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# Challenges in preparing, preserving and detecting *para*-water in bulk: overcoming proton exchange and other hurdles

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*Para*-water is an analogue of *para*-hydrogen, where the two proton spins are in a quantum state that is antisymmetric under permutation, also known as singlet state. The populations of the nuclear spin states in *para*-water are believed to have long lifetimes just like other Long-Lived States (LLSs). This hypothesis can be verified by measuring the relaxation of an excess or a deficiency of *para*-water, also known as a "Triplet–Singlet Imbalance" (TSI), *i.e.*, a difference between the average population of the three triplet states T (that are symmetric under permutation) and the population of the singlet state S. In analogy with our recent findings on ethanol and fumarate, we propose to adapt the procedure for Dissolution Dynamic Nuclear Polarization (D-DNP) to prepare such a TSI in frozen water at very low temperatures in the vicinity of 1.2 K. After rapid heating and dissolution using an aprotic solvent, the TSI should be largely preserved. To assess this hypothesis, we studied the lifetime of water as a molecular entity when diluted in various solvents. In neat liquid H<sub>2</sub>O, proton exchange rates have been characterized by spin-echo experiments on oxygen-17 in natural abundance, with and without proton decoupling. One-dimensional exchange spectroscopy (EXSY) has been used to study proton exchange rates in H<sub>2</sub>O, HDO and D<sub>2</sub>O mixtures diluted in various aprotic solvents. In the case of 50 mM H<sub>2</sub>O in dioxane-*d*<sub>8</sub>, the proton exchange lifetime is about 20 s. After dissolving, one can observe this TSI by monitoring intensities in oxygen-17 spectra of H<sub>2</sub>O (if necessary using isotopically enriched samples) where the AX<sub>2</sub> system comprising a "spy" oxygen A and two protons X<sub>2</sub> gives rise to binomial multiplets only if the TSI vanishes. Alternatively, fast chemical addition to a suitable substrate (such as an activated aldehyde or ketone) can provide AX<sub>2</sub> systems where a carbon-13 acts as a spy nucleus. Proton signals that relax to equilibrium with two distinct time constants can be considered as a hallmark of a TSI. We optimized several experimental procedures designed to preserve and reveal dilute *para*-water in bulk.

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## Introduction

Although water is a fundamental constituent of our biosphere, characterization of some of its most basic properties remains a formidable challenge. Most applications of magnetic resonance imaging (MRI), both *in vitro* and *in vivo*, are based on observing

the magnetization of the two hydrogen nuclei of water. This is actually limited to the observation of *ortho*-water. If the populations of the three symmetrical states deviate from Boltzmann equilibrium, they rapidly recover through longitudinal spin-lattice relaxation with a time constant  $T_1$ , which is typically on the order of 10 s in neat water. So far, neither NMR nor MRI has been able to exploit the properties of the invisible singlet state  $S_0$ , also known as *para*-water, by analogy to *para*-hydrogen.<sup>1,2</sup> The following linear combination of populations:

$$\frac{1}{3} [|T_{+1}\rangle\langle T_{+1}| + |T_0\rangle\langle T_0| + |T_{-1}\rangle\langle T_{-1}|] - |S_0\rangle\langle S_0| \quad (1)$$

where:

$$|T_{+1}\rangle = |\alpha\alpha\rangle \quad |T_0\rangle = \frac{1}{\sqrt{2}} [|\alpha\beta\rangle + |\beta\alpha\rangle]$$

$$|T_{-1}\rangle = |\beta\beta\rangle \quad |S_0\rangle = \frac{1}{\sqrt{2}} [|\alpha\beta\rangle - |\beta\alpha\rangle]$$

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is equivalent to a “Triplet–Singlet Imbalance” (henceforth TSI), in analogy to the expression coined by Meier *et al.*<sup>3</sup> for the “A/E imbalance” in  $^{13}\text{CH}_3$  groups, which refers to a population imbalance between symmetric and antisymmetric states belonging to the irreducible representations A and E of methyl groups. An A/E imbalance can be induced by cooling down to *ca.* 1 K molecules such as  $\gamma$ -picoline that contain methyl groups characterized by very low rotational barriers.<sup>3–8</sup> In the high-temperature approximation, *i.e.*, at spin temperatures above a few degree Kelvin, the distribution between *para*- and *ortho*-water is given by the ratio 1:3. If this ratio is perturbed, as we shall demonstrate in this paper, the resulting TSI in eqn (1) should have a lifetime that might be much longer than the spin–lattice relaxation time ( $T_{\text{TSI}} > T_1$ ). This is analogous to molecules that contain two magnetically inequivalent protons that can sustain a Long-Lived State (LLS).<sup>9</sup> Such an LLS can preserve spin order over periods that are much longer than  $T_1$ . In  $^{12}\text{CH}_2$  groups, for instance, we have shown that one can have lifetimes with ratios  $T_{\text{TSI}}/T_1 > 36$ .<sup>10</sup> The LLS can be prepared by several NMR techniques at room temperature<sup>10,11</sup> or by DNP at low temperature.<sup>12</sup> Optimization of nearly-symmetric molecules comprising pairs of  $^{13}\text{C}$  nuclei has led to very long lifetimes in solution that can exceed one hour at room temperature.<sup>13</sup> Another intriguing feature of the LLS is that they can be used to improve the sensitivity of drug screening experiments.<sup>14–16</sup> In the past, several studies have focused on the isolation of *para*-water and the characterization of its long-lived behavior.<sup>17</sup> In crystal water trapped in gypsum ( $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ ), Pake found evidence of isolated pairs of protons.<sup>18</sup> At  $T < 100$  K, the water molecules trapped in gypsum crystals cannot flip around their two-fold symmetry axis. As a result, the two protons may be at unequal distances from other protons belonging to remote hydration water molecules. Thus the two protons may experience different intermolecular dipolar couplings, and their magnetic equivalence can be lifted. It is therefore possible to populate the antisymmetric state. Eisendrath, Stone and Jeener<sup>19,20</sup> characterized *para*-water in solid gypsum. More recently, the separation of *ortho*- and *para*-water has been achieved in molecular beams travelling through inhomogeneous magnetic fields,<sup>21</sup> where a beam of *ortho*-water can be deflected and refocused in the manner of the Stern–Gerlach experiment, or in inhomogeneous electric fields.<sup>22</sup> These methods can produce a large TSI but only for very small quantities of water. Isomer enrichment of  $\text{H}_2\text{O}$  in bulk, by means of absorption in column chromatography, has been proposed<sup>23</sup> and later challenged.<sup>24</sup> In other studies, the interconversion between *para*- and *ortho*-isomers was achieved by isolating water molecules in frozen inert gases<sup>25–27</sup> or by trapping them in  $\text{C}_{60}$  cages.<sup>28</sup> In the latter case, the conversion has been monitored by a combination of infrared spectroscopy, inelastic neutron scattering and cryo-MAS NMR spectroscopy.<sup>29</sup> This allows one to study *ortho*–*para* conversion rates<sup>30</sup> of isolated water molecules. However, the confinement in  $\text{C}_{60}$  cages prevents one from monitoring interactions with surrounding molecules. In this work we shall discuss the possibility of preparing samples of non-confined water characterized by a significant TSI at concentrations on the order of a few mM. Our approach is

