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A bioinspired heterogeneous catalyst based on the model of the manganese-dependent dioxygenase for selective oxidation using dioxygen†

Jérémy Chaignon,^{ab} Marie Gourgues,^{ac} Lhoussain Khrouz,^a Nicolás Moliner,^b Laurent Bonneviot,^a Fabienne Fache,^c Isabel Castro^{*b} and Belén Albela^{*a}

A hybrid bioinspired material with manganese(II) complexes grafted on the surface of a mesostructured porous silica is investigated. The Mn sites mimic the manganese-dependent dioxygenase (MndD), which is an enzyme that catalyses the oxidation of catechol derivatives. The metal complexes were introduced in the silica using a dinuclear complex $[Mn_2L_2(Cl)_2(\mu-Cl)_2]$ as a precursor with a clickable ligand *N,N'*-bis[(pyridin-2-yl)methyl]prop-2-yn-1-amine (L). Azide moieties covalently grafted on MCM-41 type mesoporous silica were utilised to anchor the manganese complex through Huisgen cycloaddition using $CuBr(PPh_3)_3$ as a catalyst. A second functional group – trimethylsilyl or pyridine—was grafted on the silica to bring, together with nanopore size confinement, a similar metal environment as in MndD. The mesostructure of the materials was maintained after incorporation of the Mn complex. Catalytic oxidation of 3,5-di-*t*-butyl-catechol (3,5-DTBC) into quinone occurred without the need of an additional base when the metal complex was confined in the porous solid. In comparison, the oxidation of 4-*t*-butyl-catechol (4-TBC) that always required a basic media led to a total oxidation into the *ortho*-quinone, contrary to the molecular analogue.

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Introduction

Many industrial processes involve catalytic reactions, since they allow working under milder conditions with high selectivity and short reaction time, decreasing, therefore, the cost of the final product.¹ However, these enhanced processes are not a prerogative for industry, for they existed in nature long before chemists could understand them. A class of natural metalloproteins called metalloenzymes is able to carry out extremely specific reactions under physiological conditions (between 10 and 40 °C for most of them).² In particular, several metalloenzymes called oxygenases perform the oxidation of organic compounds by activating molecular dioxygen. Various metal ions can be found in these oxygenases, mainly copper, iron and manganese.^{3–9} Although the reaction with dioxygen is thermodynamically favourable, the electronic configuration of the O₂ molecule creates a strong kinetic barrier that requires

activation, which is useful to prevent any spontaneous reaction of O₂ with organic molecules.

Structural and mechanistic research on metalloenzymes in the past decades has allowed the development of numerous bioinspired systems that can be used for catalysis. Homogeneous biomimetic complexes are often not optimal for reactions involving dioxygen activation. Indeed, the activation of oxygen can partially oxidise the catalyst. Grafting on a solid support can stabilise the catalyst and confer confinement as in the enzymatic pocket.

Amongst oxygenases, a recurrent metal coordination triad is observed, composed by two histidines (His) and one carboxylic acid, coming either from glutamate (Glu) or aspartate (Asp) moieties. This widespread structural motif is called the 2-His-1-carboxylate facial triad, and is found in many metalloproteins.⁶ This is the case of catechol dioxygenase, which catalyses the cleavage of catechol derivatives. Two kinds of catechol dioxygenases can be found, which differ on the nature of the metal ion involved at the active site: Fe^{III} allows an intradiol cleavage while Fe^{II} or Mn^{II} performs the extradiol one.^{6–9} In particular, the manganese-dependent dioxygenase (MndD) is a system that has attracted much attention since its catalytic mechanism is still under discussion.^{7,10–12} Therefore, the study of bioinspired systems that mimic the structural and/or catalytic function of MndD could help to understand its reaction mechanism.

^aLaboratoire de Chimie UMR CNRS 5182, Ecole Normale Supérieure de Lyon, University of Lyon, 46, Allée d'Italie, 69364 Lyon, Cedex-07, France. E-mail: belen.albela@ens-lyon.fr

^bInstituto de Ciencia Molecular, Universitat de València, C/Catedrático José Beltrán 2, 46980 Paterna, València, Spain. E-mail: isabel.castro@uv.es

^cInstitut de Chimie et Biochimie Moléculaires et Supramoléculaires (ICBMS) UMR CNRS 5246, University of Lyon, Bd. du 11 Novembre 1918, 69622 Villeurbanne, Cedex, France

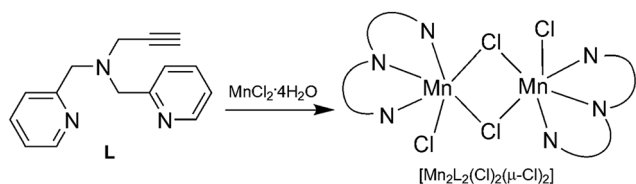
† Electronic supplementary information (ESI) available: TGA profiles of materials, magnetic measurements and N₂ sorption isotherms. See DOI: 10.1039/c7ra00514h



Molecular models that contain a manganese centre are however much less frequent than those containing iron. Only few examples in the literature report manganese(II) complexes as mimics of the catechol dioxygenase. The first reported model to our knowledge was synthesised by Sakurai *et al.* two years after the discovery of MndD.¹³ Their report describes the use of penicillamine with manganese(II) in the oxidation of catechol to the corresponding muconic acid derivatives resulting from a true extradiol dioxygenase activity. However, very low conversion rates were achieved (<2%). Ramadan and El-Mehasseb subsequently investigated the oxidative properties of manganese(II) with tridentate pyridyl hydrazone ligands.¹⁴ They reported the transformation of catechol into *o*-benzoquinone in presence of catalyst and O₂, which would correspond to an oxidase activity (oxidation without incorporation of oxygen atom in the substrate). Similar results were obtained a few years later with tetradentate ligands containing imidazole, phenolate and/or pyridine moieties.¹⁵ At the same time, the manganese(II) complex of another tetradentate ligand, *N,N'*-bis(2-pyridylmethyl)-1,2-ethanediamine (bispicen), managed to oxidise the 4-*tert*-butylcatechol into its quinone equivalent mixed with extradiol intermediates.¹⁶ Reaction conditions turned out to have an important effect on the resulting products.

