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Supporting Information

ABSTRACT: A new nitroxide-based biradical having a long electron spin–lattice relaxation time ($T_{1e}$) has been developed as an exogenous polarization source for DNP solid-state NMR experiments. The performance of this new biradical is demonstrated on hybrid silica-based mesostructured materials impregnated with 1,1,2,2-tetrachloroethane radical containing solutions, as well as in frozen bulk solutions, yielding DNP enhancement factors ($\varepsilon$) of over 100 at a magnetic field of 9.4 T and sample temperatures of $\sim$100 K. The effects of radical concentration on the DNP enhancement factors and on the overall sensitivity enhancements ($\Sigma$) are reported. The relatively high DNP efficiency of the biradical is attributed to an increased $T_{1e}$, which enables more effective saturation of the electron resonance. This new biradical is shown to outperform the polarizing agents used so far in DNP surface-enhanced NMR spectroscopy of materials, yielding a 113-fold increase in overall sensitivity for silicon-29 CPMAS spectra as compared to conventional NMR experiments at room temperature. This results in a reduction in experimental times by a factor of $>12700$, making the acquisition of 13C and 15N one- and two-dimensional NMR spectra at natural isotopic abundance rapid (hours). It has been used here to monitor a series of chemical reactions carried out on the surface functionalities of a hybrid organic–silica material.

INTRODUCTION

Nuclear magnetic resonance (NMR) is one of the richest sources of structural and dynamic information for molecular and materials sciences. However, it is traditionally limited by its intrinsic poor sensitivity. Dynamic nuclear polarization (DNP) has in principle the potential to increase NMR sensitivity by orders of magnitude by transferring polarization from unpaired electrons to nuclei. First discovered in the 1950s with low magnetic fields, high-field DNP NMR experiments are now possible with gyrotron sources, which are capable of delivering high-power high-frequency microwaves (MW). In this respect, in the past few years, DNP-enhanced solid-state NMR spectroscopy under magic angle spinning (MAS) conditions has made great progress, and DNP signal enhancement factors ($\varepsilon$) of up to $\sim$250 have been reported at high fields (up to 9.4 T) and low temperatures (<100 K) for biological solids as well as for hybrid or inorganic materials. The dramatic sensitivity gain provided by the DNP approach has thus enabled the determination of the interstrand architecture of amyloid fibrils and the elucidation of the bonding topologies at the surface of functionalized mesostructured materials. In this approach, samples are dissolved or impregnated with a solution containing stable exogenous paramagnetic centers such as nitroxides or trityls, which act as polarizing agents. Most high-field solid-state DNP NMR experiments are carried out in the presence of biradical polarizing agents as they can transfer polarization through the cross-effect (CE) mechanism, which is
currently the most efficient mechanism for DNP at high fields.20 The CE relies upon having a three-spin system consisting of two dipolar coupled electrons and a nucleus. For polarization transfer to occur from the electrons to the nucleus upon saturation of one of the EPR frequencies, the two unpaired electrons should have a significant dipolar coupling, and their EPR frequencies must differ by the nuclear Larmor frequency (here proton).18

Initial CE high-field DNP experiments were performed with solutions doped with high concentrations of monoradical species, such as TEMPO.21 Much effort has been recently devoted to the design and synthesis of more efficient radicals for DNP.21–24 Griffin and co-workers demonstrated that biradical polarizing agents yielded significant improvements in the DNP process, and therefore in the sensitivity enhancement factors.19 The success of biradicals is primarily attributed to the uniformly short distance between the two unpaired electrons in a biradical, which guarantees that all unpaired electrons are dipolar coupled to at least one other electron. With nitroxide-based biradicals, the EPR frequency difference is provided by g-anisotropy, and the difference in EPR frequencies can match the 1H Larmor frequency when the $g_{xx}$ (or $g_{yy}$) component of the $g$ tensor of one of the unpaired electrons is parallel to the $g_{zz}$ component of the other.25,26 Recently, triradicals, heterogeneous biradical species or mixtures of radical species,27,28 and metal ions29 have been proposed as alternatives to dinitroxide polarizing agents.

