@MOF₁ catalyst.



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Notes and references

- (a) H.-C. J. Zhou and S. Kitagawa, *Chem. Soc. Rev.*, 2014, **43**, 5415–5418; (b) H. Furukawa, K. E. Cordova, M. O'Keeffe and O. M. Yaghi, *Science*, 2013, **341**, 1230444.
- (a) J. Liu, L. Chen, H. Cui, J. Zhang, L. Zhang and C.-Y. Su, *Chem. Soc. Rev.*, 2014, **43**, 6011–6061; (b) T. Zhang and W. Lin, *Chem. Soc. Rev.*, 2014, **43**, 5982–5993; (c) Z. Hu, B. J. Deibert and J. Li, *Chem. Soc. Rev.*, 2014, **43**, 5815–5840; (d) A. Corma, H. García, F. X. Llabrés and i. Xamena, *Chem. Rev.*, 2010, **110**, 4606–4655.
- S. M. Cohen, *Chem. Rev.*, 2012, **112**, 970–1000.
- (a) J. D. Evans, C. J. Sumbly and C. J. Doonan, *Chem. Soc. Rev.*, 2014, **43**, 5933–5951; (b) J. Juan-Alcañiz, J. Gascon and F. Kapteijn, *J. Mater. Chem.*, 2012, **22**, 10102–10118.
- (a) J. Schneider, M. Matsuoka, M. Takeuchi, J. Zhang, Y. Horiuchi, M. Anpo and D. W. Bahnemann, *Chem. Rev.*, 2014, **114**, 9919–9986; (b) X. Qian, K. Fuku, Y. Kuwahara, T. Kamegawa, K. Mori and H. Yamashita, *ChemSusChem*, 2014, **7**, 1528–1536; (c) X.-d. Wang and O. S. Wolfbeis, *Chem. Soc. Rev.*, 2014, **43**, 3666–3761; (d) J. Feng and H. Zhang, *Chem. Soc. Rev.*, 2013, **42**, 387–410; (e) B. Yan, *RSC Adv.*, 2012, **2**, 9304–9324; (f) L. D. Carlos, R. A. S. Ferreira, V. de Zea Bermudez and S. J. L. Ribeiro, *Adv. Mater.*, 2009, **21**, 509–534; (g) P. Escribano, B. Julián-López, J. Planelles-Aragó, E. Cordoncillo, B. Viana and C. Sanchez, *J. Mater. Chem.*, 2008, **18**, 23–40.
- (a) Y. Yamada, A. Nomura, K. Ohkubo, T. Suenobu and S. Fukuzumi, *Chem. Commun.*, 2013, **49**, 5132–5134; (b) H. O. Lintang, K. Kinbara, K. Tanaka, T. Yamashita and T. Aida, *Angew. Chem., Int. Ed.*, 2010, **49**, 4241–4245; (c) R. Nakamura, A. Okamoto, H. Osawa, H. Irie and K. Hashimoto, *J. Am. Chem. Soc.*, 2007, **129**, 9596–9597; (d) K. Feng, R.-Y. Zhang, L.-Z. Wu, B. Tu, M.-L. Peng, L.-P. Zhang, D. Zhao and C.-H. Tung, *J. Am. Chem. Soc.*, 2006, **128**, 14685–14690; (e) K. Inumaru, T. Kasahara, M. Yasui and S. Yamanaka, *Chem. Commun.*, 2005, 2131–2133; (f) H. Yamashita, K. Yoshizawa, M. Ariyuki, S. Higashimoto, M. Che and M. Anpo, *Chem. Commun.*, 2001, 435–436.
- W.-P. To, G. S.-M. Tong, W. Lu, C. Ma, J. Liu, A. L.-F. Chow and C.-M. Che, *Angew. Chem., Int. Ed.*, 2012, **51**, 2654–2657.
- (a) W.-P. To, K. T. Chan, G. S. M. Tong, C. Ma, W.-M. Kwok, X. Guan, K.-H. Low and C.-M. Che, *Angew. Chem., Int. Ed.*, 2013, **52**, 6648–6652; (b) H.-M. Ko, K. K.-Y. Kung, J.-F. Cui and M.-K. Wong, *Chem. Commun.*, 2013, **49**, 8869–8871; (c) J. J. Yan, A. L.-F. Chow, C.-H. Leung, R. W.-Y. Sun, D.-L. Ma and C.-M. Che, *Chem. Commun.*, 2010, **46**, 3893–3895.
