Improved NMR transfer of magnetization from protons to half-integer spin quadrupolar nuclei at moderate and high MAS frequencies

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Abstract. Half-integer spin quadrupolar nuclei are the only magnetic isotopes for the majority of the chemical elements. 20 Therefore, the transfer of polarization from protons to these isotopes under magic-angle spinning (MAS) can provide precious insights into the interatomic proximities in hydrogen-containing solids, including organic, hybrid, nanostructured and biological solids. This transfer has recently been combined with dynamic nuclear polarization (DNP) in order to enhance the NMR signal of half-integer quadrupolar isotopes. However, the cross-polarization transfer lacks of robustness in the case of quadrupolar nuclei and we have recently introduced as an alternative technique a *D*-RINEPT (through-space refocused

- 25 insensitive nuclei enhancement by polarization transfer) scheme combining a hetero-nuclear dipolar recoupling built from adiabatic pulses and a continuous wave decoupling. This technique has been demonstrated at 9.4 T with moderate MAS frequencies, $v_R \approx 10$ -15 kHz, in order to transfer the DNP-enhanced ¹H polarization to quadrupolar nuclei. Nevertheless, polarization transfers from protons to quadrupolar nuclei are also required at higher MAS frequencies in order to improve the ¹H resolution. We investigate here how this transfer can be achieved at $v_R \approx 20$ and 60 kHz. We demonstrate that the *D*-
- 30 RINEPT sequence using adiabatic pulses still produces efficient and robust transfers, but requires large rf-fields, which may not be compatible with the specifications of most MAS probes. As an alternative, we introduce robust and efficient variants of the *D*-RINEPT and PRESTO (phase-shifted recoupling effects a smooth transfer of order) sequences using symmetry-based recoupling schemes built from single and composite π -pulses. Their performances are compared using the average Hamiltonian theory and experiments at $B_0 = 18.8$ T on γ -alumina and isopropylamine templated microporous aluminophosphate AlPO₄-14,

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35 featuring low and significant ¹H-¹H dipolar interactions, respectively. These experiments demonstrate that the ¹H magnetization can be efficiently transferred to ${}^{27}Al$ nuclei using D-RINEPT with SR4²₄(270₀90₁₈₀) recoupling, and PRESTO with R227(180₀) or R16⁶₇(270₀90₁₈₀) schemes at $v_{\rm R} = 20$ or 62.5 kHz, respectively. The *D*-RINEPT and PRESTO recoupling schemes complement each other since the latter is affected by dipolar truncation, whereas the former is not.

We also analyse the losses during these recoupling schemes, and we show how these magnetization transfers can be used at v_R = 62.5 kHz to acquire in 72 min 2D HETCOR spectra between ¹H and quadrupolar nuclei, with a Non-Uniform Sampling.

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I. Introduction

Ouadrupolar nuclei with a nuclear spin quantum number S = 3/2, 5/2, 7/2 or 9/2 are the only NMR-active isotopes for over 60% of the chemical elements of the first six periods of the periodic table, including six of the eight most abundant elements by mass in the Earth's crust: O, Al, Ca, Na, Mg and K (Ashbrook and Sneddon, 2014). A wide range of materials, including 45 organic compounds, biological macromolecules as well as nanostructured or hybrid materials, contain half-integer spin quadrupolar nuclei and protons. Proximities between these isotopes have notably been probed in solid-state NMR experiments by transferring the polarization of protons to half-integer quadrupolar nuclei through dipolar couplings under magic-angle spinning (MAS) conditions (Rocha et al., 1991; Hwang et al., 2004; Peng et al., 2007; Vogt et al., 2013; Chen et al., 2019). More recently, this polarization transfer has been combined under MAS with dynamic nuclear polarization (DNP) in order to enhance the NMR signals of half-integer spin quadrupolar nuclei (Vitzthum et al., 2012; Perras et al., 2015a; Nagashima et al., 50 2020). This approach has notably allowed the detection of insensitive quadrupolar nuclei with low natural abundance, such as ¹⁷O or ⁴³Ca, or low gyromagnetic ratio, γ , such as ^{47,49}Ti, ⁶⁷Zn or ⁹⁵Mo, near surfaces of materials (Perras et al., 2015a, 2016, 2017; Blanc et al., 2013; Hope et al., 2017; Lee et al., 2017; Nagashima et al., 2020, n.d. 2021; Li et al.).

- This transfer has originally been achieved using cross-polarization under MAS (CPMAS) (Harris and Nesbitt, 1988). 55 Nevertheless, this technique lacks robustness for quadrupolar nuclei since the spin-locking of the central transition (CT) between energy levels $\pm 1/2$ is sensitive to the strength of the quadrupole interaction, the offset, the chemical shift anisotropy (CSA) and the rf-field inhomogeneity (Vega, 1992; Amoureux and Pruski, 2002; Tricot et al., 2011), Furthermore, CPMAS experiments require a careful adjustment of the rf-field applied to the quadrupolar isotope in order to fulfill the Hartmann-Hahn conditions, $(S+1/2)v_{1S} + \varepsilon v_{1H} = nv_R$, where v_{1S} and v_{1H} denote the amplitudes of the rf-fields applied to the S quadrupolar
- isotope and to the protons, respectively, $\varepsilon = \pm 1$, $n = \pm 1$ or ± 2 , and v_R denotes the MAS frequency, while avoiding the rotary 60 resonance recoupling (\mathbb{R}^3) $v_{1S} = pv_{\mathbb{R}}/(S+1/2)$ with p = 0, 1, 2, 3 (Amoureux and Pruski, 2002; Ashbrook and Wimperis, 2009). Moreover, the magnetization of the quadrupolar nuclei cannot be spin-locked for some crystallite orientations, which leads to line-shape distortions (Barrie, 1993; Hayashi and Hayamizu, 1993; Ding and Mcdowell, 1995).
- These issues have been circumvented by the use of the PRESTO (phase-shifted recoupling effects a smooth transfer of order) scheme (Perras et al., 2015a, b; Zhao et al., 2004) and more recently, the through-space refocused INEPT (denoted 65 RINEPT hereafter) (Nagashima et al., 2020; Giovine et al., 2019). These schemes benefit from higher robustness than CPMAS

since they do not employ a spin-lock on the quadrupolar channel, but instead a limited number (two or three) of CT selective pulses. In these sequences, the dipolar interactions between protons and quadrupolar nucleus are reintroduced by applying on the ¹H channel symmetry-based recoupling sequences, such as $R18^{5}_{2}$ for PRESTO or $SR4^{2}_{1}$ for RINEPT (Zhao et al., 2001;

- 70 Brinkmann and Kentgens, 2006a). In the case of recoupling sequences built from single square π-pulses, the RINEPT sequence using SR4²₁ (denoted RINEPT-SR4²₁) is more efficient than PRESTO at $v_R \ge 60$ kHz because of its higher robustness to rffield inhomogeneity and ¹H offset and CSA. At $v_R < 20$ kHz, the PRESTO technique is more efficient since the efficiency of RINEPT-SR4²₁ is reduced by the increased losses due to ¹H-¹H interactions during the SR4²₁ recoupling and the windows used to rotor-synchronize the SR4²₁ blocks, whereas the PRESTO sequence is devoid of these windows (Giovine et al., 2019).
- Recently, we have introduced a novel variant of the RINEPT sequence employing the SR4²₁ recoupling built: (i) from tanh/tan (tt) adiabatic inversion pulses, (ii) continuous-wave (CW) irradiations during the windows, and (iii) composite $\pi/2$ and π pulses on the ¹H channel, in order to limit the losses due to ¹H-¹H interactions and improve the transfer efficiency at moderate MAS frequencies (Nagashima et al., 2020, n.d.). This novel RINEPT variant, denoted RINEPT-CWc-SR4²₁(tt), is more efficient than PRESTO and CPMAS at $\nu_R \approx 12.5$ kHz and it has been combined with DNP to detect the NMR signal of
- 80 quadrupolar nuclei with small dipolar coupling with protons, including the low-γ isotopes, such as ^{47,49}Ti, ⁶⁷Zn or ⁹⁵Mo, and unprotonated ¹⁷O nuclei. Furthermore, for quadrupolar nuclei subject to large dipolar interactions, such as ¹⁷O nuclei of OH groups, we have shown that a RINEPT-CWc-SR4²₁(tt) version with only two pulses on the quadrupolar channel is more efficient that its PRESTO counterpart (Nagashima et al., n.d. 2021).

However, several NMR experiments require the transfer of ¹H magnetization to quadrupolar nuclei at $v_R > 12.5$ kHz. In particular, MAS frequencies of $v_R \ge 20$ kHz are needed to avoid the overlap between the center-bands and the spinning sidebands of satellite transitions (ST) in ²⁷Al NMR spectra at 18.8 T. In addition, magnetization transfers at $v_R \ge 60$ kHz are advantageous to acquire through-space hetero-nuclear correlation (*D*-HETCOR) 2D spectra between protons and quadrupolar nuclei endowed with high resolution along the ¹H dimension since fast MAS averages out the ¹H-¹H dipolar couplings.

Concurrently, we have demonstrated that the efficiency of PRESTO transfers using the R16⁶₇ recoupling can be improved at $v_R = 62.5$ kHz using (270₀90₁₈₀) composite π -pulses as a basic inversion element, where the standard notation for the pulses is used: ξ_{ϕ} denotes a rectangular, resonant rf-pulse with flip angle ξ and phase ϕ in degrees (Giovine et al., 2019). More recently, SR4²₁ and R12⁵₃ recoupling schemes built from (90₋₄₅90₄₅90₋₄₅) composite π -pulses have been proposed, but they have not yet been incorporated into RINEPT transfers (Perras et al., 2019). Globally, no systematic study of the RN^v_n recouplings built from composite π -pulses has been carried out.