The water-soluble biradical TOTAPOL22 (Figure 1d) is today the most commonly employed exogenous polarizing agent in DNP solid-state NMR, in particular for biological applications.30 TOTAPOL was shown, however, to have a relatively flexible structure,25 and as a result the frequency matching condition for the CE mechanism is only properly fulfilled for the fraction of the biradicals that adopts the correct conformation. Another step forward was achieved by Griffin, Tordo, and co-workers who introduced bTbK26 (Figure 1e), a biradical functionalized nitroxide radicals,35 we show that, in 1,1,2,2-tetrachloroethane ("tetrachloroethane") at 100 K, bCTbK has a $T_{1e}$ value that is almost twice as long as that for bTbK. On a model hybrid mesostructured silica-based material,14,36 we show that this biradical yields proton and silicon-29 cross-polarization (CP) DNP enhancement factors that are between 2.5 and 4 times larger than bTbK. The reduction in experimental times provided by DNP experiments with bCTbK further pushes back the limits of solid-state NMR spectroscopy in terms of sensitivity, notably for the structural investigation of dilute surface species. This polarizing mixture is directly applicable to study a wide range of problems. We show here that with the high sensitivity provided by DNP with bCTbK, it becomes possible to expeditiously monitor the stepwise functionalization at the surface of hybrid materials by $^1$H, $^{13}$C, $^{29}$Si, as well as $^{15}$N solid-state NMR spectroscopy at natural isotopic abundance. Notably, high-quality one-dimensional $^{15}$N solid-state NMR spectra can be acquired with experiment times on the order of 1 h, and two-dimensional $^1$H–$^{15}$N HETCOR spectra can be acquired in less than 12 h.

## EXPERIMENTAL SECTION

Synthesis and characterization of the hybrid mesostructured silica materials are described in the Supporting Information. The biradical TOTAPOL was purchased in crystalline form from DyNuPol Inc., MA, and used without further purification. The solvents were purchased from Sigma-Aldrich and used without further purification.

**Synthesis of bCTbK.** bCTbK was prepared in a four-step sequence starting from 2,2,6,6-tetramethylpiperidin-4-one. The 2,6-
Figure 2. (a) Proton DNP (ε_H) and silicon DNP enhancement (ε_Si,CP). (b) The 29Si quenching factor (θ_Si) shown as a function of bCTbK concentration in 1,1,2,2-tetrachloroethane. (c) Overall (Σ_Si,CP and Σ_Si,CP) sensitivity enhancements. Note that ε_H quantifies the enhancement of the solvent and ε_Si,CP quantifies that of the surface. Experimental details and definition of the enhancement factors are given in the Supporting Information.

Table 1. DNP Efficiency Comparison and Electron Spin–Lattice Relaxation Times for the TOTAPOL, bTbK, and bCTbK Biradicals under Optimal Conditions

<table>
<thead>
<tr>
<th>biradical</th>
<th>solvent</th>
<th>ε_H</th>
<th>ε_Si,CP</th>
<th>Σ_Si,CP</th>
<th>Σ_Si,CP</th>
<th>Σ_Si,CP</th>
<th>Σ_Si,CP</th>
<th>T_1^e (μs)</th>
<th>T_2^e (μs)</th>
</tr>
</thead>
<tbody>
<tr>
<td>TOTAPOL</td>
<td>water</td>
<td>31(3)</td>
<td>15(4)</td>
<td>0.75(3)</td>
<td>554(4)</td>
<td>1.6(1)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>bTbK</td>
<td>tetrachloroethane</td>
<td>26(3)</td>
<td>21(3)</td>
<td>12(5)</td>
<td>453(4)</td>
<td>1.5(1)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>bCTbK</td>
<td>tetrachloroethane</td>
<td>105(3)</td>
<td>52(3)</td>
<td>38 (7)</td>
<td>762(4)</td>
<td>2.8(1)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

