Gadolinium Effect at High-Magnetic-Field DNP: 70% 13C Polarization of [U-13C] Glucose Using Trityl

Capozzi, Andrea; Patel, Saket; Wenckebach, W. Thomas; Karlsson, Magnus; Lerche, Mathilde H.; Ardenkjær-Larsen, Jan Henrik

Published in:
Journal of Physical Chemistry Letters

Link to article, DOI:
10.1021/acs.jpcllett.9b01306

Publication date:
2019

Document Version
Peer reviewed version

Citation (APA):

General rights
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.
Gadolinium Effect at High Magnetic Field DNP: 70% $^{13}C$ Polarization of [U-$^{13}C$]glucose Using Trityl
Andrea Capozzi, Saket Patel, Tom Wenckebach, Magnus Karlsson, Mathilde Hauge Lerche, and Jan Henrik Ardenkjæer-Larsen

J. Phys. Chem. Lett., Just Accepted Manuscript • Publication Date (Web): 03 Jun 2019

Downloaded from http://pubs.acs.org on June 3, 2019

Just Accepted

"Just Accepted" manuscripts have been peer-reviewed and accepted for publication. They are posted online prior to technical editing, formatting for publication and author proofing. The American Chemical Society provides "Just Accepted" as a service to the research community to expedite the dissemination of scientific material as soon as possible after acceptance. "Just Accepted" manuscripts appear in full in PDF format accompanied by an HTML abstract. "Just Accepted" manuscripts have been fully peer reviewed, but should not be considered the official version of record. They are citable by the Digital Object Identifier (DOI®). "Just Accepted" is an optional service offered to authors. Therefore, the "Just Accepted" Web site may not include all articles that will be published in the journal. After a manuscript is technically edited and formatted, it will be removed from the "Just Accepted" Web site and published as an ASAP article. Note that technical editing may introduce minor changes to the manuscript text and/or graphics which could affect content, and all legal disclaimers and ethical guidelines that apply to the journal pertain. ACS cannot be held responsible for errors or consequences arising from the use of information contained in these "Just Accepted" manuscripts.
Gadolinium Effect at High Magnetic Field DNP:
70% $^{13}$C Polarization of [U-$^{13}$C] Glucose Using Trityl

Andrea Capozzi,$^{1,*}$ Saket Patel,$^1$ W. Thomas Wenckebach,$^{2,3}$ Magnus Karlsson,$^1$

Mathilde H. Lerche,$^1$ and Jan Henrik Ardenkjær-Larsen.$^{1,4}$

$^1$Center for Hyperpolarization in Magnetic Resonance, Department of Health Technology, Technical University of Denmark, Building 349, 2800 Kgs Lyngby (Denmark)

$^2$Paul Scherrer Institute, CH-5232 Villigen (Switzerland)

$^3$National High Magnetic Field Laboratory, Gainesville, Florida (USA).

$^4$GE Healthcare, Park Alle 295, 2605 Brøndby (Denmark)

Corresponding Author
*Andrea Capozzi, PhD

Technical University of Denmark

Ørsted Plads, Building 349, room 110

2800 Kgs. Lyngby

T: +45 45 25 53 27 ; Email: andcapo@dtu.dk

ABSTRACT

Herein, we show that the trityl Electron Spin Resonance (ESR) features, crucial for an efficient Dynamic Nuclear Polarization (DNP) process, are sample composition dependent. Working at 6.7 T and 1.1 K with a generally applicable DNP sample solvent mixture such as water:glycerol plus trityl, addition of Gd$^{3+}$ lead to a dramatic increase of [U-$^{13}$C]glucose polarization from 37±4% to 69±3%. This is the highest value reported to date and comparable to what can be achieved on pyruvic acid. Moreover, performing ESR measurements at actual DNP conditions, we provide experimental evidence that gadolinium doping not only shortens the trityl electron spin-lattice relaxation time but also
modifies the radical g-tensor. The latter yielded a considerable narrowing of the ESR spectrum linewidth. Finally, in the frame of the spin temperature theory, we argue on how and within which boundaries these two phenomena can improve the DNP performances.

TOC GRAPHICS

KEYWORDS: dDNP, gadolinium, glucose, hyperpolarization, LOD-ESR, trityl.

