


 Cite this: *Phys. Chem. Chem. Phys.*,
 2023, 25, 822

Large ^{31}P -NMR enhancements in liquid state dynamic nuclear polarization through radical/target molecule non-covalent interaction†

 Maik Reinhard, ^{ab} Marcel Levien, ^{ab} Marina Bennati ^{ab} and
 Tomas Orlando ^{*a}

Dynamic nuclear polarization (DNP) is a method to enhance the low sensitivity of nuclear magnetic resonance (NMR) via spin polarization transfer from electron spins to nuclear spins. In the liquid state, this process is mediated by fast modulations of the electron-nuclear hyperfine coupling and its efficiency depends strongly on the applied magnetic field. A peculiar case study is triphenylphosphine (PPh_3) dissolved in benzene and doped with BDPA radical because it gives ^{31}P -NMR signal enhancements of two orders of magnitude up to a magnetic field of 14.1 T. Here we show that the large ^{31}P enhancements of BDPA/ PPh_3 in benzene at 1.2 T (i) decrease when the moieties are dissolved in other organic solvents, (ii) are strongly reduced when using a nitroxide radical, and (iii) vanish with pentavalent ^{31}P triphenylphosphine oxide. Those experimental observations are rationalized with numerical calculations based on density functional theory that show the tendency of BDPA and PPh_3 to form a weak complex via non-covalent interaction that leads to large hyperfine couplings to ^{31}P ($\Delta A_{\text{iso}} \geq 13$ MHz). This mechanism is hampered in other investigated systems. The case study of ^{31}P -DNP in PPh_3 is an important example that extends the current understanding of DNP in the liquids state: non-covalent interactions between radical and target can be particularly effective to obtain large NMR signal enhancements.

 Received 2nd September 2022,
 Accepted 29th November 2022

DOI: 10.1039/d2cp04092a

rsc.li/pccp

Introduction

Nuclear magnetic resonance (NMR) spectroscopy is one of the most widely used methods in analytical chemistry. It relies on the detection of small energy splittings associated with the nuclear Zeeman level and therefore suffers from low sensitivity. To overcome this limitation and expand the class of viable NMR experiments, several methods to enhance the NMR signal have been developed. Those are known as hyperpolarization methods, and they usually involve the spin polarization transfer from a highly polarized spin system to the target nuclei.¹ Among those methods, dynamic nuclear polarization (DNP) is one of the most versatile. It uses microwave (MW) irradiation to transfer spin polarization from unpaired electron spins, usually located on an organic radical, to the nuclear spins on a target molecule (Fig. 1a).

In the liquid state, the polarization transfer is governed by a cross-relaxation process known as Overhauser effect.^{2–5} The efficiency of this process strongly depends on the choice of the polarizing agent (PA), *i.e.*, the organic radical, and of the target molecule and nucleus, as well as on the applied magnetic field. The case of ^1H as target nucleus for DNP is rather unfavorable: ^1H -NMR signal enhancements for water at room temperature (<45 °C) are limited to $|\varepsilon| \lesssim 30$ for magnetic fields $B \gtrsim 9.4$ T.^{6,7} The reason for this is the dominant dipolar relaxation between the nuclear spin of ^1H and the electron spin of the radical, which decays rapidly for increasing magnetic fields.⁸ On the contrary, ^{13}C -DNP is much more effective because the dipolar relaxation is counterbalanced and overcome by scalar relaxation, a mechanism that originates from the Fermi contact interaction between the radical and the target molecule.⁹ ^{13}C -NMR signal enhancements in liquids can be up to $\varepsilon \sim 1000$ at a magnetic field of 3.4 T.¹⁰ Furthermore, recent reports showed that it is possible to obtain sizeable enhancements (>10) of ^{13}C -NMR signals in a variety of small molecules at high magnetic field (9.4 T),¹¹ even in water solutions.¹² Those findings sparked new interest in the method, and now several groups are committed to tackle the open challenges to make DNP in the liquid state applicable to routine NMR spectroscopy.^{13–22} Within this

^a ESR Spectroscopy Group, Max Planck Institute for Multidisciplinary Sciences, Am Fassberg 11, Göttingen, Germany. E-mail: tomas.orlando@mpinat.mpg.de

^b Department of Chemistry, Georg-August-University, Tammannstraße 4, Göttingen, Germany

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2cp04092a>



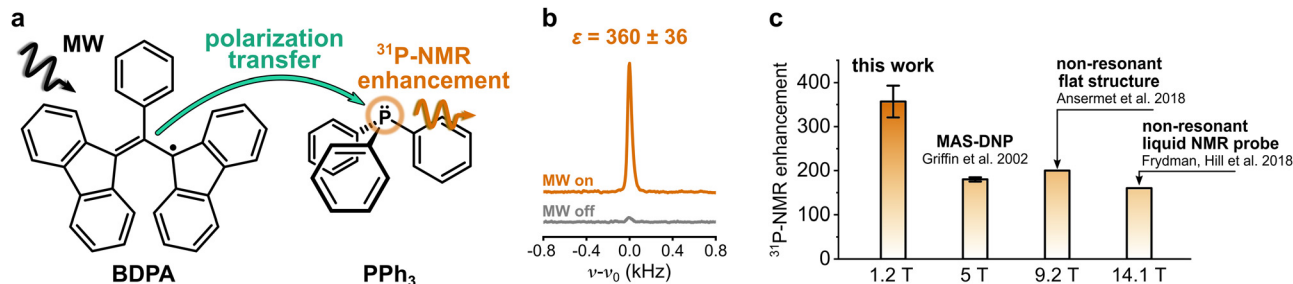
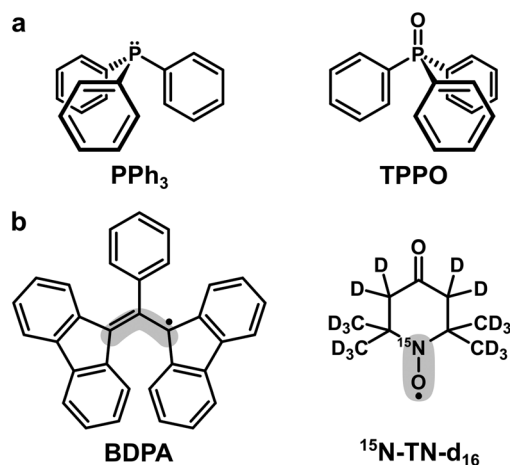