similar to our strategy for preparing hyperpolarized *para*-ethanol<sup>31</sup> and *para*-fumarate.<sup>32</sup>

## Methodology

Our approach involves three consecutive steps.

### Step 1 – preparing a TSI

A flow of populations between the triplet and singlet energy levels in water cannot be induced by intramolecular dipole–dipole (DD) interactions between the two protons belonging to the same water molecule, although it may be induced by intermolecular dipole–dipole (DD) interactions<sup>19,20</sup> or by the proton chemical shift anisotropy (CSA) in the condensed phase. In liquid  $\text{H}_2\text{O}$  diluted in  $\text{D}_2\text{O}$  at 300 K, the CSA has been determined to be 28 ppm.<sup>33</sup> In gypsum, the proton CSA of the (hopping) water molecules has been determined to be 10 ppm at 300 K.<sup>34</sup> The proton CSA of the hopping water molecules in  $\text{Ba}(\text{ClO}_3)_2 \cdot \text{H}_2\text{O}$  has also been determined to be 10 ppm at 300 K.<sup>35</sup> In glassy frozen solutions, the symmetry of the two protons in each  $\text{H}_2\text{O}$  molecule is broken by the anisotropy of the chemical shifts, except for some particular orientations, so that the singlet state  $|S_0\rangle$  is mixed with the central triplet state  $|T_{+1}\rangle$ . Depending on the coupling between the rotational and Zeeman energy levels, two situations can occur. If the energy levels are primarily determined by the rotational quantum numbers, as in the gas phase<sup>36</sup> or  $\text{C}_{60}$  cages (where the rotational levels typical for the gas phase remain a good approximation), the singlet state  $S_0$  has the lowest energy. On the other hand, if rotational quantization can be neglected, the ground state is  $|\alpha\alpha\rangle = |T_{+1}\rangle$  which belongs to the triplet manifold. In our samples, the rotation of water is believed to be hindered by hydrogen bonding with other water and/or solvent molecules. Thus, when  $\text{H}_2\text{O}$  molecules are diluted in a deuterated aprotic solvent, doped with a radical and frozen at low temperatures in a high magnetic field, DNP can be used to achieve a TSI. The ESR transitions of the radical can be saturated by microwave irradiation to populate mostly the  $|\alpha\alpha\rangle = |T_{+1}\rangle$  state (Fig. 1A). During dissolution, the magnetic equivalence of the two protons in each  $\text{H}_2\text{O}$  molecule is restored. Hence, our strategy should lead to an excess of the average population of the three triplet states compared to the population of the singlet state (Fig. 1B). This amounts to a TSI. If its life-time is longer than  $T_1$ , this may be considered as the hallmark of *para*-water.

### Step 2 – protecting the TSI during transfer

After rapid heating of the sample by injection of a hot aprotic solvent, the sample can be transferred<sup>37</sup> to an NMR or MRI system. Longitudinal  $T_1$  relaxation in a few seconds leads to the return of the triplet manifold to Boltzmann equilibrium at room temperature (Fig. 1C). However,  $T_1$  relaxation does not affect the TSI, so that the singlet state remains depleted. On a longer time-scale  $T_{\text{TSI}}$ , the populations of the triplet and singlet states will return to their Boltzmann equilibrium (Fig. 1D). In the dissolution step, all relaxation mechanisms that could



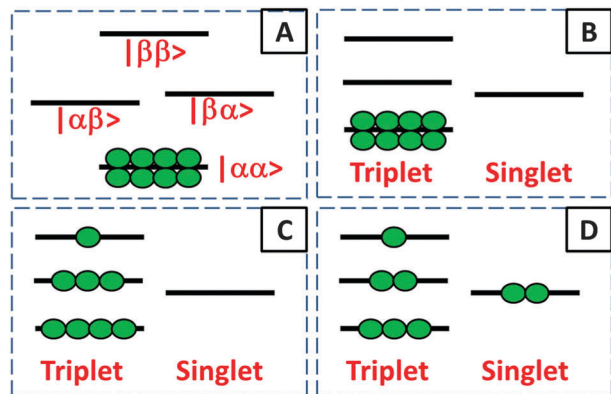


Fig. 1 (A) Schematic representation of the energy levels of the two magnetically inequivalent protons of  $\text{H}_2\text{O}$  in the solid state at 1.2 K. By lowering the spin temperature to a few mK by DNP, only the ground state  $|\alpha\alpha\rangle = |T_{+1}\rangle$  is populated. (B) After dissolution with a hot aprotic solvent, the magnetic equivalence is restored. (C) Relaxation in the triplet manifold occurs on a time-scale  $T_1$ . (D) Equilibration of the T/S imbalance occurs with a time constant  $T_{\text{TSI}}$  that is believed to be much longer than  $T_1$ .

