

General Perturb-Then-Diagonalize Model for the Vibrational Frequencies and Intensities of Molecules Belonging to Abelian and Non-Abelian Symmetry Groups

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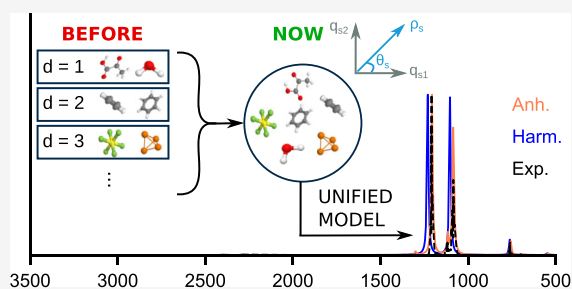


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ABSTRACT: In this paper, we show that the standard second-order vibrational perturbation theory (VPT2) for Abelian groups can be used also for non-Abelian groups without employing specific equations for two- or threefold degenerate vibrations but rather handling in the proper way all the degeneracy issues and deriving the peculiar spectroscopic signatures of non-Abelian groups (e.g., *l*-doubling) by a posteriori transformations of the eigenfunctions. Comparison with the results of previous conventional implementations shows a perfect agreement for the vibrational energies of linear and symmetric tops, thus paving the route to the transparent extension of the equations already available for asymmetric tops to the energies of spherical tops and the infrared and Raman intensities of molecules belonging to non-Abelian symmetry groups. The whole procedure has been implemented in our general engine for vibro-rotational computations beyond the rigid rotor/harmonic oscillator model and has been validated on a number of test cases.



1. INTRODUCTION

The reliability of quantum chemical (QC) models to support experimental findings is related from one side to their accuracy and from the other side to their feasibility, robustness, and ease of use.^{1,2} Concerning the accuracy, electronic structure computations of energies, geometries, and force fields are nowadays able to rival high-resolution spectroscopy for small systems and to help assignments and interpretation of all kinds of spectra for larger molecular systems, provided that nuclear motions and environmental effects are taken into proper account.^{3–8} In the present contribution, we will be concerned with molecular vibrations, which are directly sampled by different conventional [infrared (IR), Raman] and chiral (VCD, ROA) spectroscopies,^{2,9} but tune also the outcomes of other spectroscopies (e.g., distortion constants in microwave spectroscopy⁴ or line shapes in electronic spectroscopies¹⁰). Until quite recently, except for exceedingly small systems, the rigid rotor/harmonic oscillator model was nearly exclusively employed to describe molecular vibrations. However, the neglect of anharmonicity and ro-vibrational couplings can introduce significant errors, sometimes leading to even qualitatively wrong interpretations of experimental data.

Among the different approaches available to go beyond the rigid rotor/harmonic oscillator approximation,^{11–35} those based on perturbation theory applied to the expansion of the nuclear Hamiltonian in the power series of products of vibrational and rotational operators (hereafter referred to as vibrational perturbation theory, VPT) are particularly appealing for their remarkable cost/performance ratio, at least for semi-rigid

molecular systems. Moreover, some formulations of VPT, such as the Van Vleck contact transformation method,³⁶ fully justify a generalized model (GVPT2),^{37,38} allowing to couple the advantages of perturbative (for weakly coupled modes) and variational (for strongly coupled modes) treatments in a well-sound and robust framework. Actually, the GVPT2 approach belongs to the class of perturb-then-diagonalize many-body models, which, although less widely used than diagonalize-then-perturb models, have in the present context some appealing advantages from both conceptual and implementative points of view. Implementations of VPT2 approaches in general-purpose QC software are now quite widespread,^{39–50} although fully automatic and robust implementations of GVPT2 are less common. The situation is different for intensities of IR, Raman, VCD, and ROA spectra, which need to account for both mechanical and electrical/magnetic anharmonicity. To the best of our knowledge, the only general platform allowing GVPT2 computations for all the spectroscopic techniques mentioned above is the one implemented by some of the present authors in the Gaussian package.⁵¹ Extension of perturbative/variational procedures to the treatment of flexible systems is underway along different avenues including approaches based on reaction

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paths or surfaces following the decoupling of one or two large-amplitude motions from a bath of small-amplitude modes and/or the replacement of Cartesian coordinates by generalized internal coordinates.^{52–55}