Hyperpolarization of nuclear spins via dissolution Dynamic Nuclear Polarization (dDNP)\textsuperscript{1} has proven its great potential in enhancing the sensitivity of a broad variety of molecules, in particular for detection with \textsuperscript{13}C Nuclear Magnetic Resonance (NMR) and Magnetic
Resonance Imaging (MRI).\textsuperscript{2-3} Prior to dissolution, the microwave driven polarization transfer from the electron spins, in form of free radicals, to the nuclear spins of interest takes place in the solid state at cryogenic temperature (1 – 1.5 K) and moderate magnetic field (3.35 – 7 T).\textsuperscript{4} Neat [1-$^{13}$C]pyruvic acid (PYR) doped with the trityl\textsuperscript{a} free radicals has become the most studied sample because of [1-$^{13}$C]pyruvate’s undisputed usefulness in biomedical applications,\textsuperscript{5-7} the really high achievable carbon polarization (up to 70%)\textsuperscript{8-10} and its relatively long relaxation time constant after dissolution.\textsuperscript{11} Detailed studies for different trityl-based samples are scarcer, limiting the potential of other biologically interesting substrates. A crucial example is the so-called “gadolinium effect”. At the broadly used dDNP field value (i.e. 3.35 T), admixture of small amounts of Gd\textsuperscript{3+} can double the polarization enhancement of neat [1-$^{13}$C]PYR/trityl samples, leading to 40% carbon polarization at optimal conditions.\textsuperscript{12} Differently, increasing the magnetic field strength (4.6 – 7 T), where the same sample can reach $^{13}$C polarization of 60 – 70 %,

\textsuperscript{a} Discussion of other polarizing agents such as BDPA and TEMPO is beyond the scope of the present manuscript. To avoid confusion, we specify that all data, observation and comments taken from the literature are limited to the case of trityl.
almost negligible improvement or worsening of the DNP enhancement factor has been observed by adding Gd$^{3+}$. At 3.35 T, trityl based non-PYR DNP samples showed $^{13}$C polarization gain as high as 4-folds, but those samples did not excessively benefit from increasing the magnetic field strength. At magnetic fields higher than 3.35 T, very little has been published on the effect of Gd$^{3+}$, or other lanthanides, on trityl based non-PYR DNP sample preparations. The behavior between different DNP sample preparations is often explained by changes in nuclear and electron relaxation parameters. In particular, although the precise physical mechanism behind the "gadolinium effect" is still not fully understood, shortening of the electron spin-lattice relaxation time ($T_{1e}$) is the most acknowledged process to contribute to the nuclear polarization increase. Often, because of hardware limitations, ESR measurements tailored at understanding the process behind the "gadolinium effect" are performed at field strengths and/or temperatures deviating from the targeted DNP conditions. These experimental boundaries place severe restrictions on continuous efforts to better understand and improve the DNP process.
The purpose of this work was to measure the trityl main ESR features at high field DNP conditions (6.7 T) and cryogenic temperatures (1.1 K) to investigate the mechanism behind Gd-doping. Using a purpose built Longitudinal Detection (LOD) ESR probe,\textsuperscript{24} we could compare the properties of trityl embedded in a generally applicable sample matrix (glycerol-water) for [U-$^{13}$C$_7$]-D-glucose polarization, to the well-studied sample of trityl in neat [1-$^{13}$C]PYR. The choice of glucose was driven by a growing interest in the hyperpolarization community for this substrate, which is central for energy metabolic studies in all cells. Also, the use of glucose has been limited so far by the relatively low achievable carbon polarization, even at high field.\textsuperscript{20} In particular, we aimed at investigating if $T_{1e}$ reduction was the only parameter affected by Gd$^{3+}$ doping and why, for the same trityl radical concentration and experimental conditions, neat PYR samples outperform other $^{13}$C-labeled substrates at high magnetic field. We will discuss our results in the frame of spin temperature theory and the DNP mechanism known as Thermal Mixing (TM).\textsuperscript{4, 25}
In a recent publication, Ardenkjær-Larsen at al. investigated the optimal AH111501 trityl concentration as a function of the magnetic field strength to achieve efficient DNP on neat [1-$^{13}$C]PYR.$^{13}$ At 6.7 T the optimum turned out to be 30 mM. We chose [1-$^{13}$C]PYR plus 30 mM AH111501 trityl as benchmark sample formulation for our study (from now onward referred as PYR_sample). The corresponding DNP sweep overlaid to the radical LOD-ESR spectrum and $T_{1e}$ measurements are reported in Figure 1, panel A and B, respectively. The PYR_sample showed a somewhat asymmetric ESR spectrum centered at $\omega_0 = 187.99$ GHz with full-width at half-maximum (FWHM) of 84±2 MHz well matched to the $^{13}$C Larmor frequency (71.8 MHz). At this low temperature, the trityl radical g-tensor appeared rhombic ($g_{xx} = 2.00332$, $g_{yy} = 2.00311$, $g_{zz} = 2.00237$; see Supporting Information, Figure S1) rather than axial, as reported previously by Lumata et al. for measurements performed at 100 K.$^{16}$ The DNP sweep reflected well the asymmetry of the ESR spectrum, showing a slightly sharper negative lobe and zero crossing very close to $\omega_0$. The $T_{1e}$ measured 400±2 ms, less than half of the value previously reported at 3.35 T and 1.2 K for a sample of neat [1-$^{13}$C]PYR doped with 15 mM trityl (approx. 1000 ms).$^{12}$ At the optimum microwave irradiation frequency (i.e. 187.94 GHz), corresponding
to the DNP positive maximum, 70±5% carbon polarization was achieved with a buildup time constant of 1200±50 s (see Supporting Information, Figure S2). Moreover, modulation of the microwave frequency had no effect on the buildup time and DNP enhancement indicating that the $PYR_{sample}$ was characterized by fast and efficient electron spectral diffusion. The relatively short $T_{1e}$ of the $PYR_{sample}$ was not merely a radical concentration effect. For a sample containing 15 mM trityl only we measured a $T_{1e}$ of 430±3 ms (see Supporting Information, Figure S3).

