

Symmetry and Correlation in Multiple Self-Consistent-Field Solutions of Transition-Metal Complexes

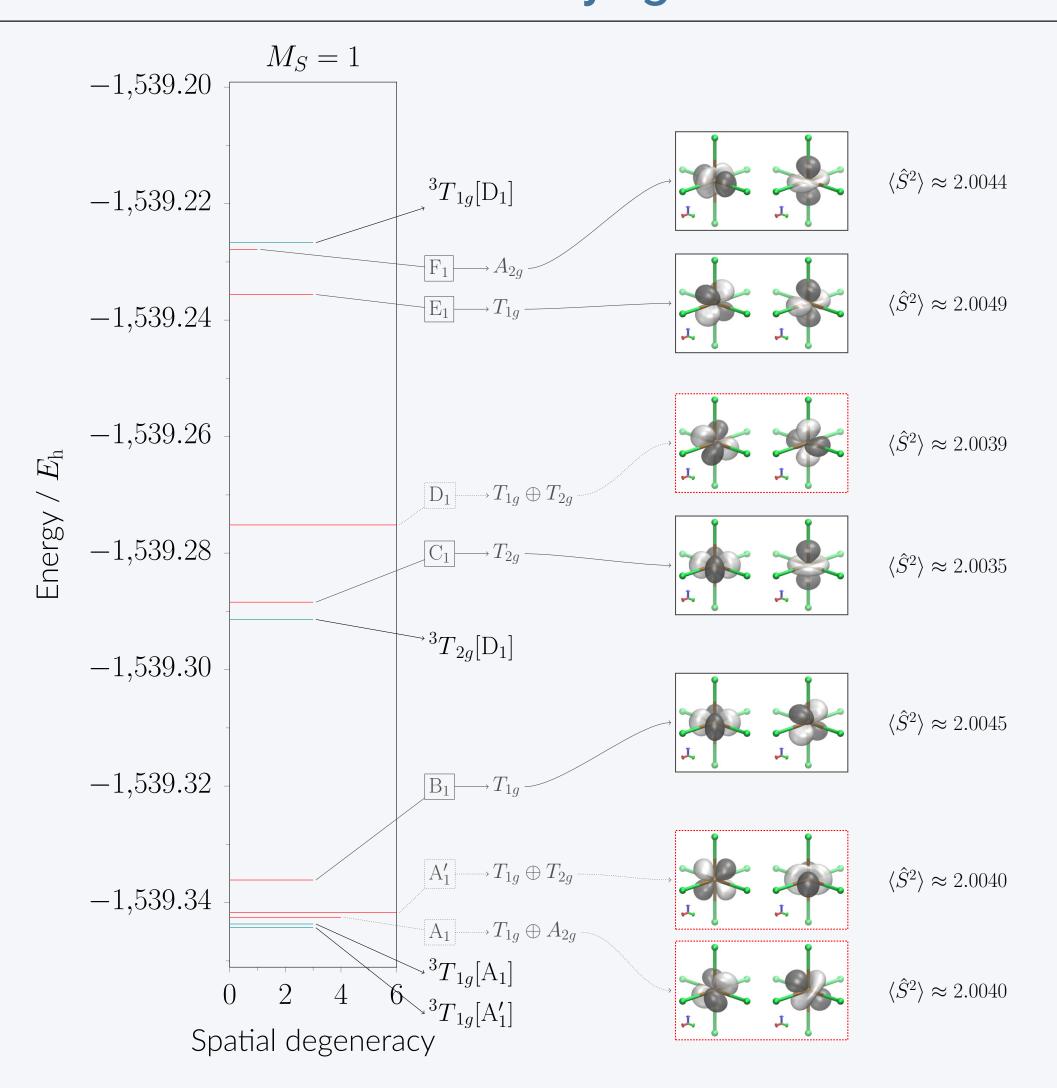
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Low-Lying UHF Solutions and NOCI Wavefunctions of Model Octahedral [VF₆]³⁻



