



NJ A-CFS: Not Just Another - Crystal Field Software

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1 Introduction

The application of crystal-field (CF) or ligand-field (LF) effective Hamiltonians is a well-established practice in the study of the magnetic properties of inorganic complexes. The history of the crystal field formalism dates back to the pioneering work of Bethe [1]. Since then, various parametrization schemes have been developed to describe the splitting of the free ion terms of a metal center in the presence of surrounding ligands. As a result, many CF/LF programs have been made available in the literature, such as BonnMag [2], SIMPRE [3, 4, 5], CONDON [6, 7], *f_electron* (available at link: https://github.com/octoYot/f_electron), and PyCrystalField [8].

In this context, we introduce NJA-CFS (Not Just Another-Crystal Field Software), with

a nod to the fact that while it may appear to be just another CF software, its underlying purpose sets it apart. NJA-CFS was designed to meet the needs of both first-time users of crystal field theory and those who require a high degree of flexibility in the choice of crystal field parameters (CFPs) formalisms (*vide infra*). Its aim is to provide an intuitive, user-friendly structure that bridges accessibility and the rigor demanded by advanced users.

NJA-CFS serves as a comprehensive toolkit for manipulating crystal field parameters from various sources and exploring how these manipulations influence magnetic properties. Its Python-based foundation ensures excellent readability and visualization capabilities, while also allowing seamless integration with external routines and workflows.

This tutorial will guide you through the program's application, alongside a detailed theoretical explanation of its functions and references to the relevant literature.

In this guide, the sections indicated as:

```
1 # example code
```

represent code listings, while those reported as:

```
$ python main.py
Output
```

represent code printouts in the execution terminal.

2 Requirements

The code folder contains two python scripts: `nja_cfs.py`, which contains all the program functions organized in classes and `test_nja.py`, containing all the examples and test functions to prove the correct functioning of program and its reliability. Additionally, all the additional tables are saved in a dedicated folder.

In order to run the codes you need to install python (version 3.12). The additional dependencies and their versions are: `numpy` (version 2.0.2)[9], `matplotlib` (version 3.9.2)[10], `scipy` (version 1.14.1)[11] and `numba` (version 0.60.0)[12]. However, the program could work also for different versions of the packages.

The functions have been tested on Ubuntu 22.04 LTS inside a dedicated Anaconda environment.

One way to set up the proper environment to run the codes is through the use of `miniconda`, available at <https://docs.anaconda.com/miniconda/>. `Miniconda` is a free, lightweight version of `Anaconda` that installs only Python and the `conda` package manager. It's a good option if you only need Python and don't need all the scientific packages that come with `Anaconda`. Once installed, it can be activated using:

```
$ conda activate
```

the confirmation of the correct activation is confirmed by the presence of `(base)` in the terminal line:

```
(base) $
```

All the packages can be in principle installed with the command:

```
(base) $ conda install <package_name>
```

Warning: Since these are simply python scripts, and NOT an actual installable package, they need always to be present in the execution folder!

3 List of functions and program structure

4 Example scripts

5 Theoretical background

The main contributions included in an effective Hamiltonian to describe the magnetic and electronic properties of small inorganic complexes (e.g. single molecule magnets) are: the electron-electron interaction ($\hat{\mathcal{H}}_{e-e}$), the spin-orbit coupling ($\hat{\mathcal{H}}_{s-o}$), the crystal field splitting ($\hat{\mathcal{H}}_{cf}$) and - if the system is subjected to the action of an external field - the term describing the interaction, e.g. the Zeeman splitting caused by the system exposure to external magnetic fields ($\hat{\mathcal{H}}_Z$):

$$\hat{\mathcal{H}} = \hat{\mathcal{H}}_{e-e} + \hat{\mathcal{H}}_{s-o} + \hat{\mathcal{H}}_{cf} + \hat{\mathcal{H}}_Z \quad (1)$$

The latter term is usually the smaller one.

The extension of the CF effect depends on the type of considered metal ion. For transition metals, the splitting contribution introduced by the ligands is generally bigger than the spin-orbit coupling contribution, while the reverse holds for lanthanoid ions. This will influence the number and classification scheme of the microstates (basis set) included in a hypothetical CF calculation. For example, in the case of f^n configurations the basis set can be reduced to include just the ground state multiplet, which will be almost independent of the coordination environment, like it is done in many known CF programs, e.g. SIMPRE. This will lead to a considerable reduction of the computational burden, especially for the calculation on lanthanoid complexes with $4 \leq n \leq 10$. However, the inclusion of all possible microstates ensures intermediate coupling and J-mixing, therefore this simplification should be evaluated case by case.

Once the Hamiltonian and the basis set classification scheme are defined, the other main point in a CF program is the formalism for the resolution of the integrals. In quantum-mechanical (QM) calculations, Hartree-Fock being the most simple example, the wavefunctions are generally represented as determinantal product states of the type:

$$K_1, K_2, \dots, K_n = (n!)^{-\frac{1}{2}} \sum_P (-1)^p \psi_1(K_1) \psi_2(K_2) \dots \psi_n(K_n) \quad (2)$$

where p is the number of permutations, n is the number of electrons in the considered configuration, ψ_i are the wavefunction basis set, generally composed by the product of an orbital part and a spin part, and K_i (with $i = 1, \dots, n$) are the set of quantum numbers used to unambiguously describe the electron in the specific orbital. Given the computational burden that the resolution of integrals in first quantization with such wavefunctions could bring, an al-

ternative way for Hamiltonian matrix elements calculation has been developed thanks to the work of Condon and Shortley [13], Racah[14], and others, on the application of group theory to the interpretation of atomic spectra.

In this approach the Hamiltonian integrals calculations are performed according to the theory of angular momenta in Racah's tensor operator formalism[15, 16, 17, 18, 19, 20] through the well-known Wigner-Eckart theorem, expressed for a generic tensor operator of rank k and order q as

$$\langle j'm'|T_q^{(k)}|jm\rangle = \langle jmkq|j'm'\rangle \langle j'\|T^{(k)}\|j\rangle \quad (3)$$

where the factors on the r.h.s. are respectively: Clebsch-Gordan coefficients $\langle jmkq|j'm'\rangle$ and the so-called Reduced Matrix Elements (RME) $\langle j'\|T^{(k)}\|j\rangle$ (*vide infra*).

According to this procedure, each operator in equation 1 has to be written in terms of Racah's tensor operators[21] as

$$\begin{aligned} \hat{\mathcal{H}} = & \left(\frac{e^2}{4\pi\epsilon_0} \right) \sum_i^n \sum_{j>i}^n \sum_{k=0}^\infty [r_{>}^{-(k+1)} \cdot r_{<}^k] \sum_{q=-k}^{+k} (-1)^q C_{-q}^{(k)}(i) C_q^{(k)}(j) + \\ & + \hbar^{-2} \zeta \{L^{(1)} \otimes S^{(1)}\}^0 + \\ & + \left(\frac{e^2}{4\pi\epsilon_0} \right) \sum_{i=1}^n \sum_L Z_L \sum_{k=0}^{2L} [r_{>}^{-(k+1)} \cdot r_{<}^k] \sum_{q=-k}^{+k} (-1)^q C_{-q}^{(k)}(L) C_q^{(k)}(i) + \\ & + \mu_B \hbar^{-1} \sum_{q=-1}^{+1} (-1)^q B_{-q}^{(1)} (\kappa L_q^{(1)} + g_e S_q^{(1)}). \end{aligned} \quad (4)$$

A detailed description of the computation of the integrals for each term is reported in following subsections.

To reduce the computational burden further, instead of using the determinantal product of states, the basis wave-vectors are combined to obtain a proper wavefunction through coefficients called: fractional parentage coefficients (cfp) $G_{n,K}^{n-1,K'}$

$$|\psi(l^n, K)\rangle = \sum_{K'} |\psi(l^{n-1}, K')\rangle G_{n,K}^{n-1,K'}. \quad (5)$$

In this equation (analogous to equation 8 of appendix ??), the sum runs over the basis functions of the $n - 1$ states. In this case, K follows the Koster-Nielson classification scheme [22], where the atomic states are constructed in the Russel-Sunders coupling scheme, in which, besides the canonical $|L, S, J, M_J\rangle$, additional quantum numbers, not of physical meaning, are introduced based on the properties of certain mathematical groups. Specifically, these groups are: R_5 in the case of d^n configurations and R_7 and G_2 for f^n configurations. Additionally, the seniority number is also used (more detailed information on cfp's are available in the appendices 5.1 and 5.4).

The Clebsch-Gordan coefficients and RME are generally tabulated. Alternatively, recursive relations are available, based on the use of 3j and 6j-symbols for the first and of cfp's for the latter (see appendix 5.1.1). Also for cfp's, a closed form has been defined by Redmond [23], however, it does not ensure the reproduction of the same coefficients obtained by group-theoretical considerations. A more complete approach, based on the use of Casimir operators, was defined by Nielson in his doctoral thesis[24] (described in appendix 5.1). All

these relations are available in NJA-CFS.

5.1 Fractional parentage coefficients

The following discussion will be limited to f^n configurations (unless otherwise specified) since analogous considerations and the complete set of coefficients for d^n configurations can be easily found in [17].

The determinantal product of states is the most common wavefunction structure used in QM calculations, however, their construction for systems with more than two electrons can be quite cumbersome. To avoid that, the tensor operator representation of Hamiltonian contributions can be exploited. This enables the application of the Wigner-Eckart theorem, whose most general form is the following:

$$\langle \alpha K M_K | T_q^{(k)} | \alpha' K' M'_K \rangle = (-1)^{K-M_K} \begin{pmatrix} K & k & K' \\ -M_K & q & M'_K \end{pmatrix} \langle \alpha K || T^{(k)} || \alpha' K' \rangle \quad (6)$$

where K and M_K are the angular momentum quantum numbers used to discriminate different states. The product of the first two terms on the r.h.s. is a vector coupling coefficient, while the remaining term is called reduced matrix element (RME).

