



Cite this: *Phys. Chem. Chem. Phys.*,  
2024, 26, 8952

# What governs magnetic exchange couplings in radical-bridged dinuclear complexes?†

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Coupling transition metal or lanthanide ions through a radical bridging ligand is a promising route to increase performances in the area of single molecular magnets. A better understanding of the underlying physical mechanisms governing the magnetic exchange couplings is thus of valuable importance to design future compounds. Here, couplings in three series of metal–radical–metal compounds based on transition metal ions are investigated by means of the decomposition/recomposition methods. This work presents the generalisation and first application of the method to systems with an arbitrary number of magnetic centres featuring several unpaired electrons. Thanks to the decomposition into the three main contributions (direct exchange, kinetic exchange, and spin polarisation) as well as a description in terms of electron–electron interactions, we study the influence of the nature of the metal centre and the radical ligand on the couplings. We combine the energetic contributions extracted with orbital and charge population analysis to rationalise the results.

Received 22nd December 2023,  
Accepted 1st February 2024

DOI: 10.1039/d3cp06243k

rsc.li/pccp

Single-molecule magnets (SMM) are molecules capable of retaining magnetisation under a certain temperature, so-called blocking temperature. In the search for compounds with high blocking temperatures, complexes with radical bridging transition metal- or lanthanide-based paramagnetic centres have emerged as one of the most promising families of molecules.<sup>1–7</sup> Indeed, the couplings occurring between each ion centre and the radical yield a strong indirect ferromagnetic coupling which may increase this property.<sup>8,9</sup> Understanding the physical mechanisms governing these couplings is thus of great interest in guiding the design of future compounds.

Theoretical and computational chemistry, where the magnetic exchange coupling  $J$  is computed as the energy difference between the spin states, has played a critical role in providing molecular magnetism with both conceptual tools and numerical analysis.<sup>10</sup> Wave-function-based methods have been specially well suited for this dual purpose where the dynamic correlation is usually accounted on top of a CASSCF wave-function,<sup>11</sup> through mostly CI<sup>12</sup> or PT2 treatment.<sup>13–18</sup> This strategy provides highly accurate results where the proper physical description of the multiconfigurational character of the spin states is essential. In addition, the resulting wave-functions are dense objects from which much information may be extracted with the combination of model Hamiltonians and

the effective Hamiltonian theory, providing a deep rationalisation of the physical mechanisms of magnetic phenomena.<sup>10,19–23</sup> However, this sophistication results in a computational cost which is most of the time prohibitive for real-life applications.

To overcome this limitation, density functional theory (DFT) in its Kohn–Sham formulation<sup>24,25</sup> has been the most used alternative with the first calculations of magnetic exchange couplings in the early 80's.<sup>26,27</sup> However, this strategy is not exempt from its own drawbacks since (i) KS-DFT is intrinsically a monodeterminantal theory and the computation of  $J$  resorts to the use of Broken-Symmetry (BS) configurations,<sup>26–35</sup> (ii) there is a strong dependence on the exchange–correlation functional which has been widely studied over the last two decades<sup>22,36–42</sup> and (iii) the fact that this approach provides an energy difference only but no information about the underlying physical mechanisms.

Ferré *et al.* have provided a solution to this last limitation with the decomposition method, which allows decomposing the magnetic exchange coupling computed in DFT in its 3 main physical contributions: the ferromagnetic direct exchange  $J_0$  between the magnetic centres, the antiferromagnetic kinetic exchange  $\Delta J_{KE}$  resulting from the relaxation of the magnetic orbitals and the spin polarisation  $\Delta J_{SP}$  due to the different field felt by the non-magnetic electrons in the different states. This decomposition based on the Heisenberg–Dirac–van Vleck (HdVV) Hamiltonian has been successfully applied in various situations with two unpaired electrons such as dinuclear complexes,<sup>43</sup> quantum spin liquids<sup>44</sup> or radical systems.<sup>45</sup> Recently, the present authors have shown the relevance of the method in dinuclear complexes with more than one unpaired electron per magnetic

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† Electronic supplementary information (ESI) available: Structures, additional orbital plots and details of calculations. See DOI: <https://doi.org/10.1039/d3cp06243k>



centre.<sup>46</sup> In addition, this method has been generalised to systems with an arbitrary number of one-unpaired-electron-based magnetic sites by employing the more general Hubbard Hamiltonian.<sup>47–49</sup> This strategy provides a solution to the long-standing question of the spin contamination of  $J$  couplings in multicentre systems by summing up these three properly spin-decontaminated contributions.<sup>49</sup>

This work presents the complete generalisation of this newly developed decomposition/recomposition method to polynuclear magnetic systems featuring several unpaired electrons per magnetic centre. This method is applied to metal–radical–metal complexes to investigate both the influence of the metal centre and the role of the bridging ligand on the magnetic exchange couplings. This paper is organised as follows. Section 1 presents the computation of the different contributions to the magnetic exchange couplings in multicentre systems with a special focus on the  $t$  and  $U$  Hubbard Hamiltonian parameters in the context of several unpaired electrons. In Section 2, we present the three series of compounds studied in this work, *i.e.* the M–tetraazalene–M (with M = Cr, Mn, Fe and Co) series,<sup>50</sup> the M–semiquinone–M (with M = Fe, Co and Ni) series<sup>51</sup> and a series based on bis(bidentate) benzosemiquinoid-based radical bridging two Cr ions.<sup>52</sup> Details of calculations are provided in Section 3. Section 4 discusses the decomposition in the three series, corroborated with orbital and charge population analysis. Finally, an open-source input files generator code that we have developed to ease the use of the method is shortly introduced in Section S1 of the ESI.†

## 1 Methodology

Magnetic exchange couplings are mostly described through the HDvV Hamiltonian,

$$\hat{H}^{\text{HDvV}} = \sum_{I,J} -2J_{IJ} \hat{S}_I \cdot \hat{S}_J \quad (1)$$

where  $J_{IJ}$  expresses the strength of the coupling between two local high-spin sites  $I$  and  $J$  and  $\hat{S}_I$  and  $\hat{S}_J$  their local spin operators, respectively. Based on the pioneering work of De Loth *et al.*<sup>53</sup> and later Calzado *et al.*<sup>19–21</sup> in wave-function theory, Ferré *et al.*<sup>54,55</sup> and recently in a more general form David *et al.*<sup>47–49</sup> decomposed in KS-DFT the magnetic exchange coupling as,

$$J_{IJ} = J_0^{\text{H}} + \Delta J_{\text{KE}}^{\text{H}} + \Delta J_{\text{SP}}^{\text{H}} \quad (2)$$

The following discussion presents the extraction of these three contributions and uses the notation introduced in ref. 49, of which the reader may refer to for further details. Let us consider a system with  $\mathcal{N}$  magnetically coupled local high-spin sites labelled  $A, B, C, \dots, N$ , bearing  $n_A, n_B, n_C, \dots, n_N$  unpaired electrons, respectively, resulting in a maximum of  $\mathcal{N}(\mathcal{N} - 1)/2$  couplings. To each magnetic centre is associated magnetic orbitals labelled with their lowercase letter  $a_\alpha, b_\beta, \dots$  and whose Greek letters are used for the summation over the number of unpaired electrons. Finally,  $i$  and  $j$  correspond to core orbitals, *i.e.* non-magnetic occupied ones, and  $r$  and  $s$  to virtual orbitals.

### 1.1 Direct exchange

The starting point of the method is the computation of the high-spin (HS) state in the restricted open-shell (RO) or in the quasi-restricted open-shell (QRO) formalism.<sup>56</sup> This determinant of a  $(n_A + n_B + n_C + \dots + n_N + 1)$  spin multiplicity defines a set of  $n_{\text{core}}$  doubly occupied core orbitals  $i$  and a set of  $n_A + n_B + n_C + \dots + n_N$  singly occupied molecular orbitals (SOMOs), of which their localisation yields,

$$\Phi_{\text{HS,RO}} = \Phi_{ABC\dots N,\text{RO}} = \left| \prod_i^{n_{\text{core}}} \bar{ii} \prod_\alpha^{n_A} a_\alpha \prod_\beta^{n_B} b_\beta \prod_\gamma^{n_C} c_\gamma \dots \prod_\nu^{n_N} n_\nu \right| \quad (3)$$

To determine the  $\mathcal{N}(\mathcal{N} - 1)/2$  direct exchange contributions,  $\mathcal{N}(\mathcal{N} - 1)/2$  energy differences between the HS and some BS determinants must be defined. One may start by considering the  $\mathcal{N}$  configurations whose all spins of a magnetic site are flipped from the HS,RO determinant without optimising any orbital such as,

$$\Phi_{\bar{A}BC\dots N,\text{NO}} = \left| \prod_i^{n_{\text{core}}} \bar{ii} \prod_\alpha^{n_A} \bar{a}_\alpha \prod_\beta^{n_B} b_\beta \prod_\gamma^{n_C} c_\gamma \dots \prod_\nu^{n_N} n_\nu \right| \quad (4)$$

$$\Phi_{A\bar{B}C\dots N,\text{NO}} = \left| \prod_i^{n_{\text{core}}} \bar{ii} \prod_\alpha^{n_A} a_\alpha \prod_\beta^{n_B} \bar{b}_\beta \prod_\gamma^{n_C} c_\gamma \dots \prod_\nu^{n_N} n_\nu \right| \quad (5)$$

$$\Phi_{AB\bar{C}\dots N,\text{NO}} = \left| \prod_i^{n_{\text{core}}} \bar{ii} \prod_\alpha^{n_A} a_\alpha \prod_\beta^{n_B} b_\beta \prod_\gamma^{n_C} \bar{c}_\gamma \dots \prod_\nu^{n_N} n_\nu \right| \quad (6)$$

