

Fast magic angle spinning for the characterization of milligram quantities of organic and biological solids at natural isotopic abundance by ^{13}C - ^{13}C correlation DNP-enhanced NMR

Adam N. Smith ^{a,1,2}, Rania Harrabi ^{a,1}, Thomas Halbritter ^b, Daniel Lee ^a, Fabien Aussenac ^c, Patrick C.A. van der Wel ^d, Sabine Hediger ^a, Snorri Th. Sigurdsson ^b, Gaël De Paëpe ^{a,*}

^a Univ. Grenoble Alpes, CEA, CNRS, IRIG, MEM, 38000, Grenoble, France

^b University of Iceland, Department of Chemistry, Science Institute, Dunhaga 3, 107, Reykjavik, Iceland

^c Bruker BioSpin, Wissembourg, France

^d Zernike Institute for Advanced Materials, University of Groningen, Nijenborgh 4, 9747 AG, Groningen, Netherlands



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ABSTRACT

We show that multidimensional solid-state NMR ^{13}C - ^{13}C correlation spectra of biomolecular assemblies and microcrystalline organic molecules can be acquired at natural isotopic abundance with only milligram quantities of sample. These experiments combine fast Magic Angle Spinning of the sample, low-power dipolar recoupling, and dynamic nuclear polarization performed with AsymPol biradicals, a recently introduced family of polarizing agents. Such experiments are essential for structural characterization as they provide short- and long-range distance information. This approach is demonstrated on diverse sample types, including polyglutamine fibrils implicated in Huntington's disease and microcrystalline ampicillin, a small antibiotic molecule.

1. Introduction

Structural and dynamical characterization of solids at the atomic scale remains a salient, but difficult task. Magic Angle Spinning solid-state NMR (MAS ssNMR) has proved to be particularly adept at characterizing amorphous to microcrystalline solids ranging from biomolecules to materials [1,2]. Two major axes of methodological development have focused on fast MAS (*i.e.* sample rotation frequencies >25 kHz) [3,4] and sensitivity enhancement through dynamic nuclear polarization (DNP) [5–11]. With the recent development of both cryogenic fast MAS probes and DNP technologies, these two axes are now combined [12–16].

Faster MAS rates provide numerous experimental benefits, including more efficient averaging of anisotropic interactions (which is particularly important for high magnetic fields), longer coherence lifetimes, and larger and more homogenous radiofrequency field strengths (*i.e.* B_1) as a result of the smaller coil dimension [4]. To achieve these fast MAS rates, a smaller sample rotor diameter (*i.e.* outer diameter <3.2 mm) is required, which restricts sample volumes (*e.g.* ~2.5 μL in a Bruker 1.3

mm rotor *vs.* ~30 μL in a Bruker 3.2 mm rotor). Such a drastic reduction in sample volume can be advantageous when the sample amount is limited to a few milligrams or less, but also clearly impacts the overall signal-to-noise ratio. This is especially true for the detection of diluted spins with low gyromagnetic ratio (*e.g.* ^{13}C , ^{15}N). The loss in sensitivity due to the smaller sample volume can in part be mitigated by using indirect detection approaches, *via* ^1H spins [17,18]. In solids, this generally requires a MAS frequency >40 kHz to substantially reduce the proton linewidths, thus increasing the signal-to-noise ratio, and then benefits from the excellent receptivity of proton spins [19–22]. As a consequence, ^1H -detected fast MAS 2D X - ^1H heteronuclear correlation (HETCOR) experiments have been implemented in recent years for samples at natural isotopic abundance (NA), while correlation experiments under fast MAS between less receptive nuclei (*e.g.* ^{13}C - ^{13}C and ^{13}C - ^{15}N) could only be applied to isotopically-enriched samples owing to the associated sensitivity limitations [23–28].

DNP is used to boost the sensitivity of ssNMR experiments by transferring the relatively larger polarization of unpaired electrons to NMR-active nuclei of interest. Currently, the most efficient high-field

* Corresponding author.

E-mail address: gael.depaeppe@cea.fr (G. De Paëpe).

¹ These authors contributed equally.

² Current affiliation: Drug Product Technologies, Process Development, Amgen, 360 Binney St, Cambridge, MA 02142, United States.