We have developed a bioinspired approach for design of the metal supported catalysts based on the model of metalloproteins.^{17–21} This approach was recently applied using the manganese dependent dioxygenase (MndD) as a model, the *N,N'*-bis[(pyridin-2-yl)methyl]prop-2-yn-1-amine (L) as a clickable ligand for grafting and [Mn₂L₂(Cl)₂(μ-Cl)₂] as a precursor metal complex (Scheme 1).¹⁸ This ligand has the particular advantage to mimic the facial triad present in the active site of the MndD and allows an efficient grafting in the silica support.^{22–25} Indeed, the alkyne side function allows grafting of the ligand in an azido-functionalised support using “click chemistry”.^{26–28} In addition, the two pyridine and the tertiary amine functions present in the ligand can coordinate to the Mn(II) ion and thus mimic the metal environment of the MndD.

In the present work we are going further in providing a comprehensive investigation of the [Mn₂L₂(Cl)₂(μ-Cl)₂] grafting, knowing that this dimer is in equilibrium with the monomer in solution.¹⁸ We will also compare two different local environments of the metal complex in the solid to test the influence of a pyridine side function meant here to mimic the histidine function neighbouring the metal active site in MndD. Both homogeneous and heterogeneous systems are tested for the first time on the oxidation of catechol derivatives.



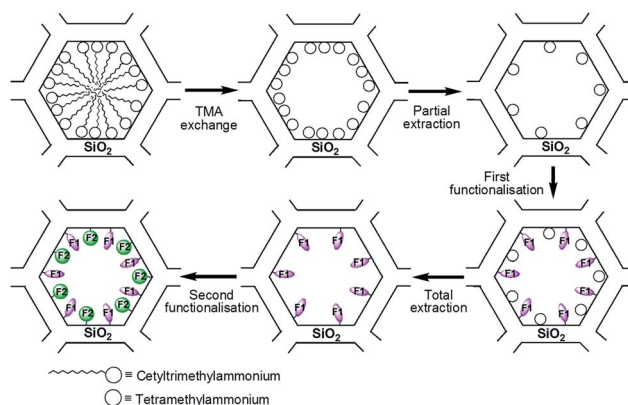
Scheme 1 Synthesis of complex [Mn₂L₂(Cl)₂(μ-Cl)₂].

Results and discussion

Incorporation of the Mn^{II} complex in the mesoporous silica: synthesis strategy to isolate the metal sites with control of their close environment

The alkyne function present in the ligand L (Scheme 1) was used to covalently graft the manganese complex into the pores of a mesoporous silica of the LUS type, which presents a 2D hexagonal structure just like the classic MCM-41 silica.²⁹ The LUS silica was synthesised using an ultra-fast microwave assisted hydrothermal synthesis that affords a high quality mesostructured porous silica.³⁰ Due to the yield and selectivity advantages of the copper(I)-catalysed Huisgen azide–alkyne cycloaddition (also known as “click chemistry”), this reaction was chosen for the grafting process.^{26–28} We had the choice to design our ligands either with an azide or an alkyne arm, the latter being preferred for two reasons. First, the incorporation of an azide function in the ligand could have been more problematic for the synthesis of the complex than the incorporation of an alkyne arm because of the coordinating potential of the azide moiety. Secondly, the synthesis of the azide tethering silane was easier than the alkyne tether, the azidopropyltriethoxysilane precursor being then synthesised starting from chloropropyltriethoxysilane upon nucleophilic substitution of the chlorine atom by an azide moiety.³¹

The molecular stencil patterning (MSP) technique was used to incorporate the complexes into the solid (Scheme 2).^{32–34} This strategy takes advantage of the cationic nature of the surfactant (cetyltrimethylammonium, CTA⁺) present in the mesoporous silica. The general procedure is a four step strategy: (i) the template is partially removed with hydrochloric acid; due to electrostatic repulsion, the remaining surfactant rearranges thus making some silica surface available, (ii) a first organosilane (F1) is grafted onto this freed surface, (iii) all of the remaining surfactant is removed, freeing silica surface once again, and (iv) the second function (F2) is grafted, providing a double functionalised material. In the present work, however, a slightly different protocol was applied where CTA⁺ was exchanged with tetramethylammonium (TMA⁺) prior to the first partial extraction (Scheme 2).³⁴ This modification allowed



Scheme 2 Molecular stencil patterning strategy. TMA = tetramethylammonium.



a better diffusion of the F1 grafting function since there was more available pore space. If the grafted functions are modelled as spheres arranged in 2D hexagonal packing on the pore surface, isolating a function F2 will result in having it surrounded by 6 functions F1. Every F1 will thus be adjacent to 3 functions F2, which corresponds to a ratio F1/F2 of 2.³²

The nomenclature used for the synthesised solids is the following. LUS stands for the as-made support containing CTA⁺ molecules inside the pores, while the material exchanged with TMA⁺ is denoted as LUS-TMA. Partial extraction is represented by an -Ex suffix where *x* is the percentage of extracted TMA⁺. For instance, a material where 15% of TMA⁺ was extracted is named LUS-TMA-E15. Totally extracted materials are simply noted LUS-E. Likewise, when the remaining TMA⁺ is removed from a LUS-TMA-Ex material, the resulting one is denoted as LUS-Ex. Grafting reactions are symbolised by prefixes. For instance, silica functionalised with azide (N₃) and trimethylsilyl (TMS) groups will be named N₃-TMS@LUS. In the case of double functionalisation, the last function incorporated is placed at the beginning of the name: a material functionalised with F1 and then with F2 will thus be called F2-F1@LUS.

