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# Spin conversion rates due to dipolar interactions in mono-isotopic quantum dots at vanishing spin-orbit coupling

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Dipolar interaction between the magnetic moments of electrons is studied as a source for electron spin decay in quantum dots or arrays of quantum dots. This magnetic interaction will govern spin decay, after other sources, such as the coupling to nuclear spins or spin orbit coupling, have been eliminated by a suitable sample design. Electron-electron (Coulomb) interactions, important for magnetic properties, are included. Decomposing the dipolar operator according to the symmetric group of electron permutations allows one to deduce vanishing decay channels as a function of electron number and spatial symmetries of the quantum dot(s). Moreover, we incorporate the possibility of rapid phonon induced spin conserving transitions which crucially affect the temperature dependence of spin decay rates. An interesting result is that a sharp increase of the spin decay rate occurs already at relatively low temperatures.

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## I. INTRODUCTION

One of the proposals to realize qubits for quantum computing<sup>1</sup> uses the electron's spin in semiconducting solid state nanostructures.<sup>2,3</sup> This approach could benefit from traditional electronics device experience and allows for straightforward scalability. Recent efforts have succeeded to demonstrate controlled preparation and detection of single<sup>4</sup> and of pairs<sup>5</sup> of electron spins in quantum dots. Aiming for long running coherent computations, however, solid state based devices somewhat suffer from relatively short spin decoherence and relaxation times as a drawback, for example, compared to nuclear spin based qubits.<sup>6</sup> Exponential decay of the upper Zeeman level population has been observed in GaAs quantum dots<sup>4</sup> over times not exceeding  $10^{-3}$  sec. Therefore, it is important to know (and, if possible, to control) any kind of mechanism causing spin relaxation in solids, and particularly in semiconductors. Previous theoretical work has valued several contributions. They can be subdivided into two classes regarding the magnetic mechanism to mix spin states as a source for spin decay: (i) spin-orbit coupling,<sup>7,8,9,10</sup> also, recently, in its interplay with the electron-electron interaction,<sup>11,12</sup> or (ii) coupling to nuclear spins. The latter can act through the spin-flip Overhauser effect by hyperfine interactions.<sup>13,14,15,16</sup> In principle, both magnetic sources for spin mixing can be eliminated by a proper device design and by the choice of the semiconductor material. Non-vanishing spin-orbit splitting can have several causes in semiconductors which we briefly address: *p*-type bands, i.e. usually valence bands, may split by spin-orbit effects, arising near the nuclei. Secondly does the lack of spatial inversion symmetry produce spin splitting even of *s*-type bands, ei-

ther by the Dresselhaus mechanism arising in the absence of crystallographic centrosymmetry as in Zincblende or Wurtzite structures. The latter particularly refers to all III-V semiconductors with GaAs being the most striking example. Also devices lacking structural inversion symmetry, e.g. near surfaces or in asymmetric quantum wells, producing internal electric fields show spin splitting due to the Rashba mechanism. Fortunately, this latter spin-orbit source may be suppressed by fine tuning suitable gate voltages.<sup>17</sup> The goal to avoid spin-orbit coupling effects therefore suggests using spins of conduction band electrons in Si or in Ge in carefully symmetrically prepared structures. Also the attempt to avoid coupling to nuclear spins favors the use of Si or Ge: their natural isotopic mixture contains nuclear spin  $I = 0$  to more than 95% (Si) or more than 92% (Ge), respectively.<sup>18</sup>

With this work we consider the effect of dipolar interactions between the magnetic moments of electron spins which, for fundamental reasons, *cannot* be removed by design. While considerably weaker<sup>19</sup> than the above quoted mechanisms this interaction unavoidably causes spin relaxation and, in the absence of other magnetic interactions, combined with the never vanishing electron-phonon coupling,<sup>20</sup> will set the ultimate limit for long time quantum computations using electron spins,<sup>3</sup> even in optimally designed structures. We study transitions between energy levels differing in their *total* spins<sup>21</sup> and disregard here effects associated with transitions between Zeeman levels (which conserve the symmetry of many body electron levels, see below) at finite magnetizations when an external magnetic field is applied. After introducing the model in Sect. II we reveal circumstances of particular spin stability w.r.t. dipolar interactions (Sect. III A), also regarding excited (many electron) states in Sect. III D, depending on the electron number and on the symmetry of the single or the ensemble of quantum dots.

We explicitly include Coulomb interactions<sup>11,12</sup> due to their importance for magnetic properties. For example,

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they can cause total ground state spins greater<sup>22,23,24,25</sup> than  $S = 0$  or  $S = 1/2$ , as expected for even or odd numbers  $N$  of non-interacting electrons. Here, we take the electrons to be confined inside one quantum dot or in different quantum dots. As a complementary approach to the non-interacting or weakly interacting regime we focus on strong Coulomb interactions at low electron densities, where pocket states,<sup>26,27,28</sup> offer a reliable description of many body states localized by Coulomb repulsion, even in single quantum dots.<sup>23</sup> Pocket states are briefly reviewed in Sect. III C for the present purpose to determine matrix elements of the dipolar interaction in Sect. III D.

Pursuing most of the foregoing theoretical work on electron spin decay in quantum dots we consider in Sect. IV A phonons (which themselves cannot change spin states) to provide the transition energy between discrete dot levels. Contrary to extended bulk situations, this transition energy is much bigger than mere magnetic energies which is one reason for the relative stability of quantum dot compared to bulk electron states, in accordance with experimental fact.<sup>29</sup> Spin changing transitions due to the combined action of dipolar energy and phonons are discussed in Sect. IV B. Generalizing previous results we account for rapid *spin conserving* excitations of the electron system induced by phonons that occur already at relatively low temperatures; these transitions turn out to govern predominantly the temperature dependence of spin decay times, discussed in Sect. IV C. Finally, we summarize and value our findings in Sect. V.

## II. MODEL

Specifically, we consider the  $N$ -electron system

$$H_0 = \sum_{i=1}^N \left( \frac{\mathbf{p}_i^2}{2m^*} + v(\mathbf{r}_i) \right) + \frac{1}{2} \sum_{i \neq j} w(|\mathbf{r}_i - \mathbf{r}_j|) \quad (1)$$

confined by the potential  $v(\mathbf{r})$  which is supposed to describe a single quantum dot or more complex situations of many quantum dots, such as for example  $N$  quantum dots each containing a single electron. To be realistic, particularly regarding magnetic properties, we include interactions between electrons  $w(r) = \frac{e^2}{\kappa} r^{-1}$ , depending on the static dielectric constant  $\kappa$  of the host material; Coulomb interactions are always considerably stronger than dipolar energies. Moderate screening, not reducing the interaction range to values smaller than the electron separation, will not affect qualitatively our results. In Eq. (1),  $\mathbf{p}_i$  and  $\mathbf{r}_i$  are  $d$ -component momentum and position vectors, depending on the dimensionality  $d$  of the quantum dot wave functions (in heterostructures,  $d = 2$ );  $m^*$  is the band electron mass.

Notice that at strong Coulomb repulsion, which is the focus of this work,  $N$  electrons Wigner crystallized<sup>23,30</sup> in a single quantum dot become in their theoretical treatment at low energies very similar to the case of  $N$  electrons localized in separate quantum dots. The essential

physics of both situations is captured by an antiferromagnetic Heisenberg lattice model.<sup>26,31</sup> Eigenstates of  $H_0$  exhibit well defined spins  $S$  and can be classified according to the eigenvalues of the  $z$ -component  $\hat{S}_z$  and the square  $\hat{S}^2$  of the total spin operator  $\hat{S} = \sum_{i=1}^N \hat{S}_i$  yielding eigenvalues  $S_z$  and  $S(S+1)$ , respectively. With  $SU(2)$  symmetry in spin space, Zeeman multiplets  $-S \leq S_z \leq +S$  are degenerate. We index eigenstates  $|\phi_n\rangle$  and eigenvalues  $E_n$  of  $H_0$  by  $n$ , taken to incorporate the values of  $S$  and  $S_z$ . Transitions between  $|\phi_n\rangle$  and  $|\phi_{n'}\rangle$  may or may not change  $S$ . The present work focuses on inelastic transitions that change the *total* spin values  $S \rightarrow S'$  rather than on transitions within a Zeeman multiplet. As already mentioned, many-electron ground states  $|\phi_{n=0}\rangle$  may exhibit total spin values  $S_0 > 1/2$  as a result of electron-electron interactions.<sup>22,23,24,25</sup>