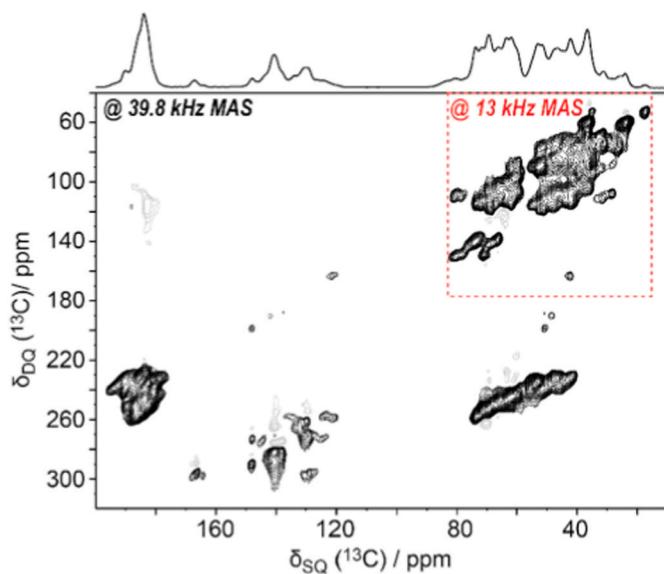


Fig. 1. DNP-enhanced ^{13}C - ^{13}C DQ-SQ dipolar correlation spectrum of ~ 1 mg of U^{13}C , ^{15}N LecA, using the recoupling sequence S_3 [48,49] at 9.4 T and 39.8 kHz MAS. The sample was prepared by impregnating the protein pellet with 40 mM cAsymPol-POK glycerol-d₈/D₂O solution (60:40 v/v). The maximum recoupling bandwidth, which would have been achieved at 13 kHz MAS frequency, is highlighted with a red dashed square. See Supporting Information for more experimental details.

MAS-DNP mechanism is the cross-effect (CE), which often relies on the use of bisnitroxide paramagnetic dopants [29–36], although efficient heteroradicals have also been introduced [37–40]. The corresponding transfer of polarization is driven through periodic energy-level (anti-) crossings of the coupled spin system composed of the two electrons and the surrounding nuclei [41,42]. As the MAS rate increases, these energy level crossings may become less efficient, which decreases the DNP efficiency and lengthens the hyperpolarization build-up time [43–45]. This is notably the case for polarizing agents (PAs) with small to moderate electron-electron couplings (dipolar coupling and/or J exchange interaction). This is one of the reasons why we recently introduced the AsymPol family of polarizing agents, which was designed to have large effective electron-electron $|2J + D|$ couplings between the two nitroxide moieties in the biradical [35]. This leads to efficient transfers of polarization at the different energy level crossings, which results in short hyperpolarization build-up time constants (T_B) [35,46]. As such, this class of PAs is particularly adapted for DNP with fast MAS and herein we illustrate their use in this regime.

We show that fast MAS, combined with DNP and PAs particularly suited to this regime, can overcome previous sensitivity limitations and enables the acquisition of multidimensional correlation spectra of low abundant nuclei at NA (e.g. ^{13}C - ^{13}C) on milligram-scale sample quantities. This is illustrated with polyglutamine (polyQ) amyloid fibrils (implicated in Huntington's disease) [47] and microcrystals of the antibiotic molecule ampicillin. Such experiments are typically orders of magnitude more demanding in terms of sensitivity than ^1H -X HETCOR-type experiments, but can provide extremely valuable information.

2. Material and methods

All experiments were performed on a 9.4 T Bruker Avance III spectrometer equipped with a 263 GHz gyrotron and a corrugated waveguide to transfer the microwave irradiation to the Bruker 1.3 mm MAS DNP probe cooled to ~ 100 K. The probe was in double resonance ^1H / ^{13}C configuration. The various DNP samples were impregnated with a DNP matrix containing 5–40 mM AsymPol-POK or cAsymPol-POK. The

detailed preparation, matrix composition and radical concentration is given in the Supporting Information for each sample. The sample amount in the 1.3 mm rotor was ~ 1 mg for LecA and $\text{D}_2\text{Q}_{15}\text{K}_2$ fibrils and ~ 2 mg for ampicillin. The DNP experiments were run at a MAS frequency between 35 and 40 kHz. All 2D ^{13}C - ^{13}C DQ/SQ dipolar correlation NMR experiments were acquired with the S_3 recoupling sequence [48,49]. The sequence and precise parameter settings for the different experiments are given in the Supporting Information.