reduce  $T_{\text{TSI}}$  must be kept under control. The radicals can be eliminated by chemical reduction with scavengers such as ascorbate.<sup>38</sup> Alternatively, porous solids containing covalently bound radicals can be filtered after dissolution.<sup>39</sup> Spin rotation (SR) is another mechanism that could cause losses of the TSI. SR is due to the coupling between the nuclear spins and the molecular magnetic dipole induced by the electric dipole moment of the  $\text{H}_2\text{O}$  molecule as it undergoes rotational diffusion. The SR mechanism of *ortho-para* conversion has been intensively studied.<sup>40,41</sup> Indeed, in the gas phase,<sup>42,43</sup> the longitudinal relaxation times were found to be on the order of  $T_1 = 20$  ms near 0.1 MPa and 373 K at 800 MHz. We believe however that SR is not an efficient mechanism for  $\text{H}_2\text{O}$  in the condensed phase where collisions on the atomic scale should make SR ineffective. This should also be the case when the rotation of a  $\text{H}_2\text{O}$  molecule is hindered because it is trapped in a cage of an aprotic solvent. Finally, one should pay attention to the exchange of protons between different water molecules. In other words, the lifetime  $\tau_{\text{EX}} = 1/k_{\text{EX}}$  of a water molecule as a molecular entity has to be longer than the lifetime of the TSI. Since the latter lifetime is unknown, the best option is to slow down the proton exchange as much as possible. In pure water the lifetime of proton exchange has been assessed to be  $\tau_{\text{EX}} \approx 1$  ms by studying either linewidths in proton spectra<sup>44</sup> or intensities of  $^{17}\text{O}$  lines with and without proton decoupling.<sup>45,46</sup> However, it can be shown by EXSY<sup>47</sup> that dilution in aprotic solvents can effectively slow down the exchange of protons and thus extend  $\tau_{\text{EX}}$ . This approach should help to preserve the TSI and thus the lifetime of *para*-water.

### Step 3 – detecting TSI relaxation

To detect the relaxation of the TSI, at least four distinct approaches can be used. (i) Direct detection by infrared (IR) spectroscopy relies on the fact that *ortho*- and *para*-water give rise to distinct IR absorption bands in the gas phase.<sup>29,36</sup>

However, in the condensed phase, in particular, when water is diluted in aprotic solvents, we found that the IR signatures are difficult to identify. (ii) Direct detection by multiplet effects in oxygen-17 NMR. In thermal equilibrium, the oxygen-17 signal of  $\text{H}_2\text{O}$  shows a normal binomial 1:2:1 triplet. If, however, one is able to populate a TSI, non-binomial multiplets could be observed like in *para*-ethanol.<sup>31</sup> (iii) If the proton signals relax back to equilibrium with two different time constants this would be a hallmark of *ortho-para* conversion. (iv) Long-lived water can be revealed indirectly by chemical addition onto a suitable substrate such as an activated aldehyde or ketone, monitored by  $^1\text{H}$  or  $^{13}\text{C}$  NMR. For this strategy to be successful, the reactivity of water molecules has to fulfill three requirements that are partly contradictory: (a) the water must be sufficiently diluted so that intermolecular  $^1\text{H}$  exchange is slowed down; (b) the rate of the chemical addition that is used to reveal the presence of *para*-water must be faster than the relaxation of the TSI; and (c) the two protons that are added onto the substrate must stem from the same water molecule. Our kinetic measurements show that, under suitable conditions, these requirements may indeed be satisfied.

## Results and discussion

### Step 1 – preparing a TSI

Unlike Jeener and coworkers<sup>19,20</sup> who postulated that the two protons of a  $\text{H}_2\text{O}$  molecule can have different environments because of intermolecular dipole-dipole interactions in a magnetic field of 0.7 T, we shall assume that the proton chemical shift anisotropy (CSA) makes significant contributions to the breaking of the symmetry at 6.7 T. We diluted water in deuterated dimethylsulfoxide ( $\text{DMSO-d}_6$ ), doped it with *ca.* 50 mM

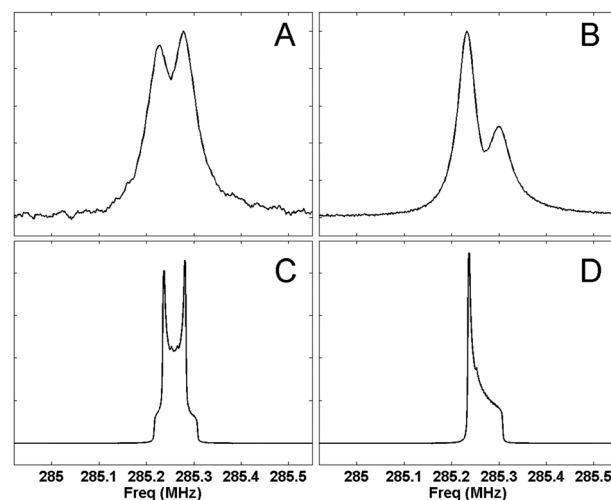


Fig. 2 Experimental proton spectra of  $\text{H}_2\text{O}$  diluted in  $\text{DMSO}$  doped with 50 mM TEMPOL and frozen at *ca.* 1.2 K in a field of 6.7 T, with (A) and without (B) irradiation with a resonant microwave field at 187.9 GHz. Simulations by using SIMPSON<sup>48</sup> (see text for details) for initial density operators  $\rho = I_z + S_z$  (C) or  $\rho = (1/2)[1/2I_z + I_z + S_z + 2I_zS_z]$  (D). The experimental spectra A and B have larger linewidths than the simulated C and D because of intermolecular dipolar couplings and paramagnetic species in the frozen glass.



