



Cite this: *Phys. Chem. Chem. Phys.*,
2024, 26, 17666

Pulsed dynamic nuclear polarization: a comprehensive Floquet description†

Gian-Marco Camenisch, ^a Nino Wili, ^b Gunnar Jeschke ^{*a} and
Matthias Ernst ^{*a}

Dynamic nuclear polarization (DNP) experiments using microwave (mw) pulse sequences are one approach to transfer the larger polarization on the electron spin to nuclear spins of interest. How the result of such experiments depends on the external magnetic field and the excitation power is part of an ongoing debate and of paramount importance for applications that require high chemical-shift resolution. To date numerical simulations using operator-based Floquet theory have been used to predict and explain experimental data. However, such numerical simulations provide only limited insight into parameters relevant for efficient polarization transfer, such as transition amplitudes or resonance offsets. Here we present an alternative method to describe pulsed DNP experiments by using matrix-based Floquet theory. This approach leads to analytical expressions for the transition amplitudes and resonance offsets. We validate the method by comparing computations by these analytical expressions to their numerical counterparts and to experimental results for the XiX, TOP and TPPM DNP sequences. Our results explain the experimental data and are in very good agreement with the numerical simulations. The analytical expressions allow for the discussion of the scaling behaviour of pulsed DNP experiments with respect to the external magnetic field. We find that the transition amplitudes scale inversely with the external magnetic field.

Received 30th April 2024,
Accepted 5th June 2024

DOI: 10.1039/d4cp01788a

rsc.li/pccp

1 Introduction

Dynamic nuclear polarization (DNP) has become a routine method to increase the nuclear polarization of solid samples by transferring the higher electron polarization to the nuclei of interest typically at cryogenic temperatures.^{1,2} Two different implementations of DNP are currently in wide-spread use: (i) magic-angle spinning (MAS) DNP operated typically at around 100 K to investigate materials, surfaces, and biomolecules.^{2–5} (ii) Dissolution DNP operated typically at around 1 K to generate highly polarized small molecules for metabolic imaging or spectroscopic applications.^{6–11} Both modes of DNP use continuous-wave (CW) irradiation of microwaves (mw) on the sample to generate a high polarization on the nuclei. In MAS DNP, polarization transfer to protons is the most common mode followed by a subsequent polarization-transfer step to other nuclei. In dissolution DNP, direct enhancement of ¹³C is quite common but transfer to protons with subsequent

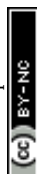
polarization transfer to other nuclei has also been implemented. A number of different mechanisms (Overhauser effect (OE),¹² solid effect (SE),^{13–16} cross effect (CE),^{17–19} and thermal mixing (TM)^{20,21}) have been identified that explain the polarization transfer. They all have in common that increasing static magnetic fields lead to a decrease of the polarization transfer efficiency. Typically, they are described in an incoherent way using arguments that rely on the saturation of certain transitions and relaxation properties.

Some time ago, the nuclear orientation *via* electron spin locking (NOVEL) sequence was introduced where the microwave nutation frequency is matched to the nuclear Zeeman frequency.^{22–25} Both fields modulate the pseudo-secular part of the hyperfine coupling leading to a time-independent effective Hamiltonian if the two frequencies are matched. A number of additional pulsed polarization-transfer techniques for DNP have been introduced recently that can also be described by a coherent effective time-averaged Hamiltonian in a suitable frame of reference.^{26–30} They all rely on modulation by a pulse sequence that in some way matches the nuclear Larmor frequency, thus generating a time-independent part of the Hamiltonian from the pseudo-secular part of the hyperfine interaction that promotes polarization transfer. Different irradiation schemes have been proposed but this principal mode of operation is the same for all of them.

^a Department of Chemistry and Applied Biosciences, ETH Zürich, Zürich, Switzerland. E-mail: gjeschke@ethz.ch, maer@ethz.ch

^b Interdisciplinary Nanoscience Center (iNANO) and Department of Chemistry, Aarhus University, Aarhus, Denmark

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4cp01788a>



In this publication, we analyze pulsed DNP sequences in more detail, using two different approaches. In a first approach, we use operator-based Floquet theory to calculate effective Hamiltonians at the resonance conditions that drive the double-quantum or zero-quantum polarization transfer.^{31,32} This provides theoretical understanding under which conditions we can expect DNP to happen and allows to estimate the efficiency. In a second, complementary approach, we use matrix-based Floquet theory^{33–36} to connect the Fourier coefficients of the pulse sequence to the various resonance conditions and to get a better understanding of the scaling behavior of pulsed DNP as a function of the static magnetic field.

2 Theoretical concepts

We will first discuss pulsed DNP sequences in the framework of operator-based Floquet theory (Section 2.1) which gives effective Hamiltonians that promote polarization transfer at resonance conditions. These effective Hamiltonians can be calculated only numerically due to the complexity of the interaction-frame transformation that hides the direct connection between the Fourier coefficients of the microwave irradiation and the transition probabilities. In a complementary treatment in the framework of classical matrix-based Floquet theory (Section 2.3), analytical diagonalization in connection with perturbation theory can be used to establish this connection. This treatment allows a better understanding of transition amplitudes and their dependence on the experimental parameters like microwave amplitude and static magnetic field.

2.1 Operator-based Floquet theory

In this section, we present an operator-based Floquet theory formalism³¹ to describe pulsed DNP. The formalism introduced in this section is similar to the one presented in ref. 29 and we repeat some of the results but in a slightly different notation. The two-spin Hamiltonian for a spin system consisting of one electron spin (S) in the Zeeman rotating frame and one nuclear spin (I) in the laboratory frame is given as

$$\hat{\mathcal{H}}(t) = \Omega_{0,S}\hat{S}_z + A_{zz}\hat{S}_z\hat{I}_z + B\hat{S}_z\hat{I}_x + \omega_{0,I}\hat{I}_z + \mathcal{H}_{\mu w}^{(S)}(t), \quad (1)$$

where $\Omega_{0,S} = \omega_{0,S} - \omega_{\mu w}$ is the offset of the microwave irradiation, A_{zz} the secular and $B = \sqrt{A_{zx}^2 + A_{zy}^2}$ the pseudo-secular part of the hyperfine coupling. The electron resonance frequency is given by $\omega_{0,S} = g_{zz}\frac{\mu_B}{\hbar}B_0$ and for the nucleus by $\omega_{0,I} = -\gamma_I B_0$ while $\omega_{\mu w}$ is the microwave frequency and B_0 is the external static magnetic field applied along the laboratory z -axis. The orientation-dependent g_{zz} value determines the frequency offset from the mw irradiation. For low static magnetic fields, the isotropic value g_{iso} can be used instead. The Hamiltonian $\mathcal{H}_{\mu w}^{(S)}(t)$ refers to the Hamiltonian of the μw irradiation and can be written as

$$\mathcal{H}_{\mu w}^{(S)}(t) = \omega_{1,S}(t)\{\cos[\phi(t)]\hat{S}_x + \sin[\phi(t)]\hat{S}_y\}, \quad (2)$$

where $\omega_{1,S}(t)$ is the time-dependent pulse amplitude and $\phi(t)$ the time-dependent phase. To describe the effect of the microwave

irradiation, it is best to transform the Hamiltonian of eqn (1) into an interaction frame with respect to $\Omega_{0,S}\hat{S}_z + \mathcal{H}_{\mu w}^{(S)}(t)$. Such a transformation can be analytically written as

$$\begin{aligned} \mathcal{H}^I(t) &= \hat{U}_S^{-1} \left(\hat{\mathcal{H}} - \Omega_{0,S}\hat{S}_z - \mathcal{H}_{\mu w}^{(S)}(t) \right) \hat{U}_S \\ &= \sum_{\chi=x,y,z} a_{z\chi}^{(S)}(t) \hat{S}'_{\chi} (A_{zz}\hat{I}_z + B\hat{I}_x) + \omega_{0,I}\hat{I}_z, \end{aligned} \quad (3)$$

with $\hat{U}_S = \hat{T} \exp \left(-i \int_0^t \Omega_{0,S}\hat{S}_z + \mathcal{H}_{\mu w}^{(S)}(\tau) d\tau \right)$ where $\hat{T}$ is the Dyson time-ordering operator³⁷ and $a_{z\chi}^{(S)}(t)$ are the time-dependent elements of a 3×3 rotation matrix. Note that in the following we drop the “dash” on the S spin operators when referring to the interaction frame. We consider time-periodic irradiation schemes with a modulation period τ_m with $\mathcal{H}_{\mu w}^{(S)}(t) = \mathcal{H}_{\mu w}^{(S)}(t + \tau_m)$. This results in a modulation frequency

$$\omega_m = \frac{2\pi}{\tau_m}. \quad (4)$$

Operator-based Floquet theory can then be used to describe the effect of the irradiation. The interaction-frame transformation, as given in eqn (3), corresponds to rotation around an effective field axis in the spin space of the electron spin. The effective frequency is

$$\omega_{\text{eff},S}(\Omega_{0,S}) = \frac{\beta_{\text{eff},S}}{\tau_m}, \quad (5)$$

where $\beta_{\text{eff},S}$ describes the rotation angle around the effective field axis within one period τ_m . Note the functional dependence of $\omega_{\text{eff},S}$ on the offset frequency $\Omega_{0,S}$. The Fourier coefficients of the Hamiltonian can be calculated from the rotation-matrix elements $a_{z\chi}^{(S)}(t)$ as

$$a_{z\chi}^{(k,\ell)} = \frac{1}{\tau_m} \int_{-\infty}^{\infty} a_{z\chi}^{(S)}(t) e^{-ik\omega_m t} e^{-i\ell\omega_{\text{eff},S} t} dt. \quad (6)$$

Now we transform the Hamiltonian into a frame rotating with the nuclear Larmor frequency $\omega_{0,I}$ and insert the Fourier coefficient of eqn (6) for the electron spin. This leads to

$$\begin{aligned} \mathcal{H}^I(t) &= \sum_{k=-\infty}^{\infty} \sum_{\ell=-1}^1 \left\{ \left[\frac{A_{zz}}{2} \left(a_+^{(k,\ell)} \hat{S}^+ + a_-^{(k,\ell)} \hat{S}^- + 2a_{zz}^{(k,\ell)} \hat{S}_z \right) \hat{I}_z \right. \right. \\ &\quad \left. \left. + \frac{B}{4} \left(a_+^{(k,\ell)} \hat{S}^+ + a_-^{(k,\ell)} \hat{S}^- + 2a_{zz}^{(k,\ell)} \hat{S}_z \right) \right. \right. \\ &\quad \left. \left. \times \left(\hat{I}^+ e^{-i[\omega_{0,I} t]} + \hat{I}^- e^{i[\omega_{0,I} t]} \right) \right] e^{ik\omega_m t} e^{i\ell\omega_{\text{eff},S} t} \right\} \\ &= \sum_{n=-1}^1 \sum_{k=-\infty}^{\infty} \sum_{\ell=-1}^1 \mathcal{H}^{\hat{n},(k,\ell)} e^{in[\omega_{0,I} t]} e^{ik\omega_m t} e^{i\ell\omega_{\text{eff},S} t} \end{aligned} \quad (7)$$

where $a_{\pm}^{(k,\ell)} = a_{zx}^{(k,\ell)} \mp ia_{zy}^{(k,\ell)}$ and $\hat{S}^{\pm} = \hat{S}_x \pm i\hat{S}_y$. Please note that we assumed a negative Larmor frequency for the nuclear spin corresponding to a positive gyromagnetic ratio. The Hamiltonian in eqn (7) contains three fundamental frequencies $|\omega_{0,I}|$, $\omega_{\text{eff},S}$ and ω_m and we can write it as a Fourier series (see last line in eqn (7)). The first-order effective Hamiltonian based on triple-mode



Floquet theory is, thus, given by

$$\begin{aligned}\mathcal{H}_{\text{eff}}^{(1)} &= \mathcal{H}^{(0,0,0)} + \sum_{n_0, k_0, \ell_0} \mathcal{H}^{(n_0, k_0, \ell_0)} \\ &= \frac{A_{zz}}{2} \left(a_+^{(0,0)} \hat{S}^+ + a_-^{(0,0)} \hat{S}^- + 2a_{zz}^{(0,0)} \hat{S}_z \right) \hat{I}_z \\ &\quad + \sum_{n_0, k_0, \ell_0} \mathcal{H}^{(n_0, k_0, \ell_0)}\end{aligned}\quad (8)$$

In eqn (8) n_0 , k_0 and ℓ_0 sum over only the values where the resonance condition

$$n_0 |\omega_{0,I}| + k_0 \omega_m + \ell_0 \omega_{\text{eff},S} = 0 \quad (9)$$

is fulfilled. Polarization transfer between the electron spin (S) and the nuclear spin (I) can only be mediated by the double-quantum (DQ) terms ($\hat{S}^\pm \hat{I}^\pm$) or the zero-quantum (ZQ) terms ($\hat{S}^\pm \hat{I}^\mp$). Therefore, if we now define a tuple $(-1, k_0, \ell_0)$ and its negative $(1, -k_0, -\ell_0)$, we obtain a first-order effective Hamiltonian for the ZQ and DQ-transfer given by

$$\mathcal{H}_{\text{ZQ}}^{(1)} = \frac{B}{4} \left(a_-^{(k_0, \ell_0)} \hat{S}^- \hat{I}^+ + a_+^{(-k_0, -\ell_0)} \hat{S}^+ \hat{I}^- \right) \quad (10)$$

$$\mathcal{H}_{\text{DQ}}^{(1)} = \frac{B}{4} \left(a_+^{(k_0, \ell_0)} \hat{S}^+ \hat{I}^+ + a_-^{(-k_0, -\ell_0)} \hat{S}^- \hat{I}^- \right) \quad (11)$$

Note that for a negative gyromagnetic ratio (positive Larmor frequency) for the nuclear spin the sign for the indices k_0 and ℓ_0 changes, e.g., $a_-^{(k_0, \ell_0)} \hat{S}^- \hat{I}^+$ in case of a negative Larmor frequency is $a_-^{(-k_0, -\ell_0)} \hat{S}^- \hat{I}^+$ in case of a positive Larmor frequency. The polarization transfer $\rho_0 \rightarrow \hat{I}_z$ can be calculated following ref. 29 by

$$\begin{aligned}\langle \hat{I}_z(t) \rangle &= \frac{\gamma_e}{\gamma_I} \langle \rho_0 | \hat{S}_z \rangle \text{Tr} \{ U_{\text{ZQ/DQ}} \hat{S}_z U_{\text{ZQ/DQ}}^{-1} \hat{I}_z \} / \text{Tr} \{ \hat{I}_z^2 \} \\ &= \pm \frac{\gamma_e}{\gamma_I} \langle \rho_0 | \hat{S}_z \rangle \sin^2 \left(\frac{B}{4} \sqrt{a_+^{(\mp k_0, \mp \ell_0)} a_-^{(\pm k_0, \pm \ell_0)}} t \right),\end{aligned}\quad (12)$$

where $U_{\text{ZQ/DQ}} = \exp(-i\mathcal{H}_{\text{ZQ/DQ}}^{(1)}t)$ and the upper binary operator in this equation refers to the ZQ case and the lower one to the DQ case, respectively. Again, eqn (12) is valid for the case of a positive gyromagnetic ratio of the nuclear spin (negative Larmor frequency). For the case of a negative gyromagnetic ratio of the nuclear (positive Larmor frequency) the sign for the indices k_0 and ℓ_0 changes, e.g. for the ZQ case $a_+^{(-k_0, -\ell_0)}$ would change to $a_+^{(+k_0, +\ell_0)}$. Also note that in case of a positive gyromagnetic ratio of the nuclear spin the ZQ enhancement gets negative and the DQ enhancement positive due to the ratio $\frac{\gamma_e}{\gamma_I}$.