⋮

where NO stands for non-optimised. This leads to the following energy differences,

$$E[\Phi_{\bar{A}BC\dots N,\text{NO}}] - E[\Phi_{ABC\dots N,\text{RO}}] = \sum_{K=1, K \neq A}^{\mathcal{N}} 4S_A S_K J_0^{\text{AK}} \quad (7)$$

$$E[\Phi_{A\bar{B}C\dots N,\text{NO}}] - E[\Phi_{ABC\dots N,\text{RO}}] = \sum_{K=1, K \neq B}^{\mathcal{N}} 4S_B S_K J_0^{\text{BK}} \quad (8)$$

$$E[\Phi_{AB\bar{C}\dots N,\text{NO}}] - E[\Phi_{ABC\dots N,\text{RO}}] = \sum_{K=1, K \neq C}^{\mathcal{N}} 4S_C S_K J_0^{\text{CK}} \quad (9)$$

⋮

with  $S_A, S_B, \dots$  the spin of the magnetic site  $A, B, \dots$ , respectively. The left  $\mathcal{N}(\mathcal{N} - 3)/2$  energy differences may be defined through the BS determinants with two magnetic sites flipped as,

$$\Phi_{\bar{A}\bar{B}C\dots N,\text{NO}} = \left| \prod_i^{n_{\text{core}}} \bar{ii} \prod_\alpha^{n_A} \bar{a}_\alpha \prod_\beta^{n_B} \bar{b}_\beta \prod_\gamma^{n_C} c_\gamma \dots \prod_\nu^{n_N} n_\nu \right| \quad (10)$$

$$\Phi_{\bar{A}B\bar{C}\dots N,\text{NO}} = \left| \prod_i^{n_{\text{core}}} \bar{ii} \prod_\alpha^{n_A} \bar{a}_\alpha \prod_\beta^{n_B} b_\beta \prod_\gamma^{n_C} \bar{c}_\gamma \dots \prod_\nu^{n_N} n_\nu \right| \quad (11)$$



$$\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} = \left| \prod_i^{n_{\text{core}}} i \bar{i} \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} \bar{c}_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (12)$$

$$\vdots$$

yielding,

$$E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] - E[\Phi_{ABC\dots N, \text{RO}}] = \sum_{K=1, K \neq A, B}^{\mathcal{N}} 4S_A S_K J_0^{AK} + 4S_B S_K J_0^{BK} \quad (13)$$

$$E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] - E[\Phi_{ABC\dots N, \text{RO}}] = \sum_{K=1, K \neq A, C}^{\mathcal{N}} 4S_A S_K J_0^{AK} + 4S_C S_K J_0^{CK} \quad (14)$$

$$E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] - E[\Phi_{ABC\dots N, \text{RO}}] = \sum_{K=1, K \neq B, C}^{\mathcal{N}} 4S_B S_K J_0^{BK} + 4S_C S_K J_0^{CK} \quad (15)$$

$$\vdots$$

The  $\mathcal{N}(\mathcal{N}-1)/2$  direct exchange contributions may be extracted from these two sets of energy differences by solving the set of linear equations.

## 1.2 Spin polarisation contribution

The spin polarisation contribution corresponds to the different responses felt by the core orbitals due to the different fields created by the magnetic electrons in the high and low spin states. Its extraction proceeds through the relaxation of the core orbitals in the previously generated HS(Q)RO and BS,NO determinants and results in a substantially similar set of equations to solve as for the direct exchange. For the sake of conciseness, only the main working equations are presented and the reader should refer to ref. 49 for further discussions. Starting from the HS,RO determinant, the relaxation of the core orbitals is allowed while the magnetic orbitals are kept frozen,

$$\Phi_{ABC\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} b_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (16)$$

where FM stands for frozen magnetic orbitals and the prime for the relaxed orbitals. Relaxing the core orbitals of the BS determinants with one magnetic centre flipped,

$$\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} \bar{a}_{\alpha} \prod_{\beta}^{n_B} b_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (17)$$

$$\Phi_{A\bar{B}\bar{C}\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (18)$$

$$\Phi_{ABC\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} b_{\beta} \prod_{\gamma}^{n_C} \bar{c}_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (19)$$

$$\vdots$$

and of those with the spin of two magnetic centres flipped,

$$\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} \bar{a}_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (20)$$

$$\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} \bar{a}_{\alpha} \prod_{\beta}^{n_B} b_{\beta} \prod_{\gamma}^{n_C} \bar{c}_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (21)$$

$$\Phi_{A\bar{B}\bar{C}\dots N, \text{FM}} = \left| \prod_i^{n_{\text{core}}} i' i'' \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} \bar{c}_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \quad (22)$$

$$\vdots$$

result in a set of  $\mathcal{N}(\mathcal{N}-1)/2$  energy differences from which the direct exchange is retrieved,

$$(E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) - (E[\Phi_{ABC\dots N, \text{FM}}] - E[\Phi_{ABC\dots N, \text{RO}}]) = \sum_{K=1, K \neq A}^{\mathcal{N}} 4S_A S_K \Delta J_{\text{SP}}^{AK} \quad (23)$$

$$(E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) - (E[\Phi_{ABC\dots N, \text{FM}}] - E[\Phi_{ABC\dots N, \text{RO}}]) = \sum_{K=1, K \neq B}^{\mathcal{N}} 4S_B S_K \Delta J_{\text{SP}}^{BK} \quad (24)$$

$$(E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) - (E[\Phi_{ABC\dots N, \text{FM}}] - E[\Phi_{ABC\dots N, \text{RO}}]) = \sum_{K=1, K \neq C}^{\mathcal{N}} 4S_C S_K \Delta J_{\text{SP}}^{CK} \quad (25)$$

$$\vdots$$

$$(E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) - (E[\Phi_{ABC\dots N, \text{FM}}] - E[\Phi_{ABC\dots N, \text{RO}}]) = \sum_{K=1, K \neq A, B}^{\mathcal{N}} 4S_A S_K \Delta J_{\text{SP}}^{AK} + 4S_B S_K \Delta J_{\text{SP}}^{BK} \quad (26)$$

$$(E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) - (E[\Phi_{ABC\dots N, \text{FM}}] - E[\Phi_{ABC\dots N, \text{RO}}]) = \sum_{K=1, K \neq A, C}^{\mathcal{N}} 4S_A S_K \Delta J_{\text{SP}}^{AK} + 4S_C S_K \Delta J_{\text{SP}}^{CK} \quad (27)$$



$$\begin{aligned}
& (E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) - (E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{FM}}] \\
& - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{RO}}]) = \sum_{K=1, K \neq B, C}^N 4S_B S_K \Delta J_{\text{SP}}^{BK} + 4S_C S_K \Delta J_{\text{SP}}^{CK} \\
& \vdots
\end{aligned} \quad (28)$$

### 1.3 Kinetic exchange

The kinetic exchange contribution is the main contribution to spin-decontaminate and this aspect is particularly challenging in systems involving several couplings since the Yamaguchi formula is no longer valid, whilst Shoji *et al.* provided an extension to multicentre systems based on spin correlation functions.<sup>49,57</sup> To overcome the use of a spin-decontamination formula, some of the present authors proposed to proceed through the extraction of the  $t$  and  $U$  Hubbard parameters associated with one pair of magnetic centres.<sup>49</sup> Considering a pair of magnetic centres  $A$  and  $B$ , this is done by relaxing  $A$  to  $B$ , and independently  $B$  to  $A$ . It results in two sets of  $t$  and  $U$ , which are equal in highly symmetric situations.

Let us consider the extraction of the Hubbard Hamiltonian parameters to determine the kinetic exchange contribution to the  $J_{AB}$  coupling. Four relaxations may be considered (i) the occupied magnetic orbitals of  $A$  to the virtual magnetic orbitals  $B$  in the  $\bar{A}\bar{B}\dots N$  situation, (ii) the same relaxation in the  $\bar{A}\bar{B}\dots N$  situation, (iii) the occupied magnetic orbitals of  $B$  to the virtual magnetic orbitals of  $A$  in the  $\bar{A}\bar{B}\dots N$  situation and (iv) the same relaxation in the  $\bar{A}\bar{B}\dots N$  situation. As discussed in ref. 49, (i) and (ii) provide similar  $t$  and  $U$  parameters while the same is true for (iii) and (iv).