In the present article, we investigate the use of RINEPT-CWc using an adiabatic recoupling scheme at the higher MAS frequencies of $v_R = 20$ and 62.5 kHz. We demonstrate using numerical simulations of spin dynamics and experiments on γ -alumina and isopropylamine templated microporous aluminophosphate AlPO₄-14 (hereafter AlPO₄-14) that the rf requirement of this technique increases with the ¹H-¹H dipolar interactions. In practice, this rf requirement is not compatible with the specifications of most MAS probes at $v_R \ge 20$ kHz, even for moderate ¹H-¹H dipolar interactions. As an alternative, we

100 introduce variants of the PRESTO and RINEPT sequences by selecting with average Hamiltonian (AH) theory the recoupling schemes built from single rectangular or composite π -pulses. Finally, using experiments on γ -alumina and AlPO₄-14, which feature small and moderate ¹H-¹H dipolar interactions respectively, we identify the most robust and efficient PRESTO and RINEPT transfers at B₀ = 18.8 T with v_R = 20 and 62.5 kHz.



115 Figure 1: ${}^{1}H \rightarrow {}^{27}Al$ (a,c) PRESTO-RN^v_n and (b,c) *D*-RINEPT-CWc-RN^v_n pulse sequences. Those applied to ${}^{1}H$ and ${}^{27}Al$ channels are shown in (a,b) and (c), respectively. The narrow and broad black bars represent $\pi/2$ and π -pulses, respectively. The acquisition of the FIDs (indicated with the vertical dashed line) starts after (a) the end of the RN^v_n block in the case of PRESTO or (b) on top of the echo shifted with $\tau_R/2$ with respect to the end of the last recoupling block in the case of RINEPT.

II. Pulse sequences and theory

120 II.1. PRESTO

II-1-1. Single-quantum hetero-nuclear dipolar recoupling

A RN_n^v sequence, where *N* is an even positive integer and *n* and *v* are integers, consists of *N*/2 pairs of elements $\mathcal{R}_{\phi} \mathcal{R}_{-\phi}^{*}$, with $\phi = \pi v/N$ radians an overall phase shift. \mathcal{R}_{ϕ} is an inversion pulse with a duration of nT_R/N , where $T_R = 1/v_R$ is the rotor period, and $\mathcal{R}_{-\phi}^{*}$ is an inversion pulse derived from \mathcal{R} by changing the sign of all phases. \mathcal{R} and \mathcal{R} are identical when they

125 are amplitude-modulated, i.e. all phase shifts are multiple of π . The rf-field requirement of RN_n^{ν} is equal to:

$$\nu_1 = \frac{N}{n} \frac{\xi^{tot}}{2\pi} \nu_R \tag{1}$$

where $\xi^{tot} = \sum_{i=1}^{P} \xi^{i}$ is the sum of the flip angles of the *P* individual pulses of the *R* element.

In the PRESTO sequence (Fig. 1a), symmetry-based γ -encoded RN_n^{ν} schemes applied to the ¹H channel reintroduce the |m|= 2 space components and the single-quantum (SQ) terms of the hetero-nuclear dipolar couplings between the protons and the 130 quadrupolar nuclei, as well as the ¹H CSA, while they suppress the contributions of ¹H isotropic chemical shifts, the heteronuclear *J*-couplings with protons, and the ¹H-¹H dipolar couplings to the first-order AH (Zhao et al., 2004). The hetero-nuclear dipolar interaction is characterized by a space rank *l* and a spin rank λ . A γ -encoded |m| = 2 SQ hetero-nuclear dipolar recoupling must selectively reintroduce the two components $\{l, m, \lambda, \mu\} = \{2, 2, 1, \mu\}$ and $\{2, -2, 1, -\mu\}$ of the hetero-nuclear dipolar coupling and ¹H CSA with $\mu = \pm 1$, while all other components must be suppressed.

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During these recoupling schemes, the contribution of the dipolar coupling between $I = {}^{1}H$ and *S* nuclei to the first-order Hamiltonian is equal to (Zhao et al., 2004):

$$\bar{H}_{D,IS}^{(1)} = \omega_{D,IS} S_z [I^+ \exp(i2\varphi) + I^- \exp(-i2\varphi)],$$
⁽²⁾

where $I^{\pm} = I_x \pm iI_y$ are the shift operators, and the magnitude and phase of the recoupled *I-S* dipolar coupling are given by

$$\omega_{D,IS} = -\kappa \frac{\sqrt{3}}{2} b_{IS} \sin^2 \left(\beta_{PR}^{D,IS} \right) \tag{3}$$

140 and

$$\varphi = \gamma_{PR}^{D,IS} - \omega_R t^0, \tag{4}$$

respectively, where b_{1S} is the dipolar coupling constant in rad/s, and κ is the scaling factor of the recoupled hetero-nuclear dipolar interaction, which depends on the RN_n^{ν} symmetry and the \mathcal{R} element. The Euler angles $\{0, \beta_{PR}^{D,IS}, \gamma_{PR}^{D,IS}\}$ relate the *I-S* vector to the MAS rotor frame, and t^0 refers to the starting time of the recoupling. The norm of $\overline{H}_{D,IS}^{(1)}$ does not depend on the 145 $\gamma_{PR}^{D,IS}$ angle, since these recoupling schemes are γ -encoded (Pileio et al., 2007; Martineau et al., 2012). The Hamiltonian of Eq.2 does not commute among different spin pairs, and hence, the PRESTO sequence is affected by dipolar truncation, *i.e.*, the transfer to distant nuclei is attenuated by the stronger couplings with nearby spins (Bayro et al., 2009).

As mentioned above, the SQ hetero-nuclear dipolar recoupling schemes also reintroduce the ¹H CSA with the same scaling factor κ , but without commuting with the recoupled ¹H-S dipolar interactions. Therefore, in the case of large ¹H CSA, for instance at high magnetic fields, this interaction can interfere with the ¹H-S dipolar couplings, especially with the small ones. These interferences can be limited by the use of the PRESTO-III variant, depicted in Fig.1a,c (Zhao et al., 2004), in which three CT-selective pulses are applied to the *S* channel. Indeed, the CT-selective π -pulses partly refocus the ¹H CSA, which limits these interferences.

II-1-2. Selection of the recoupling sequence

155 On the basis of the AH and spin dynamics simulations, the R18⁷₁ and R18⁵₂ schemes built from single rectangular π -pulses were selected for hetero-nuclear dipolar recoupling at moderate MAS frequencies, $v_R \approx 10$ kHz (Zhao et al., 2001), while more recently, sequences based on symmetries R12⁴₅, R14⁵₆, R16⁶₇, R14⁵₈, R18⁷₈, R16⁶₉, R20⁸₉ and R18⁷₁₀ using (270₀90₁₈₀) as inversion element were chosen for the measurement of ¹H CSA at fast MAS frequencies, $v_R \approx 60-70$ kHz (Pandey et al., 2015). We also transferred the proton polarization to ²⁷Al nuclei at $v_R = 62.5$ kHz using PRESTO with R16²₃ recoupling built from a 160 single rectangular π -pulse (Giovine et al., 2019). We screened here the RN_n^{ν} schemes built from single rectangular and composite π -pulses to achieve γ -encoded |m| = 2 hetero-nuclear SQ dipolar recoupling at $v_R = 20$ or 62.5 kHz. Dipolar recoupling at $v_R \ge 60$ kHz is useful to correlate the signals of quadrupolar nuclei with high-resolution ¹H spectra without using homo-nuclear dipolar decoupling. We tested the three following composite π -pulses: (1) (270₀90₁₈₀), which is offset-compensated and amplitude-modulated and has been employed

- 165 in several RN_n^{ν} sequences (Giovine et al., 2019; Carravetta et al., 2000; Levitt, 2002; Pandey et al., 2015), (2) (90₀240₉₀90₀), which compensates both rf inhomogeneity and offset (Freeman et al., 1980; Duong et al., 2019), and (3) (90₋₄₅90₋₄₅), which has homo-nuclear decoupling properties (Madhu et al., 2001). Adiabatic pulses cannot be employed for SQ heteronuclear dipolar recoupling since they yield vanishing scaling factors for the rotational components with $\mu \neq 0$ (Nagashima et al., 2018).
- 170 A total of 109 RN_n^v symmetries with $2 \le N \le 30$, $2 \le n \le 7$ and $1 \le v \le 11$ were found which recouple the $\{2, \pm 2, 1, \pm 1\}$ or $\{2, \mp 2, 1, \pm 1\}$ rotational components of the ¹H-S dipolar coupling and ¹H CSA. We selected the RN_n^v recouplings based on those symmetries with rf-field limited to $v_1 \le 120$ and 190 kHz for $v_R = 20$ and 62.5 kHz, respectively. We only considered the RN_n^v symmetries with $45 \le \phi \le 135^\circ$ since sequences with ϕ close to 90° are better compensated for rf field errors and inhomogeneities (Brinkmann and Kentgens, 2006b). The scaling factor, κ , of the recoupled ¹H-S dipolar interaction was
- 175 calculated using the 'C and R symmetries' Mathematica package (Carravetta et al., 2000; Brinkmann and Levitt, 2001; Brinkmann et al., 2000; Brinkmann and Edén, 2004).