## III. TRANSITION MATRIX ELEMENTS

In order to satisfy the Pauli principle an  $N$ -fermion state  $\langle \mathbf{r}_1, s_1, \dots, \mathbf{r}_N, s_N | \phi_n \rangle$  must belong to the  $A_2 \equiv [1^N]$  representation of the symmetric (permutational) group  $S_N$  with respect to permutations  $p \in S_N$  of the particle enumeration,  $\{1, \dots, N\} \rightarrow \{p(1), \dots, p(N)\}$ , see Ref. 32. When permuting only spin coordinates  $\{s_1, \dots, s_N\} \rightarrow \{s_{p(1)}, \dots, s_{p(N)}\}$  the state  $|\phi_n\rangle$  transforms according to the irreducible representation (partition)  $\Gamma = [N/2 + S, N/2 - S]$  of  $S_N$  for spin- $\frac{1}{2}$  Fermions<sup>27,28</sup> at given  $S = \begin{cases} 0 \\ 1/2 \end{cases}, \dots, N/2$  for  $\begin{cases} \text{even} \\ \text{odd} \end{cases} N$ . Correspondingly, when permuting only positions  $\{\mathbf{r}_1, \dots, \mathbf{r}_N\} \rightarrow \{\mathbf{r}_{p(1)}, \dots, \mathbf{r}_{p(N)}\}$ ,  $|\phi_n\rangle$  transforms according to  $\bar{\Gamma} = [2^{N/2-S}, 1^{2S}]$  (with  $\Gamma \times \bar{\Gamma}$  containing the  $A_2$ -representation). We notice that total spin changing transitions require altering the wave function's symmetry, which necessitates operators acting simultaneously in position and in spin space (by contrast, transitions within a Zeeman multiplet leave unaltered the symmetries of wave functions).

### A. Dipolar energy

Here, we investigate the dipolar interaction  $H^D$  between electrons. As seen in Eq. (3) below, it contains products of position and spin operators and, indeed, mixes spin states. However, it is by far too weak to provide the energy

separating quantum dot eigenlevels. Focusing on Si, we consider in Section IV acoustical deformation potential phonons<sup>33</sup> to supply the necessary transition energy. Unaided electron-phonon coupling, though, does not mix spin states and thus leaves spins unaltered. Eventually, it turns out that dipolar interaction, as a result of its smallness, ensues considerably smaller transition rates at low temperatures than, for instance, nuclear spin induced spin mixing.<sup>13</sup>

The operator of the dipolar energy

$$H^D = \frac{1}{2} \sum_{i \neq j} H_{ij}^D \quad (2)$$

is, as required for identical particles, invariant with respect to permuting the electron enumeration; however,  $H^D$  can be decomposed into parts that are *not* invariant under permuting coordinates  $\mathbf{r}_i$  or spins  $\hat{\mathbf{S}}_i$  separately. Let us first recap the interaction between a pair of magnetic moments

$$H_{ij}^D = \frac{\gamma^2}{r_{ij}^5} \left[ r_{ij}^2 \hat{\mathbf{S}}_i \cdot \hat{\mathbf{S}}_j - 3(\mathbf{r}_{ij} \cdot \hat{\mathbf{S}}_i)(\mathbf{r}_{ij} \cdot \hat{\mathbf{S}}_j) \right] \quad (3)$$

where  $\gamma = ge\hbar/2mc$  ( $c$  is the velocity of light and the  $g$ -factor for dot carriers which even in few electron quantum dots is found to take basically bulk values).<sup>5,34</sup> Its Heisenberg-like first part is manifestly  $SU(2)$ -invariant in spin space and commutes with  $\hat{\mathbf{S}}^2$ . This part neither changes  $S$  nor  $S_z$  and just renormalizes the energies slightly. It therefore can be ignored in view of the smallness of dipolar energies compared to the dot level separations. In Eq. (3) we abbreviate  $\mathbf{r}_{ij} := \mathbf{r}_i - \mathbf{r}_j$  and  $r_{ij} := |\mathbf{r}_{ij}|$ . The second part of  $H_{ij}^D$  can be decomposed as

$$\frac{1}{r_{ij}^5} (\mathbf{r}_{ij} \cdot \hat{\mathbf{S}}_i)(\mathbf{r}_{ij} \cdot \hat{\mathbf{S}}_j) = \left[ H_{ij}^{(0)} + H_{ij}^{(1)} + H_{ij}^{(2)} \right] \quad (4)$$

where the three terms

$$H_{ij}^{(0)} = \frac{|\varrho_{ij}|^2}{4} (\hat{S}_{+i}\hat{S}_{-j} + \hat{S}_{-i}\hat{S}_{+j}) + \zeta_{ij}^2 \hat{S}_{zi}\hat{S}_{zj} \quad (5)$$

$$H_{ij}^{(1)} = \frac{\zeta_{ij}}{2} \left[ \varrho_{-ij} (\hat{S}_{+i}\hat{S}_{zj} + \hat{S}_{zi}\hat{S}_{+j}) + \varrho_{+ij} (\hat{S}_{-i}\hat{S}_{zj} + \hat{S}_{zi}\hat{S}_{-j}) \right] \quad (6)$$

$$H_{ij}^{(2)} = \frac{\varrho_{+ij}^2}{4} \hat{S}_{-i}\hat{S}_{-j} + \frac{\varrho_{-ij}^2}{4} \hat{S}_{+i}\hat{S}_{+j}; \quad (7)$$

are responsible to alter symmetries and spins after carrying out summation over ( $i \neq j$ ). In Eqs. (5–7) they change  $S_z$  by 0,  $\pm 1$ , and  $\pm 2$ , respectively.  $\hat{S}_{\pm} := \hat{S}_x \pm i\hat{S}_y$  denote usual rising or lowering operators in spin space,  $\varrho_{\pm ij} := (x_{ij} \pm iy_{ij})/r_{ij}^{5/2}$  is a complex (angular momentum generating) coordinate in the plane perpendicular to the axes of spin quantization, taken as the  $z$ -axes, and  $\zeta_{ij} := z_{ij}/r_{ij}^{5/2}$ .

The spin changing part Eqs. (5–7) of  $H^D$  can now further be decomposed according to partitions  $\Gamma$  of the symmetric group  $S_N$ ,

$$\begin{aligned} & \sum_{i \neq j} (H_{ij}^{(0)} + H_{ij}^{(1)} + H_{ij}^{(2)}) \\ &= H^{\Gamma=[N]} + H^{\Gamma=[N-1,1]} + H^{\Gamma=[N-2,2]}. \end{aligned} \quad (8)$$

This latter representation is particularly useful to deduce non-zero transition matrix elements between quantum dot eigenstates of different *total* spins. No other partitions occur since  $H_{ij}^D$  transforms as a product of two vector operators in position as well as in spin space, cf. Eq. (3), i.e. as a tensor of rank two. In Eq. (8)  $H^{\Gamma=[N-1,1]}$  changes the total spin  $S$  of dot by  $\pm 1$  and  $H^{\Gamma=[N-2,2]}$  by  $\pm 2$ , where the latter occurs only for  $N \geq 4$  while the former already for  $N \geq 3$ .  $H^D$  cannot achieve spin changes by more than  $\pm 2$ . For example, we can conclude already at this stage that the rate for *direct* transitions of an excited  $S = 3$  quantum dot state into the (assumed)  $S = 0$  singlet ground state will be of the order  $\mathcal{O}((H^D)^4)$  and therefore will be very small. All properly symmetrized operators  $H^{\Gamma=[N-1,1]}$  and  $H^{\Gamma=[N-2,2]}$  for  $N = 3$  and  $N = 4$  are listed in Appendix A.