In the present work two materials containing a manganese(II) complex were studied using [Mn₂L₂(Cl)₂(μ-Cl)₂]¹⁸ as metal precursor: a new solid {Mn}-Py@LUS-E15 (S1), where pyridine (Py) functions are in the vicinity of the metal complexes, and the already reported {Mn}-TMS@LUS-E (S2),¹⁸ where the manganese complexes are surrounded by hydrophobic trimethylsilyl (TMS) functions (Scheme 3). In this nomenclature, the {Mn} function is the product of the reaction between [Mn₂L₂(Cl)₂(μ-Cl)₂] and the azide (N₃) tether present in both host materials. The TMS function was chosen to mimic some hydrophobic residues present in the enzymatic pocket of the metalloprotein, whereas Py function was selected to mimic the His200 residue present in the second coordination sphere of the Mn active site, which is known to help in the activation of molecular oxygen during the catalytic cycle of MnDd.^{7,35,36}

Optimisation of the “click chemistry” grafting of the Mn complex

Even though click chemistry is known to be efficient, one must be careful about the conditions implied in those reactions. Indeed, the copper-catalysed Huisgen cycloaddition has proven to be sensitive to coordinating molecules when the classical copper catalyst is used. In principle, copper(II) sulphate or acetate is directly used and reduced to copper(I) *in situ*, most of the time by treatment with sodium ascorbate. For instance,



Scheme 3 Host materials for the grafting of the Mn(II) complex: N₃-TMS@LUS-E and N₃-Py@LUS-E15.

reaction with L and benzyl azide has been reported and performed under these conditions.³⁷ Instead of sodium ascorbate, methanol can also be used as reductant.³⁸ Some preliminary tests between the ligand L and azidopropyltrimethoxysilane were performed in order to optimise the conditions for the click reaction on the solid support. Under these conditions, coordination of copper to the ligand L was observed in all cases, even when copper(I) bromide was used. A tris(triphenylphosphine) copper(I) bromide complex was required as catalyst to avoid this problem. This change of catalyst was inspired by organic syntheses of triethoxysilanes using the microwave-assisted Huisgen copper-catalysed reaction reported in the literature.³⁹ Given that triethoxysilanes are particularly sensitive to water and to many chemical functions such as alcohols or carboxylic acids, this kind of synthesis was quite a challenge. However, using microwave irradiation and CuBr(PPh₃)₃ as catalyst, the desired organosilane was obtained in a yield typical for click chemistry (above 90%).

Then, several tests were performed to click the manganese(II) complex [Mn₂L₂(Cl)₂(μ-Cl)₂] in the azide-functionalised silica N₃@LUS-E. After an overnight reaction with CuBr(PPh₃)₃, the IR spectrum of the resulting material still exhibited a band around 2100 cm⁻¹, which is characteristic of the presence of azide moieties. After a one-week reaction, no absorption occurred in this region, proving that all the azide moieties had reacted. This long reaction time can be explained by the porous nature of the material. All the reactive sites at the silica surface are not as accessible as in solution, and diffusion inside the porous material becomes a non-negligible parameter. Removal of the copper catalyst was performed by washing with methanol. This technique was efficient enough to remove the catalyst, as confirmed by EDX microscopy, EPR and elemental analysis: no trace of copper could be found beyond the detectable threshold. Therefore, this technique was applied to incorporate the manganese complex into the host materials N₃-Py@LUS-E15 and N₃-TMS@LUS-E, affording solids {Mn}-Py@LUS-E15 (S1), {Mn}-TMS@LUS-E (S2), respectively.

Characterisation of the hybrid materials

The porous structure of the host materials was checked using powder X-ray diffraction (XRD) during the different steps of the synthesis. The original 2D hexagonal porous structure was maintained during the functionalisation of the silica and the incorporation of the manganese complex (Fig. 1).

Thermogravimetric analyses (TGA) indicate that after exchange of CTA⁺ surfactant molecules present in the as-made silica by TMA⁺, almost 2/3 of the internal surface of the silica was covered by TMA⁺ cations, which corresponds to 2.1 mmol of TMA⁺ per gram of SiO₂. Starting from this solid, two intermediate materials were synthesised: TMS@LUS-TMA and Py@LUS-TMA-E15. In order to obtain a F1/F2 molar ratio of 2, TMS functionalisation was performed without any partial extraction. Indeed, TMS is grafted to the support with only one covalent bond and the ammonia generated by the aperture of the azasilane used for the functionalisation displaces a small part of the TMA⁺. However, in the case of pyridine, which possesses



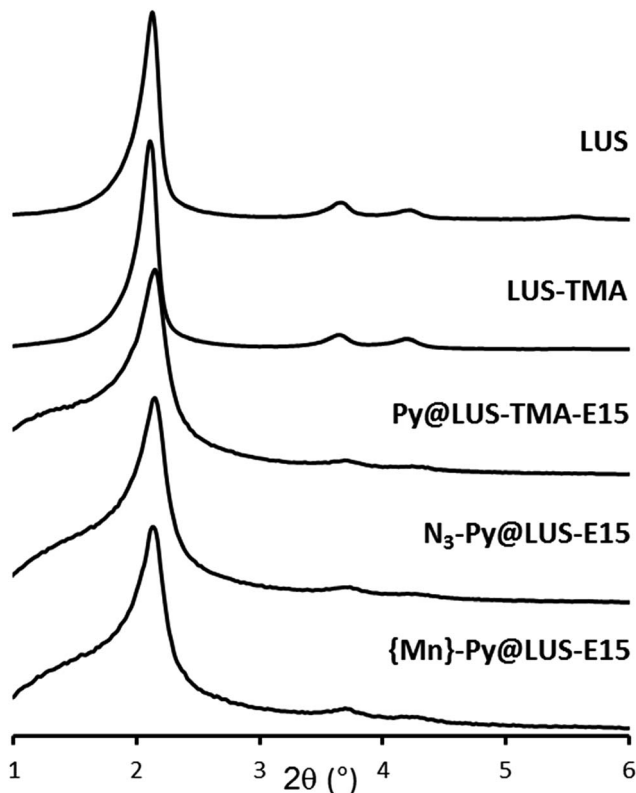


Fig. 1 Powder XRD patterns of the support LUS-E and the materials LUS-TMA, Py@LUS-TMA-E15, N₃-Py@LUS-E15 and {Mn}-Py@LUS-E15 (S1).

a triethoxysilane tether function, 15% of TMA⁺ was extracted before the functionalisation reaction in order to favour the grafting of the function. In other words, TMS functionalisation was performed on LUS-TMA material while pyridine functionalisation was carried out with LUS-TMA-E15 material. The F1/F2 molar ratio was calculated from TGA, elemental analyses, and quantitative solid ²⁹Si-NMR (Table 1 and Fig. S1†). Py/N₃ and TMS/N₃ molar ratios of 1.9 and 1.8 can be deduced, and these are close to the desired value of 2.0 required to insure site isolation. Note that the TMS amount cannot be precisely deduced neither from TGA – the mass loss is very low and occurs in a broad temperature range – nor from elemental analyses due to the presence of carbon atoms in TMA⁺.