Let us now focus only on one of these extractions with the first relaxation to present the procedure in the context of several unpaired electrons per magnetic centre. This relaxation results in a reorganisation of the electronic distribution which may be thought of as slightly emptying the orbitals of  $A$  to fill the virtual orbitals of  $B$ ,

$$\begin{aligned}
\Phi_{\bar{A}'\bar{B}\bar{C}\dots N, \text{FC}} &= \left| \prod_i^{n_{\text{core}}} \bar{i} \bar{i} \prod_{\alpha}^{n_A} a'_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \\
&= C_0 \left| \prod_i^{n_{\text{core}}} \bar{i} \bar{i} \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| \\
&\quad + \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l} \hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \\
&\quad \times \left| \prod_i^{n_{\text{core}}} \bar{i} \bar{i} \prod_{\alpha}^{n_A} a_{\alpha} \prod_{\beta}^{n_B} \bar{b}_{\beta} \prod_{\gamma}^{n_C} c_{\gamma} \dots \prod_{\nu}^{n_N} n_{\nu} \right| + \mathcal{O}(2) \\
&= C_0 \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} + \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l} \hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} \\
&\quad + \mathcal{O}(2)
\end{aligned} \quad (29)$$

where  $C_0$  and  $C_{b_k a_l}$  correspond to the normalized coefficients with  $C_0^2 + \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l}^2 = 1$  and  $\mathcal{O}(2)$  to some higher-order

expansion determinants. The entry of these ionic-like forms in the wavefunction results in two sets of energetic contributions,

$$\begin{aligned}
& E[\Phi_{\bar{A}'\bar{B}\bar{C}\dots N, \text{FC}}] \\
&= C_0^2 E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] + \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l}^2 E[\hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] \\
&\quad + 2C_0 \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l} \langle \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} | \hat{H} | \hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} \rangle + \mathcal{O}(2) \\
&= \left( 1 - \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l}^2 \right) E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] \\
&\quad + \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l}^2 E[\hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] \\
&\quad + 2C_0 \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l} \langle \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} | \hat{H} | \hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} \rangle + \mathcal{O}(2) \\
&= E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] \\
&\quad + \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l}^2 (E[\hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) \\
&\quad + 2C_0 \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l} \langle \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} | \hat{H} | \hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} \rangle + \mathcal{O}(2)
\end{aligned} \quad (30)$$

The second term of the right-hand equation is homogeneous with the  $U$  on-site repulsion energy Hubbard parameter while the last is homogeneous with the hopping integral  $t$ . These terms correspond to some individual contributions from all  $A$  orbital- $B$  orbital pairs. To avoid a laborious extraction of all these terms, let us define some overall effective  $\tilde{t}$  and  $\tilde{U}$  parameters capturing the relevant physics of the relaxation mechanism of the effective spin centre  $A$  over  $B$  as,

$$\begin{aligned}
E[\Phi_{\bar{A}'\bar{B}\bar{C}\dots N, \text{FC}}] &= E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] + (1 - C_0^2) \tilde{U} \\
&\quad + 2C_0 \sqrt{1 - C_0^2} \tilde{t}
\end{aligned} \quad (31)$$

with,

$$\begin{aligned}
(1 - C_0^2) \tilde{U} &= \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l}^2 (E[\hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}] \\
&\quad - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}])
\end{aligned} \quad (32)$$

$$\sqrt{1 - C_0^2} \tilde{t} = \sum_k^{n_A} \sum_l^{n_B} C_{b_k a_l} \langle \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} | \hat{H} | \hat{a}_{b_k}^{\dagger} \hat{a}_{a_l} \Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}} \rangle \quad (33)$$

Hence, it results in a similar set of equations as for the one electron per centre case where both terms may be extracted by,

$$\tilde{U} = (E[\Phi_{\bar{A}'\bar{B}\bar{C}\dots N, \text{FC}}] - E[\Phi_{\bar{A}\bar{B}\bar{C}\dots N, \text{NO}}]) \frac{2\alpha - 1}{\alpha - 1} \quad (34)$$



$$\tilde{t} = (E[\Phi_{A'BC...N,FC}] - E[\Phi_{A\bar{B}C...N,NO}])\sqrt{\frac{\alpha}{1-\alpha}} \quad (35)$$

where,

$$\alpha = \frac{\langle \hat{S}^2 \rangle_{A'BC...N,FC} - \langle \hat{a}_{b_k}^\dagger \hat{a}_{a_l} \Phi_{A\bar{B}C...N,NO} | \hat{S}^2 | \hat{a}_{b_k}^\dagger \hat{a}_{a_l} \Phi_{A\bar{B}C...N,NO} \rangle}{\langle \hat{S}^2 \rangle_{A\bar{B}C...N,NO} - \langle \hat{a}_{b_k}^\dagger \hat{a}_{a_l} \Phi_{A\bar{B}C...N,NO} | \hat{S}^2 | \hat{a}_{b_k}^\dagger \hat{a}_{a_l} \Phi_{A\bar{B}C...N,NO} \rangle} \quad (36)$$

Pointing out that  $\langle \hat{a}_{b_k}^\dagger \hat{a}_{a_l} \Phi_{A\bar{B}C...N,NO} | \hat{S}^2 | \hat{a}_{b_k}^\dagger \hat{a}_{a_l} \Phi_{A\bar{B}C...N,NO} \rangle = \langle \hat{S}^2 \rangle_{A\bar{B}C...N,NO} - 1$ ,  $\alpha$  may be written in a simpler form as,

$$\alpha = 1 - \left( \langle \hat{S}^2 \rangle_{A\bar{B}C...N,NO} - \langle \hat{S}^2 \rangle_{A'BC...N,FC} \right) \quad (37)$$

Finally, the kinetic exchange contribution can be obtained by summing the Hubbard parameters resulting from both relaxations as,

$$8S_A S_B \Delta J_{KE} = \frac{\tilde{t}_{A \rightarrow B}^2}{\tilde{U}_{A \rightarrow B}} + \frac{\tilde{t}_{A \leftarrow B}^2}{\tilde{U}_{A \leftarrow B}} \quad (38)$$

## 2 Molecular systems

To investigate both roles of the bridging ligand and of the metal centre in M–radical–M complexes, three series of compounds have been studied. The first series consists of 4 complexes presented in Fig. 1(a),  $[(\text{TPyA})_2\text{Cr}_2^{\text{III}}(\text{NMePh}_2\text{L}^{3-\bullet})]^{3+}$  and  $[(\text{TPyA})_2\text{M}_2^{\text{II}}(\text{NMePh}_2\text{L}^{3-\bullet})]^{2+}$  (with M = Mn<sup>II</sup>, Fe<sup>II</sup> and Co<sup>II</sup>), where  $\text{NMePh}_2\text{LH}_2 = N,N,N,N$ -tetra(2-methylphenyl)-2,5-diamino-1,4-diiminobenzoquinone; TPyA = tris(2-pyridyl-methyl)amine, labelled **1a**, **2a**, **3a** and **4a**, respectively.<sup>50</sup> The couplings between the

**Table 1** Formula, electronic configuration of the M centre and experimental magnetic exchange coupling in cm<sup>−1</sup> of the M–radical–M complexes

Complex	M	Configuration	$J_{\text{exp}}^{\text{M-Rad}}$
<b>1a</b>	$[(\text{TPyA})_2\text{Cr}_2^{\text{III}}(\text{NMePh}_2\text{L}^{3-\bullet})]^{3+}$	Cr <sup>III</sup> d <sup>3</sup>	−626
<b>2a</b>	$[(\text{TPyA})_2\text{Mn}^{\text{II}}(\text{NMePh}_2\text{L}^{3-\bullet})]^{2+}$	Mn <sup>II</sup> d <sup>5</sup>	−157
<b>3a</b>	$[(\text{TPyA})_2\text{Fe}^{\text{II}}(\text{NMePh}_2\text{L}^{3-\bullet})]^{2+}$	Fe <sup>II</sup> d <sup>6</sup>	−307
<b>4a</b>	$[(\text{TPyA})_2\text{Co}^{\text{II}}(\text{NMePh}_2\text{L}^{3-\bullet})]^{2+}$	Co <sup>II</sup> d <sup>7</sup>	−396
<b>3b</b>	$[(\text{Me}_6\text{tren})_2\text{Fe}_2^{\text{II}}(\text{C}_6\text{H}_4\text{O}_2^{2-\bullet})]^{3+}$	Fe <sup>II</sup> d <sup>6</sup>	−144
<b>4b</b>	$[(\text{Me}_6\text{tren})_2\text{Co}_2^{\text{II}}(\text{C}_6\text{H}_4\text{O}_2^{2-\bullet})]^{3+}$	Co <sup>II</sup> d <sup>7</sup>	−252
<b>5b</b>	$[(\text{Me}_6\text{tren})_2\text{Ni}_2^{\text{II}}(\text{C}_6\text{H}_4\text{O}_2^{2-\bullet})]^{3+}$	Ni <sup>II</sup> d <sup>8</sup>	< −600
<b>1c<sup>O</sup></b>	$[(\text{TPyA})_2\text{Cr}_2^{\text{III}}(\text{O}^{\text{L}^{3-\bullet}})]^{3+}$	Cr <sup>III</sup> d <sup>3</sup>	−352
<b>1c<sup>OS</sup></b>	$[(\text{TPyA})_2\text{Cr}_2^{\text{III}}(\text{OS}^{\text{L}^{3-\bullet}})]^{3+}$	Cr <sup>III</sup> d <sup>3</sup>	−401
<b>1c<sup>S</sup></b>	$[(\text{TPyA})_2\text{Cr}_2^{\text{III}}(\text{S}^{\text{L}^{3-\bullet}})]^{3+}$	Cr <sup>III</sup> d <sup>3</sup>	−487

radical moiety and the metal centres have been experimentally determined by fitting the  $\chi T$  versus  $T$  magnetic susceptibility curve using the following HDvV model Hamiltonian,

$$\hat{H} = -2J\hat{S}_{\text{rad}} \cdot (\hat{S}_{\text{M}_1} + \hat{S}_{\text{M}_2}) \quad (39)$$