α Measured on Mat-PhOH impregnated with a 16 mM biradical solution. Estimated errors are given in parentheses. β Recorded on a triple resonance CPMAS probe at 99 K. The difference of DNP efficiency dependence on the biradical concentration on this probe. This explains the discrepancy between ε_H and ε_Si for bCTbK in tetrachloroethane. γ Recorded on a double resonance CPMAS probe at 102 K. δ In this case, it is not possible to quantify the overall enhancement of the carbon due to the fact that it is impossible to obtain a MW off carbon spectrum with high enough S/N ratio to calculate reliable integrals in a reasonable experimental time. However, it is possible to compare the relative efficiency of the different biradicals for enhancing the carbon by comparing the integrated intensities per unit of mass (Σ_Si). Here, the Σ_Si values of the aromatic resonances of Mat-PhOH impregnated with the various biradicals containing solutions are displayed. The concentration of the bTbK and TOTAPOL solutions was 12 mM, and the bCTbK one was a 16 mM solution (concentration giving optimal sensitivity for each biradical). The Σ_Si of Mat-PhOH impregnated with a 16 mM solution of bCTbK in tetrachloroethane was taken as a reference. ε Electron spin–lattice (T_1^e) and spin–spin (T_2^e) relaxation times were measured on the edge of the EPR line in a frozen bulk solution at 90 K with 8 mM biradical concentration at an EPR resonance frequency of 94 GHz. For TOTAPOL, glycerol was added to water to form a glass (60%/40% glycerol/H_2O by volume). f The 1H resonance of water is too broad to calculate a reliable proton DNP enhancement.
RESULTS AND DISCUSSION

bCTbK DNP Efficiency. The DNP efficiency of bCTbK was evaluated by impregnating a model mesoporous silica-based material Mat-PhOH\(^\text{16}\) containing phenolic ligands incorporated at the surface of the pores ((Figure 1a–c). The material was impregnated with bCTbK containing tetrachloroethane solutions of various concentrations. Tetrachloroethane was used because we have previously shown that this solvent provides optimal DNP enhancements (\(\varepsilon\)) for bTbK.\(^\text{31}\) Figure 2a displays the dependence of the proton (\(\varepsilon_H\)) and silicon (\(\varepsilon_{Si,CP}\)) DNP enhancement on the radical concentration (where \(\varepsilon\) is defined as the ratio of the integrated intensities of the spectra acquired with and without microwave irradiation under otherwise identical conditions). \(\varepsilon_H\) is found to peak at 65 for a radical concentration of 15.8 mM (similar to optimum concentrations previously observed for other biradicals).\(^\text{17,31}\) \(\varepsilon_{Si,CP}\) peaks at 52 for a concentration of 19.8 mM. The difference between the silicon enhancement at 15.8 and 19.8 mM is small, so the offset between the optimal concentration for proton and silicon is within experimental error. Note these curves were obtained with a double-resonance probe at 102 K. Higher enhancements for proton were obtained with a triple resonance probe at 99 K (see Table 1, footnote b, Figure 3, and (the calculation of these factors here is detailed in Table S3).\(^\text{17}\) \(\Sigma\) is an estimation of the overall gain in sensitivity obtained by carrying out low-temperature DNP solid-state NMR experiments instead of conventional room-temperature experiments.\(^\text{17}\)

As expected, and as previously observed on other systems, \(\theta\) diminishes as the radical concentration increases (Figure 2b).\(^\text{17,31}\) Here, however, quenching appears to be less than for water/TOTAPOL with \(\theta = 0.39\) for Mat-PhOH impregnated with a 12 mM solution of TOTAPOL in water and 0.54 for a 12 mM solution of bCTbK in tetrachloroethane. Figure 2c displays the overall sensitivity enhancements for silicon with (\(\Sigma_{Si,CP}\)) and without (\(\Sigma_{SLCP}\)) the Boltzmann factor as a function of the radical concentration.