3. Results and discussion

3.1. DNP-enhanced ^{13}C - ^{13}C dipolar correlation experiments with fast MAS and low-power recoupling

Access to faster MAS alone has extra built-in benefits: e.g. it enhances the use of low-power ^{13}C - ^{13}C dipolar recoupling sequences. Importantly, this allows the safe use of long mixing periods compatible with both long-distance polarization transfers and current radio-frequency (RF) hardware limitations, while keeping a relatively large ^{13}C excitation bandwidth. This is illustrated in Fig. 1, which shows a ^{13}C - ^{13}C double-quantum single-quantum (DQ-SQ) dipolar correlation experiment obtained on the uniformly labeled protein LecA (12.76 kDa). This experiment was recorded using the low-power dipolar recoupling sequence S_3 , for which the recoupling condition requires the RF field strength to match half the MAS frequency [48], thus limiting the spectral bandwidth on which the recoupling is effective to about that value [49]. As illustrated in Fig. 1, access to the faster spinning of about 40 kHz enables the acquisition of a broadband DQ-SQ S_3 correlation spectrum encompassing all ^{13}C resonances for the protein, which is not possible at lower MAS rates. Indeed, at 13 kHz MAS, the S_3 sequence would only achieve a recoupling bandwidth of ~ 6.5 kHz [48,49], prohibiting observation of broadband correlations from carbonyl/aromatic to aliphatic carbons. The DNP enhancement for the experiment of Fig. 1 was obtained using cAsymPol-POK as PA, as it was recently shown to be very efficient at polarizing proton-dense molecular systems [46]. Consistent with the characteristics of the AsymPol biradical family [35, 46], a high DNP enhancement of 130 was obtained at 9.4 T and 39.8 kHz MAS, accompanied by a very short build-up time, T_B , of 0.5 s. The spectrum presented in Fig. 1 was recorded in about 6 h with ~ 1 mg of protein sample. It can be noted that this experiment could also be collected in tens of minutes with a reasonable sensitivity. As is the case for ssNMR studies of proteins at room temperature, ^{13}C - ^{13}C correlation spectra are one of the cornerstones of biomolecular DNP, particularly for the study of large systems or the observation of cryo-trapped intermediates. This approach can be combined with specific and sparse labeling, as well as targeted and selective DNP approaches to compensate for the limited resolution resulting from low temperature DNP-NMR measurements [50,51].

3.2. Application on natural isotopic abundance fibrils

Considering the excellent sensitivity of the fast MAS-DNP spectrum obtained with cAsymPol-POK on only 1 mg of ^{13}C / ^{15}N -labeled protonated biomolecule, we wondered if it was even necessary to isotopically enrich the sample? Recently, the first steps towards multidimensional ^{13}C - ^{13}C and ^{13}C - ^{15}N experiments at NA have been demonstrated on samples containing tens of milligrams. This was achieved at 9.4 T using MAS-DNP with a relatively large sample holder (i.e. 3.2 mm outer diameter) and spinning at MAS frequencies less than 15 kHz [30,46, 52–65]. Through-bond (i.e. J based) [66] and through-space (i.e. dipolar coupling based) [67] experiments were reported and used, for instance, to determine the *de novo* assignment of a self-assembled 2'-deoxyguanosine derivative presenting two different molecules in the asymmetric crystallographic unit cell [61]. Beyond the fact that NA NMR is of interest for systems that cannot be easily isotopically labeled, additional benefits of working at NA were further illustrated. More specifically, it