However, for a negative gyromagnetic ratio of the nuclear spin the ZQ enhancement is positive and the DQ enhancement is negative. The factor $\langle \rho_0 | \hat{S}_z \rangle$ takes into account that only the part of the electron polarization that is along the effective field can be transferred to the nuclei. For small times t and using $a_\pm^{(k_0, \ell_0)} = \left(a_\mp^{(-k_0, -\ell_0)} \right)^*$ the performance of a certain pulse scheme can be roughly estimated by²⁹

$$\varepsilon_{\text{ZQ/DQ}} = \mp \langle \rho_0 | \hat{S}_z \rangle a_{\text{eff}} \quad (13)$$

where

$$a_{\text{eff}} = \begin{cases} \left| a_+^{(-k_0, -\ell_0)} \right|, & \text{for ZQ condition} \\ \left| a_+^{(k_0, \ell_0)} \right|, & \text{for DQ condition} \end{cases} \quad (14)$$

For CW irradiation (time-independent pulse amplitude $\omega_{1,S}$ and phase offset ϕ in eqn (2)), we can calculate the Fourier coefficients of the Hamiltonian of eqn (7) analytically which leads to an effective Hamiltonian relevant for DNP of the form

$$\frac{B}{4} \sin(\theta) (\hat{S}^- \hat{I}^+ + \hat{S}^+ \hat{I}^- - \hat{S}^+ \hat{I}^+ - \hat{S}^- \hat{I}^-), \quad (15)$$

where $\theta = \arctan\left(\frac{\omega_{1,S}}{\Omega_{0,S}}\right)$ describes the direction of the effective field in the electron spin coordinate system (see Fig. S1 in ESI† Section A). In case of the solid effect¹⁶ we can invoke the small-angle approximation ($\Omega_{0,S} \gg \omega_{1,S}$), where $\sin(\theta) \approx \frac{\omega_{1,S}}{\Omega_{0,S}}$ leads to a

prefactor of $\frac{B\omega_{1,S}}{4\omega_{0,I}}$. For NOVEL^{23,25} $\theta = \frac{\pi}{2}$ and the prefactor is $\frac{B}{4}$.

Section A of the ESI† contains the derivation to calculate the Fourier coefficients of the Hamiltonian analytically starting from eqn (6). The effective Hamiltonian of eqn (15) as well as the scaling factor for NOVEL and SE were already reported in ref. 38.

2.2 Microwave irradiation scheme

So far we did not specify the microwave irradiation scheme characterized by $\mathcal{H}_{\text{mw}}^{(S)}(t)$ in eqn (1) in detail. In order to apply Floquet theory the only requirement is that the mw irradiation is time periodic with a modulation period τ_m . When we limit the discussion to amplitude-modulated monochromatic pulses ($\phi(t) = \phi$), the pulse amplitude $\omega_{1,S}(t)$ becomes the only parameter that has to fulfill the time-periodic restriction. In the following we set $\phi = 0$ for simplicity of the notation. Any time-periodic amplitude modulation can then be represented as a half-range Fourier expansion of the form

$$\begin{aligned}\omega_{1,S}(t) &= \omega_{1,S}^{(\text{max})} \left[\frac{a_0}{2} + \sum_{q=1}^{\infty} \{ a_q \cos(q\omega_m t) + b_q \sin(q\omega_m t) \} \right] \\ &= \frac{\omega_{1,S}^{(\text{max})}}{2} \left[a_0 + \sum_{q=1}^{\infty} \{ (a_q - ib_q) e^{iq\omega_m t} + (a_q + ib_q) e^{-iq\omega_m t} \} \right],\end{aligned}\quad (16)$$

where a_0 , a_q and b_q are the Fourier coefficients of the microwave irradiation scheme and $q \in \mathbb{N}$. $\omega_{1,S}^{(\text{max})}$ is the maximum value of the time-dependent pulse amplitude $\omega_{1,S}(t)$. Note the difference between the Fourier coefficients of the microwave irradiation scheme (a_q and b_q) as given in eqn (16) and the Fourier coefficients of the interaction-frame trajectory ($a_{z,I}^{(k,\ell)}$) as calculated in eqn (6). For CW irradiation all Fourier coefficients in eqn (16) are zero except for a_0 , which is equal to 2 and $\omega_{1,S}(t) = \omega_{1,S}^{(\text{max})}$. For an even pulse amplitude function $\omega_{1,S}(t)$ all odd Fourier coefficients (b_q) are 0. In contrast for an odd function



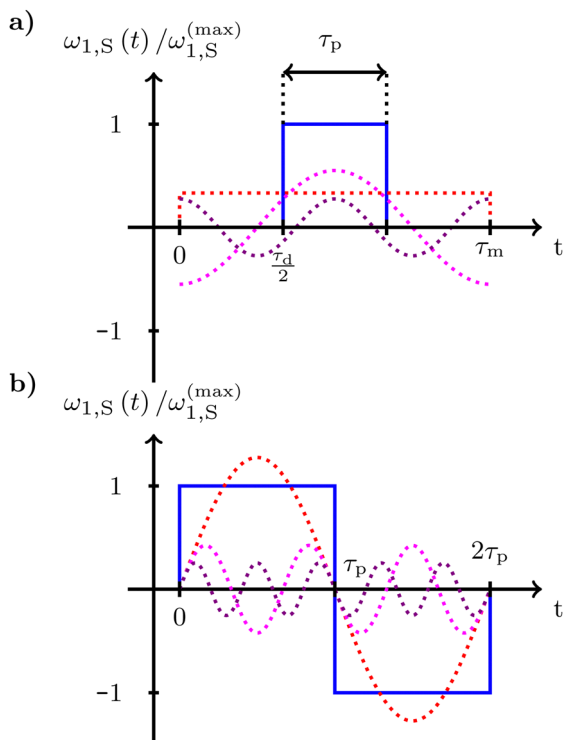


Fig. 1 Half-range Fourier expansion of (a) the TOP DNP pulse scheme and (b) the XiX DNP pulse scheme according to eqn (16). The DNP pulse sequence is represented with a blue solid line. The TOP DNP sequence consists of a pulse with length τ_p and a delay τ_d , whereas the XiX DNP sequence consists of two pulses with opposite phase. The different Fourier modes are shown with dashed lines. In the TOP sequence the unmodulated Fourier mode described by the Fourier coefficient a_0 is represented in dashed red. The first Fourier mode a_1 and the second mode a_2 are represented by magenta and violet dashed lines respectively. A fraction $\frac{\tau_d}{\tau_m} = \frac{2}{3}$ is used for this particular TOP sequence (see Table 1). In the XiX case, the first Fourier mode b_1 , the third mode b_3 and the fifth mode b_5 are represented by red, magenta and violet dashed lines respectively.

$\omega_{1,S}(t)$ all even coefficients are 0, e.g. a_0 and a_q . In the following three sections we will discuss pulsed DNP schemes, first the Time-optimized Pulsed (TOP) DNP²⁷ sequence, second the XiX (X-inverse-X) DNP sequence^{28,29} and last the Two-Pulse Phase Modulation (TPPM) DNP sequence.³⁰ We represent them by the half-range Fourier expansion as given in eqn (16).

2.2.1 TOP pulse scheme. The TOP DNP sequence is presented in Fig. 1(a) and consists of a delay τ_d and a pulse with length τ_p .²⁷ In this representation, the TOP DNP sequence is an even irradiation scheme in the sense of the half-range Fourier expansion which means that all odd Fourier coefficients b_q are 0. The value of a_0 is given for arbitrary values of τ_d and τ_p as

$$a_0 = \frac{2\tau_p}{\tau_p + \tau_d} = \frac{2\tau_p}{\tau_m} \quad (17)$$

The Fourier coefficients a_q for the TOP sequence are obtained as

$$a_q = -\frac{2}{q\pi} \sin\left(\frac{q\pi\tau_d}{\tau_m}\right) \quad (18)$$

One can clearly see from eqn (17) and (18) that the Fourier coefficients of the microwave irradiation scheme depend on the

Table 1 Fourier coefficients for different duty cycles for the TOP DNP sequence. For certain combinations of τ_d , τ_p and q some coefficients a_q are 0. The first entry represents the case where $\tau_d = 0$ e.g. CW irradiation. The Fourier coefficient a_0 is scaled by the maximal possible value obtained by CW which is two

τ_d/τ_p	τ_p/τ_m	τ_d/τ_m	$a_0/2$	a_q
0	1	0	1	0 for all q
1	1/2	1/2	1/2	0 for all $q = 2, 4, 6, 8, \dots$
2	1/3	2/3	1/3	0 for all $q = 3, 6, 9, 12, \dots$
3	1/4	3/4	1/4	0 for all $q = 4, 8, 12, 16, \dots$

duty cycle τ_p/τ_m of the TOP sequence. The coefficient a_0 describes the average field and can have values between zero (no pulse) and two (cw irradiation). The Fourier coefficients a_q for $q \in \mathbb{N}$ exhibit a much more complex behaviour. For certain combinations of the parameters τ_d , τ_p and q certain coefficients a_q can be 0. This is summarized in Table 1 and has important consequences for the discussion in Section 3.3.

2.2.2 XiX pulse scheme. The XiX DNP sequence is shown schematically in Fig. 1(b).^{28,29} Although XiX is a phase-modulated pulse scheme, it can be considered as well as an amplitude modulation scheme with equal pulse length τ_p and inverted amplitude. Such an amplitude modulation can be described by an odd half-range Fourier expansion. The odd coefficients b_q are given by

$$b_q = \begin{cases} \frac{4}{q\pi}, & \text{for odd } q \\ 0, & \text{for even } q \end{cases} \quad (19)$$

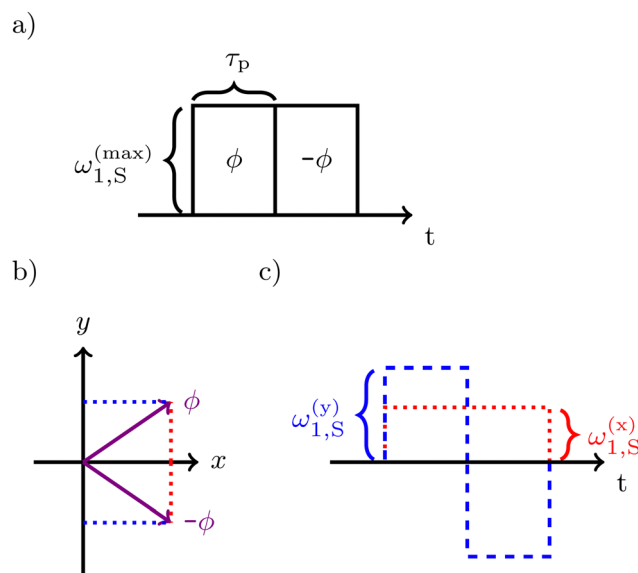


Fig. 2 (a) Schematic representation of the TPPM pulse sequence consisting of two pulses of length τ_p with opposite phases ϕ and $-\phi$, respectively. (b) Cartesian representation of the opposite phases in a TPPM experiment. The x-part of the phase of the two pulses is identical. The y-part of the phase of the two pulses changes sign and results in an XiX type irradiation. (c) The TPPM sequence can be viewed as a combination of an amplitude-modulated XiX pulse sequence with amplitude $\omega_{1,S}^{(y)}$ and phase y (blue dashed lines) and an additional orthogonal CW irradiation with amplitude $\omega_{1,S}^{(x)}$ and phase x (red dotted lines).



and are independent of the pulse length and modulation period. Another difference to the TOP sequence is that the Fourier coefficients b_q are non-zero only for odd q .

2.2.3 TPPM pulse scheme. The TPPM pulse scheme consists of two consecutive pulses with phases $\pm\phi$ (see Fig. 2(a) and (b)). This scheme can be considered as an amplitude-modulated XiX sequence with an additional orthogonal CW part as indicated in Fig. 2(c). This can be easily understood by starting from the Hamiltonian $\hat{\mathcal{H}}_{\mu w}^{(S)}(t)$ as given in eqn (2). By inserting a function $\phi(t)$ switching from $+\phi$ to $-\phi$ at a modulation frequency ω_m we obtain

$$\hat{\mathcal{H}}_{\mu w}^{(S)}(t) = \omega_{1,S}^{(x)} \hat{S}_x + \omega_{1,S}^{(y)}(t) \hat{S}_y \quad (20)$$

with

$$\omega_{1,S}^{(x)} = \cos(\phi) \omega_{1,S}^{(\max)} \quad (21)$$

and

$$\begin{aligned} \omega_{1,S}^{(y)}(t) &= \sin(\phi) \omega_{1,S}^{(\max)} \sum_{q=1}^{\infty} b_q \sin(q\omega_m t) \\ &= \omega_{1,S}^{(\max)} \sum_{q=1}^{\infty} \tilde{b}_q \sin(q\omega_m t) \end{aligned} \quad (22)$$

The coefficients b_q in eqn (22) are the same as for the XiX case as given in eqn (19). Please note that in the TPPM case we multiply coefficients b_q by a factor $\sin(\phi)$ to account for the phase modulation. This results in a new definition $\tilde{b}_q = \sin(\phi)b_q$ as performed in the last step of eqn (22). The amplitude of the CW contribution is given by eqn (21). Hence, due to the effective CW part in TPPM, the a_0 Fourier coefficient of the microwave irradiation is

$$\frac{a_0}{2} = \cos(\phi). \quad (23)$$

This coefficient is not zero unless $\phi = \frac{\pi}{2}$, which corresponds to the XiX limit of TPPM with phase y . Therefore, the TPPM sequence can be viewed as a combination of the CW irradiation and an orthogonal XiX DNP sequence. The time-dependent modulation can be treated in a similar way as in the XiX sequence with the exception that there is an additional time-independent CW part.