Here, we tackle a different problem, related to the treatment of molecular systems with non-Abelian point group symmetries, namely, linear, symmetric, and spherical tops. As a matter of fact, a significant ensemble of molecular systems, ranging from small to large sizes and of interest in various research fields, belong to these classes, including, for instance, organic and organometallic compounds like coronene and ferrocene^{30,56–58} or acetylene derivatives.^{59–70} The presence of degenerate modes raises multiple practical and theoretical issues, which have been only marginally addressed until now. A possible workaround is to lift the degeneracies by reducing the symmetry of the molecular system to the closest Abelian group symmetry,⁴² but this can lower the accuracy of the results. The rotational problem is actually simpler for non-Abelian groups because the rigid rotor approximation leads to analytical solutions, but a proper account of the degeneracy for the vibrational problem requires a careful check of the relative orientations of the degenerate modes to ensure that the anharmonic force fields and property surfaces are correctly built and an alternative derivation able to account for the couplings involving the degenerate modes. In a previous work,⁷¹ we have presented a complete framework implementing different equations for the vibrational frequencies based on the symmetry for linear and symmetric tops and taking into proper account both intrinsic and accidental degeneracies, leading to additional terms in the Hamiltonian or to singularities in the perturbative expansion, respectively. However, the current situation is unsatisfactory at several levels. First, the formulation derived from the work of Plíva⁷² cannot be extended straightforwardly to spherical tops, and to the best of our knowledge, no complete and correct derivation has been proposed. Second, the current implementation is particularly intricate due to the constant case switch depending on the symmetry and degeneracy of the modes in the calculation of the quantities of interest for the vibrational energies. Finally, it requires the use of complex algebra in the variational treatment, which leads to complex eigenvectors of the variational matrix, and thus the transition moments, even if the final intensities remain, of course, real.

Based on these premises, a unified treatment of Abelian and non-Abelian symmetries would be more efficient and general, provided that the results can be easily transformed to the more standard representation for compatibility and/or interpretative purposes. Furthermore, the simplicity of the new approach could allow a more straightforward implementation of new strategies based on VPT2⁷³ not only for linear and symmetric tops but also for spherical tops. Finally, a general and robust framework can be set for the calculation of both IR and Raman intensities for all point groups without any need of complex algebra. This has convinced us to follow a different route, that is, to employ the asymmetric-top formulation also for the other cases, handling in the proper way all the degeneracy issues and deriving the customary spectroscopic signatures of non-Abelian groups (e.g., *l*-type doubling) by a posteriori transformations of the eigenvectors. As will be shown in the following, the results for frequencies are exactly the same as those delivered by our previous conventional implementation (including resonance contributions), but we are now able to compute also intensities for both IR and Raman spectra.

The paper is organized as follows: In the **Theory** section, we start with deriving energy expressions for linear and symmetric tops, including resonance-free expressions for the zero-point energy, and then move to a general treatment of resonances and to intensities for all vibrational spectroscopies. The Results and Discussion section is organized in the same way but also considers different levels of electronic structure theory, starting from Hartree–Fock (HF) and second-order Møller–Plesset perturbation theory (MP2), which do not involve any underlying noise connected to numerical integration, and then moving to methods rooted in the density functional theory (DFT) up to double hybrids. The main results, remaining challenges, and perspectives are shortly outlined in the concluding section.

2. THEORY

2.1. Framework. In order to set up the framework for our discussion, let us consider a system of N vibrational normal modes with a non-Abelian symmetry. These modes will generally be identified by the indexes i, j, k, l . Where relevant, and except if specified otherwise, different sets of indexes will be used to distinguish degenerate modes (s, t, u , followed by a subscript number for each mode) from the non-degenerate ones (m, n, o). As an example, modes s_1 and s_2 are two degenerate modes with the same harmonic wavenumber ω_s , while m and n are two non-degenerate modes.