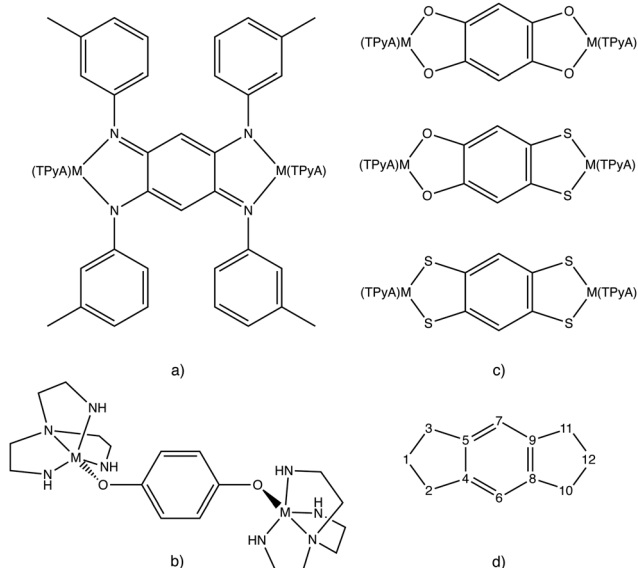
These complexes present large antiferromagnetic couplings ranging from −157 cm<sup>−1</sup> for **2a** to −626 cm<sup>−1</sup> for **1a** and are detailed in Table 1. This series of tetraazalene radical-based compounds has been previously studied by Singh using DFT calculations.<sup>58</sup>

The second series (Fig. 1(b)) is based on semiquinone radical with a  $[(\text{Me}_6\text{tren})_2\text{M}_2^{\text{II}}(\text{C}_6\text{H}_4\text{O}_2^{2-\bullet})]^{3+}$  architecture with Me<sub>6</sub>tren = tris(2-dimethylaminoethyl)amine and M = Fe, Co, Ni, labelled **3b**, **4b** and **5b**, respectively.<sup>51</sup> Here again, radical–metal couplings have been determined by fitting the same model Hamiltonian over  $\chi T$  versus  $T$  curve, resulting in strong antiferromagnetic couplings presented in Table 1.

Finally, a third series of three complexes consisting in Cr<sup>III</sup> ions linked through a bis(bidentate) benzosemiquinoid-based radical,  $[(\text{TPyA})_2\text{Cr}_2^{\text{III}}(\text{L}^{3-\bullet})]^{3+}$ , is presented in Fig. 1(c).<sup>52</sup> It results in two symmetrical 1,2,4,5-tetrahydroxybenzene **1c<sup>O</sup>** and 1,2,4,5-tetrathiobenzene **1c<sup>S</sup>** radicals and an asymmetrical intermediate 1,2-dithio-4,5-dihydroxybenzene **1c<sup>OS</sup>**. Variable-temperature dc magnetic susceptibility data reveal the presence of relatively strong antiferromagnetic couplings between metal ions and the radical moieties at −352, −401 and −487 cm<sup>−1</sup> for **1c<sup>O</sup>**, **1c<sup>OS</sup>** and **1c<sup>S</sup>**, respectively.

## 3 Computational details

For the first series (**1a–4a**), molecular structures from ref. 58 have been used and correspond to the optimised geometries at the B3LYP<sup>59–62</sup>/TZV<sup>63,64</sup> level of theory. The molecular structures of the second (**3b–5b**) and third (**1c<sup>O,OS,S</sup>**) series have been obtained from the Cambridge Structural Database (CSD),<sup>65</sup> of which the position of hydrogen atoms have been optimised at the B3LYP/def2-SVP level of theory with ORCA 4.2.0.<sup>66</sup> All decompositions of magnetic exchange couplings have been performed using the B3LYP functional with the def2-TZVP basis set for all atoms except the metal centres where the def2-QZVPP basis sets have been used.<sup>67</sup> Due to the use of the RIJCOSX



**Fig. 1** Schematic representations of (a) the tetraazulene-based series M = Cr<sup>III</sup>, Mn<sup>II</sup>, Fe<sup>II</sup> and Co<sup>II</sup>, (b) the semiquinone-based series with M = Fe<sup>II</sup>, Co<sup>II</sup> and Ni<sup>II</sup>, (c) the benzosemiquinoid-based series with M = Cr<sup>III</sup>, and (d) atom numbering of the benzosemiquinoid-based series of Table 8.





approximation, quasi-restricted open-shell formalism has been used to compute HS states.<sup>56</sup> The selective relaxation of the orbitals is permitted by the LSCF method,<sup>68</sup> implemented in the ORCA package since the version 4.2. All figures of orbitals have been done using Jmol.<sup>69</sup>

## 4 Results and discussion

Table 2 presents the decomposition into the three contributions ( $J_0$ ,  $\Delta J_{KE}$  and  $\Delta J_{SP}$ ) as well as their sum ( $J_\Sigma$ ) for the couplings of all complexes. Considering first the total magnetic exchange couplings between the metal centre and the radical bridge,  $J_\Sigma$  properly reproduces the antiferromagnetic nature and the order of magnitude of the experimental determination for the first series of compounds, with  $J_\Sigma$  at  $-665$ ,  $-132$ ,  $-176$  and  $-257$  cm<sup>-1</sup> for **1a**, **2a**, **3a** and **4a**, respectively, compared to the experimental values at  $-626$ ,  $-157$ ,  $-307$  and  $-396$  cm<sup>-1</sup>. For the second series, the same situation is observed with theoretical evaluations at  $-47$ ,  $-278$  and  $\sim -445$  cm<sup>-1</sup> for **3b**, **4b** and **5b**, respectively. For the third series, the sign and the magnitude of couplings in **1c<sup>O</sup>** and **1c<sup>S</sup>** are correctly

reproduced with  $-280$  and  $-370$  cm<sup>-1</sup> instead of  $-401$  and  $-487$  experimentally, respectively. As mentioned in Section 2, **1c<sup>OS</sup>** presents an asymmetric radical ligand and it then should result in two different couplings, considering the coupling between the radical moiety and the Cr ion closer to the oxygen atoms (labelled  $M_1(O)$ ) or closer to the sulfur atoms ( $M_2(S)$ ). Experimentally, the model Hamiltonian used considered the same coupling for both interactions (eqn (39)) and results in  $J$  at  $-401$  cm<sup>-1</sup>. DFT calculations depict a completely different story and both couplings turn out to be very different at  $-29$  and  $-629$  cm<sup>-1</sup> for the  $J^{M_1(O)-Rad}$  and  $J^{M_2(S)-Rad}$ , respectively. It may be worth noting that averaging both theoretical evaluations result in a value of  $-329$  cm<sup>-1</sup> which is between both  $J_\Sigma$  of **1c<sup>O</sup>** and **1c<sup>S</sup>**, as the experimental determination. Finally, these calculations confirm the expected negligible coupling between both metal ions for the three series of compounds.

Focusing now on the decomposition, in all series, the spin polarisation is negligible regarding the magnitude of both  $J_0$  and  $\Delta J_{KE}$  contributions and the overall coupling and, for instance, the largest value of  $\Delta J_{SP}$  is obtained for **1a** with  $-24$  cm<sup>-1</sup> for a total coupling of  $-665$  cm<sup>-1</sup>. Whilst the spin polarisation contribution may be essential in the energy splitting of polyradicals, it is negligible in transition metal-based compounds because of the very local nature of magnetic orbitals. Hence, it is worth noting that in the present type of coupling between both a radical centre and a metal ion, the spin polarisation is negligible, regardless of the radical ligand or the metal considered.

Hence, the nature of the metal–radical couplings results from the competition between the ferromagnetic direct exchange and the antiferromagnetic kinetic exchange. The latter is always larger and yields antiferromagnetic couplings for all complexes. First, let us now focus on a detailed discussion of both contributions and the physics governing them in the context of varying the metal centres with the series **1–4a** and **3–5b**. We will follow the discussion by investigating the change in the bridging radical ligands in the third series.

### 4.1 Impact of the metal centre

**4.1.1 Direct exchange contribution.** The first series of compounds presents three similar direct exchange contributions with  $119$  cm<sup>-1</sup> for **3a** and the two very close  $144$  and  $140$  cm<sup>-1</sup> for **1a** and **4a**, respectively, as well as a lower contribution at  $67$  cm<sup>-1</sup> for **2a**. To rationalise these values, the usual orbital analysis may be corroborated with a decomposition of the direct exchange into its electron–electron interaction, presented in Table 3 and whose methodology has been introduced in ref. 46. This method is based on treating a coupling between two centres *A* and *B* with  $n_A$  and  $n_B$  unpaired electrons, respectively, as a system of  $n_A + n_B$  centres with one unpaired electron to determine all electron–electron contributions. It is worth noting that in Table 3, all sums of the  $J_0^{d-rad}$  are equal to  $4S_M S_{rad} J_0$  and then both extractions are consistent.