\(\Sigma_{SLCP}\) and \(\Sigma_{SLCP}\) peak at 15.8 mM with a value of 38 and 113, respectively. As expected, \(\Sigma_{SLCP}\) is always lower than \(\Sigma_{SLCP}\) as it accounts for the decrease in \(\theta\) and \(T_1(1H)\) with increasing radical concentration (Figure S3), the latter meaning that more scans can be acquired in the same amount of time. A value of \(\Sigma_{Si,CP}\) = 113 translates into a reduction in experimental time by a factor of 12.700 as compared to ordinary room-temperature experiments.

Performance in Frozen Bulk Solutions and Other Solvents. bCTbK also provides high enhancements in frozen bulk solution (\(\varepsilon_H = 63, \varepsilon_{Si,CP} = 67\), Figure S4), as compared to \(\varepsilon_H = 22\) for bTbK.\(^\text{31}\) The high enhancements provided by bCTbK also mean that reasonable \(\varepsilon\) of \(\sim 30\) can be obtained for nonoptimal solvents (e.g., \(\varepsilon_H = 23\) and \(\varepsilon_{Si,CP} = 26\) for a 12 mM solution of bCTbK in mesitylene as opposed to \(\varepsilon_H = 6\) for a 10 mM solution of bTbK in mesitylene,\(^\text{31}\) both impregnated in Mat-PhOH, Figure S5). This therefore extends the number of solvents that can be used for DNP experiments, which is a necessity if chemically reactive systems are to be studied by DNP SENS in the future.

Comparison with Previously Used Biradicals. Under the same experimental conditions, both TOTAPOL and bCTbK are found to be typically less efficient than bCTbK (\(\varepsilon_{Si,CP} = 31\) for TOTAPOL/water and \(\varepsilon_{Si,CP} = 21\) for bTbK/tetrachloroethane, both impregnated within Mat-PhOH, Table 1). With \(\varepsilon_{Si,CP} = 52\), Mat-PhOH impregnated with bCTbK solutions provides the best enhancement obtained so far under any conditions for this model material. This corresponds to an improvement in the DNP efficiency by a factor of about 2.5 with respect to bTbK in tetrachloroethane.

We attribute the improved performance of the bCTbK biradical mainly to more favorable electron relaxation times. The DNP efficiency of biradicals has been shown to be significantly dependent on \(T_{1e}\) as a longer \(T_{1e}\) allows for a better saturation of the EPR line, therefore increasing the DNP effect.\(^\text{21,33,32}\) The effect of higher electron saturation on the DNP enhancement has previously been demonstrated by Thurber et al. by using field-modulation during the polarization buildup.\(^\text{25}\) To the best of our knowledge, biradicals have not been previously developed for use as DNP polarizing agents to specifically have long relaxation times. The methyl groups around each nitroxide center in bTbK were replaced with spirocyclohexyl groups in bCTbK, increasing the size and rigidity of the radical. Both the low-temperature and the room-temperature EPR spectra of bTbK and bCTbK dissolved in tetrachloroethane are similar, which suggests that the g and A tensors of both biradicals are similar.

Figure 3. Comparison between the X-ray structure of bTbK and the DFT optimized geometry of bCTbK (optimized at the B3LYP/6-31G(d,p) level).

Figure S2), with \(\varepsilon_H\) peaking at 105 for Mat-PhOH impregnated with a 16 mM bCTbK solution in tetrachloroethane (note that an enhancement of 105 corresponds to a 1% proton polarization at 400 MHz and 99 K, as compared to 0.0097% for the thermal equilibrium polarization, without DNP). The increased \(\varepsilon\) observed with a different probe was attributed to the slightly lower sample temperatures as well as to slightly higher MW power at the sample.