2.3 Matrix-based Floquet theory

While operator-based Floquet theory can be used to calculate effective Hamiltonians that describe the polarization transfer in DNP, the direct connection to the various harmonics of the mw irradiation is often lost in the complexities of the interaction-frame transformation. In principle, the connection can be recovered through an analytical Bessel function formulation of the interaction-frame transformation³⁹ but insights are limited due to the complexity of the expressions. In this section we consider matrix-based Floquet theory as an alternative description that provides a direct connection between transition probabilities and the microwave irradiation scheme. The Hamiltonian as defined in eqn (1) can be rewritten with the

amplitude modulation of the mw irradiation schemes as discussed in Section 2.2 to

$$\begin{aligned} \hat{\mathcal{H}}(t) &= \Omega_{0,S} \hat{S}_z + A_{zz} \hat{S}_z \hat{I}_z + B \hat{S}_z \hat{I}_x + \omega_{0,I} \hat{I}_z + \omega_{1,S}^{(\max)} \frac{a_0}{2} \hat{S}_x \\ &+ \frac{\omega_{1,S}^{(\max)}}{2} \sum_{q=1}^{\infty} [(a_q - ib_q) e^{iq\omega_m t} \\ &+ (a_q + ib_q) e^{-iq\omega_m t}] \hat{S}_x. \end{aligned} \quad (24)$$

For the TOP- and XiX DNP sequence $\chi = x$. The analytical expressions for the Fourier coefficients of the TOP sequence can be found in eqn (17) and (18) and for the XiX sequence in eqn (19), respectively. As shown in Fig. 2 the TPPM sequence can also be regarded as an amplitude-modulated pulse sequence with phase $+y$ with an orthogonal CW part with phase $+x$. Therefore, $\chi = y$ in eqn (24) for the TPPM case. In TPPM $\frac{a_0}{2} = \cos(\phi)$ and $a_q = 0 \forall q \neq 0$. The Fourier coefficients are $b_q = \sin(\phi) \frac{4}{q\pi}$ for odd values of q . The Hamiltonian in eqn (24) can be expressed as a Fourier expansion

$$\hat{\mathcal{H}}(t) = \sum_{q'=-\infty}^{+\infty} \hat{\mathcal{H}}^{(q')} e^{iq'\omega_m t} \quad (25)$$

with the Fourier components

$$\hat{\mathcal{H}}^{(0)} = \Omega_{0,S} \hat{S}_z + A_{zz} \hat{S}_z \hat{I}_z + B \hat{S}_z \hat{I}_x + \omega_{0,I} \hat{I}_z + \omega_{1,S}^{(\max)} \frac{a_0}{2} \hat{S}_x \quad (26)$$

and

$$\hat{\mathcal{H}}^{(q')} = \hat{\mathcal{H}}^{(\pm q)} = \frac{1}{2} \omega_{1,S}^{(\max)} (a_{|q'|} \mp ib_{|q'|}) \hat{S}_x \quad q' \in \mathbb{Z}. \quad (27)$$

Be aware that in contrast to the expansion in eqn (16) and (24) the Fourier index q' of the Hilbert space Hamiltonian in eqn (25) can also be negative while $q = |q'|$ is always limited to positive values. Now we can use the Fourier components in eqn (26) and (27) to express the Floquet matrix $\hat{\mathcal{H}}_{\mathcal{F}}$. We thus obtain the elements of the infinite-dimensional Floquet matrix $\hat{\mathcal{H}}_{\mathcal{F}}$ as³⁶

$$\langle \mu, \nu, c_1 | \hat{\mathcal{H}}_{\mathcal{F}} | \kappa, \eta, c_2 \rangle = \hat{\mathcal{H}}_{\mu, \nu, \kappa, \eta}^{(c_1 - c_2)} + c_1 \omega_m \delta_{c_1 c_2} \delta_{\mu \kappa} \delta_{\nu \eta}. \quad (28)$$

In eqn (28) δ_{ij} represents the Kronecker delta. $\hat{\mathcal{H}}_{\mu, \nu, \kappa, \eta}^{(c_1 - c_2)}$ is a specific element of the Floquet matrix, which can be obtained from eqn (26) and (27). In the upcoming sections, we use Greek letters to represent the eigenvalues of the spin operator in Hilbert-space and Latin letters to denote the Fourier harmonics of amplitude modulation of the μw irradiation scheme. Be aware that in eqn (28) the difference $c_1 - c_2$ between two Fourier indices in Floquet space has to match the Fourier index q' in eqn (25)–(27). According to eqn (28) the central blocks of the infinite-dimensional Floquet matrix (blocks where $q' = c_1 - c_2 = 0$) contain the Fourier component $\hat{\mathcal{H}}^{(0)}$ (eqn (26)) with frequency offsets $c_1 \omega_m \mathbb{1}$ (last term of eqn (28)). The off-diagonal blocks (blocks where $q' = c_1 - c_2 \neq 0$) contain the Fourier



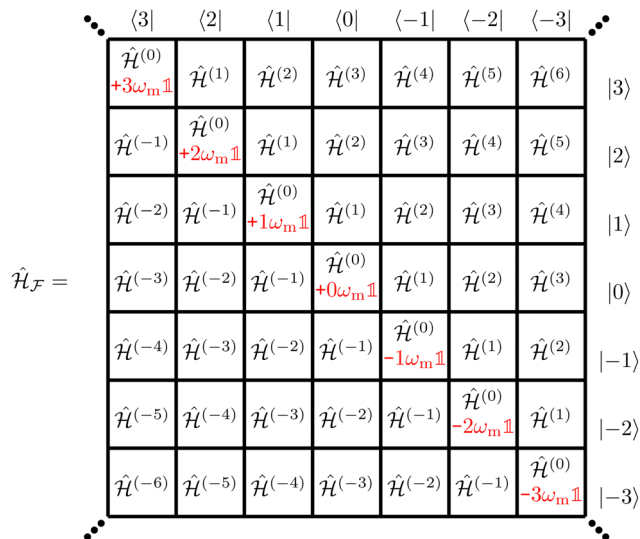


Fig. 3 Schematic representation of the infinite-dimensional Floquet matrix. Each block is represented by the corresponding Fourier component $\hat{\mathcal{H}}^{(q')}$, which is in case of a two-spin system consisting of spin 1/2 particles a 4×4 matrices. The expression $q'\omega_m \mathbb{1}$ in red on the central block $\langle q' | \hat{\mathcal{H}}^{(q')} | q' \rangle$ results from last term of eqn (28).

components $\hat{\mathcal{H}}^{(q')}$ as given in eqn (27). In the upcoming sections we will use the Fourier index q' of the Fourier expansion in Hilbert space to denote connected Fourier states in Floquet space. For a two-spin 1/2 system, the Fourier components $\hat{\mathcal{H}}^{(q')}$ are 4×4 matrices within the basis set $|\kappa, \eta\rangle$ where $\kappa \in \{\alpha_S, \beta_S\}$ and $\eta \in \{\alpha_n, \beta_n\}$. Part of the infinite-dimensional Floquet matrix $\hat{\mathcal{H}}_{\mathcal{F}}$ is presented in Fig. 3.

As outlined in Section 2.2.1 in the TOP DNP sequence the even Fourier coefficients a_0 and a_q are non-zero (see eqn (17) and (18)). Therefore, eqn (27) simplifies to

$$\hat{\mathcal{H}}^{(q')} = \frac{1}{2} \omega_{1,S}^{(\max)} a_{|q'|} \hat{S}_x \quad \forall q' \in \mathbb{Z}. \quad (29)$$

All diagonal blocks in the infinite-dimensional Floquet matrix for the TOP DNP sequence contain the Fourier component $\hat{\mathcal{H}}^{(0)}$ as given in eqn (26). In case of the XiX DNP sequence (see Section 2.2.2) the odd Fourier coefficients b_q are the non-zero ones (see eqn (19)) resulting in

$$\hat{\mathcal{H}}^{(0)} = \Omega_{0,S} \hat{S}_z + A_{zz} \hat{S}_z \hat{I}_z + B \hat{S}_z \hat{I}_x + \omega_{0,I} \hat{I}_z \quad (30)$$

for eqn (26) and in

$$\hat{\mathcal{H}}^{(q')} = \hat{\mathcal{H}}^{(\pm q)} = \begin{cases} \mp \frac{i}{2} \omega_{1,S}^{(\max)} \frac{4}{|q'| \pi} \hat{S}_x, & \text{for odd } q' \\ 0, & \text{for even } q' \end{cases} \quad (31)$$

for eqn (27). According to Section 2.2.3, the TPPM sequence can be regarded as a combination of the CW irradiation scheme and the XiX DNP sequence. The diagonal and off-diagonal blocks in Floquet space are given by the Fourier components in eqn (26) and (27) with phase $+\gamma$, respectively. Inserting the

Fourier coefficients for the TPPM case leads to

$$\hat{\mathcal{H}}^{(0)} = \Omega_{0,S} \hat{S}_z + A_{zz} \hat{S}_z \hat{I}_z + B \hat{S}_z \hat{I}_x + \omega_{0,I} \hat{I}_z + \omega_{1,S}^{(\max)} \cos(\phi) \hat{S}_x \quad (32)$$

for the diagonal block and

$$\hat{\mathcal{H}}^{(q')} = \hat{\mathcal{H}}^{(\pm q)} = \begin{cases} \mp \frac{i}{2} \omega_{1,S}^{(\max)} \sin(\phi) \frac{4}{|q'| \pi} \hat{S}_y, & \text{for odd } q' \\ 0, & \text{for even } q' \end{cases}, \quad (33)$$

for the off-diagonal block, respectively. Compared to the XiX sequence, in the TPPM the phase of the off-diagonal block changes from x to y and the block contains an additional factor $\sin(\phi)$ as mentioned in Section 2.2.3. The diagonal block $\hat{\mathcal{H}}^{(0)}$ in the infinite-dimensional Floquet matrix is given in eqn (26) in its most general analytical form. $\hat{\mathcal{H}}^{(0)}$ is similar for the TOP and the TPPM sequence (except for the definition of Fourier coefficient a_0) and the XiX case differs by a missing term with the $\hat{S}_x$ operator due to $a_0 = 0$. However, we still can define a general diagonalization of $\hat{\mathcal{H}}^{(0)}$ that is valid for all three pulse schemes. This will be further discussed in Section 2.4.

2.4 Diagonalization of the central Floquet block

In order to apply perturbation theory for the side-diagonal parts of the Floquet Hamiltonian that originate from the time-dependent mw irradiation, we first diagonalize the spin-degrees of freedom of the central blocks of the Floquet Hamiltonian $\hat{\mathcal{H}}^{(0)}$ which have the form

$$\hat{\mathcal{H}}^{(0)} = \Omega_{0,S} \hat{S}_z + A_{zz} \hat{S}_z \hat{I}_z + B \hat{S}_z \hat{I}_x + \omega_{0,I} \hat{I}_z + \omega_{1,S}^{(\max)} \frac{a_0}{2} \hat{S}_x. \quad (34)$$

The offset terms $q'\omega_m \mathbb{1}$ do not influence diagonalization, since the unit operator is invariant under rotation. Direct diagonalization of eqn (34) leads to results that are very complex and not easy to interpret. Therefore, $\hat{\mathcal{H}}^{(0)}$ is approximately diagonalized in a two-step approach like outlined in ref. 40. In the first step, the reduced Hamiltonian

$$\hat{\mathcal{H}}_1 = \Omega_{0,S} \hat{S}_z + A_{zz} \hat{S}_z \hat{I}_z + B \hat{S}_z \hat{I}_x + \omega_{0,I} \hat{I}_z \quad (35)$$

is diagonalized exactly and the additional term $\omega_{1,S}^{(\max)} \frac{a_0}{2} \hat{S}_x$ (contained in eqn (34)) is subjected to this first transformation leading to a term of the form $\hat{S}_x \cos(\eta) + 2 \hat{S}_y \hat{I}_y \sin(\eta)$. From this resulting term $\hat{S}_x \cos(\eta)$ is treated as a perturbation in a second unitary transformation (see Fig. S2 in ESI,[†] Section B). All the details of this two consecutively applied unitary transformations are given in ESI,[†] Section B in much more detail and were adapted from ref. 40. Here we present only the relevant results needed for further discussion. The same set of transformations is then applied to the off-diagonal parts of the Floquet Hamiltonian $\hat{\mathcal{H}}^{(q')}$ with $q' \neq 0$ (see eqn (27)). In a final step, we can then apply second-order perturbation theory to calculate the transition frequencies and probabilities enabled by the Fourier components of the various mw irradiation schemes.



The first unitary transformation that diagonalizes the Hamiltonian in eqn (35) is given by

$$\hat{U}_1 = \exp(-i(\eta_x \hat{S}_x \hat{I}_y + \eta_\beta \hat{S}_\beta \hat{I}_y)). \quad (36)$$

Since the first diagonalization takes place in the I -spin sub space, we have to distinguish between a positive Larmor frequency $\omega_{0,I}$ and a negative one to diagonalize eqn (35). In the following we restrict our discussion to negative Larmor frequencies (see Fig. S2 in ESI,† Section B). The angles η_x and η_β describing the first unitary transformation can be obtained from Fig. S2 in ESI,† Section B and are given as

$$\eta_x = \arctan\left(\frac{-B}{2\omega_{0,I} + A_{zz}}\right) \quad (37)$$

$$\eta_\beta = \arctan\left(\frac{-B}{2\omega_{0,I} - A_{zz}}\right). \quad (38)$$

The angles are chosen such that $0^\circ \leq \eta_x, \eta_\beta \leq 90^\circ$. The Hamiltonian $\hat{\mathcal{H}}_1$ of eqn (35) after the first unitary transformation is given by

$$\hat{\mathcal{H}}_1 = \Omega_{0,S} \hat{S}_z + \omega'_{0,I} \hat{I}_z + A' \hat{S}_z \hat{I}_z \quad (39)$$

with

$$\omega'_{0,I} = (\omega_{12} + \omega_{34})/2 \quad (40)$$

$$A' = \omega_{12} - \omega_{34} \quad (41)$$

and the nuclear frequencies

$$\omega_{12} = \left(\omega_{0,I} + \frac{A_{zz}}{2}\right) \cos(\eta_x) - \frac{B}{2} \sin(\eta_x) \quad (42)$$

$$\omega_{34} = \left(\omega_{0,I} - \frac{A_{zz}}{2}\right) \cos(\eta_\beta) - \frac{B}{2} \sin(\eta_\beta). \quad (43)$$

The full diagonal block of the Floquet Hamiltonian (see eqn (34)) contains in addition the spin operator $\hat{S}_x$ which transforms into

$$\hat{U}_1^{-1} \hat{S}_x \hat{U}_1 = \hat{S}_x \cos(\eta) + 2\hat{S}_y \hat{I}_y \sin(\eta), \quad (44)$$

with $\eta = \frac{1}{2}(\eta_x + \eta_\beta)$. The angle 2η is shown in Fig. S2 (ESI†) and can take values from 0° to 180° . The case $2\eta < 90^\circ$ is obtained for spin systems in the weak coupling regime, where $|A_{zz}|, |B| < |\omega_{0,I}|$. $2\eta > 90^\circ$ is attained in the strong-coupling regime, meaning that $|A_{zz}| > |\omega_{0,I}|$. Near the weak-coupling limit, we have $2\eta \approx 0^\circ$ and near the strong-coupling limit, we have $2\eta \approx 180^\circ$. In both these cases, $\sin \eta \approx 0$, meaning the excitation of formally forbidden electron-nuclear transitions is weak. The total Hamiltonian in eqn (34) after the first unitary transformation is thus given by

$$\begin{aligned} \hat{\mathcal{H}}^{(0)'} &= \Omega_{0,S} \hat{S}_z + \omega'_{0,I} \hat{I}_z + A' \hat{S}_z \hat{I}_z \\ &+ \omega_{1,S}^{(\max)} \frac{A_0}{2} (\hat{S}_x \cos(\eta) + 2\hat{S}_y \hat{I}_y \sin(\eta)) \end{aligned} \quad (45)$$

Using eqn (27) and (44) the off-diagonal blocks containing the Fourier components $\hat{\mathcal{H}}^{(q')}$ with $q' \neq 0$ after the first

diagonalization step can be written as

$$\begin{aligned} \hat{\mathcal{H}}^{(q)'} &= \hat{\mathcal{H}}^{(\pm q)'} \\ &= \frac{1}{2} \omega_{1,S}^{(\max)} (a_{|q'|} \mp ib_{|q'|}) (\hat{S}_x \cos(\eta) + 2\hat{S}_y \hat{I}_y \sin(\eta)) \\ &= \frac{1}{2} \omega_{1,S}^{(\max)} (a_{|q'|} \mp ib_{|q'|}) (\hat{S}_x \cos(\eta) \\ &\quad + \frac{1}{2} (\hat{S}^+ \hat{I}^- + \hat{S}^- \hat{I}^+ - \hat{S}^+ \hat{I}^+ - \hat{S}^- \hat{I}^-) \sin(\eta)). \end{aligned} \quad (46)$$