The same concentration of AH111501 trityl radical (i.e. 30 mM) was used in the preparation of a sample containing 2 M [U-$^{13}$C,d$_7$]-D-glucose in glycerol:water 50:50 (v:v) (from now onward referred as $GLU0_{sample}$). For this sample, the maximum achievable DNP enhancement was lower than the $PYR_{sample}$. At the optimum microwave irradiation frequency (i.e. 187.93 GHz), corresponding to the DNP positive maximum, 37±4% carbon polarization was achieved with a buildup time constant of 1330±50 s (see Table 1). Also in this case modulation of the microwave frequency had no effect. The $GLU0_{sample} T_{1e}$ was 2.5 times longer and the LOD-ESR spectrum/$^{13}$C DNP sweep 20
broader than the PYR_sample (see Figure 1, panel C and D, respectively). Indeed, the sample matrix composition modified the radical g-tensor that showed even more evident rhombic nature ($g_{xx} = 2.00354$, $g_{yy} = 2.00305$, $g_{zz} = 2.00223$; see Supporting Information, Figure S4). The different glassing sample matrix composition can also justify the 925±5 ms long $T_{1e}$ since the phonon spectral distribution, at the origin of the electron spin-lattice relaxation in the solid-state at this low temperature, depends on the stiffness and density of the latter.
Figure 1 LOD-ESR spectrum overlaid with $^{13}$C DNP sweep and $T_{1e}$ measurement at 6.7 T and 1.1 K for neat $[1^{-13}$C]pyruvic acid (PYR) with 30 mM trityl (panel A and B) and 2 M $[U^{-13}$C,d$_7$]-D-glucose in glycerol:water 50:50 (v:v) with 30 mM trityl (panel C and D). The ESR spectrum intensity was maximized according to the panel dimension.