Note that \(\varepsilon\) quantifies the observed DNP efficiency, but it does not take into account factors such as the reduction in NMR signal intensity or changes in longitudinal relaxation times caused by the introduction of a paramagnetic radical in the sample.\(^\text{17,25}\) We have recently shown how the equivalent fraction of NMR active nuclei that contributes to the NMR signal (the quenching factor \(\theta\)) can be experimentally measured, and we introduced overall sensitivity enhancement factors \(\Sigma\) and \(\Sigma\), which take into account these parameters.\(^\text{17}\) The factor \(\Sigma\) also takes into account the additional sensitivity enhancement (Boltzmann) factor of \(\sim 3\) obtained by carrying out the experiments at 100 K rather than at room temperature.
Because bCTbK has not yet yielded crystals suitable for diffraction, to confirm that the structures of these two spiro radicals are similar, the geometry of both radicals was optimized with DFT calculations (Figure 3, Table 2). The accuracy of the calculations was confirmed by noting that the calculated geometry of bTbK closely matches that obtained from its single-crystal X-ray diffraction structure (Table 2). DFT calculations show that bTbK and bCTbK share analogous structures: (i) the calculated angles \( \psi \) between the mean planes \( \text{C}_6\text{N}_1\text{O}_2\text{C}_4 \) and \( \text{C}_7\text{N}_5\text{O}_6\text{C}_8 \) planes, (ii) the intramolecular nitrogen–nitrogen (N–N) distances in bTbK and bCTbK are predicted to differ by only 0.044 Å. This suggests that the electron–electron dipolar couplings are on the order of 29 MHz and are nearly identical for both radicals.

Having confirmed that the major structural features and EPR tensor parameters for both bTbK and bCTbK are essentially the same, it is then clear that the improved DNP enhancements obtained with bCTbK can be mainly attributed to its slower electron relaxation. \( T_{1e} \) is found to be 1.7 times higher for bCTbK (762 \( \mu \)s) than for bTbK (453 \( \mu \)s) (Table 1) under otherwise identical conditions. It is known that for radicals in glassy organic solvents around 100 K, Raman processes are the dominant relaxation mechanism, suggesting that \( T_{1e} \) is likely to increase as the rigid size/molecular mass of the radical is increased.\(^{34}\) It has also been observed that methyl groups lead to shorter transverse relaxation times.\(^{35}\) We also note that the \( T_{1e} \) of bCTbK in tetrachloroethane is longer than that of TOTAPOL in water (554 \( \mu \)s), despite the fact that electron relaxation times of nitroxides in frozen solutions are known to increase with the polarity of the solvent.\(^{34}\) A water-soluble version of bCTbK is expected to have an even longer \( T_{1e} \) possibly leading to higher enhancements. The high DNP efficiency of bCTbK can thus be partially attributed to both its favorable orientation of the TEMPO moieties and its long \( T_{1e} \).\(^{34}\)

Note that \( T_{1e} \) is known to be anisotropic along the EPR line, and it has been measured in this study at the high-field edge of the spectrum, corresponding to the zone irradiated during DNP experiments.

Improved DNP efficiency may also arise from longer spin–spin relaxation times (\( T_{2e} \)), as \( T_{2e} \) has also been hypothesized to help maintain the frequency match condition for the CE.\(^{21}\)

In summary, these results show the influence of electron relaxation times on the efficiency of polarizing agents and thus add another parameter to consider for further rational design of polarization agents for DNP experiments.

**Monitoring Chemical Reactions on a Surface.** Developing functional materials has become a major field of research in recent years due to their application in various areas such as catalysis, drug delivery, or gas purification. Understanding the structures and the chemical transformation of their surface functionalities at a molecular level is essential to provide reliable structure–property relationships, which are essential to the design and the controlled synthesis of the targeted functional materials. In this respect, solid-state NMR would be a method of choice, as it allows one to access structures, conformations, and dynamics of molecules, including surface species. However, the applicability of NMR is typically limited by its inherent low sensitivity, with the low concentration of the surface functionalities (typically <10 mmol/g), and the low natural abundance of most of the relevant NMR active nuclei (1% for carbon-13 and 0.3% for nitrogen-15).\(^{39}\)