The formalism introduced by Plíva for symmetric and linear tops⁷² relies on a special set of coordinates for degenerate modes based on a complex combination

$$q_s^\pm = q_{s_1} \pm iq_{s_2} \quad (1)$$

where q_i represents the dimensionless normal coordinate associated to mode i . This form can actually be directly related to the definition of the harmonic vibrational wave function ψ . Thus, it is convenient to first recall its representation for a system with N' non-degenerate modes and N'' sets of degenerate modes ($N'' = 0$ for asymmetric tops). In the canonical representation (C), ψ , associated to the vibrational state $|v\rangle$, is given as a product of one-dimensional functions

$$\psi_v^C(\mathbf{q}) = \prod_{i=1}^{N'} \varphi_{v_i}^C(q_i) = \prod_{m=1}^{N'} \varphi_{v_m}^C(q_m) \prod_{s=1}^{N''} \prod_{\sigma=1}^{N_s''} \varphi_{v_{s\sigma}}^C(q_{s\sigma}) \quad (2)$$

where N_s'' is the degeneracy order associated to degenerate mode s , v_i is the number of vibrational quanta corresponding to mode i , and $\varphi_{v_i}^C(q_i)$ are the well-known one-dimensional harmonic oscillator wave functions. To make the discussion simpler, we will consider only molecules with at most doubly degenerate modes, that is, linear and symmetric tops. Extension to spherical tops is deferred to a dedicated section. Equation 2 can thus be written more explicitly as

$$\psi_v^C(\mathbf{q}) = \prod_{m=1}^{N'} \varphi_{v_m}^C(q_m) \prod_{s=1}^{N''} \varphi_{v_{s_1}}^C(q_{s_1}) \varphi_{v_{s_2}}^C(q_{s_2}) \quad (3)$$

An alternative way to treat the degenerate coordinates is through the polar representation,²⁵ in which the wave function, namely, $\psi_v^P(\mathbf{q})$, assumes the following form:

$$\psi_v^P(\mathbf{q}) = \prod_{m=1}^{N'} \varphi_{v_m}^C(q_m) \prod_{s=1}^{N''} \varphi_{v_s, l_s}^P(\rho_s, \theta_s) \quad (4)$$

where ν_s and $l_s = \{-\nu_s, -\nu_s + 2, \dots, \nu_s - 2, \nu_s\}$ are respectively the principal and angular quantum numbers, while $\varphi_{\nu_s, l_s}^p(\rho_s, \theta_s)$ is the harmonic wave function of the two-dimensional isotropic harmonic oscillator Hamiltonian expressed in terms of the polar coordinates ρ_s and θ_s arising from a pair of degenerate modes.

It is worth recalling that both representations, canonical and polar, have the same eigenvalues and hence vibrational energies but different eigenstates. From a theoretical perspective, the polar representation is preferable since it leads to an explicit quantization of the vibrational angular momentum stemming from each pair of degenerate vibrations. The formulation proposed by Pliva takes these properties into account through a transformation of the degenerate normal coordinates and the associated force field.^{72,74}

Having confirmed that the presence of degenerate modes does not preclude the development of VPT2 equations for symmetric and linear tops in the canonical representation (see Appendix A), the main task is to define a suitable transformation between the states obtained in each representation. At the harmonic level, it is possible to build a linear transformation between sets of degenerate states³⁸

$$|\psi_\nu^P\rangle = \mathbf{P}_\nu^T |\psi_\nu^C\rangle \quad (5)$$

where $|\psi_\nu^P\rangle$ and $|\psi_\nu^C\rangle$ are the column vectors containing the states with the same harmonic energy, which implies in practice that ν_m and $\nu_s = \nu_{s_1} + \nu_{s_2}$ are constant, and \mathbf{P}_ν is a unitary matrix connecting the two sets of states. The complete set of rotation matrices required for the conversion of fundamentals, first overtones, and binary (1 + 1) combinations from the vibrational ground state is reported in Section S1 of the Supporting Information. Furthermore, following the recent extension of our computational framework to the inclusion of three-quanta states,⁷⁵ the full set of the corresponding rotations is reported for the sake of completeness.

Therefore, a unified framework can be set up, in which calculations are run in two steps:

1. anharmonic calculations in the canonical framework;
2. if degenerate modes are present, application of the necessary rotations to switch to the polar representation.

