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Nanoparticle-assisted dynamic nuclear polarization in liquids

Beatrice Bernadette Mascitti,^{id a} Giordano Zanoni,^{id a} Alex van der Ham,^{id b} Luming Yang,^{id b} Lorenzo Franco,^{id a} Fabrizio Mancin,^{id a} Federico Rastrelli^{id *a} and Tomas Orlando^{id *c}

Gold nanoparticles (AuNPs) protected with self-assembled organic monolayers represent versatile nanostructures with applications in catalysis, molecular sensing, and recognition. The ordered arrangement of ligands within the monolayer promotes specific orientations and proximities of functional groups, facilitating weak yet selective interactions with small molecules in solution. In the context of nuclear magnetic resonance (NMR), AuNPs can act as magnetization reservoirs, allowing selective transfer to interacting molecules and making them promising chemosensors. While “NMR chemosensing” typically exploits nucleus–nucleus magnetization transfer mechanisms, we explored the potential of electron–nucleus interactions to allow selectively enhanced Overhauser effect dynamic nuclear polarization (OE-DNP). To this aim, we designed AuNPs simultaneously functionalized with two different thiols: a radical-bearing thiol as a polarizing agent for OE-DNP and an organic thiol tailored with different head groups to impart selective recognition. To evaluate both the polarization transfer efficiency and the selectivity, we performed OE-DNP experiments on a mixture of chloroform and methanol. Our results show sizeable enhancements on chloroform, possibly limited only by the mobility of the radical. On the other hand, the affinity for methanol does not correspond to greater enhancements on this target molecule. Although the system, as designed, does not provide efficient DNP on methanol as expected, our results confirm that AuNPs may be used as polarizing agents at magnetic fields up to 9.4 T. We foresee that future modifications of the nanoparticles’ monolayer will allow the combination of spin polarization transfer efficiency with molecular recognition, offering a novel strategy to boost hyperpolarization techniques and expanding their application towards NMR detection of specific analytes.

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Introduction

Gold nanoparticles (AuNPs) protected by monolayers of organic molecules are self-organized and multivalent entities able to perform different functions.^{1–3} Their applications in a variety of contexts have been extensively demonstrated and reviewed, spanning fields as diverse as bioimaging, nanomedicine, sensing and catalysis.^{4–10}

The large interest in this class of molecular systems arises because the self-assembly of a monolayer of ligands onto the surface of a metal core offers a distinct pathway for achieving ordered and rather complex molecular architectures. Within

the monolayer, in fact, the ligands are aligned and kept in close proximity, which restricts the possible configurations. These arrangements enable cooperative interactions that can be exploited to execute defined tasks such as, for example, molecular recognition.

The combination of NMR and AuNPs endowed with pre-defined binding affinity for specific classes of molecules lies at the basis of “NMR chemosensing”.¹¹ In fact, the reduced rotational diffusion constants of NPs provide a magnetization reservoir within the monolayer that can be selectively transferred *via* nuclear Overhauser effect (NOE) only to molecules interacting with the monolayer. As such, when employed in the analysis of complex mixtures, NMR chemosensing typically provides an edited NMR spectrum showing only the signals of specific target molecules.

In recent years, NMR chemosensing has established remarkable achievements in terms of selectivity, and the advent of “high-power” water saturation transfer difference (STD) experiments allowed to partly overcome the inherent sensitivity

^a Department of Chemical Sciences, University of Padova, Padova, Italy.

E-mail: federico.rastrelli@unipd.it

^b ESR Spectroscopy Group, Max Planck Institute for Multidisciplinary Sciences, Göttingen, Germany

^c National High Magnetic Field Laboratory and Florida State University, Tallahassee, FL, USA. E-mail: torlando@magnet.fsu.edu

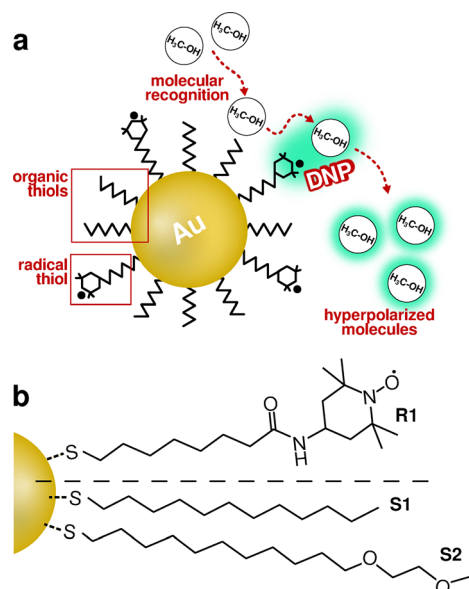



Fig. 1 (a) Cartoon of the DNP-chemosensing experiment. (b) Gold nanoparticles functionalized with the deprotected radical thiol **R1** and coated with organic thiols **S1** or **S2**.