Starting with **1a**, Cr<sup>III</sup> ion features three unpaired electrons occupying  $t_{2g}$  orbitals due to a near octahedral arrangement. These orbitals as well as the  $\pi^*$  orbital of the radical moiety are

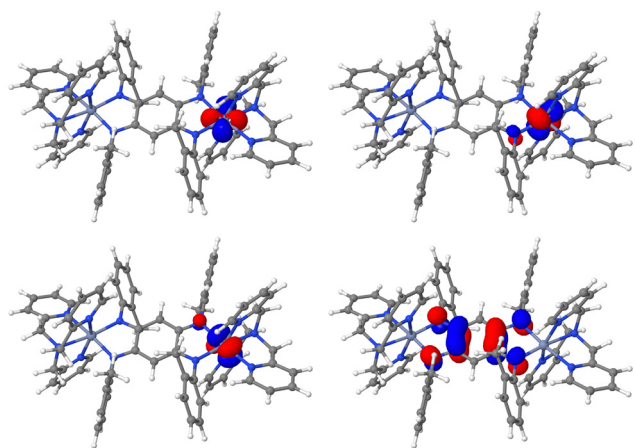
Table 2 Decomposition ( $J_0$ ,  $\Delta J_{KE}$  and  $\Delta J_{SP}$ ), recomposition ( $J_\Sigma$ ) and experimental value ( $J_{exp}$ ) for all compounds in cm<sup>-1</sup>

		$J_0$	$\Delta J_{KE}$	$\Delta J_{SP}$	$J_\Sigma$	$J_{exp}$
<b>1a</b>	$J^{M_1-Rad}$	144	-786	-24	-665	-626
	$J^{M_2-Rad}$	144	-786	-24	-665	
	$J^{M_1-M_2}$	6	-15	0	-9	
<b>2a</b>	$J^{M_1-Rad}$	67	-202	3	-132	-157
	$J^{M_2-Rad}$	67	-202	3	-132	
	$J^{M_1-M_2}$	1	-1	0	-1	
<b>3a</b>	$J^{M_1-Rad}$	119	-312	17	-176	-307
	$J^{M_2-Rad}$	119	-313	17	-176	
	$J^{M_1-M_2}$	2	-3	1	0	
<b>4a</b>	$J^{M_1-Rad}$	140	-411	14	-257	-396
	$J^{M_2-Rad}$	140	-411	14	-257	
	$J^{M_1-M_2}$	3	-1	1	3	
<b>3b</b>	$J^{M_1-Rad}$	298	-366	21	-47	-144
	$J^{M_2-Rad}$	298	-366	21	-46	
	$J^{M_1-M_2}$	2	0	0	2	
<b>4b</b>	$J^{M_1-Rad}$	109	-391	3	-278	-252
	$J^{M_2-Rad}$	109	-391	3	-278	
	$J^{M_1-M_2}$	0	0	0	0	
<b>5b</b>	$J^{M_1-Rad}$	111	-559	1	-447	< -600
	$J^{M_2-Rad}$	111	-554	1	-442	
	$J^{M_1-M_2}$	1	-1	0	0	
<b>1c<sup>O</sup></b>	$J^{M_1-Rad}$	132	-420	8	-280	-352
	$J^{M_2-Rad}$	132	-420	8	-280	
	$J^{M_1-M_2}$	2	-6	0	-4	
<b>1c<sup>S</sup></b>	$J^{M_1-Rad}$	107	-461	-16	-370	-487
	$J^{M_2-Rad}$	107	-461	-16	-370	
	$J^{M_1-M_2}$	3	-8	-1	-6	
<b>1c<sup>OS</sup></b>	$J^{M_1(O)-Rad}$	71	-101	1	-29	-401
	$J^{M_2(S)-Rad}$	135	-741	-23	-629	
	$J^{M_1-M_2}$	2	-4	0	-2	



**Table 3** Individual radical-d orbital contributions ( $J_0^{\text{d-rad}}$ ), the sum of the individual contributions ( $4S_{\text{M}}S_{\text{rad}}J_0$ ) and the total direct exchange ( $J_0$ ) for the first and second series of compounds (**1–4a** and **3–5b**) in  $\text{cm}^{-1}$

		$J_0^{\text{d-rad}}$	$4S_{\text{M}}S_{\text{rad}}J_0$	$J_0$
<b>1a</b>	$d_{xy}$	18	432	144
	$d_{xz}$	201		
	$d_{yz}$	213		
<b>2a</b>	$d_{xy}$	12	337	67
	$d_{z^2}$	36		
	$d_{x^2-y^2}$	67		
	$d_{yz}$	111		
	$d_{xz}$	111		
<b>3a</b>	$d_{z^2}$	72	476	119
	$d_{x^2-y^2}$	82		
	$d_{yz}$	157		
	$d_{xz}$	165		
<b>4a</b>	$d_{z^2}$	73	419	140
	$d_{x^2-y^2}$	128		
	$d_{yz}$	219		
<b>3b</b>	$d_{yz}$	249	1186	296
	$d_{z^2}$	274		
	$d_{x^2-y^2}$	324		
	$d_{xy}$	339		
<b>4b</b>	$d_{yz}$	69	324	109
	$d_{z^2}$	108		
	$d_{x^2-y^2}$	146		
<b>5b</b>	$d_{z^2}$	64	222	111
	$d_{x^2-y^2}$	158		



**Fig. 2** Isosurface of the  $d_{xy}$  (top left),  $d_{yz}$  (top right),  $d_{yz}$  (bottom left) and  $\pi^*$  radical (bottom right) magnetic orbitals of **1a**. Blue grey = chromium, blue = nitrogen, grey = carbon, white = hydrogen. Isovalue = 0.5 a.u.

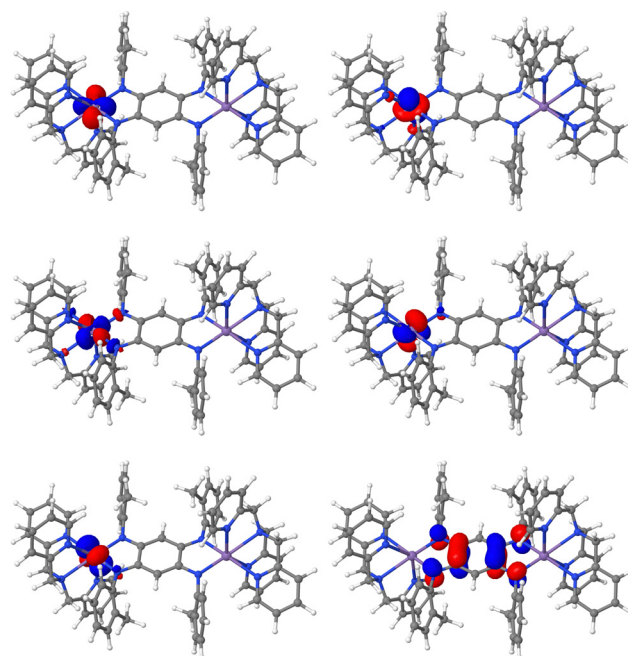
presented in Fig. 2. Both  $d_{xz}$  and  $d_{yz}$  orbitals have the lobes pointing towards the ligands and would have a strong interaction with the radical centre due to its  $\pi$  arrangement. It results in two similar direct exchange contributions at 201 and 213  $\text{cm}^{-1}$ . As the opposite, the  $d_{xy}$  orbital is in the plane of the radical ligand and its lobes do not point towards the ligand, yielding a much smaller contribution at 18  $\text{cm}^{-1}$ . It may be worth noting that some authors arrived at a slightly different

interpretation based on the computation of the overlap between the metal orbitals and the radical moiety from BS unrestricted calculations.<sup>58</sup> However, the physical content of the magnetic orbitals may differ between these two works, resulting in different conclusions. The second compound **2a** presents an interesting example for the orbital analysis since all the d orbitals are occupied in the  $\text{Mn}^{\text{II}}$  ion, presented in Fig. 3. Like **1a**, the strongest exchange interactions are provided with the  $d_{xz}$  and  $d_{yz}$ , both at 111  $\text{cm}^{-1}$ , while the weakest results from the interaction with the radical centre and the  $d_{xy}$  orbital at 12  $\text{cm}^{-1}$ . The two  $e_g$  orbitals provide intermediate contributions. The  $d_{x^2-y^2}$ -rad interaction provides a slightly smaller contribution than the  $d_{xz}$  and  $d_{yz}$  at 67  $\text{cm}^{-1}$  since, while the lobes also point towards the ligands, they are in a  $\sigma$  arrangement. Finally, the  $d_{z^2}$  interaction is relatively small and is between  $d_{x^2-y^2}$  and  $d_{xy}$  ones at 36  $\text{cm}^{-1}$ .

Since the orbital analysis of compounds **3a** and **4a** are similar, they would not be detailed in the discussion but the orbitals of both complexes may be found in Fig. S2 and S3 (ESI†).

Let us turn now to the second series of compounds featuring a semiquinone radical ligand. In this series, **3b** presents a much stronger direct exchange interaction at 296  $\text{cm}^{-1}$  than **4b** and **5b** at 109 and 111  $\text{cm}^{-1}$ , respectively. To explain this situation, a first observation may be done by comparing the orbital of the radical centre of the three compounds presented Fig. 4. Hence, the **3b** radical orbital shows a significantly greater delocalisation tail over both magnetic centres, from which may result a stronger interaction between the radical centre and metal electrons.