As outlined later in more detail, for spin systems near the weak-coupling limit, the angle η is very small. Therefore, we perform the second diagonalization step only with respect to the $\hat{S}_x$ term, leaving the term $2\hat{S}_y \hat{I}_y \sin(\eta)$ in eqn (45) as a remaining off-diagonal perturbation. We thus diagonalize the Hamiltonian

$$\hat{\mathcal{H}}_{\text{tr}}^{(0)'} = \Omega_{0,S} \hat{S}_z + \omega'_{0,I} \hat{I}_z + A' \hat{S}_z \hat{I}_z + \omega'_{1,S} \hat{S}_x \quad (47)$$

with $\omega'_{1,S} = \omega_{1,S}^{(\max)} \frac{A_0}{2} \cos(\eta)$. Diagonalization of eqn (47) takes place in the subspace of the S -spin. Thus, we need to take into account the sign of $\Omega_{0,S}$ (see Fig. S2 in ESI,† Section B). The upper binary operator in the following equations corresponds to the $+\Omega_{0,S}$ case and the lower one to the $-\Omega_{0,S}$ case, respectively. The unitary transformation to diagonalize eqn (47) is given by

$$\hat{U}_2 = \exp(-i(\pm\theta_x \hat{S}_y \hat{I}^z \pm \theta_\beta \hat{S}_y \hat{I}^\beta)). \quad (48)$$

The angles θ_x and θ_β as shown in Fig. S2 (ESI†) can be calculated as

$$\theta_x = \arctan\left(\frac{\pm\omega'_{1,S}}{\Omega_{0,S} + \frac{A'}{2}}\right) \quad (49)$$

$$\theta_\beta = \arctan\left(\frac{\pm\omega'_{1,S}}{\Omega_{0,S} - \frac{A'}{2}}\right) \quad (50)$$

and can take any values from 0° to 90° . In the doubly tilted frame the truncated Hamiltonian can be expressed as

$$\hat{\mathcal{H}}_{\text{tr}}^{(0)''} = \hat{U}_2^{-1} \hat{\mathcal{H}}_{\text{tr}}^{(0)'} \hat{U}_2 = \Omega''_{0,S} \hat{S}_z + \omega'_{0,I} \hat{I}_z + A'' \hat{S}_z \hat{I}_z \quad (51)$$

with

$$\Omega''_{0,S} = (\omega_{13} + \omega_{24})/2 \quad (52)$$

$$A'' = \omega_{13} - \omega_{24} \quad (53)$$

and the electronic frequencies

$$\omega_{13} = \left(\Omega_{0,S} + \frac{A'}{2}\right) \cos(\theta_x) \pm \omega'_{1,S} \sin(\theta_x) \quad (54)$$

$$\omega_{24} = \left(\Omega_{0,S} - \frac{A'}{2}\right) \cos(\theta_\beta) \pm \omega'_{1,S} \sin(\theta_\beta). \quad (55)$$

The angle $2\theta = (\theta_x - \theta_\beta)$ is depicted in Fig. S2 (ESI†). The spin



operator $\hat{S}_x$ from the off-diagonal parts of the Floquet Hamiltonian $\mathcal{H}^{(q')}$ in the doubly tilted frame can be expressed as

$$\begin{aligned}\mathcal{H}^{(q')''} &= \mathcal{H}^{(\pm q)''} \\ &= \frac{1}{2}\omega_{1,S}^{(\max)}(a_{|q'|} \mp ib_{|q'|})[\cos(\eta) \\ &\quad \times \{\cos(\theta_\alpha)\hat{S}_x\hat{I}^\alpha + \cos(\theta_\beta)\hat{S}_x\hat{I}^\beta \\ &\quad \pm \sin(\theta_\alpha)\hat{S}_z\hat{I}^\alpha \pm \sin(\theta_\beta)\hat{S}_z\hat{I}^\beta\} \\ &\quad + \sin(\eta)(2\hat{S}_y\hat{I}_y\cos(\theta) \pm \hat{I}_x\sin(\theta))]\end{aligned}\quad (56)$$

In this doubly tilted frame, the total Hamiltonian including also the term $2\hat{S}_y\hat{I}_y\sin(\eta)$ as given in eqn (45) can be expressed as

$$\begin{aligned}\mathcal{H}^{(0)''} &= \hat{U}_2^{-1} \left(\mathcal{H}_{\text{tr}}^{(0)'} + 2\hat{S}_y\hat{I}_y\sin(\eta) \right) \hat{U}_2 \\ &= \Omega_{0,S}'\hat{S}_z + \omega_{0,I}'\hat{I}_z + A''\hat{S}_z\hat{I}_z \\ &\quad + \omega_{1,S}'\sin(\eta) \left[\frac{1}{2}(\hat{S}^+\hat{I}^- + \hat{S}^-\hat{I}^+ - \hat{S}^+\hat{I}^+ - \hat{S}^-\hat{I}^-)\cos(\theta) \pm \hat{I}_x\sin(\theta) \right]\end{aligned}\quad (57)$$

Eqn (57) can be used to derive the Fourier coefficients of the Hamiltonian for DNP experiments like solid effect DNP¹⁶ and NOVEL^{23,25} (see eqn (15) and ESI,[†] Section B).

In case of TPPM, the diagonal blocks of the Floquet Hamiltonian in the doubly tilted frame have the same form as given in eqn (57). The modulated part contained in the off-diagonal parts of the Floquet Hamiltonian is phase shifted by 90° and

contains the spin operator $\hat{S}_y$ (see eqn (33)). In the doubly tilted frame we obtain

$$\begin{aligned}\mathcal{H}^{(q')''} &= \mathcal{H}^{(\pm q)''} = \hat{U}_2^{-1}\hat{U}_1^{-1}\mathcal{H}^{(q')}\hat{U}_1\hat{U}_2 \\ &= \mp \frac{i}{2}b_{|q'|}\omega_{1,S}\sin(\phi)\{\cos(\eta)\hat{S}_y \\ &\quad - \sin(\eta)[\cos(\theta^{(+)})2\hat{S}_x\hat{I}_y \\ &\quad + \sin(\theta^{(+)})2\hat{S}_z\hat{I}_y]\}\end{aligned}\quad (58)$$

with $\theta^{(+)} = \frac{1}{2}(\theta_\alpha + \theta_\beta)$. Please note, that we have to set $\frac{a_0}{2} = \cos(\phi)$ for the term $\omega_{1,S}' = \omega_{1,S}^{(\max)}\frac{a_0}{2}\cos(\eta)$ in the TPPM case, *e.g.*, in eqn (49) and (50) for the angles θ_α and θ_β and all other equations containing $\omega_{1,S}'$. The ZQ- and DQ terms are contained in the operator $2\hat{S}_x\hat{I}_y = \frac{i}{2}(\hat{S}^+\hat{I}^- - \hat{S}^-\hat{I}^+ - \hat{S}^+\hat{I}^+ + \hat{S}^-\hat{I}^-)$. In the ESI,[†] Section B we present a detailed step-by-step derivation of this entire section.

3 Results and discussion

3.1 Fourier synthesized DNP

In the following sections we use the Fourier expansion of eqn (16) and the matrix representation of the Floquet Hamiltonian in the electron rotating frame to explain the occurrence of certain resonance peaks in the DNP profile. We artificially set some Fourier coefficients of the microwave irradiation scheme to 0 and see which resonance peaks are left/unchanged compared to the original sequence. For this Fourier-synthesized

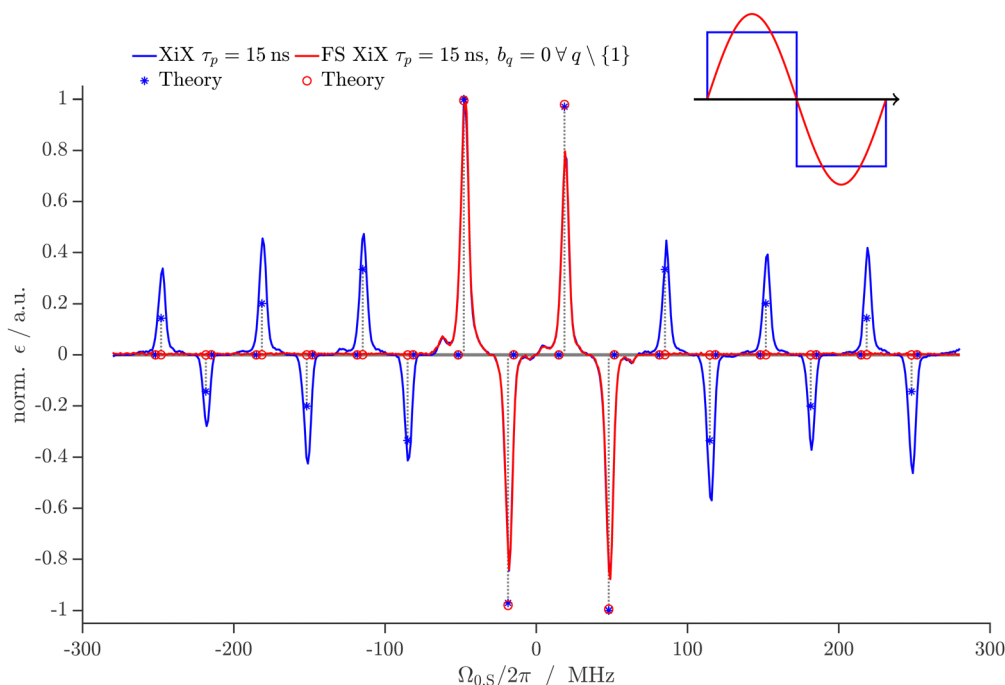


Fig. 4 DNP profile for XiX (blue line) and FS-XiX (red line) recorded at X-band (0.35 T) with a Rabi frequency of $\frac{\omega_{1,S}}{2\pi} = 4$ MHz and a pulse length $\tau_p = 15$ ns (further experimental parameters are given in Section 5). The dots and asterisks represent the estimated enhancement $\epsilon_{\text{ZQ/DQ}}$ as given in eqn (13).



(FS) DNP sequences we use the name FS-XiX in case where the original sequence is the XiX sequence or FS-TOP for the TOP sequence, respectively. In the following Section the original sequence will be always represented by blue color and the FS case by red color. All experiments were conducted with the parameters/procedures as given in Section 5. As a chemical system we used the trityl radical Ox063 dissolved in DNP juice at 80 K which gives high enhancement factors as first described in ref. 24. We discuss the details for the XiX sequence and only give a more summary description for the TOP and TPPM sequence since many of the properties are very similar.

3.2 XiX sequence

3.2.1 DNP profile. The experimental comparison of the DNP profile of the standard XiX DNP sequence with the FS-XiX case for $b_q = 0 \forall q \setminus \{1\}$ at X-band (0.35 T) using $\tau_p = 15$ ns and $\frac{\omega_{1,S}}{2\pi} = 4$ MHz is shown in Fig. 4. Examples for pulse lengths of $\tau_p = 5, 9$ and 45 ns can be found in ESI,[†] Section J. We only discuss the case where we set all Fourier coefficients of the microwave irradiation to 0 except the one for $q = 1$. All others can be converted to this case by scaling the modulation frequency of the XiX DNP sequence to the considered harmonic.

In Fig. 4 we see that only some select resonances remain visible for the FS-XiX sequence. This is confirmed by calculating the transfer efficiency using operator-based Floquet theory. The theoretical values were calculated numerically (see Section 2.1 or ESI,[†] Section L) and are indicated by circles or asterisks in Fig. 4. We can group the resonance conditions ($\Omega_{0,S}$ for which eqn (9) is fulfilled) obtained from numerical calculations using operator-based Floquet theory in three groups, *i.e.*, resonance conditions that are present (non-zero) in the XiX case and the FS-XiX case like $\frac{\Omega_{0,S}}{2\pi} \approx \pm 18$ or ± 48 MHz, resonance conditions that are non-zero in the XiX case, but 0 in the FS-XiX case like $\frac{\Omega_{0,S}}{2\pi} \approx \pm 115$ MHz and resonance conditions that are zero in both cases, *e.g.*, $\frac{\Omega_{0,S}}{2\pi} \approx \pm 81$ MHz. This behavior can be better explained by using the matrix representation of the infinite-dimensional Floquet Hamiltonian as described in Section 2.3. In case of XiX we only need the first diagonalization step from Section 2.4 because $a_0 = 0$. The four eigenvalues can be calculated according to eqn (39) and are given as

$$\varepsilon_{\alpha,\alpha}^{(c_1)} = c_1 \omega_m + \frac{1}{4} \sqrt{1 + \frac{B^2}{(A_{zz} + 2\omega_{0,I})^2}} (A_{zz} + 2\omega_{0,I}) + \frac{\Omega_{0,S}}{2} \quad (59)$$

$$\varepsilon_{\alpha,\beta}^{(c_1)} = c_1 \omega_m - \frac{1}{4} \sqrt{1 + \frac{B^2}{(A_{zz} + 2\omega_{0,I})^2}} (A_{zz} + 2\omega_{0,I}) + \frac{\Omega_{0,S}}{2} \quad (60)$$

$$\varepsilon_{\beta,\alpha}^{(c_1)} = c_1 \omega_m + \frac{1}{4} \sqrt{1 + \frac{B^2}{(-A_{zz} + 2\omega_{0,I})^2}} (-A_{zz} + 2\omega_{0,I}) - \frac{\Omega_{0,S}}{2} \quad (61)$$

$\langle 1 $	$\langle 0 $	$\langle -1 $	
$\begin{pmatrix} \epsilon_{\alpha,\alpha}^{(1)} & 0 & 0 & 0 \\ 0 & \epsilon_{\alpha,\beta}^{(1)} & 0 & 0 \\ 0 & 0 & \epsilon_{\beta,\alpha}^{(1)} & 0 \\ 0 & 0 & 0 & \epsilon_{\beta,\beta}^{(1)} \end{pmatrix}$	$\begin{pmatrix} 0 & 0 & I_{SQ}^{(1)} I_{DQ}^{(1)} \\ 0 & 0 & I_{ZQ}^{(1)} I_{SQ}^{(1)} \\ I_{SQ}^{(1)} I_{ZQ}^{(1)} & 0 & 0 \\ I_{DQ}^{(1)} I_{SQ}^{(1)} & 0 & 0 \end{pmatrix}$	$\begin{pmatrix} 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \end{pmatrix}$	$ 1\rangle$
$\begin{pmatrix} 0 & 0 & I_{SQ}^{(-1)} I_{DQ}^{(-1)} \\ 0 & 0 & I_{ZQ}^{(-1)} I_{SQ}^{(-1)} \\ I_{SQ}^{(-1)} I_{ZQ}^{(-1)} & 0 & 0 \\ I_{DQ}^{(-1)} I_{SQ}^{(-1)} & 0 & 0 \end{pmatrix}$	$\begin{pmatrix} \epsilon_{\alpha,\alpha}^{(0)} & 0 & 0 & 0 \\ 0 & \epsilon_{\alpha,\beta}^{(0)} & 0 & 0 \\ 0 & 0 & \epsilon_{\beta,\alpha}^{(0)} & 0 \\ 0 & 0 & 0 & \epsilon_{\beta,\beta}^{(0)} \end{pmatrix}$	$\begin{pmatrix} 0 & 0 & I_{SQ}^{(1)} I_{DQ}^{(1)} \\ 0 & 0 & I_{ZQ}^{(1)} I_{SQ}^{(1)} \\ I_{SQ}^{(1)} I_{ZQ}^{(1)} & 0 & 0 \\ I_{DQ}^{(1)} I_{SQ}^{(1)} & 0 & 0 \end{pmatrix}$	$ 0\rangle$
$\begin{pmatrix} 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \end{pmatrix}$	$\begin{pmatrix} 0 & 0 & I_{SQ}^{(-1)} I_{DQ}^{(-1)} \\ 0 & 0 & I_{ZQ}^{(-1)} I_{SQ}^{(-1)} \\ I_{SQ}^{(-1)} I_{ZQ}^{(-1)} & 0 & 0 \\ I_{DQ}^{(-1)} I_{SQ}^{(-1)} & 0 & 0 \end{pmatrix}$	$\begin{pmatrix} \epsilon_{\alpha,\alpha}^{(-1)} & 0 & 0 & 0 \\ 0 & \epsilon_{\alpha,\beta}^{(-1)} & 0 & 0 \\ 0 & 0 & \epsilon_{\beta,\alpha}^{(-1)} & 0 \\ 0 & 0 & 0 & \epsilon_{\beta,\beta}^{(-1)} \end{pmatrix}$	$ -1\rangle$