limitations.^{9,12} In this context, it seems plausible to investigate whether the nucleus–nucleus dipolar interactions at the basis of the chemosensing phenomenon can be paralleled by electron–nucleus interactions, which would open the pathway to hyperpolarized chemosensing experiments *via* electron–nuclear polarization transfer (Fig. 1a).

Among the various existing hyperpolarization techniques¹³ – some of which successfully used for selective chemosensing,^{14,15} dynamic nuclear polarization based on the Overhauser effect (OE-DNP) in the liquid state seems the most appropriate for our applications. In OE-DNP, high electron spin polarization is transferred to nearby nuclear spins *via* electron–nuclear cross-relaxation enabled by microwave irradiation, thereby enhancing their NMR signal.^{16–19} The efficiency of OE-DNP at high magnetic fields depends on two factors:²⁰ (i) the presence of a large (a few MHz) electron–nuclear coupling,^{20–23} often favoured by non-covalent interactions, such as hydrogen bonds,^{24,25} halogen bonds,^{26–28} or van der Waals interactions.²⁹ (ii) The presence of fast dynamics (enabled by molecular diffusion, collisions, structural reorientations, *etc.*)^{22,30–33} that modulate the electron–nuclear couplings on the picoseconds or sub-picoseconds timescale.^{34,35} In the context of ¹³C-OE-DNP, those conditions are satisfied in systems constituted by an organic radical such as nitroxide as a polarizing agent (PA) and small molecules as OE-DNP targets. Although small molecules in low viscosity media perform significantly better than systems characterized by slower dynamics,^{36–39} it has been recently proven that OE-DNP can be effective in viscous matrices like lipid bilayers,^{40–42} and even on solid surfaces.⁴³ Similarly, bulky polarizing agents such as BDPA radical^{29,44,45} or substrates functionalized with nitroxide^{33,46–48} could be used as PAs in DNP, and their efficacy

relies not only on OE-DNP but also on other mechanisms that work in liquids under specific circumstances.^{40,41,49–51} In the same context, other groups have explored the idea of grafting a PA on solid beads to polarize *via* OE-DNP a flowing liquid,^{46,47,52,53} or to use OE-DNP as a tool to explore the complex surface dynamics.^{54–56}

With the aim of achieving DNP-enhanced chemosensing (Fig. 1a) and considering the requirements for efficient OE-DNP, we designed and synthesized gold nanoparticles (AuNPs) <2 nm in diameter functionalized with two surface ligands (Fig. 1b). The first ligand is a thiol (**R1**) featuring a nitroxide radical (one of the most efficient PAs for ¹³C-DNP of small molecules in liquids) as a head group, and a C₇ aliphatic spacer (Fig. 1b), which provides enough molecular mobility to favour fast dynamics and limit the downsides arising from the large size of the AuNPs. The second ligand is a thiol that either acts as an inert “dilutant” (**S1**) or can form non-covalent interactions (namely hydrogen bonds) with suitable target molecules (**S2**). On the OE-DNP side, we selected two target molecules: chloroform, which gives high DNP enhancements at high magnetic field, and methanol, which is prone to interact with the oxomethylene groups of thiol **S2**. Their performance was compared to that of the free nitroxide radical TEMPOL and TEMPONE. Our results show that AuNPs can be used as PAs for OE-DNP at high magnetic fields (9.4 T). Although the molecular recognition properties do not offer a significant advantage to boost OE-DNP performance, AuNPs remain promising systems for OE-DNP, and their versatility in terms of chemical functionalization make these systems an optimal platform to achieve selective NMR hyperpolarization in liquids.

Materials and methods

Synthesis and functionalization of AuNPs

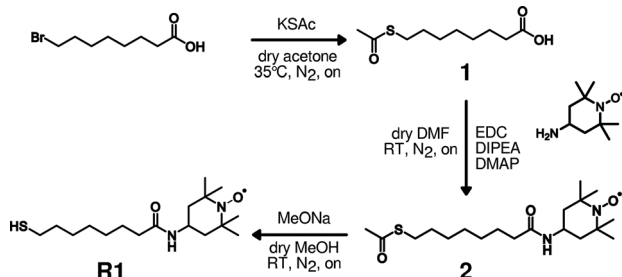
The AuNPs in this work were prepared following a two-step procedure involving the spontaneous formation of gold cores in solution through the reduction of gold salts (see SI). In general, a weak ligand (dioctylamine) is initially employed and then exchanged with a thiol: this procedure is known to improve the size control of the final nanoparticles.