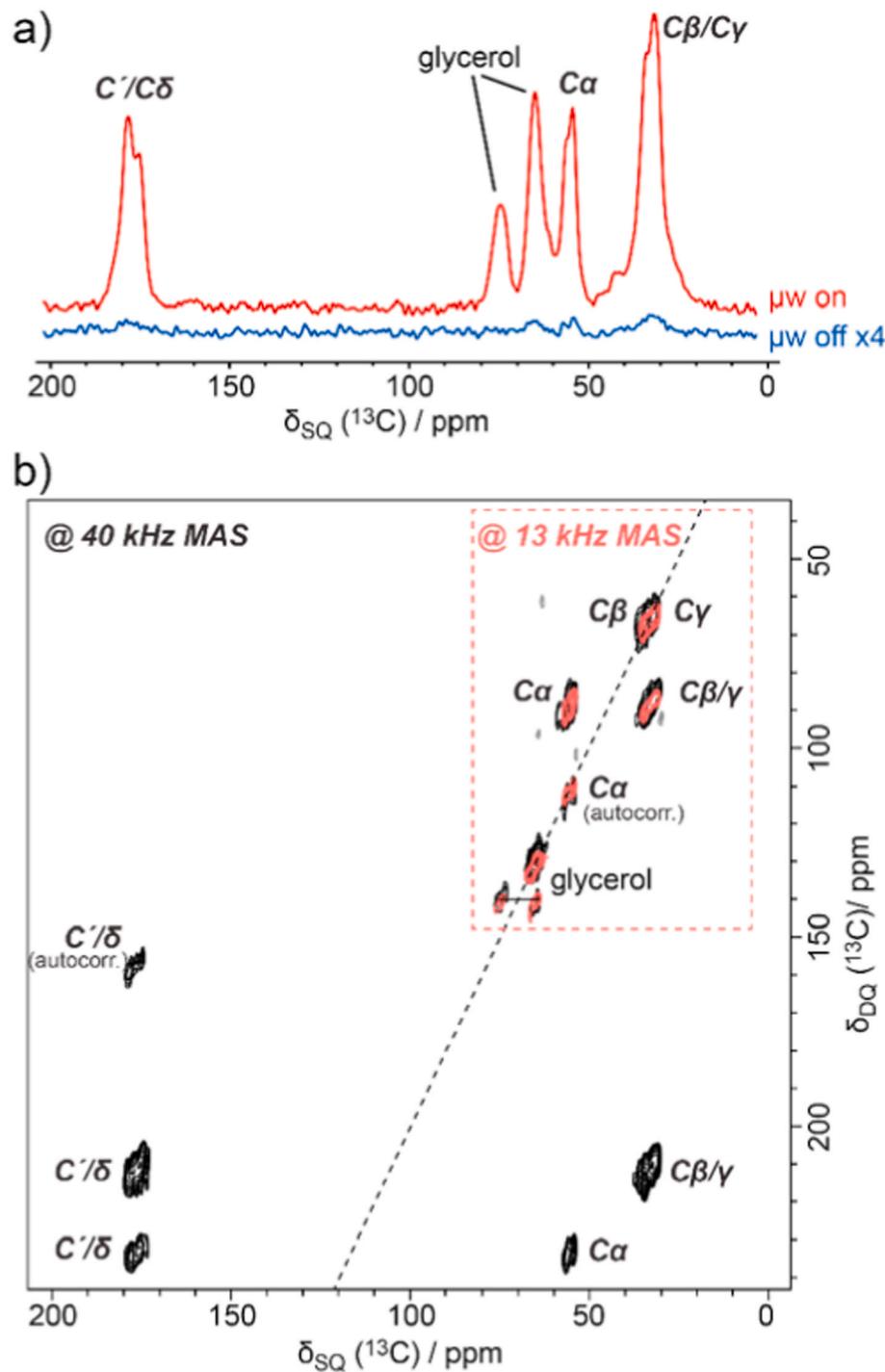


Fig. 2. (a) ^1H - ^{13}C CP spectra at a MAS frequency of 40 kHz on ~ 1 mg of polyQ amyloid fibrils formed by $\text{D}_2\text{Q}_{15}\text{K}_2$ with (red top trace) and without microwave (μw) (blue bottom trace) irradiation. The spectra were scaled to compensate for different numbers of co-added transients, to allow for intensity comparison. The sample was prepared with 5 mM AsymPol-POK glycerol-d₆/D₂O/H₂O solution (30:60:10 v/v/v). The DNP enhancement, measured by taking the ratio of signal intensity with and without μw irradiation, is 75. The DNP build-up time, T_B , was 3.1 s. (b) Superimposed DNP-enhanced ^{13}C - ^{13}C DQ-SQ dipolar correlation spectra on the same sample as in (a), recorded using 13.55 kHz (red) and 40 kHz (black) MAS frequencies (3.2 and 1.3 mm probes, respectively). Assignment for the ^{13}C nuclei from the glutamine residues in the polyQ amyloid core is given in the figure, based on our previous work [52]. Long-range inter-residue C α and C'/ δ autocorrelation peaks are clearly present. Note that the C'/ δ autocorrelation peak is folded in the indirect dimension. All the spectra were acquired at 9.4 T. See Supporting Information for more experimental details.