Fig. 1 (a) Sketch of the polarization transfer process between the PA (BDPA radical) and the target molecule (PPh_3). (b) ^{31}P -NMR spectra of PPh_3 in benzene doped with BDPA recorded at 1.2 T with and without MW irradiation. (c) ^{31}P -NMR enhancements reported at different magnetic fields for PPh_3 in benzene (fluorobenzene at 9.2 T) doped with BDPA radical.

context, it is important to design targets and PAs whose properties are specifically tailored to maximize the attainable NMR signal enhancements. To this aim, it is essential to identify the physical mechanisms that make the polarization transfer particularly effective.

DNP studies on target nuclei other than ^1H and ^{13}C have been only rarely reported. Here we consider the case of ^{31}P , a target nucleus that is often used in NMR studies of phospholipids (e.g. mixtures, morphology), in metabolomics (e.g. ATP monitoring), and in structural investigations of DNA.^{23–25} Most studies of ^{31}P -DNP date back to the early 80s, when MW and NMR technology limited the DNP measurements to magnetic fields below 1.2 T.^{26–30} The reported results show a strong dependence of ^{31}P -DNP on the chemical environment of ^{31}P ; specifically, they observed enhancements dominated by scalar relaxation for sterically exposed trivalent phosphorus, while pentavalent phosphorus compounds show dipolar dominated enhancement.³⁰ A unique case is the compound triphenylphosphine (PPh_3) (Fig. 1a): when PPh_3 is dissolved in benzene and doped with α,γ -Bisdiphenylene- β -phenylallyl radical (BDPA), ^{31}P -NMR signal enhancements of two orders of magnitude are readily reachable, even at larger magnetic fields. ^{31}P -NMR enhancements $\epsilon > 150$ were reported for PPh_3 at $B \approx 5$ T,³¹ and more recently, up to $\epsilon \approx 160$ at 14.1 T (Fig. 1c).^{13,14} Notably, those enhancements show almost no dependence on the external magnetic field.

Here we compare the performance of two polarizing agents, BDPA and the nitroxide radical TEMPONE (TN) for enhancing ^{31}P -NMR signal of PPh_3 , and of its oxidized counterpart, *i.e.* triphenylphosphine oxide (TPPO) (Scheme 1). Those systems were also tested in different organic solvents, which have a significant effect on the DNP outcome. DNP measurements performed on a hybrid EPR/NMR setup operating at 1.2 T allowed us to quantify the efficiency of the Overhauser effect and show that ^{31}P -DNP is particularly favourable only for the system BDPA/ PPh_3 in benzene. With numerical calculations based on density functional theory (DFT), we rationalize this observation showing that BDPA and PPh_3 form a weak complex that favors large hyperfine couplings. Our findings corroborate the model that the formation of a complex between radical and target molecule that leads to large hyperfine coupling values is crucial to obtain large NMR signal enhancements on small molecules in the liquid state.



Scheme 1 (a) Trivalent (PPh_3) and pentavalent (TPPO) phosphorus compounds used as target molecules. (b) Organic radicals used as PAs. The grey areas mark the sites with the largest electron spin density: $\sim 40\%$ for BDPA and $\sim 90\%$ for ^{15}N -TN- d_{16} .¹⁸

Experimental methods

Triphenylphosphine (PPh_3) and triphenylphosphine oxide (TPPO) were used as target molecules. As polarizing agents we utilized two organic radicals: BDPA (α,γ -Bisdiphenylene- β -phenylallyl, in 1 : 1 complex with benzene) and TEMPONE (TN) in its deuterated and ^{15}N -labeled version (^{15}N -TN- d_{16} , 4-Oxo-2,2,6,6-tetramethylpiperidine- d_{16} , 1- ^{15}N -1-oxyl). Both targets and PAs were purchased from Sigma-Aldrich and used without further purification. Organic solvents chloroform (CHCl_3), tetrachloromethane (CCl_4), benzene, pyridine and DMSO were purchased from Merck KGaA.

The concentration of phosphorous compounds in the chosen solvent ranged between 1 M and 4 M. TPPO was soluble in CHCl_3 , pyridine and DMSO (1 M), while its solubility was too low in the less polar organic solvents benzene and CCl_4 and prevented NMR detection at 1.2 T. The radical concentration was ~ 8 –10 mM and was calibrated for each sample with CW-EPR experiments. All samples were prepared in stocks of 100–550 μL , which were then degassed by freeze–pump–thaw cycles (three to five) to remove dissolved oxygen. The solutions were transferred into a glove box, where 4–4.5 μL were used to fill a quartz tube (1.6 mm outer diameter, Wilmad-LabGlass),



which was then sealed with a flame. The sample preparation procedure results in an error in the radical concentration of 10% for tetrachloromethane, benzene, pyridine, and DMSO samples and 15% for chloroform samples, the latter due to a lower boiling point. Radical stability after sealing was monitored with CW-EPR. Radicals were stable in the tested solvents for the whole duration of the DNP, NMR, and EPR measurements (~ 12 h).