The ligand TM-thiol **R1** carrying the TEMPO headgroup was synthesized with the procedure outlined in Scheme 1, involving the condensation of 8-(acetylthio)octanoic acid with 4-amino TEMPO. The insertion of TM-thiol **R1** in AuNPs covered with homogeneous monolayers of **S1** was realized by chemical exchange (see SI for details) to obtain nanoparticles **R1S1-AuNP**. Nanoparticles **R1S2-AuNP** were obtained from direct synthesis, meaning that both **R1** and **S2** were added in the desired proportions to form a mixed monolayer. Similar AuNPs functionalized with radicals have been prepared with a different protocol by Chechik *et al.*, 2004.⁵⁷

Sample preparation for magnetic resonance spectroscopy

Solvents (CHCl₃, ¹³C-methanol, methanol) and radicals (TEMPOL: 4-Hydroxy-TEMPO and TEMPONE: 4-Oxo-TEMPO) were





Scheme 1 Synthetic scheme of TM-thiol **2** which is then grafted on the particle as radical thiol **R1**.

used as received from the supplier, while $^{13}\text{C-CHCl}_3$ was stabilized with 1% of ethanol and K_2CO_3 was added to neutralize acid species in solution. Samples for magnetic resonance measurements were prepared by dissolving the PA (nitroxide or **R1SX-AuNPs**) in the organic solvent used as target for DNP experiments. To reduce the content of dissolved oxygen, all samples were degassed with at least three freeze-pump-thaw cycles and then transferred in quartz tubes that were either flame-sealed or closed with air-tight caps.

Electron paramagnetic resonance (EPR)

CW-EPR was used to probe whether the radical thiols are bound to the AuNPs, quantify their actual concentration, and evaluate their mobility compared to the unbound nitroxide radical.

Electron paramagnetic resonance experiments were performed on a Bruker Elexys E580 operating at ~ 9.6 GHz and 0.34 T. Modulation was 100 kHz and modulation amplitude was kept in the range 0.02–0.05 mT, which is below the radical linewidth. The EPR spectra were fitted with the Matlab package Easyspin.^{58,59} Other characterizations and the calibration experiments in Fig. 2c were performed on a Bruker ECS106 spectrometer operating at 9.6 GHz (see SI for details).

Dynamic nuclear polarization: experiments

DNP measurements at 1.2 T were performed on a Bruker Elexys E580 operating at 34 GHz with 40 W microwave (MW) power interfaced with a Bruker Avance III console for NMR detection.²⁶ The double resonance (electron and ^{13}C) probe consists of a cavity for MW irradiation, and a custom-made copper coil for radiofrequency irradiation and detection.²⁶ In the case of ^{13}C , the excitation and detection are at ~ 13 MHz. Samples were ~ 7 μL in volume sealed in a 1.6 mm outer diameter quartz tube. The lack of active cooling in combination with the small sample volume prevented the use of neat $^{13}\text{C-MeOH}$ due to its high MW absorption.

The DNP instrument at 9.4 T consists of a 263 GHz gyrotron, an NMR magnet, and a $^1\text{H-}^{13}\text{C}$ double-resonance probe with MW irradiation specifically designed for experiments in liquids.²⁷ Samples were prepared by degassing *via* freeze-pump-thaw cycles ~ 60 – 70 μL of sample and transferring it in a 5 mm outer-diameter quartz tube. A 4 mm outer diameter tube was used as an insert to decrease the sample thickness along the MW propagation direction, which is orthogonal to the tube axis.²⁷ The temperature of the