Note, that the property of  $H_{ij}^{(0-2)}$  to change  $S_z$  by 0,  $\pm 1$ ,  $\pm 2$ , respectively, is unrelated to their respective capability to change  $S$ . In the absence of further symmetries of the quantum dot shape, all three operators  $H_{ij}^{(0-2)}$  contain both,  $H^{\Gamma=[N-1,1]}$  and  $H^{\Gamma=[N-2,2]}$ . In cases of frozen electron motion in  $z$ -direction, as it applies to quantum dots (or arrays of quantum dots) fabricated on the basis of semiconducting hetero-structures<sup>35</sup> all of the above contributions involving  $\zeta_{ij}$  vanish. Then  $H^{(0)}$  simplifies and  $H^{(1)}$  vanishes entirely, so that  $S_z$  can either remain unaltered (through  $H^{(0)}$ ) or change by  $\pm 2$  (through  $H^{(2)}$ ).

## B. Two electrons

Let us first focus on two electrons, i.e.  $N = 2$ . This is relevant, for example, for double dots containing one electron on either side to realize the basic entity of coupled qubits.<sup>3,36</sup> As already mentioned in the previous section, non- $A_1$  symmetric partitions Eq. (8) of  $H^D$  occur only for  $N \geq 3$ . Further,  $H^D$  does not contain the  $A_2$  partition for any  $N$ . Therefore, spin conversion transitions from a triplet excited state into the singlet ground state<sup>37</sup> will never be mediated by  $H^D$ . In the related physics context of nuclear spin conversion of  $H_2$  molecules<sup>38</sup> the stability of ortho-hydrogen (even over weeks) is traced back<sup>39</sup> to parity symmetry of both, the molecule and the magnetic dipolar interactions between the two protons (of actually close proximity which enhances dipolar forces) to prevent the transition from the odd-parity ortho  $S = 1$  state into the (by 80 Kelvin lower) even-parity para  $S = 0$  ground state. In this case, spins refer to the protons. In the context of quantum dots we can generalize this finding: Irrespective of the shape of the quantum dot confining potential and of the functional form of the electron-electron interaction  $w(\mathbf{r}_1 - \mathbf{r}_2)$  the dipolar interaction will not change (triplet or singlet) spin states as a result of permutational symmetry and quantum mechanical particle identity of  $N = 2$  electrons. This statement is not restricted to the lowest (golden rule) order  $\mathcal{O}((H^D)^2)$  but even holds true to any order of  $H^D$ . As one neat corollary

we conclude that two electrons in square shaped quantum dots (in the absence of other magnetic mechanisms) will stay in their respective spin states. This supports a corresponding proposal for quantum computations based on superpositions of states where the two electrons occupy either of the two equivalent electrostatic energy minimum positions at diagonally opposite corners in a square.<sup>40</sup>

### C. Strong interaction, pocket states

In case of more than two electrons we focus on strong Coulomb forces at low carrier densities, i.e. at large values of the electron gas parameter  $r_s \gg 1$ . Then, the kinetic energy is small and the electron system lowers its energy by Wigner localizing<sup>30</sup> the charge density near electrostatically favorable places. Precursors of Wigner crystallization have been found already at  $r_s \approx 4$  in two-dimensional quantum dots.<sup>23</sup> A similar localization of charge density arises when the external confining potential separates the electrons, such as in the case of  $N$  quantum dots, each containing a single electron. In either case, at strong Coulomb interactions, eigenstates  $|\phi_n\rangle$  of  $H_0$  Eq. (1) are well described by pocket states,<sup>26,27,28</sup> which exploit the electron localization. They allow to estimate the spin dependent low energy spectrum to exponential accuracy with increasing  $r_s$ , or with increasing dot separation.

In the Wigner crystal state, electrons vibrate about electrostatic energy minimum positions. Linearizing the (Coulomb and external) forces yields the plasmon spectrum of the confined  $N$ -electron system. Energy level separations  $\omega_{\text{pl}}^2 \sim \omega_0^2 + Ar_s^{-3}$  can be estimated from the dynamical matrix with a prefactor  $A$  depending on  $N$  and on the dot lay-out;  $\omega_0$  is the confining frequency of the quantum dot(s). Due to the electron spin each plasmon level is  $2^N$ -fold degenerate. Quantum corrections (partly) split this degeneracy into sub-levels, with all exhibiting well defined total spins  $S = \left\{ \begin{smallmatrix} 0 \\ 1/2 \end{smallmatrix} \right\}, \dots, N/2$  for  $\left\{ \begin{smallmatrix} \text{even} \\ \text{odd} \end{smallmatrix} \right\} N$  [of  $(2S+1)$ -fold Zeeman degeneracy, by spin rotation invariance], according to Sect. II. The ground state (in more than one spatial dimension) need not be of minimal spin  $S_0 = 0$  or  $S_0 = 1/2$ .<sup>22,24,25,27</sup> A given spin  $S$  may appear more than once in such a spin split plasmon level; examples of spectra are discussed in Refs. 26,27,28. The splitting arises due to permutational electron exchanges by quantum mechanical tunneling through the electrostatic barrier consisting of the  $v$ -term plus the  $w$ -term in Eq. (1). In the simplest case there are  $N!$  different, but all energetically precisely equivalent, possibilities to arrange the localized electrons; this defines  $1 \leq p \leq N!$  pocket states  $|p\rangle$ . The width of each pocket state corresponds to plasmonic zero point oscil-

lations and scales roughly as  $\omega_{\text{pl}}^{-1/2}$  in  $Nd$ -dimensional configuration space ( $d$  being the spatial dimensionality of the quantum dot, often<sup>35</sup>  $d = 2$  but also  $d = 1$  is realized, for example in rods of carbon nanotubes<sup>41</sup>). The energy scale  $\Delta$  for spin splittings of plasmon levels through quantum mechanical electron exchanges is tuned by the magnitude of overlap integrals  $\langle p' | H_0 | p \rangle$  between two different arrangements  $p$  and  $p'$ . This latter quantity can be estimated semiclassically<sup>27,28,31</sup> to read  $\Delta \sim \langle p' | H_0 | p \rangle \sim \omega_{\text{pl}} \exp(-\sqrt{r_s})$  so that  $\Delta/\omega_{\text{pl}} \ll 1$ . Numerically obtained quantum dot spectra<sup>23,27,42</sup> indeed nicely follow this behavior. For example, it exhibits the predicted<sup>24</sup> crossover into a spin polarized  $S = 3/2$  ground state in a spherical two dimensional quantum dot containing  $N = 3$  electrons at sufficiently low electron density.<sup>23,25</sup> As a result, all eigenstates

$$|\phi_n\rangle = \frac{1}{N_n} \sum_p c_{np} |p\rangle \quad (9)$$

belonging to the plasmon ground multiplet can approximately be expressed through the set  $\{|p\rangle\}$ . The (real) coefficients  $c_{np}$ , appearing in Eq. (9), ensue from the irreducible representation  $\Gamma$  of the permutational group  $S_N$ , according to the wave functions symmetry which at the same time fixes the total spin  $S$  of  $|\phi_n\rangle$ ;  $N_n = \sqrt{\sum_{pp'} c_{np} c_{np'}} \langle p' | p \rangle$  ensures normalization,  $\langle \phi_n | \phi_n \rangle = 1$ .

### D. Dipolar matrix elements

Pocket states allow to conveniently estimate the matrix elements  $\langle \phi_n | H^D | \phi_{n'} \rangle$  of the dipolar energy since, to leading order, electron positions may be taken as being well localized,  $\delta$ -function like, for  $\varrho_{ij}$  and  $\zeta_{ij}$  in Eqs. (5–7) or in the already symmetrized expressions in Appendix A. This leads to a finite lattice spin problem. Having constructed symmetrized spin states, the matrix elements of  $H^D$  for  $N > 4$  follow straightforwardly from Eq. (8).

We demonstrate our approach for the particularly symmetric cases of  $N = 3$  and  $N = 4$  electrons occupying equilateral electrostatic equilibrium positions, as in a two-dimensional spherical quantum dot,<sup>23,24</sup> in triangularly or square shaped quantum dots,<sup>35</sup> or in equilateral triangular or square arrangements of single electron quantum dots. We assume frozen motion in  $z$ -direction, as in heterostructures, so that terms involving  $\zeta_{ij}$  or  $\zeta_{ij}^2$  are irrelevant in Eqs. (5–7). Symmetrized, non-trivial spin states of minimal  $S_z$ -components are presented in Table I.