^{31}P -DNP measurements at 1.2 T were performed on a hybrid EPR/NMR system which combines a Bruker ElexSys E580 EPR spectrometer and a Bruker AVANCE III NMR console. We used a Bruker ER-5106QT/W resonator and a home built copper coil wrapped around the Q-Band quartz tube with 4 to 5 turns for NMR detection.¹¹ With this arrangement, microwave and radio-frequency can be applied simultaneously, allowing for EPR, NMR, and DNP measurements on the same sample. The MW power was adjusted to avoid severe heating during MW irradiation.

Optimized geometries of the radical/target molecule pairs were computed with DFT using Orca 5.0.2³² with B3LYP as functional, def2-TZVPP as basis set, and the dispersion correction D3BJ. Several structures (≥ 12) with different starting orientations of the PA and target molecule were computed for each system; the starting orientations were chosen to take into account different approaches of the two molecules (ESI†). For each optimized geometry, we assessed the non-covalent interactions by calculating the interaction energy E_{int} , defined as the difference between the electronic energy of the dimer and the electronic energy of the two monomers.³³ The values were corrected for the basis set superposition error with the Boys-Bernardi procedure (ESI†).^{34–37} The calculations were performed in vacuum as well as with the implicit solvation model C-PCM^{38,39} for benzene and chloroform, using the same starting geometries. We calculated the isotropic hyperfine coupling to ^{31}P using EPR-III basis set⁴⁰ for H, C, N, and O atoms and IGLO-II⁴¹ for P atom.

Results and discussion

^{31}P -NMR signals of PPh_3 and TPPO in various solvents were recorded with (DNP) and without MW irradiation (thermal) at a magnetic field of 1.2 T. ^{31}P -NMR signals were integrated to calculate the enhancements $\varepsilon = I_{\text{DNP}} \cdot n_{\text{thermal}} / (I_{\text{thermal}} \cdot n_{\text{DNP}})$, where I is the integral and n is the number of scans. Fig. 2a reports the enhancements of the investigated systems. In PPh_3 , the highest ^{31}P -NMR signal enhancement, *i.e.* $^{31}\text{P}\varepsilon = 360 \pm 36$, has been obtained in benzene and when using BDPA. The radical $^{15}\text{N-TN-d}_{16}$ is not as efficient, and gives ^{31}P -NMR signal enhancements that are a factor of 5 to 10 smaller, with a maximum of $^{31}\text{P}\varepsilon = 21 \pm 3$ for PPh_3 in benzene. No ^{31}P -NMR signal enhancements were observed on the TPPO molecule with all utilized solvents and PAs. Earlier reports³⁰ suggest that the large enhancements observed in PPh_3 could be due to the lone pair of the trivalent ^{31}P , which tends to coordinate with the unpaired electron of the radical. On the contrary, ^{31}P in TPPO is in the pentavalent configuration, which hampers this mechanism.



Fig. 2 (a) ^{31}P -NMR enhancements recorded at 1.2 T and (b) coupling factors calculated with eqn (1) for both PPh_3 and TPPO doped with BDPA and $^{15}\text{N-TN-d}_{16}$ in different solvents.

The enhancements give only partial insight into the efficiency of ^{31}P -DNP. Indeed, $^{31}\text{P}\varepsilon$ is determined by several parameters, as given by the Overhauser equation:^{2,4}

$$\varepsilon = 1 - s_{\text{eff}} \cdot f \cdot \xi \cdot \frac{\gamma_e}{\gamma_n} \quad (1)$$

where ε is directly proportional to: (i) the effective saturation factor s_{eff} , quantifying how much of the EPR line is saturated; (ii) the leakage factor f , which stands for the amount of paramagnetic relaxation over the total nuclear relaxation; (iii) the ratio of electron (γ_e) and nuclear (γ_n) gyromagnetic ratios, which is $\gamma_e/\gamma_n \sim 1626$ for ^{31}P ; (iv) the coupling factor ξ that accounts for cross-relaxation transition rates between the electron and the nuclei and ultimately gives the efficiency of the polarization transfer.^{5,42,43} Therefore, to compare the DNP efficiency among the tested systems, it is important to quantitatively access the coupling factor ξ with eqn (1) once f and s are known.