TEMPOL radicals and froze it to form a glassy state at *ca.* 1.2 K and 6.7 T. In Fig. 2 we can see proton spectra before (A) and after (B) saturation of the ESR transitions by microwave irradiation at 187.9 GHz with frequency modulation.<sup>50</sup> Simulations of H<sub>2</sub>O powder spectra have been carried out by using the SIMPSON program<sup>48</sup> on a spin system made up of two protons with shielding anisotropy  $\Delta_{\text{CS}} = -16.19$  ppm, asymmetry  $\eta_{\text{CS}} = 0.17$  and a dipolar coupling  $d = -30.314$  kHz. 4180 crystallite orientations were considered. The relative orientations of the relevant shielding tensors, expressed in a common crystal frame, were given by the Euler angles  $\Omega^{\text{H}(1)} = \{-165.11^\circ, 115.33^\circ, 36.75^\circ\}$  and  $\Omega^{\text{H}(2)} = \{-14.89^\circ, 115.33^\circ, -36.75^\circ\}$ , with the dipolar coupling tensor between the two protons oriented according to  $\Omega^{\text{H}(1,2)} = \{180.00^\circ, 58.83^\circ, 0.00^\circ\}$ . These parameters have been calculated in previous work on water molecules in barium chlorate monohydrate<sup>35</sup> by means of DFT and planewave-pseudopotential methods as implemented in the CASTEP code.<sup>51</sup> The shielding ellipsoids have their main components aligned along the OH bonds. A realistic pulse was used, with an rf-field strength  $\nu_1 = 50$  kHz and nutation angles  $\beta = 90^\circ$  and  $1.8^\circ$ , in Fig. 2C and D, respectively. The initial density operator was either  $\rho(0) = I_z + S_z$  to describe the high-temperature approximation

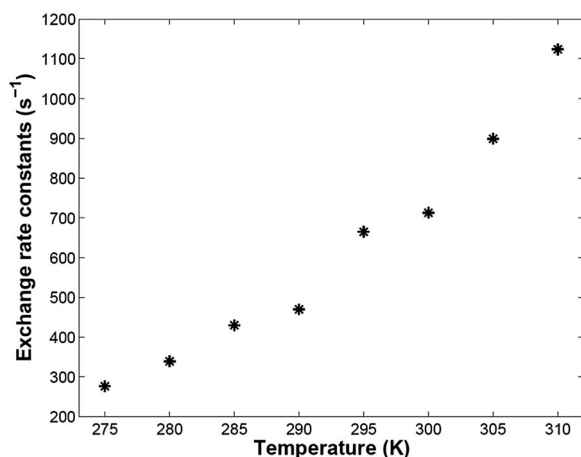


Fig. 3 Measurement of fast proton exchange rates  $k_{\text{EX}}$  in pure water. Proton exchange rates as a function of temperature at pH = 5.93 were extracted from ratios of peak heights of oxygen-17 echoes observed without and with proton decoupling.<sup>45,46</sup> The lifetimes  $\tau_{\text{EX}} = 1/k_{\text{EX}}$  range from 3.6 to 0.9 ms between 275 and 310 K.

(Fig. 2C) or  $\rho(0) = (1/2)[1/2E + I_z + S_z + 2I_zS_z]$  which corresponds to a pure state where only the lowest-lying state is populated at very low spin temperatures (Fig. 2D). The experimental spectra in Fig. 2A and B have larger linewidths than the simulated spectra in Fig. 2C and D: this can be ascribed to the presence of paramagnetic species in solution (TEMPOL) and to intermolecular dipolar couplings that were not considered in the simulations. Finally this evidence suggests that DNP indeed allows one to drive the populations towards the lowest-lying state: since there is some mixing between the central triplet state and the singlet state, a very low spin temperature (on the order of 10 mK in our experiments) is sufficient to generate a TSI.

## Step 2 – protecting the TSI during transfer

Proton exchange in water could represent the major source of losses of TSI in our experiment. Hence, we studied the exchange in water in bulk and dilute solutions. The combined effects of proton exchange and proton  $T_1$  relaxation on the transverse  $T_2$  relaxation of  $^{17}\text{O}$  nuclei in H<sub>2</sub><sup>17</sup>O can be characterized by using multiple refocusing of transverse  $^{17}\text{O}$  magnetization in the manner of Carr, Purcell, Meiboom and Gill (CPMG). One may compare  $^{17}\text{O}$  echo decays in the presence or absence of proton decoupling, in analogy with a similar work carried out on  $^{15}\text{N}$ .<sup>45,46</sup> Our observations shown in Fig. 3 are in agreement with pioneering studies by Meiboom.<sup>44</sup> We measured lifetimes  $\tau_{\text{EX}} = 1/k_{\text{EX}}$  of a few milliseconds in pure water at pH 5.93 and different temperatures. The lifetime of pure water as a molecular entity is clearly too short for our purposes but, as mentioned above, dilution in aprotic solvents can be used to extend this lifetime. In dilute solutions, the proton spectra of mixtures of H<sub>2</sub>O and HDO feature two distinct resonances. In order to observe HDO triplets due to  $^1J_{\text{HD}}(^1\text{H}, ^{17}\text{O}) = 80$  Hz, the lifetime of HDO has to be  $\tau_{\text{EX}} > 1/^1J_{\text{HD}}(^1\text{H}, ^{17}\text{O}) = 12.5$  ms. Slow exchange rates can be quantified using selective 1D or 2D exchange spectroscopy (EXSY).<sup>47</sup> The pulse sequence used is shown in Fig. 5. Our samples consisted of a mixture of H<sub>2</sub>O, HDO and D<sub>2</sub>O diluted in various organic solvents at different concentrations (see Table 1 and Fig. 4). The concentrations  $[\text{H}_2\text{O}]$  and  $[\text{HDO}]$  have been determined by NMR within  $\pm 10\%$ , by scaling their peak intensities to an external reference with a concentration that is known *a priori*. In nitromethane and dioxane,  $[\text{H}_2\text{O}]$  and  $[\text{HDO}]$  were increased with respect to other solvents since, at low concentrations, the proton exchange was too slow to be monitored *via* 1D-EXSY.