Fig. 5 Part of the infinite-dimensional Floquet matrix of Fig. 3 for the Fourier states $\langle 1 |$, $\langle 0 |$ and $\langle -1 |$. We consider systems consisting of four sub-blocks in Fourier space. The system comprising the Fourier states 0 and -1 is indicated by an orange frame. There arise four different transitions marked in yellow, greenish-blue, brown and purple. These transitions can be further grouped in two ZQ- and two DQ-transitions (see Section 3.2.1). In the system comprising the Fourier states $\langle 1 |$ and $\langle -1 |$ (light-green frames), the four sub-blocks in Fourier space are not connected.

$$\varepsilon_{\beta,\beta}^{(c_1)} = c_1 \omega_m - \frac{1}{4} \sqrt{1 + \frac{B^2}{(-A_{zz} + 2\omega_{0,I})^2}} (-A_{zz} + 2\omega_{0,I}) - \frac{\Omega_{0,S}}{2} \quad (62)$$

In eqn (59)–(62) the superscript in *e.g.* $\varepsilon_{\alpha,\alpha}^{(c_1)}$ denotes the Fourier index, the first subscript index refers to the electron spin state and the second to the nuclear spin state, respectively. For weak irradiation, meaning that $\omega_{1,S} \ll \omega_m$, the off-diagonal blocks that are relevant are the ones that connect almost degenerate Floquet states in the diagonal blocks (see Fig. 5). These off-diagonal blocks contain the transition amplitudes of the corresponding transition. For DQ transitions in Floquet space, *e.g.* ($|\alpha_S, \alpha_I, c_1\rangle \leftrightarrow |\beta_S, \beta_I, c_2\rangle$), this means that

$$\varepsilon_{\alpha,\alpha}^{(c_1)} - \varepsilon_{\beta,\beta}^{(c_2)} = 0 \quad (63)$$

has to hold resulting in the following expression for the microwave offset at resonance

$$\Omega_{0,S}^{(\text{res,DQ})} = (c_2 - c_1) \omega_m - \frac{1}{4} \left[\sqrt{1 + \frac{B^2}{(A_{zz} + 2\omega_{0,I})^2}} (A_{zz} + 2\omega_{0,I}) + \sqrt{1 + \frac{B^2}{(-A_{zz} + 2\omega_{0,I})^2}} (-A_{zz} + 2\omega_{0,I}) \right] \quad (64)$$



For ZQ transitions, we need to consider ($|\alpha_S, \beta_I, c_1\rangle \leftrightarrow |\beta_S, \alpha_I, c_2\rangle$) and we obtain

$$\Omega_{0,S}^{(\text{res,ZQ})} = (c_2 - c_1)\omega_m + \frac{1}{4} \left[\sqrt{1 + \frac{B^2}{(A_{zz} + 2\omega_{0,I})^2}} (A_{zz} + 2\omega_{0,I}) + \sqrt{1 + \frac{B^2}{(-A_{zz} + 2\omega_{0,I})^2}} (-A_{zz} + 2\omega_{0,I}) \right] \quad (65)$$

For spin systems in the weak coupling regime, where $|A_{zz}|, |B| \ll |\omega_{0,I}|$, the resonance offset in eqn (64) and (65) can be approximated by

$$\Omega_{0,S}^{(\text{res,DQ})} \approx (c_2 - c_1)\omega_m - |\omega_{0,I}| \quad (66)$$

and

$$\Omega_{0,S}^{(\text{res,ZQ})} \approx (c_2 - c_1)\omega_m + |\omega_{0,I}|, \quad (67)$$

respectively. The results in eqn (66) and (67) have two important consequences. First of all, we can assign the different resonance peaks in the experimental DNP profile in Fig. 4 to a specific Fourier index q' of the pulse sequence. So each resonance peak can be viewed as a modulation side band resulting from the fundamental modulation period of the applied pulse sequence. Second, according to eqn (66) and (67) we observe that the resonance offset $\Omega_{0,S}$ for $c_2 - c_1 = q' = \pm 1$ (the resonance offset with largest transition amplitude in the DNP profile see discussion later) shifts towards larger offsets when going towards larger magnetic fields. *E.g.* for a XiX sequence with $\tau_p = 1$ ns corresponding to $\frac{\omega_m}{2\pi} = 500$ MHz at a magnetic field of 28 T ($\frac{\omega_{0,I}}{2\pi} \approx -1200$ MHz), this would lead to resonance offset $\frac{\Omega_{0,S}}{2\pi}$ for $c_2 - c_1 = q' = \pm 1$ of ∓ 700 MHz and ± 1700 MHz.

This can have important consequences for the hardware design of a pulsed DNP spectrometer. The relation between operator-based Floquet theory and matrix-based Floquet theory in the weak irradiation limit and weak coupling regime is discussed in ESI† Section K.

At this point we need to consider the matrix elements that connect the ZQ/DQ transitions for the Fourier states c_1 and c_2 , because these elements represent the transition amplitudes. A transition $|c_1\rangle \leftrightarrow |c_2\rangle$ in Fourier space is connected by the off-diagonal blocks $\hat{\mathcal{H}}^{(q')}$ with $c_2 - c_1 = q'$ as indicated in Fig. 5 by the orange frame for the case $|0\rangle \leftrightarrow |-1\rangle$ and green frames for $|1\rangle \leftrightarrow |-1\rangle$, respectively. In such a constellation one can observe a total of four different transitions (two ZQ- and two DQ-transitions) as marked with yellow and purple for the DQ case and greenish-blue and brown for the ZQ case, respectively. The transition amplitudes can be obtained from the prefactor of the DQ- and ZQ-operators terms in the off-diagonal elements

$\hat{\mathcal{H}}^{(q')}$ in eqn (46) and are given by

$$I_{\text{ZQ/DQ}}^{(q')} = \begin{cases} \mp \text{sign}(q') \omega_{1,S}^{(\text{max})} \frac{i}{4} b_{|q'|} \sin(\eta), & \text{for odd } q' \\ 0, & \text{for even } q' \end{cases} \quad (68)$$

In the XiX case, whenever $q' = c_2 - c_1$ is odd, the off-diagonal block in the Floquet matrix is non-zero and, therefore, the transition amplitudes are non-zero as can be seen by eqn (68) and Fig. 5. This results in a resonance in the DNP profile that we observe experimentally by a polarization transfer. These resonance conditions are also confirmed by a non-zero Fourier coefficient for the Hamiltonian $a_{\pm}^{(k_0, \ell_0)}$ arising in numerical simulations using operator-based Floquet theory, *e.g.* for $\frac{\Omega_{0,S}}{2\pi} \approx \pm 18, \pm 48$ or ± 115 MHz in Fig. 4. Whenever $q' = c_2 - c_1$ is even, we do not observe a resonance peak in the experimental DNP profile. In numerical simulation using operator-based Floquet theory we obtain a match in the resonance condition, but with the Fourier coefficient of the Hamiltonian $a_{\pm}^{(k_0, \ell_0)}$ equal to 0. This represent cases like $\frac{\Omega_{0,S}}{2\pi} \approx \pm 81$ MHz in Fig. 4.

In the FS-XiX case, all Fourier coefficients of the microwave irradiation scheme are 0 except for $|q'| = 1$ and all off-diagonal blocks in the matrix-based Floquet representation are 0 except for the $\hat{\mathcal{H}}^{(\pm 1)}$ block. Therefore, in the FS-XiX case only those resonance peaks remain in the DNP profile for which $c_2 - c_1 = q' = \pm 1$ holds, *i.e.*, $\frac{\Omega_{0,S}}{2\pi} \approx \pm 18$ MHz and ± 48 MHz in Fig. 4. By using eqn (66) and (67), these four resonance peaks can be grouped into two DQ transitions ($\frac{\Omega_{0,S}}{2\pi} \approx -48$ MHz and $+18$ MHz) and two ZQ transitions ($\frac{\Omega_{0,S}}{2\pi} \approx -18$ MHz and $+48$ MHz). They are indicated in Fig. 5 by yellow colour for the DQ transition at $\frac{\Omega_{0,S}}{2\pi} \approx +18$ MHz, by purple colour for the DQ transition at $\frac{\Omega_{0,S}}{2\pi} \approx -48$ MHz, by greenish-blue colour for the ZQ transition at $\frac{\Omega_{0,S}}{2\pi} \approx +48$ MHz and by brown colour for the ZQ transition at $\frac{\Omega_{0,S}}{2\pi} \approx -18$ MHz. All other resonances resulting from $|c_2 - c_1| = |q'| \neq 1$ are absent in the experimental DNP profile. Numerical simulations using operator-based Floquet theory results in zero transition amplitude for these resonances.

As outlined above, we can group the resonance peaks in Fig. 4 according to the Fourier index q' (see Fig. 6 and Fig. S14 in ESI† Section M). In Fig. 6 and Fig. S14 (ESI†) we compare the approach using the infinite-dimensional Floquet Hamiltonian, which results in analytical expressions for the resonance offset $\Omega_{0,S}$ and transition amplitudes, with the numerical approach using operator-based Floquet theory for the DQ and ZQ resonance conditions. Both these approaches agree well with each other in the weak coupling regime ($|A_{zz}|, |B| \ll |\omega_{0,I}|$). The resonance offsets obtained by numerical simulation using operator-based Floquet theory are shown in dashed lines in



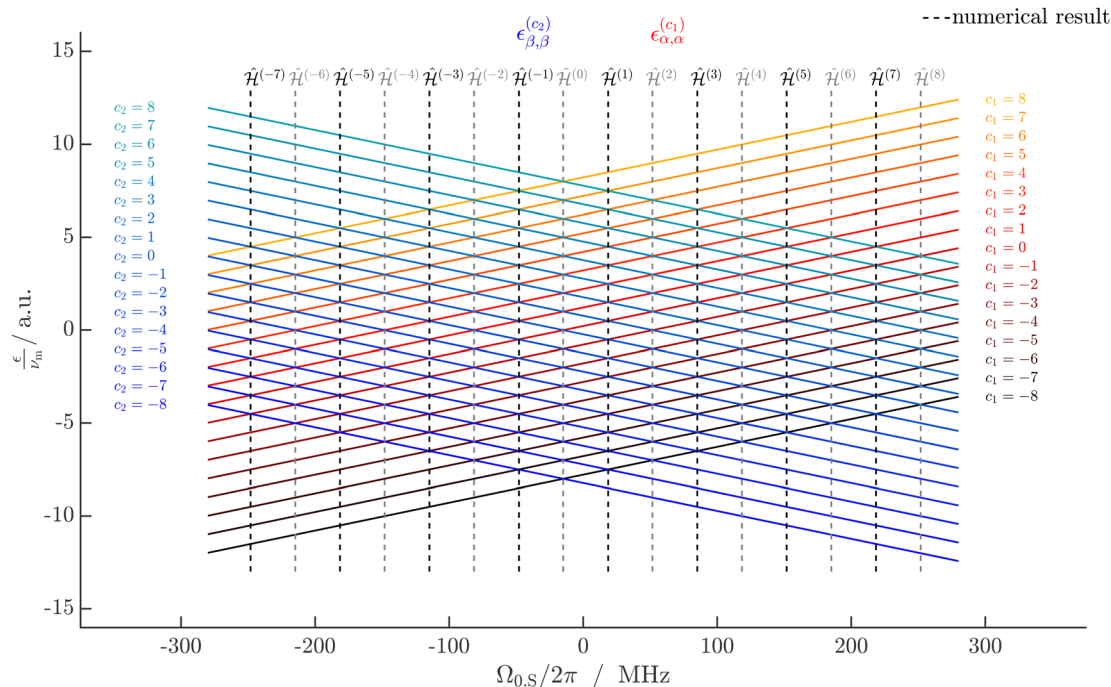


Fig. 6 Comparison between the eigenvalues of a specific diagonal block in the infinite-dimensional Floquet Hamiltonian as given in eqn (59) (red color code) and eqn (62) (blue color code) with the numerical simulation using operator-based Floquet theory (dashed lines) for the XiX DNP sequence at 0.35 T and a pulse length τ_p of 15 ns. Shown here are resonance conditions for DQ transitions. A resonance condition occurs at a mw offset $\Omega_{0,S}$, where the eigenvalues are equal i.e. $\epsilon_{\alpha,\alpha}^{(c_1)} = \epsilon_{\beta,\beta}^{(c_2)}$ with $c_2 - c_1 = q'$. The Fourier components $\hat{H}^{(q')}$ with Fourier index q' are indicated above the dashed lines. In the weak irradiation regime ($\omega_{1,S} \ll \omega_m$), each resonance condition can be assigned to a Fourier index of the infinite Floquet Hamiltonian and therefore to a mode in the Fourier expansion of the mw irradiation. The color code for the dashed lines indicates the strength of the resonance condition (see eqn (68)). For the analytical calculations of the eigenvalues according to eqn (59) and (62) the following parameters were used: $\frac{\omega_m}{2\pi} = 33.333$ MHz, $\frac{\omega_{0,I}}{2\pi} = -14.83$ MHz, $A_{zz} = B = 1$ MHz. Hyperfine constants are given here in linear frequency, whereas they are expressed in angular frequency in eqn (59)–(62).

those two figures. They overlap with the intersection points obtained by eqn (59) and (62) for DQ resonances and eqn (60)

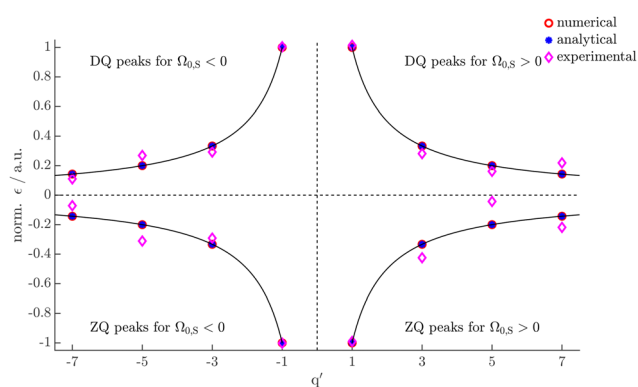


Fig. 7 Comparison between the transition amplitudes obtained by the analytical expression of eqn (68) obtained by matrix-based Floquet theory (red dots), the numerically calculated Fourier coefficients of the Hamiltonian in operator-based Floquet theory as given in eqn (14) (blue asterisks) and the experimental enhancements (magenta diamonds). The experimental values were obtained from a non-linear fit of the DNP profile presented already in Fig. 4. The black line is plotted to indicate the inverse scaling with respect to the Fourier index q of the analytical and numerical values.

and (61) for ZQ resonances, respectively. The dashed lines are color-coded according to transitions between Floquet states that have a non-zero transition amplitude (black) and a zero transition amplitude (gray) according to eqn (68).

A closer look at Fig. 4 reveals that there are some small peaks visible at the foot of the large assigned enhancements. As already explained in ref. 41 and 29 these small peaks correspond to three-spin electron- ^1H - ^1H transitions. One can use the same formalism described above for the two-spin case to obtain analytical expressions for the resonance offset $\Omega_{0,S}$ and the transition amplitudes for the three-spin case (see ESI,[†] Section N).