This method can be applied as it is only for configurations comprising two equivalent electrons. For l^n with $n > 2$ the possibility of the occurrence of more than one term with a given K , e.g. with the same L and S , raises the problem of defining a state. So, a classification like the following should be used:

$$|l^n \alpha S L M_S M_L \rangle \quad (7)$$

where α would be the additional quantum number used for terms' differentiation.

Considering that S , L , M_L , M_S are group-theoretical in origin, since e.g. the $2L + 1$ components of a multiplet (with given S and M_S) transform under rotation according to the irreducible representation \mathfrak{D}_L of R_3 , the same principle could be used for other continuous groups that include the operations of R_3 . The method for the individuation of such groups and the application of tensor operators to the theory of continuous groups are due to Racah[25].

Racah found that the group that could serve for such scope are (in order): $R_3 \subset G_2 \subset R_7 \subset U_7$. The quantum numbers associated to each groups are labels for their irreducible representations. G_2 is represented by $U = (u_1, u_2)$ and R_7 is represented by $W = (w_1, w_2, w_3)$. The representation of R_3 (and its subgroups) are the usual orbital angular momentum quantum numbers and the use of irreducible representations of U_7 (represented as $[\lambda_1, \lambda_2, \dots, \lambda_n]$) is equivalent to the specification of the spin quantum number S . The decomposition schemes for the reductions $U_7 \rightarrow R_7$, $R_7 \rightarrow G_2$ and $G_2 \rightarrow R_3$ are reported in tables 5-1, 5-2 and 5-3 of [19] respectively.

However, for few cases, also these classification scheme is not enough, and Racah introduced the seniority quantum number[18], that is (for a certain S, W pair) represented by an integer ν such that f^ν is the first configuration in which that S, W appears.

The immediate advantage of the group theoretical classification of states is that many states can be uniquely without going to the lengths of calculating the particular linear combination of determinantal product state to which they correspond. In order to sfruttare le

potenzialità di questo approccio it is necessary to find a different way for the evaluation of the matrix elements, not requiring the construction of such linear combinations. Racah[17], through the elaboration of an idea of Bacher and Goudsmit[26], developed an elegant solution to this problem.

Assuming that Ω refers to the set of quantum numbers describing the state of l^n configuration, $\bar{\Omega}$ the set of quantum numbers for the state of l^{n-1} and ω the quantum numbers for the one-electron state. Since every determinantal product state for an n -electron system is a sum of terms[19]:

$$|\Omega\rangle = \sum_{\bar{\Omega}, \omega} \langle \bar{\Omega}; \omega | \Omega \rangle |\bar{\Omega}\rangle |\omega_n\rangle \quad (8)$$

where $\langle \bar{\Omega}; \omega | \Omega \rangle$ are the coefficients of our interest and the subscript n indicates that the last state in the equation refers to the n th electron.

Using this equation, the matrix elements for a generic single-particle operator $F = \sum_i f_i$, between two states of the l^n configuration, can be expressed as:

$$\langle \Omega | F | \Omega' \rangle = n \sum_{\bar{\Omega}, \omega, \omega'} \langle \Omega | \bar{\Omega}; \omega \rangle \langle \omega_n | f_n | \omega_n \rangle \langle \bar{\Omega}; \omega' | \Omega' \rangle \quad (9)$$

while, for a two-particles operator $G = \sum_{i>j} g_{ij}$:

$$\langle \Omega | G | \Omega' \rangle = [n/(n-2)] \sum_{\bar{\Omega}, \bar{\Omega}', \omega} \langle \Omega | \bar{\Omega}; \omega \rangle \langle \bar{\Omega} | \sum_{j<i \neq n} g_{ij} | \bar{\Omega}' \rangle \langle \bar{\Omega}'; \omega | \Omega' \rangle \quad (10)$$

The procedure used for the computation of the cfp is taken from the Nielson PhD thesis[24].

According to the group-theoretical classification of states:

- the single-electron wavefunction, with quantum numbers: $S = 1/2$, $W = (100)$, $U = (10)$, $L = 3$, m_l , m_s , correspond to the irreducible representation Γ_f ,
- the antisymmetric states of $n-1$ electrons, with quantum numbers: S' , W' , U' , α' , L' , M'_L , M'_S , constitute the basis of the irreducible representation $\Gamma_A^{(n-1)}$,
- the antisymmetric states of n electrons, with quantum numbers: S , W , U , α , L , M_L , M_S , constitute the basis of the irreducible representation $\Gamma_A^{(n)}$.

The objective is to find the reduction coefficients of the Kronecker product $\Gamma_A^{(n-1)} \times \Gamma_f$ relative to the antisymmetric part $\Gamma_A^{(n)}$, indicated in Nielson's notation as:

$$b_n(S, W, U, \alpha, L, M_L, M_S; S', W', U', \alpha', L', M'_L, M'_S, m_l, m_s) \quad (11)$$

These coefficients can be factorized according to the Racah's theorem as follows:

$$b_n(S, W, U, \alpha, L, M_L, M_S; S', W', U', \alpha', L', M'_L, M'_S, m_l, m_s) = C_{n, W'}^{S' \ 1/2 \ S} C_{U' \ (10) \ U}^{W' \ (100) \ W} C_{\alpha' L' \ 3 \ \alpha L}^{U' \ (10) \ U} C_{M'_L \ m_l \ M_L}^{L' \ 3 \ L} C_{M'_S \ m_s \ M_S}^{S' \ 1/2 \ S} \quad (12)$$

where the last two coefficients are Clebsch-Gordon coefficients and the product of the other

three compose the cfp:

$$a_n(S, W, U, \alpha, L; S', W', U', \alpha', L') = C_{n, W' (100) W}^{S' 1/2 S} C_{U' (10) U}^{W' (100) W} C_{\alpha' L' 3 \alpha L}^{U' (10) U} \quad (13)$$

In Racah's notation (used from now on):

$$(I^{n-1}(\alpha' \nu' S' L') | SL | \} I^n \alpha \nu SL) \equiv a_n(S, W, U, \alpha, L; S', W', U', \alpha', L') \quad (14)$$

$$(f^{n-1} \nu' S' + f | \} f^n \nu S) \equiv C_{n, W' (100) W}^{S' 1/2 S} \quad (15)$$

$$(W' U' + f | WU) \equiv C_{U' (10) U}^{W' (100) W} \quad (16)$$

$$(U' \alpha' L' + f | U \alpha L) \equiv C_{\alpha' L' 3 \alpha L}^{U' (10) U} \quad (17)$$

For the coefficients $(f^{n-1} \nu' S' + f | \} f^n \nu S)$ a closed form is available (equations 52a-52b-56 of [18]). The other coefficients $(W' U' + f | WU)$ and $(U' \alpha' L' + f | U \alpha L)$, which represent the reduction coefficients of the Kronecker product of groups R_7 and G_2 respectively, can be computed via the diagonalization of the corresponding Casimir operators (G) (equivalent of a diagonalization of L^2 operator for the reduction of a representation of the group R_3). The Racah's expression of Casimir operators are:

$$G(R_7) = \frac{3}{5}(U^1)^2 + \frac{7}{5}(U^3)^2 + \frac{11}{5}(U^5)^2 \quad (18)$$

$$G(G_2) = \frac{3}{4}(U^1)^2 + \frac{11}{4}(U^5)^2 \quad (19)$$

where U^1, U^3 and U^5 are unit tensor operators, with known eigenvalues:

$$\lambda(W) = w_1(w_1 + 5)/2 + w_2(w_2 + 3)/2 + w_3(w_3 + 1)/2 \quad (20)$$

$$\lambda(U) = (u_1^2 + u_1 u_2 + u_2^2 + 5u_1 + 4u_2)/12 \quad (21)$$

The matrix elements of $G(G_2)$ between the states arising from the coupling of a single f electron to parent states of quantum numbers U', L', M'_L to give the final states of angular momentum L, M_L , can be expressed as:

$$\begin{aligned} & \langle U' \alpha' L' + f \rightarrow LM_L | G(G_2) | U' \alpha'' L'' + f \rightarrow LM_L \rangle = \\ & = (-1)^{L+3+L''} \frac{11}{2} \left\{ \begin{matrix} L & 3 & L' \\ 5 & L'' & 3 \end{matrix} \right\} (U' \alpha' L' \| U_{n-1}^5 \| U' L'' \alpha'') + \\ & + \delta_{\alpha' \alpha''} \delta_{L' L''} \left[\lambda(U') + \lambda(10) + \frac{1}{112} (L(L+1) - L'(L'+1) - 3(3+1)) \right] \end{aligned} \quad (22)$$

Once these matrix elements are evaluated for a chosen L and U' , its diagonalization produces eigenvalues for U containing L and known to occur in the reduction $(U') \times (10)$. Each (normalized) eigenvector is a whole set of $(U' \alpha' L' + f | U \alpha L)$ with L' varying along the row.

The RME of U^5 are computed iteratively, from the previous cycle of calculation. In this case the labels $n, n-1, n-2$ are used only to represent the hierarchy of computation, these coefficients are general and do not depend on the particular number of electrons considered.