**Table 1** Proton exchange rates for mixtures of H<sub>2</sub>O + HDO diluted in aprotic deuterated organic solvents at 800 MHz and 300 K. The concentrations, protonation fractions  $\alpha = [\text{H}]/([\text{H}] + [\text{D}])$  and chemical shifts  $\nu$  of H<sub>2</sub>O and HDO in the liquid phase are reported. The parameters  $\alpha$ ,  $R_{\text{eff}}$  and  $k_{\text{EX}}$  were estimated by global fitting of the four curves in Fig. 5. In order to compare exchange lifetimes, normalized  $\tau_{\text{EX}}$  have been calculated for  $[\text{H}_2\text{O}] + [\text{HDO}] = 50$  mM, assuming a linear dependence of the exchange rate on the concentration<sup>49</sup>

Solvent	Experiments@800 MHz					Fitting@800 MHz			$\tau_{\text{EX}}$ (s) normalized to $[\text{H}_2\text{O}] + [\text{HDO}] = 50$ mM
	$\alpha$	$[\text{H}_2\text{O}]$ (mM)	$\nu(\text{H}_2\text{O})$ (ppm)	$[\text{HDO}]$ (mM)	$\nu(\text{HDO})$ (ppm)	$\alpha$	$R_{\text{eff}}$ (s)	$k_{\text{EX}}$ (s <sup>-1</sup> )	
Dioxane-d <sub>8</sub>	0.28	105	~2.71	169	~2.67	0.30	0.19	$0.14 \pm 0.03$	20
Nitromethane-d <sub>3</sub>	0.26	36	~2.13	50	~2.10	0.40	0.08	$0.30 \pm 0.03$	3
Acetonitrile-d <sub>3</sub>	0.19	5	~2.18	4	~2.15	0.35	0.15	$0.23 \pm 0.02$	0.4
DMSO-d <sub>6</sub>	0.06	40	~3.36	11	~3.33	0.10	0.55	$1.7 \pm 0.2$	0.06
Acetone-d <sub>6</sub>	0.22	12	~2.88	14	~2.85	0.30	0.17	$4.0 \pm 0.5$	0.04
Dichloromethane-d <sub>2</sub>	0.73	2	~1.63	21	~1.60	0.77	0.15	$4.6 \pm 0.4$	0.001





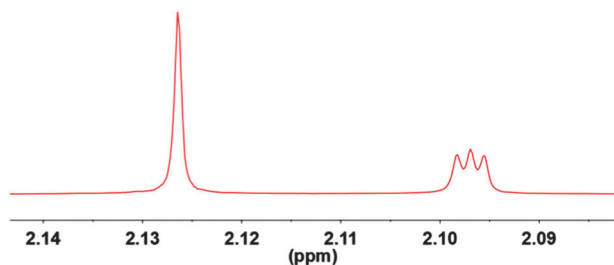


Fig. 4 Proton spectrum of 36 mM H<sub>2</sub>O (singlet on left-hand side) and 50 mM HDO (triplet due to  $J(\text{H},\text{D})$  on the right-hand side) in nitromethane- $\text{d}_3$  at 300 K and 800 MHz.

The build-up and decay curves shown in Fig. 5 can be fitted to the following functions (see the Appendix for details):

$$D_{\text{H}_2\text{O}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} [\cosh(k_{\text{EX}}t) + (1 - 2\alpha) \sinh(k_{\text{EX}}t)]$$

$$C_{\text{H}_2\text{O}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} (2 - 2\alpha) \sinh(k_{\text{EX}}t)$$

$$D_{\text{HDO}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} [\cosh(k_{\text{EX}}t) + (2\alpha - 1) \sinh(k_{\text{EX}}t)]$$

$$C_{\text{HDO}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} 2\alpha \sinh(k_{\text{EX}}t)$$

The parameters  $\alpha$ ,  $R_{\text{eff}}$  and  $k_{\text{EX}}$  have been determined by global fitting to the curves in Fig. 5 and are shown in Table 1.