The longer $T_{1e}$ of trityl dissolved in glycerol:water and glucose, compared to neat PYR at 6.7 T, encouraged us to investigate the effect of Gd$^{3+}$ doping at different concentrations,
i.e. \([\text{Gd}^{3+}] = 0.5\ \text{mM, 1 mM, 2 mM, 4 mM and 8 mM}\) (from now onward the different glucose samples will be referred as \(\text{GLU}[\text{Gd}^{3+}]_\text{sample}\)). Already at 0.5 mM Gd\(^{3+}\) the \(T_{1e}\) dropped to 538\(\pm\)19 ms. As one would expect in a system containing a slow relaxing spin species (i.e. the trityl) and a dilute fast relaxing spin species (i.e. the Gd\(^{3+}\)), the spin-lattice relaxation rate (\(1/T_{1e}\)) increased linearly with the Gd\(^{3+}\) concentration (see Figure 2 panel B).\(^4, 26-27\) Only the \(\text{GLU8}_\text{sample}\) deviated from this trend suggesting that at 8 mM Gd\(^{3+}\) concentration the interaction between the fast relaxing spins themselves was not negligible. Surprisingly, we also observed a narrowing of the ESR spectra especially on the high frequency hand side: the FWHM decreased to 85\(\pm\)2 MHz for the \(\text{GLU0.5}_\text{sample}\) and continued monotonically to 65\(\pm\)2 MHz for the \(\text{GLU8}_\text{sample}\) (see Figure 2 panel A and panel D); the g-tensor gradually evolved from rhombic to axial and the principal values obtained by Lumata et al from measurements performed at 100 K\(^{16}\) resulted in a very good fit for the \(\text{GLU8}_\text{sample}\) ESR spectrum (see Supporting Information, Figure S4). The origin of the narrowing is still not clear. Less efficient spectral diffusion, caused by \(T_{1e}\) shortening, can generate some narrowing of the ESR spectrum due to more difficult electron spins saturation at the extreme edges of the latter. Nevertheless, while trityl \(1/T_{1e}\)
increases linearly upon Gd$^{3+}$ doping of the glucose samples, the ESR FWHM appears to reach a minimum of approx. 60 MHz. Therefore, the contribution of a different phenomenon involving a modification of the radical $g$-anisotropy may be involved. To clarify this point, more extensive work involving simulations of the influence spectral diffusion on the LOD signal will be required.

The glucose samples series showed $^{13}$C DNP sweeps with increasingly closer maxima as a function of the Gd$^{3+}$ concentration (see Figure 2, panel C). This characteristic behavior was reported previously$^{17}$ and a purely $T_{1e}$ reduction could justify it.$^{25, 28-29}$ Nevertheless, in our case the DNP sweeps consistently followed the ESR spectra appearance modification: besides the narrowing, they became more and more asymmetric revealing a growing negative enhancement for increasing Gd$^{3+}$ concentration. All concerned numerical values are summarized in columns 2 to 5 of Table 1; data for individual samples are reported in Supporting Information, Figure S5 and S6.
**Figure 2** Effect of adding increasing concentrations of Gd$^{3+}$ (color coded) to a 2 M [U-$^{13}$C,$^{1}$H$_2$]-D-glucose in glycerol:water 50:50 (v:v) sample doped with 30 mM trityl, at 6.7 T and 1.1 K. LOD-ESR spectra with integral value normalized to 1 (panel A). Electron spin-lattice relaxation rate; experimental points are represented by the black circles, while the dotted red curve shows a linear fit (panel B). DNP microwave sweeps (panel C). Measured FWHM (panel D). In panel B and D values for the PYR$_{sample}$ have been added for comparison (red circles).
Gadolinium doping had a strong effect on the $^{13}$C spin polarization of the glucose samples. At 0.5 mM Gd$^{3+}$ concentration the nuclear spin polarization increased from $37\pm4\%$ to $59\pm4\%$; the \textit{GLU1_sample} achieved the maximum value (i.e. $69\pm3\%$) and for higher Gd$^{3+}$ concentration the polarization started to decrease (see Figure 3, panel A).

Moreover, the buildup time constant ($T_b$) remained basically unchanged (approx. 1400 s) for all measured Gd$^{3+}$ concentrations except for the \textit{GLU8_sample} where a longer $T_b$ was measured (see Figure 3, panel B). Solid-state polarization values and buildup time constants are reported in column 6 and 9 of Table 1.

**Figure 3** Solid-state $^{13}$C polarization (panel A) and buildup time constant (panel B) as a function of the Gd$^{3+}$ concentration in the glucose samples series (black circles) and the \textit{PYR_sample} (red circle). For each sample the polarization was measured at best microwave irradiation conditions.
Our results show a strong correlation between the maximum achievable carbon polarization on one side and two trityl electron spin parameters on the other: $T_{1e}$ and FWHM. Indeed, it is interesting to notice that [U-\(^{13}\)C,d\(_7\)]-D-glucose could reach a solid-state polarization value similar to the one achieved by the \textit{PYR-sample} (i.e. 70\%) when Gd\(^{3+}\) doping modified the aforementioned trityl radical parameters to match the ones observed in a pyruvic acid matrix (see position of red circles in Figure 3 panel B and D).