$N$	$S$	$S_z$	index	
3	$\frac{3}{2}$	$\frac{1}{2}$	A	$\frac{1}{\sqrt{3}}( \uparrow\uparrow\downarrow\rangle +  \uparrow\downarrow\uparrow\rangle +  \downarrow\uparrow\uparrow\rangle)$
3	$\frac{1}{2}$	$\frac{1}{2}$	E <sub>a</sub>	$\frac{1}{\sqrt{3}}( \uparrow\uparrow\downarrow\rangle + e^{i2\pi/3} \uparrow\downarrow\uparrow\rangle + e^{-i2\pi/3} \downarrow\uparrow\uparrow\rangle)$
3	$\frac{1}{2}$	$\frac{1}{2}$	E <sub>b</sub>	$\frac{1}{\sqrt{3}}( \uparrow\uparrow\downarrow\rangle + e^{-i2\pi/3} \uparrow\downarrow\uparrow\rangle + e^{i2\pi/3} \downarrow\uparrow\uparrow\rangle)$
4	2	0	A	$\frac{1}{\sqrt{6}}( \uparrow\uparrow\uparrow\downarrow\rangle +  \uparrow\uparrow\downarrow\uparrow\rangle +  \uparrow\downarrow\uparrow\uparrow\rangle +  \uparrow\downarrow\uparrow\downarrow\rangle +  \uparrow\downarrow\downarrow\uparrow\rangle +  \uparrow\downarrow\downarrow\downarrow\rangle)$
4	1	0	T <sub>x</sub>	$\frac{1}{2}( \uparrow\uparrow\uparrow\downarrow\rangle - i \uparrow\uparrow\downarrow\uparrow\rangle -  \uparrow\uparrow\downarrow\downarrow\rangle + i \uparrow\downarrow\uparrow\uparrow\rangle)$
4	1	0	T <sub>y</sub>	$\frac{1}{2}( \uparrow\uparrow\uparrow\downarrow\rangle + i \uparrow\uparrow\downarrow\uparrow\rangle -  \uparrow\uparrow\downarrow\downarrow\rangle - i \uparrow\downarrow\uparrow\uparrow\rangle)$
4	1	0	T <sub>z</sub>	$\frac{1}{\sqrt{2}}( \uparrow\uparrow\uparrow\downarrow\rangle -  \uparrow\downarrow\uparrow\uparrow\rangle)$
4	0	0	E <sub>1</sub>	$\frac{1}{2}( \uparrow\uparrow\uparrow\downarrow\rangle -  \uparrow\uparrow\downarrow\uparrow\rangle +  \uparrow\uparrow\downarrow\downarrow\rangle -  \uparrow\downarrow\uparrow\uparrow\rangle)$
4	0	0	E <sub>2</sub>	$\frac{1}{\sqrt{8}}( \uparrow\uparrow\uparrow\downarrow\rangle +  \uparrow\uparrow\downarrow\uparrow\rangle +  \uparrow\uparrow\downarrow\downarrow\rangle +  \uparrow\downarrow\uparrow\uparrow\rangle - 2 \uparrow\downarrow\uparrow\downarrow\rangle - 2 \uparrow\downarrow\downarrow\uparrow\rangle)$

TABLE I: Symmetrized, non-trivial spin states of minimal  $S_z$ -component for  $N = 3$  and  $N = 4$ . Spin states of larger  $|S_z|$  are obtained easily.

### 1. $N = 3$

In this case  $\sum_{i \neq j} \varrho_{ij}^2 = 0$  and, in Appendix A, we replace  $|\varrho_{ij}|^2$  by  $r^2$  for  $N = 3$  where  $r$  is the mean inter-electron separation. Then, the only non-vanishing term  $H^{(2)[2,1]}$  takes the value

$\sum_{i \neq j} H_{ij}^{(2)} = \frac{r^2}{4} \sum_{i \neq j} e^{i2\vartheta_{ij}} \hat{S}_{-i} \hat{S}_{-j} + \text{h.c.}$  where  $\vartheta_{ij} = 0, \frac{2\pi}{3}, \frac{-2\pi}{3}$  is the azimuthal angle of  $\mathbf{r}_{ij}$ .

Thus,  $H^D$  necessarily changes  $S_z$  by  $\pm 2$  and has non-vanishing matrix elements only between the A and the E states  $\{S = 3/2, S_z = 3/2\} \leftrightarrow \{S = 1/2, S_z = -1/2\}$  and  $\{S = 3/2, S_z = -3/2\} \leftrightarrow \{S = 1/2, S_z = +1/2\}$  of Table I. Their value emerges as:

$$\langle \phi_{\{A, S_z = \pm 3/2\}} | H^D | \phi_{\{E_{a,b}, S_z = \mp 1/2\}} \rangle = -\frac{3\sqrt{3}}{4} \frac{\gamma^2}{r^3}.$$

In particular, this means that the not Zeeman aligned states of  $S = 3/2$  with  $S_z = \pm 1/2$  remain unaffected from dipolar decay.

### 2. $N = 4$

For a square arrangement of  $N = 4$  electrons two distances occur:  $r$  along one edge and  $\sqrt{2}r$  across the diagonal. Inspecting all the terms  $H^{(0)[3,1]} \dots H^{(2)[2,2]}$  for  $N = 4$  in Appendix A reveals that only contributions  $\frac{1}{16} \sum_{i \neq j} [\varrho_{+ij}^2 \pm \varrho_{+\bar{i}\bar{j}}^2] [\hat{S}_{-i} \hat{S}_{-j} \pm \hat{S}_{-\bar{i}} \hat{S}_{-\bar{j}}] + \text{h.c.}$  remain non-vanishing. Again,  $S_z$  has to change by  $\pm 2$ . The term  $\sim [\varrho_{+ij}^2 + \varrho_{+\bar{i}\bar{j}}^2]$  vanishes across the diagonal of the square, while  $[\varrho_{+ij}^2 - \varrho_{+\bar{i}\bar{j}}^2] = 0$  along any edge. An evaluation yields the non-zero matrix elements between symmetrized spin states of Table I, i.e.

$$\begin{aligned} \langle \phi_{\{A, S_z = \pm 2\}} | H^D | \phi_{\{T_z, S_z = 0\}} \rangle &= -\frac{3}{\sqrt{2} \cdot 32} \frac{\gamma^2}{r^3} \\ \langle \phi_{\{A, S_z = \pm 1\}} | H^D | \phi_{\{T_z, S_z = \mp 1\}} \rangle &= -\frac{3}{64} \frac{\gamma^2}{r^3} \end{aligned}$$

$$\langle \phi_{\{A, S_z = \pm 2\}} | H^D | \phi_{\{E_1, S_z = 0\}} \rangle = -\frac{3}{32} \frac{\gamma^2}{r^3}.$$

All other matrix elements vanish identically. In particular, states of symmetries T<sub>x</sub>, T<sub>y</sub>, E<sub>2</sub>, and A with  $S_z = 0$  do not exhibit dipolar decay.

### E. Mixed spin states

Non-vanishing elements of  $H^D$  slightly mix eigenstates  $|\phi_n\rangle$  of  $H_0$ . Here we are interested in admixtures to spin states  $S$

$$|\psi_{n_S}\rangle = |\phi_{n_S}\rangle + \sum_{S' \neq S} \sum_{n_{S'}} \frac{\langle \phi_{n_{S'}} | H^D | \phi_{n_S} \rangle}{E_{n_S} - E_{n_{S'}}} |\phi_{n_{S'}}\rangle \quad (10)$$

arising from other spins  $S' \neq S$ . In Eq. (10) we have disregarded the very unlikely case of accidental degeneracy between eigenlevels of  $H_0$  (cf. (1)) of different spins.<sup>43</sup>

Eventually, this mixing will cause spin changing transitions and thus spin relaxation. We disregard dipolar admixtures from other states  $|\phi_{n_{S'}}\rangle$  of the same spin in Eq. (10) as those occur much more efficiently by phonons, see in the subsequent Section. To this end, we take  $\{|\psi_{n_S}\rangle\}$  as exact eigenstates of  $H_0 + H^D$ . From now on we denote by  $n_S$  the subset of  $n$ -values enumerating eigenstates of  $H_0$  that belong to the definite spin  $S$ .