The leakage factor f depends on the nuclear relaxation times of the target nucleus measured with ($T_{1,n}$) and without radical ($T_{1,\text{dia}}$), and $f = 1 - T_{1,n}/T_{1,\text{dia}}$. f depends ultimately on the concentration of the PA (c_{PA}) and for our samples is $f \sim 0.8$ or larger for $c_{\text{PA}} \sim 8\text{--}10$ mM (Table 1). The effective saturation factor s_{eff} depends on the type of radical chosen as PA and can vary significantly. The EPR spectrum of BDPA shows a hyperfine structure that arises from the coupling to the protons on the diphenylene and phenyl rings (ESI† Fig. S2). At 34 GHz and



Table 1 Overhauser parameters of ^{31}P -DNP of triphenylphosphine (PPh_3) for the two different PAs in different organic solvents at 1.2 T. The leakage factor f was calculated as $f = 1 - T_{1,n}/T_{1,\text{dia}}$ using $T_{1,\text{dia}} = 10.9$ s for PPh_3 in CHCl_3 , 19.0 s in CCl_4 , 20.5 s in benzene and 21.6 s in pyridine, respectively. Errors are 10% for c_{PA} (15% for CHCl_3 samples), 10% for c_{PPh_3} (15% for CCl_4), 10% for $T_{1,n}$, T_{Buildup} , f , s_{eff} , and ϵ , and is 15% for ζ (15% for ϵ and 20% for ζ in pyridine). Errors are increased for ϵ and ζ to 15% and 25%, respectively if $T_{1,n} \neq T_{\text{Buildup}}$, which indicates sample heating during MW irradiation

PA	Solvent	c_{PA} (mM)	c_{PPh_3} (M)	$T_{1,n}$ (s)	T_{Buildup} (s)	f	s_{eff}	ϵ	ζ
BDPA	CHCl_3	10	4	1.2	1.1	0.89	0.91	51	-0.038
BDPA	Pyridine	10	2	3.7	3.8	0.83	0.82	127	-0.114
BDPA	CCl_4	10	2	3.8	4.5	0.80	0.90	150	-0.129
BDPA	Benzene	10	2	6.1	8.4	0.70	1.00	360	-0.312
^{15}N -TN- d_{16}	CHCl_3	10	4	2.3	1.3	0.79	0.34	14	-0.030
^{15}N -TN- d_{16}	CCl_4	10	2	2.3	2.0	0.88	0.30	17	-0.037
^{15}N -TN- d_{16}	Benzene	8	2	3.5	3.4	0.83	0.16 ^a	21	-0.092

^a For this sample, the MW power was reduced to avoid excessive sample heating.

$c_{\text{PA}} \sim 10$ mM the lines are merged into one and the total spectral width is ~ 1 mT; this ensures an almost complete saturation of the EPR line when MW irradiation is applied on resonance with the center of the line. On the contrary, the spectrum of ^{15}N -TN- d_{16} consists of two sharp lines separated by ~ 2 mT stemming from the hyperfine coupling with the ^{15}N nucleus. When one of the two lines is irradiated on resonance (as in a DNP experiment), the other line is only partially saturated *via* a mechanism known as ELDOR effect.^{44–46} The saturation factors were measured with an ELDOR (electron double resonance) experiment (ESI⁺) and, under similar experimental conditions, s_{eff} is $s_{\text{eff}} > 0.8$ for BDPA, while it is limited to $s_{\text{eff}} < 0.35$ for ^{15}N -TN- d_{16} (Table 1). The leakage and the saturation factors together cannot explain the difference in enhancements among the tested systems.

With those values of f and s_{eff} and eqn (1), one can calculate the coupling factor $^{31}\text{P}\zeta$ (Fig. 2b). The quantification of this experimental parameter leads to three important observations: (1) BDPA is a better PA than ^{15}N -TN- d_{16} for ^{31}P and is characterized by a more efficient polarization transfer, *i.e.* $|^{31}\text{P}\zeta_{\text{BDPA}}| > |^{31}\text{P}\zeta_{\text{TN}}|$; this is rather surprising because nitroxide radicals are superior PAs both for ^1H and ^{13}C DNP;¹⁸ (2) no enhancements are observed on ^{31}P -TPPO at 1.2 T and $^{31}\text{P}\zeta \sim 0$ independently of the solvent; (3) the solvent influences the ^{31}P -DNP performance: in particular, benzene is the best solvent, while CCl_4 , pyridine, and chloroform follow in this order, for both BDPA and TN.

In the following, we rationalize those observations and investigate the interactions between target molecule and PA that could affect the polarization transfer process. From previous studies^{5,43} it is known that, when the enhancements are positive ($^{31}\text{P}\epsilon > 0$, and therefore $^{31}\text{P}\zeta < 0$), the scalar (or Fermi contact) interaction between the electron and the nucleus is dominant over the dipolar one. The scalar interaction drives electron-nuclear cross-relaxation through modulation of the isotropic hyperfine coupling (A_{iso}) between the electron spin of the radical and the nuclear spin of the target. In the case of small molecules interacting with organic radicals in liquids at room temperature, those modulations arise from a collisional process.⁴⁷ Particularly, one has $A_{\text{iso}} \neq 0$ during the collision and $A_{\text{iso}} = 0$ when the two molecules diffuse apart. The collisional process is described by the following spectral density:^{47,48a}

$$J(\omega_e, \tau_i) = \sum_i \frac{4\pi^2 \langle A_{\text{iso},i}^2 \rangle}{\tau_{p,i}} [\tau_i \exp(-\tau_i \omega_e)]^2 \quad (2)$$

where the index i stands for the i -type of collision. The duration of each collision is τ , the collision rate is $1/\tau_p$, and the hyperfine coupling is A_{iso} (in Hz in eqn (2)). This analytical model is a good approximation of more sophisticated numerical simulations^{48a,b} and can be effectively used to interpret DNP data. Furthermore, as earlier proposed by the group of Dorn,^{9c,d} the hyperfine coupling calculated with DFT correlates with the scalar enhancement observed experimentally in a variety of compounds, provided that the timescale of the collisional process ($\tau_i, \tau_{p,i}$) is comparable. Here we utilize a similar methodology to examine the interaction between target molecules (PPh_3 and TPPO) and the radicals (BDPA and TN).