For these three compounds, a similar orbital analysis may be realised whilst the numerical contributions of every



**Fig. 3** Isosurface of the  $d_{xy}$  (top left),  $d_{z^2}$  (top right),  $d_{x^2-y^2}$  (middle left),  $d_{yz}$  (middle right),  $d_{xz}$  (bottom left) and  $\pi^*$  radical (bottom right) magnetic orbitals of **2a**. Purple = manganese, blue = nitrogen, grey = carbon, white = hydrogen. Isovalue = 0.5 a.u.



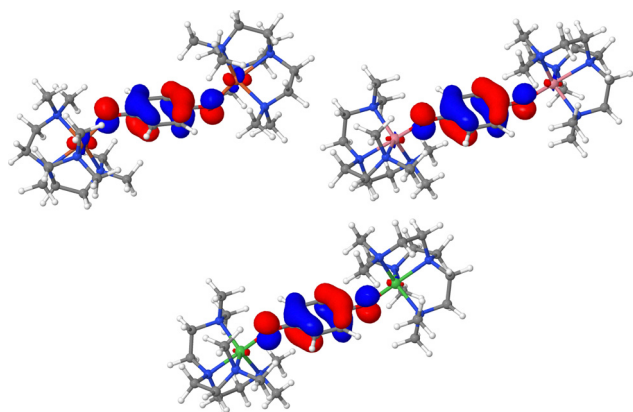


Fig. 4 Isosurface of the radical  $\pi^*$  orbitals of **3b** (top left), **4b** (top right) and **5b** (bottom). Orange = iron, pink = cobalt, green = nickel, blue = nitrogen, red = oxygen, grey = carbon, white = hydrogen. Isovalue = 0.5 a.u.

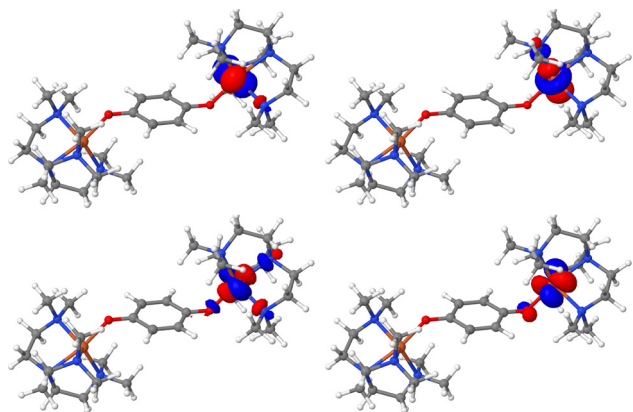


Fig. 5 Isosurface of the  $d_{yz}$  (top left),  $d_{z^2}$  (top right),  $d_{x^2-y^2}$  (bottom left) and  $d_{xy}$  (bottom right) magnetic orbitals of **3b**. Orange = iron, blue = nitrogen, red = oxygen, grey = carbon, white = hydrogen. Isovalue = 0.5 a.u.

electron–electron interaction is presented in Table 3. For the sake of conciseness, let us focus on **3b** since the conclusions are similar for **4b** and **5b**. In **3b**, two  $t_{2g}$  and two  $e_g$  orbitals are singly populated due to the trigonal bipyramidal arrangement of the  $\text{Fe}^{\text{II}}$  ions. The two stronger exchange contributions are provided with the  $d_{xy}$  and  $d_{x^2-y^2}$  orbitals due to the orientation of their lobes towards the ligand and a delocalisation tail over

the oxygen atom of the bridging ligand (Fig. 5), resulting in contributions of 339 and 324  $\text{cm}^{-1}$ , respectively. Hence, with their orientation perpendicular to the metal–oxygen bond, the  $d_{z^2}$  and  $d_{yz}$  lead to two weaker contributions at 274 and 249  $\text{cm}^{-1}$ , respectively.

**4.1.2 Kinetic exchange.** The kinetic exchange is the dominant contribution in both series of compounds, resulting in the antiferromagnetic nature of the metal–radical coupling. This coupling takes place between two magnetic centres of a very different nature: a radical centre bearing one unpaired electron in a rather delocalised  $\pi^*$  orbital, and a metal ion with several unpaired electrons occupying d orbitals with a very local character. The kinetic exchange of a coupling is computed by considering two relaxations, the metal-to-radical and radical-to-metal, which thus correspond to two very different situations. Table 4 presents for both series the effective  $\tilde{t}$  and  $\tilde{U}$  parameters and the associated energetic contribution related to both metal-to-radical and radical-to-metal relaxations.

Starting with the first series, the radical-to-metal relaxation provides the strongest contribution to the total kinetic exchange with, for instance,  $-606 \text{ cm}^{-1}$  for metal  $\leftarrow$  rad and  $-179 \text{ cm}^{-1}$  for metal  $\rightarrow$  rad in the case of **1a**. For the four complexes, this difference is explained by both, a change in the  $t$  and  $U$  values. For the former, the metal  $\leftarrow$  rad  $t$  value is always stronger and this reflects the fact that it is easier for the electron to jump from the radical moiety to the metal ion than the opposite. For the  $U$  value, this parameter is for all compounds of the series smaller for metal  $\leftarrow$  rad relaxation, and the on-site repulsion energy is then smaller on the metal ion. Hence, both parameters favour the metal  $\leftarrow$  rad relaxation and explain the large discrepancy between both situations.

Regarding the energetic contribution, the second series shows different trends. Qualitatively, whilst **5b** follows the trend of the first series with a larger metal  $\leftarrow$  rad relaxation, it is the opposite for **3b** and both contributions are equal in **4b**. Quantitatively, the metal  $\leftarrow$  rad and metal  $\rightarrow$  rad relaxations provide contributions of the same order of magnitude, around a few hundred of  $\text{cm}^{-1}$ . Going further in the Hubbard parameters, for **3b** the  $t$  values from both relaxations are similar whilst both  $U$  are very different, explaining the difference between the contribution from metal  $\rightarrow$  rad ( $-220 \text{ cm}^{-1}$ ) and metal  $\leftarrow$  rad ( $-146 \text{ cm}^{-1}$ ). For **5b** the situation is the opposite since both  $U$  are similar and the different  $t$  values ( $-3.5 \times 10^{-3}$  and  $-4.4 \times 10^{-3} \text{ cm}^{-1}$ ) explain the discrepancy between the contribution from both relaxations. It is interesting to point out that both **4b** and **5b**  $U$

Table 4 Hubbard Hamiltonian  $\tilde{t}$  and  $\tilde{U}$  parameters and their energetic contribution ( $-(8S_A S_B)^{-1} \tilde{t}^2 / \tilde{U}$ ) for both metal  $\rightarrow$  rad and metal  $\leftarrow$  rad relaxations and the resulting total kinetic exchange ( $\Delta J_{\text{KE}}$ ) for all compounds in  $\text{cm}^{-1}$

	Metal $\rightarrow$ rad			Metal $\leftarrow$ rad			$\Delta J_{\text{KE}}$
	$\tilde{t} \times 10^{-3}$	$\tilde{U} \times 10^{-3}$	$-(8S_A S_B)^{-1} \tilde{t}^2 / \tilde{U}$	$\tilde{t} \times 10^{-3}$	$\tilde{U} \times 10^{-3}$	$-(8S_A S_B)^{-1} \tilde{t}^2 / \tilde{U}$	
<b>1a</b>	−4.7	40.5	−179	−6.3	21.9	−606	−786
<b>2a</b>	−3.3	42.2	−51	−5.2	34.3	−152	−202
<b>3a</b>	−3.2	48.4	−53	−4.7	21.6	−259	−312
<b>4a</b>	−2.6	53.6	−40	−4.1	15.0	−371	−411
<b>3b</b>	−4.7	23.5	−220	−5.0	42.4	−146	−366
<b>4b</b>	−4.1	27.6	−197	−4.7	38.5	−194	−391
<b>5b</b>	−3.7	28.8	−241	−4.4	30.7	−318	−559





**Table 5** Hubbard Hamiltonian  $t$  and  $U$  parameters resulting from the radical-d orbital interaction, their energetic contribution ( $t^2/U$ ), the sum of both  $d \rightarrow \text{rad}$  and  $d \leftarrow \text{rad}$  ( $\sum t^2/U$ ) and the total energetic contribution to the coupling  $(-8S_A S_B)^{-1} \sum t^2/U$  for the second series of compounds (**3–5b**) in  $\text{cm}^{-1}$