The necessary relations are the following:

$$\begin{aligned}
& (U'''L''' + f \rightarrow L' \| U_{n-1}^5 \| U'''L'^v + f \rightarrow L'') = \\
& = (2L' + 1)^{1/2} (2L'' + 1)^{1/2} \left[(-1)^{L''+L'} (U'''L''' \| U_{n-2}^5 \| U'''L'^v) \right. \\
& \times \left\{ \begin{matrix} L''' & L' & 3 \\ L'' & L'^v & 5 \end{matrix} \right\} + (-1)^{L'^v+L'} \delta_{L'''L'^v} \delta_{\alpha''' \alpha'^v} \left\{ \begin{matrix} 3 & L' & L''' \\ L'' & 3 & 5 \end{matrix} \right\} \left. \right] \quad (23)
\end{aligned}$$

and

$$\begin{aligned}
& (U'\alpha'L' \| U_{n-1}^5 \| U'\alpha''L'') = \\
& = \sum_{\alpha''', L''', \alpha'^v, L'^v} (U''' \alpha''' L''' + f | U'\alpha'L') (U'^v \alpha'^v L'^v + f | U'\alpha'L') \\
& \times (U'''L''' + f \rightarrow L' \| U_{n-1}^5 \| U'''L'^v + f \rightarrow L'') \quad (24)
\end{aligned}$$

An analogous procedure can be applied (subsequently) for the computation of $(W'U' + f | WU)$. The matrix elements of $G(R_7)$ are expressed as:

$$\begin{aligned}
& \langle W'U' + f \rightarrow ULM_L | G(R_7) | W'U'' + f \rightarrow ULM_L \rangle = \\
& = \sum_{L', L''} \left[(U'L' + f | UL) (U''L'' + f | UL) (-1)^{L''+3+L} \right. \\
& \times \frac{14}{5} \left\{ \begin{matrix} L & 3 & L' \\ 3 & L'' & 3 \end{matrix} \right\} (W'U'L' \| U_{n-1}^3 \| W'U''L'') \left. \right] + \\
& + \delta_{U'U''} \left[\frac{4}{5} \lambda(U) + \lambda(W') - \frac{4}{5} \lambda(U') + \lambda(100) - \frac{4}{5} \lambda(10) \right] \quad (25)
\end{aligned}$$

In this case, the W that have to be considered are those that occur in the reduction $W' \times (100)$. The iterative procedure for the computation of the RME for U^3 is similar to the one described for U^5 , with equations:

$$\begin{aligned}
& (W'U'L' \| U_{n-2}^3 \| W'U''L') = \\
& = \sum_{U''', U'^v} (W'''U''' + f | W'U') (W'''U'^v + f | W'U'') \\
& \times \left[\sum_{L''', L'^v} (U'''L''' + f | U'L') (U'^v L'^v + f | U''L'') \right. \\
& \times (W'''U'''L''' + f \rightarrow L' \| U_{n-1}^3 \| W'''U'^v L'^v + f \rightarrow L'') \left. \right] \quad (26)
\end{aligned}$$

and:

$$\begin{aligned}
& (W'''U''' + f \rightarrow L''' \| U_{n-1}^3 \| W'''U'^v + f \rightarrow L'') = \\
& = (-1)^{L''' + 3 + L'' + 3} (2L' + 1)^{1/2} (2L'' + 1)^{1/2} \\
& \times \left\{ \begin{matrix} L''' & L' & 3 \\ L'' & L'^v & 3 \end{matrix} \right\} (W'''U'''L''' \| U_{n-2}^3 \| W'''U'^v L'^v) + \\
& + (-1)^{L''' + 3 + L' + 3} \delta_{L'''L'^v} \delta_{U'''U'^v} (2L' + 1)^{1/2} (2L'' + 1)^{1/2} \left\{ \begin{matrix} 3 & L' & L''' \\ L'' & 3 & 3 \end{matrix} \right\} \quad (27)
\end{aligned}$$

5.1.1 Reduced matrix elements

The RMEs for the different tensor operators used in the integrals computation for the physical operators (vide infra) can be computed using the following equations, using the calculated cfps, for a generic rank k and configuration l^n :

$$\begin{aligned}
& (\alpha LS \| U^{(k)} \| \alpha' L' S') = \delta_{SS'} n [(2L + 1)(2L' + 1)]^{1/2} \\
& \times \sum_{\alpha'', L'', S''} (l^{n-1} (\alpha'' L'' S'') l SL | \} l^n \alpha LS) (l^{n-1} (\alpha'' L'' S'') l SL | \} l^n \alpha' L' S') \\
& \times (-1)^{K+L+l+L''} \left\{ \begin{matrix} l & L & L'' \\ L' & l & k \end{matrix} \right\} \quad (28)
\end{aligned}$$

and:

$$\begin{aligned}
& (\alpha LS \| V^{(1k)} \| \alpha' L' S') = \\
& = \frac{3}{2} [(2L + 1)(2L' + 1)(2S + 1)(2S' + 1)]^{1/2} \\
& \times \sum_{\alpha'', L'', S''} (l^{n-1} (\alpha'' L'' S'') l SL | \} l^n \alpha LS) (l^{n-1} (\alpha'' L'' S'') l SL | \} l^n \alpha' L' S') \\
& \times (-1)^{k+L+l+L''} \left\{ \begin{matrix} l & L & L'' \\ L' & l & k \end{matrix} \right\} (-1)^{1+S+1/2+S''} \left\{ \begin{matrix} 1/2 & S & S'' \\ S' & 1/2 & 1 \end{matrix} \right\} \quad (29)
\end{aligned}$$

In both equations, the 6-j symbol $\left\{ \begin{matrix} a & b & c \\ A & B & C \end{matrix} \right\}$ represents the coupling of three angular momenta, and can be solved using the Racah's equation[16]:

$$\begin{aligned}
& \left\{ \begin{matrix} a & b & c \\ A & B & C \end{matrix} \right\} = (-1)^{a+b+A+B} f(abc) f(ABc) f(AbC) f(aBC) \\
& \sum_{n=n_{\min}}^{n_{\max}} (-1)^n (a+b+A+B+1-n)! / [n!(a+b-c-n)! \\
& \times (A+B-c-n)!(a+B-C-n)!(A+b-C-n)! \\
& \times (-a-A+c+C+n)!(-b-B+c+C+n)!] \quad (30)
\end{aligned}$$

with:

$$f(abc) = \left[\frac{(a+b-c)!(a-b+c)!(-a+b+c)!}{(a+b+c+1)!} \right]^{1/2}$$

and:

$$n_{\min} = \max\{0; a + A - c - C; b + B - c - C\}$$

$$n_{\max} = \min\{a + b + A + B + 1; a + b - c; A + B - c; a + B - C; A + b - C\}$$

5.2 Crystal field coefficients formalism

The perturbation of the electron cloud of a metal ion, subjected to the influence of a coordination environment, can be effectively described by a crystal-field Hamiltonian of the type:

$$\hat{\mathcal{H}}_{cf} = -e \sum_{i=1}^n V(r_i) \quad (31)$$

where e is the elementary charge, the summation i runs over the n electrons in the open-shell orbitals and $V(r_i)$ is the potential felt by the electron at the position r_i . The crystal field potential, in presence of a field independent charge distribution $\rho(R)$, can be expressed as:

$$V(r_i) = \int \frac{\rho(R)}{|R - r_i|} d\tau \quad (32)$$

where R indicates a general point in the ligand framework and $d\tau$ is a volume element. Alternatively, considering a discrete distribution of charges represented in a point charge model (PCM), the following expression is valid:

$$V(r_i) = \sum_L \frac{(-Ze)_L}{|R_L - r_i|} \quad (33)$$

where Z is the fraction of negative charge relative to the L th ligand.

The potential in equation 32 can be expanded as follows:

$$\frac{1}{|R - r_i|} = \sum_{k=0}^{\infty} \frac{r_{<}^k}{r_{>}^{k+1}} P_k(\cos \omega) \quad (34)$$

where $r_{<}$ and $r_{>}$ are the shortest and longest distances respectively and $P_k(\cos \omega)$ are the Legendre polynomials relative to the angle ω between R and r_i . Generally, the ligands are more distant than the electron of the metal ion itself, therefore equation 34 becomes:

$$\frac{1}{|R - r_i|} = \sum_{k=0}^{\infty} \frac{r_i^k}{R^{k+1}} P_k(\cos \omega) \quad (35)$$

$P_k(\cos \omega)$ can be further expanded according to the spherical harmonics addition theorem [27]. According to that, ω can be decomposed in the polar angles: (θ, ϕ) , which describe the distribution of ligand charges, and (θ_i, ϕ_i) , which describe the position of the i th-electron:

$$P_k(\cos \omega) = \sqrt{\frac{4\pi}{2k+1}} \sum_{q=-k}^k Y_k^{q*}(\theta, \phi) Y_k^q(\theta_i, \phi_i) \quad (36)$$

leading to the final expression:

$$V(r_i) = \sum_{k=0}^{\infty} \sqrt{\frac{4\pi}{2k+1}} \left[\sum_{q=-k}^k Y_k^q(\theta_i, \phi_i) \int \rho(R) \frac{r_i^k}{R^{k+1}} Y_k^{q*}(\theta, \phi) d\tau \right] \quad (37)$$

or in the PCM:

$$V(r_i) = \sum_{k=0}^{\infty} \sqrt{\frac{4\pi}{2k+1}} \left[\sum_{q=-k}^k Y_k^q(\theta_i, \phi_i) \sum_L (-Ze)_L \frac{r^k}{R_L^{k+1}} Y_k^{q*}(\theta_L, \phi_L) \right] \quad (38)$$

The radial part of this expression and the angular part describing the ligand distribution are usually factorized in the so-called crystal field parameters (CFPs), with general expression [28]:

$$V(r_i) = \sum_{kq} f_{kq}(R) Y_k^q(\theta_i, \phi_i) \quad (39)$$

This factorization can be performed in many ways, giving rise to the multitudes of formalism available in literature (vide infra). I implemented some of them - and their conversion factors - in NJA-CFS.

Throughout the whole discussion on CFPs I will use the representation formalism employed by Ryabov and Rudowicz in their numerous reviews on the topic (referenced below).