**Table 2. Distances, Angle, and Dipolar Coupling for bTbK and bCTbK**

<table>
<thead>
<tr>
<th>compound</th>
<th>( d_{\text{N}−\text{O}} ) (Å)</th>
<th>( d_{\text{N}−\text{N}} ) (Å)</th>
<th>( \psi ) (°)</th>
<th>( D^* ) (MHz)</th>
</tr>
</thead>
<tbody>
<tr>
<td>bTbK (X-ray)</td>
<td>1.283</td>
<td>13.03</td>
<td>79.57</td>
<td>30.9</td>
</tr>
<tr>
<td></td>
<td>1.289</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>bTbK (DFT(^{\ddag}))</td>
<td>1.285</td>
<td>13.342</td>
<td>80.36</td>
<td>28.8</td>
</tr>
<tr>
<td></td>
<td>1.286</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>bCTbK (DFT(^{\ddag}))</td>
<td>1.285</td>
<td>13.249</td>
<td>77.56</td>
<td>29.3</td>
</tr>
<tr>
<td></td>
<td>1.286</td>
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</tr>
</tbody>
</table>

\(^{\ddag}\)N–O bond length of the two nitroxides. \(^{\ddag}\)O1−O5 distance. \(^{\ddag}\)N1−N6 distance. \(^{\ddag}\)Angle between the mean \( \text{C}_6\text{N}_1\text{O}_2\text{C}_4 \) and \( \text{C}_7\text{N}_5\text{O}_6\text{C}_8 \) planes.

\(^{\ddag}\)From the dipole point approximation (\( D = 52 \times 160/d_{1e}^2 \) where \( d_{1e} \) is the mean electron–electron distance). \(^{\ddag}\)Optimized at the B3LYP/6-31G(dp) level.

**Figure 4.** (a) Structure, (b) \(^1\text{H}−^{13}\text{C} \) CP spectra, and (c) \(^1\text{H}−^{15}\text{N} \) spectra of compounds Mat-PrN3, Mat-PrNH2, and Mat-PrIm. The most probable assignments based on the observed chemical shift resulting from this spectrum are shown. The asterisks indicate spinning sidebands, and the open circles note the presence of byproducts or incomplete conversion from the previous synthetic steps. The proton enhancements with a sample spinning frequency of 12 kHz are also reported. Experimental details are described in the text and in the Supporting Information.

We recently demonstrated that DNP SENS makes it possible to probe in acceptable experimental times (hours) the surfaces of various materials such as pure as oxides (silica,\(^{16}\) alumina\(^{40}\)), functionalized hybrid organic–inorganic silica materials,\(^{14}\) functionalized silica nanoparticles,\(^{15}\) and metal organic frameworks (MOFs)\(^{41}\) at natural isotopic abundance. However, DNP enhancement factors rarely exceeded 40 in our hands. To demonstrate the potential of the large enhancements provided by the bCTbK polarizing mixture, we investigate the stepwise chemical transformation and the formation of imidazolium-containing materials (Mat-PrIm), which can be used as permanently immobilized “ionic liquids”\(^{42}−^{46}\) or as precursors to N-heterocyclic carbenes,\(^{47}−^{48}\) an important class of ligands for stable single-site catalysts (Figure 4a).