- D. T. Genna, A. G. Wong-Foy, A. J. Matzger and M. S. Sanford, *J. Am. Chem. Soc.*, 2013, **135**, 10586–10589.
- X. Zhang, F. X. Llabrés, i. Xamena and A. Corma, *J. Catal.*, 2009, **265**, 155–160.
- J. Muzart, *Tetrahedron*, 2009, **65**, 8313–8323.
- V. A. Blatov and M. V. Peskov, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2006, **62**, 457–466.
- A. Spek, *J. Appl. Crystallogr.*, 2003, **36**, 7–13.
- K. S. W. Sing, D. H. Everett, R. A. W. Haul, L. Moscou, R. A. Pierotti, J. Rouquérol and T. Siemieniewska, *Pure Appl. Chem.*, 1985, **57**, 603–619.
- C.-Y. Sun, C. Qin, C.-G. Wang, Z.-M. Su, S. Wang, X.-L. Wang, G.-S. Yang, K.-Z. Shao, Y.-Q. Lan and E.-B. Wang, *Adv. Mater.*, 2011, **23**, 5629–5632.
- Y.-A. Li, S.-K. Ren, Q.-K. Liu, J.-P. Ma, X. Chen, H. Zhu and Y.-B. Dong, *Inorg. Chem.*, 2012, **51**, 9629–9635.
- (a) C. L. Whittington, L. Wojtas and R. W. Larsen, *Inorg. Chem.*, 2014, **53**, 160–166; (b) G.-S. Yang, M.-N. Li, S.-L. Li, Y.-Q. Lan, W.-W. He, X.-L. Wang, J.-S. Qin and Z.-M. Su, *J. Mater. Chem.*, 2012, **22**, 17947–17953; (c) R. W. Larsen, J. Miksovská, R. L. Musselman and L. Wojtas, *J. Phys. Chem. A*, 2011, **115**, 11519–11524.
- H. Yersin, A. F. Rausch, R. Czerwieńiec, T. Hofbeck and T. Fischer, *Coord. Chem. Rev.*, 2011, **255**, 2622–2652.
- (a) P. Duan, N. Yanai and N. Kimizuka, *J. Am. Chem. Soc.*, 2013, **135**, 19056–19059; (b) J.-H. Kim, F. Deng, F. N. Castellano and J.-H. Kim, *Chem. Mater.*, 2012, **24**, 2250–2252.
- (a) G. Bort, T. Gallavardin, D. Ogden and P. I. Dalko, *Angew. Chem., Int. Ed.*, 2013, **52**, 4526–4537; (b) S. Sumalekshmy and C. J. Fahrni, *Chem. Mater.*, 2011, **23**, 483–500; (c) M. Pawlicki, H. A. Collins, R. G. Denning and H. L. Anderson, *Angew. Chem., Int. Ed.*, 2009, **48**, 3244–3266.
- (a) H.-H. Fang, Q.-D. Chen, J. Yang, H. Xia, B.-R. Gao, J. Feng, Y.-G. Ma and H.-B. Sun, *J. Phys. Chem. C*, 2010, **114**, 11958–11961; (b) J. Yu, Y. Cui, H. Xu, Y. Yang, Z. Wang, B. Chen and G. Qian, *Nat. Commun.*, 2013, **4**, 2719–2725.
- Q. Zhao, C. Huang and F. Li, *Chem. Soc. Rev.*, 2011, **40**, 2508–2524.
- (a) Q.-C. Xue, J. Xie, H.-M. Jin, Y.-X. Cheng and C.-J. Zhu, *Org. Biomol. Chem.*, 2013, **11**, 1606–1609; (b) T. N. Zehnder, O. Blacque and K. Venkatesan, *Dalton Trans.*, 2014, **43**, 11959–11972.
- (a) D. M. Schultz and T. P. Yoon, *Science*, 2014, **343**, 1239176; (b) C. K. Prier, D. A. Rankic and D. W. C. MacMillan, *Chem. Rev.*, 2013, **113**, 5322–5363.
- P.-K. Chow, G. Cheng, G. S. M. Tong, W.-P. To, W.-L. Kwong, K.-H. Low, C.-C. Kwok, C. Ma and C.-M. Che, *Angew. Chem., Int. Ed.*, 2015, **54**, 2084–2089.