These RN_n^{ν} symmetries eliminate the contribution of ¹H-¹H dipolar interactions to the first-order Hamiltonian, but not their contribution to the second-order. The cross-terms between ¹H-¹H interactions in the second-order Hamiltonian can be written (Brinkmann and Edén, 2004):

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$$\overline{H}^{(2),DD_1 \times DD_2} = \frac{1}{\nu_R} \sum_{\{1,2\}} \kappa^{DD_1 \times DD_2}_{\{1,2\}} \left[A^{DD_2}_{l_2 m_2} \right]^R \left[A^{DD_1}_{l_1 m_1} \right]^R \exp[i(m_1 + m_2)\omega_R t^0] \left[T^{DD_2}_{\lambda_2 \mu_2}, T^{DD_1}_{\lambda_1 \mu_1} \right]$$
(5)

where the sum is taken over all second-order cross-terms {1,2} between the { $l_1, m_1, \lambda_1, \mu_1$ } and { $l_2, m_2, \lambda_2, \mu_2$ } rotational components of DD₁ and DD₂ ¹H-¹H dipolar interactions, respectively. $\kappa_{\{1,2\}}^{DD_1 \times DD_2}$ is the scaling factor of this cross-term, $\left[A_{l_im_i}^{DD_i}\right]^R$ and $T_{\lambda_i\mu_i}^{DD_i}$ denote the component m_i of the l_i th rank spatial irreducible spherical tensor A^{DD_i} in the MAS rotor-fixed frame and the component μ_i of the λ_i th rank spin irreducible spherical tensor operator T^{DD_i} . Eq.5 indicates that the amplitude of the second-order Hamiltonian decreases at higher MAS frequencies. The magnitude of the cross-terms between ¹H-¹H interactions was estimated by calculating the Euclidean norm (Hu et al., 2009; Gansmüller et al., 2013):

$$\left\|\kappa_{\{1,2\}}^{DD_1 \times DD_2}\right\|_2 = \sqrt{\sum_{\{1,2\}} \left|\kappa_{\{1,2\}}^{DD_1 \times DD_2}\right|^2}.$$
(6)

For each basic element \mathcal{R} , we selected the RN_n^v schemes with the highest ratio $\kappa / \left\| \kappa_{\{1,2\}}^{DD_1 \times DD_2} \right\|_2$ in order to minimize the interference of ¹H-¹H dipolar interactions with the ¹H-*S* dipolar recoupling. Besides ¹H-¹H dipolar interactions, other cross-terms involving ¹H CSA and offset can also interfere with the ¹H-*S* dipolar recoupling. These cross-terms can be expressed by Eq.5, in which DD₁ and DD₂ indexes are substituted by other interactions, such as ¹H CSA or isotropic chemical shift (δ_{iso}).

For the selected symmetries, we estimated the magnitude of the cross-terms between ¹H CSA or offset by calculating the Euclidean norms $\|\kappa_{\{1,2\}}^{CSA \times CSA}\|_2$ and $\|\kappa_{\{1,2\}}^{\delta iso \times \delta iso}\|_2$ given by Eq.6.

The corresponding selected RN_n^{ν} sequences are listed in Tables S1 and S2 for $\nu_R = 20$ and 62.5 kHz, respectively.

- For $v_R = 20$ kHz, according to the AH, the RN_n^{ν} sequence with the highest robustness to ${}^{1}H{}^{-1}H$ dipolar interactions is $R22_2^{7}(180_0)$. However, this recoupling is slightly less robust to ${}^{1}H$ CSA and offset than $R18_2^{5}(180_0)$, which has already been reported. For this MAS frequency, the RN_n^{ν} schemes using the chosen composite pulses either required rf-fields greater than 120 kHz, e.g. $v_1 = 130$ and 173 kHz for the $R26_3^7$ schemes built from $(90_{-45}90_{45}90_{-45})$ and (270_090_{180}) pulses, or did not suppress efficiently the second-order cross-terms between ${}^{1}H{}^{-1}H$ interactions because of small rf-field ($v_1 \le 62.5 \text{ kHz}$).
- For $v_R = 62.5$ kHz, the RN_n^{ν} sequences using composite π -pulses recouple the ¹H-*S* dipolar interaction with a higher scaling factor than those built from single π -pulses. According to AH, the (90₀240₉₀90₀) basic element leads to the highest robustness to ¹H-¹H interferences. Even if the amplitude of the cross-terms is inversely proportional to the MAS frequency (Eq.5), the amplitude of these terms is lower at $v_R = 20$ than 62.5 kHz. The (270₀90₁₈₀) element is less robust to ¹H-¹H interferences, but benefits from a high robustness to offset. The selected RN_n^{ν} symmetries for this element include R14⁵₆ and R16⁶₇, which have
- 205 already been employed for the measurement of ¹H CSA and the transfer of ¹H polarization to half-integer quadrupolar nuclei at $v_R \ge 60$ kHz.(Giovine et al., 2019; Pandey et al., 2015) The scaling factors κ of the ¹H-*S* dipolar interaction of the RN_n^{ν} schemes built from single π -pulses are small with $45 \le \phi \le 135^{\circ}$, and hence we also selected in Table S3 those with an extended ϕ range of 20-160°. These recoupling schemes are less robust to offset than the RN_n^{ν} schemes built from (270₀90₁₈₀) element.

II-2. D-RINEPT

210 II-2-1. Zero-quantum hetero-nuclear dipolar recoupling

In the *D*-RINEPT sequence, the ¹H-*S* dipolar interactions are reintroduced under MAS by applying non- γ -encoded twospin order dipolar recoupling to the ¹H channel. These schemes reintroduce the |m| = 2 space components and the zeroquantum (0Q) terms of the ¹H-*S* dipolar interaction and ¹H CSA, *i.e.*, the rotational components $\{l, m, \lambda, \mu\} = \{2, \pm 2, 1, 0\}$, while they suppress the contributions of ¹H isotropic chemical shifts, the hetero-nuclear *J*-couplings with protons, and the ¹H-¹H dipolar couplings to the first-order AH (Brinkmann and Kentgens, 2006a, b). The contribution of the ¹H-*S* dipolar coupling to this Hamiltonian is equal to (Giovine et al., 2019; Brinkmann and Kentgens, 2006a; Lu et al., 2012):

$$\overline{H}_{D,IS}^{(1)} = 2\omega_{D,IS}I_z S_z \tag{7}$$

where

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$$\omega_{D,IS} = \kappa b_{IS} \sin^2 \left(\beta_{PR}^{D,IS} \right) \cos(2\varphi). \tag{8}$$

220 The norm of $\overline{H}_{D,IS}^{(1)}$ depends on the φ phase, given by Eq.4, and hence on the $\gamma_{PR}^{D,IS}$ angle. Therefore, these two-spin order dipolar recoupling schemes are non- γ -encoded. The Hamiltonian of Eq.7 commutes among different spin pairs and hence, these recoupling schemes are not affected by dipolar truncation. Similarly, the recoupled ¹H CSA contribution to the first-

order Hamiltonian is proportional to I_z and hence, also commutes with the recoupled ¹H-S dipolar interactions and does not interfere with the hetero-nuclear dipolar recoupling.

225 II-2-2. Selection of the recoupling sequence

Different RN_n^v sequences have been proposed to achieve non- γ -encoded |m| = 2 two-spin order dipolar recoupling, including: (i) symmetries $R(4n)_n^{2n-1} = R12_3^5$, $R16_4^7$, $R20_5^9$, $R24_6^{11}$, $R28_7^{13}$ and $R32_8^{15}$ for n = 3, 4, 5, 6, 7 and 8 using single π -pulses as basic element, which have been employed to measured ¹H-¹⁷O dipolar couplings at $v_R = 50$ kHz (Brinkmann and Kentgens, 2006b), (ii) SR4₁² recoupling built from a single π -pulse, which corresponds to the [$R4_1^2R4_1^{-2}$]_0[$R4_1^2R4_1^{-2}$]_{120}[$R4_1^2R4_1^{-2}$]_{240} sequence and has been employed in the RINEPT scheme (Nagashima et al., n.d.; Giovine et al., 2019), (iii) R12₃⁵ and SR4₁² schemes using a (90₋₄₅90₄₅90₋₄₅) composite π -pulse as a basic element, which have been incorporated into *D*-HMQC at $v_R = 36$ kHz (Perras et al., 2019), and (iv) SR4₁² schemes built from a (tt) adiabatic pulse, which have been used in the RINEPT sequence (Nagashima et al., n.d., 2020). During the (tt) pulse, the instantaneous rf-amplitude is equal to:

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$$\omega_{1}(t) = \omega_{1,\max} \begin{cases} \tanh\left[\frac{8\xi t}{T_{\rm R}}\right] & 0 \le t < T_{\rm R}/8\\ \tanh\left[2\xi\left(1 - \frac{4t}{T_{\rm R}}\right)\right] & T_{\rm R}/8 \le t < T_{\rm R}/4 \end{cases}$$
(9)

where $\omega_{1,\text{max}}$ is the peak amplitude of the rf-field, *t* refers to the time since the start of the pulse, which lasts $T_R/4$ when incorporated into the SR4²₁ recoupling scheme. The parameter ξ determines the rise and fall times of the pulse. Hence, in the frequency-modulated (FM) frame,(Garwood and DelaBarre, 2001) the frequency offset is

$$\phi_I(t) = \frac{\Delta v_{0,\max}}{2\theta \tan(\theta)} \ln\left\{ \cos\left[\theta\left(1 - 8\frac{t}{T_R}\right)\right] \right\}.$$
(10)

240 where $\Delta v_{0,max}$ is the peak amplitude of the carrier frequency modulation and θ determines the frequency sweep rate in the center of the pulse. Here, we employed $\xi = 10$ and $\theta = 87^{\circ} = atan(20)$ (Kervern et al., 2007; Nagashima et al., 2018, 2020).