Table 1 Data deduced from N₂ sorption isotherms, TGA, elemental analyses and quantitative solid ²⁹Si NMR^a

Material	V _p ^b (cm ³ g ⁻¹)	M _n (wt%)	F1/Si ^c	F2/Si ^d	Cl/Mn ^e
LUS-E	0.86	—	—	—	—
N ₃ -TMS@LUS-E	0.26	—	0.046	0.025	—
N ₃ -Py@LUS-E15	0.28	—	0.064	0.034	—
S1 ¹⁸	0.31	1.7	—	—	0.7
S2	0.22	2.4	—	—	0.5

^a {Mn}-Py@LUS-E15 = S1; {Mn}-TMS@LUS-E = S2. ^b Porous volume deduced from sorption N₂ isotherms using *t*-plot, ±0.01 cm³ g⁻¹. ^c Molar ratio, F1 = TMS or Py. ^d Molar ratio, F2 = N₃. ^e Molar ratio, deduced from chemical analyses.

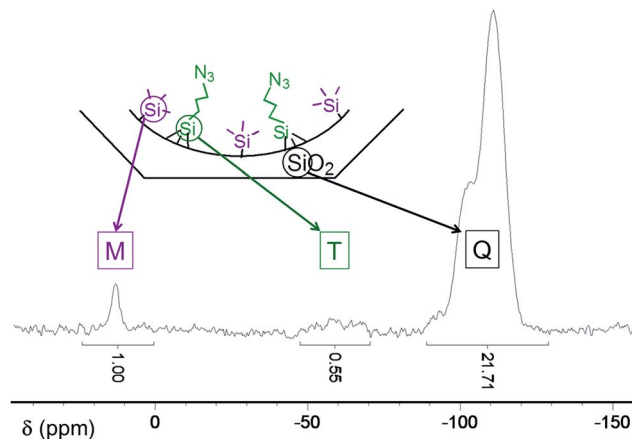


Fig. 2 Solid ²⁹Si HPDEC NMR spectrum of N₃-TMS@LUS-E.

Integration of M and Q masses from solid ²⁹Si-NMR could however give a precise value of the quantity of grafted TMS functions per gram of SiO₂ (Fig. 2).

The two hybrid materials {Mn}-Py@LUS-E and {Mn}-TMS@LUS-E present a manganese weight loading of about 2%, 1.7 and 2.4%, respectively (Table 1). The chloride counter-anions present in the initial manganese complex are only partially retained in the final materials. The Cl/Mn molar ratio in the grafted materials (Cl/Mn = 0.7 and 0.5 for {Mn}-Py@LUS-E and {Mn}-TMS@LUS-E, respectively) is approximately 25% of the original chloride anions present in the dinuclear precursor (Cl/Mn = 2). This diminution tends to prove that the chloride bridges are partially lost during the grafting process questioning about the presence of mononuclear species.

Nitrogen sorption isotherms were performed to measure the remaining porous volume of the final materials (Table 1 and Fig. 3). The pore volume, measured using the *t*-plot method, was reduced from 0.86 cm³ g⁻¹ in LUS-E down to 0.2–0.3 cm³ g⁻¹ in the final materials, indicating a partial filling of

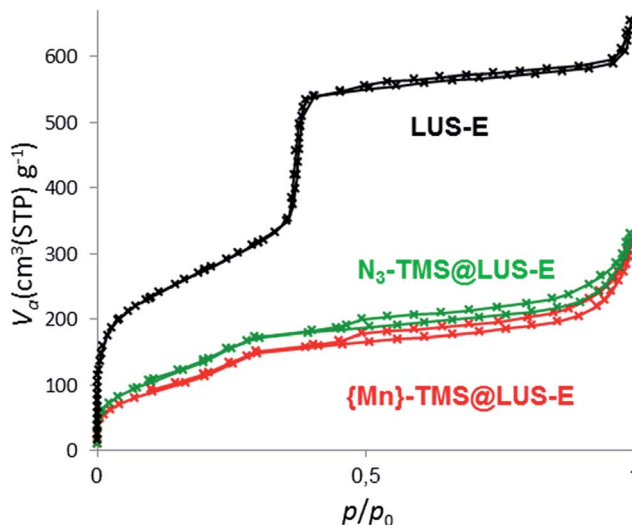


Fig. 3 Nitrogen sorption isotherms at 77 K of LUS-E, N₃-TMS@LUS-E, {Mn}-TMS@LUS-E (S2).



the pores after the incorporation of the complex. However, there was still enough pore volume to allow the substrate diffusion and its access to the manganese(II) centres for catalytic applications.

Nature of the metal complexes inside the solid

In order to determine the relative amount of mono- and dinuclear species, magnetic and EPR measurements were performed for both solids {Mn}-Py@LUS-E15 (**S1**), {Mn}-TMS@LUS-E (**S2**).

The magnetic properties of **S1** and **S2** are shown in Fig. 4 and **S2**[†] in the form of the $\chi_M T$ vs. T plot (χ_M being the molar magnetic susceptibility per Mn^{II} ion and T the absolute temperature). They are qualitatively similar to those of the dinuclear precursor [Mn₂L₂(Cl)₂(μ -Cl)₂].¹⁸ $\chi_M T$ remains constant from room temperature down to ca. 100 K and it abruptly decreases upon further cooling, suggesting the occurrence of weak antiferromagnetic interactions between the Mn^{II} ions within the grafted metal complex. The least-squares fits of the experimental data through the appropriate expression derived from the spin Hamiltonian for an antiferromagnetically coupled dinuclear manganese(II) complex ($H = -JS_1S_2$),¹⁸ assuming the presence of a variable amount of mononuclear manganese(II) species (α), gave $J = -1.5$ cm⁻¹ and $g = 2.0$ for both **S1** and **S2** with $\alpha = 0.25$ (**S1**) and 0.38 (**S2**) (solid lines in Fig. 4 and **S2**[†]). The calculated α values indicate the presence of a relatively larger amount of mononuclear Mn^{II} species resulting from the loss of the two bridging chlorides in **S2**, in agreement with the slightly smaller value of the Cl/Mn ratio, as discussed above.