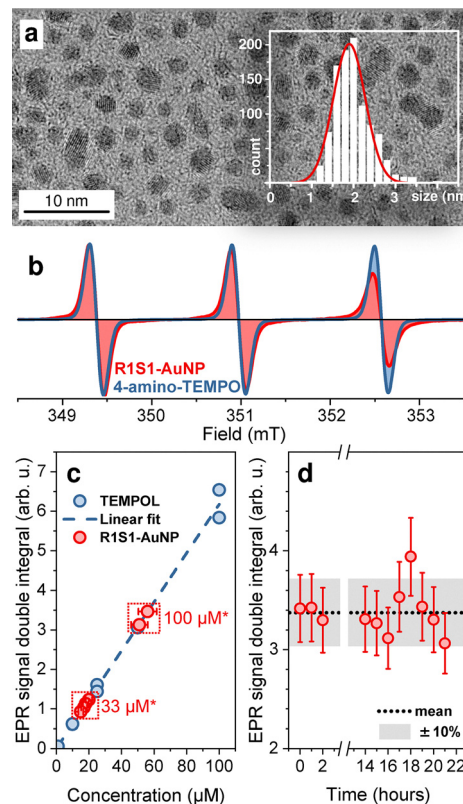


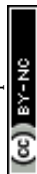
Fig. 2 (a) TEM image of the sample **R1S1-AuNP**, and histogram of the gold core size distribution. The curve was fitted with a Gaussian distribution with mean and standard deviation $d_{\text{Au}} = 1.90 \pm 0.38$ nm. (b) CW-EPR spectra of 4-Amino-TEMPO and **R1S1-AuNP** in solutions recorded at 9 GHz. The intensities have been normalized. The area under the peaks (highlighted in color) is calculated by taking the double-integral of the spectrum. (c) Calibration curve for the radical concentration obtained for 4-amino-TEMPO (blue) and calibrated data for sample **R1S1-AuNP**. The numbers marked with * indicate the nominal concentration of **R1** estimated from structural data. (d) Double-integral value of CW-EPR spectra recorded over ~ 24 hours on a sample of **R1S1-AuNP** dispersed in CHCl_3 . The grey bands indicate $\pm 10\%$ around the mean value (dotted line).

sample under MW irradiation was controlled with a flow of cold nitrogen gas. Despite the sample being exposed to high-power MW irradiation, we did not notice any degradation of the AuNPs, which could arise if the temperature of the sample is not optimally controlled.

Dynamic nuclear polarization: theory

The spin polarization transfer between electron and nuclei is driven by the modulation of the electron–nuclear hyperfine couplings, both scalar and dipolar, which give counteractive contributions. Molecular mobility, and in particular the relative motions between the PA and the target molecule, determines the ratio of the two: molecular diffusion (translation and rotation) modulates the dipolar coupling, while molecular collisions modulate the scalar one. The NMR enhancements ε is given by the Overhauser equation:

$$\varepsilon = 1 - s \cdot f \cdot \xi \cdot \frac{|\gamma_e|}{\gamma_{^{13}\text{C}}} \quad (1)$$



where: $|\gamma_e|$ and γ_{13C} are the electronic and nuclear gyromagnetic ratio, respectively; s is the saturation factor, namely the degree of saturation of the electron resonance; f (leakage factor) is the contribution of the paramagnetic relaxation over the total nuclear relaxation; and ξ is the coupling factor, which defines the efficiency of the spin polarization transfer process.

When experiments are performed under similar conditions (microwave irradiation, sample formulation) the DNP efficiency, and therefore the enhancement, depends mainly on how the molecular motions modulate the electron nuclear hyperfine coupling, *i.e.* on ξ . The coupling factor ξ can be written as a function of the spectral densities $J(\omega)$ associated with given molecular motions:

$$\xi = \frac{5}{7} \cdot \left(1 - \frac{7k_D J_D(\omega_n, \tau_D) + 7k_{rot} J_{rot}(\omega_n, \tau_{rot})}{R_1} \right) + \frac{12}{5} \cdot \left(\frac{0.5J_{sc}(\omega_e, \tau_{sc})}{R_1} \right), \quad (2)$$

where ω_n and ω_e are the nuclear and electron Larmor frequencies, respectively; τ_i are the correlation times; k_i are constants. The subscript “D” indicates the terms describing the translational diffusion, “rot” the rotational diffusion, and “sc” the scalar coupling. The term R_1 is the sum of all contributions:

$$R_1 = \frac{1}{2} J_{sc}(\omega_e, \tau_{sc}) + \sum_{i=D,rot} k_i [7J_i(\omega_e, \tau_i) + 3J_i(\omega_n, \tau_i)]. \quad (3)$$

Results and discussion

Structural characterization

The size and size-distribution of **R1SX-AuNPs** was estimated by transmission electron microscopy (TEM) images taken after the synthesis (Fig. 2a and SI). The average size of the gold core is $d_{Au} \lesssim 1.9$ nm, while the size distribution follows a narrow Gaussian profile (Fig. 2a and Table 1). UV-vis spectra validate these results and confirm that the average size of the gold core is $d_{Au} < 2$ nm (SI).