We screened here the RN_n^{ν} schemes built from (180₀), (270₀90₁₈₀), (90₀240₉₀90₀) and (90₋₄₅90₄₅90₋₄₅) inversion elements. A total of 58 RN_n^{ν} symmetries with $2 \le N \le 30$, $2 \le n \le 7$ and $1 \le \nu \le 11$ were found which recouple the {2, ±2, 1, 0} rotational components of the ¹H-*S* dipolar coupling and ¹H CSA. We only considered the RN_n^{ν} symmetries with $60 \le \phi \le 120^{\circ}$ since the currently employed non- γ -encoded |m| = 2 two-spin order hetero-nuclear dipolar recoupling schemes have $75 \le \phi \le 90^{\circ}$.

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We calculated the scaling factor of the recoupled ¹H-S dipolar interaction and the Euclidean norm $\|\kappa_{\{1,2\}}^{DD_1 \times DD_2}\|_2$ of the cross-terms between ¹H-¹H interactions using the 'C and R symmetries' Mathematica package. For each basic element \mathcal{R} , we selected the RN_n^{ν} schemes with the highest ratios $\kappa / \|\kappa_{\{1,2\}}^{DD_1 \times DD_2}\|_2$. The selected RN_n^{ν} sequences are listed in Table S4, along with the parameters of the SR4²₁ schemes built from the different basic elements \mathcal{R} for the sake comparison. For these sequences, we calculated the Euclidean norms, $\|\kappa_{\{1,2\}}^{CSA \times CSA}\|_2$ and $\|\kappa_{\{1,2\}}^{\delta iso \times \delta iso}\|_2$, in order to estimate the magnitudes of the cross-terms between ¹H CSA and offset.

According to the AH, the $(90_0240_{90}90_0)$ composite π -pulse yields the highest robustness to ¹H-¹H dipolar interactions. However, the rf-field requirement of the RN_n^{ν} sequences built from this composite pulse, $v_1 = 1.16Nv_R/n$, is not compatible at $v_R = 62.5$ kHz with most 1.3 mm MAS probes (e.g. $v_1 = 291$ kHz for $SR4_1^2$). Furthermore, the highest robustness to ¹H CSA and offset is achieved using the (270_090_{180}) composite π -pulse. The $SR4_1^2$ schemes benefit from the highest robustness to ¹H CSA, because of the three-step multiple-quantum super-cycle (Brinkmann and Edén, 2004; Brinkmann and Kentgens, 2006a). Contrary to the RN_n^{ν} with |m| = 2 SQ hetero-nuclear dipolar recouplings, the rf-field of the RN_n^{ν} with |m| = 2 two-spin order schemes is always higher than $2v_R$ since these symmetries with 2n > N, such as $R12_9^5$, have smaller κ scaling factors for the basic elements employed here.

In the case of the adiabatic RN_n^{ν} (tt) sequences, the determination of the scaling factors of the first- and second-order terms of the effective Hamiltonian is more cumbersome since they depend on the $\nu_{1,max}$, $\Delta\nu_{0,max}$, ξ and θ parameters (Nagashima et al., 2018). For example, the scaling factor of the R12⁵₃ and SR4²₁ schemes is $\kappa = 0.31$ for $\nu_{1,max}/\Delta\nu_{0,max} = 0.685$, $\xi = 10$ and $\theta =$ 87°, and this value monotonously decreases for increasing $\nu_{1,max}/\Delta\nu_{0,max}$ ratios.

II-2-3. D-RINEPT-CWc sequence

- The *D*-RINEPT-CWc sequence is displayed in Fig.1b,c. The ¹H-*S* dipolar couplings are reintroduced by applying the RN_n^{ν} schemes listed in Table S4 during the defocusing and refocusing delays τ , which are identical in this article, even if distinct delays can improve the transfer efficiency (Nagashima et al., n.d.). As the two-spin order recoupling schemes are non- γ -encoded, they must be rotor-synchronized. We used here a delay of T_R between two successive RN_n^{ν} blocks. In the *D*-RINEPT-CWc sequence, a CW irradiation is applied during these delays in order to limit the losses due to ¹H-¹H dipolar interactions 270 (Nagashima et al., n.d.). The nutation during this CW irradiation is eliminated by employing CW irradiations with opposite
- phases. Furthermore, the robustness to ¹H rf-field inhomogeneity is improved by replacing the first π and second $\pi/2$ pulses by composite (90₀180₉₀90₀) and (90₉₀90₀) pulses, respectively, the CW irradiation being applied between the individual pulses (Freeman et al., 1980; Levitt and Freeman, 1979).

III. Numerical simulations

275 III-1. Simulation parameters

All simulations were performed using the version 4.1.1 of the SIMPSON package (Bak et al., 2000). The powder average was performed using 462 { α_{MR} , β_{MR} , γ_{MR} } Euler angles relating the molecular and rotor frames. This set of angles was obtained by considering 66 { α_{MR} , β_{MR} } pairs and 7 γ_{MR} angles. The { α_{MR} , β_{MR} } values were selected according to the REPULSION algorithm (Bak and Nielsen, 1997), while the γ_{MR} angles were regularly stepped from 0 to 360°.

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To accelerate the simulations, we used a ${}^{1}\text{H} \rightarrow {}^{15}\text{N}$ RINEPT transfer instead of the ${}^{1}\text{H} \rightarrow {}^{27}\text{Al}$ one, because the computing time is proportional to the cube of the size of the density matrix. Furthermore, in RINEPT experiments, only CT-selective

pulses are applied to the quadrupolar nuclei and hence, the contribution of the STs to the signal can be disregarded. The ${}^{1}H\rightarrow{}^{15}N$ RINEPT transfer was simulated for a ${}^{15}N^{1}H_{4}$ spin system. A similar approach has already been applied for the simulation of the RINEPT transfer from protons to quadrupolar nuclei (Nagashima et al., n.d.; Giovine et al., 2019). This

- ¹⁵N¹H₄ spin system comprises a tetrahedron of four protons with a ¹⁵N nucleus on one of its symmetry axis. The dipolar coupling constants between protons are all equal to $|b_{HH}|/(2\pi) = 1$, 7 or 15 kHz. The dipolar coupling between the ¹⁵N nucleus and its closest ¹H neighbor is $|b_{HN}|/(2\pi) = 2575$ Hz, corresponding to a ¹H-²⁷Al distance of 2.3 Å, typical of the distance between the protons of hydroxyl groups and the Al atoms of the first surface layer of hydrated γ -alumina (Lee et al., 2014). All protons were subject to a CSA of 6 kHz, *i.e.*, 7.5 ppm at 18.8 T, with a null asymmetry parameter (Liang et al., 2018). We
- 290 simulated the ¹H → ¹⁵N RINEPT-CWc sequences incorporating either SR4²₁(tt) or R12⁵₃(tt) recoupling schemes. We used a static magnetic field of 18.8 T, for which the ¹H and ¹⁵N Larmor frequencies were equal to 800 and 81 MHz, respectively, and MAS frequencies of v_R = 20 or 62.5 kHz. The defocusing and refocusing periods were both equal to their optimal values τ = 650 or 640 µs at v_R = 20 or 62.5 kHz, respectively. The rf-field nutation frequency on the ¹H channel was equal to 200 kHz during the π/2 and π-pulses that do not belong to the recoupling sequence, as well as the CW irradiation, whereas the pulses
- applied to $S = {}^{15}$ N nuclei were considered as ideal Dirac pulses. For the (tt) adiabatic pulses, the simulations were performed with $v_{1,max}/v_R$ and $\Delta v_{0,max}/v_R$ ratios ranging from 0.5 to 10 and from 10 to 200, respectively. All other pulses were applied on resonance. The density matrix before the first pulse was equal to $I_{1z} + I_{2z} + I_{3z} + I_{4z}$. We normalized the transfer efficiency of the 1 H \rightarrow 15 N RINEPT sequences to the maximal signal for a 1 H \rightarrow 15 N through-bond RINEPT sequence made of ideal Dirac pulses in the case of a 15 N- 1 H spin system with a *J*-coupling constant of 150 Hz.



Figure 2: (a-d) Simulated transfer efficiency of ${}^{1}H \rightarrow {}^{15}N D$ -RINEPT-SR4 ${}^{2}_{1}$ (tt) sequence for a ${}^{15}N{}^{1}H_{4}$ spin system as function of $\nu_{1,max}/\nu_{R}$ and $\Delta\nu_{0,max}/\nu_{R}$ for $\nu_{R} = 20$ and 62.5 kHz and $b_{HH}/(2\pi) =$ (a) 1, (b,d) 7 and (c) 15 kHz. (e,f) Experimental ${}^{1}H \rightarrow {}^{15}N D$ -RINEPT-SR4 ${}^{2}_{1}$ (tt) signal of L-histidine-HCl as function of $\nu_{1,max}/\nu_{R}$ and $\Delta\nu_{0,max}/\nu_{R}$ at 18.8 T with $\nu_{R} =$ (e) 40 or (f) 62.5 kHz. In (a-d) the white star indicates recoupling conditions with minimal rf field leading to maximal transfer efficiency, and the white vertical line mimics the rf-field distribution within the coil.