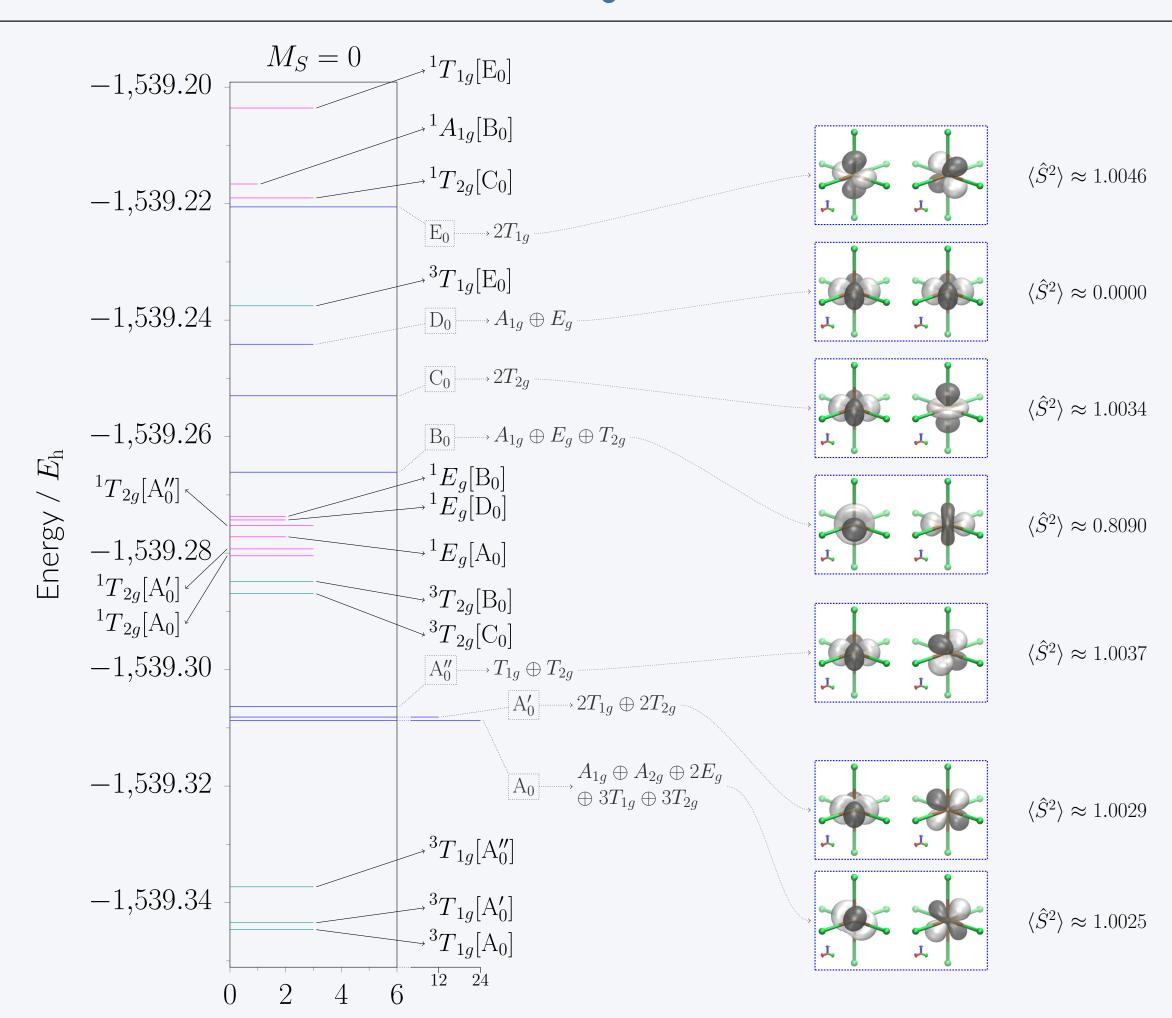
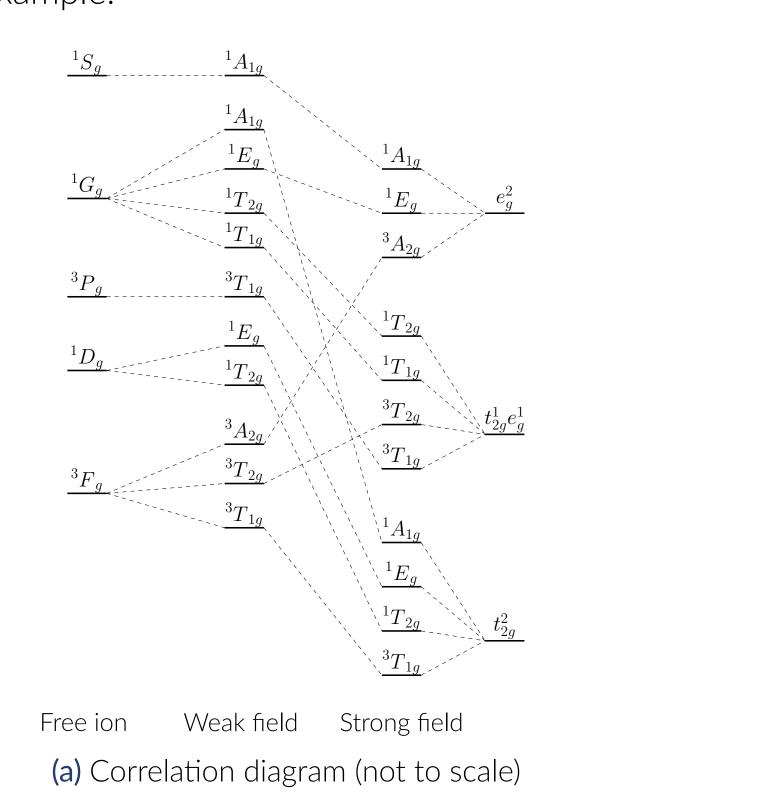