With the morphological data obtained from TEM and thermogravimetric analysis (TGA), and considering the amount of radical thiol **S1** added in the synthesis step (SI), we estimated the average composition of **R1SX-AuNPs**. However, to precisely estimate the amount of radical thiol **R1** grafted on the Au-NPs surface, we utilized CW-EPR (Fig. 2b). Specifically, we quantified the number of unpaired electron spins in the sample, which is proportional to the double integral of the EPR spectrum (Fig. 2b). A calibration curve was obtained by dissolving 4-

Amino-TEMPO in dichloromethane with precise concentrations ranging from 1 μ M to 100 μ M (Fig. 2b). Using this same procedure and ensuring consistency in solvent, sample tubes, and sample volume, we prepared samples of **R1SX-AuNP** with known concentration of radical thiol **R1** derived from the structural characterization obtained with TEM and TGA (Fig. 2a and Fig. S15). From this analysis we estimated that the amount of radical thiol **R1** over the total amount of thiols grafted onto the gold surface is $\sim 5\%$ for **R1S1-AuNP**, and $\sim 7.5\%$ for **R1S2-AuNP** (Table 1). Notably, AuNPs without **R1** functionalization do not show any EPR signal.

To verify the stability of **R1SX-AuNPs** in organic solvents, we conducted spin counting experiments with CW-EPR as a function of time. As representative sample, we selected **R1S1-AuNP** dissolved in $CHCl_3$ and monitored it over approximately 24 hours. The results (Fig. 2c) show that the double-integral values fluctuate by $\sim 10\%$ around the mean value, possibly due to slightly different experimental conditions (temperature and sample positioning in the resonator), but there is no indication of radical quenching. Importantly, the shape of the CW-EPR spectrum does not change over time, indicating that the radical thiol **R1** does not detach from the AuNPs.

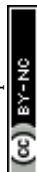
Radical mobility *via* EPR

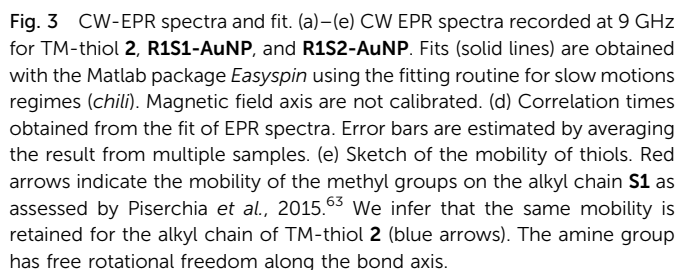
The CW-EPR spectrum of a nitroxide radical dispersed in low-viscosity organic solvents (*e.g.* $CHCl_3$, CH_2Cl_2) shows three lines separated by the isotropic hyperfine coupling A_{iso} , which are equally intense due to the complete averaging of the g-tensor due to the fast tumbling of the molecule (Fig. 2b). This is also the case for radical thiol 2 (Fig. 3a). When the mobility of the radical is constrained, as in the case of **R1** bound to AuNPs, the g-tensor is only partially averaged. This causes the lines to broaden, and the high-field line to appear less intense (Fig. 3b and c). Considering the difference in EPR spectra between radicals bound to the AuNPs and those that remain unbound, the spectra of samples **R1S1-AuNP** and **R1S2-AuNP** (Fig. 3b and c) indicate that all radicals are bound to the AuNPs, as there is no contribution from fast-tumbling nitroxides that might have remained unbound during the synthesis. This is further corroborated by the fact that the spectra can be fitted with a single component.

The rotational correlation time τ_c was estimated by fitting the CW EPR spectra. Nitroxide radicals TEMPOL and TEM-PONE as well as radical thiol 2 have correlation times of few picoseconds (~ 7 ps and ~ 5 ps, respectively, Table 2). When the radical thiol is grafted onto the AuNP surface (**R1**), its mobility is drastically reduced, as evidenced by the reduced intensity and broadening of the high field line in the CW-EPR

Table 1 Structural properties of AuNPs systems. The diameter of the gold core d_{Au} was obtained by TEM measurements, and the values indicate the mean of the Gaussian distribution of sizes and its standard deviation. The thiols surface density and the total number of thiols per single AuNP were obtained by TEM and TGA (Supplementary Information), while the amount of radical thiol **R1** was quantified with CW-EPR