III-2. Optimal adiabatic recoupling

The transfer efficiency of RINEPT using RN_n^{ν} schemes built from adiabatic (tt) pulses depends on $v_{1,max}$ and $\Delta v_{0,max}$ parameters. For a similar ¹⁵N¹H₄ spin system with $|b_{HN}|/(2\pi) = 2.575$ and $|b_{HH}|/(2\pi) = 7$ kHz, spinning at $v_R = 12.5$ kHz, we showed using numerical simulations that a maximal transfer efficiency was achieved provided that $v_{1,max} = 0.07\Delta v_{0,max}$ and $v_{1,max}/v_R \ge 8$ (Nagashima et al., n.d.). In practice, we used $v_{1,max} = 11v_R = 137$ kHz and $\Delta v_{0,max} = 160v_R = 2$ MHz.

Similar simulations were performed here for $v_R = 20$ or 62.5 kHz. As seen in Fig.2a-c, at a given MAS frequency, higher ¹H-¹H dipolar couplings require higher rf-field and broader carrier frequency sweep so that the (tt) pulses remain adiabatic in spite of the modulation of the ¹H-¹H dipolar couplings by MAS (Nagashima et al., n.d.; Kervern et al., 2007). For $|b_{HH}|/(2\pi)$

315 = 7 kHz, the minimal v_{1,max}/v_R ratio decreases for higher MAS frequencies (compare Figs.2b and d) since the contribution of the modulation of ¹H-¹H dipolar couplings by MAS to the first adiabaticity factor is proportional to (v_{1,max})²/v_R and hence, v_{1,max} values proportional to √v_R, i.e. v_{1,max}/v_R ratio inversely proportional to √v_R, are sufficient to maintain the adiabaticity of the pulses (Kervern et al., 2007). Nevertheless, Fig.2d indicates that the SR4²₁(tt) recoupling requires v_{1,max} ≥ 313 kHz for v_R = 62.5 kHz, which is hardly compatible with the specifications of most 1.3 mm MAS probes. Similar transfer efficiencies were simulated for the RINEPT sequence with R12⁵₃(tt) recoupling scheme (not shown).

IV. NMR experiments

IV-1. Samples and experimental conditions

L-[U-¹⁵N]-histidine HCl (hereafter referred to as "histidine") and isotopically unmodified γ -alumina were purchased from Merck, and AlPO₄-14 was prepared as described previously (Antonijevic et al., 2006).

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All ¹H \rightarrow S RINEPT-CWc and PRESTO-III experiments were performed at $B_0 = 18.8$ T on Bruker BioSpin Avance NEO spectrometers equipped with double-resonance ¹H/X probes.

 ${}^{1}\text{H} \rightarrow {}^{15}\text{N}$ RINEPT-CWc-SR4²₁(tt) experiments on histidine were performed with 1.3 and 0.7 mm MAS probes spinning at $v_R = 40$ or 62.5 kHz, with defocusing and refocusing delays equal to $\tau = 375$ or 384 µs, respectively. The rf-field of the ${}^{1}\text{H} \pi/2$ and π pulses, which do not belong to the recoupling scheme, was equal to 200 kHz, that of the continuous wave irradiation to

- 330 100 kHz, and that of the ¹⁵N pulses to 62 kHz. ¹H decoupling with an rf-field of 16 kHz was applied during the acquisition. The pulses on the ¹H channel were applied on resonance, whereas those on the ¹⁵N channel were applied at the isotropic chemical shift of the ¹⁵NH^τ signal (172 ppm). These 1D spectra resulted from averaging 8 transients with a relaxation delay of 3 s. The ¹⁵N isotropic chemical shifts were referenced to an aqueous saturated solution of NH₄NO₃ using [¹⁵N]-glycine as a secondary reference.
- ¹H→²⁷Al RINEPT-CWc and PRESTO-III experiments on γ-alumina and AlPO₄-14 were performed with a 1.3 mm MAS probe spinning at v_R = 20 (to test the RN^ν_n schemes with large rf-field requirement) or 62.5 kHz. The tested recoupling schemes are listed in Tables 1 and 2 for v_R = 20 kHz and Tables 3 and 4 for v_R = 62.5 kHz. The rf-field of the ¹H π/2 and π pulses, which do not belong to the recoupling scheme, was equal to 208 kHz, that of the continuous wave irradiation to 147 kHz, and the ²⁷Al CT-selective one for π/2 and π pulses to 10 kHz. The defocusing and refocusing delays τ are given in Tables 1 to 4.
 The pulses on the ¹H channel were applied on resonance, whereas those on ²⁷Al channel were applied (i) on resonance with
- AlO₆ signal of γ -alumina in Figs.4 and 7, Tables 1 and 3, as well as in Figs.5 and 8 when the offset is null, (ii) on resonance with AlO₄ signal of AlPO₄-14 in Figs.S2 and S4, Tables 2 and 4 as well as in Figs.S3 and S5 when the offset is null, and (iii) in the middle of the AlO₄ and AlO₆ peaks for the 1D spectra shown in Figs.3 and 6. These differences in offset explain some changes in the relative efficiencies of the recoupling between the figures. These 1D spectra resulted from averaging 64

transients with a relaxation delay of 1 s. The ²⁷Al isotropic chemical shifts were referenced at 0 ppm to 1 mol.L⁻¹ [Al(H₂O)₆]³⁺ solution.

We also measured the decay of the transverse proton magnetization of AlPO₄-14 during a spin echo sequence, in which the refocusing π -pulse was identical to that used in the defocusing part of the RINEPT-CWc sequences (Fig. 1b). This decay was measured at $v_R = 20$ and 62.5 kHz either without any recoupling or by applying a SR4²₁ recoupling built from (180₀), (270₀90₁₈₀) and tt pulses during the delays of the spin echo sequence. The rf fields during the recoupling two blocks were equal

to their optimal values given in Tables 2 and 4. We also acquired several 2D ¹H→²⁷Al *D*-HETCOR spectra of AlPO₄-14 using RINEPT-CWc-SR4²₁ with (180₀), (270₀90₁₈₀) and tt pulses as well as PRESTO-R16⁶₇(270₀90₁₈₀). These 2D spectra were acquired using a non-uniform sampling (NUS) with an exponentially biased sampling retaining 25% of the points with respect to uniform sampling. The 2D spectra

resulted from 8 transients for each of the 500 t_1 increments with a recycle delay of 1 s, i.e. an acquisition time of 72 min.

IV-2. Optimal adiabatic recoupling

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Figs.2e and f show the efficiency of the ${}^{1}\text{H} \rightarrow {}^{15}\text{N}$ RINEPT-SR4²₁(tt) transfer for histidine as function of the $\nu_{1,max}/\nu_R$ and $\Delta\nu_{0,max}/\nu_R$ ratios for $\nu_R = 40$ or 62.5 kHz, respectively. These experimental data indicate that at higher MAS frequencies, an efficient adiabatic recoupling can be achieved for lower $\nu_{1,max}/\nu_R$ and $\Delta\nu_{0,max}/\nu_R$ ratios. This result agrees with the numerical simulations of Figs.2b and d.



Figure 3: ²⁷Al 1D spectra of γ -alumina at 18.8 T with $v_R = 20$ (a) and 62.5 (b) kHz acquired using ¹H \rightarrow ²⁷Al transfers with RINEPT-CWc and SR4²₁(tt), SR4²₁(270₀90₁₈₀) and R12⁵₃ (270₀90₁₈₀), or PRESTO and (a) R22⁷₂(180₀) or R18⁵₂ (180₀), or (b) R16⁶₇ (270₀90₁₈₀) and R14⁵₆ (270₀90₁₈₀). The τ delays and $v_1/v_{1,max}$ rf-fields were fixed to their optimum values given in Tables 1 and 3.

365 IV-3. PRESTO and RINEPT performances for $v_R = 20$ kHz

IV-3-1. γ-alumina

The 1D spectra of γ -alumina acquired using ¹H \rightarrow ²⁷Al RINEPT and PRESTO sequences, shown in Fig.3, exhibit two resonances at 70 and 10 ppm, assigned to tetra- (AlO₄) and hexa-coordinated (AlO₆) resonances, respectively (Morris and Ellis, 1989). The signal of penta-coordinated (AlO₅) sites, which are mainly located in the first surface layer, is barely detected because of the lack of sensitivity of conventional solid-state NMR spectroscopy (Lee et al., 2014). The most intense peak, AlO₆, was used to compare the transfer efficiencies of RINEPT and PRESTO sequences with different recoupling schemes. Table 1 lists the measured performances of ¹H \rightarrow ²⁷Al RINEPT-CWc and PRESTO transfers using various recoupling for γ alumina at v_R = 20 kHz. We notably compared the PRESTO sequences using R22⁷₂(180₀) and R18⁵₂(180₀) recoupling with the RINEPT-CWc scheme using SR4²₁ and R12⁵₃ with: single (180₀), composite (270₀90₁₈₀) and (90₋₄₅90₄₅90₋₄₅) or (tt) adiabatic

375 pulses. A low transfer efficiency was obtained for RINEPT-CWc-SR4²₁(90₀240₉₀90₀) because of its low scaling factor, $\kappa = 0.131$, and hence its performances are not reported in Table 1. We also tested the recoupling schemes based on the symmetry

SC2₁⁰, corresponding to the $[C2_1^0]_0[C2_1^0]_{120}[C2_1^0]_{240}$ sequence with a basic element $C = (90_{45}90_{135}90_{45}90_{225}90_{315}90_{225})$, or C6₃⁰ built from $C' = (90_{30}90_{120}90_{30}90_{240}90_{330}90_{240})$. These basic elements, which derive from $(90_{-45}90_{45}90_{-45})$, have recently been proposed (Perras et al., 2019). As seen in Table 1 and Fig.3a, the sequences yielding the highest transfer efficiencies are by decreasing order RINEPT-CWc with SR4₁²(tt) or R12₃⁵(tt) > PRESTO-R22₂⁷(180₀) > RINEPT-CWc-SR4₁²(270₀90₁₈₀) \approx

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PRESTO-R18 $_{2}^{5}(180_{0})$ > RINEPT-CWc-R12 $_{3}^{5}(270_{0}90_{180})$. Figs. 4 and 5 display the signal intensity of these sequences as function of the rf-field amplitude and offset, respectively.