		$t \times 10^{-3}$	$U \times 10^{-3}$	$t^2/U$	$\sum t^2/U$	$(-8S_A S_B)^{-1} \sum t^2/U$
<b>3b</b>	$d_{yz} \rightarrow \text{rad}$	—	—	—	—	—
	$d_{yz} \leftarrow \text{rad}$	−0.2	37.8	−1	−1	—
	$d_{xy} \rightarrow \text{rad}$	−0.7	40.0	−12	−12	—
	$d_{xy} \leftarrow \text{rad}$	−0.8	31.9	−19	−31	−367
	$d_{z^2} \rightarrow \text{rad}$	−2.5	28.5	−214	−471	—
	$d_{z^2} \leftarrow \text{rad}$	−3.0	34.5	−257	—	—
	$d_{x^2-y^2} \rightarrow \text{rad}$	−4.2	29.7	−582	−968	—
<b>4a</b>	$d_{x^2-y^2} \leftarrow \text{rad}$	−3.9	39.5	−386	—	—
	$d_{yz} \rightarrow \text{rad}$	—	—	—	—	—
	$d_{yz} \leftarrow \text{rad}$	—	—	—	—	—
	$d_{z^2} \rightarrow \text{rad}$	−2.2	43.5	−111	−284	−396
	$d_{z^2} \leftarrow \text{rad}$	−2.4	33.4	−173	—	—
	$d_{x^2-y^2} \leftarrow \text{rad}$	−3.6	25.8	−495	−903	—
	$d_{x^2-y^2} \rightarrow \text{rad}$	−4.4	48.0	−408	—	—
<b>5b</b>	$d_{z^2} \rightarrow \text{rad}$	−1.6	46.6	−52	−171	—
	$d_{z^2} \leftarrow \text{rad}$	−2.0	33.1	−119	—	—
	$d_{x^2-y^2} \rightarrow \text{rad}$	−3.5	27.7	−452	−558	—
	$d_{x^2-y^2} \leftarrow \text{rad}$	−4.5	40.4	−493	−946	—

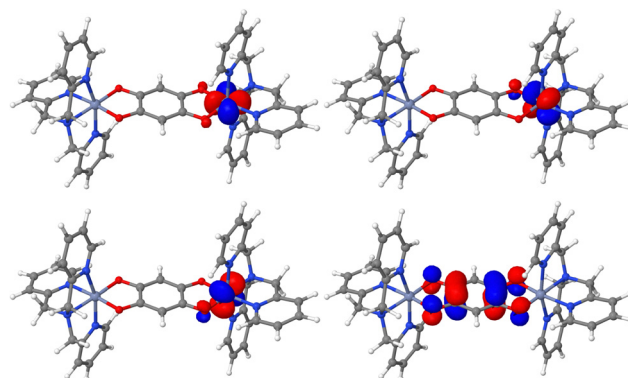
values of the metal  $\rightarrow \text{rad}$  relaxation are very similar whilst the **3b** one is slightly smaller. This may be readily related to the shape of the SOMO localised on the radical ligand in Fig. 4. Indeed, as mentioned, the  $\pi^*$  orbital of **3b** is more diffused than for the other compounds and would result in weaker on-site repulsion energy.

To end this discussion, the electron–electron decomposition may be used in the context of the kinetic exchange contribution to extract the individual  $t$  and  $U$  parameters, of which the results are presented for the second series of compounds in Table 5. Let us focus on **3b** since it represents a good example among all studied compounds. Unlike the direct exchange, the strongest contribution to  $\Delta J_{\text{KE}}$  is provided by the interaction between the radical centre and the  $d_{x^2-y^2}$  at  $-968 \text{ cm}^{-1}$ , followed by the interaction with the  $d_{z^2}$  orbital at  $-471 \text{ cm}^{-1}$ . The interaction with the  $d_{xy}$  orbital, which is the strongest for the direct exchange, is eventually the third interaction with an order of magnitude lower at  $-31 \text{ cm}^{-1}$ . This may readily be explained with the plot of the orbital in Fig. 4 and 5 for the radical ligand and the metal orbitals, respectively. Here, the lobes of the  $d_{xy}$  orbital are in a perpendicular  $\pi$  arrangement to the  $\pi^*$  of the radical centre. This situation results in an unfavourable pathway for the electron to jump from one site to the other, and this is numerically confirmed with some negligible  $t$  values below  $-10^{-3} \text{ cm}^{-1}$ . Finally, it may be worth noting that summing all electron–electron contributions provides an evaluation of  $\Delta J_{\text{KE}}$  in very good agreement with those from the effective  $\tilde{t}$  and  $\tilde{U}$  and confirms the relevance of the presented analysis.

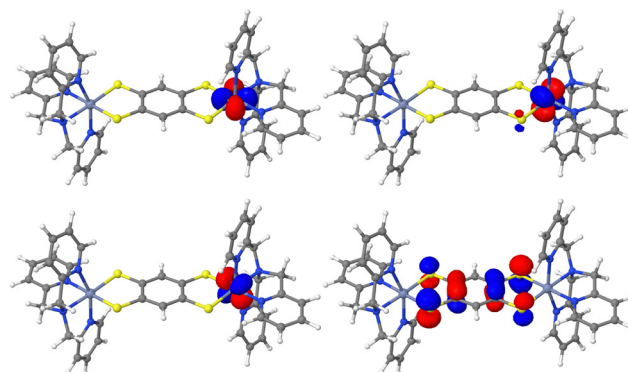
#### 4.2 Modification of the radical-bridging ligands in the $1c^R$ series

The third series of compounds with  $1c^{\text{OS,OS}}$  corresponds to an interesting situation with small variations of the benzosemiquinoid-based radical ligand bridging both  $\text{Cr}^{\text{III}}$  ions.

Let us start with the main decomposition presented in Table 2 and a comparison between the computed  $J_{\Sigma}$  couplings of  $1c^{\text{O}}$  at  $-280 \text{ cm}^{-1}$  and  $1c^{\text{S}}$  at  $-370 \text{ cm}^{-1}$ . The coupling in both compounds result from a strong direct exchange and an even larger kinetic exchange, yielding the antiferromagnetic character. However, the larger antiferromagnetic coupling in  $1c^{\text{S}}$  results at the same time from a weaker ferromagnetic direct exchange ( $107$  instead of  $132 \text{ cm}^{-1}$  in  $1c^{\text{O}}$ ), and a stronger antiferromagnetic kinetic exchange contribution ( $-461$  instead of  $-420 \text{ cm}^{-1}$ ). As already mentioned, whilst an averaging coupling has experimentally been determined for  $1c^{\text{OS}}$ , theoretically two very distinct couplings are found at  $-29 \text{ cm}^{-1}$  for the coupling on the side of the oxygen atoms and  $-629 \text{ cm}^{-1}$  on the side of the sulfur ones, labelled  $\text{Cr}(\text{O})\text{Rad}$  and  $\text{Cr}(\text{S})\text{Rad}$  hereafter, respectively. This corresponds to a strong quenching of the coupling compared to  $1c^{\text{O}}$  and conversely, a strong enhancement regarding  $1c^{\text{S}}$ . Despite this very large discrepancy, the direct exchange contributions show more moderate variations with values at  $71$  and  $135 \text{ cm}^{-1}$  for  $\text{Cr}(\text{O})\text{Rad}$  and  $\text{Cr}(\text{S})\text{Rad}$ , respectively. The kinetic exchange is then mainly responsible of this



**Fig. 6** Isosurface of the  $d_{xy}$  (top left),  $d_{xz}$  (top right),  $d_{yz}$  (bottom left) and radical (bottom right) magnetic orbitals of  $1c^{\text{O}}$ . Blue grey = chromium, blue = nitrogen, red = oxygen, grey = carbon, white = hydrogen. Isovalue =  $0.5 \text{ a.u.}$



**Fig. 7** Isosurface of the  $d_{xy}$  (top left),  $d_{yz}$  (top right),  $d_{xz}$  (bottom left) and radical (bottom right) magnetic orbitals of  $1c^{\text{S}}$ . Blue grey = chromium, blue = nitrogen, yellow = sulfur, grey = carbon, white = hydrogen. Isovalue =  $0.5 \text{ a.u.}$



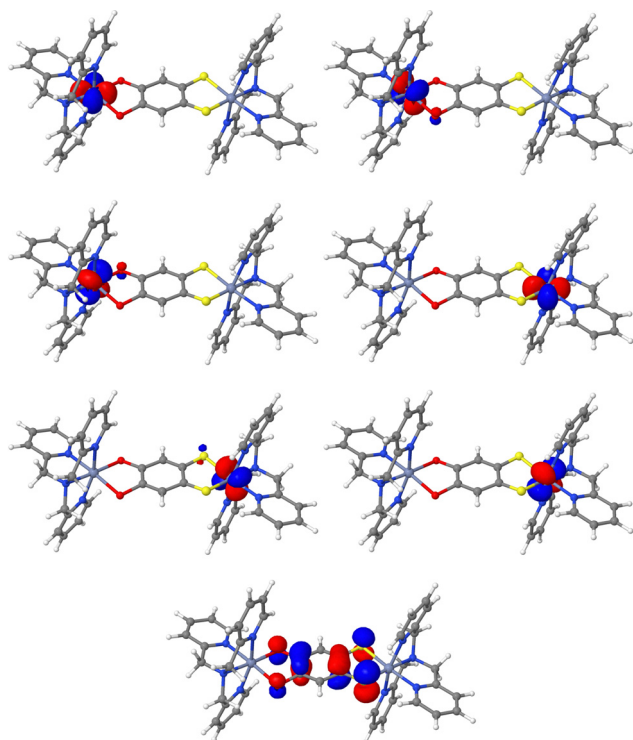


Fig. 8 Isosurface of the  $d_{xy}$  (top left),  $d_{xz}$  (top right) and  $d_{yz}$  (middle left) magnetic orbitals of  $M_1(O)$ , the  $d_{xy}$  (middle right),  $d_{xz}$  (bottom left) and  $d_{yz}$  (bottom right) magnetic orbitals of  $M_1(S)$ , and radical orbital (bottom) in  $1c^{OS}$ . Blue grey = chromium, blue = nitrogen, red = oxygen, yellow = sulfur, grey = carbon, white = hydrogen. Isovalue = 0.5 a.u.