3.2.2 Transition amplitudes. The transition amplitudes according to eqn (68) can be compared to the numerically calculated Fourier coefficients of the Hamiltonian, as given in eqn (13), and to the experimental enhancements. This is graphically represented in Fig. 7. Note that, unlike the transition amplitudes, the numerical estimate for the enhancement as given in eqn (13) also contains the projection of ρ_0 onto the effective field (factor $\langle \rho_0 | \hat{S}_z \rangle$ in eqn (13)). However, for the pulsed DNP experiments presented in this work and for pulsed DNP experiments using low to moderate Rabi frequencies $\omega_{1,S}$, such that $\omega_{1,S} \ll \omega_m$, this factor is close to 1 since the resonance offsets are large and the irradiation is weak.



The experimental values were obtained by taking the maximal or minimal intensity of the DNP profile of Fig. 4. They were scaled linearly so that they agreed best with the theoretical black curve that represents the function $1/q$. As seen from Fig. 7, both, the transition amplitudes obtained by matrix-based Floquet theory (analytical) and the Fourier coefficients of the Hamiltonian obtained from operator-based Floquet theory (numerical), scale with the inverse of the Fourier coefficient q' (see black line in Fig. 7). For the analytical transition amplitudes this follows from inserting eqn (19) into eqn (68). There are a number of reasons for the observed differences between theory and experiments. In the theory so far we completely neglect relaxation effects, electron/proton concentrations or nuclear spin diffusion processes. These effects can contribute to the efficiency of the polarization transfer and therefore to the enhancement observed in experiments. Imperfections of the spectrometer, such as limited width of the microwave resonator mode or differences in non-linearity of the traveling wave tube amplifier at different frequencies can also influence the polarization enhancement. We tried to minimize effects of spectrometer imperfections during the acquisition by the techniques described in Section 5, and the ESI,† Sections E and F.

3.2.3 Field dependence of DNP sequences. In order to discuss the field dependence of a particular DNP sequence we need to emphasize some important points. We use the Fourier coefficients of the effective Hamiltonian obtained from operator-based Floquet theory (especially eqn (12) and (13)) for the analysis of the performance of a certain sequence.^{27–30} In the previous Section we presented analytical expressions for transition amplitudes obtained from the off-diagonal blocks in the infinite-dimensional Floquet matrix and showed that they are identical with the numerically calculated counterparts obtained from operator-based Floquet theory. Such an analysis does not cover any experimental parameters such as relaxation times, electron and proton concentration, nuclear spin diffusion, repeated application of the DNP sequence, or electronic imperfections. However, these parameters can play an important role in the polarization transfer and in the magnitude of the enhanced signal that is measured.

We restrict our discussion to the weak-irradiation regime, *i.e.*, $\omega_{1,S} \ll \omega_m$. In this regime the transition amplitude and, therefore, the transition probability between two almost degenerate Floquet states can be described by the matrix elements in the off-diagonal block of the Floquet matrix connecting the almost degenerate states. For a typical XiX sequence with short pulses τ_p , the modulation frequency ω_m becomes large, which results for the sequence to be in the weak irradiation regime even for large Rabi frequencies $\omega_{1,S}$. The transition amplitudes as given by eqn (68) for non-zero resonance conditions (for odd values of $c_2 - c_1 = q'$) are

$$I_{ZQ/DQ}^{(q')} = \frac{i}{4} b_{|q'|} \omega_{1,S}^{(\max)} \sin(\eta) \approx \frac{i}{4} b_{|q'|} \omega_{1,S}^{(\max)} \frac{B}{2\omega_{0,I}} \quad (69)$$

First we notice that the transition amplitude is independent of the

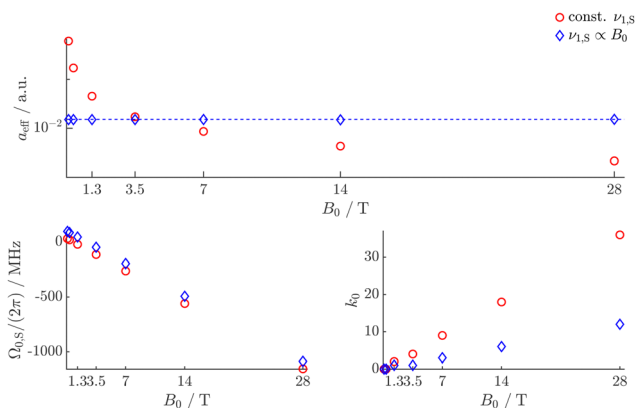


Fig. 8 Calculated values for a_{eff} , $\Omega_{0,S}$ and k_0 at resonance for different magnetic fields (0.1, 0.35, 1.3, 3.5, 7, 14, 28) T for DQ peaks with $c_2 - c_1 = 1$. The red circles represent a XiX DNP sequence with a fixed Rabi frequency $\frac{\omega_{1,S}}{2\pi} = 4$ MHz and pulse length $\tau_p = 15$ ns. XiX DNP sequences with $\tau_p = 5$ ns and Rabi frequencies that increase linearly with the external B_0 -field *e.g.* (0.1 MHz, 0.1 T), (0.35 MHz, 0.35 T), (1.3 MHz, 1.3 T), (3.5 MHz, 3.5 T), (7 MHz, 7 T), (14 MHz, 14 T) and (28 MHz, 28 T) for the Rabi frequency and the external magnetic field, respectively, are shown in blue diamonds. The linear behaviour of $\Omega_{0,S}$ with respect to the external magnetic field can be explained by the approximation in eqn (66). Due to eqn (9) also k_0 shows a similar linear behaviour. For the red circles, a fraction of a_{eff} for two different magnetic fields scale with the inverse fraction of the corresponding magnetic fields, respectively. In contrast, a_{eff} is constant over the different magnetic fields for a Rabi frequency that scales linearly with the magnetic field (blue diamonds). This is indicated by the dashed blue line. Please note the logarithmic scale in the y axis in the plot for a_{eff} .

pulse length τ_p (remember that $b_q = \frac{4}{q\pi}$ for odd q). The rightmost approximate expression results from the definitions of the angles $\eta_\alpha = \arctan\left(\frac{\pm B}{2\omega_{0,I} + A_{zz}}\right)$, $\eta_\beta = \arctan\left(\frac{\pm B}{2\omega_{0,I} - A_{zz}}\right)$ and $\eta = \frac{1}{2}(\eta_\alpha + \eta_\beta)$. The argument inside the arctan function scales with the inverse of the Larmor frequency of the nuclei and, therefore, with the inverse of the external magnetic field. In the weak coupling regime, where $|A_{zz}|, |B| \ll |\omega_{0,I}|$, we can invoke the small angle approximation and simplify $\sin(\eta) \approx \frac{B}{2\omega_{0,I}}$ in eqn (69). This approximation was already used to derive the prefactors for the solid-effect DNP mechanism in operator-based Floquet theory. We again obtain an unfavorable inverse scaling of the transition probability with larger magnetic fields. As eqn (69) contains also $\omega_{1,S}^{(\max)}$ as a pre-factor, the transition amplitude can be kept field-independent by increasing the Rabi frequency linearly with the external B_0 -field. This is the same scaling as required in CW DNP schemes.

To demonstrate our conclusion, we plotted in Fig. 8 the Fourier coefficient a_{eff} , the offset $\Omega_{0,S}$ and k_0 obtained from operator-based Floquet theory at the resonance condition for two different cases. In the first case we look at a XiX DNP sequence with a pulse length $\tau_p = 15$ ns and a fixed Rabi frequency $\frac{\omega_{1,S}}{2\pi} = 4$ MHz but for different magnetic fields. This case is shown in Fig. 8 in red circles. In blue diamonds in Fig. 8 we look at a XiX DNP sequence with a pulse length $\tau_p = 5$ ns



$\left(\frac{\omega_m}{2\pi} = 100 \text{ MHz}\right)$ for the following nutation frequency and external magnetic field combinations *i.e.* $\left(\frac{\omega_{1,S}}{2\pi}, B_0\right) \in \{(0.1 \text{ MHz}, 0.1 \text{ T}), (0.35 \text{ MHz}, 0.35 \text{ T}), (1.3 \text{ MHz}, 1.3 \text{ T}), (3.5 \text{ MHz}, 3.5 \text{ T}), (7 \text{ MHz}, 7 \text{ T}), (14 \text{ MHz}, 14 \text{ T}), (28 \text{ MHz}, 28 \text{ T})\}$. The pulse length of the second case was chosen to ensure that the weak irradiation regime ($\omega_{1,S} \ll \omega_m$) is fulfilled for all combinations. For both cases we discuss and compare the DQ resonances for the difference in Fourier indices $c_2 - c_1 = 1$. The ZQ case with $c_2 - c_1 = 1$ is treated in Fig. S16 in ESI,[†] Section O. Note that the cases ZQ $c_2 - c_1 = +1$ and DQ $c_2 - c_1 = -1$ are equivalent (except for the sign of a_{eff} and $\Omega_{0,S}$) due to the point symmetry of the ideal DNP profile.

From the first case with fixed Rabi frequency $\omega_{1,S}$, but varying magnetic field (red circles in Fig. 8) we can confirm the unfavorable scaling of the transition probability with larger magnetic fields. The numerically calculated Fourier coefficients of the Hamiltonian a_{eff} as given in eqn (14) scale with the inverse of the magnetic field *e.g.* the fraction $\frac{a_{\text{eff}}(B_0 = 7 \text{ T})}{a_{\text{eff}}(B_0 = 0.35 \text{ T})} = \frac{0.35}{7}$. This relation is true for all comparisons within the used magnetic fields and is predicted by eqn (69). In Fig. 8 one can nicely observe the linear increase of the resonance offset $\Omega_{0,S}$ for resonant DNP as already outlined in eqn (66) and (67). We find that k_0 also depends linearly on the magnetic field within the quantization error of a discrete parameter. This can be explained by the increase of $\omega_{0,I}$ in the resonance condition of eqn (9). k_0 selects the Fourier coefficients $a_{\pm}^{(k_0, \ell_0)}$ of the effective Hamiltonian relevant for DNP (see eqn (10) and (11)). The Fourier coefficients $a_{\pm}^{(k_0, \ell_0)}$ tend to decrease for larger values of k_0 , which can be seen from eqn (6) and which is the actual reason for the decreased transition probability towards larger magnetic fields.

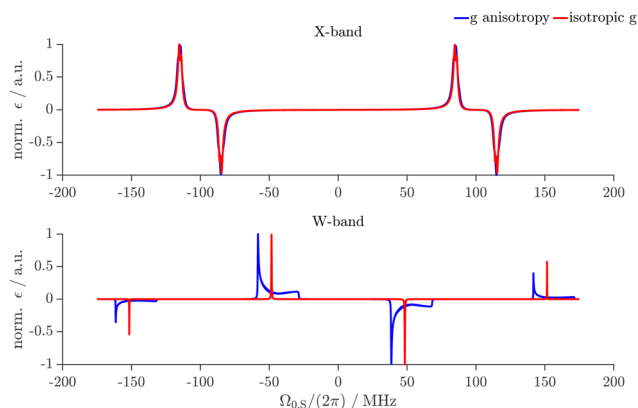


Fig. 9 Full numerical simulation of the field profile for XiX DNP with $\tau_p = 5 \text{ ns}$ and $\frac{\omega_{1,S}}{2\pi} = 4 \text{ MHz}$ at 80 K for B_0 fields of 0.35 T (X-band) top and 3.5 T (W-band) bottom for an isotropic g -tensor (red) and with an axial g -tensor with a slight g -anisotropy (blue). At X-band the difference between the simulations with and without g -anisotropy are very minor due to the small magnitude of the g -anisotropy. However, at W-band the resonance peaks in the DNP profile using a slight g -anisotropy are much broader than for the isotropic counterpart and resemble the g -tensor of the electron spin. All details of the numerical simulation can be found in ESI,[†] Section Q.

Increasing the Rabi frequency linearly with the external B_0 -field keeps the transition amplitude field independent as one can see from the blue diamonds in Fig. 8. a_{eff} is constant over the seven different combinations used in our simulations. Again, this behaviour can be explained with the right-hand side of eqn (69). Assuming that the strength of the pseudo-secular term B does not change (we assume that we polarize the same proton), the only terms that change within the seven combinations are $\omega_{1,S}^{(\text{max})}$ and $\omega_{0,I}$. The ratio $\frac{\omega_{1,S}^{(\text{max})}}{\omega_{0,I}}$ is kept constant here. Accordingly, the numerically calculated Fourier coefficients of the effective Hamiltonian in operator-based Floquet theory are the same.

We discussed here the three parameters a_{eff} , $\Omega_{0,S}$ and k_0 to reveal the behaviour of the XiX DNP pulse sequence when increasing the external magnetic field. As outlined in Section 2.1 the parameters $\omega_{\text{eff},S}$ and $\langle \rho_0 | \hat{S}_z \rangle$ at resonance condition also influence the performance of the XiX DNP sequence. However, the behaviour of $\omega_{\text{eff},S}$ and $\langle \rho_0 | \hat{S}_z \rangle$ with respect to different magnetic fields is difficult to predict/explain due to the merging of different effects (see Fig. S17 and S18 in ESI,[†] Section O). Cases for $c_2 - c_1 = \pm 3$ or even higher values are equivalent and are not discussed explicitly here.

3.2.4 Linewidth in the DNP profile. So far our theory as outlined in Sections 2.1 and 2.3 does not include a line width provided by either relaxation of the electrons or the orientation dependence of the g value. In this situation, our theory predicts a DNP profile with infinitely narrow lines, since the resonance condition in eqn (9) and the analytical expression for the microwave offset at resonance (see eqn (64) and (65)) are fulfilled for only a single $\Omega_{0,S}$. However, the experimental DNP profile (see Fig. 4) clearly shows broadened lines. The fit of a field-swept EPR spectrum at X band (9.8 GHz) to a Gaussian line shape gave a FWHM of 6.23 MHz for the allowed transition (see Fig. S4 in ESI,[†] Section D). The resonances in the DNP profile in Fig. 4 were also fitted to a Gaussian line shape (see Fig. S19 in ESI,[†] Section P) and the FWHM of each individual peak was extracted (Table S2 in ESI,[†] Section P) with an average width of about 5.59 MHz.

A full numerical simulation using GAMMA⁴² showed that the line width in the DNP profile is influenced by the g -anisotropy of the electron and dominated by this effect in W band. The resulting simulated field profiles for XiX with $\tau_p = 5 \text{ ns}$ and $\frac{\omega_{1,S}}{2\pi} = 4 \text{ MHz}$ at 80 K for B_0 fields of 0.35 and 3.5 T are plotted in Fig. 9. We simulated the field profile using a two-spin electron-proton system with an electron-proton distance $r_{\text{en}} = 4.5 \text{ \AA}$ with either an axial g -tensor with a slight g -anisotropy of $g_{xx} = g_{yy} = 2.00319$ and $g_{zz} = 2.00258$ or an isotropic g -tensor with $g_{\text{iso}} = 2.00299$.⁴³ A more detailed description of the GAMMA simulations can be found in ESI,[†] Section Q. At 0.35 T one can barely see a difference between the field profile with g -anisotropy and with isotropic g -tensor because of the small g -anisotropy. However, at 3.5 T the field profile of the simulation using a g -anisotropy show resonance peaks that resemble the powder lineshape for a g -tensor with axial symmetry.



3.3 TOP sequence

The discussion of the TOP DNP sequence in terms of its Fourier coefficients is not as straightforward as in the XiX case because the Fourier coefficients of the mw irradiation scheme now depend on pulse length τ_p and delay τ_d (see eqn (17) and (18)). In addition, the unmodulated Fourier coefficient a_0 is non-zero. Therefore, we need two consecutively applied unitary transformations to approximately diagonalize $\mathcal{H}^{(0)}$ (see Section 2.4 or ESI,† Section B for more details). As a consequence of this second unitary transformation the transition amplitudes now depend on the mw offset. However, the resonance conditions in the TOP DNP sequence can still be assigned to a certain Fourier index q' as will become clear later in this section.