As previously described, the (electric) crystal field of an open-shell ion in a crystalline environment can be described by a potential function which satisfies the Laplace equation. The solution of this equation can be expressed in a series of spherical harmonics. Stevens (1952) was the first one to show that the matrix elements of these spherical harmonics, within the same angular momentum manifold, could be calculated by replacing them by operator equivalents having the similar transformation properties. Since then, tensor operators (TOs) of angular momenta [16] have been widely used in the construction of crystal field Hamiltonians [29] and (surely too) many conventions and formalism have been presented since then[30]. Two classes of TOs can generally be identified: the spherical tensor operators (STOs) and the tesseral tensor operators (TTOs), which are the operator equivalents to the spherical and tesseral harmonics respectively. The conventional Stevens operators (CSOs) [31] and the extended Stevens operators (ESOs)[30] belongs to the TTO class. Both of them are generally indicated as B_k^q , where $q \geq 0$ for the CSOs and $-k \leq q \leq k$ for the ESOs. The first are not used anymore since they do not transform consistently under a general rotation of coordinates for some crystal symmetries, as first pointed out by Buckmaster (1962) [32]. Nowadays, the most used operator equivalents are the Stevens ESOs one[33]. According to this approach, the crystal field hamiltonian in a single $J(f^n)$ -multiplet or $L(d^n)$ -multiplet can be expressed as:

$$\hat{\mathcal{H}}_{\text{cf}}^{\text{Stevens}} = \sum_{k,q} B_k^q O_k^q(J \text{ or } L) \quad (40)$$

where O_k^q are the ESOs, either defined in terms of total angular momentum J or orbital angular momentum L . The B_k^q defined herein should not be confused with the B_k^q from the $\hat{\mathcal{H}}_{\text{ZFS}}$, where the O_k^q are defined in terms of the spin angular momentum S . The CFPs defined in this way are real-valued. Another widespread notation used to express the crystal field

Table 1: Values of radial integrals $\langle r^k \rangle$ (for $k = 2, 4, 6$) computed by Edvardsson and Klintenberg [34].

configuration	2	4	6
f^1	1.456	5.437	42.26
f^2	1.327	4.537	32.65
f^3	1.222	3.875	26.12
f^4	1.135	3.366	21.46
f^5	1.061	2.964	17.99
f^6	0.997	2.638	15.34
f^7	0.942	2.381	13.36
f^8	0.893	2.163	11.75
f^9	0.849	1.977	10.44
f^{10}	0.810	1.816	9.345
f^{11}	0.773	1.677	8.431
f^{12}	0.740	1.555	7.659
f^{13}	0.710	1.448	7.003

Table 2: Stevens coefficients α , β and γ for f^n configurations[31].

configuration	α	β	γ
f^1	$-\frac{2}{35}$	$\frac{2}{7 \cdot 45}$	0
f^2	$-\frac{11 \cdot 15^2}{32}$	$-\frac{55 \cdot 33 \cdot 3}{8 \cdot 17}$	$\frac{17 \cdot 16}{7 \cdot 11^2 \cdot 13 \cdot 5 \cdot 3^4}$
f^3	$-\frac{33^2}{14}$	$-\frac{11^2 \cdot 13 \cdot 297}{952}$	$-\frac{13^2 \cdot 11^3 \cdot 3^3 \cdot 7}{17 \cdot 19 \cdot 5}$
f^4	$\frac{11^2 \cdot 15}{13}$	$\frac{13 \cdot 3^3 \cdot 11^3 \cdot 5}{26}$	$\frac{11^2 \cdot 13^2 \cdot 3 \cdot 63}{2584}$
f^5	$\frac{13}{7 \cdot 45}$	$\frac{33 \cdot 7 \cdot 45}{26}$	0
f^6	0	0	0
f^7	0	0	0
f^8	$-\frac{1}{99}$	$\frac{2}{11 \cdot 1485}$	$-\frac{1}{13 \cdot 33 \cdot 2079}$
f^9	$-\frac{9 \cdot 35}{2}$	$-\frac{11 \cdot 45 \cdot 273}{8}$	$\frac{11^2 \cdot 13^2 \cdot 3^3 \cdot 7}{5}$
f^{10}	$-\frac{1}{30 \cdot 15}$	$-\frac{1}{11 \cdot 2730}$	$-\frac{13 \cdot 33 \cdot 9009}{8}$
f^{11}	$\frac{45 \cdot 35}{4}$	$\frac{11 \cdot 15 \cdot 273}{8}$	$\frac{13^2 \cdot 11^2 \cdot 3^3 \cdot 7}{5}$
f^{12}	$\frac{1}{99}$	$\frac{3 \cdot 11 \cdot 1485}{2}$	$-\frac{13 \cdot 33 \cdot 2079}{4}$
f^{13}	$\frac{2}{63}$	$-\frac{77 \cdot 15}{2}$	$\frac{13 \cdot 33 \cdot 63}{4}$

hamiltonian, generally used for f^n multiplets, in terms of ESOs is the following:

$$\hat{\mathcal{H}}_{\text{cf}}^{\text{Stevens}} = \sum_{k,q} A_k^q \langle r^k \rangle \theta_k O_k^q \quad (41)$$

where A_k^q are the CFPs (like the B_k^q), $\langle r^k \rangle$ are the expectation value of r^k (listed in table 1 for f^n , analogous values are available also for d^n configurations) and the θ_k are the multiplicative Stevens factors, indicated as α , β and γ in the original Stevens paper from '52 [31] for the rank $k = 2, 4, 6$ respectively. The properties of TTOs differ from the STOs. An example of the latter is represented by the Wybourne operators.

According to Wybourne (1965)[35, 36, 5], the crystal field hamiltonian can be defined as:

$$\hat{\mathcal{H}}_{\text{cf}}^{\text{Wybourne}} = \sum_{k,q,i} B_q^k C_q^{(k)}(\theta_i, \phi_i) \quad (42)$$

where the sum over i runs on the electrons in the chosen (open-shell) configuration and $-k \leq q \leq k$ [36, 37]. $C_q^{(k)}$ (also called Racah's harmonics[16]) are Wybourne's spherical tensor operators $T_q^{(k)}$ equivalent to the renormalized spherical harmonics:

$$C_q^{(k)} = \sqrt{\frac{4\pi}{2k+1}} Y_k^q \quad (43)$$

Alternatively, by rewriting equation 36 as:

$$P_k(\cos \omega) = \frac{4\pi}{2k+1} \left[Y_k^0 Y_k^0(i) + \sum_{q=1}^k (Y_k^{-q} Y_k^q(i) + Y_k^q Y_k^{-q}(i)) \right] \quad (44)$$

where the spherical harmonics with (i) refers to i th-electron position, while the others refer to the ligands position. The Hamiltonian can be expressed as:

$$\begin{aligned} \hat{\mathcal{H}}_{\text{cf}}^{\text{Wybourne}} = & -e \sum_{k,i} \left[B_0^k C_0^{(k)}(i) \right. \\ & \left. + \sum_{q=1}^k (B_q^k (C_{-q}^{(k)}(i) + (-1)^q C_q^{(k)}(i)) + B_q'^k (C_{-q}^{(k)}(i) - (-1)^q C_q^{(k)}(i))) \right] \end{aligned} \quad (45)$$

where k has values 2 and 4 for d^n and 2, 4 and 6 for f^n configurations. The B_q^k and $B_q'^k$ are the so-called crystal field coefficients (CFCs), that can be converted in the corresponding crystal field parameters B_q^k and $B_q'^k$, when solving the integral, by bringing them outside the matrix element, together with the radial parts of the wavefunctions $R_{nl}(r)$. Therefore, the CFPs contains the electron charge $-e$, a radial and an angular part. In this way, the tensor operators are left acting solely on the angular part. The use of the same notation for both coefficients and parameters is confusing but, unfortunately, it is what is done in literature.

Due to the operators' properties, the Stevens CFPs B_k^q (or A_k^q) in equations 40 and 41 are real-valued, whereas the Wybourne CFPs B_q^k are in general complex. All operators of the STO class are simply related to each other by specific numerical coefficients arising from the Wigner-Eckhart theorem while the same is not true for the operators of the TTO class [37]. Furthermore, while equations 40 and 41 are valide only in J (or L)-constants manifolds, expression 42 is applicable to the complete set of microstates.

In order to derive the relationship between the two set of operators we can express the O_k^q in terms of $T_q^{(k)}$:

$$O_k^0 = \lambda_{k0} T_0^{(k)} \quad (46)$$

$$O_k^q = \lambda_{kq} (T_{-q}^{(k)} + (-1)^q T_q^{(k)}) \quad (47)$$

$$O_k^{-q} = i\lambda_{kq} (T_{-q}^{(k)} - (-1)^q T_q^{(k)}) \quad (48)$$

The conversion coefficients ($\lambda_{kq} = B_q^k / A_k^q \langle r^k \rangle$) are listed in table 3.