Here, the imidazolium unit was introduced by a stepwise postfunctionalization of Mat-PrN3 prepared by a direct
synthesis approach. This included a Staudinger reduction providing an aminopropyl unit in Mat-PrNH₂ followed by its conversion into Mat-PrIm via reaction with formaldehyde, glyoxal, and benzylamine (Figure 4, see the Supporting Information for additional experimental details). Note that Mat-PrN₃ was prepared by a direct synthesis approach, that is, by a cohydrolysis co-condensation of (EtO)₃SiPrN₃ and (EtO)₄Si in the presence of a structure-directing agent, here Pluronic-P123. This allows the formation of high-surface area mesostructured materials as evidenced by TEM and N₂ adsorption experiments (see the Supporting Information) as well as the controlled site distribution of surface functionalities. Moreover, the nonpolar nature of azido group enables one to access a broad range of concentrations, including high concentrations, and makes it possible to prepare a variety of materials via postfunctionalization or click chemistry. To monitor these synthetic steps at the surface of materials with conventional room-temperature solid-state NMR would be prohibitively time-consuming (experiments typically take between 12 and 24 h for a one-dimensional carbon-13 CPMAS spectra; see the Supporting Information). In particular, acquisition of 2D correlation spectra or nitrogen-15 spectra at natural isotopic abundance is not conceivable even when they would be necessary for unambiguous assignment.

The interpretation and assignment of the DNP enhanced solid-state NMR spectra are now discussed in detail. The ¹³C CPMAS spectra of Mat-PrN₃ display three carbon resonances with chemical shifts of 8.1, 21.1, and 53.0 ppm, as expected from its proposed structure. They correlate with protons whose chemical shifts are 1.1, 2.15, and 3.2 ppm, respectively (Figure 5b). Upon conversion to Mat-PrNH₂ with the polarizing agent, high-quality 1D ¹³C CPMAS spectra (10 min, Figure 4b) and a 2D ¹²C-¹³C correlation spectra (3.5 h, Figure 5a–d) may be acquired and allow for the complete characterization of the carbon skeletons. The nitrogen-containing functionalities were identified with ¹³C-¹⁵N CPMAS spectra (14 h each, Figure 4c). These high-quality ¹⁵N CPMAS spectra even allow for minor reaction byproducts to be observed. Note that spectra with a signal-to-noise ratio of 20, high enough for routine characterization, can be acquired in 40 min (see Figure S6). A 2D ¹³C-¹⁵N correlation spectrum of Mat-PrIm was also acquired overnight (14 h, Figure 5e, vide infra). None of these would have been possible without the gain in sensitivity provided by DNP SENS. These results are all the more impressive considering that ¹⁵N is an extremely unreceptive NMR nucleus due to its low magnetogyric ratio and low natural abundance. Note that with bTbK, experimental lengths 6 times longer would have been necessary to get the same sensitivity.
Staudinger reaction, the $^{13}$C resonance of C$_{6}$ is shifted from 53.0 to 41.5 ppm, consistent with conversion of the azide group into an amine.

The nitrogen spectrum now displays one main resonance at 34.3 ppm, characteristic of the NH$_{2}$ group (Figure 4c). The byproducts observed on the $^{15}$N spectrum at 89 ppm are attributed to a small amount of unconverted starting azide units (ca. 10%). One interpretation for the incomplete conversion is the inaccessibility of some of these organic fragments on the surface. The carbon-13 spectrum of Mat-PrIm, obtained subsequently upon reaction with formaldehyde, glyoxal, and benzylamine, displays resonances with chemical shifts characteristic of aromatic groups (129.2 ppm $^{13}$C/6.8 ppm $^{1}$H) and (122.1 ppm $^{13}$C/6.8 ppm $^{1}$H), as well as the three expected aliphatic resonances (Figures 4b and 5d). The $^{15}$N spectrum confirms the presence of imidazolium nitrogen atoms with the most intense resonance centered at 185 ppm (the two nitrogen resonances being not resolved, Figure 4c).