The highest transfer efficiencies are obtained with the RINEPT-CWc sequence incorporating a (tt) adiabatic pulse. This recoupling also leads to the highest robustness to offset and rf inhomogeneity, and $SR4_1^2$ (tt) and $R12_3^5$ (tt) yield identical transfer efficiency and robustness. Hence, the three-step multiple-quantum super-cycle of the $SR4_1^2$ symmetry does not improve the

robustness in the case of a (tt) basic element. However, these recoupling schemes require maximum rf fields of $v_{1,max} \ge 8v_R = 160$ kHz, which may exceed the rf power specifications of most 3.2 mm MAS probes.

The PRESTO sequences using $R22_2^7(180_0)$ and $R18_2^5(180_0)$ recoupling also result in good transfer efficiencies, but lower than RINEPT-CWc-SR4_1^2(tt). However, they use rf-fields of $v_1/v_R = 5.5$ and 4.5, which are compatible with the specifications of 3.2 mm MAS probes. The higher transfer efficiency of $R22_2^7(180_0)$ with respect to $R18_2^5(180_0)$ stems from its weaker

second-order cross-terms between ¹H-¹H interactions (Table S1).

The efficiency of the RINEPT-CWc-SR4²₁(270₀90₁₈₀) sequence, with $v_1 = 4v_R$, is comparable to that of PRESTO-R18⁵₂(180₀), but with a higher robustness to offset and rf inhomogeneity. We can notice that amplitude modulated recoupling schemes, for which the phase shifts are equal to 180°, such as SR4²₁(270₀90₁₈₀) and SR4²₁(180₀), exhibit a high robustness to rf

field maladjustments (Fig.5) (Carravetta et al., 2000). The use of (270_090_{180}) composite pulses with SR4²₁ symmetry, instead of single π pulses, improves its transfer efficiency as well as its robustness to offset and rf field inhomogeneity.

In summary, for $v_R = 20$ kHz in γ -alumina, the RINEPT-CWc-SR4²₁(270₀90₁₈₀) sequence achieves efficient and robust transfers of magnetization from protons to ²⁷Al nuclei using a moderate rf field of $v_1 = 4v_R$. For ¹H spectra with a width smaller than 20 kHz and MAS probes with a good rf-homogeneity, PRESTO-R22⁷₂(180₀) can result in slightly higher transfer efficiencies.



Figure 4: ²⁷AlO₆ on-resonance signal of γ -alumina at $v_R = 20$ kHz as function of v_1 or $v_{1,max}$ for PRESTO-R22⁷₂(180₀) and -R18⁵₂(180₀) as well as RINEPT-CWc-SR4²₁ (tt), -SR4²₁ (270₀90₁₈₀) and -R12⁵₃ (270₀90₁₈₀). For each curve, τ was fixed to its optimum value given in Table 1.



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Figure 5: ²⁷AlO₆ signal of γ -alumina at $\nu_R = 20$ kHz as function of offset for PRESTO-R22⁷₂(180₀) and -R18⁵₂(180₀) as well as RINEPT-CWc-SR4²₁ (tt), -SR4²₁ (270₀90₁₈₀) and -R12⁵₃ (270₀90₁₈₀). For each curve, τ and ν_1 or $\nu_{1,max}$ were fixed to their optimum values given in Table 1.

PRESTO /RINEPT	Recoupling	τ /µs	$\frac{\nu_{l}/\nu_{l,max}}{/kHz}$	AlO ₆ ^a	$\Delta v_0{}^b$ /kHz	$\Delta v_0 / v_1$	$\Delta v_1^{\ c}$ /kHz	$\Delta \nu_{l}/\nu_{l}$
RINEPT	$SR4_1^2(tt)$	400	160	1	110	0.68	> 100 ^d	> 0.62
	R12 ⁵ ₃ (tt)	400	160	1	110	0.68	$> 100^{d}$	> 0.62
PRESTO	$R22_{2}^{7}(180_{0})$	400	110	0.73	30	0.27	39	0.35
RINEPT	SR4 ² ₁ (270 ₀ 90 ₁₈₀)	400	80	0.63	50	0.63	44	0.55
PRESTO	$R18_{2}^{5}(180_{0})$	400	90	0.61	28	0.31	27	0.30
	$R12_{3}^{5}(270_{0}90_{180})$	400	80	0.50	40	0.50	35	0.44
	$SR4_1^2(90_{-45}90_{45}90_{-45})$	400	63	0.42	14	0.22	14	0.22
	$SR4_1^2(180_0)$	400	45	0.40	17	0.38	24	0.53
RINEPT	$R12_{3}^{5}(180_{0})$	400	45	0.35	10	0.22	15	0.33
	$R12_{3}^{5}(90_{-45}90_{45}90_{-45})$	400	66	0.35	11	0.17	18	0.27
	$SC2_1^0(C)$	400	63	0.31	14	0.22	45	0.71
	C6 ⁰ ₃ (<i>C</i> ')	400	66	0.28	10	0.15	40	0.60

Table 1. Comparison of the performances of ${}^{1}\text{H} \rightarrow {}^{27}\text{Al}$ RINEPT-CWc and PRESTO transfers using various recouplings for AlO₆ signal of γ -alumina at $v_R = 20$ kHz.

^a AlO₆ signal normalized to that with ${}^{1}\text{H}\rightarrow{}^{27}\text{Al}$ RINEPT-CWc-SR4²₁(tt). ^b FWHM of the robustness to offset. ^c FWHM of the robustness to rf-field. ^d Only a lower bound of rf-field could be determined due to probe rf specifications (Fig.4).

415 IV-3-2. AIPO4-14

Fig.6a shows the ${}^{1}\text{H} \rightarrow {}^{27}\text{Al}$ RINEPT and PRESTO 1D spectra of AlPO₄-14 recorded with $v_R = 20$ kHz. They exhibit three ${}^{27}\text{Al}$ resonances at 43, 21 and -2 ppm assigned to AlO₄, AlO₅ and AlO₆ sites, respectively.(Ashbrook et al., 2008) The AlO₅ and AlO₆ sites are directly bonded to OH groups. The ${}^{1}\text{H}$ MAS spectrum is shown in Fig.S1. According to the literature, the ${}^{27}\text{AlO}_4$ signal subsumes the resonances of two AlO₄ sites with quadrupolar coupling constants $C_Q = 1.7$ and 4.1 MHz, whereas those of AlO₅ and AlO₆ sites are equal to 5.6 and 2.6 MHz, respectively (Fernandez et al., 1996; Antonijevic et al., 2006). The ${}^{1}\text{H} {}^{-1}\text{H}$ dipolar couplings within the isopropylamine template molecule are larger than in γ -alumina. We used the most intense peak, AlO₄, to compare the ${}^{1}\text{H} \rightarrow {}^{27}\text{Al}$ transfer efficiencies of RINEPT-CWc and PRESTO sequences with different recoupling schemes, and the results are given in Table 2. The six sequences yielding the highest transfer efficiencies are the same as for γ -alumina and their relative efficiencies are comparable for the AlO₄ peak of AlPO₄-14 and the AlO₆ signal of γ -alumina.

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Figure 6: ²⁷Al 1D spectra of AlPO₄-14 at 18.8 T with $v_R = 20$ (a) and 62.5 (b) kHz acquired using ¹H \rightarrow ²⁷Al transfers with RINEPT-CWc and SR4²₁(tt), SR4²₁(270₀90₁₈₀) and R12⁵₃(270₀90₁₈₀), or PRESTO and (a) R22⁷₂(180₀) and R18⁵₂ (180₀), or (b) R16⁶₇ (270₀90₁₈₀) and R14⁵₆ (270₀90₁₈₀). The τ delays and $v_1/v_{1,max}$ rf fields were fixed to their optimal values given in Tables 2 and 4. The resonance at *ca*. 11 ppm in (a) is due to an impurity.

Nevertheless, the rf requirement of the SR4²₁(tt) and R12⁵₃(tt) schemes is higher for AlPO₄-14 than for γ -alumina because of the larger ¹H-¹H dipolar couplings, in agreement with the numerical simulations of Figs.2a-c. This rf requirement prevents the use of these adiabatic recoupling schemes at v_R = 20 kHz with most 3.2 mm MAS probes. That of the other sequences, and their robustness to offset and rf-fields homogeneity are similar for both samples (Table 2 and Figs.S2 and S3).

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In the case of AlPO₄-14, PRESTO yields a higher efficiency than RINEPT for AlO₅ and AlO₆, contrary to the AlO₄ resonance, since (i) these Al sites are directly bonded to OH groups and (ii) $R22_2^7(180_0)$ and $R18_2^5(180_0)$ schemes are subject to dipolar truncation (section II-1-1), which prevents to transfer the ¹H magnetization of these OH groups to ²⁷AlO₄ nuclei.