The formalism actually used in the integrals computation is the Wybourne one. The coefficients can either be fed into the function *MatrixH* (that performs the computation of energy levels and eigenfunctions) directly, through the dictionary *dic_bkq*, or using a PCM model, via

Table 3: The conversion factors λ_{kq} between the Stevens CFPs and the Wybourne CFPs, for $k = 2, 4, 6$ [5].

q	2	4	6
0	2	8	16
1	$1/\sqrt{6}$	$2/\sqrt{5}$	$8/\sqrt{42}$
2	$2/\sqrt{6}$	$4/\sqrt{10}$	$16/\sqrt{105}$
3		$2/\sqrt{35}$	$8/\sqrt{105}$
4		$8/\sqrt{70}$	$16/3\sqrt{14}$
5			$8/3\sqrt{77}$
6			$16/\sqrt{231}$

the PCM item, applying the following equations[5]:

$$B_0^k = \sqrt{\frac{4\pi}{2k+1}} \langle r^k \rangle \sum_L \frac{Z_L e^2}{R_L^{k+1}} Y_k^0(\theta_L, \phi_L) \quad (49)$$

$$B_q^k = \sqrt{\frac{4\pi}{2k+1}} (-1)^q \langle r^k \rangle \sum_L \frac{Z_L e^2}{R_L^{k+1}} \text{Re} Y_k^q(\theta_L, \phi_L) \quad (50)$$

$$B_q^k = \sqrt{\frac{4\pi}{2k+1}} (-1)^q \langle r^k \rangle \sum_L \frac{Z_L e^2}{R_L^{k+1}} \text{Im} Y_k^q(\theta_L, \phi_L) \quad (51)$$

Alternatively, the B_q^k can be computed from $A_k^q \langle r^k \rangle$ using the λ_{kq} factors, using the *from_Akqrk_to_Bkq*. A function for the calculation of the Stevens coefficients is also implemented in *calc_Akqrk* from PCM:

$$A_q^0 = \sqrt{4\pi 2k+1} \sum_L \frac{Z_L e^2}{R_L^{k+1}} Z_{k0}(\theta_L, \phi_L) p_{k0} \quad (52)$$

$$A_k^q = \sqrt{4\pi 2k+1} \sum_L \frac{Z_L e^2}{R_L^{k+1}} Z_{kq}^c(\theta_L, \phi_L) p_{kq} \quad (53)$$

$$A_k^{-q} = \sqrt{4\pi 2k+1} \sum_L \frac{Z_L e^2}{R_L^{k+1}} Z_{kq}^s(\theta_L, \phi_L) p_{kq} \quad (54)$$

where p_{kq} are tesseral hamronics' numerical prefactors (listed in table 4) and:

$$Z_{k0} = Y_k^0 \quad (55)$$

$$Z_{kq}^c = \frac{1}{\sqrt{2}} \text{Re}(Y_k^{-q} + (-1)^q Y_k^q) \quad (56)$$

$$Z_{kq}^s = -\frac{1}{\sqrt{2}} \text{Im}(Y_k^{-q} - (-1)^q Y_k^q) \quad (57)$$

All the equation presented so far have proven that it is possible to express the ligand-field potential as linear combination of spherical harmonics multiplied by coefficients, bearing information on radial part of the electronic wavefunction and ligand's wavefunction. This is true also at the one-electron level:

$$\langle l, m_l | V | l, m_l' \rangle = \sum_j c_{kq}^j \langle l, m_l | Y_k^q | l, m_l' \rangle \quad (58)$$

Table 4: Tesseral harmonics' prefactors p_{kq} , for $k = 2, 4, 6$ [5].

q	2	4	6
0	$\frac{1}{4}\sqrt{\frac{5}{\pi}}$	$\frac{3}{16}\sqrt{\frac{1}{\pi}}$	$\frac{1}{32}\sqrt{\frac{13}{\pi}}$
1	$\frac{1}{2}\sqrt{\frac{15}{\pi}}$	$\frac{3}{4}\sqrt{\frac{5}{2\pi}}$	$\frac{1}{8}\sqrt{\frac{273}{4\pi}}$
2	$\frac{1}{4}\sqrt{\frac{15}{\pi}}$	$\frac{3}{8}\sqrt{\frac{5}{\pi}}$	$\frac{1}{64}\sqrt{\frac{2730}{\pi}}$
3		$\frac{3}{8}\sqrt{\frac{70}{\pi}}$	$\frac{1}{32}\sqrt{\frac{2730}{\pi}}$
4		$\frac{3}{16}\sqrt{\frac{35}{\pi}}$	$\frac{21}{32}\sqrt{\frac{13}{7\pi}}$
5			$\sqrt{\frac{9009}{512\pi}}$
6			$\frac{231}{64}\sqrt{\frac{26}{231\pi}}$

where j runs over the ligand indices and $\langle l, m_l | V | l, m_l' \rangle$ are one-electron ligand field integrals in the complex basis, elements of the one-electron ligand field matrix (V^{LF}). Since the $\langle l, m_l | Y_k^q | l, m_l' \rangle$ integrals can be easily solved using the tensor operator formalism (vide infra), what we get is a series of equations that express the c_{kq}^j coefficients in terms of $\langle l, m_l | V | l, m_l' \rangle$. This can be expressed in matrix form as:

$$\mathbf{c} = \mathbf{C} \mathbf{M} \quad (59)$$

where \mathbf{c} and \mathbf{M} are vectors containing the $(l+1)(2l+1)$ independent coefficients and V^{LF} independent matrix elements respectively, and \mathbf{C} is the $(l+1)(2l+1) \times (l+1)(2l+1)$ coefficient matrix. This implementation allows easily also the reverse process, i.e. compute the ligand field matrix from the coefficients via $\mathbf{M} = \mathbf{C}^{-1}\mathbf{c}$. In order for the procedure to properly work, the elements in \mathbf{c} , \mathbf{C} and \mathbf{M} have to be coherently ordered.

Since, for NJA-CFS, we are interested in real-valued CF coefficients, I implemented this procedure (in the *from_Vint_to_Bkq_2* function) using the ligand-field matrix elements in the real basis and the \mathbf{C} matrix obtained exploiting the relations between the real basis and the complex basis:

$$|0\rangle = |\sigma\rangle \quad (60)$$

$$|1\rangle = -\frac{i}{\sqrt{2}}|\pi_s\rangle - \frac{1}{\sqrt{2}}|\pi_c\rangle \quad (61)$$

$$|-1\rangle = -\frac{i}{\sqrt{2}}|\pi_s\rangle + \frac{1}{\sqrt{2}}|\pi_c\rangle \quad (62)$$

$$|2\rangle = \frac{i}{\sqrt{2}}|\delta_s\rangle + \frac{1}{\sqrt{2}}|\delta_c\rangle \quad (63)$$

$$|-2\rangle = -\frac{i}{\sqrt{2}}|\delta_s\rangle + \frac{1}{\sqrt{2}}|\delta_c\rangle \quad (64)$$

$$|3\rangle = -\frac{i}{\sqrt{2}}|\phi_s\rangle - \frac{1}{\sqrt{2}}|\phi_c\rangle \quad (65)$$

$$|-3\rangle = -\frac{i}{\sqrt{2}}|\phi_s\rangle + \frac{1}{\sqrt{2}}|\phi_c\rangle \quad (66)$$

where the complex basis is indicated as $|l, m_l\rangle = |m_l\rangle$, while the real basis is indicated with

$|l, u\rangle = |u\rangle$ (with $u = \sigma, \pi_s, \pi_c, \delta_s, \delta_c, \phi_s, \phi_c$). I chose this convention for the real orbitals since it is the one used in the papers on the subject. The expression are the same as the one reported in table 2 of [38] for d^n configurations and table 3 of [39] for f^n configurations.

The V^{LF} matrix elements can be computed ab initio through what is called: ab initio ligand field theory (AILFT). The one-electron ligand field matrices of a AILFT from an ORCA CASSCF calculation are organized as follows: for d^n configurations ($l = 2$), the elements are displayed with $m_l = [0, 1, -1, 2, -2]$, or, using the ORCA's orbital labels, as $[d_{z^2}, d_{xz}, d_{yz}, d_{x^2-y^2}, d_{xy}]$, while for f^n configurations ($l = 3$), the elements are displayed with $m_l = [0, 1, -1, 2, -2, 3, -3]$, or, using the ORCA's orbital labels, as $[f_0, f_{+1}, f_{-1}, f_{+2}, f_{-2}, f_{+3}, f_{-3}]$. In the notation used in equations 60, the orbitals are ordered as: $[\sigma, \pi_c, \pi_s, \delta_c, \delta_s, \phi_c, \phi_s]$. An important note: the phases of f_{+3} and f_{-3} orbitals in ORCA are different from the ones generally used, so in NJA-CFS a sign change is performed for the mixed integrals involving those orbitals. Those matrix are obtained using the *actorb* flag in the ORCA *%casscf* block (see the ORCA manual for further details).

Another alternative to the (electrostatically-based) crystal field models, is the angular overlap model (AOM). Analogously to the other ligand-field treatment, also this one is based on one-electron operators. It assumes that the antibonding-orbital energies (E^*) are determined by covalent perturbation weak enough to be proportional to the square of appropriate overlap integrals:

$$E^* = (A_t^d)^2 e_t \quad (67)$$

where d indicates the metal d orbital and t the symmetry of the ligand orbital (i.e. σ, π, \dots). A_t^d is an angular factor expressing the fact that the chosen global coordinate frame of the metal atom is not generally coincident with the local frame to which the symmetry of the ligand functions are referred. e_t is the parameter enclosing the information on the overlap integral between the metal and ligand orbitals and it is different for each ligand type and for each bonding mode. Contributions arising from the different ligands to this antibonding energy are considered additive. A general expression for the one-electron matrix elements within the AOM model is the following:

$$\langle l, u | V | l, v \rangle = \sum_j^{\text{ligands}} \sum_t^{\text{modes}} D_{ut}^l(j) D_{vt}^l(j) e_t(j) \quad (68)$$

where D_{ut}^l are the matrix elements of the transformation of the real metal orbitals into each ligand coordinate frame. In direction cosines, the D matrix elements for the modes e_σ , e_{π_c} and e_{π_s} is the following, for $l = 2$ (equation 9 of [38]):

$$\begin{bmatrix} (1/2)(3\gamma_3^2 - 1) & \sqrt{3}\alpha_3\gamma_3 & \sqrt{3}\beta_3\gamma_3 \\ \sqrt{3}\gamma_1\gamma_3 & \alpha_1\gamma_3 + \alpha_3\gamma_1 & \beta_1\gamma_3 + \beta_3\gamma_1 \\ \sqrt{3}\gamma_2\gamma_3 & \alpha_2\gamma_3 + \alpha_3\gamma_2 & \beta_2\gamma_3 + \beta_3\gamma_2 \\ \sqrt{3}\gamma_1\gamma_2 & \alpha_1\gamma_2 + \alpha_2\gamma_1 & \beta_1\gamma_2 + \beta_2\gamma_1 \\ (1/2)\sqrt{3}(\gamma_1^2 - \gamma_2^2) & \alpha_1\gamma_1 - \alpha_2\gamma_2 & \beta_1\gamma_1 - \beta_2\gamma_2 \end{bmatrix} \quad (69)$$