According to the TM model and spin temperature interpretation of the DNP mechanism, within the correct boundaries, a reduction of both parameters can increase the maximum achievable nuclear polarization. To qualitatively clarify this point, it is useful to look at the analytical stationary solution of the extended Provotorov equations for the inverse electron non-Zeeman spin temperature $\beta_{NZ} = \hbar/k_B T_{NZ}$. This is available in the high temperature approximation\(^{25}\), where it is found to be

\[
\frac{\beta_{NZ}}{\beta_L} = - \frac{2W_g(\omega)T_{1e}\omega_0(\omega - \omega_0)}{2W_g(\omega)T_{1e}[(\omega - \omega_0)^2 + D^2] + D^2} \tag{1}
\]

and

\[
\beta_L = \frac{\hbar}{k_B T_{NZ}} \tag{2}
\]
\[
\lim_{s \to \infty} \frac{\beta_{NZ}}{\beta_L} = -\frac{\omega_0(\omega - \omega_0)}{(\omega - \omega_0)^2 + D^2}
\]

Here \( \beta_L = \frac{\hbar}{k_B T_L} \) is the inverse lattice temperature, \( \omega \) the microwave irradiation frequency; \( D^2 \) the second moment of the ESR spectrum related to its linewidth; \( W_g(\omega) \) the rate of transitions induced by the microwaves and \( s = W_g(\omega)T_{1e} \) the saturation parameter. As far as the \(^{13}\)C Larmor frequency is smaller than the width of the ESR line, narrowing of the latter (i.e. smaller \( D \)) reduces the energy that can be stored in the non-Zeeman reservoir yielding directly to higher \( \beta_{NZ} \) that will eventually be transferred to the nuclei increasing their polarization enhancement. It is easy to see that when a strong microwave field is applied (\( s >> 1 \)), Eq. (2) has maxima \( \pm \omega_0 2D \) at \( \omega = \omega_0 \pm D \). If the ESR line becomes too narrow, the thermal contact between the electron non-Zeeman and nuclear Zeeman reservoirs decreases and the nuclear polarization is expected to drop.

Understanding the effect of \( T_{1e} \) shortening on the nuclear polarization is not as straightforward as for the ESR linewidth. Eq. (1), or its generalization beyond the high temperature approximation,\(^{29}\) represents an ideal case where the nuclei can relax to the
lattice only via the electron non-Zeeman reservoir. In that case shortening \( T_{1e} \) is equivalent to reduce the microwave power, thus the saturation factor \( s \). The consequence is a reduction of \( \beta_{NZ} \) and eventually a drop of the nuclear polarization.\(^{30}\) Differently, if the system presents nuclear leakage (relaxation pathways different from the coupling to the electron non-Zeeman reservoir), a moderate reduction of \( T_{1e} \) can be beneficial. Indeed, the rate at which \( \beta_{NZ} \) is achieved depends on \( T_{1e} \) and shortening of the latter can make leakage relatively less important.\(^ {25}\)

The already short \( T_{1e} \) for the \textit{PYR} \textit{sample} at 6.7 T can, at least partially, explain why addition of Gd\(^{3+} \) to DNP samples composed by neat [1-\(^{13}\)C]PYR plus trityl has a negligible effect at magnetic field values higher than 3.35 T.\(^ {8, 10, 14}\) At dDNP temperatures, the \( T_{1e} \) of trityl dissolved in a glassing matrix is dominated by the Orbach and direct process.\(^ {31}\) The Orbach process requires a hitherto unidentified excited state of the radical and involves two phonon modes coupling the two spin states to that excited state. The resulting relaxation rate is independent of the ESR frequency \( (\omega_{0S}) \), but depends exponentially on the ratio between the ground state – excited state energy difference \( (\omega_{a}) \).
and lattice temperature \((T_{L})\) such as 
\[ \frac{1}{T_{1e,0rb} \sim (\omega_{a})^3 \left( e^{\hbar \omega_{a}/k_{B}T_{L}} - 1 \right)} \]
where \(\hbar\) and \(k_{B}\) are Planck’s constant and Boltzman factor. Differently, the direct process involves only one phonon mode and its rate depends strongly on \(\omega_{0S}\), while being independent on the temperature under typical DNP conditions: 
\[ \frac{1}{T_{1e,dir} \sim (\omega_{0S})^5 \coth \left( \frac{\hbar \omega_{0S}}{2k_{B}T_{L}} \right)} \]
Thus, one would expect the Orbach process to prevail at low magnetic field, because more phonon modes can participate to the radical relaxation, and the direct process to provide a stronger contribution at high magnetic field. Our observation is in good agreement with Lumata et al.\(^{16}\) They reported no difference for the \(T_{1e}\) of trityl between 9.5 GHz and 95 GHz, while a considerable reduction of the latter was measured at 240 GHz.