The calculated $-J$ value in **S1** and **S2** materials are twenty five-fold larger than that found for the dinuclear precursor ($J = -0.059$ cm⁻¹).¹⁸ The stronger antiferromagnetic coupling for

the grafted metal complex could be explained by the bending of the Mn-Cl-Mn angle (θ) within the double-bridged Mn₂(μ -Cl)₂ diamond core with respect to that found in the dinuclear precursor ($\theta = 97.5^\circ$).¹⁸ This situation agrees with the general trend found in related double-bridged di- μ -chlorodimanganese(II) complexes reported in the literature, which exhibit weak ferro- ($J = 0.66$ – 1.1 cm⁻¹ with $\theta = 93.5$ – 96.4°)^{40–42} to moderate antiferromagnetic couplings ($-J = 4.8$ – 5.2 cm⁻¹ with $\theta = 90.2$ – 90.7°).^{43,44} Overall, these results suggest that the molecular structure of the dinuclear precursor was likely modified upon grafting within the pore walls of the mesoporous silica.

EPR spectra of **S1** and **S2** solids confirmed the presence of both monomeric and dimeric Mn species. In both cases the spectra were simulated assuming the monomeric ratio α deduced from the magnetic measurements (Fig. 4 and **S2**[†]). The shape of the simulated signals matched well that of the experimental signals assuming one type of monomer and two types of dimers, a dimer 1 with $D = 0.0333$ cm⁻¹ and a dimer 2 with $D = 0.0733$ cm⁻¹. The calculated parameters for the different species were: for **S1**, 62.5% of dimer 1 with $g = [1.990 \ 1.990 \ 2.120]$, 12.5% of dimer 2 with $g = [2.000 \ 2.000 \ 2.150]$, 25% of monomer with $g = [2.055 \ 2.018 \ 1.940]$, and for **S2**, 28% of dimer 1 with $g = [1.990 \ 2.000 \ 2.130]$, 34% of dimer 2 with $g = [1.990 \ 2.100 \ 2.200]$, 38% of monomer with $g = [2.080 \ 2.010 \ 1.940]$ (see ESI[†] for additional parameters). The presence of two different dimers revealed by EPR is not surprising given the various possibilities of interaction with the functionalised material or the triazole cycle generated during the Huisgen cycloaddition. The Cl/Mn molar ratio deduced from the elemental analysis for **S1** and **S2** (0.7 and 0.5, respectively) led us to suppose that the solid likely contains a dimer 1 with two chloride bridges, a dimer 2 with one chloride bridge and a monomer without any chloride ion in the first coordination sphere (Scheme 4 and Fig. **S2**[†]). Considering this hypothesis, the calculated Cl/Mn molar ratios would be 0.68 and 0.45 for **S1** and **S2**, respectively, which are within the margin of error of elemental analysis. In addition, the difference of g values for dimer 2 in both solids could be explained either by the presence of a second bridge provided by the different supports or by a different geometry around the metal ion induced by the solid matrix.

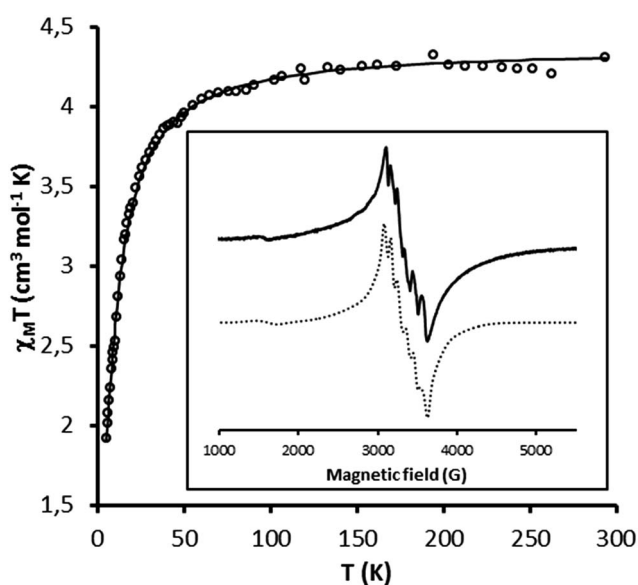
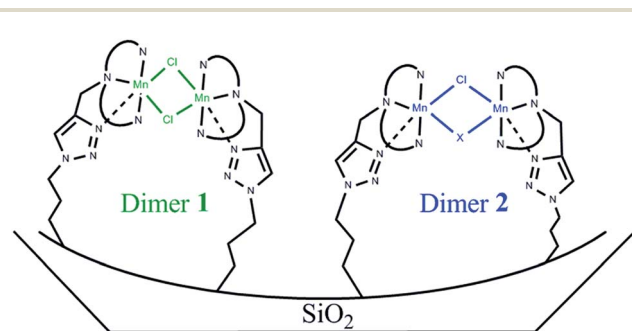


Fig. 4 Temperature dependence of $\chi_M T$ for **S1**. The solid line represents the best fit discuss in the text. Insert: solid-state EPR spectra of **S1** at 120 K and its simulation (dotted line).



Scheme 4 Representation of dimers 1 and 2. X represents a possible second bridge for dimer 2. Dashed bonds are possible bonds created after the click reaction.