Figure 1. Energy and symmetry of low-lying UHF solutions and NOCI wavefunctions constructed from them in octahedral $[VF_6]^{3-}$. $[S_{M_S}]$: symmetry-conserved UHF set S with \hat{S}_z eigenvalue M_S . $[S_{M_S}]$: spatial or spin symmetry-broken UHF set S with \hat{S}_z eigenvalue M_S . $[S_{M_S}]$: a specific NOCI set of symmetry Γ constructed from UHF set S with \hat{S}_z eigenvalue M_S .

Introduction

Transition-metal complexes are **strongly correlated** as they have many low-energy electronic states that exhibit **high degrees of degeneracy**. Figure 2 gives such states for octahedral d^2 as an example.



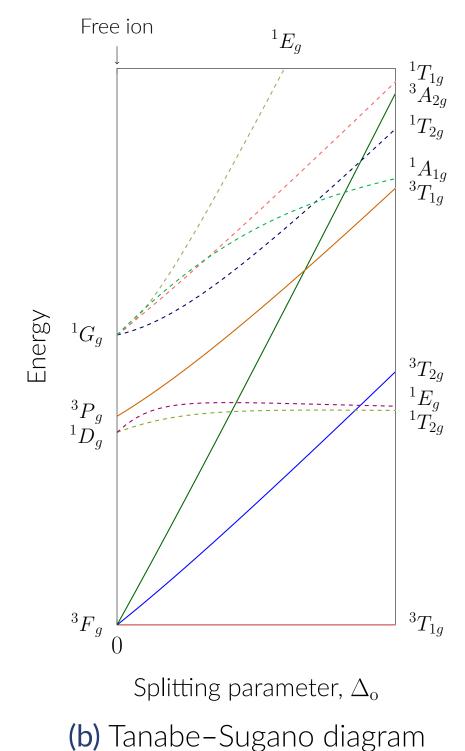


Figure 2. All electronic terms of a true d^2 system in an octahedral field.

The non-linear HF equations for these complexes therefore admit multiple low-lying solutions that are degenerate or nearly degenerate.

We have located these solutions using SCF metadynamics and investigated their symmetry properties in a model octahedral d^2 system, $[VF_6]^{3-}$ (Figure 1).

Symmetry Breaking in HF

Degenerate eigenfunctions of the spinless Hamiltonian *must* transform as a single irreducible representation (irrep) of the underlying point group \mathcal{B} , the spin rotation group $\mathrm{SU}(2)$, or the time reversal group \mathcal{T} .