We computed optimized geometries (up to 15 optimized structures for each complex for each solvent) and isotropic hyperfine couplings of the complexes BDPA/ PPh_3 , TN/ PPh_3 , BDPA/TPPO, and TN/TPPO. The results are summarized in Fig. 3, where each point represents A_{iso} for each optimized structure in vacuum, benzene, and chloroform. When PPh_3 interacts with the BDPA radical, A_{iso} spans the largest range,



Fig. 3 Hyperfine coupling A_{iso} calculated for each optimized structure of the investigated complexes. 15 structures were computed for both PPh_3 systems, and 12 for TPPO. The circles are calculated values and they partially overlap. Color bands are visual aids.



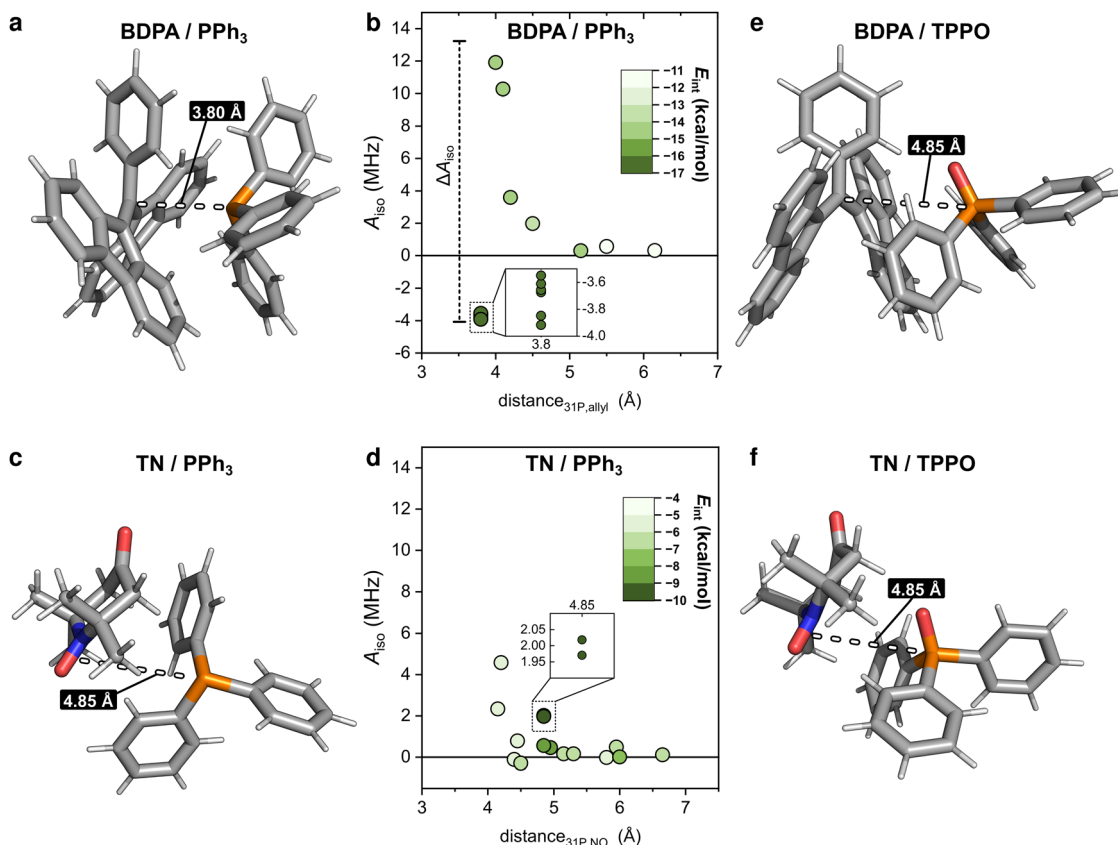


Fig. 4 (a) Structure of the complex BDPA/PPh₃ in benzene corresponding to the minimum interaction energy $E_{\text{int}} = -16.3$ kcal mol⁻¹ and with $A_{\text{iso}} = -3.7$ MHz where the lone pair of the ³¹P atom is pointing to the allyl group of the BDPA. (b) Hyperfine coupling A_{iso} calculated for each of the optimized structures in benzene and plotted as a function of the distance between ³¹P and the allyl group of the BDPA. The distance is the mean of the distances between ³¹P and the two closest carbons of the allyl group of BDPA. The color map shows the interaction energy E_{int} of the complex. (c) Structure of the complex TN/PPh₃ in benzene with minimum interaction energy $E_{\text{int}} = -9.5$ kcal mol⁻¹ and with $A_{\text{iso}} = 2.02$ MHz. (d) Hyperfine coupling A_{iso} calculated for each optimized structure in benzene and plotted as a function of the distance between ³¹P and the NO group of TN. (e) Structure of the complex BDPA/TPPO in CHCl₃ with minimum interaction energy $E_{\text{int}} = -15.1$ kcal mol⁻¹ and with $A_{\text{iso}} = -0.07$ MHz. (f) Structure of the complex TN/TPPO in chloroform with minimum interaction energy $E_{\text{int}} = -9.4$ kcal mol⁻¹ and with $A_{\text{iso}} = -0.02$ MHz. Atom colour code: H white, C grey, N blue, O red, P orange.