was demonstrated that the ^{13}C - ^{13}C spin dynamics of small organic molecules at NA can be treated as a superposition of individual dipolar build-up contributions from intra- and inter-molecular spin pairs [62], which enables internuclear distances of up to ~ 7 Å to be probed [64]. This would not be possible for fully isotopically-enriched systems. Subsequently, the approach was applied to disease-relevant protein aggregates, for which spectral fingerprints as well as dipolar build-ups were demonstrated [52], and extended to magnetic fields larger than 10 T [38].

Despite the utility of these studies, made with larger sample holders and lower spinning frequencies, there are numerous advantages to pursuing similar experiments under fast MAS with 1.3 mm (and smaller)

rotors for the reasons mentioned above. In particular, it addresses the structural characterization of mass-limited samples that cannot be obtained in large amounts and/or for which isotopic enrichment is not feasible, as for example synthetic and natural small molecules. This is illustrated in Fig. 2 on NA polyglutamine (polyQ) amyloid fibrils composed of the polyQ peptide $\text{D}_2\text{Q}_{15}\text{K}_2$, which features an analogous atomic structure to the aggregated state of disease-causing polyQ proteins. In this DQ-SQ ^{13}C - ^{13}C dipolar correlation spectrum, acquired at 40 kHz MAS on only ~ 1 mg of fibrils, we choose a long mixing time (23 ms) to allow polarization transfer over long distances (up to about 7 Å [64]) within the polyQ amyloid core. At NA, ^{13}C - ^{13}C spin pairs are only 0.01% abundant, and as such long-range distance restraints can be