Sample	Synthesis method	Formula	d_{Au} (nm)	Thiols surface density (nm^{-2})	Total thiols per AuNP	R1 /total thiols
R1S1-AuNP	Chemical exchange	$Au_{211}R_{17}S_{1125}$	1.90 ± 0.38	11.6	132	$5.3 \pm 0.3\%$
R1S2-AuNP	Direct synthesis	$Au_{134}R_{15}S_{267}$	1.63 ± 0.30	8.6	72	$7.5 \pm 0.7\%$



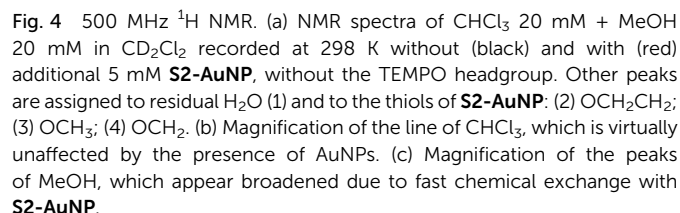


These correlation times are comparable to those reported for fullerenes (C₆₀) functionalized with nitroxide derivatives, which have also been used as PAs for OE-DNP.³³ In our case, the correlation time τ_c of the radical thiol **R1** is shorter than the

Radical	τ_c (ps)	$\tau_{c, NP}$ (ps)
TEMPOL	7(1)	—
TEMPONE	5(1)	—
TM thiol 2	5(1)	—
R1S1-AuNP	183(33)	3.7×10^3
R1S2-AuNP	565(113)	4.6×10^3

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The recognition capabilities of **R1S2-AuNP** were tested by ^1H -NMR. To avoid the effect of paramagnetic broadening of the NMR lines caused by the TEMPO headgroup, we tested AuNPs bearing only the group **S2** (**S2-AuNP**). The AuNPs were dissolved in CD_2Cl_2 along with CHCl_3 and MeOH, both at 20 mM. Fig. 4 shows a comparison of this spectrum with that obtained from a similar system without nanoparticles. There is a marked broadening of the MeOH signals in the presence of **S2-AuNPs** (Fig. 4c), whereas the signals of CHCl_3 are unaffected (Fig. 4b). This effect likely arises from the tendency of **S2-AuNPs** to act as hydrogen bond acceptors for MeOH molecules, thus concentrating them in the monolayer and increasing the exchange rate of OH protons. It is worth pointing out that such evidence could not be corroborated by saturation transfer experiments typically used in NMR chemosensing, due to the overlap between the MeOH signals and those arising from the oxomethylene headgroups of **S2** (~ 3.5 ppm in Fig. 4).



Nonetheless, as a control experiment, we acquired ^1H NMR spectra at varying temperatures (298 K, 283 K, and 278 K). These spectra exhibit changes predominantly limited to the chemical shifts of the peaks (see SI), thereby suggesting that the line broadening observed in presence of **S1-AuNPs** is unlikely due to be due to a decrease in correlation time but rather due to chemical exchange.

We performed OE-DNP experiments both at low (1.2 T) and high magnetic fields (9.4 T) recording the ^{13}C enhancements ($^{13}\text{C}_\epsilon$) on CHCl_3 and on mixtures of CHCl_3 and MeOH doped with **R1SX-AuNPs** or nitroxide radicals. The results are summarized in Fig. 5. At 1.2 T, $^{13}\text{C}_\epsilon$ on CHCl_3 doped with **R1S1-AuNP** is $^{13}\text{C}_\epsilon \sim 145$, which is only $\sim 25\%$ smaller than $^{13}\text{C}_\epsilon = 195$ obtained with TEMPOL (Fig. 5a). At 9.4 T, we tested a mixture of 500 mM ^{13}C - CHCl_3 and 500 mM of ^{13}C -MeOH dispersed in CHCl_3 . The analyte concentration was selected as in ref. 27 in order to optimize the ^{13}C sensitivity and the acquisition time on the DNP instrument. Once again, the enhancements obtained with TEMPOL ($^{13}\text{C}_\epsilon \sim 60$ on ^{13}C - CHCl_3 and $^{13}\text{C}_\epsilon \sim 5$ on ^{13}C -MeOH) are larger than those obtained with **R1SX-AuNPs**: on CHCl_3 , **R1S1-AuNPs** provide $^{13}\text{C}_\epsilon \sim 24$, which is further reduced to $^{13}\text{C}_\epsilon \sim 5$ when using **R1S2-AuNPs** (Fig. 5b). Although **R1S2-AuNPs** recognize MeOH molecules, we do not observe an enhancement on ^{13}C -MeOH.