$\Delta A_{\text{iso}} \sim 17$ MHz, going from negative ($A_{\text{iso}} \sim -4$ MHz) to positive ($A_{\text{iso}} > 10$ MHz) in vacuum. The range is smaller for the system TN/PPh₃, $\Delta A_{\text{iso}} \sim 14$ MHz, and it is further reduced to $\Delta A_{\text{iso}} \sim 5$ MHz when the solvent model is used. For TPPO, the hyperfine couplings are more than 10 times smaller and span the range -0.4 MHz $< A_{\text{iso}} < 0.7$ MHz.

To clarify why the two radicals lead to a different A_{iso} to the ³¹P of PPh₃, we took a closer look at the calculated geometries. By considering the pairs BDPA/PPh₃, one notices that large hyperfine couplings ($|A_{\text{iso}}| > 3$ MHz) are observed when the lone pair of ³¹P is pointing directly to the allyl group of BDPA, which carries the majority of the electron spin density ($\sim 40\%$) (Fig. 4a and Scheme 1).¹⁸ This configuration is unusual because the accessibility of the allyl group by small solvent molecule is limited.¹⁸ The interaction energy E_{int} calculated for each BDPA/PPh₃ optimized structure show that complexes with short ³¹P-allyl group distances ($d \leq 4$ Å) and large hyperfine couplings ($|A_{\text{iso}}| > 3$ MHz) are energetically favored over other geometries where ³¹P is further away from the allyl group (Fig. 4b and Fig. S4, ESI[†]). The same is found for calculations performed in vacuum, benzene, and chloroform.

We performed a similar analysis for the system TN/PPh₃. In terms of distance, ³¹P cannot get as close to the electron spin density localized on the NO group of TN ($d > 4.2$ Å in benzene) (Fig. 4c). The interaction energy E_{int} has a minimum at distances $d \sim 4.85$ Å, which corresponds to a hyperfine coupling $|A_{\text{iso}}| \sim 2$ MHz (Fig. 4d and Fig. S4, ESI[†]). Therefore, TN/PPh₃ show a non-covalent interaction that is less efficient in terms of hyperfine coupling. Additional calculations of E_{int} were performed with fixed values for the distance between ³¹P and the electron spin density (both for BDPA and TN), and confirm the tendency of PPh₃ to approach BDPA, and specifically near the allyl group, while this is not the case for the pair TN/PPh₃ (Fig. S6a, ESI[†]). Those results indicate that the large ³¹P-NMR enhancements observed on PPh₃ doped with BDPA are facilitated by a non-covalent interaction that leads to large values of hyperfine coupling A_{iso} , which in turn renders the scalar relaxation from the collisional process more efficient (eqn (2)).

In regard to TPPO, the absence of enhancements must be a consequence of particularly small hyperfine couplings (Fig. 3). The oxygen atom bound to ³¹P hampers a close approach between the ³¹P of TPPO and BDPA (Fig. 4e) and leads to low



hyperfine coupling values ($|A_{\text{iso}}| < 0.7$ MHz). In the case of TN, TPPO reaches a distance ^{31}P -NO group $d \geq 4.2$ Å in both vacuum and chloroform, which is similar to the distances obtained for PPh₃ (Fig. 4f). Nevertheless, the hyperfine coupling $|A_{\text{iso}}|$ remains below 0.7 MHz. As already observed for ^{13}C -DNP of carbonyl groups,^{9c,d,10} the oxygen atom might be responsible for withdrawing electron spin density from ^{31}P ,⁴⁹ which results in lower A_{iso} values thus decreasing the attainable enhancements.

The last remark concerns the role of the solvent. Recent experimental and theoretical studies on DNP mechanisms have been focused on the interaction of only two moieties, i.e. the radical and a small molecule that is at the same time the solvent and the target molecule.^{7,9–11,48} Our experimental data show that a third player, i.e. the solvent, has a considerable effect in ^{31}P -DNP and possibly contributes to the spin polarization transfer. In the context of DNP, the solvent (i) determines the degree of MW absorption, and (ii) when diffusivity increases, dipolar and scalar relaxations increase.⁵⁰ The diffusion coefficient for benzene at 298 K is $D = 2.2 \times 10^5 \text{ cm}^2 \text{ s}^{-1}$,⁵¹ while it is lower for CHCl₃ ($D = 2.14 \times 10^5 \text{ cm}^2 \text{ s}^{-1}$),¹⁷ CCl₄ ($D = 1.4 \times 10^5 \text{ cm}^2 \text{ s}^{-1}$),⁵¹ and pyridine ($D = 1.54 \times 10^5 \text{ cm}^2 \text{ s}^{-1}$).⁵² Although a faster collision rate enabled by a larger D might favor ^{31}P -DNP in benzene, the minor differences in D cannot explain the large differences observed in coupling factors. The solvent might affect the kinetics of the interaction between radical and target molecule, either stabilizing a non-covalent interaction or preventing its formation. Despite the presented numerical analysis considers the effect of the solvent through an implicit solvation model, it is not sufficient to gain insight in these mechanisms. To the best of our knowledge, more sophisticated simulations tools based on molecular dynamics and used for other systems⁴⁸ are not yet available for large radicals like BDPA. Such investigations will be the subject of future works.