The complete and detailed characterization of Mat-PrIm was further obtained by the acquisition of a 2D $^{1}$H−$^{15}$N correlation spectrum (Figure 5e), which shows three resolved cross peaks, all consistent with the structure of Mat-PrIm. The cross peak at (185.6 ppm $^{15}$N/8.9 ppm $^{1}$H) corresponds to a correlation between the nitrogens and aromatic protons, notably the C$_{6}$−H proton. A shoulder at (131.1 ppm $^{13}$C/8.9 ppm $^{1}$H) can be discerned in the $^{1}$H−$^{13}$C HETCOR spectra corresponding to a correlation between C$_{6}$ and this proton. The cross peak at (185.3 ppm $^{15}$N/5.4 ppm $^{1}$H) corresponds to the correlations between N$_{v}$ and N$_{c}$, and the CH$_{x}$ protons of C$_{v}$ and C$_{c}$. The peak at (187.2 ppm $^{15}$N/1.9 ppm $^{1}$H) indicates the correlation between the protons of C$_{v}$ and N$_{v}$, which is the only nitrogen close enough for magnetization transfer with this CP contact time (1 ms). The $^{15}$N chemical shift for this cross peak is slightly different from those of the previous two, confirming the presence of two distinct imidazolium nitrogens in the molecule. This analysis is confirmed by the lower intensity of this peak as compared to the other two, indicating a longer distance from the protons correlating with the other nitrogen, once again compatible with the molecular structure of Mat-PrIm.

The conformation of the organic moieties with respect to the surface is also important to help understand the folding of functionalities and the derived properties. A 2D $^{1}$H−$^{29}$Si correlation spectrum of Mat-PrIm could be acquired in 10 min. The cross peak at (−64.7 ppm $^{29}$Si/1.2 ppm $^{1}$H) shows the expected correlations between the T site silicon atoms (silicon bonded to the carbon of the organic moiety) and the aliphatic protons of carbon C$_{v}$ and C$_{c}$. The cross peak at (−99.8 ppm $^{29}$Si/7.6 ppm $^{1}$H) shows a correlation between the Q sites silicon atoms and the aromatic protons. This correlation indicates that the imidazolium ring is in close proximity to the surface of the material and that the tether folds to maximize the interaction between the aromatic groups and the surface functionalities.\textsuperscript{15}

In summary, the combination of the multidimensional proton, carbon, nitrogen, and silicon spectra clearly provides the mean to unambiguously and rapidly characterize these surface compounds at a molecular level. This demonstrates how, due to the large enhancements provided by the bCTbK biradical, DNP SENS can now be routinely applied to characterize materials and surface compounds.

**CONCLUSION**

Here, we introduced a new biradical, bCTbK, as a polarization source for DNP NMR experiments. This derivative of the well-known bTbK provides high DNP and sensitivity enhancements due to its rigid structure and absence of methyl groups, leading to slow electron relaxation and enhanced saturation of the EPR line. We expect that the very high enhancements will make this radical the polarization source of choice for the characterization of materials. As bCTbK is also a very efficient polarization source for bulk frozen solution, its applications are not limited to the study of surfaces, and it could be used for the study of proteins in lipid environments, polymers, or in dissolution DNP experiments, for instance. Design and testing of water-soluble versions of bCTbK for biological applications are currently underway.

The high efficiency of this radical was demonstrated for DNP SENS by following a series of chemical reactions on the surface of materials, monitoring the $^{13}$C, $^{29}$Si, $^{1}$H, and $^{15}$N chemical shifts at natural isotopic abundance using multidimensional correlation experiments. The reduction in experiment time by DNP SENS experiments with bCTbK is estimated to be on the order of 12 700 as compared to room-temperature experiments. These experiments would have previously been impossible to undertake at natural isotopic abundance in reasonable time. This study demonstrates that DNP SENS can be used routinely to characterize materials and surface compounds, as solution-state NMR is used today in molecular chemistry.

This work also illustrates that there is still much room for improvement of the polarization sources in DNP experiments and adds a new parameter to consider in the rational design of biradicals for DNP. Rigid radicals with bulkier functionalities are known; such radicals could have even longer electron relaxation times, which could lead to higher DNP enhancements.

**ASSOCIATED CONTENT**

Supporting Information

Additional experimental details of preparation of materials and radicals, measurement of quenching and enhancement factors, comparison of radical efficiency, and $^{15}$N NMR spectra of Mat-PrN$_{x}$, Mat-PrNH$_{2}$, and Mat-PrIm. This material is available free of charge via the Internet at http://pubs.acs.org.

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