## IV. SPIN RELAXATION RATES

Typically, the electron-phonon interaction  $H^{\text{el-ph}}$  establishes thermal equilibrium between electron and lattice reservoirs on time-scales short compared to the times on which spin changing transitions occur. This is so because the latter cannot be achieved directly by  $H^{\text{el-ph}}$  (cf. Sect. IV A), so that equilibrium will be established rapidly only among dot levels of given total spins. This

suggests to divide the total Hilbert space:

$$\mathcal{H} = \bigoplus_S \mathcal{H}_S$$

of *coupled* electron-phonon states into orthogonal subspaces  $\mathcal{H}_S$ , labeled according to the *electron* spin  $S$ . Transitions among subspaces occur only slowly by the action of  $H^D$  while thermal equilibrium resides within each of the subspaces after much shorter times  $\tau^{\text{el-ph}}$  at the lattice temperature  $(k_B\beta)^{-1}$ . Consider a certain electronic spin state  $S'$ , as it may have been prepared, for example, using electronic transport techniques.<sup>2,44,45</sup> Then, the rate

$$R_{S \leftarrow S'} = \frac{d}{dt} \langle P_S(t) \rangle_{S'} \Big|_{t \gtrsim \tau^{\text{el-ph}}} \quad (11)$$

for its decay into a particular spin  $S \neq S'$  is given as the temporal increase of the spin  $S$ -population  $\langle P_S(t) \rangle_{S'}$ , assuming an initial (i.e. after intra- $\mathcal{H}_{S'}$  equilibration has taken place)  $S'$  thermal equilibrium state,

$$P_{S'} e^{-\beta H} P_{S'} / \text{Tr} \{ P_{S'} e^{-\beta H} P_{S'} \}.$$

Here,  $P_S = \sum_{n_S} |\phi_{n_S}\rangle \langle \phi_{n_S}| \otimes \mathbb{1}^{\text{ph}}$  projects onto  $\mathcal{H}_S$ ,  $\mathbb{1}^{\text{ph}}$  denotes a unit operator on the phonon space. Transition rates  $R_{S \leftarrow S'}$  observe the detailed balance condition, ensuring one vanishing (stationary) eigenvalue of the matrix  $M_{SS'} = R_{S' \leftarrow S} - \tau_{S'}^{-1} \delta_{SS'}$ , which governs the rate dynamics. In the present context we are primarily interested in the total decay rate of the initial spin  $S'$  population; i.e.,

$$\tau_{S'}^{-1} = \sum_{S \neq S'} R_{S \leftarrow S'}. \quad (12)$$

In Eq. (11) the time evolution refers to the Hamiltonian  $H = H_0 + H^D + H^{\text{ph}} + H^{\text{el-ph}}$  where the electron-phonon interaction  $H^{\text{el-ph}}$  will be discussed next. This approach, in principle, accounts for rapid thermalizing spin conserving multi-phonon transitions within subspaces  $\mathcal{H}_S$ .

### A. Coupling to phonons

Electron-phonon coupling in semiconductors has been studied intensively in the 1950-ies and 1960-ies. For homopolar semiconductors, such as Si or Ge, deformation potential coupling<sup>46</sup> has been established. It can be expressed as<sup>47</sup>

$$H^{\text{el-ph}} = \sum_{\mathbf{q}} g_{\mathbf{q}} \rho(\mathbf{q}) (b_{\mathbf{q}} + b_{-\mathbf{q}}^{\dagger}) \quad (13)$$

where we have suppressed the phonon branch index. Considerably below room temperature, pertinent to possible quantum computing, optical phonons don't contribute so that  $b_{\mathbf{q}}^{\dagger}$  in Eq. (13) is meant to create a longitudinal acoustical phonon of momentum  $\mathbf{q}$ . For excitations of the electronic system most relevant are phonon

wave lengths  $2\pi c_s/\omega_{\text{pl}} = 50$  nm or  $2\pi c_s/\Delta = 500$  nm, assuming<sup>35</sup>  $\omega_{\text{pl}} \simeq 3$  meV and  $\Delta \simeq 0.3$  meV, respectively, cf. Sect. III C for the definitions of the energies  $\Delta$  and  $\omega_{\text{pl}}$ . At this wave lengths intra-valley scattering dominates. Its strength

$$g_{\mathbf{q}}^2 = \frac{E_2^2}{2\rho_M V c_s} |\mathbf{q}| \quad (14)$$

mainly is regulated by the deformation potential constant  $E_2$  for longitudinal coupling which takes values of about<sup>47,48</sup> 10 eV in Si. Further,  $g_{\mathbf{q}}^2$  depends on the mass density  $\rho_M$ , the normalization volume  $V$  for the phonon modes, and on the sound velocity  $c_s$ . In Eq. (13) the operator

$$\rho(\mathbf{q}) = N \sum_{n_S, n'_S} \rho_{n_S n'_S}(\mathbf{q}) |\phi_{n'_S}\rangle \langle \phi_{n_S}| \quad (15)$$

of the total electron density may excite the correlated electron system at non-zero  $\mathbf{q}$ , though at conserved total spin  $S$  (and conserved  $z$ -component  $S_z$ ). It can be decomposed into the basis  $\{|\phi_n\rangle\}$ , where  $|\phi_n\rangle$  and  $|\phi_{n'}\rangle$  have same spin  $S$ , i.e.,

$$\rho_{n_S n'_S}(\mathbf{q}) = \int d\mathbf{r} e^{i\mathbf{q}\mathbf{r}} \int d\mathbf{r}_2 \dots d\mathbf{r}_N \langle \mathbf{r}, \mathbf{r}_2, \dots, \mathbf{r}_N | \phi_{n_S} \rangle \times \langle \phi_{n'_S} | \mathbf{r}, \mathbf{r}_2, \dots, \mathbf{r}_N \rangle. \quad (16)$$

At small  $q \equiv |\mathbf{q}|$  these coefficients are expanded,  $\rho_{nn'}(\mathbf{q}) = \delta_{nn'} + \alpha(q\ell)^{\nu}$  where, to lowest non-vanishing order,  $\nu = 1$  unless the electron charge density distribution of the quantum dot or of the ensemble of quantum dots is parity symmetric, in which case  $\nu = 2$ . The quantity  $\ell$  either equals the typical distance between electrons if  $n$  and  $n'$  belong to the same plasmon multiplet, or  $\ell \simeq (m\omega_{\text{pl}})^{-1/2}$  for  $n$  and  $n'$  from different plasmon multiplets. The magnitude of  $\alpha$  can be estimated by inserting Eq. (9) into Eq. (16) and using, for convenience, the density distribution  $\rho_{nn'}(\mathbf{r})$  in real space. This reveals that  $\alpha$  is proportional to the maximum overlap between unequal pocket states, i.e.  $\max_{p, p'; p \neq p'} \langle p | p' \rangle$ , a quantity which, in turn, is proportional<sup>27</sup> to  $\Delta/\omega_{\text{pl}}$ .

### B. Transition rates

We are now in the position to calculate the phonon mediated transition rate Eq. (11) as a result of spin mixing, see Eq. (10). Assuming a not too strong electron-phonon coupling, use of standard time dependent perturbation theory with respect to  $H^{\text{el-ph}}$ , as explicated in Appendix B, yields to leading order the rate  $R_{S \leftarrow S'}$ , cf. Eq. (11), reading

$$R_{S \leftarrow S'} = \frac{2\pi}{Z_{S'}} \sum_{n_S, n_{S'}} e^{-\beta E_{n_{S'}}} J_{n_S, n_{S'}} (|E_{n_{S'}} - E_{n_S}|) \times [n(|E_{n_{S'}} - E_{n_S}|) + \Theta(E_{n_{S'}} - E_{n_S})], \quad (17)$$

where we have defined the (temperature independent) coupled density of phonon states for transitions between spins  $S$  and  $S'$ , respectively, as

$$J_{n_{S'}, n_S}(\omega) = \sum_{\mathbf{q}} g_{\mathbf{q}}^2 \delta(\omega - |\mathbf{q}|c_s) \times \left| \sum_{n'_S} \rho_{n_S n'_S}(\mathbf{q}) \frac{\langle \phi_{n'_S} | H^D | \phi_{n_{S'}} \rangle}{E_{n_{S'}} - E_{n'_S}} - \sum_{n'_{S'}} \rho_{n'_{S'} n_{S'}}(\mathbf{q}) \frac{\langle \phi_{n_S} | H^D | \phi_{n'_{S'}} \rangle}{E_{n'_{S'}} - E_{n_S}} \right|^2. \quad (18)$$

In Eq. (17)  $Z_S = \sum_{n_S} e^{-\beta E_{n_S}}$  denotes the partition function inside the subspace  $\mathcal{H}_S$ ,  $n(\omega) = (e^{\beta\omega} - 1)^{-1}$  the Bose function, and  $\Theta(x)$  the Heavyside step function. Straightforwardly, higher order terms regarding  $H^{\text{el-ph}}$  can also be considered for  $R_{S \leftarrow S'}$ , although the corresponding explicit expressions are rather lengthy. In Eq. (10) we have assumed that phonon states and energy eigenvalues remain unaffected by the weak dipolar mixing.