Hence, at $v_R = 20$ kHz, for both AlPO₄-14 and γ -alumina, the RINEPT-CWc-SR4²₁(270₀90₁₈₀) and PRESTO-R22⁷₂(180₀) sequences are the best choices to transfer the ¹H magnetization to ²⁷Al nuclei.

PRESTO	Recoupling	τ	$\tau = v_1/v_{1,max}$		Intensity ^a			A., /	Δv_1	A., /
/RINEPT		/µs	/kHz	AlO ₆	AlO ₅	AlO ₄	/kHz	$\Delta v_0 / v_1$	/kHz	$\Delta v_1 / v_1$
DINEDT	$SR4_1^2(tt)$	800	208	1	1	1	120	0.58	_b	_ b
KINLF I	R12 ⁵ ₃ (tt)	800	208	0.99	0.99	0.98	120	0.58	_ b	_ b
PRESTO	$R22_{2}^{7}(180_{0})$	600	114	1.54	1.07	0.67	26	0.23	38	0.33
RINEPT	SR4 ² ₁ (270 ₀ 90 ₁₈₀)	800	77	0.72	0.65	0.67	45	0.58	48	0.62
PRESTO	$R18_{2}^{5}(180_{0})$	600	94	1.45	1.03	0.62	25	0.27	26	0.28
	$R12_{3}^{5}(270_{0}90_{180})$	800	77	0.58	0.50	0.48	46	0.60	36	0.47
	$SR4_1^2(180_0)$	600	43	0.64	0.45	0.36	14	0.33	23	0.53
	$SR4_1^2(90_{-45}90_{45}90_{-45})$	800	61	0.56	0.43	0.25	16	0.26	20	0.32
RINEPT	$SC2_1^0(C)$	800	68	0.54	0.41	0.24	18	0.26	52	0.73
	$R12_3^5(90_{-45}90_{45}90_{-45})$	600	61	0.43	0.30	0.21	8	0.13	18	0.29
	$R12_{3}^{5}(180_{0})$	600	45	0.34	0.28	0.21	8	0.18	18	0.40
	$C6_{3}^{0}(C')$	600	68	0.52	0.36	0.21	10	0.15	42	0.61

Table 2. Comparison of the performances of ${}^{1}H \rightarrow {}^{27}Al$ RINEPT-CWc and PRESTO transfers with AlPO₄-14 at v_R = 20 kHz.

^a Intensities of AlO₆, AlO₅ and AlO₄ resonances normalized to their intensities with ¹H \rightarrow ²⁷Al RINEPT-CWc-SR4²₁(tt). ^b FWHM of the robustness to rf-field was not measured for RINEPT-SR4²₁(tt) and -R12⁵₃(tt) (Fig. S2).

IV-4. PRESTO and RINEPT performances for $v_R = 62.5$ kHz

Similar comparisons of the performances of the various RINEPT-CWc and PRESTO sequences were performed for γ -alumina and AlPO₄-14 at $\nu_R = 62.5$ kHz.

IV-4-1. γ-alumina

450 The corresponding data for γ -alumina are given in Table 3. The sequences yielding the highest transfer efficiencies are by decreasing order: RINEPT-CWc with SR4²₁(tt) or R12⁵₃(tt) > RINEPT-CWc-SR4²₁(270₀90₁₈₀) \approx PRESTO-R16⁶₇(270₀90₁₈₀) > PRESTO-R14⁵₆(270₀90₁₈₀) > RINEPT-CWc-R12⁵₃(270₀90₁₈₀).

The nominal rf requirements of the RINEPT sequences using adiabatic or (270_090_{180}) composite π -pulses correspond to $v_{1max} \approx 5v_R$ (313 kHz: Fig.2d) or $4v_R$ (250 kHz), which exceed the specifications of our 1.3 mm MAS probe, and the sequences were

tested only up to $v_{1max} = 208$ kHz (Fig.7). This suboptimal rf field may limit the transfer efficiencies of these sequences. The PRESTO-R16⁶₇(270₀90₁₈₀) and -R14⁵₆(270₀90₁₈₀) sequences yield transfer efficiencies comparable to those of RINEPT-CWc-SR4²₁(270₀90₁₈₀), but with a significantly lower rf field, $v_1 \approx 137$ kHz $\approx 2.3v_R$. Furthermore, the robustness to offset of these PRESTO sequences is comparable to that of RINEPT-CWc-SR4²₁(270₀90₁₈₀) (Fig.8). PRESTO-R22³₄(180₀) and



Figure 7: ²⁷AlO₆ on-resonance signal of γ -alumina at $\nu_R = 62.5$ kHz as function of ν_1 or $\nu_{1,max}$ for PRESTO-R16⁶₇(270₀90₁₈₀) and -R14⁵₆(270₀90₁₈₀) as well as RINEPT-CWc-SR4²₁ (tt), -SR4²₁ (270₀90₁₈₀) and -R12⁵₃ (270₀90₁₈₀). For each curve, τ was fixed to its optimum value given in Table 3.



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Figure 8: ²⁷AlO₆ signal of γ -alumina at $\nu_R = 62.5$ kHz as function of offset for PRESTO-R16⁶₇(270₀90₁₈₀) and -R14⁵₆(270₀90₁₈₀) as well as RINEPT-CWc-SR4²₁ (tt), -SR4²₁ (270₀90₁₈₀) and -R12⁵₃ (270₀90₁₈₀). For each curve, τ and ν_1 or $\nu_{1,max}$ were fixed to their optimum values given in Table 3.

	PRESTO/RINEPT	Recoupling	τ /µs	$\nu_1/\nu_{1,max}$ /kHz	AlO ₆ ^a	Δv_0 /kHz	$\Delta v_0 / v_1$	Δv_1 /kHz	$\Delta v_1 / v_1$
475		$SR4_1^2(tt)$	256	208	1	74	0.36	_b	- ^b
	RINEPT	R12 ⁵ ₃ (tt)	256	208	1	74	0.36	_ b	_ b
		SR4 ² ₁ (270 ₀ 90 ₁₈₀)	320	208	0.92	96	0.46	_ b	_ b
480	DDESTO	R16 ⁶ ₇ (270 ₀ 90 ₁₈₀)	448	137	0.91	90	0.66	42	0.31
-00	PRESIO	$R14_{6}^{5}(270_{0}90_{180})$	384	146	0.86	100	0.68	38	0.26
	RINEPT	$R12_{3}^{5}(270_{0}90_{180})$	320	208	0.82	86	0.41	_ ^b	_ b
495		$SR4_1^2(180_0)$	320	125	0.75	52	0.42	88	0.70
485		$R12_{3}^{5}(180_{0})$	288	125	0.74	16	0.13	85	0.68
	DDESTO	$R22_4^3(180_0)$	256	157	0.67	68	0.43	20	0.13
	PRESIO	$R16_3^2(180_0)$	384	155	0.51	48	0.31	40	0.26
490		$SC2_1^0(C)$	256	186	0.34	50	0.27	84	0.45
	DINIEDT	$C6_{3}^{0}(C')$	256	186	0.34	43	0.23	76	0.41
495	KINEF I	$SR4_1^2(90_{-45}90_{45}90_{-45})$	256	186	0.32	47	0.25	70	0.38
		$R12_3^5(90_{-45}90_{45}90_{-45})$	256	186	0.32	40	0.22	70	0.38

470	Table 3. Comparison of the performances of ${}^{1}H \rightarrow {}^{27}Al$ RINEPT-CWc and PRESTO transfers using various recouplings for the AlO
	signal of γ -alumina at $v_R = 62.5$ kHz.

AlO₆ signal normalized to that with ${}^{1}\text{H} \rightarrow {}^{27}\text{Al}$ RINEPT-CWc-SR4²₁(tt).

^b FWHM of the robustness to rf-field was not measured for RINEPT-SR4²₁(tt) and -R12⁵₃(tt) (Fig.7).

500 **IV-4-2. AIPO**₄**-1**4

In the case of AlPO₄-14, the relative transfer efficiencies for 27 AlO₄ species follow a similar order as for γ -alumina, except that the transfer efficiencies of PRESTO-R16 ${}^{6}_{7}(270_{0}90_{180})$ and -R14 ${}^{5}_{6}(270_{0}90_{180})$ are significantly lower than that of RINEPT-CWc-SR4 ${}^{2}_{1}(270_{0}90_{180})$ (Table 4). This decreased efficiency of the PRESTO schemes for AlO₄ stems notably from the dipolar truncation, which prevents the transfer of magnetization from the OH groups bonded to AlO₅ and AlO₆ sites to AlO₄, since

505 these ${}^{27}\text{AlO}_4$ nuclei are significantly more distant to protons (see Table S5). Furthermore, the amplitude-modulated SR4 ${}^2_1(270_090_{180})$ recoupling benefits from a higher robustness to rf-field inhomogeneity than the PRESTO schemes (Fig.S4). Conversely, the robustness to offset of these three sequences are comparable (Fig.S5), whereas the rf requirements of R16 ${}^6_7(270_090_{180})$ and R14 ${}^5_6(270_090_{180})$ are much lower than that of SR4 ${}^2_1(270_090_{180})$.

In summary, at $v_R = 62.5$ kHz, for both γ -alumina and isopropylamine-templated AlPO₄-14, PRESTO-R16⁶₇(270₀90₁₈₀) and

510 RINEPT-CWc-SR4 $_{1}^{2}(270_{0}90_{180})$ are the best methods to transfer the polarization of protons to quadrupolar nuclei. However, the first sequence requires a much lower rf-field than the second does.