HF wavefunctions do not have to obey this due to their approximate nature. Thus, consider a set of degenerate HF solutions $S = \{ {}^w\Psi \mid w = 1, 2, \ldots \}$ and a particular group \mathcal{G} :

- if S spans a single irrep in \mathcal{G} , then S is symmetry-conserved in \mathcal{G} ;
- if S spans multiple irreps in \mathcal{G} , then S is symmetry-broken in \mathcal{G} .

HF solutions break symmetry to become lower in energy and possibly recover some electron correlation. Restoring symmetry of symmetry-broken HF solutions allows us to form physically meaningful wavefunctions while incorporating said correlation.

Restoring Symmetry: Non-Orthogonal CI (NOCI)

For a complete symmetry-broken set $S = \{^w \Psi \mid w = 1, 2, \ldots\}$, solving the generalised eigenvalue equation

 $m{H}m{A} = m{S}m{A}m{E}$ where $(m{H})_{wx} = \langle ^w\Psi | \hat{\mathscr{H}} | ^x\Psi \rangle$ and $(m{S})_{wx} = \langle ^w\Psi | ^x\Psi \rangle$ gives coefficients A_{wm} such that the NOCI wavefunctions

$$^{m}\Phi = \sum^{w} \Psi A_{wm}$$

conserve symmetry and can be used to approximate corresponding electronic terms.

UHF vs. NOCI: Jahn-Teller Distortion

Consider the $T_{1q} \otimes e_q$ Jahn-Teller distortion in octaheral $[VF_6]^{3-}$. Figure 3 shows that:

• the UHF A_1 and A_1' solutions fail to exhibit the expected energy minima;

Spatial & time-reversal degeneracy

• the ${}^3T_{1g}[A_1 \oplus A_1']$ **NOCI wavefunctions** do show the expected stabilisation and give the correct degeneracy splitting upon symmetry descent.