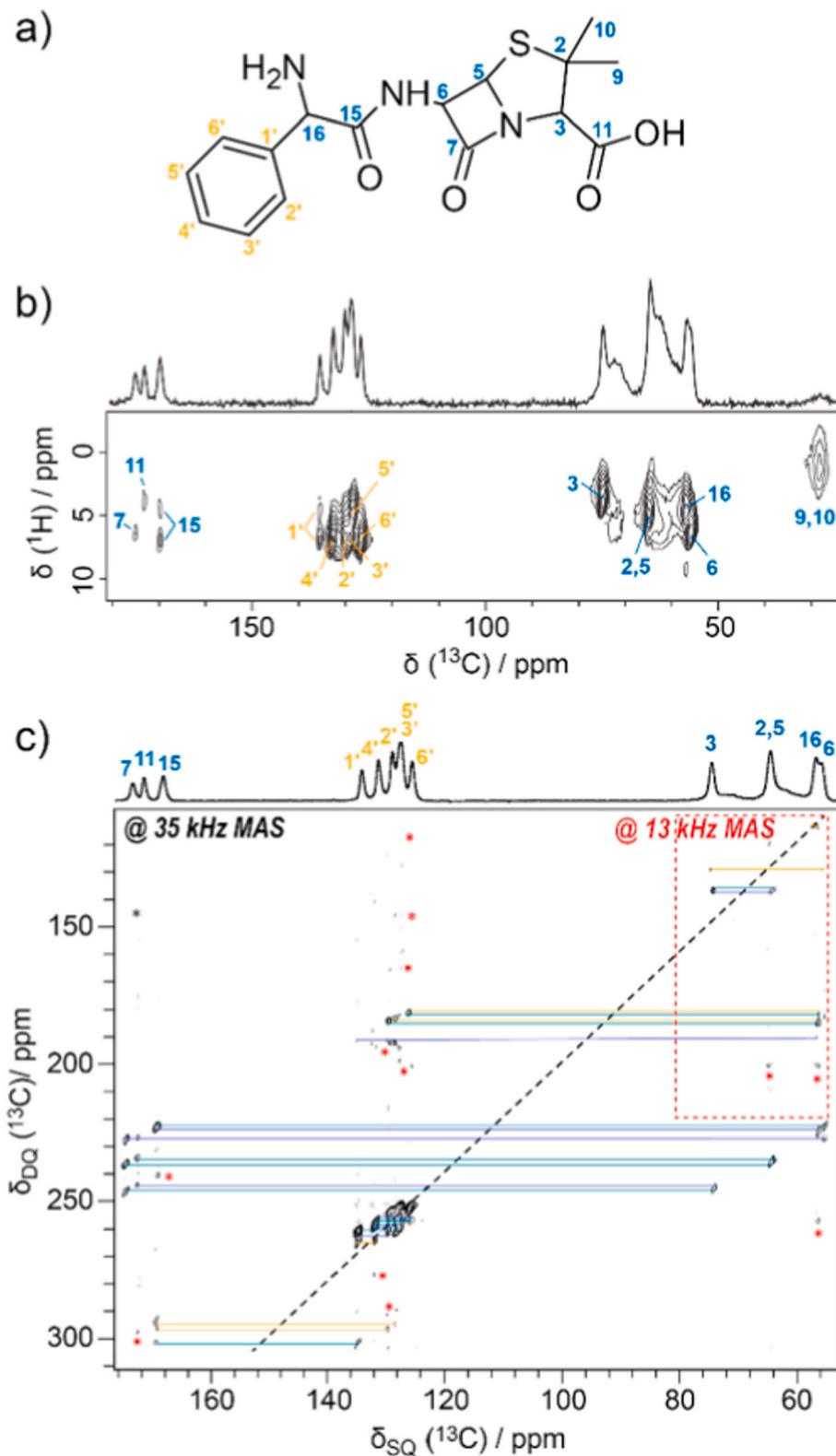


Fig. 3. (a) Chemical structure of ampicillin. (b) DNP-enhanced 2D ^1H - ^{13}C HETCOR dipolar correlation spectrum of ~ 2 mg of microcrystalline ampicillin at NA. A 200 μs CP contact time was used to ensure the observation of only short-range correlations. The MAS frequency was 40 kHz. The ampicillin powder was impregnated with 5 mM of AsymPol-POK in glycerol-d₈/D₂O/H₂O solution (30:60:10 v/v/v). The ^{13}C assignment given in the spectrum is based on the published assignment [69]. (c) DNP-enhanced 2D ^{13}C - ^{13}C DQ-SQ dipolar correlation spectrum using the recoupling sequence S₃ [48,49] at 35 kHz MAS on a similar sample of ampicillin impregnated with 40 mM of cAsymPol-POK in glycerol-d₈/D₂O/H₂O solution (60:30:10 v/v/v). The ^{13}C assignment is mapped out in the 2D spectrum, which yields short-range (equivalent to one-bond and two-bond in blue and cyan, respectively) and long-range (three-bonds and more in yellow) correlations (based on the known assignment [69]). Signals from spinning sidebands, t₁-noise, and solvent are indicated with red asterisks. The interference of the methyl rotation frequency at 100 K with RF pulses led to a reduced sensitivity for the methyl resonances. (see projection on top of the spectrum in panel b), and as a result homonuclear correlations with the methyl carbons are not visible (region therefore not shown). The maximum recoupling bandwidth, which would have been achieved at 13 kHz MAS frequency, is highlighted with a red dashed square. All spectra were acquired at 9.4 T. See Supporting Information for more experimental details.

probed without the interfering effects of dipolar truncation [54]. This type of correlation spectrum is ideal for differentiating between polymorphic structures [52]. For example, we clearly observe C α -C α and C-C' correlations from the polyQ backbone, which report on the β -strand configuration of the amyloid core, and C α -C'/C δ correlations, which report on side chain conformations [52]. Access to such long-distance constraints directly necessitates the ability to apply RF

irradiation for long recoupling times. The use of low-power sequences like S₃ [48,49] while maintaining an efficient recoupling over the entire spectral bandwidth is essential in this case [64] and only possible with fast MAS.