At low temperatures,  $T \ll \Delta/k_B$  compared to the typical distance  $\Delta$  between dot levels of same or of different total spins inside the lowest plasmon multiplet, the Bose factor  $n(|\Delta|) \ll 1$  is small and only the ground level  $n_{0S}$  will be occupied within each subspace  $\mathcal{H}_S$ . In this temperature regime thermalization into the global ground state  $n = 0_{S_0}$  of spin  $S_0$  will take place exclusively through the direct process by emission of a resonant phonon of energy  $\Delta$  so that the coupled density of states  $J_{n_{0S} 0_{S_0}}(\Delta)$  controls the relaxation rate  $R_{S_0 \leftarrow S}$ . Still, a summation over excited levels  $n'_{S_0} > 0$  and  $n'_{0S} > n_{0S}$  appears in Eq. (18), as the lowest terms  $n'_{S_0} = 0$  and  $n'_{0S} = n_{0S}$  cancel exactly. In Section IV A it has been estimated that non-diagonal coefficients  $\rho_{nn'}(\mathbf{q}) \sim (\Delta/\omega_{\text{pl}})(r\Delta/c_s)^\nu$  for low energy and long wave lengths transitions; here  $r$  denotes the distance between electrons and  $\nu = 1$  or  $\nu = 2$  in the absence or presence of parity symmetry. For the electron-phonon coupling Eq. (14) this results in a spin transition rate at zero temperature through the direct process, reading

$$R_{S_0 \leftarrow S} = 2\pi J_{n_{0S} 0_{S_0}}(\Delta) \quad (19)$$

with

$$J_{n_{0S} 0_{S_0}}(\Delta) = \frac{E_2^2 \gamma^4}{c_s^7 \rho_M \pi^2 \omega_{\text{pl}}^2} N^2 n_a^2 \Delta^5,$$

unless this transition is not suppressed entirely for cases discussed in Sect. III D. In Eq. (19) we have assumed for simplicity that level separations  $E_{n'_{S_0}} - E_{0_{S_0}} \sim \Delta$  and  $E_{n'_{0S}} - E_{n_{0S}} \sim \Delta$  both are of the order<sup>49</sup>  $\Delta$ . Also, we have inserted the areal density  $n_a = r^{-2}$  of electrons, focusing on the measured quantity in two-dimensional samples.

In Si the rate Eq. (19) appears to be very small at zero temperature,  $\sim 10^{-7} \text{ s}^{-1}$  for three electrons at densities corresponding to  $r_s = 1$ , and considering a quantum dot<sup>35</sup> of  $\omega_{\text{pl}} = 3 \text{ meV}$  and  $\Delta = 0.3 \text{ meV}$ . However, this number strongly varies with parameters as seen in Eq. (19). Parity symmetric quantum dots (where  $\nu = 2$ ) would suppress this decay rate even further at small transition energies due to  $J_{n_0}(\Delta) \sim \Delta^7$  in this case. These values are, of course, considerable smaller than the decay rates estimated from spin orbit effects,<sup>7,8,9</sup> if present.

They are also smaller than the rates estimated from the hyperfine interaction with nuclei of non-zero spin.<sup>13,14,16</sup> Particularly in Ref. 13 a hybrid mechanism is considered which is closely related to the one presented here in combining the electron-phonon coupling with a spin-mixing interaction. Transitions between total spin  $S = 1$  and  $S = 0$  of a two-electron quantum dot are investigated. The low temperature rate has been estimated<sup>13</sup> to  $\sim 10^{-2} \text{ s}^{-1}$  for similar quantum dot parameters as above, assuming a two-dimensional dot fabricated on the basis of heterostructures. This rate is proportional to the number  $N_n$  of non-vanishing nuclear spins covered by the electron wave function. In GaAs almost every nucleus has spin  $I = 3/2$ . It is instructive to determine from this result<sup>13</sup> a critical concentration  $\tilde{C}_n$  of  $^{29}\text{Si}$  nuclei in silicon, the only ones of non-vanishing spin  $I = 1/2$ , beyond which the here described dipolar mechanism should prevail over the spin decay via nuclear spins. For a quantum dot of the same excitation energy, the electron wave function in natural silicon covers only about  $N_n^{\text{Si}} \approx 10^3$  of the  $^{29}\text{Si}$  nuclei while<sup>14</sup> in GaAs  $N_n^{\text{GaAs}} \approx 10^5$ . Two further important differences between Si and GaAs have to be taken into account. Firstly, the type of electron-phonon coupling which is piezo-elastic in GaAs while we have deformation potential coupling in Si. Accidentally, for the here considered quantum dot parameters (and assuming again heterostructures and now laterally parabolic confining potential)  $J_{nn'}(\omega)$  in Si is only by 0.6 smaller than in GaAs. Secondly, the nuclear spin  $I^{\text{Si}} = 1/2$  of  $^{29}\text{Si}$  as compared to  $I^{\text{GaAs}} = 3/2$  in GaAs which reduces the coupling by  $I^{\text{Si}}(I^{\text{Si}} + 1)/I^{\text{GaAs}}(I^{\text{GaAs}} + 1) = 1/5$ . This yields  $\tilde{C}_n^{\text{Si}} \approx 2 \times 10^{-4} \text{ nm}^{-3}$  (note that  $C_n^{\text{Si}} \approx 2.5 \times 10^{-3} \text{ nm}^{-3}$  has been reported<sup>50</sup> experimentally). This value is less stringent than the isotopic purification required for the quantum computer,<sup>6</sup> based on the nuclear spins of  $^{31}\text{P}$  donors, where  $C_n$  should be smaller than  $N^{-1} 10^{-4} \text{ nm}^{-3}$  in Si with  $N$  being the number of qubits.

## C. Temperature Dependence

Through the marked increase of  $J_{n_0}(\omega) \sim \omega^{4+2\nu}$  as a function of transition energy  $\omega$ , relaxation can take advantage from spin conserving thermal excursions to plasmonic excited levels and accomplish the spin transition at an elevated energy. In NMR-theory this possibility is called the ‘Orbach process’<sup>51</sup> and shows up in a steeply increasing relaxation rate with temperature. Our formu-



lation, Eq. (17), of the transition rate explicitly incorporates such thermal excursions. They turn out to influence considerably the temperature dependence of  $R_{S_0 \leftarrow S}$  which only at low temperatures follows the Bose-behavior  $\sim J_{n0}(\Delta)n(\Delta)$  of direct transitions. Already at temperatures not much exceeding  $\Delta$  the severely stronger increase  $\sim J_{n0}(\omega_{pl}) \exp(-\omega_{pl}/k_B T)$ , following from Eq. (17), can easily enhance the transition rate by three orders of magnitudes, depending on system parameters. A similarly pronounced increase of (nuclear) spin relaxation rates has been discussed in detail<sup>52</sup> in the context of quantum rotating molecules: substantial increases in spin changing transition rates by more than six orders of magnitudes are depicted with Figure 3b of Ref. 52. Again, the stronger increasing density of coupled phonon states in quantum dots of parity symmetry should lead to even more pronounced temperature sensitivity.

## V. RESUMÉ

We have investigated dipolar interactions between the magnetic moments of electrons confined to one or to several quantum dots and studied the rate of inelastic total spin changing transitions. As compared to the coupling to nuclear spins<sup>13,14,15,16,53</sup> and to spin-orbit induced decay,<sup>8,9,10,11,12</sup> dipolar spin decay turns out as much weaker. However, either of the earlier studied mechanisms can, at least in principle, be eliminated by a suitable sample design. It is therefore possible that the dipolar interactions between electronic spin moments, together with the coupling to lattice modes,<sup>20</sup> as discussed in the present work, will ultimately limit long time quantum computations, even when devices become optimally designed. Experimentally, the dipolar mechanism should show up most directly by observing at low temperatures the dependence on the electron density, cf. Eq. (19).