For the glucose samples we cannot exclude a priori some nuclear leakage (e.g. paramagnetic impurities like \(O_2\) or trityl radical clusters), so up to 1 mM of \(Gd^{3+}\) concentration the reduction of the two radical parameters discussed above both improve the glucose polarization. Determining whether \(T_{1e}\) shortening or ESR line narrowing is more crucial in enhancing the glucose \(^{13}C\) polarization in the range 0 – 1 mM of \(Gd^{3+}\) doping will need further investigation and is beyond the scope of the present work. For
higher concentrations, the DNP enhancement decreases mainly because of the drastic reduction of $T_{1e}$ that prevents efficient ESR line saturation working at constant microwave power.

<table>
<thead>
<tr>
<th>$[\text{Gd}^{3+}]$ (mM)</th>
<th>$T_{1e}$ (ms)</th>
<th>ESR FWHM (MHz)</th>
<th>DNP maxima delta (MHz)</th>
<th>$\varepsilon_+$/$\varepsilon_-$</th>
<th>SS $T_b$ (s)</th>
<th>LS $T_1$ (s)</th>
<th>LS pol (%)</th>
<th>SS pol (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>925±4</td>
<td>108±2</td>
<td>120±5</td>
<td>1.124</td>
<td>1330±50</td>
<td>19.2±0.2</td>
<td>22±2</td>
<td>37±4</td>
</tr>
<tr>
<td>0.5</td>
<td>538±19</td>
<td>85±2</td>
<td>110±5</td>
<td>1.010</td>
<td>1368±28</td>
<td>19.2±0.1</td>
<td>35±2</td>
<td>59±4</td>
</tr>
<tr>
<td>1</td>
<td>279±11</td>
<td>79±2</td>
<td>90±5</td>
<td>0.930</td>
<td>1357±58</td>
<td>19.1±0.1</td>
<td>41±1</td>
<td>69±3</td>
</tr>
<tr>
<td>2</td>
<td>205±6</td>
<td>74±2</td>
<td>90±5</td>
<td>0.877</td>
<td>1390±98</td>
<td>19.0±0.2</td>
<td>38±2</td>
<td>64±4</td>
</tr>
<tr>
<td>4</td>
<td>112±3</td>
<td>69±2</td>
<td>70±5</td>
<td>0.796</td>
<td>1360±90</td>
<td>18.7±0.2</td>
<td>36±1</td>
<td>61±2</td>
</tr>
<tr>
<td>8</td>
<td>80±2</td>
<td>65±2</td>
<td>70±5</td>
<td>0.770</td>
<td>1680±40</td>
<td>17.4±0.3</td>
<td>31±2</td>
<td>55±2</td>
</tr>
</tbody>
</table>

Table 1 $^{13}$C spin polarization of [U-$^{13}$C$_6$-d$_7$]-D-glucose at increasing Gd$^{3+}$ concentrations. In solid state at 6.7 T and 1.1 K (SS pol) and 10 s after dissolution in 40 mM phosphate buffer pH 7.3, at 9.4 T and 50 °C (LS pol). SS $T_b$ solid-state polarization build-up time constant. LS $T_1$ Liquid state relaxation constant. LS pol liquid state polarization measured at 50 °C and 9.4 T 10 s after dissolution. SS pol solid-state polarization, back calculated given the liquid state relaxation constant and the sample transfer time from polarizer to NMR magnet.
The liquid-state $^{13}$C polarization, measured in a 9.4 T vertical NMR magnet 10 s after dissolution and transport, nicely followed the trend of the solid-state values (see Table 1).