3.3. Application to natural isotopic abundance microcrystals

Efficient hyperpolarization of the fibrils benefits from the fact that one of the fibril dimensions is on the nanometer scale. The electron polarization, initially transferred from the PAs to the surface protons of the system, can easily be transferred across the width of the fibrils by proton-proton spin-diffusion, ensuring a homogeneous polarization of the nanofibrils [68]. In contrast, hyperpolarizing microcrystals of small organic molecules poses a greater challenge and requires the use of PAs that are particularly suited to polarize proton-dense systems, such as the ones of the AsymPol family [35,46]. The efficient transfer of polarization of these radicals is supported by relatively short build-up times, even under fast MAS conditions, which allows hyperpolarization to diffuse through the particles in a time shorter than the intrinsic nuclear relaxation time, T_{1n} . Thus, using AsymPol-POK and cAsymPol-POK, we were able to obtain DNP-enhanced ^1H - ^{13}C heteronuclear (HETCOR) and ^{13}C - ^{13}C DQ-SQ dipolar correlation experiments at 40 kHz MAS on only 2 mg of ampicillin at NA (see Fig. 3). Although ^1H - ^{13}C HETCOR experiments on organic microcrystalline solids can usually be recorded with standard ssNMR, DNP is needed for mass-limited synthetic small molecules and very useful to shorten experimental time, especially in the case of long ^1H T_1 relaxation times. Extending 2D ssNMR on NA samples also to ^{13}C - ^{13}C correlation experiments is orders of magnitude more challenging in terms of sensitivity. From the spectra presented in Fig. 3, we were able to find all expected ^{13}C resonances of the molecule (assignment based on Ref. [69]), with the exception of the two methyl groups (carbons 9, 10) and carbons 2 and 5, which are not resolved in the 2D experiments.

4. Conclusion

In conclusion, we have shown that ^{13}C multidimensional correlation spectra can be acquired with only milligram quantities of sample at NA. More specifically, broadband ^{13}C - ^{13}C dipolar correlation spectra, only possible under fast MAS, were acquired on biomolecular aggregates and microcrystalline small molecules, both at NA. The newly developed PAs of the AsymPol family, such as AsymPol-POK and cAsymPol-POK [35, 46], made these experiments feasible in a reasonable amount of time, thanks to their short hyperpolarization build-up times which increase the overall sensitivity of these DNP experiments under fast MAS.

In addition to the added spectroscopic benefits of performing experiments at fast MAS, the demonstrated feasibility of such experiments for small amounts of sample at NA also opens avenues for the characterization of sample types that were, so far, not amenable to ssNMR analysis. In particular, the characterization of *ex-vivo* amyloid fibrils from patient- or animal-derived systems is an intriguing aspect of research associated with numerous neurodegenerative disorders [70, 71]. As it is well known that amyloid fibrils adopt polymorphic structures and that the cellular milieu (*i.e.* molecular crowding, interaction with chaperones, post-translation modifications, etc.) can dictate the adopted conformation, it is imperative to study aggregates that are formed under native conditions [72]. With the method demonstrated here, one can now envision the study of *ex-vivo* amyloids at NA in the milligram quantities available from these systems.

The work presented here should also prove impactful for the study of natural products that are of great importance for numerous medicinal purposes. Indeed, a continued hurdle in the use and development of natural products, *e.g.* as new drug candidates, is their characterization, because they are typically isolated in small quantities. The structural information that can be gained from ssNMR is also invaluable for the characterization of other small molecules and their formulations. Finally, applications to other types of organic or hybrid (nano)materials, including thin films, can be enabled by these types of 2D NA ssNMR studies. The ability to rapidly characterize milligram quantities with multidimensional correlation experiments, as demonstrated here, will enable more routine use of ssNMR in these endeavors.

Declaration of competing interest

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: Gaël De Paëpe has patent Dinitroxide biradical compounds as polarizing agents pending to US. Snorri Sigurdsson has patent Dinitroxide biradical compounds as polarizing agents pending to US.

Data availability

Data will be made available on request.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.jssnmr.2022.101850>.

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