Catalytic properties

We first tested the oxidation of 3,5-di-*t*-butylcatechol (3,5-DTBC), known to be easier to study if compared to catechol, as it leads to more stable intermediates and thus to fewer by-products, due to the presence of 3- and 5-*t*-butyl blocked positions.⁴⁵ In order to appreciate the effect of the manganese complex anchoring into the mesoporous silica, we systematically compared the heterogeneous catalyst and its homogeneous analogue. We mainly identified one product, named A, which is the quinone equivalent of 3,5-DTBC (Scheme 5) and confirmed the oxidase activity of our catalysts (Table 2).¹⁴

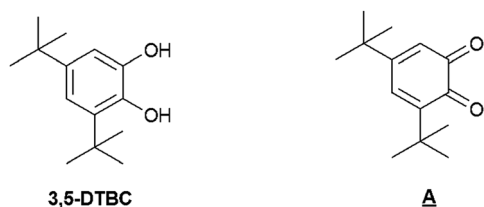
Without any catalyst and in the presence of a base (Table 2, entry 1), a mixture of unidentified products and the starting material was obtained. In the absence of base, no reaction occurred, without or with MnCl₂ as catalyst (entries 0 and 2, respectively). In homogeneous conditions, with a catalytic amount of base (10%), complete conversion was observed and product A was isolated exclusively whatever the catalyst used, *i.e.* MnCl₂ or [Mn₂L₂(Cl)₂(μ-Cl)₂] (entries 3 and 4). Working under reflux did not change the reactivity (entry 5).

Heterogeneous catalysts led also to product A even in the absence of an additional base (entries 6 and 7 for {Mn}-Py@LUS-E15 and {Mn}-TMS@LUS-E, respectively). It is not surprising since it was already observed that pyridine has a negligible effect on this reaction contrary to triethylamine.⁴⁶ Nevertheless, our heterogeneous catalysts turned out to be as efficient as the homogeneous counterpart in the case of 3,5-DTBC, with the advantage of being easier to separate and potentially recyclable.

Table 2 Oxidation of the 3,5-DTBC with Mn-based catalysts^a

Entry	Catalyst	Base (%)	Conversion (%)	Selectivity ^b (%)
0	—	—	0	0
1	—	200	~20 ^c	0
2	MnCl ₂	0	100	100
3	MnCl ₂	10	100	100
4	[Mn]	10	100	100
5	[Mn] ^d	10	100	100
6	S1	0	100	100
7	S2	0	100	100

^a [Mn₂L₂(Cl)₂(μ-Cl)₂] = [Mn]; {Mn}-Py@LUS-E15 = S1; {Mn}-TMS@LUS-E = S2. Standard reaction conditions: 50 mL MeOH, 0.25 mmol 3,5-DTBC, 10% catalyst, O₂, room temperature, 5 h; base = 2,2,6,6-tetramethylpiperidine. ^b Selectivity for product A. ^c Presence of unknown products. ^d T = 65 °C.



Scheme 5 3,5-DTBC and its quinone equivalent A.

We then turned our attention to a more complex substrate, the 4-*t*-butyl catechol (4-TBC), for which we mainly identified two products, named B¹⁶ and C (Scheme 6).^{47,48} Without catalyst and in the presence of base, a total conversion of 4-TBC to a mixture of unidentified products was obtained (Table 3, entry 1), whereas without base but in presence of [Mn₂L₂(Cl)₂(μ-Cl)₂] no reaction occurred (entry 2). With both catalyst and base, a mixture of products B and C were isolated with a 100% total conversion (entry 3).

Changing the nature of oxidant (air instead of oxygen) led to a slight modification of the B/C ratio in favour of B product (entries 3 and 4), whereas heating under reflux had almost no significant effect on this ratio (entries 4 and 5). Reducing the reaction time from 5 h (entry 4) to 3 h (entry 7) diminished the amount of B in favour of C. We thus assume that there is equilibrium between B and C depending on the reaction conditions. Finally, using a catalytic amount of base considerably diminished the activity of the catalyst (entries 4 and 6). Concerning the nature of the catalyst, with either Mn(NO₃)₂ (entry 8) or MnCl₂ (entry 9) no reaction occurred, which proves that the ligand L is required for the activity of the catalyst (entry 10). In the absence of base, the heterogeneous catalysts had almost no activity in the case of 4-TBC (entries 11 and 12), whereas it showed oxidase activity with 3,5-DTBC. In the former case, an addition of stoichiometric amount of base was required to go to completion, leading selectively to only one compound, product B (entry 13). Indeed 3,5-DTBC is easier to oxidise since the intermediates are more stable.^{49,50} It is noteworthy that in all cases no notable changes were observed in the N₂ sorption isotherms, before and after catalytic tests, leading us to believe that the material do not undergo major modifications throughout the whole catalytic process (Fig. S3†). Nonetheless it remains to study the recyclability of these bio-inspired catalysts.

Experimental

Chemicals

All chemicals were used without further purification. Solvents were dried using molecular sieves if needed. Propargylamine, 2-picoly chloride hydrochloride and 3-chloropropyltriethoxysilane were purchased from Sigma-Aldrich, tetrabutylammonium iodide and 2,2,6,6-tetramethylpiperidine were purchased from Acros Organics, tetramethylammonium bromide, hexamethyldisilazane were purchased from Alfa Aesar, 2-(4-pyridylethyl)triethoxysilane was provided by ABCR and bromotris(triphenylphosphine)copper bromide was purchased from Santa Cruz Biotechnology.

Synthesis of the ligand and the azide tether

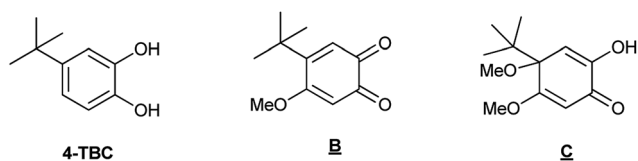
Synthesis of the ligand L.⁵¹ A mixture of propargylamine (1.50 g, 27.3 mmol) and potassium carbonate (22.8 g, 163.8 mmol) was stirred in 140 mL of acetonitrile for 5 min. Then 2-picoly chloride hydrochloride (9.84 g, 60.0 mmol) dissolved in 140 mL of acetonitrile was added. The resulting solution was stirred under reflux for 5 days. The solution was filtered and the filtrate was evaporated under reduced pressure. The resulting brown oil was