Table 6 Individual radical-d orbital contributions ( $J_0^{d-rad}$ ), the sum of the individual contributions ( $4S_M S_{rad} J_0$ ) and the total direct exchange ( $J_0$ ) for the third series of compounds ( $1c^R$ ) in  $cm^{-1}$

		$J_0^{d-rad}$	$4S_M S_{rad} J_0$	$J_0$
$1c^O$	$d_{xy}$	33		
	$d_{xz}$	178	396	132
	$d_{yz}$	185		
$1c^S$	$d_{xy}$	31		
	$d_{yz}$	137	323	108
	$d_{xz}$	155		
$1c^{OS}$	$d_{xy}(O)$	13		
	$d_{xz}(O)$	92	215	72
	$d_{yz}(O)$	110		
	$d_{xy}(S)$	51		
	$d_{xz}(S)$	175	405	135
	$d_{yz}(S)$	179		

Table 7 Hubbard Hamiltonian  $\tilde{t}$  and  $\tilde{U}$  parameters and their energetic contribution ( $-(8S_A S_B)^{-1} \tilde{t}^2 / \tilde{U}$ ) for both metal  $\rightarrow$  rad and metal  $\leftarrow$  rad relaxations and the resulting total kinetic exchange ( $\Delta J_{KE}$ ) for all compounds in  $cm^{-1}$

	Metal $\rightarrow$ rad			Metal $\leftarrow$ rad			$\Delta J_{KE}$
	$\tilde{t} \times 10^{-3}$	$\tilde{U} \times 10^{-3}$	$-(8S_A S_B)^{-1} \tilde{t}^2 / \tilde{U}$	$\tilde{t} \times 10^{-3}$	$\tilde{U} \times 10^{-3}$	$-(8S_A S_B)^{-1} \tilde{t}^2 / \tilde{U}$	
$1c^O$	−3.9	29.0	−174	−4.8	30.8	−246	−420
$1c^S$	−3.9	27.8	−178	−5.4	34.9	−283	−461
$1c^{OS}(O)$	−1.8	39.2	−29	−2.4	26.5	−73	−101
$1c^{OS}(S)$	−4.8	31.5	−246	−6.4	27.3	−495	−741

strong variation in  $J_S$  with values at  $-101$  and  $-741$   $cm^{-1}$  for Cr(O)Rad and Cr(S)Rad, respectively.

**4.2.1 Direct exchange contribution.** As for **1a**, the octahedral coordination in these series of Cr<sup>III</sup>-based compounds results in occupying the  $t_{2g}$  orbitals, presented in Fig. 6, 7 and 8 for  $1c^O$ ,  $1c^S$  and  $1c^{OS}$ , respectively. Table 6 presents the decomposition of the direct exchange into electron–electron interactions, from which the orbital analysis may be done. Regarding the similitude with **1a**, only relevant aspects are detailed here. For all couplings, the interactions between the radical centre and the  $d_{xz}$  and  $d_{yz}$  largely dominate the overall  $J_0$  for the same reasons as for **1a**. Looking at these orbitals in Fig. 6 and 7, in  $1c^O$  they show more pronounced delocalisation tails over the oxygen atoms, explaining their larger interactions at about  $180$   $cm^{-1}$  than in  $1c^S$  at  $137$  and  $155$   $cm^{-1}$ . For  $1c^{OS}$ , the decomposition of  $J_0$  confirms a quenching of Cr(O)Rad compared to the  $1c^O$  coupling ( $72$  instead of  $132$   $cm^{-1}$ ) and an enhancement of Cr(S)Rad compared to the  $1c^S$  coupling ( $135$  instead of  $108$   $cm^{-1}$ ).

**4.2.2 Kinetic exchange contribution.** Let us turn now to the dominant kinetic exchange contribution, of which the detail of the metal-to-radical and radical-to-metal contributions and the associated  $\tilde{t}$  and  $\tilde{U}$  are presented in Table 7 for the three compounds. For the four involved couplings, the radical-to-metal contribution dominates the overall kinetic exchange and is approximately twice the radical-to-metal one.

Let us consider first the two symmetrical compounds  $1c^O$  and  $1c^S$ . The metal-to-radical relaxations provide similar energetic contributions at  $-174$  and  $-178$   $cm^{-1}$ , resulting from similar  $\tilde{t}$  and  $\tilde{U}$  parameters. Whilst the  $\tilde{t}$  cannot be rationalised from the shape of the orbitals only, both radical orbitals seem relatively similar between  $1c^O$  and  $1c^S$ , which could result in similar  $\tilde{U}$ . The radical-to-metal relaxation is slightly different between  $1c^O$  and  $1c^S$  and comes from different  $\tilde{t}$  and  $\tilde{U}$  values. Here, the  $t_{2g}$  orbitals of  $1c^O$  show more pronounced delocalisation tails over the chalcogen atoms than in  $1c^S$ , resulting in a lower  $\tilde{U}$  at  $30.8 \times 10^{-3}$ .

Let us now focus this discussion by considering the impressive discrepancy with the kinetic exchange of both Cr(O)Rad and Cr(S)Rad couplings in  $1c^{OS}$ , at  $-101$  and  $-741$   $cm^{-1}$ . The small value of the Cr(O)Rad coupling results from a quenching of both metal-to-radical and radical-to-metal relaxations and oppositely, an enhancement of both makes the strong value of Cr(S)Rad. This quenching and this enhancement may be attributed to the  $\tilde{t}$  values, driving these changes. For the



closer to the sulfur atoms side of the complex, the hopping should be eased while the hopping between the radical centre and the other Cr-based centre should be more difficult. This is numerically confirmed with, for instance values at  $-2.4 \times 10^{-3} \text{ cm}^{-1}$  for Cr(S)Rad instead of  $-6.4 \times 10^{-3} \text{ cm}^{-1}$  for Cr(O)Rad in the context of radical-to-metal mechanism. Finally, this final discussion is also in line with the value of direct exchange contributions of  $1\mathbf{c}^{\text{OS}}$  since the Cr(O)Rad  $J_0$  at  $72 \text{ cm}^{-1}$  reflects the interaction between two more distant centres than Cr(S)Rad at  $135 \text{ cm}^{-1}$ .

## 5 Conclusion

Firstly, we applied this method to study the couplings in two series of radical-bridged dinuclear compounds, based on tetraazalene and semiquinone radical ligands, and we have shown the predominance of the direct and kinetic exchange contributions. We get insights into both contributions by decomposing them into their electron–electron interactions between the radical centre and the different d orbitals of the metal ions. This has allowed us to provide orbital analysis with numerical evaluations and we have shown the different roles of the d magnetic orbitals due to the different geometrical arrangements of both series. In addition, we have shown that in the tetraazalene-based compounds, the radical-to-metal is a much more favourable relaxation for kinetic exchange, whilst the situation is very nuanced for the semiquinone-based series.

Secondly, we used the method to investigate the influence of chemical substitution in a bis(bidentate) benzosemiquinoid radical bridging ligand in a series of chromium-based complexes. In addition to the orbital analysis, we have corroborated this study using charge population. Considering first the two symmetrical complexes, we have shown the impact of the nature of the chalcogen atoms on the definition of the magnetic centres and on the resulting coupling between them. Then, we highlighted a large discrepancy between the experimental determination and the theoretical evaluation of the asymmetrical compound. We explained the impact of the asymmetry of radical bridging ligand on the definition of the magnetic centres and the couplings between them.

**Table 8** Atomic Löwdin charge population of the Cr ion and the main atoms of the third series computed at the high spin RO stage in atomic units

Atoms	$1c^O$		$1c^S$		$1c^{OS}$	
1	Cr	-2.27	Cr	-2.72	Cr	-1.89
2	O	0.13	S	0.62	O	-0.11
3	O	0.13	S	0.64	O	-0.16
4	C	0.15	C	-0.09	C	0.17
5	C	0.16	C	-0.07	C	0.14
6	C	-0.05	C	-0.04	C	-0.04
7	C	-0.05	C	-0.04	C	-0.03
8	C	0.16	C	-0.07	C	-0.14
9	C	0.15	C	-0.09	C	-0.14
10	O	0.13	S	0.64	S	0.71
11	O	0.13	S	0.62	S	0.80
12	Cr	-2.27	Cr	-2.72	Cr	-2.78

## Author contributions

Grégoire David carried out the conceptualisation, the implementation, performed simulations, produced the data, the formal analysis, the methodology, the supervision and all the writing process. Gwenhaél Duplaix-Rata carried out the implementation, performed simulations, produced the data and carried out the investigation. Boris Le Guennic carried out the supervision, and the validation and reviewed the paper.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

G. D. is grateful to Dr Mukesh Kumar Singh for having provided the optimised structures of the first series of compounds. The authors thank the French GENCI/IDRIS-CINES centres for high-performance computing resources. G. D. received research funding from the European Unions 843 Horizon 2020 Research and Program under Marie Skłodowska-Curie Grant Agreement No. 899546.

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