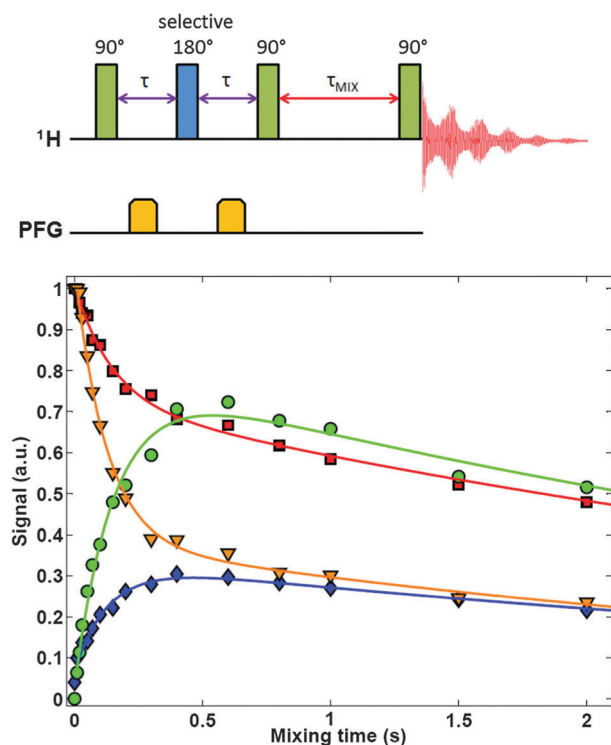


Fig. 5 [top] Pulse sequence for selective 1D-EXSY experiments. [bottom] Typical build-up and decay curves measured using 1D-EXSY experiments used for the measurement of the lifetime of water as a molecular entity, when diluted in aprotic solvents (acetonitrile- $\text{d}_3$  in this example). Red squares and orange triangles represent intensities of diagonal peaks of H<sub>2</sub>O and HDO, respectively; green circles describe magnetization transfer from HDO to H<sub>2</sub>O, while blue rhombi describe the reverse reaction. The rates measured for different solvents are reported in Table 1.

Table 2 Longitudinal relaxation rates  $R_1$  of H<sub>2</sub>O diluted in aprotic deuterated organic solvents at 600 MHz and 300 K. The rates  $R_1$  were determined by inversion recovery. Their values can be compared to  $R_{\text{eff}}$  in Table 1 but only in a qualitative way since  $R_{\text{eff}}$  depends on [H<sub>2</sub>O], [HDO] and  $k_{\text{EX}}$  and for different experimental conditions

Solvent	Experiments@600 MHz	
	$R_1^{\text{H}_2\text{O}}$ (s)	[H <sub>2</sub> O] (mM)
Dioxane- $\text{d}_8$	0.26	358
Nitromethane- $\text{d}_3$	0.08	91
Acetonitrile- $\text{d}_3$	0.15	9
DMSO- $\text{d}_6$	0.30	42
Acetone- $\text{d}_6$	0.17	10
Dichloromethane- $\text{d}_2$	0.11	6

The rates  $R_1 = 1/T_1$  of H<sub>2</sub>O have been determined experimentally by inversion recovery (see Table 2).  $R_{\text{eff}}$  is an average of  $R_1(\text{H}_2\text{O})$  and  $R_1(\text{HDO})$  weighted by their concentrations. Hence, a comparison between  $R_{\text{eff}}$  in Table 1 and  $R_1$  in Table 2 can only be qualitative because of different experimental conditions. However, in most cases the values are similar. In order to facilitate comparisons, in Table 1 we estimated the lifetimes for a concentration [H<sub>2</sub>O] + [HDO] = 50 mM, assuming that the rates vary linearly with concentration.<sup>49</sup> It is evident that, at concentrations below 50 mM, dioxane allows one to extend the lifetime of water as a molecular entity up to a few minutes. Dioxane, therefore, seems a good solvent for dissolution DNP. However, it has a much higher viscosity and lower heat capacity than water, so that our dissolution apparatus has to be re-designed thoroughly.

### Step 3 – detecting TSI relaxation

To observe a TSI in H<sub>2</sub>O at room temperature after dissolution, we have resorted to a chemical reaction, inspired by the PASADENA<sup>1</sup> and ALTADENA<sup>52</sup> methods. We have shown recently<sup>32</sup> that one can lift the degeneracy of the two protons in fumarate ( $^-\text{OOCCH}=\text{CHCOO}^-$ ) by addition of D<sub>2</sub>O to produce malate ( $^-\text{OOCCHDCHODCOO}^-$ ), a reaction that is catalyzed by fumarase. In our work on *para*-ethanol,<sup>31</sup> instead, the detection was possible, without any chemical reactions, by monitoring non-binomial multiplets in an AX<sub>2</sub> system. Following a suggestion by Jean-Maurice Mallet we have explored the addition of water (though not yet of *para*-water) on aldehydes, *i.e.*,  $\text{RCHO} + \text{H}_2\text{O} \rightarrow \text{RCH}(\text{OH})_2$ . The C=O double bond of the aldehyde can be activated by substituents such as R = CCl<sub>3</sub>, as in chloral (CCl<sub>3</sub>CHO, see Fig. 6). When the reaction is carried out in a dilute solution in acetonitrile, the two water protons (highlighted by stars in Fig. 6) may be assumed to end up on the same hydrate molecule. The two OH protons that stem from

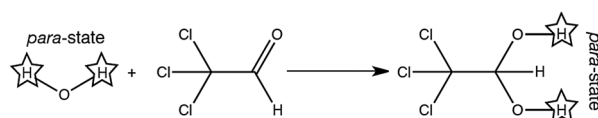


Fig. 6 The addition of H<sub>2</sub>O to chloral gives chloral hydrate CCl<sub>3</sub>CH(OH)<sub>2</sub>, where the two protons highlighted by stars can be assumed to stem from one and the same water molecule if the solution is sufficiently dilute.