Table 3 Oxidation of 4-TBC with Mn-based catalysts^a

Entry	Catalyst	Base (%)	Oxidant	Conversion (%)	Selectivity (%)	
					B	C
1	—	200	O ₂	100 ^b	0	0
2	[Mn]	0	O ₂	0	0	0
3	[Mn]	200	O ₂	100	60	40
4	[Mn]	200	Air	100	74	26
5	[Mn]	200	Air ^c	100	66	34
6	[Mn]	10	Air	48	100	0
7	[Mn]	200	Air ^d	100	38	62
8	Mn(NO ₃) ₂	10	O ₂	0	0	0
9	MnCl ₂	10	O ₂	0	0	0
10	[Mn]	10	O ₂	70	100	0
11	S1	0	O ₂	<5	<5	0
12	S2	0	O ₂	<5	<5	0
13	S1	200	O ₂	100	100	0

^a [Mn₂L₂(Cl)₂(μ-Cl)₂] = [Mn]; {Mn}-Py@LUS-E15 = S1; {Mn}-TMS@LUS-E = S2. Standard reaction conditions: 50 mL MeOH, 0.25 mmol 4-TBC, 10% catalyst, O₂, room temperature, 5 h; base = 2,2,6,6-tetramethylpiperidine. ^b Mixture of unidentified products. ^c T = 65 °C. ^d 3 h.



Scheme 6 4-TBC and the two main oxidation products B and C.

dissolved in 100 mL of distilled water and extracted with dichloromethane (3 times). The organic phases were gathered and dried over sodium sulphate. The solvent was finally evaporated to afford L in a 94% yield. ¹H NMR (CDCl₃, 300 MHz): δ (ppm) 8.55 (ddd, *J* = 4.80, 1.80, 0.90 Hz, 2H), 7.64 (td, *J* = 7.50, 1.80 Hz, 2H), 7.50 (d, *J* = 7.80 Hz, 2H), 7.15 (ddd, *J* = 7.50, 4.80, 1.20 Hz, 2H), 3.91 (s, 4H), 3.41 (d, *J* = 2.40 Hz, 2H), 2.29 (t, *J* = 2.40 Hz, 1H). ¹³C NMR (300 MHz, CDCl₃): δ (ppm) 158.76, 149.26, 136.47, 123.14, 122.10, 78.32, 73.62, 59.44, 42.54. IR (KBr): 3294 (w), 3062 (w), 3011 (w), 2921 (w), 2839 (w), 2095 (m, C≡C) cm⁻¹.

Synthesis of 3-azidopropyltriethoxysilane.³¹ Sodium azide (27 g, 0.41 mol) was added portionwise to a solution of 3-chloropropyltriethoxysilane (20 g, 83 mmol) and tetrabutylammonium iodide (153 mg, 0.42 mmol) in 220 mL of butanone. The mixture was then stirred at reflux for 5 d under nitrogen. The solution was then filtered over celite and the solvent was removed under reduced pressure. The resulting oil was redissolved in 400 mL CH₂Cl₂ and quickly washed twice with 20 mL of distilled water. The organic phase was dried over Na₂SO₄ and the solvent was removed under reduced pressure to afford the 3-azidopropyltriethoxysilane (84 mg, 0.34 mmol) as a pale yellow oil in a 80% yield. ¹³C NMR (200 MHz, CDCl₃): δ (ppm) 58.16, 53.54, 22.39, 17.97, 7.33.

Synthesis of the grafted complexes

[Mn₂L₂(Cl)₂(μ-Cl)₂] was used to synthesise the hybrid materials.¹⁸ The complex was incorporated in a LUS mesoporous

silica where azide functions were previously grafted using the so-called molecular stencil patterning technique.³⁴ A Berghof speedwave oven was used for the synthesis of the mesoporous silica LUS using the procedure described elsewhere.^{18,30} It provides a microwave irradiation at 2450 MHz with a power of 1450 W. The experimental conditions used were: *T* = 180 °C, ramp = 1 min, plateau = 9 min (total heating time = 10 min).³⁰

Synthesis of LUS-TMA.³⁴ Tetramethylammonium bromide (1.7 g, 11 mmol) was dissolved in 150 mL of EtOH 70% and stirred for 10 min at 40 °C. The LUS mesoporous silica (3 g) was added and the resulting suspension was stirred for 45 min at 40 °C. The solid was filtered on Büchner and washed twice with 50 mL of EtOH 70% and twice with 50 mL of acetone. This procedure was repeated three times to ensure a total exchange. The solid LUS-TMA obtained was finally dried in at 80 °C overnight.

Synthesis of {Mn}-Py@LUS-E15 (S1)

LUS-TMA-E15. Hydrochloric acid (0.25 mL of 1 M solution) was added to a suspension of LUS-TMA (1 g) in 200 mL of technical EtOH and stirred at RT for 1 h. The solid was then filtered and washed twice with 100 mL of technical EtOH, before being dried overnight at 80 °C.

Py@LUS-TMA-E15. LUS-TMA-E15 (500 mg) was pretreated as at 130 °C under argon for 2 h, then at 130 °C under vacuum for 2 h. After cooling under argon, a solution of 2-(4-pyridylethyl)triethoxysilane (459 mg, 1.7 mmol) dissolved in 10 mL of dry toluene was added. The suspension was stirred for 1 h, 20 mL of toluene were added, and the mixture was stirred at 80 °C during 18 h. After filtration and washing twice with 20 mL of toluene, 20 mL of technical EtOH and 20 mL of acetone, the solid was finally dried overnight at 80 °C.

Py@LUS-E15. Hydrochloric acid (2 mL of 1 M solution) was added to a suspension of Py@LUS-TMA-E15 (1 g) in 200 mL of technical EtOH and stirred at RT for 1 h. The solid was then



filtered and washed twice with 100 mL of technical EtOH, before being dried overnight at 80 °C.

N₃-Py@LUS-E15. Py@LUS-E15 (500 mg) was pretreated following the same procedure used in the synthesis of Py@LUS-TMA-E15. After cooling under argon, a solution of 3-azidopropyltriethoxysilane (371 mg, 1.5 mmol) dissolved in 10 mL of dry toluene was added. The suspension was stirred for 1 h, 20 mL of toluene were added, and the mixture was stirred at 80 °C during 18 h. After filtration and washing twice with 20 mL of toluene, 20 mL of technical EtOH and 20 mL of acetone, the solid was finally dried overnight at 80 °C. Elemental analysis: C: 9.78%, H: 1.99%, N: 2.14%.