Conclusions

We investigated large ^{31}P -NMR enhancements that have been observed from 1.2 T (this study) up to 14.1 T^{13,14,31} on the system BDPA/PPh₃. Our DNP data recorded on a hybrid EPR/NMR instrument operating at 1.2 T show that those enhancements are attenuated when TEMPONE radical is used as a PA, and vanish when triphenylphosphine oxide (TPPO) is the target molecule. The experimental observations were interpreted in the context of the collisional model for scalar relaxation with the support of DFT calculations and show that PPh₃ forms a weak complex with the polarizing agent BDPA, which favors large hyperfine couplings. This mechanism is precluded when ^{31}P is in the pentavalent configuration or when a nitroxide is used. This finding provides a novel aspect within the context of DNP in the liquid state and shows how the choice of the optimal PA is strongly dependent on the chemical environment of the target nucleus. We foresee that the design of future PAs for liquid DNP, possibly supported by DFT calculations, will use non-covalent interactions between target molecule and PA as an effective strategy to boost the NMR enhancements.

Data availability

Original data associated with Fig. 2 (ELDOR spectra, NMR spectra, NMR relaxation data), and output files of all geometries optimizations shown in Fig. 3 and 4 can be downloaded free of charge from the Göttingen Research Online website (DOI: [10.25625/A5WDZW](https://doi.org/10.25625/A5WDZW)).

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors acknowledge Dr. Igor Tkac, Dr. Markus Hiller, and Dr. Andreas Meyer for the fruitful discussions. Financial support has been provided by the Max Planck Society, the ERC advanced grant Bio-enMR 101020262, and the Deutsche Forschungsgemeinschaft (DFG) – Project number 455993474. This work used the Scientific Compute Cluster at GWDG, the joint data center of Max Planck Society for the Advancement of Science (MPG) and University of Göttingen.

Notes and references

- J. H. Ardenkjaer-Larsen, G. S. Boebinger, A. Comment, S. Duckett, A. S. Edison, F. Engelke, C. Griesinger, R. G. Griffin, C. Hilty, H. Maeda, G. Parigi, T. F. Prisner, E. Ravera, P. J. M. van Bentum, S. Vega, A. Webb, C. Luchinat, H. Schwalbe and L. Frydman, *Angew. Chem.*, 2015, **54**, 9162–9185.
- A. W. Overhauser, *Phys. Rev.*, 1953, **92**, 411–415.
- T. R. Carver and C. P. Slichter, *Phys. Rev.*, 1953, **92**, 212–213.
- I. Solomon, *Phys. Rev.*, 1955, **99**, 559–565.
- H. H. Hausser and D. Stehlik, Dynamic nuclear polarization in liquids. In J. S. Waugh, ed., *Advances in Magnetic Resonance*, Academic Press, 1968, vol. 3, 79–139.
- C. Griesinger, M. Bennati, H. M. Vieth, C. Luchinat, G. Parigi, P. H. Engelke, S. J. Glaser, V. Denysenkov and T. F. Prisner, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2012, **64**, 4–28.
- V. P. Denysenkov and T. F. Prisner, *eMagRes*, 2019, 41–54.
- M. Bennati, C. Luchinat, G. Parigi and M.-T. Tuerke, *Phys. Chem. Chem. Phys.*, 2010, **12**, 5902–5910.
- (a) M. Bennati and T. Orlando, *eMagRes*, 2019, **8**, 11–18; (b) M. D. Lingwood and S. Han, *J. Magn. Reson.*, 2009, **201**, 137–145; (c) X. Wang, W. C. Isley, S. Salido, Z. Sun, L. Song, C. J. Cramer and H. Dorn, *Chem. Sci.*, 2015, **6**, 6482–6495; (d) J. L. Russ, J. Gu, K.-H. Tsai, T. Glass, J. C. Duchamp and H. C. Dorn, *J. Am. Chem. Soc.*, 2007, **129**, 7018–7027.
- G. Liu, M. Levien, N. Karschin, G. Parigi, C. Luchinat and M. Bennati, *Nat. Chem.*, 2017, **9**, 676–680.
- T. Orlando, R. Dervisoglu, M. Levien, I. Tkach, T. F. Prisner, L. Andreas, V. Denysenkov and M. Bennati, *Angew. Chem., Int. Ed.*, 2019, **58**, 1402–1406.