To better visualize the strong carbon polarization enhancement achieved via Gd$^{3+}$ doping, we report in Figure 4 the solid-state polarization buildup and liquid-state relaxation for the $GLU0_{\text{sample}}$ and the $GLU1_{\text{sample}}$. Addition of 1 mM Gd$^{3+}$ allowed us to achieve $41\pm1\%$ liquid-state $^{13}$C polarization, the highest value reported to date. Moreover, the liquid-state nuclear $T_1$ remained unchanged within the experimental errors ($19.2\pm0.2$ s).

Liquid-state polarization and relaxation time constant values of the other glucose samples are reported in Table 1 columns 7 and 8.

![Figure 4](image-url)  
*Figure 4* $^{13}$C solid-state polarization at 6.7 T and 1.1 K (left panel) and relaxation after dissolution at 9.4 T and 313 K (right panel) of $GLU0_{\text{sample}}$ (filled black circles) and $GLU1_{\text{sample}}$ (open grey circles). The
middle panel indicates the 10 s sample transfer time from solid to liquid-state; the inset of the right panel shows the hyperpolarized $^{13}$C glucose NMR spectrum.

It is very clear from this study that the sample matrix has a profound impact on the features of the trityl radical. The $PYR_{sample}$ at high magnetic field is a “lucky” combination of these properties yielding $^{13}$C polarization as high as 70%. Indeed, at 6.7 T and 1.1 K, the measured $T_{1e}$ of trityl in a neat frozen PYR matrix was less than half of what it was in a glycerol/water mixture and the FWHM was 20 % narrower. When glycerol/water was employed as glassing solvent Gd$^{3+}$ addition was used to fine tune the radical properties and significantly increase the glucose $^{13}$C DNP enhancement matching the polarization value achieved by [1-$^{13}$C]pyruvic acid. The gadolinium not only decreased the trityl $T_{1e}$, as expected, but also generated a narrowing of the ESR spectrum.

ASSOCIATED CONTENT

This material is available free of charge via the Internet at http://pubs.acs.org.
• Experimental methods

• Additional results

AUTHOR INFORMATION/ CORRESPONDING AUTHOR

*Andrea Capozzi, PhD

Technical University of Denmark

Department of Health Technology

Ørsted Plads, Building 349, room 110

2800 Kgs. Lyngby

T: +45 45 25 53 27

Email: andcapo@dtu.dk

ORCID: 0000-0002-2306-9049

NOTES

The authors declare no conflict of interest.

ACKNOWLEDGMENT
We thank Dr Jacques van der Klink and Dr Olivier Ouari for fruitful discussions. The research leading to these results has received funding from the Danish National Research Foundation (DNRF124); the European Union's Horizon 2020 research and innovation programme under the Marie Sklodowska-Curie grant agreement no. 713683 (COFUNDfellowsDTU).

REFERENCES


Ernst, M. Dissolution DNP using trityl radicals at 7 T field. *Phys Chem Chem Phys* 2017,
19(29), 19196-19204.

15. Walker, S. A.; Edwards, D. T.; Siaw, T. A.; Armstrong, B. D.; Han, S.
Temperature dependence of high field C-13 dynamic nuclear polarization processes
with trityl radicals below 35 Kelvin. *Phys Chem Chem Phys* 2013, 15(36), 15106-
15120.

Song, L. K.; Merritt, M. E. Electron spin resonance studies of trityl OX063 at a

on DNP of [1-C-13]Pyruvate Doped with Trityl OX063, BDPA, or 4-Oxo-TEMPO. *J Phys

Sherry, A. D.; Lumata, L. Transition Metal Doping Reveals Link between Electron T-1


28. Serra, S. C.; Filibian, M.; Carretta, P.; Rosso, A.; Tedoldi, F. Relevance of
electron spin dissipative processes to dynamic nuclear polarization via thermal mixing.


29. Wenckebach, W. T. Dynamic nuclear polarization via thermal mixing: Beyond the

30. Jannin, S.; Comment, A.; van der Klink, J. J. Dynamic Nuclear Polarization by

31. Chen, H. J.; Maryasov, A. G.; Rogozhnikova, O. Y.; Trukhin, D. V.; Tormyshev,
V. M.; Bowman, M. K. Electron spin dynamics and spin-lattice relaxation of trityl radicals

32. Abragam, A.; Bleaney, B. *Electron paramagnetic resonance of transition ions.*