First let us take a closer look on how the four eigenvalues for the TOP DNP case look like. The four eigenvalues can be obtained from eqn (51) and are given as

$$\varepsilon_{\alpha,\alpha}^{(c_1)} = c_1 \omega_m + \frac{1}{4} \sqrt{1 + \left(\frac{2\omega'_{1,S}}{A' + 2\Omega_{0,S}} \right)^2} (A' + 2\Omega_{0,S}) + \frac{\omega'_{0,I}}{2} \quad (70)$$

$$\varepsilon_{\alpha,\beta}^{(c_1)} = c_1 \omega_m - \frac{1}{4} \sqrt{1 + \left(\frac{2\omega'_{1,S}}{A' - 2\Omega_{0,S}} \right)^2} (A' - 2\Omega_{0,S}) - \frac{\omega'_{0,I}}{2} \quad (71)$$

$$\varepsilon_{\beta,\alpha}^{(c_1)} = c_1 \omega_m - \frac{1}{4} \sqrt{1 + \left(\frac{2\omega'_{1,S}}{A' + 2\Omega_{0,S}} \right)^2} (A' + 2\Omega_{0,S}) + \frac{\omega'_{0,I}}{2} \quad (72)$$

$$\varepsilon_{\beta,\beta}^{(c_1)} = c_1 \omega_m + \frac{1}{4} \sqrt{1 + \left(\frac{2\omega'_{1,S}}{A' - 2\Omega_{0,S}} \right)^2} (A' - 2\Omega_{0,S}) - \frac{\omega'_{0,I}}{2} \quad (73)$$

The exact analytical expressions for $\omega'_{0,I}$ and A' can be found in eqn (40) and (41), respectively. We can now invoke the same steps as for the XiX case and solve

$$\varepsilon_{\alpha,\alpha}^{(c_1)} - \varepsilon_{\beta,\beta}^{(c_2)} = 0 \quad (74)$$

and

$$\varepsilon_{\alpha,\beta}^{(c_1)} - \varepsilon_{\beta,\alpha}^{(c_2)} = 0 \quad (75)$$

for $\Omega_{0,S}^{(\text{res,DQ})}$ and $\Omega_{0,S}^{(\text{res,ZQ})}$, respectively. This leads to simplified expressions for these conditions in the weak coupling regime ($|A_{zz}|, |B| \ll |\omega_{0,I}|$). However, compared to the XiX case, we need to make an additional approximation. If the condition

$$|\omega'_{1,S}| = \omega_{1,S}^{(\text{max})} \frac{a_0}{2} \cos(\eta) \ll |\Omega_{0,S}| \quad (76)$$

is fulfilled, we find the same approximate resonance conditions as in the XiX case (e.g. eqn (66) and (67)), namely

$$\Omega_{0,S}^{(\text{res,DQ})} \approx (c_2 - c_1) \omega_m - |\omega_{0,I}| \quad (77)$$

and

$$\Omega_{0,S}^{(\text{res,ZQ})} \approx (c_2 - c_1) \omega_m + |\omega_{0,I}|. \quad (78)$$

Note that one needs to check for each case if the condition eqn (76) is fulfilled for all resonances. If so, the transition

Table 2 Fourier coefficients of the TOP DNP sequence according to eqn (17) and (18) for $\frac{\tau_d}{\tau_p} = 2$ and the expected resonance offsets according to eqn (77) and (78) calculated for different $c_2 - c_1 = q'$. The values for the resonance offsets are compared to a DNP profile measured from $\frac{\Omega_{0,S}}{2\pi} = -30$ MHz to $+\frac{\Omega_{0,S}}{2\pi} = +30$ MHz at 0.35 T with $\frac{\omega_{1,S}}{2\pi} = 4$ MHz, $\tau_p = 28$ ns, $\tau_d = 56$ ns as reported in ref. 27. The italic entries are those resonances which were also observed experimentally in ref. 27. The bold entries were reported as being observed by numerical simulations using operator-based Floquet theory, but gave zero enhancement in the experimental DNP profile (see Table 1 for $\frac{\tau_d}{\tau_p} = 2$)

$c_2 - c_1 = q'$	$a_{ q' }$	$(c_2 - c_1)\nu_m + \nu_{0,I} $ [MHz]	$(c_2 - c_1)\nu_m - \nu_{0,I} $ [MHz]
-4	0.138	-32.789	-62.449
-3	0	-20.884	-50.544
-2	-0.276	-8.980	-38.640
-1	0.551	2.925	-26.735
0	0.667	<i>14.830</i>	<i>-14.830</i>
1	-0.551	26.735	-2.925
2	0.276	38.640	<i>8.980</i>
3	0	50.544	20.884
4	-0.138	62.449	32.789

amplitudes for the TOP case are given according to eqn (56)

$$I_{\text{ZQ/DQ}}^{(q')} = \begin{cases} \mp \omega_{1,S}^{(\text{max})} \frac{a_0}{4} \cos(\theta) \cos(\eta) \sin(\eta), & \text{if } q' = 0 \\ \mp \omega_{1,S}^{(\text{max})} \frac{a_{|q'|}}{4} \cos(\theta) \sin(\eta), & \text{else} \end{cases} \quad (79)$$

where $\theta = \frac{1}{2}(\theta_\alpha - \theta_\beta)$ (see eqn (49) and (50) for analytical expression for θ_α and θ_β). As expected also the transition amplitudes for the TOP DNP sequence contain the term $\sin(\eta)$. As explained in Section 3.2.3 this term scales unfavorably with the inverse of B_0 when invoking the weak coupling regime. For low to moderate $\omega_{1,S}^{(\text{max})}$ e.g. the weak irradiation regime, θ is small and therefore $\cos(\theta) \approx 1$ holds.

We can now test the theory derived for TOP DNP against experimental observations reported in ref. 27. Tan *et al.* performed a TOP DNP experiment at 0.35 T (corresponding to $\nu_{0,I} \approx -14.83$ MHz) using trityl radical Ox063 with the following parameters: $\frac{\omega_{1,S}}{2\pi} = 4$ MHz, $\tau_p = 28$ ns, $\tau_d = 56$ ns. This results in $\frac{\tau_d}{\tau_p} = 2$ and a linear modulation frequency of $\frac{\omega_m}{2\pi} = 11.905$ MHz. They measured the DNP profile for this particular TOP DNP experiment from $\frac{\Omega_{0,S}}{2\pi} = -30$ MHz to +30 MHz and found the following values for the resonance offsets: $\frac{\Omega_{0,S}}{2\pi} \approx \pm 3, \pm 9, \pm 15$ and ± 27 MHz.²⁷ They also found a

DNP enhancement at $\frac{\Omega_{0,S}}{2\pi} \approx \pm 21$ MHz in numerical calculations using operator-based Floquet theory but with zero DNP enhancement in experiments. They explained the absence of DNP enhancement at this particular offset by assuming that the “ZQ- and DQ terms” (Fourier coefficients $a_{\pm}^{(k_0, \ell_0)}$ in eqn (10) and (11) in our notation) “have almost equal sizes” (and different sign). We can now use eqn (77)–(79) to explain the occurrence



and absence of resonance peaks in the TOP DNP profile reported in ref. 27. In Table 2 we list the Fourier indices for the TOP DNP experiment for the fraction $\frac{\tau_d}{\tau_p} = 2$ (see eqn (17) and (18)) along with the estimated values according to eqn (77) and (78) for different values of $c_2 - c_1 = q'$. We see that the approximations in eqn (77) and (78) give very accurate predictions for the microwave offsets $\Omega_{0,S}$, where DNP enhancements are expected. Eqn (79) together with eqn (17) and (18) fairly well predicts the amount of polarization transfer in the DNP profile presented in Fig. 2 in ref. 27. The resonances at $\frac{\Omega_{0,S}}{2\pi} \approx 15$ and 27 MHz show the largest enhancement, whereas the peak at $\frac{\Omega_{0,S}}{2\pi} \approx 9$ MHz shows only a small enhancement, as predicted from theory. However, the resonance offset $\frac{\Omega_{0,S}}{2\pi} \approx \pm 3$ MHz should in theory result in a similar enhancement as $\frac{\Omega_{0,S}}{2\pi} \approx \pm 27$ MHz. The data shown in Fig. 2 in ref. 27 indicate a slightly smaller enhancement for $\frac{\Omega_{0,S}}{2\pi} \approx \pm 3$ MHz than for $\frac{\Omega_{0,S}}{2\pi} \approx \pm 27$ MHz. We tentatively attribute this difference between simulation and experiment to the fact that the ZQ- and DQ enhancements occur at a frequency difference that is within the FWHM (~ 6 MHz) of the EPR line of the trityl radical Ox063. This case is similar to the enhancements close to $\frac{\Omega_{0,S}}{2\pi} = 0$ MHz in the XiX field profile for $\tau_p = 45$ ns shown in Fig. S13 (ESI†). As discussed in Section 3.2.4, the width of the experimental DNP profile at 0.35 T resembles the width of the EPR line. This may result in a partial cancellation of the ZQ- and DQ-enhancement due to the overlap of both peaks in the DNP profile and the different sign of their amplitudes. Using the theory outlined here, we can also explain the absence of DNP enhancement for $\frac{\Omega_{0,S}}{2\pi} \approx \pm 21$ MHz. According to eqn (17) and (18) and Table 1 all Fourier coefficients of the pulse sequence are non-zero except for Fourier indices of $q = 3, 6, 9, 12, \dots$. Therefore, for $c_2 - c_1 = q' = \pm 3$ we can calculate (by using eqn (77) and (78)) one pair of the offset to be $\frac{\Omega_{0,S}}{2\pi} \approx \pm 21$ MHz. This means that the enhancement at this resonance offset is zero because the Fourier coefficient a_3 is zero.

3.4 TPPM sequence

The analytical expressions for the diagonal blocks of the Floquet matrix after the two diagonalization steps ($\hat{\mathcal{H}}^{(0)''}$ in eqn (57)) are similar for the TPPM and TOP sequences. Hence, we can adapt the results obtained for the TOP DNP sequence in Section 3.3 to the TPPM case. The only difference is the definition of the Fourier coefficient $\frac{a_0}{2}$ (compare eqn (17) and (23)). In both cases $\frac{a_0}{2}$ cannot exceed one. Therefore, the condition in eqn (76) is also valid in the TPPM case, resulting in the same approximations for $\Omega_{0,S}$ as in the TOP DNP and XiX

Table 3 Comparison of transition amplitudes of different DNP sequences

Experiment	Transition amplitude	Comment
NOVEL	$\frac{B}{4}$	$\omega_{1,S}^{(\max)} = \omega_{0,I}$
SE	$\frac{B\omega_{1,S}^{(\max)}}{4\omega_{0,I}}$	
XiX	$\frac{B\omega_{1,S}^{(\max)}}{4\omega_{0,I}} \frac{ib_{ q' }}{2}$	For odd q' , $b_q = \frac{4}{q\pi}$
TOP	$\frac{B\omega_{1,S}^{(\max)}}{4\omega_{0,I}} \frac{a_0}{2}$	$a_0 = \frac{2\tau_p}{\tau_m}$
	$\frac{B\omega_{1,S}^{(\max)}}{4\omega_{0,I}} \frac{a_{ q' }}{2}$	$a_q = -\frac{2}{q\pi} \sin\left(\frac{q\pi\tau_d}{\tau_m}\right)$
TPPM	$\frac{B\omega_{1,S}^{(\max)}}{4\omega_{0,I}} \cos(\phi)$	
	$\frac{B\omega_{1,S}^{(\max)}}{4\omega_{0,I}} \frac{ib_{ q' }}{2} \sin(\phi)$	For odd q' , $b_q = \frac{4}{q\pi}$

DNP case, e.g. eqn (77) and (78). Therefore, in the weak irradiation regime the resonance offsets $\Omega_{0,S}$ of the three DNP sequences (TOP, XiX and TPPM) occur at the same spectral position when using the same modulation period ω_m . Using eqn (57) and (58) the transition amplitudes between the Fourier states c_1 and c_2 for the TPPM case are then given as

$$I_{ZQ/DQ}^{(q')} = \begin{cases} \mp \frac{B}{4} \cos(\phi) \frac{\omega_{1,S}^{(\max)}}{\omega_{0,I}}, & \text{if } q' = 0 \\ \mp \frac{Bib_{|q'|}}{4} \frac{\sin(\phi) \cos(\theta^{(+)}) \omega_{1,S}^{(\max)}}{2\omega_{0,I}}, & \text{if } q' \text{ is odd} \\ 0, & \text{if } q' \text{ is even} \end{cases} \quad (80)$$

where $q' = c_2 - c_1$ and $\theta^{(+)} = \frac{1}{2}(\theta_\alpha + \theta_\beta)$. For the angles θ_α and θ_β one needs to insert $\omega'_{1,S} = \omega_{1,S}^{(\max)} \cos(\phi) \cos(\eta)$ into in eqn (49) and (50). For the weak irradiation regime, i.e., for low to moderate $\omega_{1,S}^{(\max)}$ and sufficient large ω_m , $\cos(\theta^{(+)}) \approx 1$ holds. Thus for $\phi \rightarrow 0$ we obtain the same transition amplitude as for the SE and for $\phi \rightarrow \frac{\pi}{2}$ the transition amplitude as for the XiX sequence, respectively.

In Table 3 we compare the transition amplitudes for different DNP sequences applying the weak irradiation regime and the weak coupling regime. Please note that we do not distinguish between ZQ- and DQ-polarization transfer in that table and that $q = |q'| = |c_2 - c_1|$. Pulsed DNP sequences like XiX, TPPM or TOP in general show an inverse scaling of transfer efficiency with respect to the external magnetic field B_0 : the same as the CW experiments like SE.

4 Conclusions

Combining operator- and matrix-based Floquet theory for the description of a general amplitude-modulated pulsed DNP sequence, we can explain the origin of the resonance offsets



where enhancement occurs and the corresponding enhancement factors for pulsed DNP experiments TOP,²⁷ XiX^{28,29} or TPPM³⁰ sequences. We show experimentally that the resonances in the XiX DNP case can be excited by a simple sine wave and originate from a single frequency mode of the Fourier expansion. The TPPM sequence can be viewed as an XiX sequence with an additional orthogonal CW component and will, therefore, have a similar excitation profile. Using matrix-based Floquet theory, we obtain analytical expressions for the mw offsets that correspond to resonance conditions as well as expressions for the corresponding transition amplitudes. The latter quantities allow a better understanding of the scaling behavior of pulsed DNP experiments with respect to the external magnetic B_0 -field.

The frequency offset at the resonance conditions and the magnitude of the transition amplitudes agree very well with the numerical results obtained using operator-based Floquet theory. Both complementary approaches are able to predict experimental observations, *e.g.*, the mw offsets corresponding to resonance conditions and the magnitude of polarization transfer in an experimental DNP profile. However, important effects that influence experimental enhancements, such as relaxation, spin diffusion processes, experimental parameters like concentration of the radical and imperfections in mw irradiation are not covered by the presented theory. Regardless, the numerical results using effective Hamiltonians in operator-based Floquet theory as well as the analytical expressions resulting from matrix-based Floquet theory reveal an unfavorable scaling of the transition amplitudes with magnetic field. This result is not unexpected, since the effectiveness of the pseudo-secular $B\hat{S}_z\hat{J}_x$ term in the Hamiltonian of eqn (1) for mixing tends to zero when the external magnetic field approaches the high-field limit for the nuclear spin.