2.2. Vibrational Energies. Full derivations for the VPT2 energies of asymmetric^{11,32,42} and symmetric or linear^{71,72,74} tops have been reported elsewhere. Here, we will focus on the novel aspects using refs^{48,71,76} as a basis. Based on Pliva's formalism, the vibrational energy for symmetric and linear tops can be written

$$\begin{aligned} \varepsilon_{\nu, l}^P &= \varepsilon_0^P + \sum_{i=1}^N \omega_i \nu_i + \sum_{i=1}^N \sum_{j=i}^N \chi_{ij}^P \left(\nu_i \nu_j + \frac{\nu_i}{2} d_j + \frac{\nu_j}{2} d_i \right) \\ &+ \sum_{s=1}^{N''} \sum_{t=s}^{N''} g_{st}^P I_{st} \end{aligned} \quad (6)$$

where ε_0^P is the zero-point vibrational energy (ZPVE) in the polar representation, d_i is the degeneration of the i th normal mode, and χ^P and \mathbf{g} are respectively the anharmonic constant matrix in polar representation and the matrix containing the anharmonic contributions from the angular momenta, given in ref 71 and reported in Section S2 of the Supporting Information.

Let us consider a set of degenerate harmonic states in the canonical ($|\psi_\nu^C\rangle$) and polar ($|\psi_\nu^P\rangle$) representations. The associated blocks of the contact-transformed vibrational

Hamiltonian are respectively $\tilde{\mathbf{H}}_{\nu, \nu}^C$ and $\tilde{\mathbf{H}}_{\nu, \nu}^P$ where the subscript “ ν ” indicates that only matrix elements between states with the same principal quantum numbers ν are included in the block. These blocks will be simply referred to as diagonal blocks in the following. By construction, only the diagonal elements of the corresponding matrices contribute to the anharmonic energies. Using eq 5, the following identity can be written

$$\tilde{\mathbf{H}}_{\nu, \nu}^P = \mathbf{P}_\nu^T \tilde{\mathbf{H}}_{\nu, \nu}^C \mathbf{P}_\nu \quad (7)$$

Since the trace of a matrix is invariant under similarity transformation, we have

$$\sum_{s=1}^{N''} \sum_{l_s} \varepsilon_{\nu, l_s}^P = \sum_{s=1}^{N''} \sum_{\sigma=1}^{N''} \sum_{\nu_s=0}^{\nu_s} \varepsilon_{\nu}^C; \quad \nu_s = \nu_s - \sum_{\alpha=1}^{\sigma-1} \nu_{s_\alpha} \quad (8)$$

Equation 7 can thus be systematically used to obtain the energies in the polar representation, starting from both energies and resonances (see later) evaluated within the canonical representation.

After demonstrating that the expression of the resonance-free ZPVE for asymmetric tops^{77–79}

$$\begin{aligned} \varepsilon_0^C &= \sum_{i=1}^N \frac{\omega_i}{2} + \sum_{i=1}^N \sum_{j=1}^N \frac{f_{ijj}}{32} \\ &- \sum_{i=1}^N \sum_{j=1}^N \sum_{k=1}^N \left[\frac{f_{iik} f_{ijk}}{32 \omega_k} + \frac{f_{ijk}^2}{48(\omega_i + \omega_j + \omega_k)} \right] \\ &- \sum_{\tau} \frac{B_{\tau}^{\text{eq}}}{4} \left[1 - \sum_{i=1}^{N-1} \sum_{j=i+1}^N \{ \zeta_{ij, \tau} \}^2 \frac{(\omega_i - \omega_j)^2}{\omega_i \omega_j} \right] \end{aligned} \quad (9)$$

can also be used for linear and symmetric tops (see Section S3 of the Supporting Information for details and notation), that is,

$$\varepsilon_0^C = \varepsilon_0^P \quad (10)$$

Equation 7 can be applied to calculate the energies in the polar representation for one- and two-quanta states involving at least one degenerate mode. Combining the rotation matrices reported in Section S1 of the Supporting Information with the transformation of the Hamiltonian given in eq 7, it is straightforward to prove that the anharmonic fundamental energies do not vary under the change of representation, a property true for any excited state involving only one degenerate mode excited with a single quantum ($\nu_s = 1$), for instance, $|\psi_{1_m 1_j, \pm 1_s}^P\rangle$, $|\psi_{1_m 1_n 1_l, \pm 1_s}^P\rangle$, $|\psi_{2_m 1_l, \pm 1_s}^P\rangle$, and so on.