Table 4. Comparison of the performances of ${}^{1}\text{H} \rightarrow {}^{27}\text{Al}$ RINEPT-CWc and PRESTO transfers using various recouplings for AlPO₄-14 at $v_R = 62.5$ kHz.

PRESTO	Decoupling	τ	$v_1/v_{1,max}$	Intensity ^a			Δv_0	A /	Δv_1	A /
/RINEPT	Recoupting	/µs	/kHz	AlO ₆	AlO ₅	AlO ₄	/kHz	$\Delta \mathbf{v}_0 / \mathbf{v}_1$	/kHz	$\Delta v_1/v_1$
	$SR4_1^2(tt)$	480	208	1	1	1	48	0.23	_ b	_ b
DINIEDT	R12 ⁵ ₃ (tt)	480	208	1.07	1	1.06	44	0.21	_ b	_ ^b
KINEPI	SR4 ² ₁ (270 ₀ 90 ₁₈₀)	480	208	1.05	0.95	0.97	85	0.41	90	0.43
	$R12_{3}^{5}(270_{0}90_{180})$	480	208	0.91	0.84	0.91	80	0.38	68	0.33
PRESTO	R16 ⁶ ₇ (270 ₀ 90 ₁₈₀)	672	146	1.71	1.21	0.76	80	0.55	50	0.34
	$R14_{6}^{5}(270_{0}90_{180})$	576	146	1.72	1.27	0.76	86	0.59	45	0.31
DUEDE	$SR4_1^2(180_0)$	480	129	0.84	0.79	0.75	48	0.37	64	0.49
KINEF I	$R12_{3}^{5}(180_{0})$	480	136	0.72	0.67	0.74	18	0.13	54	0.40
DDESTO	$R22_4^3(180_0)$	512	157	1.47	1.18	0.69	60	0.38	20	0.33
PRESIO	$R16_3^2(180_0)$	480	147	1.17	0.83	0.52	64	0.44	20	0.31
	$R12_3^5(90_{-45}90_{45}90_{-45})$	256	190	0.48	0.27	0.14	32	0.17	75	0.39
RINEPT	$C6^0_3(C')$	256	193	0.47	0.28	0.14	28	0.15	78	0.40
	$SR4_1^2(90_{-45}90_{45}90_{-45})$	256	196	0.48	0.14	0.14	36	0.18	77	0.39
	$SC2_1^0(C)$	256	188	0.53	0.25	0.14	44	0,23	80	0.43

^a Intensities of AlO₆, AlO₅ and AlO₄ resonances normalized to their intensities with ${}^{1}\text{H} \rightarrow {}^{27}\text{Al RINEPT-CWc-SR4}_{1}^{2}(tt)$. ^b FWHM of the robustness to rf-field was not measured for RINEPT-SR4}{}^{2}(tt) and -R12}{}^{5}(tt) (Fig.S4).

IV-5. Decay of transverse ¹H magnetization during recoupling

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We also measured the decay of the ¹H transverse magnetization during a spin-echo experiment, in which the refocusing π pulse was the composite one employed in the defocusing part of the RINEPT-CW sequence shown in Fig. 1b. We performed these experiments on AlPO₄-14 since the ¹H-¹H dipolar interactions are larger in this sample than in γ -alumina. This decay was measured either in the absence of any recoupling or under a **SR4**²₁ recoupling built from (180₀), (270₀90₁₈₀) or tt inversion element. The three ¹H signals featured a mono-exponential decay with a time constant *T*₂' reported in Table 5. At v_R = 20 kHz, the *T*₂' constants are significantly shorter under **SR4**²₁(180₀) and **SR4**²₁(270₀90₁₈₀) than without recoupling.

This faster decay can stem from the reintroduction of ¹H-¹H dipolar interactions in the second- and higher-order terms of the

525 AH by the recoupling as well as the effect of pulse transients (Wittmann et al., 2016). Conversely, the T_2 ' constants under **SR4**²₁(tt) are much longer than without recoupling, showing that the adiabatic pulses using large rf field efficiently decouple the ¹H-¹H dipolar interactions, whereas the continuous variation of the phase and amplitude during these pulses minimizes the transients.

At $v_R = 62.5$ kHz, the T_2 ' constants without recoupling are lengthened with respect to those at $v_R = 20$ kHz since faster 530 MAS better averages the ¹H-¹H dipolar interactions (Mao et al., 2009). Conversely, the T_2 ' constants under SR4²₁(270₀90₁₈₀) recoupling are shorter at $v_R = 62.5$ than at 20 kHz. This counter-intuitive reduction may stem from the shorter pulse lengths at $v_R = 62.5$ kHz, which results in a larger number of transients. For the same reason, the T_2 ' constants under SR4²₁(180₀) are only slightly longer at high MAS frequency. The T_2 ' constants under SR4²₁(tt) recoupling are much shorter at $v_R = 62.5$ than at 20 kHz because the adiabaticity criterion is not fulfilled at $v_R = 62.5$ kHz and hence, the elimination of ¹H-¹H dipolar interactions

535 is less effective (Figs.2f and S4).

Table 5. ¹H T_2 ' values of AlPO₄-14 without recoupling or with SR4²₁ recoupling built from (180₀), (270₀90₁₈₀) or tt inversion element. The estimated error bars are equal to 7 %.

v _R (kHz)		20			62.5	
<i>T</i> ₂ ' (ms)	$\mathrm{NH_{3}^{+}}$	СН	$CH_3 + OH$	$\mathrm{NH_{3}^{+}}$	СН	$CH_3 + OH$
No recoupling	1.6	1.6	1.4	4.0	4.2	4.4
(180 ₀)	0.6	0.4	0.6	0.9	0.5	0.7
(270_090_{180})	0.8	0.6	0.9	0.5	0.3	0.4
tt	52	1000	170	2.2	2.7	2.1



Figure 9: ¹H-²⁷Al *D*-HETCOR 2D spectrum of AlPO₄-14, along with its skyline projections, at $B_0 = 18.8$ T and $v_R = 62.5$ kHz acquired in only 72 min with only ca. 2.5 µL of active volume with NUS 25% using RINEPT-CWc-SR4²₁(270₀90₁₈₀) transfer.

Fig. 9 demonstrates the possibility to acquire 2D ${}^{1}\text{H}{}^{27}\text{Al}$ *D*-HETCOR spectra using RINEPT-CWc-SR4²₁(270₀,90₁₈₀) transfer at v_R = 62.5 kHz. This spectrum was recorded using a NUS scheme retaining 25% of the t₁ points, which would be acquired using uniform sampling. In this spectrum, the CH proton only correlates with the AlO₄ site since it is too distant from AlO₅ and AlO₆ sites (see Table S5). The other two ${}^{1}\text{H}$ signals correlate with the three Al environments. Similar 2D spectra (not shown) were acquired using RINEPT-CWc transfer based on SR4²₁(180₀) and SR4²₁(tt) recoupling as well as PRESTO-R16⁶₇(270₀90₁₈₀). Their skyline projections are shown in Figs. S6 and S7.

575 V. Conclusions

In this work, we have introduced novel symmetry-based hetero-nuclear dipolar recoupling schemes, which can be incorporated into the RINEPT and PRESTO sequences to transfer the magnetization from protons to half-integer quadrupolar nuclei at v_R = 20 or 62.5 kHz. These new recouplings have been compared to the existing ones. We have shown that the RINEPT-CWc-SR4²₁(tt) sequence with adiabatic pulses, which produces efficient and robust transfers at $v_R \approx 10-15$ kHz,(Nagashima et al., 2020) requires rf-fields incompatible with the specifications of most MAS probes for $v_R \ge 20$ kHz. Conversely, the introduced 580 RINEPT-CWc-SR4 $_{1}^{2}(270_{0}90_{180})$ and PRESTO-R22 $_{2}^{2}(180_{0})$ techniques with rf-fields of ca. $4v_{R}$ and $5.5v_{R}$, respectively, are the methods of choice at $v_R = 20$ kHz to transfer the magnetization from protons to quadrupolar nuclei. At $v_R = 62.5$ kHz, the RINEPT-CWc-SR4²₁ (270₀90₁₈₀) and PRESTO-R16⁶₇ (270₀90₁₈₀) sequences with rf-requirements of ca. 4v_R and 2.3v_R, respectively, result in the most robust and efficient transfers. At both MAS frequencies, the RINEPT and PRESTO techniques 585 complement each other since the latter is dipolar truncated, whereas the former is not. As a result, the RINEPT sequences must be chosen to observe simultaneously protonated and unprotonated sites, whereas the PRESTO schemes can be employed for the selective observation of quadrupolar nuclei in proximity to protons. These techniques are expected to be useful for transferring the DNP-enhanced magnetization of protons to quadrupolar nuclei in indirect MAS DNP experiments at $v_R \ge 20$ kHz, notably used at high magnetic fields (Nagashima et al., n.d., 2020; Rankin et al., 2019; Berruver et al., 2020), We also 590 show that they can be used to correlate the NMR signals of protons and quadrupolar nuclei at high MAS frequencies.

Author contributions: JSG, AGMR and JT carried out the NMR experiments on γ-alumina and AlPO₄-14. YT performed the spin dynamics simulations and carried out the NMR experiments on l-histidine HCl. OL derived average Hamiltonian theory for the investigated recoupling sequences. OL and JPA wrote the manuscript. All the authors contributed to the editing of the manuscript.

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