The performance of a particular pulsed DNP sequence can be maintained if and only if the Rabi frequency on the electron spin is increased linearly with the external B_0 -field. Therefore, pulsed DNP at high B_0 -fields requires the use of high-power mw devices. Such devices are not yet readily available at higher mw frequencies.

An alternative approach is simultaneous irradiation of electrons and protons by mw and rf, respectively. Such techniques, reminiscent of Hartmann–Hahn cross polarization, do not rely on the pseudo-secular part of the hyperfine coupling. In these schemes, mixing is achieved by the modulation of the secular part of the hyperfine coupling $A_{zz}\hat{S}_z\hat{I}_z$. A possible disadvantage of this approach would be that the transfer rate of polarization needs to compete with relaxation of the electrons and nuclei in the modulated spin-lock frame rather than with longitudinal relaxation of electrons and nuclei.

As confirmed by numerical simulations, part of the line width observed in experimental DNP profiles can be attributed to the anisotropy of the g -tensor. As expected, the g tensor contribution becomes dominant at larger magnetic fields. At higher fields, this effect could lead to an overlap of ZQ- and DQ-transfer resulting in a diminished polarization transfer if the bandwidth of the sequence is large. However, sequences

currently in use have a quite narrow DNP matching profile and will result in orientation selection for the g tensor.

The observations in this publication have implications for the future design of new pulsed DNP sequences to circumvent the unfavorable scaling of the effective Hamiltonian with respect to the external B_0 -field. Moreover, we were able to show that an enhanced NMR signal can be achieved by a simple cosine or sine wave. This opens new possibilities since those trigonometric functions are already implemented on most AWG and are less prone to electronic imperfections (pulse transients, finite rise times) than rectangular pulses. On the other hand, limited rise time of rectangular pulses corresponds to an attenuation of higher harmonics of the modulation frequency and is, thus, not expected to substantially reduce performance of pulsed DNP schemes.

5 Materials and methods

5.1 Sample preparation

In all experiments a 5 mM sample of OX063 trityl radical dissolved in DNP juice (glycerol- d_8 : D_2O : H_2O , 6:3:1 by volume) was used. The sample was prepared by the following procedure: 1.69 mg of the trityl ($MW = 1359 \text{ g mol}^{-1}$) were dissolved in 24.15 mg water ($MW = 18.02 \text{ g mol}^{-1}$, $\rho_0 = 0.997 \text{ g mL}^{-1}$) and 80.93 mg D_2O ($MW = 20.03 \text{ g mol}^{-1}$, $\rho_0 = 1.11 \text{ g mL}^{-1}$). This gives a $\sim 1.2 \mu\text{mol}$ solution of trityl dissolved in a D_2O : H_2O matrix of 3:1 by volume. From that resulting solution 48.6 μL were added to 99.92 mg glycerol- d_8 ($MW = 100.14 \text{ g mol}^{-1}$, $\rho_0 = 1.371 \text{ g mL}^{-1}$) to give the desired 5 mM solution of trityl in DNP juice. 40 μL of that final solution were added to a 3 mm OD quartz capillary. All masses were weighted in with a AT261 Delta Range scale with a precision of 0.01 mg. A pipetman from GILSON (P100) with volumes adjustable to 0.1 μL precision was used to transfer and measure the volumes.

All experiments were conducted at 80 K and an external magnetic field of 0.35 T (X-band). The sample was flash frozen in liquid nitrogen before measuring in the resonator.

5.2 EPR/NMR spectroscopy

All experiments were acquired on a home-built X-band spectrometer similar to the spectrometer as described in ref. 44. A constant temperature of 80 K was achieved with closed cycle cryogen-free cryostat from Cryogen Limited. An arbitrary waveform generator (Keysight model M8190A) was used to generate mw pulses and the pulses were amplified using a 1 kW traveling wave tube (TWT) amplifier. The Hahn echoes were recorded using a digitizer running at 1.8 GSa s^{-1} (SP devices ADQ412). A Bruker EN4118A-MD4 resonator was used with a home-built external rf tuning and matching box. NMR experiments were performed with an OpenCore spectrometer.^{45,46} A schematic representation of a general DNP experiment used in this work can be seen in Fig. 10. Each DNP sequence was started with a ^1H saturation pulse train consisting of eleven 100° pulses spaced by 1 ms to destroy any polarization on the protons. Afterwards, each basic DNP block was repeated k times to give a



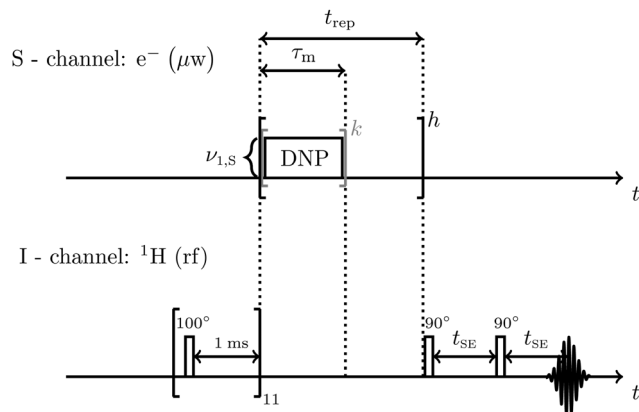


Fig. 10 General DNP pulse sequence used in the realm of this work. The saturation train on the ^1H channel is followed by a repeated DNP cycle (h -loop). The actual DNP sequence, either XiX or FS-XiX is repeated k -times. The enhanced ^1H signal is afterwards recorded with a solid echo with two 90° pulses of $2.5\ \mu\text{s}$ length separated by $t_{\text{SE}} = 20\ \mu\text{s}$. A eight-step phase cycle was used for the solid echo with $\{x, x, y, y, -x, -x, -y, -y\}$ for the first pulse and detection and $\{y, -y, x, -x, y, -y, x, -x\}$ for the second pulse.

total contact time $t_{\text{con}} = k \cdot \tau_m$ (gray loop in Fig. 10). The total DNP experiment was then repeated h times to give a total build-up time $t_{\text{DNP}} = h \cdot t_{\text{rep}}$ (outer black loop in Fig. 10). The enhanced proton signal was finally read out by a solid echo consisting of two 90° pulses of $2.5\ \mu\text{s}$ length separated by $t_{\text{SE}} = 20\ \mu\text{s}$. For the solid echo a eight step phase cycle was used with $\{x, x, y, y, -x, -x, -y, -y\}$ for the first pulse and detection and $\{y, -y, x, -x, y, -y, x, -x\}$ for the second pulse. For most of the experiments the following values were used: $t_{\text{con}} = 5\ \mu\text{s}$, $t_{\text{rep}} = 2\ \text{ms}$ and $h = 1000$. To record a DNP profile, the mw frequency on the EPR spectrometer was adjusted accordingly and a total of 8 scans were recorded for every mw frequency. The build-up curves were recorded by incrementing either h or k at a given resonance offset and keeping all other parameters constant. The reference experiment was recorded the same way with the exception that the mw irradiation was turned off and a delay of $180\ \text{s} \approx 5 \cdot T_{1,\text{H}}$ was used between two consecutive scans. A total of 512 scans were recorded and accumulated for the reference experiment. To process the data, a cosine-squared apodization function was used and time-domain data was zero-filled to twice the number of recorded data points. After Fourier transformation of the resulting time-domain data, the peak was fitted by a Lorentzian function and integrated using a standard library function in MATLAB over a symmetric integration window. The range of the integration window was constant for all individual recordings as well as for the reference experiment. The experimental enhancement was calculated by dividing the integral of the enhanced signal by the integral of the reference signal normalized by the number of scans e.g. normalized by 8 for a DNP experiment and by 512 for the reference experiment.

An echo-detected nutation experiment was used to compensate for the limited width of the microwave resonator mode and differences in non-linearity of the traveling wave tube amplifier at different frequencies.⁴⁷ The result of those experiments can be found in the ESI,[†] Sections E and F. All experimental spectra

and plots for the DNP, EPR and NMR experiments together with the complete experimental parameters can be found in ESI,[†] Sections C–I.

Data availability

The data that support the findings of this study are available at <https://doi.org/10.3929/ethz-b-000675865>.

Author contributions

M. E. and G. J. designed the research. Simulation and data evaluation script were written by G. M. C., N. W. and M. E., G. M. C. performed all experimental measurements. G. M. C. and M. E. wrote the first draft and all authors were involved in editing the final manuscript.

Conflicts of interest

The authors declare that they have no conflicts of interest.

Acknowledgements

Daniel Klose and Rene Tschaggelar are acknowledged for building the EPR spectrometer and Kazuyuki Takeda for help and discussion about the Opencore NMR spectrometer. We thank Mohammed M. Albannay for help in installing the Opencore NMR spectrometer. Oliver Oberhänsli is acknowledged for mechanical support. This research has been supported by the ETH Zürich and the Schweizerischer Nationalfonds zur Förderung der Wissenschaftlichen Forschung (grant no. 200020_188988 and 200020_219375).

Notes and references

- 1 A. Abragam and M. Goldman, *Rep. Prog. Phys.*, 1978, **41**, 395–467.
- 2 A. S. Lilly Thankamony, J. J. Wittmann, M. Kaushik and B. Corzilius, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2017, **102**, 120–195.
- 3 L. R. Becerra, G. J. Gerfen, R. J. Temkin, D. J. Singel and R. G. Griffin, *Phys. Rev. Lett.*, 1993, **71**, 3561–3564.
- 4 B. Corzilius, A. A. Smith and R. G. Griffin, *J. Chem. Phys.*, 2012, **137**, 21–27.
- 5 T. Biedenbänder, V. Aladin, S. Saeidpour and B. Corzilius, *Chem. Rev.*, 2022, **122**, 9738–9794.
- 6 J. H. Ardenkjær-Larsen, B. Fridlund, A. Gram, G. Hansson, L. Hansson, M. H. Lerche, R. Servin, M. Thaning and K. Golman, *Proc. Natl. Acad. Sci. U. S. A.*, 2003, **100**, 10158–10163.
- 7 F. Jähnig, G. Kwiatkowski and M. Ernst, *J. Magn. Reson.*, 2016, **264**, 22–29.
- 8 F. Jähnig, G. Kwiatkowski, A. Däpp, A. Hunkeler, B. H. Meier, S. Kozierke and M. Ernst, *Phys. Chem. Chem. Phys.*, 2017, **19**, 19196–19204.



- 9 D. Kurzbach and S. Jannin, *eMagRes*, 2018, **7**, 117–132.
- 10 S. Jannin, J. N. Dumez, P. Giraudeau and D. Kurzbach, *J. Magn. Reson.*, 2019, **305**, 41–50.
- 11 S. J. Elliott, Q. Stern, M. Ceillier, T. El Daraï, S. F. Cousin, O. Cala and S. Jannin, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2021, **126–127**, 59–100.
- 12 A. W. Overhauser, *Phys. Rev.*, 1953, **92**, 411–415.
- 13 C. D. Jeffries, *Phys. Rev.*, 1957, **106**, 164–165.
- 14 A. Abragam and W. G. Proctor, *Une nouvelle methode de polarization dynamique des noyaux atomiques dans les solides*, Comptes Rendus Hebdomadaires des Seances de l'Academie des Sciences, 246th edn, 1958, pp. 2253–2256.
- 15 C. D. Jeffries, *Phys. Rev.*, 1960, **117**, 1056–1069.
- 16 W. T. Wenckebach, *Appl. Magn. Reson.*, 2008, **34**, 227–235.
- 17 C. F. Hwang and D. A. Hill, *Phys. Rev. Lett.*, 1967, **18**, 110–112.
- 18 C. F. Hwang and D. A. Hill, *Phys. Rev. Lett.*, 1967, **19**, 1011–1014.
- 19 A. Equbal, A. Leavesley, S. K. Jain and S. Han, *J. Phys. Chem. Lett.*, 2019, **10**, 548–558.
- 20 B. N. Provotorov, *Sov. Phys. – JETP*, 1962, **14**, 1126–1131.
- 21 M. Borghini, *Phys. Rev. Lett.*, 1968, **20**, 419–421.
- 22 A. Henstra, P. Dirksen, J. Schmidt and W. T. Wenckebach, *J. Magn. Reson.*, 1988, **77**, 389–393.
- 23 T. V. Can, J. J. Walish, T. M. Swager and R. G. Griffin, *J. Chem. Phys.*, 2015, **143**, 1–8.
- 24 G. Mathies, S. Jain, M. Reese and R. G. Griffin, *J. Phys. Chem. Lett.*, 2016, **7**, 111–116.
- 25 T. V. Can, R. T. Weber, J. J. Walish, T. M. Swager and R. G. Griffin, *J. Chem. Phys.*, 2017, **146**, 1–7.
- 26 K. O. Tan, S. Jawla, R. J. Temkin and R. G. Griffin, *eMagRes*, 2019, **8**, 339–352.
- 27 K. O. Tan, C. Yang, R. T. Weber, G. Mathies and R. G. Griffin, *Sci. Adv.*, 2019, **5**, 1–7.
- 28 V. S. R. Redrouthu and G. Mathies, *J. Am. Chem. Soc.*, 2022, **144**, 1513–1516.
- 29 N. Wili, A. B. Nielsen, L. A. Völker, L. Schreder, N. C. Nielsen, G. Jeschke and K. O. Tan, *Sci. Adv.*, 2022, **8**, 1–13.
- 30 V. S. R. Redrouthu, S. Vinod-Kumar and G. Mathies, *J. Chem. Phys.*, 2023, **159**, 1–12.
- 31 I. Scholz, J. D. Van Beek and M. Ernst, *Solid State Nucl. Magn. Reson.*, 2010, **37**, 39–59.
- 32 M. Leskes, P. K. Madhu and S. Vega, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2010, **57**, 345–380.
- 33 J. H. Shirley, *Phys. Rev.*, 1965, **138**, B979–B987.
- 34 A. D. Bain and R. S. Dumont, *Concepts Magn. Reson.*, 2001, **13**, 159–170.
- 35 S. Vega, *eMagRes*, 2007, **2007**, 1–16.
- 36 K. L. Ivanov, K. R. Mote, M. Ernst, A. Equbal and P. K. Madhu, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2021, **126–127**, 17–58.
- 37 F. J. Dyson, *Phys. Rev.*, 1949, **75**, 486–502.
- 38 S. K. Jain, G. Mathies and R. G. Griffin, *J. Chem. Phys.*, 2017, **147**, 1–13.
- 39 M. Ernst, H. Geen and B. H. Meier, *Solid State Nucl. Magn. Reson.*, 2006, **29**, 2–21.
- 40 G. Jeschke and A. Schweiger, *J. Chem. Phys.*, 1996, **105**, 2199–2211.
- 41 K. O. Tan, M. Mardini, C. Yang, J. H. Ardenkjær-Larsen and R. G. Griffin, *Sci. Adv.*, 2019, **5**, 1–7.
- 42 S. A. Smith, T. O. Levante, B. H. Meier and R. R. Ernst, *J. Magn. Reson., Ser. A*, 1994, **106**, 75–105.
- 43 L. Lumata, Z. Kovacs, A. D. Sherry, C. Malloy, S. Hill, J. Van Tol, L. Yu, L. Song and M. E. Merritt, *Phys. Chem. Chem. Phys.*, 2013, **15**, 9800–9807.
- 44 A. Doll and G. Jeschke, *J. Magn. Reson.*, 2017, **280**, 46–62.
- 45 K. Takeda, *Rev. Sci. Instrum.*, 2007, **78**, 1–6.
- 46 K. Takeda, *J. Magn. Reson.*, 2008, **192**, 218–229.
- 47 A. Doll, S. Pribitzer, R. Tschaggelar and G. Jeschke, *J. Magn. Reson.*, 2013, **230**, 27–39.