These results can be interpreted with analytical models that describe the spin dynamics of OE-DNP in the liquid state (eqn 2).^{19,25,33,34,62} All $^{13}\text{C}_\epsilon$ measured on ^{13}C - CHCl_3 are positive, indicating that the electron–nuclear scalar coupling is larger than the competing dipolar one.³⁵ The rotational correlation time of the radical can contribute to the electron–nuclear dipolar coupling when for $\tau_c > 10$ –20 ps, effectively counteracting the scalar coupling and therefore reducing the enhancements.^{33,62}

Using the approach discussed in ref. 33, eqn (1) and (2) with $\tau_{\text{rot}} = \tau_c$, we extrapolated the $^{13}\text{C}_\epsilon$ enhancements as a function of the rotational correlation time τ_c of the radical (Fig. 5c), which is included in the term $J_{\text{rot}}(\omega, \tau_c)$ in eqn (2), while considering

unchanged all the other elements in the equations (s and f in eqn (1), $J_{\text{D}}(\omega, \tau_{\text{D}})$ and $J_{\text{sc}}(\omega, \tau_{\text{sc}})$ in eqn (2)). Despite the crude approximation, the simulations reproduce the trend observed in the experimental data and suggest that the low $^{13}\text{C}_\epsilon$ observed for **R1SX-AuNP** can be mainly ascribed to the slow rotation of the radical. By way of comparison, nitroxides grafted on silica surfaces show τ_c of the radical on the order of few nanoseconds,⁴⁷ which would lead to lower DNP performances at high magnetic fields as compared to AuNPs.

Finally, we will discuss why MeOH is not an optimal target for NP mediated OE-DNP, despite **R2S2-AuNPs** effectively working as molecular recognition agents for it. As suggested by our simulations on ^{13}C - CHCl_3 data, the enhancement $^{13}\text{C}_\epsilon \sim 5$ obtained with TEMPOL should be reduced by $\geq 60\%$ in presence of **R1S2-AuNPs** solely due to the radical mobility making it undetectable under our experimental conditions. Second, the requirements for an optimal agent for chemosensing and an optimal PA for OE-DNP are different. The spin polarization transfer in OE-DNP is very effective when there are modulations of the electron–nuclear hyperfine coupling on the order of the electron Larmor frequency ($J(\omega_e)$, eqn (2)): these arise from fast molecular dynamics (collisions, vibrations) on the order of picoseconds and lower. In contrast, as far as NMR chemosensing is concerned, effective nuclear–nuclear cross-relaxation requires the analyte's residence time within the monolayer to be comparable to, or exceed, the nanoparticle's rotational correlation time ($\tau_{\text{c,NP}}$), which is in the order of nanoseconds. This discrepancy in the dynamics might explain the limited efficacy of **R1SX-AuNPs** in polarizing the recognized target molecule *via* OE-DNP.

To possibly overcome these limitations and effectively enable molecular recognition, it appears necessary to modify the organic thiol and/or the linker to the nitroxide to adjust the mobility. AuNPs are an ideal platform for this task, due to the versatile chemistry of their monolayers, which can be controlled by adjusting the length and nature of the spacer between the Au core and