{Mn}-Py@LUS-E15 (S1). N₃-Py@LUS-E15 (300 mg) was stirred in a mixture MeOH/MeCN (60 mL/20 mL). After 10 min, complex [Mn₂L₂(Cl)₂(μ-Cl)₂] (176 mg, 0.5 mmol) was added and the suspension was stirred during 1 h, before addition of CuBr(PPh₃)₃ (93 mg, 0.10 mmol). Then the mixture was stirred at 60 °C for 6 days before filtration. The pale brown solid was washed with 200 mL of MeOH and dried overnight at 80 °C. Elemental analysis: C: 9.86%, H: 1.84%, N: 1.68%, Mn: 1.66%, Cl: 0.74%.

Synthesis of {Mn}-TMS@LUS-E (S2)¹⁸

TMS@LUS-TMA.³⁴ Before any reaction, the solid was pretreated as follows: the LUS-TMA solid (1.5 g) was stirred at 130 °C under argon for 2 h, then at 130 °C under vacuum for 2 h. The solid was let to cool to room temperature under argon, before addition of hexamethyldisilazane (2.42 mg, 15.0 mmol) in 20 mL of dry toluene. The suspension was further stirred at room temperature for 1 h, and 60 mL of toluene were then added. The suspension was stirred under argon at 80 °C for 18 h, before being filtered and washed twice with 20 mL of toluene, 20 mL of technical EtOH and 20 mL of acetone. The solid was dried overnight at 80 °C to afford the TMS@LUS-TMA silica.

TMS@LUS-E. Hydrochloric acid (2 mL of 1 M solution) was added to a suspension of TMS@LUS-TMA (1 g) in 200 mL of technical EtOH and stirred at RT for 1 h. The solid was then filtered and washed twice with 100 mL of technical EtOH, before being dried overnight at 80 °C.

N₃-TMS@LUS-E and {Mn}-TMS@LUS-E (S2). The functionalisation of TMS@LUS-E with azide functions and the incorporation of the Mn(II) complex [Mn₂L₂(Cl)₂(μ-Cl)₂] in N₃-TMS@LUS-E was performed as described elsewhere.¹⁸ CuBr(PPh₃)₃ was used as catalyst and the reaction temperature and reaction time for the click reaction were 60 °C and 6 days, respectively.

General procedure for the catalytic tests

The substrate (0.25 mmol) and the base (2,2,6,6-tetramethylpiperidine) were dissolved in MeOH (50 mL) in a single-necked round-bottom flask. To this solution was added the catalyst and the gas was passed through the reaction mixture for 5 h with magnetic stirring.

Homogeneous catalysis. The resulting solution was concentrated to dryness and the resulting mixture was extracted

with Et₂O after addition of 2 N HCl (until pH 4–5). The organic layer was dried over MgSO₄. After evaporation of Et₂O, the crude mixture was analysed by ¹H NMR (300 MHz). Silica gel flash-chromatography allowed separating the reaction products (80/20: petroleum ether/AcOEt).

Heterogeneous catalysis. Filtration of the reaction mixture allowed recovering of the catalyst for further analyses. The filtrate was concentrated to dryness, analysed by ¹H NMR (300 MHz) and the reaction products were recovered by silica gel flash-chromatography (80/20: petroleum ether/AcOEt).

Physico-chemical characterisation

Infrared spectra were recorded from KBr pellets using a Mattson 3000 IRTF spectrometer. Nitrogen sorption isotherms at 77 K were determined with a volume device Belsorp Max on solids that were dried at 80 °C under vacuum overnight. Low angle X-ray powder diffraction (XRD) experiments were carried out using a Bruker (Siemens) D5005 diffractometer using Cu K monochromatic radiation. Liquid NMR spectra were recorded on a Bruker AC 200 spectrometer. Solid NMR spectra were measured on an Advance III Bruker 500 MHz Wide Bore spectrometer, with a CP/MAS DVT (double resonance, variable temperature) probe 4 mm. The delay time between pulses (pulse width 5 μs) in HPDEC experiments was set to 500 s. Spinning speed was set to about 5 kHz and the number of scans was 5000. Frequency ¹³C = 125.76 MHz, ²⁹Si = 99.36 MHz. EPR spectra were recorded on a Bruker Elexsys e500 X-band (9.4 GHz) spectrometer with a 4102ST cavity (modulation: 2 G, power: 4 mW). The simulated spectra were calculated using the Easyspin toolbox from Matlab. Variable-temperature magnetic susceptibility measurements were made using a Quantum Design MPMS-XL-5 SQUID susceptometer, using an applied field of 1000 G.

Conclusions

The *N,N'*-bis[(pyridin-2-yl)methyl]prop-2-yn-1-amine (L) ligand is a versatile molecule allowing both coordination to a metal ion and grafting to a solid support using click chemistry. The functionalisation method herein reported is not restricted to this particular complex and can be applied to any metal complex provided that the ligand contains a free alkyne group. The site isolation is ensured by the molecular stencil patterning strategy that allows homogeneous grafting of two different functions in a precise ratio. The azide anchor function is surrounded by pyridyl or TMS functions. Once again, the method is not limited to these particular groups and can be extended to other isolating functions. We have demonstrated that the use of CuBr(PPh₃)₃ as catalyst for the Huisgen cycloaddition avoids copper complexation to L allowing therefore a better yield for the click reaction. The Mn hybrid materials here presented have been tested in the catalytic oxidation of two catechol derivatives (3,5-DTBC and 4-TBC). Their activity has been compared with that of the molecular analogue. With both catechol derivatives tested, oxidation could be achieved using molecular dioxygen as oxidant. This reaction can be compared to the natural oxidase



enzymes. Nevertheless, the heterogeneous catalysts were more advantageous as no base was necessary in the case of 3,5-DTBC and only one product was obtained in the case of 4-TBC. Therefore, once grafted in a confined and controlled environment, the metal complex does exhibit a different and more advantageous behaviour than the free complex. The broad range of isolating functions available offers numerous options to control the activity of any inserted complex, which is not possible for complexes in solution. Controlling the environment at a larger range than the sole ligand might allow tuning the catalyst activity, in a similar fashion to the natural enzymes.

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