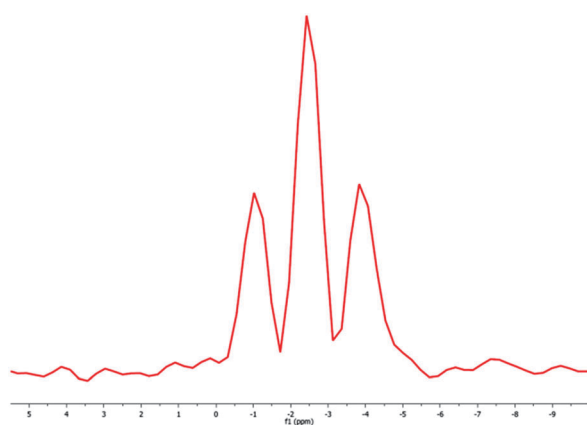


*para*-water give rise to degenerate doublets in the vicinity of 6.4 ppm. The lone proton near 5.3 ppm that stems from the aldehyde gives rise to a triplet due to two equal long-range couplings  $^3J_{\text{HH}} \approx 6$  Hz to the two OH protons. Clearly, as discussed above, deviations from the binomial 1 : 2 : 1 distribution of this triplet can be used for a quantitative determination of the TSI.<sup>31</sup>

The relative intensities of the lines in these multiplets can give a measure of the TSI in H<sub>2</sub>O, *i.e.*, of the relative populations of *ortho*- and *para*-water, much as in the <sup>17</sup>O spectrum of H<sub>2</sub><sup>17</sup>O, or in the <sup>13</sup>C spectrum of the <sup>13</sup>CH<sub>2</sub> group in partially deuterated ethanol CD<sub>3</sub><sup>13</sup>CH<sub>2</sub>OD. We therefore explored the possibility of using reactions like in Fig. 6 to observe the binomial distribution of the triplet of the CH<sub>2</sub> protons. We explored the kinetics of the addition of H<sub>2</sub>O onto two different substrates: chloral diluted in acetonitrile and 1,3-dichlorotetrafluoroacetone diluted in dioxane. The main requirement is that the reaction must be faster than the relaxation of the TSI and its decay due to proton exchange. The rate constants found using pseudo-first-order kinetic equations are reported in Table 3, albeit without DNP and thus without TSI. In acetonitrile-d<sub>3</sub>, with a 5-fold excess of chloral with respect to H<sub>2</sub>O, a pseudo first-order rate constant  $k_{\text{pfo}} = 0.002$  s<sup>-1</sup> was observed. This reaction is too slow to be useful for detecting the TSI characteristic of *para*-water. However, the addition of water onto the more reactive compound (1,3-dichlorotetrafluoroacetone, in dioxane-d<sub>8</sub>) was so fast that

**Table 3** Kinetic pseudo-first-order rate constant  $k_{\text{pfo}}$  for hydration of activated C=O bonds observed at 800 MHz and 300 K

Reactant	Solvent	[Reactant] (mM)	[H <sub>2</sub> O] (mM)	$k_{\text{pfo}}$ (s <sup>-1</sup> )
Chloral	Acetonitrile-d <sub>3</sub>	1000	200	0.002
1,3-Dichlorotetrafluoroacetone	Dioxane-d <sub>8</sub>	275	55	> 0.1



**Fig. 7** Experimental <sup>17</sup>O spectrum of 20% enriched water, 55 mM in dioxane-d<sub>8</sub>, acquired on a 400 MHz NMR spectrometer. The binomial 1 : 2 : 1 distribution of the triplet due to  $J(^1\text{H}, ^{17}\text{O}) \sim 80$  Hz is characteristic of an equilibrium Boltzmann distribution. Deviations from this binomial multiplet are expected if the TSI does not vanish, and could be a hallmark of *para*-water. Isotopic enrichment of 20% <sup>17</sup>O made it possible to use an acquisition time of a few seconds, making this method suitable for detection of a TSI.

an accurate determination of the rate was difficult by NMR. A reaction rate  $k_{\text{pfo}} > 0.1$  s<sup>-1</sup> makes it a good candidate as a “revealing agent” of TSI in *para*-water. As an alternative method for detection, one can simply measure the <sup>17</sup>O spectrum in 0.037% natural abundance or with partial isotopic enrichment to improve sensitivity. The <sup>17</sup>O spectrum in Fig. 7 shows a triplet due to  $J(^1\text{H}, ^{17}\text{O}) = 80$  Hz of 55 mM H<sub>2</sub>O in dioxane-d<sub>8</sub> at 298 K and 400 MHz. Using water enriched to 20% <sup>17</sup>O, we were able to acquire an <sup>17</sup>O spectrum in a few seconds, making this method a valid alternative to the use of a chemical reaction. Again, deviations from the binomial distribution of the intensities of the spectral lines should provide the information needed to assess the lifetime of the TSI and hence of *para*-water.

We have optimized the most critical aspects of the experimental scheme suggested in this paper. However, we were not yet able to perform complete experiments since our setup needs to be significantly adapted in order to support dissolution with dioxane.

## Conclusions

We have proposed an experimental strategy to produce *para*-water on a macroscopic scale (*i.e.* with concentrations in the mM range). Provided that the CSA of the protons is sufficient to lift the degeneracy of the two spins in a frozen sample, we demonstrated that one can use DNP to enhance the population of the ground state of water molecules, thus generating a Triplet-Singlet Imbalance (TSI) that is expected to be a long-lived state analogous to *para*-water. We proved that the lifetime of water as a molecular entity can be extended up to a few minutes by dilution in aprotic solvents. Several detection strategies can be used either by performing a “revealing” reaction such as the addition of water to an aldehyde or another suitable substrate or by observing the <sup>17</sup>O NMR spectrum of water itself. It is also possible to monitor proton magnetization of water relaxing to equilibrium with two clearly distinct time constants to assess the lifetime of the TSI and hence the amount of *para*-water in the sample. Similar information can, in principle, be obtained by infrared spectroscopy. The detection of the *para*-water signal with long lifetimes may open the way to study slow transport phenomena such as flow, diffusion, and electrophoretic mobility.

## Appendix

Let us consider a solution of pure H<sub>2</sub>O. Let  $p$  be the frequency at which a proton is exchanged with another proton belonging to a different water molecule:

$$p \propto [\text{H}_2\text{O}]$$

Hence, the proton exchange rate  $k_{\text{EX}}$  can be defined as:

$$k_{\text{EX}} = 2p$$

The factor 2 reflects the fact that a proton belonging to a H<sub>2</sub>O molecule can exchange with either of the two protons of another H<sub>2</sub>O molecule.