- 12 D. Dai, X. Wang, Y. Liu, X. Yang, C. Glaubitz, V. P. Denysenkov, X. He, T. Prisner and J. Mao, *Nat. Commun.*, 2021, **12**, 6880.
- 13 D. Yoon, A. I. Dimitriadis, M. Soundararajan, C. Caspers, J. Genoud, S. Alberti, E. de Rijk and J.-P. Ansermet, *Anal. Chem.*, 2018, **90**, 5620–5626.
- 14 T. Dubroca, A. N. Smith, K. J. Pike, S. Froud, R. Wylde, B. Trociewitz, J. McKay, F. Mentink-Vigier, J. van Tol, S. Wi, W. Brey, J. R. Long, L. Frydman and S. Hill, *J. Magn. Reson.*, 2018, **289**, 35–44.
- 15 T. Dubroca, S. Wi, J. van Tol, L. Frydman and S. Hill, *Phys. Chem. Chem. Phys.*, 2019, **21**, 21200–21204.
- 16 V. P. Denysenkov, D. Dai and T. F. Prisner, *J. Magn. Reson.*, 2022, **337**, 107185.
- 17 M. Levien, M. Hiller, I. Tkach, M. Bennati and T. Orlando, *J. Chem. Phys. Lett.*, 2020, **11**, 1629–1635.
- 18 M. Levien, M. Reinhard, M. Hiller, I. Tkach, M. Bennati and T. Orlando, *Phys. Chem. Chem. Phys.*, 2021, **23**, 4480–4485.
- 19 M. G. Concilio, M. Soundararajan, L. Frydman and I. Kuprov, *J. Magn. Reson.*, 2021, **326**, 106940.
- 20 M. G. Concilio, I. Kuprov and L. Frydman, *Phys. Chem. Chem. Phys.*, 2022, **24**, 2118–2125.
- 21 (a) A. A. Kuzhelev, D. Dai, V. P. Denysenkov, I. A. Kirilyuk, E. G. Bagryanskaya and T. F. Prisner, *J. Phys. Chem. C*, 2021, **125**, 25651–25659; (b) A. A. Kuzhelev, D. Dai, V. P. Denysenkov and T. F. Prisner, *J. Am. Chem. Soc.*, 2022, **144**, 1164–1168.
- 22 T. J. Keller and T. Maly, *Magn. Reson.*, 2021, **2**, 117–128; Y. Rao, A. Venkatesh, P. Moutzouri and L. Emsley, *J. Phys. Chem. Lett.*, 2022, **13**, 7749–7755.
- 23 J. Schiller, M. Müller, B. Fuchs, K. Arnold and D. Huster, *Curr. Anal. Chem.*, 2007, **3**, 283–301.
- 24 J. Abi-Ghanem, B. Heddi, N. Foloppe and B. Hartmann, *Nucleic Acids Res.*, 2010, **38**, 3.
- 25 F. Bhinderwala, P. Evans, K. Jones, B. R. Laws, T. G. Smith, M. Morton and R. Powers, *Anal. Chem.*, 2020, **92**, 9536–9545.
- 26 R. A. Dwek and R. E. Richards, *Chem. Commun.*, 1966, 581–582.
- 27 P. W. Atkins, R. A. Dwek, J. B. Reid and R. E. Richards, *Mol. Phys.*, 1967, **13**, 175–180.
- 28 J. A. Potenza, P. J. Caplan and E. H. Poindexter, *J. Chem. Phys.*, 1968, **49**, 2461.
- 29 E. H. Poindexter, R. A. Dwek and J. A. Potenza, *J. Phys. Chem.*, 1969, **51**, 628.
- 30 J. A. Potenza, E. H. Poindexter, P. J. Caplan and R. A. Dwek, *J. Am. Chem. Soc.*, 1969, **91**(16), 4356–4360.
- 31 N. M. Loening, M. Rosay, V. Weis and R. G. Griffin, *J. Am. Chem. Soc.*, 2002, **124**, 8808–8809.
- 32 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**, 73–78.
- 33 J. Rezáč and P. Hobza, *Chem. Rev.*, 2016, **116**, 5038–5071.
- 34 P. Politzer, J. S. Murray and T. Clark, *Phys. Chem. Chem. Phys.*, 2013, **15**, 11178.
- 35 J. Lapp and S. Scheiner, *J. Phys. Chem. A*, 2021, **125**, 5069–5077.
- 36 S. Scheiner and S. Hunter, *ChemPhysChem*, 2022, 202200011.
- 37 S. F. Boys and F. Bernardi, *Mol. Phys.*, 1970, **19**, 553.
- 38 J. Tomasi, B. Mennucci and R. Cammi, *Chem. Rev.*, 2005, **105**(8), 2999–3094.
- 39 V. Barone and M. Cossi, *J. Phys. Chem. A*, 1998, **102**(11), 1995–2001.
- 40 F. Neese, *Wiley Interdiscip. Rev.: Comput. Mol. Sci.*, 2012, **2**(1), 73–78.
- 41 W. Kutzelnigg, U. Fleischer and C. van Wüllen, Shielding Calculations: IGLO Method, in *eMagRes*, ed. R. K. Harris and R. L. Wasylshen, 2007.
- 42 M. Bennati, C. Luchinat, G. Parigi and M. Türke, *Phys. Chem. Chem. Phys.*, 2010, **12**, 5902–5910.
- 43 E. Ravera, C. Luchinat and G. Parigi, *J. Magn. Reson.*, 2016, **264**, 78–87.
- 44 J. S. Hyde, J. C. W. Chien and J. H. Freed, *J. Chem. Phys.*, 1968, **48**, 4226–4233.
- 45 M. T. Türke and M. Bennati, *Phys. Chem. Chem. Phys.*, 2011, **13**, 3630.
- 46 N. Enkin, G. Liu, M. del, C. Gimenez-Lopez, K. Porfyrikis, I. Tkach and M. Bennati, *Phys. Chem. Chem. Phys.*, 2015, **17**, 11144–11149.
- 47 W. Müller-Warmuth, R. Vilhjalmsón, P. Gerlof, J. Smidt and J. Trommel, *Mol. Phys.*, 1976, **31**, 1055–1067.
- 48 (a) T. Orlando, I. Kuprov and M. Hiller, *J. Magn. Reson. Open*, 2022, **10–11**, 100040; (b) S. E. Küçük and D. Sezer, *Phys. Chem. Chem. Phys.*, 2016, **18**, 9353–9357.
- 49 M. Bennati, *eMagRes*, 2017, **6**, 271–282.
- 50 S. G. J. van Meerten, M. C. D. Tayler, A. P. M. Kentgens and P. J. M. van Bentum, *J. Magn. Reson.*, 2016, **267**, 30–36.
- 51 A. F. Collings and R. Mills, *Trans. Faraday Soc.*, 1970, **66**, 2761–2766.
- 52 M. Holz, X. Mao, D. Seiferling and A. Sacco, *J. Chem. Phys.*, 1996, **104**, 669–679.