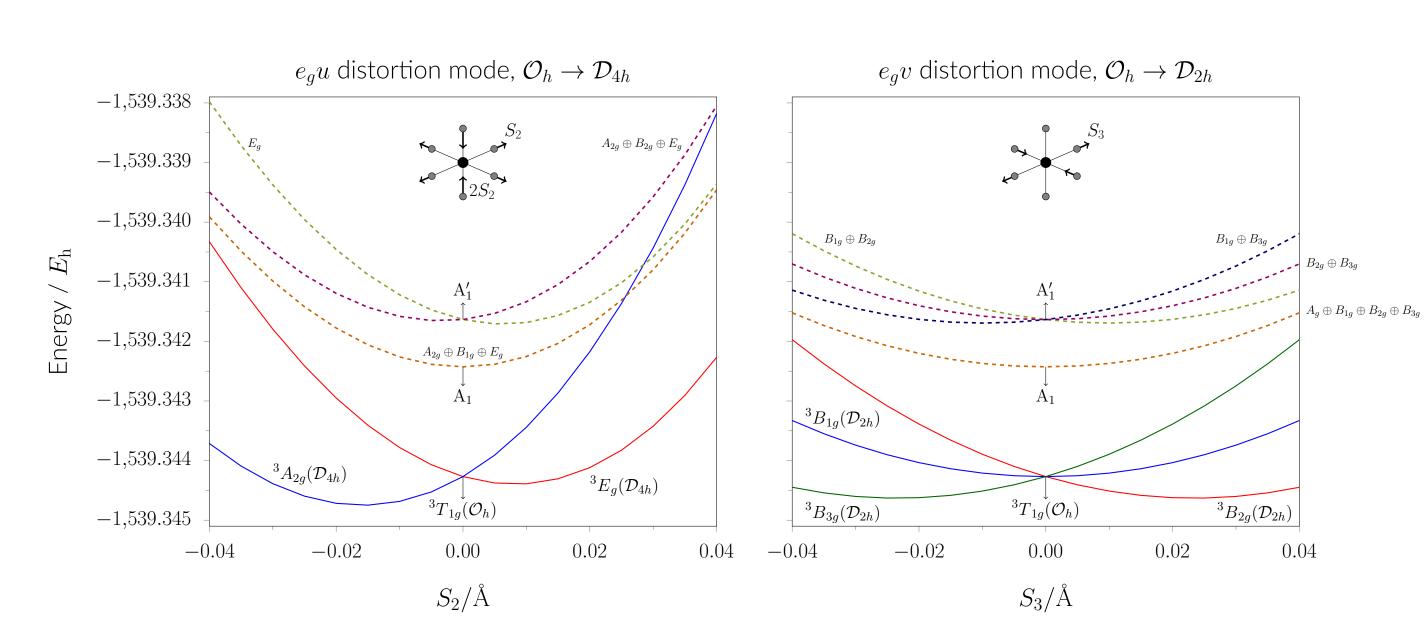


Figure 3. State energy in the $T_{1g} \otimes e_g$ Jahn-Teller distortion of octahedral $[VF_6]^{3-}$. Dashed curves: symmetry-broken UHF A_1 or A_1' solutions. Solid curves: ${}^3T_{1g}[A_1 \oplus A_1']$ NOCI wavefunctions.

Solution Topology: Euclidean Realisation of State Distances

By realising the distance matrices between symmetry-broken HF solutions to give **polytopes** showing their **arrangements in Euclidean space** (Figure 4), we hope to gain insight into the nature of their symmetry breaking.

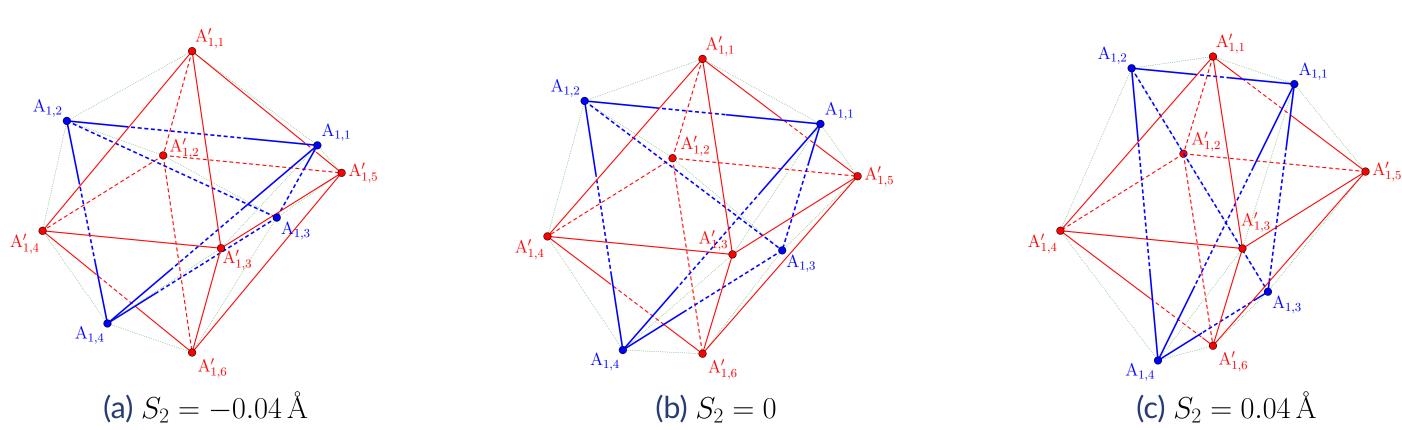


Figure 4. Three-dimensional projections of polytopes of A_1 and A_1' solutions along the e_gu distortion.

In a Nutshell

- There exist many low-lying HF solutions in transition-metal complexes.
- Some solutions break symmetry, spin or spatial or both.
- NOCI can restore symmetry to give proper physical behaviours, e.g., vibronic coupling.
- There are **clear patterns** in symmetry breaking that need further understanding.

References

- Thom, A. J. W. & Head-Gordon, M. *Physical Review Letters* **101,** 193001 (November 2008).
- 2. Thom, A. J. W. & Head-Gordon, M. *The Journal of Chemical Physics* **131**, 124113 (September 2009).
- 3. Huynh, B. C. & Thom, A. J. W. (Manuscript in preparation).