Upon generalizing previous approaches we incorporate here electron-electron interactions<sup>11,12</sup> in the quantum dot(s) which, additionally, are important for magnetic features.<sup>22,23,24,25,31</sup> For example, ground state spins may exceed the values  $S_0 = 0$  or  $S_0 = 1/2$  expected for non-interacting electrons. We focus on the limit of strong interactions, where electronic many-body wave functions can be described as ‘pocket states’<sup>26,27,28</sup> and where the spectrum exhibits spin-split plasmon multiplets. The dipolar interaction is decomposed according to the symmetric group and non-vanishing matrix elements are determined in their dependence on spatial parts of the collective electron wave functions. As we have shown, particular spatial symmetries of the quantum dot(s) can reduce the number of non-zero elements.

One important result is the stability of  $N = 2$  electron spins. Irrespective of the dimensionality, the shape of the quantum dot(s), or of the electron-electron interaction strength, the decay of triplet states into the singlet ground state is always suppressed. Any dipolar

decay channel will require participation of further electrons. This is important, for example, for two coupled qubits, the basic (gate)-element for quantum information processing.

Also at larger electron numbers, non-ground state spins can be stable with respect to dipolar interactions. We have discussed the case of  $N = 3$  electrons on an equilateral triangle. Here,  $S_z = \pm 1/2$  states of the  $S = 3/2$  sub-manifold prove robust against decay into any  $S = 1/2$  state. Only spin polarized  $S_z = \pm 3/2$  states decay into the  $S = 1/2$  sub-manifold. Further,  $N = 4$  electrons on a square shaped quantum dot exhibit robust  $S = 1$  states of  $T_x$  and  $T_y$  symmetry, and  $S = 0$  states of  $E_2$  symmetry.

Owing to the smallness of magnetic and in particular dipolar energies, compared to dot level separations, the energy accompanied with an actual spin transition has to be provided by the reservoir of lattice vibrations. As in previous approaches<sup>7,8,9,10,13</sup> we have considered the coupling to acoustic phonons. Parity symmetric dots are weaker coupled to phonons, which further suppresses spin decay in this case. Additionally, we have accounted for rapid thermal excursions of the system *within* electron-phonon subspaces of given (many electron) dot spins. This enables one to deduce the dependence of spin relaxation over a wider range of temperatures as compared to the resonant direct process. As a result we found a very striking increase of the spin decay rate. This rate grows with temperature considerably steeper than the naively expected proportionality to the Bose function describing direct processes: It occurs already at temperatures that barely exceed the energy difference between the lowest levels of different spins, but is still considerably smaller than the energy for plasmon excitations. Although we find amazingly stable spin configurations at low temperatures this marked temperature sensitivity restricts the operation temperatures of quantum computing dots (unless quantum computation can be confined to the stable spin configurations) to values that are not exceeding much the lowest level separations.

Because the same phonon energy reservoir is considered in previous work<sup>7,8,9,10,13,14</sup> for spin decay, a similar scenario regarding the dot symmetries and the spin conserving phonon induced excursions should apply also to magnetic mechanisms of the spin-orbit or of the hyperfine type. We expect therefore a similarly striking temperature sensitivity as obtained here for these mechanisms as well.

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**APPENDIX A: SYMMETRIZED OPERATORS INDUCING SPIN CHANGING TRANSITIONS FOR  
 $N = 3$  AND  $N = 4$  ELECTRONS**

The following explicit form of dipolar operators Eq. (8) contain non-vanishing elements for  $N = 3$ :

$$\begin{aligned}
 H^{(0)[2,1]} &= \sum_{i \neq j} H_{ij}^{(0)} - \frac{1}{6} \left\{ \left[ \frac{1}{4} \sum_{i \neq j} |\varrho_{ij}|^2 \right] \left[ \sum_{i \neq j} (\hat{S}_{+i} \hat{S}_{-j} + \hat{S}_{-i} \hat{S}_{+j}) \right] \right. \\
 &\quad \left. + \left[ \sum_{i \neq j} \zeta_{ij}^2 \right] \left[ \sum_{i \neq j} \hat{S}_{zi} \hat{S}_{zj} \right] \right\} \\
 H^{(1)[2,1]} &= \sum_{i \neq j} H_{ij}^{(1)} - \frac{1}{12} \left\{ \left[ \sum_{i \neq j} \zeta_{ij} \varrho_{-ij} \right] \left[ \sum_{i \neq j} (\hat{S}_{+i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{+j}) \right] \right. \\
 &\quad \left. + \left[ \sum_{i \neq j} \zeta_{ij} \varrho_{+ij} \right] \left[ \sum_{i \neq j} (\hat{S}_{-i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{-j}) \right] \right\} \\
 H^{(2)[2,1]} &= \sum_{i \neq j} H_{ij}^{(2)} - \frac{1}{24} \left\{ \left[ \sum_{i \neq j} \varrho_{+ij}^2 \right] \left[ \sum_{i \neq j} \hat{S}_{-i} \hat{S}_{-j} \right] + \left[ \sum_{i \neq j} \varrho_{-ij}^2 \right] \left[ \sum_{i \neq j} \hat{S}_{+i} \hat{S}_{+j} \right] \right\}
 \end{aligned}$$

and for  $N = 4$ :

