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QCPMG-MAS NMR of Half-Integer Quadrupolar Nuclei

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By combination of fast magic-angle spinning (MAS) and detection of the free-induction decay during a rotor-synchronized quadrupolar Carr–Purcell–Meiboom–Gill (QCPMG) train of refocusing pulses, the sensitivity of quadrupolar-echo MAS NMR spectra for the central transition of half-integer quadrupolar nuclei exhibiting large quadrupolar couplings may be significantly enhanced. Enhancements by an order of magnitude may easily be realized while maintaining information about the anisotropic interactions. In the present study the so-called QCPMG-MAS experiment is demonstrated experimentally and by numerical simulations for the two 87Rb sites in Rb2SO4.

Considerable interest in unraveling the combined effect of different anisotropic nuclear spin interactions in solid-state NMR spectroscopy has characterized the development of new experimental methods in recent years. In spectra of quadrupolar (I > ½) nuclei the combined effect of the quadrupole coupling and chemical shielding interaction may appear. Both of these interactions reflect the local electronic environment of the quadrupolar nucleus. This situation has created renewed interest in the development of new and improved experimental and numerical methods for accurate determination of magnitudes and relative orientation for the two interactions, in particular for powders. For weak quadrupolar couplings (CQ/(2I – 1) < 0.5 MHz, where CQ is the quadrupole coupling constant) accurate parameters characterizing the two tensors may be obtained from one-pulse magic-angle spinning (MAS) NMR spectra of powders with special attention being paid to, e.g., symmetric excitation/detection of all transitions and highly stabilized spinning speed (1, 2). In the case of stronger quadrupole coupling (0.5 MHz < CQ/(2I – 1) < 5 MHz) similar information may be obtained from the second-order quadrupolar-broadened central (½, ½) transition for half-integer quadrupolar nuclei in static powder (3–6) or MAS (7, 8) experiments. For larger quadrupolar couplings second-order broadening for the central transition on the order of hundreds of kilohertz generally reduces the spectral sensitivity.

We recently introduced the QCPMG experiment, i.e., the quadrupolar version (9, 10) of the Carr–Purcell–Meiboom–Gill (CPMG) (11, 12) experiment, as a means to increase the sensitivity of quadrupolar-echo (QE) spectra (13–15) for static powders. Increases in sensitivity by an order of magnitude have been observed while maintaining information about magnitudes and relative orientation of chemical shielding and quadrupole coupling tensors (16). Previously, this experiment has been used to distinguish inhomogeneous from homogeneous interactions in solid-state NMR (9, 10, 17–25), where the latter category include homonuclear dipolar coupling and molecular dynamics. For broad powder patterns induced by inhomogeneous second-order quadrupolar coupling, it should be possible to further improve the sensitivity and resolution by combining the QCPMG experiment with MAS. The obvious reason is that the second-order quadrupolar interaction is partially averaged by MAS—for infinitely fast spinning the width of the central transition lineshape is reduced by a factor of 72(1 + ηQ/6)2/(7(ηQ2 + 22ηQ + 25)) (ranging from 0.29 to 0.41), where ηQ is the asymmetry parameter of the quadrupole coupling tensor (26). To exploit this possibility, this Communication introduces the so-called QCPMG-MAS experiment as a method for improving the sensitivity of MAS NMR of half-integer quadrupolar nuclei while retaining information about the tensors for the anisotropic interactions.

Figure 1 shows the timing diagram of the QCPMG-MAS pulse sequence. The pulse sequence consist of three basic elements. Part A is a standard rotor-synchronized solid echo (13–15) through which the decaying part of the initial spin echo is acquired to form the first part of the free-induction decay (FID). Part B represents the QCPMG part of the pulse sequence which through interrupted sampling appends the full spin-echo envelopes generated by the train of M refocusing π pulses to the FID. Finally, part C extends the sampling period to allow full decay of the FID. To ensure coincidence of rotary and spin echoes (i.e., an integral number of rotor periods per spin-echo period) the pulse sequence must be rotor synchronized. This is accomplished using τ1 = τπ −...
FIG. 1. Timing scheme of the QCPMG-MAS pulse sequence for sensitivity-enhanced QE-MAS NMR of nuclei exhibiting large anisotropic interactions. Following an initial solid echo with sampling of the decaying part of the echo (A), the remaining part of the free-induction decay (FID) is sampled during a rotor-synchronized train of $M \pi$ pulses (B) ending with an extended acquisition period (C) ensuring full decay of the FID. For selective operation on the $(\frac{1}{2}, -\frac{1}{2})$ transition for half-integer quadrupolar nuclei exhibiting large quadrupole coupling interactions the pulse flip angles are scaled by $1 / (1 / 2)$ relative to the nominal flip angles written above the pulses. Rotor synchronization requires $T_y = T_r + \tau_s$ and $2N_T = T_z$, where $N$ is an integer, $T_r$ is the rotor period, and $\tau_s$ and $\tau_v$ are the durations of selective $\pi/2$ and $\pi$ pulses.