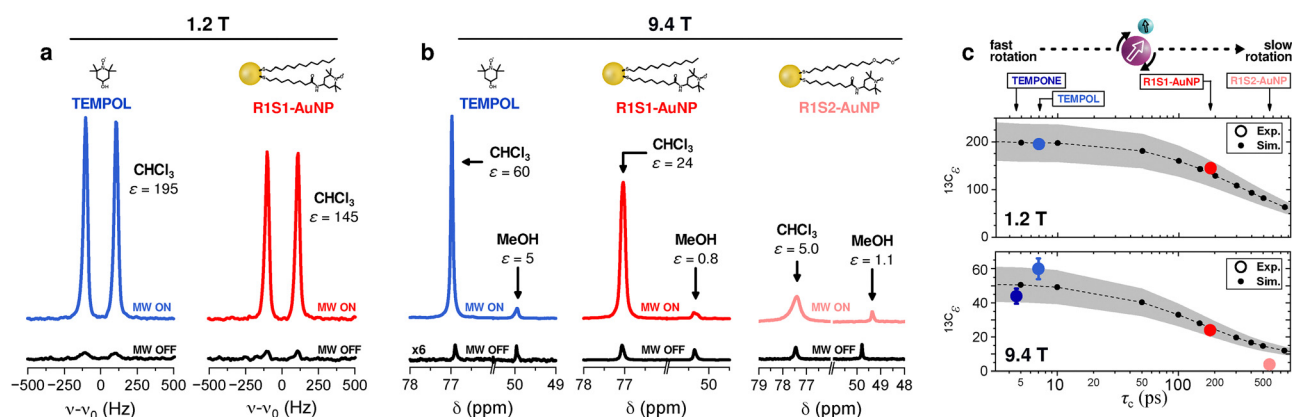


Fig. 5 (a) MW on and MW off ^{13}C DNP-NMR spectra of CHCl_3 doped with TEMPOL and **R1S1-AuNPs**. Spectra were rescaled to the same noise level. (b) MW on and MW off ^{13}C DNP-NMR spectra of ^{13}C - CHCl_3 (500 mM) and ^{13}C -MeOH (500 mM) dissolved in CHCl_3 and doped with (c) TEMPOL, and **R1S1-AuNPs**, and **R1S2-AuNPs**. Spectra were acquired with ^1H decoupling. The vertical axis is in arbitrary. (c) Experimental and predicted ^{13}C -NMR enhancements of ^{13}C - CHCl_3 as a function of the PA correlation time. $^{13}\text{C}_\epsilon$ at 1.2 T and 9.4 T plotted as a function of the rotational correlation time τ_c estimated from the EPR measurements. units. The colored band represents an error bar of 20% on the simulated points.



the headgroup, as well as by leveraging inter-chain interactions. Alongside this “direct” approach, where the nanoparticle headgroup carries the unpaired electron, we are currently investigating the feasibility of a “reverse” approach, wherein the nanoparticle headgroup is functionalized with a suitable moiety to be polarized through interaction with a radical species present in solution.

Conclusions

In this study, we investigated the feasibility of achieving selective OE-DNP in the liquid state by designing monolayer-protected nanoparticles that combine molecular recognition properties with the ability to function as polarizing agents for OE-DNP. Specifically, we designed and synthesized AuNPs (<2 nm in diameter) functionalized both with nitroxide radicals to act as PAs and with organic thiols that favour the formation of hydrogen bond networks. The morphological characterization shows that functionalized AuNPs are soluble and stable in organic solvents, and that the radical headgroup retains fast mobility (correlation time $\tau_c \sim 200$ –600 ps) thanks to the flexibility of the alkyl chain. Furthermore, AuNPs act as a molecular recognition agents for MeOH, as demonstrated by the increased line broadening due to its interaction with the monolayer. With OE-DNP measurements performed at low (1.2 T) and high (9.4 T) magnetic fields, we showed that AuNPs can be used as PAs. Although the interaction between MeOH and the functionalized AuNPs does not lead to a significantly enhanced polarization efficiency—likely due to the restrained mobility of the nitroxide headgroup—our results validate these nanostructures as viable polarizing agents in the context of OE-DNP. Furthermore, under optimal conditions, they might enable DNP-enhanced NMR-chemosensing experiments in solution for targeted selective DNP of small molecules. While silica has been suggested as a substrate to graft PAs, the modular and tunable surface chemistry of AuNPs establishes them as an ideal platform for designing DNP systems with additional chemosensing capabilities in solution. By exploiting diverse interactions, including hydrophobic forces, ion pairing, and metal–ligand coordination, these nanoreceptors can be precisely engineered to provide highly specific binding sites across a broad range of substrates. Indeed, the extensive variety of monolayers that can be assembled enables fine-tuning not only toward analyte selectivity but also over the strength and potentially over the dynamic features of the analyte binding process. Together, these features make AuNPs a uniquely controllable platform for precise modulation of analyte binding thermodynamics and kinetics, enabling the development of highly versatile chemosensors.

Author contributions

FR, FM, and TO conceived the study; BBM and GZ synthesized and characterized the compounds; BBM, LF, and TO performed EPR measurements and analysis; LY, AvdH, and TO performed the DNP measurements; BBM and FR performed the NMR measurements; TO and FR analysed the data. FR and TO wrote

the original draft and the final version was approved by all the authors.

Conflicts of interest

There are no conflicts to declare.

Data availability

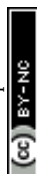
The data supporting this article have been included as part of the SI: Synthesis of and functionalization of AuNPs, morphological characterization, NMR and EPR characterization, DNP results. See DOI: <https://doi.org/10.1039/d5cp02465j>

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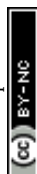
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