and for $l = 3$ (table 1 of [39]):

$$\begin{bmatrix}
 (1/2)\gamma_3(5\gamma_3^2 - 3) & (1/2)\sqrt{3/2}\beta_3(5\gamma_3^2 - 1) & (1/2)\sqrt{3/2}\alpha_3(5\gamma_3^2 - 1) \\
 (1/4)\sqrt{6}\gamma_2(5\gamma_3^2 - 1) & (1/4)\beta_2(5\gamma_3^2 - 1) + (5/2)\gamma_2\gamma_3\beta_3 & (1/4)\alpha_2(5\gamma_3^2 - 1) + (5/2)\gamma_2\gamma_3\alpha_3 \\
 (1/4)\sqrt{6}\gamma_1(5\gamma_3^2 - 1) & (1/4)\beta_1(5\gamma_3^2 - 1) + (5/2)\gamma_1\gamma_3\beta_3 & (1/4)\alpha_1(5\gamma_3^2 - 1) + (5/2)\gamma_1\gamma_3\alpha_3 \\
 \sqrt{15}\gamma_1\gamma_2\gamma_3 & \sqrt{(5/2)}(\beta_1\gamma_2\gamma_3 + \beta_2\gamma_1\gamma_3 + \beta_3\gamma_1\gamma_2) & \sqrt{(5/2)}(\alpha_1\gamma_2\gamma_3 + \alpha_2\gamma_1\gamma_3 + \alpha_3\gamma_1\gamma_2) \\
 (1/2)\sqrt{15}\gamma_3(\gamma_1^2 - \gamma_2^2) & \sqrt{5/8}(\gamma_1^2\beta_3 - \gamma_2^2\beta_3 + 2\beta_1\gamma_1\gamma_3 - 2\beta_2\gamma_2\gamma_3) & \sqrt{5/8}(\gamma_1^2\alpha_3 - \gamma_2^2\alpha_3 + 2\alpha_1\gamma_1\gamma_3 - 2\alpha_2\gamma_2\gamma_3) \\
 (1/4)\sqrt{10}\gamma_2(3\gamma_1^2 - \gamma_2^2) & (1/4)\sqrt{15}\beta_2(\gamma_1^2 - \gamma_2^2) + (1/2)\sqrt{15}\gamma_1\gamma_2\beta_1 & (1/4)\sqrt{15}\alpha_2(\gamma_1^2 - \gamma_2^2) + (1/2)\sqrt{15}\gamma_1\gamma_2\alpha_1 \\
 (1/4)\sqrt{10}\gamma_1(\gamma_1^2 - 3\gamma_2^2) & (1/4)\sqrt{15}\beta_1(\gamma_1^2 - \gamma_2^2) + (1/2)\sqrt{15}\gamma_1\gamma_2\beta_2 & (1/4)\sqrt{15}\alpha_1(\gamma_1^2 - \gamma_2^2) + (1/2)\sqrt{15}\gamma_1\gamma_2\alpha_2
 \end{bmatrix} \quad (70)$$

and θ , ϕ and χ are the ligand angles. Based on this model, we can derive an additional way to compute the CFPs. Using equation 68 one can compute the V^{LF} matrix (implemented in *from_AOM_to_Vint*), and from there one can proceed with equation 59.

5.3 Crystal field rotation

Since the (complex) crystal field coefficients in the Wybourne convention can be expressed in terms of real or complex spherical harmonics, they maintain the same transformation properties. Consequently, an arbitrary rotation of the CFPs in NJA-CFS is performed using the Wigner D-matrix (\mathbf{D}), either in Euler angles (with *rota_LF* function) or in quaternions (with *rota_LF_quat* function), by applying the general formula for the spherical harmonics Y_m^l subjected to an arbitrary rotation \mathcal{R} :

$$Y_m^l = \sum_{m'=-l}^{+l} D_{m'm}^l(\mathcal{R}) Y_{m'}^l \quad (71)$$

For a 3-dimensional rotation defined by the Euler angles α , β and γ , in the z-y-z convention and active interpretation, the elements of the unitary square matrix of dimension $2k + 1$ are defined as:

$$D_{q'q}^k(\alpha, \beta, \gamma) = e^{-iq'\alpha} d_{q'q}^k(\beta) e^{-iq\gamma} \quad (72)$$

where q runs from $-k$ to k and $d_{q'q}^k(\beta)$ is the element of the Wigner's (small) d-matrix, defined as:

$$d_{q'q}^k(\beta) = [(k+q')!(k-q')!(k+q)!(k-q)!]^{1/2} \times \sum_{s=s_{\min}}^{s_{\max}} \left[\frac{(-1)^{q'+q+s} \left(\cos \frac{\beta}{2}\right)^{2k+q-q'-2s} \left(\sin \frac{\beta}{2}\right)^{q'-q+2s}}{(k+q-s)!s!(q'-q+s)!(k-q'-s)!} \right] \quad (73)$$

where the sum over s runs over such values that the factorials are nonnegative, i.e. $s_{\min} = \max(0, q - q')$ and $s_{\max} = \min(k + q, k - q')$.

The analogous of this equations, in quaternions, are computed as described by Lynden-Bell and Stone in [40]. In brief, the derivation starts from the definition of the simplest Wigner D-matrix, for $k = 1/2$. Expressing $D^{1/2}$ as:

$$\mathbf{D}^{1/2} = \begin{pmatrix} A & B \\ -B^* & A^* \end{pmatrix} \quad (74)$$

where the elements are complex functions of α , β and γ , which satisfy:

$$AA^* + BB^* = 1 \quad (75)$$

The connection among these matrix elements and the quaternion description can be found by expanding the matrix as:

$$\mathbf{D}^{1/2} = A_{\text{Re}} \mathbf{I} - B_{\text{Im}} \mathbf{i} - B_{\text{Re}} \mathbf{j} - A_{\text{Im}} \mathbf{k} \quad (76)$$

where \mathbf{I} is the 2×2 unit matrix, A_{Re} and A_{Im} (B_{Re} and B_{Im}) are the real and imaginary parts

of A (B) respectively, and the 2×2 matrices \mathbf{i} , \mathbf{j} , \mathbf{k} are:

$$\mathbf{i} = \begin{pmatrix} 0 & -i \\ -i & 0 \end{pmatrix} \quad (77)$$

$$\mathbf{j} = \begin{pmatrix} 0 & -i \\ i & 0 \end{pmatrix} \quad (78)$$

$$\mathbf{k} = \begin{pmatrix} -i & 0 \\ 0 & i \end{pmatrix} \quad (79)$$

which satisfy the multiplication rules:

$$\mathbf{i}^2 = \mathbf{j}^2 = \mathbf{k}^2 = -\mathbf{I} \quad (80)$$

The parameters of a normalized quaternion (q_0, \mathbf{q}) are related to A and B as:

$$A = q_0 - iq_3 \quad (81)$$

$$B = -q_2 - iq_1 \quad (82)$$

Given that any rotation matrix can be built from $\mathbf{D}^{1/2}$. Since an eigenfunction of angular momentum with $l = m = L$ (where $L = k$ in this case) can be constructed from $2L$ spin = $1/2$ functions, the top left element of the Wigner rotation matrix in the customary arrangement (labeling the rows and columns in the order $k, k-1, \dots, -k$) is given by:

$$D_{kk}^k = A^{2k} \quad (83)$$

and the rest of the matrix can be constructed using the operators $\hat{\mathfrak{R}}$ and $\hat{\mathfrak{B}}$ for the generation of the matrix elements on the right and beneath respectively:

$$\hat{\mathfrak{R}} D_{q'q}^k = (k(k+1) - q(q+1))^{1/2} D_{q'q-1}^k \quad (84)$$

$$\hat{\mathfrak{B}} D_{q'q}^k = (k(k+1) - q'(q'+1))^{1/2} D_{q'-1q}^k \quad (85)$$

The effect of $\hat{\mathfrak{R}}$ and $\hat{\mathfrak{B}}$ on A , A^* , B and B^* can be deduced from the matrix in 74. Additionally, since both operators are differential operators, they obey to the usual rules for differentiation of products. The following relation was also used in the matrices construction:

$$D_{-q'-q}^k = (-1)^{q'-q} (D_{q'q}^k)^* \quad (86)$$

The matrices for $k = 2$ and $k = 3$ are reported in tables 5 and 6, while matrices for $k = 4$ and $k = 6$ are saved in dedicated files, i.e. *tab_wignerDquat.txt* and *tab_wignerDquat_coeff_t.txt*, in the NJA-CFS's *tables* folder.

In order to apply this matrices to the different orders of CFPs, the real-valued Wigner coefficients are organized in a complex array (\mathbf{B}^k) :

$$\mathbf{B}^k = [\mathcal{B}_k^k, \mathcal{B}_{k-1}^k, \dots, \mathcal{B}_{-k+1}^k, \mathcal{B}_{-k}^k] \quad (87)$$

where, the complex-valued Wigner coefficients are defined as:

$$\mathcal{B}_q^k = \begin{cases} (B_{|q|}^k + iB_{|q|}^{'k}) & \text{for } q < 0 \\ B_0^k & \text{for } q = 0 \\ (-1)^{|q|}(B_{|q|}^k - iB_{|q|}^{'k}) & \text{for } q > 0 \end{cases} \quad (88)$$

The rotation matrix is applied (according to equation 71) as follows:

$$\mathbf{B}^{\text{rot},k} = \mathbf{D}^k \mathbf{B}^k \quad (89)$$

where the elements of \mathbf{D}^k are organized according to the elements of \mathbf{B}^k . The real-valued rotated CFPs B_q^k and $B_q^{'k}$ are defined as real and imaginary part respectively of the $q = -k, \dots, 0$ elements of $\mathbf{B}^{\text{rot},k}$.