$T_{s/2}$ and adjustment of the duration of the repeating unit according to $2N_T = T_s + 2 \tau_s + \tau_v$, where $N$ is an integer, $\tau_s = 2\pi / \omega_l$, is the rotor period, $\tau_v$ is a delay preventing increased further at the expense of details in the spectral gain focused on in this work result from additional manipulation of the spin part of the Hamiltonian by a train of selective $\pi$ pulses. The actual sensitivity gain and the “digitilization” of the envelope of the rotary sidebands containing the information on the anisotropic interaction parameters may be specified experimentally by adjusting the integral number of rotor echoes within each spin-echo period $T_z$. To demonstrate the performance of the QCPMG-MAS experiment, Fig. 2 compares $^{87}$Rb ($I = \frac{3}{2}$) spectra of the two central transitions for Rb$_2$SO$_4$ corresponding to standard QE-MAS and QCPMG-MAS experiments. The experimental spectra were recorded on a Varian UNITY-400 spectrometer (9.4 T, 130.8 MHz for $^{87}$Rb) using a homebuilt high-speed MAS probe (31) with a 5-mm Si$_3$N$_4$ rotor. The experiments used a spinning frequency $\omega_s / 2\pi = 13,645$ Hz and for the QCPMG-MAS experiment the spin-echo parameters were adjusted to split the second-order spinning sideband powder patterns into spin-echo sidebands separated by 250 Hz. In this case the QCPMG-MAS experiment provides a gain in signal intensity by a factor of 7 relative to the standard QEMAS experiment while the essential features of the original centerband and spinning sideband lineshapes for the two rubidium sites in Rb$_2$SO$_4$ are clearly maintained as envelopes of spin-echo sidebands. Obviously, the sensitivity can be increased further at the expense of details in the spectral envelope. In addition to the direct gain in sensitivity, it is noted that the QCPMG-MAS experiment clearly facilitates discrimination of true sideband signals from noise/background as compared to the standard MAS or QE-MAS methods (see insets in Fig. 2).

X-ray diffraction shows that Rb$_2$SO$_4$ has orthorhombic crystal structure with space group $Pnam$ (32) and thus has two magnetically nonequivalent rubidium sites in agreement with the spectra in Fig. 2. Parameters describing magnitudes and relative orientation of the quadrupole coupling and chemical shielding tensors for the two sites have previously been determined using static-powder QE (5), MAS (8), and single-crystal (33) NMR. It should be emphasized, however, that the weak chemical shielding interaction is easily averaged by MAS, implying that accurate information about this interaction and the relative tensor orientations in this case requires application of single-crystal NMR. Information about the anisotropic interactions (for high-speed MAS specifically the quadrupolar coupling) may be obtained from the sideband manifolds observed for the two sites in the QCPMG-MAS spectrum of Fig. 2b. This is demonstrated by the simulated MAS (or QE-MAS) and QCPMG-MAS spectra in Figs. 2c and 2d calculated using the parameters determined by single-crystal NMR (33) and under the assumption of ideal RF pulse performance. The MAS and QCPMG-MAS spectra were calculated on a Silicon Graphics Onyx computer using a MAS version of the software earlier described for the calculation of static-powder QCPMG spectra (16). Clearly, the simulated MAS and QCPMG-MAS...
In conclusion, we have demonstrated that the QCPMG technique, recently proposed for sensitivity enhancement of static-powder spectra (16), may advantageously be combined with fast MAS. The resulting spectra show significant sensitivity enhancements relative to standard MAS or QE-MAS spectra while maintaining sufficient spectral information to extract accurate parameters for the anisotropic interaction tensors. The actual sensitivity gain (here a factor of 7) may be increased considerably for larger quadrupolar coupling interactions, where standard MAS experiments even for nuclei with high relative sensitivity and short longitudinal relaxation times (such as $^{87}$Rb) become increasingly time demanding. Relying on detection of the central transition alone, it is anticipated that the QCPMG-MAS experiment is experimentally feasible even in the regime of very large quadrupolar coupling interactions. In this case, however, numerical analysis of the QCPMG-MAS spectra need consideration of finite RF pulse effects as will be described elsewhere.

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**REFERENCES**