$$\begin{aligned}
H^{(0)[3,1]} &= \frac{1}{4} \sum_{i \neq j} \left\{ \left[ \frac{|\varrho_{ij}|^2 - |\varrho_{\bar{i}\bar{j}}|^2}{4} \right] \left[ (\hat{S}_{+i} \hat{S}_{-j} + \hat{S}_{-i} \hat{S}_{+j}) - (\hat{S}_{+\bar{i}} \hat{S}_{-\bar{j}} + \hat{S}_{-\bar{i}} \hat{S}_{+\bar{j}}) \right] \right. \\
&\quad \left. + \left[ \zeta_{ij}^2 - \zeta_{\bar{i}\bar{j}}^2 \right] \left[ \hat{S}_{zi} \hat{S}_{zj} - \hat{S}_{z\bar{i}} \hat{S}_{z\bar{j}} \right] \right\} \\
H^{(1)[3,1]} &= \frac{1}{8} \sum_{i \neq j} \left\{ \left[ \zeta_{ij} \varrho_{-ij} - \zeta_{\bar{i}\bar{j}} \varrho_{-\bar{i}\bar{j}} \right] \left[ (\hat{S}_{+i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{+j}) - (\hat{S}_{+\bar{i}} \hat{S}_{z\bar{j}} + \hat{S}_{z\bar{i}} \hat{S}_{+\bar{j}}) \right] \right. \\
&\quad \left. + \left[ \zeta_{ij} \varrho_{+ij} - \zeta_{\bar{i}\bar{j}} \varrho_{+\bar{i}\bar{j}} \right] \left[ (\hat{S}_{-i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{-j}) - (\hat{S}_{-\bar{i}} \hat{S}_{z\bar{j}} + \hat{S}_{z\bar{i}} \hat{S}_{-\bar{j}}) \right] \right\} \\
H^{(2)[3,1]} &= \frac{1}{16} \sum_{i \neq j} \left\{ \left[ \varrho_{+ij}^2 - \varrho_{+\bar{i}\bar{j}}^2 \right] \left[ \hat{S}_{-i} \hat{S}_{-j} - \hat{S}_{-\bar{i}} \hat{S}_{-\bar{j}} \right] + \left[ \varrho_{-ij}^2 - \varrho_{-\bar{i}\bar{j}}^2 \right] \left[ \hat{S}_{+i} \hat{S}_{+j} - \hat{S}_{+\bar{i}} \hat{S}_{+\bar{j}} \right] \right\} \\
H^{(0)[2,2]} &= \frac{1}{4} \sum_{i \neq j} \left\{ \left[ \frac{|\varrho_{ij}|^2 + |\varrho_{\bar{i}\bar{j}}|^2}{4} \right] \left[ (\hat{S}_{+i} \hat{S}_{-j} + \hat{S}_{-i} \hat{S}_{+j}) + (\hat{S}_{+\bar{i}} \hat{S}_{-\bar{j}} + \hat{S}_{-\bar{i}} \hat{S}_{+\bar{j}}) \right] \right. \\
&\quad \left. + \left[ \zeta_{ij}^2 + \zeta_{\bar{i}\bar{j}}^2 \right] \left[ \hat{S}_{zi} \hat{S}_{zj} + \hat{S}_{z\bar{i}} \hat{S}_{z\bar{j}} \right] \right\} \\
&\quad - \frac{1}{12} \left\{ \left[ \frac{1}{4} \sum_{i \neq j} |\varrho_{ij}|^2 \right] \left[ \sum_{i \neq j} (\hat{S}_{+i} \hat{S}_{-j} + \hat{S}_{-i} \hat{S}_{+j}) \right] + \left[ \sum_{i \neq j} \zeta_{ij}^2 \right] \left[ \sum_{i \neq j} \hat{S}_{zi} \hat{S}_{zj} \right] \right\} \\
H^{(1)[2,2]} &= \frac{1}{8} \sum_{i \neq j} \left\{ \left[ \zeta_{ij} \varrho_{-ij} + \zeta_{\bar{i}\bar{j}} \varrho_{-\bar{i}\bar{j}} \right] \left[ (\hat{S}_{+i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{+j}) + (\hat{S}_{+\bar{i}} \hat{S}_{z\bar{j}} + \hat{S}_{z\bar{i}} \hat{S}_{+\bar{j}}) \right] \right. \\
&\quad \left. + \left[ \zeta_{ij} \varrho_{+ij} + \zeta_{\bar{i}\bar{j}} \varrho_{+\bar{i}\bar{j}} \right] \left[ (\hat{S}_{-i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{-j}) + (\hat{S}_{-\bar{i}} \hat{S}_{z\bar{j}} + \hat{S}_{z\bar{i}} \hat{S}_{-\bar{j}}) \right] \right\} \\
&\quad - \frac{1}{24} \left\{ \left[ \sum_{i \neq j} \zeta_{ij} \varrho_{-ij} \right] \left[ \sum_{i \neq j} (\hat{S}_{+i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{+j}) \right] \right. \\
&\quad \left. + \left[ \sum_{i \neq j} \zeta_{ij} \varrho_{+ij} \right] \left[ \sum_{i \neq j} (\hat{S}_{-i} \hat{S}_{zj} + \hat{S}_{zi} \hat{S}_{-j}) \right] \right\} \\
H^{(2)[2,2]} &= \frac{1}{16} \sum_{i \neq j} \left\{ \left[ \varrho_{+ij}^2 + \varrho_{+\bar{i}\bar{j}}^2 \right] \left[ \hat{S}_{-i} \hat{S}_{-j} + \hat{S}_{-\bar{i}} \hat{S}_{-\bar{j}} \right] + \left[ \varrho_{-ij}^2 + \varrho_{-\bar{i}\bar{j}}^2 \right] \left[ \hat{S}_{+i} \hat{S}_{+j} + \hat{S}_{+\bar{i}} \hat{S}_{+\bar{j}} \right] \right\} \\
&\quad - \frac{1}{48} \left\{ \left[ \sum_{i \neq j} \varrho_{+ij}^2 \right] \left[ \sum_{i \neq j} \hat{S}_{-i} \hat{S}_{-j} \right] + \left[ \sum_{i \neq j} \varrho_{-ij}^2 \right] \left[ \sum_{i \neq j} \hat{S}_{+i} \hat{S}_{+j} \right] \right\}.
\end{aligned}$$

In the above expression,  $(\bar{i}, \bar{j})$  take the two values out of  $1, \dots, 4$  that are both different from  $(i, j)$ .

## APPENDIX B: DERIVATION OF EQ. (17)

In a perturbative expansion w.r.t.  $H^{\text{el-ph}}$  of either of the two time evolution operators appearing in Eq. (11) we write

$$\begin{aligned}
e^{-iHt} &= e^{-i\tilde{H}_0 t} \left[ \mathbb{1}^{\text{ph}} - i \int_0^t dt' e^{i\tilde{H}_0 t'} H^{\text{el-ph}} e^{-i\tilde{H}_0 t'} \right. \\
&\quad \left. - \int_0^t dt' \int_0^{t'} dt'' e^{i\tilde{H}_0 t'} H^{\text{el-ph}} e^{-i\tilde{H}_0(t'-t'')} H^{\text{el-ph}} e^{-i\tilde{H}_0 t''} + \dots \right].
\end{aligned} \tag{B1}$$

Here,  $\tilde{H}_0 = H_0 + H^D + H^{\text{ph}}$  with  $H_0$  defined in Eq. (1) and the eigenstates of  $H_0 + H^D$  taken according to Eq. (10). To second (i.e. lowest non-vanishing) order in  $H^{\text{el-ph}}$  only two of the second terms in the square bracket of (B1) contribute to Eq. (11), yielding with (13)

$$R_{S \leftarrow S'} = \frac{1}{Z_{S'}} \sum_{\mathbf{q}} g_{\mathbf{q}}^2 \sum_{n_S n_{S'}} e^{-\beta E_{n_{S'}}} \frac{d}{dt} \int_0^t dt' \int_0^t dt'' i \langle \phi_{n_{S'}} | e^{i(H_0 + H^D)t'} \rho(\mathbf{q}) e^{-i(H_0 + H^D)t'} | \phi_{n_S} \rangle$$

$$\times (-i) \langle \phi_{n_S} | e^{i(H_0 + H^D)t''} \rho(\mathbf{q}) e^{-i(H_0 + H^D)t''} | \phi_{n_{S'}} \rangle \left[ \bar{n}_{\mathbf{q}} e^{i c_s |\mathbf{q}|(t' - t'')} + (\bar{n}_{\mathbf{q}} + 1) e^{-i c_s |\mathbf{q}|(t' - t'')} \right].$$

Here, the Bose factors  $\bar{n}_{\mathbf{q}} = (e^{\beta c_s |\mathbf{q}|} - 1)^{-1}$  result after thermal averaging over phonon modes. Inserting now eigenstates  $|\psi_n\rangle$  of  $(H_0 + H^D)$  for the  $|\phi_n\rangle$ , according to (10), and carrying out the long time limit  $t \rightarrow \infty$ , yields for the decay rate

$$R_{S \leftarrow S'} = \frac{2\pi}{Z_{S'}} \sum_{\mathbf{q}} g_{\mathbf{q}}^2 \sum_{n_S n_{S'}} e^{-\beta E_{n_{S'}}}$$

$$\times \left[ i \left( \langle \psi_{n_{S'}} | - \sum_{n_{S''}} \frac{\langle \phi_{n_{S'}} | H^D | \phi_{n_{S''}} \rangle}{E_{n_{S'}} - E_{n_{S''}}} \langle \psi_{n_{S''}} | \right) \rho(\mathbf{q}) \right.$$

$$\times \left( | \psi_{n_S} \rangle - \sum_{n_{S''}} \frac{\langle \phi_{n_{S''}} | H^D | \phi_{n_S} \rangle}{E_{n_S} - E_{n_{S''}}} | \psi_{n_{S''}} \rangle \right) \Big]$$

$$\times \left[ -i \left( \langle \psi_{n_S} | - \sum_{n_{S''}} \frac{\langle \phi_{n_S} | H^D | \phi_{n_{S''}} \rangle}{E_{n_S} - E_{n_{S''}}} \langle \psi_{n_{S''}} | \right) \rho(\mathbf{q}) \right.$$

$$\times \left( | \psi_{n_{S'}} \rangle - \sum_{n_{S''}} \frac{\langle \phi_{n_{S''}} | H^D | \phi_{n_{S'}} \rangle}{E_{n_{S'}} - E_{n_{S''}}} | \psi_{n_{S''}} \rangle \right) \Big]$$

$$\times [\bar{n}_{\mathbf{q}} \delta(E_{n_{S'}} - E_{n_S} + c_s |\mathbf{q}|) + (\bar{n}_{\mathbf{q}} + 1) \delta(E_{n_{S'}} - E_{n_S} - c_s |\mathbf{q}|)],$$

which is readily brought into the form (17) with (18), after employing (15) and using the orthogonality of spin states. Further calculational details can be found in Ref. 52.

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