Alternatively, the rotation can be performed on the $(2k+1) \times (2k+1)$ one-electron ligand field matrix, i.e. the AILFT matrix from an ORCA CASSCF calculation, with the *rotate_dicV* function. Also in this case the Wigner D-matrices are used. The elements of V^{LF} are real-valued, while the elements of \mathbf{D}^k are complex. Therefore, for the application of a rotation to V^{LF} , \mathbf{D}^k are unitary transformed, using:

$$\mathbf{U} = \begin{pmatrix} 1/\sqrt{2} & 0 & 0 & 0 & -i/\sqrt{2} \\ 0 & -1/\sqrt{2} & 0 & i/\sqrt{2} & 0 \\ 0 & 0 & 1 & 0 & 0 \\ 0 & 1/\sqrt{2} & 0 & i/\sqrt{2} & 0 \\ 1/\sqrt{2} & 0 & 0 & 0 & i/\sqrt{2} \end{pmatrix} \quad (90)$$

for $k = 2$ (for d^n configurations), and:

$$\mathbf{U} = \begin{pmatrix} -1/\sqrt{2} & 0 & 0 & 0 & 0 & 0 & i/\sqrt{2} \\ 0 & 1/\sqrt{2} & 0 & 0 & 0 & -i/\sqrt{2} & 0 \\ 0 & 0 & -1/\sqrt{2} & 0 & i/\sqrt{2} & 0 & 0 \\ 0 & 0 & 0 & 1 & 0 & 0 & 0 \\ 0 & 0 & 1/\sqrt{2} & 0 & i/\sqrt{2} & 0 & 0 \\ 0 & 1/\sqrt{2} & 0 & 0 & 0 & i/\sqrt{2} & 0 \\ 1/\sqrt{2} & 0 & 0 & 0 & 0 & 0 & i/\sqrt{2} \end{pmatrix} \quad (91)$$

for $k = 3$ (for f^n configurations), as:

$$\mathcal{D}^k = \mathbf{U}^\dagger \mathbf{D}^k \mathbf{U} \quad (92)$$

At this point, each element of V^{LF} can be converted, for an arbitrary rotation \mathcal{R} either defined in terms of Euler angles or quaternions, as follows:

$$V_{MM'}^{\text{LF,rot},k} = \sum_{m'} \sum_{m''} [\mathcal{D}_{Mm'}^k(\mathcal{R})][\mathcal{D}_{M'm''}^k(\mathcal{R})^*] V_{m'm''}^{\text{LF},k} \quad (93)$$

for M, m', M' and m'' in range $-k, \dots, k$, derived from the application of equation 71.

Table 5: Wigner D matrix in quaternions, for $k = 2$. The missing elements can be derived with equation 86. $Z = q_0^2 - q_1^2 - q_2^2 + q_3^2 = AA^* - BB^*$.

q'	q	matrix element
2	2	A^4
2	1	$2A^3B$
2	0	$\sqrt{6}A^2B^2$
1	2	$-2A^3B^*$
1	1	$A^2(2Z - 1)$
1	0	$\sqrt{6}ABZ$
0	2	$\sqrt{6}A^2B^{*2}$
0	1	$-\sqrt{6}AB^*Z$
0	0	$1/2(3Z^2 - 1)$
-1	2	$-2AB^{*3}$
-1	1	$B^{*2}(2Z + 1)$
-2	2	B^{*4}
-2	1	$-2A^*B^{*3}$

Table 6: Wigner D matrix in quaternions, for $k = 3$. The missing elements can be derived with equation 86. $Z = q_0^2 - q_1^2 - q_2^2 + q_3^2 = AA^* - BB^*$.

q'	q	matrix element
3	3	A^6
3	2	$\sqrt{6}BA^5$
3	1	$\sqrt{15}B^2A^4$
3	2	$2\sqrt{5}B^3A^3$
2	3	$-\sqrt{6}A^5B^*$
2	2	$A^4(3Z - 2)$
2	1	$(1/2)\sqrt{10}BA^3(3Z - 1)$
2	0	$\sqrt{30}B^2A^2Z$
1	3	$\sqrt{15}A^4B^{*2}$
1	2	$(1/2)\sqrt{10}A^3B^*(1 - 3Z)$
1	1	$(1/4)A^2(15Z^2 - 10Z - 1)$
1	0	$(1/2)\sqrt{3}AB(5Z^2 - 1)$
0	3	$-2\sqrt{5}A^3B^{*3}$
0	2	$\sqrt{30}A^2B^{*2}Z$
0	1	$(1/2)\sqrt{3}AB^*(1 - 5Z^2)$
0	0	$(1/2)(5Z^3 - 3Z)$
-1	3	$\sqrt{15}A^2B^{*4}$
-1	2	$-(1/2)\sqrt{10}AB^{*3}(1 + 3Z)$
-1	1	$(1/4)B^{*2}(15Z^2 + 10Z - 1)$
-2	3	$-\sqrt{6}AB^{*5}$
-2	2	$B^{*4}(3Z + 2)$
-2	1	$-(1/2)\sqrt{10}A^*B^{*3}(3Z + 1)$
-3	3	B^{*6}
-3	2	$-\sqrt{6}A^*B^{*5}$
-3	1	$\sqrt{15}B^{*4}A^{*2}$

5.4 Integrals computation

5.4.1 Electron-electron interaction

The matrix elements are J and M_J independent[21]:

$$\langle l^n \nu LS J M_J | \hat{\mathcal{H}}_{e-e} | l^n \nu' L' S' J' M_J' \rangle = \langle l^n \nu LS || \hat{\mathcal{H}}_{e-e} || l^n \nu' L' S' \rangle \delta_{J,J'} \delta_{M_J', M_J} \quad (94)$$

where the seniority quantum number ν is made explicit among the possible α and:

$$\begin{aligned} \hat{\mathcal{H}}_{e-e} &= \left(\frac{e^2}{4\pi\epsilon_0} \right) \sum_i^n \sum_{j>i}^n \frac{1}{r_{ij}} \\ &= \left(\frac{e^2}{4\pi\epsilon_0} \right) \sum_i^n \sum_{j>i}^n \sum_{k=0}^\infty [r_{>}^{-(k+1)} \cdot r_{<}^k] (C_1^{(k)} \cdot C_2^{(k)}) \end{aligned} \quad (95)$$

with $k = 0, 2, 4$ for d^n configurations and $k = 0, 2, 4, 6$ for f^n configurations.

For a two-electron system the equation is simplified in:

$$\langle l^2 \nu LS \| \hat{\mathcal{H}}_{e-e} \| l^2 \nu' L' S' \rangle = \sum_{k=0}^{\infty} F_{ll}^k [\langle l \| C^{(k)} \| l \rangle]^2 (-1)^L \begin{Bmatrix} l & l & k \\ l & l & L \end{Bmatrix} \quad (96)$$

where

$$F_{ll}^k = \int \int \frac{r_{\leq}^k}{r_{>}^{k+1}} R_l^2(r_1) R_l^2(r_2) r_1^2 r_2^2 dr_1 dr_2 \quad (97)$$

and the reduced matrix elements of the Racah operator are defined as:

$$\langle l \| C^{(k)} \| l \rangle = (-1)^l [(2l+1)(1l'+1)]^{1/2} \begin{pmatrix} l & k & l' \\ 0 & 0 & 0 \end{pmatrix}. \quad (98)$$

The 3-j symbol $\begin{pmatrix} a & b & c \\ A & B & C \end{pmatrix}$ represent the coupling of two angular momenta and can be computed using the Racah's closed form[16]:

$$\begin{aligned} \begin{pmatrix} a & b & c \\ A & B & C \end{pmatrix} &= (-1)^{a-b-C} \left[\frac{(a+b-c)!(b+c-a)!(c+a-b)!}{(a+b+c+1)!} \right] \\ &\times [(a+A)!(a-A)!(b+B)!(b-B)!(c+C)!(c-C)!]^{1/2} \\ &\times \sum_{n=n_{\min}}^{n_{\max}} (-1)^n [n!(c-b+n+A)!(c-a+n-B)!(a+b-c-n)! \\ &\times (a-n-A)!(b-n+B)!]^{-1} \end{aligned} \quad (99)$$

with:

$$\begin{aligned} n_{\min} &= \max\{0; -c+b-A; -c+a+B\} \\ n_{\max} &= \min\{a+b-c; b+B; a-A\} \end{aligned}$$

For $n > 2$ and d^n configurations, the equation is the following:

$$\langle l^2 \nu LS \| \hat{\mathcal{H}}_{e-e} \| l^2 \nu' L' S' \rangle = \delta_{L,L'} \delta_{S,S'} \sum_{k=0,2,4} F_{ll}^k \cdot c^k(l^n \nu \nu' LS) \quad (100)$$

with the angular coefficients, for $k > 0$:

$$\begin{aligned} c^k(l^n \nu \nu' LS) &= \frac{1}{2} \langle l \| C^{(k)} \| l \rangle^2 \\ &\times \left\{ \frac{1}{2L+1} \sum_{\nu'' L''} \langle l^n \nu LS \| U^{(k)} \| l^n \nu'' L'' S \rangle \langle l^n \nu' LS \| U^{(k)} \| l^n \nu'' L'' S \rangle - \frac{n}{2l+1} \delta_{\nu, \nu'} \right\} \end{aligned} \quad (101)$$

and for $k = 0$:

$$c^0(l^n \nu \nu' LS) = \frac{n(n-1)}{2} \delta_{\nu, \nu'} \quad (102)$$

The parameters that have to be passed to the program are the Slater-Condon parameters

F^k . Care must be taken for the superscript k , since:

$$\begin{aligned} F^0 &= F_0 \\ F^2 &= 49F_2 \\ F^4 &= 441F_4 \end{aligned} \tag{103}$$

for d^n and:

$$\begin{aligned} F^0 &= F_0 \\ F^2 &= 225F_2 \\ F^4 &= 1089F_4 \\ F^6 &= 184041F_6 \end{aligned} \tag{104}$$

for f^n .