Let us now consider a mixture of H<sub>2</sub>O, HDO and D<sub>2</sub>O. With the ratio  $\alpha = [\text{H}]/([\text{H}] + [\text{D}])$ , their concentrations can be expressed as:

$$[\text{H}_2\text{O}] = \alpha^2 C_{\text{TOT}}$$

$$[\text{D}_2\text{O}] = (1 - \alpha)^2 C_{\text{TOT}}$$

$$[\text{HDO}] = 2\alpha(1 - \alpha)C_{\text{TOT}}$$

where:

$$C_{\text{TOT}} = [\text{H}_2\text{O}] + [\text{HDO}] + [\text{D}_2\text{O}]$$

The equations for the exchange of proton magnetization between two distinct sites can be written in the general form:

$$\frac{d\bar{M}(t)}{dt} = \bar{\bar{K}}\bar{M}(t)$$

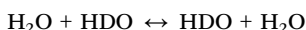
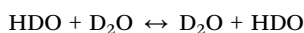
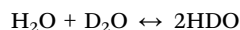
where:

$$\bar{M}(t) = \begin{pmatrix} M_{\text{H}_2\text{O}}(t) \\ M_{\text{HDO}}(t) \end{pmatrix}$$

$$\bar{\bar{K}} = \begin{pmatrix} -k & k' \\ k & -k' \end{pmatrix}$$

$$k' = k \frac{M_{\text{H}_2\text{O}}|_{\text{EQ}}}{M_{\text{HDO}}|_{\text{EQ}}} = k \frac{2[\text{H}_2\text{O}]}{[\text{HDO}]} = k \frac{2(1 - \alpha)^2}{2\alpha(1 - \alpha)} = k \frac{(1 - \alpha)}{\alpha}$$

Let us consider the following reactions involving the exchange of a proton:



If we focus attention on the magnetization transfer from H<sub>2</sub>O to HDO, there are three possibilities:

(1) H<sub>2</sub>O meets another H<sub>2</sub>O. The exchange of two protons does not lead to any transfer of magnetization between different environments with distinct chemical shifts.

(2) H<sub>2</sub>O meets D<sub>2</sub>O. There are four possible exchange processes. Each process leads to the creation of two HDO molecules, and there are two protons that transfer their magnetization between different environments:

$$k_2 = 8p[\text{D}_2\text{O}] = 8p\alpha^2$$

(3) H<sub>2</sub>O meets HDO. Again, four possible exchange processes can take place, but there is only one proton that transfers its magnetization between different environments:

$$k_3 = 4p[\text{HDO}] = 8p\alpha(1 - \alpha)$$

If we consider that we should count an exchange process not only for the proton that hops but also for its neighbor, the total

exchange rate is:

$$k = \frac{k_2 + k_3}{2} = 4\alpha p = 2\alpha k_{\text{EX}}$$

If we now consider magnetization transfer from HDO to H<sub>2</sub>O, there are again three possibilities:

(4) HDO meets H<sub>2</sub>O. This is symmetric to case (3) above:

$$k_1 = 4p[\text{H}_2\text{O}] = 4p(1 - \alpha)^2$$

(5) HDO meets D<sub>2</sub>O. Four possible exchange processes can again occur but none of them leads to any magnetization transfer between different environments.

(6) HDO meets HDO. Again, four possible exchange processes are possible, two of which swap a proton with another proton but do not lead to any magnetization transfer. Two processes swap a proton and a deuteron to create two H<sub>2</sub>O molecules, which is accompanied by a transfer of magnetization between different environments:

$$k_3 = 4p[\text{HDO}] = 8p\alpha(1 - \alpha)$$

In this case we have counted the exchange processes twice: when a molecule i meets a molecule j the effect is of course the same as the case when j meets i. Hence the total rate is:

$$k' = k_1 + \frac{k_3}{2} = 4p(1 - \alpha) = 2(1 - \alpha)k_{\text{EX}}$$

By including longitudinal relaxation, we find:

$$\bar{\bar{K}} = - \begin{bmatrix} R_1^{\text{H}_2\text{O}} + 2\alpha k_{\text{EX}} & -(2 - 2\alpha)k_{\text{EX}} \\ -2\alpha k_{\text{EX}} & R_1^{\text{HDO}} + (2 - 2\alpha)k_{\text{EX}} \end{bmatrix}$$

This matrix can be diagonalized as:

$$\bar{M}(t) = \bar{U}^{-1} e^{-\bar{\bar{D}}t} \bar{U} \bar{M}(0)$$

$$\bar{\bar{D}} = \bar{U} \bar{\bar{K}} \bar{U}^{-1}$$

so that

$$\bar{M}(t) = \begin{bmatrix} D_{\text{H}_2\text{O}} & C_{\text{HDO}} \\ C_{\text{H}_2\text{O}} & D_{\text{HDO}} \end{bmatrix} \bar{M}(0)$$

With the assumption that  $R_1^{\text{H}_2\text{O}} = R_1^{\text{HDO}} = R_1^{\text{eff}}$ , we finally find:

$$D_{\text{H}_2\text{O}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} [\cosh(k_{\text{EX}}t) + (1 - 2\alpha) \sinh(k_{\text{EX}}t)]$$

$$C_{\text{H}_2\text{O}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} (2 - 2\alpha) \sinh(k_{\text{EX}}t)$$

$$D_{\text{HDO}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} [\cosh(k_{\text{EX}}t) + (2\alpha - 1) \sinh(k_{\text{EX}}t)]$$

$$C_{\text{HDO}} = e^{-(k_{\text{EX}} + R_1^{\text{eff}})t} 2\alpha \sinh(k_{\text{EX}}t)$$

In these calculations we have neglected possible kinetic isotope effects on the rates.



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