The integrals computation for $n > 2$ and f^n configurations is more complex[18, 35]. Equation 100 is usually rewritten as:

$$\langle l^2 \nu LS \| \hat{\mathcal{H}}_{e-e} \| l^2 \nu' LS \rangle = \sum_{k=0,2,4,6} f_k F^k = \sum_{k=0,2,4,6} f^k F_k \tag{105}$$

with the F^k defined as 104, and $f^k = D_k f_k$, with D_k are those that relate F^k to F_k . Although the operator $(C_1^{(k)} \cdot C_2^{(k)})$ in equation 95 is scalar with respect to R_3 , it does not have the transformation properties of the group R_7 and G_2 (used for the states classification, see appendix 5.1) and therefore we have to use linear combinations of such operators. The matrix elements are re-defined as:

$$\hat{\mathcal{H}}_{e-e} = \sum_{k=0,1,2,3} e_k E^k \tag{106}$$

where e_k are related to the f^k by the expressions:

$$\begin{aligned} e_0 &= f^0 = n(n-1) \\ e_1 &= \frac{9}{7}f^0 + \frac{1}{42}f^2 + \frac{1}{77}f^4 + \frac{1}{462}f^6 \\ e_2 &= \frac{143}{42}f^2 - \frac{130}{77}f^4 + \frac{35}{462}f^6 \\ e_3 &= \frac{11}{42}f^2 + \frac{4}{77}f^4 - \frac{7}{462}f^6 \end{aligned} \tag{107}$$

whereas the E^k s are linear combinations of the F_k s:

$$\begin{aligned} E^0 &= F_0 - 10F_2 - 33F_4 - 286F_6 \\ E^1 &= \frac{70F_2 + 231F_4 + 2002F_6}{9} \\ E^2 &= \frac{F_2 - 3F_4 + 7F_6}{9} \\ E^3 &= \frac{5F_2 + 6F_4 - 91F_6}{3} \end{aligned} \tag{108}$$

5.4.2 Spin-orbit coupling

The spin-orbit coupling contribution can be simply computed as:

$$\langle l^n \nu LSJM_J | \hat{\mathcal{H}}_{so} | l^n \nu' L' S' J' M_J' \rangle = \kappa \langle l^n \nu LS || \hat{\mathcal{H}}_{so} || l^n \nu' L' S' \rangle \delta_{J,J'} \delta_{M_J', M_J} \quad (109)$$

This reduced matrix element can be computed as:

$$\begin{aligned} \langle l^n \nu LS || \hat{\mathcal{H}}_{so} || l^n \nu' L' S' \rangle &= \zeta (-1)^{J+L+S'} [l(l+1)(2l+1)]^{1/2} \\ &\times \left\{ \begin{matrix} L & L' & 1 \\ S' & S & J \end{matrix} \right\} \langle l^n \nu LS || V^{(11)} || l^n \nu' L' S' \rangle \end{aligned} \quad (110)$$

where the RMEs $\langle l^n \nu LS || V^{(11)} || l^n \nu' L' S' \rangle$ are computed according to equation 29.

In this case the parameters to be defined are the spin-orbit coupling coefficient ζ and the orbital reduction factor κ .

5.4.3 Crystal-Field splitting

In this case, even if different kind of parameters can be fed into the program (see appendix 5.2), the formalism used for the integral resolution is the Wybourne one[35]. The crystal-field integrals in the Racah's tensor operator formalism, according to equation 45, can be expressed as:

$$\begin{aligned} \langle \alpha SLJM_J | \hat{\mathcal{H}}_{cf} | \alpha' S' L' J' M_J' \rangle &= \langle \alpha SLJM_J | -e \sum_{i,k} \left[B_0^k C_0^{(k)}(i) + \right. \\ &\left. + \sum_{q=1}^k (B_q^k (C_{-q}^{(k)}(i) + (-1)^q C_q^{(k)}(i)) + B_q'^k (C_{-q}^{(k)}(i) - (-1)^q C_q^{(k)}(i))) \right] | \alpha' S' L' J' M_J' \rangle \end{aligned} \quad (111)$$

where α represent the additional set of quantum numbers used and k has values 2 and 4 for d^n and 2, 4 and 6 for f^n configurations. The B_q^k and $B_q'^k$ are the so-called crystal field coefficients (CFCs), that can be converted in the corresponding crystal field parameters B_q^k and $B_q'^k$, when solving the integral, by bringing them outside the matrix element, together with the radial parts of the wavefunctions $R_{nl}(r)$. In this way, the tensor operators are left acting solely on the angular part. The use of the same notation for both coefficients and parameters is confusing but, unfortunately, it is what is done in literature.

The CFPs are defined as:

$$B_0^k = \int_{r=0}^{\infty} R_{nl}^2(r) r^k dr \sqrt{\frac{4\pi}{2k+1}} \sum_L Z_{k0}^c(\theta_L) \frac{Z_L e^2}{R_L^{k+1}} \quad (112)$$

$$B_q^k = \int_{r=0}^{\infty} R_{nl}^2(r) r^k dr \sqrt{\frac{4\pi}{2k+1}} \frac{1}{\sqrt{2}} \sum_L Z_{kq}^c(\theta_L, \phi_L) \frac{Z_L e^2}{R_L^{k+1}} \quad (113)$$

$$B_q'^k = \int_{r=0}^{\infty} R_{nl}^2(r) r^k dr \sqrt{\frac{4\pi}{2k+1}} \frac{1}{\sqrt{2}} \sum_L Z_{kq}^s(\theta_L, \phi_L) \frac{Z_L e^2}{R_L^{k+1}} \quad (114)$$

where the sum over L runs over the ligands, represented in the point charge model. The $Z_{kq}(\theta_L, \phi_L)$ are the tesseral harmonics of degree k and order q , defined at the polar (colati-

tudinal) angle θ_L and azimuthal (longitudinal) angle ϕ_L at the L -th ligand position. The very same equations can be expressed in term of the (complex) spherical harmonics Y_k^q . Z_L and R_L are the point charge and the distance from the metal ion of the L -th ligand respectively. For a continuous distribution of charges, an integral over the charge density can be used. The integrals over the radial part, dedicated parametrizations are available, generally represented as $\langle r^k \rangle$.

The angular parts (containing the electron coordinates) of the matrix elements expressed in equation 111 can be calculated using the tensor operators properties, by rewriting the the summation over i in terms of unit tensor U_q^k :

$$\langle \alpha SLJM_J | \sum_i C_q^{(k)}(i) | \alpha' S' L' J' M'_J \rangle = \langle \alpha SLJM_J | U_q^{(k)} | \alpha' S' L' J' M'_J \rangle \langle l \| C^{(k)} \| l' \rangle \quad (115)$$

The U_q^k matrix elements are diagonal in S .

Applying the Wigner-Eckart theorem, and solving the RME of $C^{(k)}$, equation 115 becomes:

$$\begin{aligned} & \langle \alpha SLJM_J | \sum_i C_q^{(k)}(i) | \alpha' S' L' J' M'_J \rangle = \\ & (-1)^{2J-M_J+S+L'+k+l} [(2J+1)(2J'+1)]^{1/2} [(2l+1)(2l'+1)]^{1/2} \\ & \times \begin{pmatrix} l & k & l' \\ 0 & 0 & 0 \end{pmatrix} \begin{pmatrix} J & k & J' \\ -M_J & q & M'_J \end{pmatrix} \begin{Bmatrix} J & J' & k \\ L & L' & S \end{Bmatrix} \langle \alpha SL \| U^{(k)} \| \alpha' S' L' \rangle \end{aligned} \quad (116)$$

The RMEs $\langle \alpha SL \| U^{(k)} \| \alpha' S' L' \rangle$ can be solved using the cfps and the equations in appendix 5.1.1.

5.4.4 Zeeman interaction

The contribution from the Zeeman interaction with the magnetic field vector is expressed as:

$$\begin{aligned} & \langle l^n \nu LSJM_J | \hat{\mathcal{H}}_Z | l^n \nu' L' S' J' M'_J \rangle = \mu_B \hbar^{-1} \sum_{q=-1}^1 (-1)^q B_{-q}^{(1)} \\ & \times (-1)^{J-M} \begin{pmatrix} J & 1 & J' \\ -M_J & q & M'_J \end{pmatrix} \langle l^n \nu LSJ \| (\kappa L^{(1)} + g_e S^{(1)}) \| l^n \nu' L' S' J' \rangle \end{aligned} \quad (117)$$

where the RMEs are computed as:

$$\begin{aligned} & \langle l^n \nu LSJ \| (\kappa L^{(1)} + g_e S^{(1)}) \| l^n \nu' L' S' J' \rangle = \delta_{L,L'} \delta_{S,S'} \hbar [(2J+1)(2J'+1)]^{1/2} \\ & \times \left\{ k [L(L+1)(2L+1)]^{1/2} (-1)^{L+S+J+1} \begin{Bmatrix} J' & J & 1 \\ L & L & S \end{Bmatrix} + g_e [S(S+1)(2S+1)]^{1/2} \right. \\ & \left. \times (-1)^{L+L'+2S'+J+J'} (-1)^{L+S+J+1} \begin{Bmatrix} J' & J & 1 \\ S & S & L \end{Bmatrix} \right\} \end{aligned} \quad (118)$$

with:

$$\begin{aligned}B_{+1}^{(1)} &= -\frac{1}{\sqrt{2}}(B_x + \mathrm{i}B_y) \\B_{-1}^{(1)} &= \frac{1}{\sqrt{2}}(B_x - \mathrm{i}B_y) \\B_0^{(1)} &= B_z\end{aligned}$$

In this case the parameters needed are the magnetic field vector $\mathbf{B} = [B_x, B_y, B_z]$ and the orbital reduction factor κ .

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