Following the procedure outlined for the fundamental states, the rotation-based framework yields the expressions for the polar energies of the first overtone associated to a degenerate mode s

$$\varepsilon_{2_s, \pm 2_s}^P = \frac{1}{4} \left[\varepsilon_{2_s} + \varepsilon_{1_{s_1} 1_{s_2}}^C - 2 \left\langle \psi_{2_{s_1}}^C \left| \tilde{\mathcal{H}} \right| \psi_{2_{s_2}}^C \right\rangle \right] \quad (11a)$$

$$\varepsilon_{2_s, 0}^P = \frac{1}{2} \left[\varepsilon_{2_s} - \varepsilon_{1_{s_1} 1_{s_2}}^C + 2 \left\langle \psi_{2_{s_1}}^C \left| \tilde{\mathcal{H}} \right| \psi_{2_{s_2}}^C \right\rangle \right] \quad (11b)$$

where $\tilde{\mathcal{H}}$ is the contact-transformed Hamiltonian,²⁴ and

$$\varepsilon_{2_s} = \varepsilon_{2_{s_1}}^C + \varepsilon_{2_{s_2}}^C + \varepsilon_{1_{s_1} 1_{s_2}}^C \quad (12)$$

as well as those of the binary (1+1) combination bands involving two degenerate modes, namely, s and t

$$\varepsilon_{1,1,\pm 1,\pm 1}^P = \frac{1}{4} \left[\varepsilon_{st} - 2 \left\langle \psi_{1_1 1_1}^C \left| \tilde{\mathcal{H}} \right| \psi_{1_2 1_2}^C \right\rangle + 2 \left\langle \psi_{1_1 1_2}^C \left| \tilde{\mathcal{H}} \right| \psi_{1_2 1_1}^C \right\rangle \right] \quad (13a)$$

$$\varepsilon_{1,1,\pm 1,\mp 1}^P = \frac{1}{4} \left[\varepsilon_{st} + 2 \left\langle \psi_{1_1 1_1}^C \left| \tilde{\mathcal{H}} \right| \psi_{1_2 1_2}^C \right\rangle - 2 \left\langle \psi_{1_1 1_2}^C \left| \tilde{\mathcal{H}} \right| \psi_{1_2 1_1}^C \right\rangle \right] \quad (13b)$$

where

$$\varepsilon_{st} = \varepsilon_{1_1 1_1}^C + \varepsilon_{1_2 1_2}^C + \varepsilon_{1_1 1_2}^C + \varepsilon_{1_2 1_1}^C \quad (14)$$

Contrary to fundamental bands, the energy terms in polar representation cannot be directly obtained from the canonical representation but require additional, off-diagonal terms, collectively called Darling–Dennison coupling terms, which will be illustrated later. Finally, three-quanta transitions are treated in the Section S1 of the [Supporting Information](#).

2.3. Resonances and the Variational Correction. In the following, we will consider three models for the calculation of VPT2 energies. In the pure VPT2 approach (simply named VPT2), resonances are ignored, in the sense that all terms involved in the calculation of the quantities of interest are systematically included. As a result, this approach has a tendency to break down very quickly with the system size, often leading to unphysical results. An improvement is offered by the deperturbed VPT2 (DVPT2) scheme, where resonances are identified, usually through a multi-step procedure,^{42,76,79,80} and the related terms are discarded. Strong discrepancies are prevented, but the selective removal of terms can result in an unbalanced account of the anharmonic contributions, with varying intensity. In the most refined version, the generalized VPT2 (GVPT2) scheme,^{32,42,76,81} the resonant terms from DVPT2 are introduced back through an additional variational step, often together with other terms, broadly referred to as Darling–Dennison interaction terms.⁸² For this reason, the latter should be systematically considered for spectroscopic applications like those here. The general construction and use of polyads will be reviewed together with GVPT2.

2.3.1. Fermi Resonances and DVPT2 Energies. At the VPT2 level, the energies in the polar representation can be systematically obtained through linear combinations of canonical energies and Darling–Dennison resonances. Let us first discuss the extension of the theoretical framework developed so far to the DVPT2 scheme, where each potentially resonant term in the χ matrix is analyzed by applying specific criteria, and those which are identified as resonant are discarded from the calculation. In the present work, the identification of Fermi resonances is done through a two-step procedure, which first considers the energetic proximity of the interacting states, namely, $\omega_i \approx 2\omega_j$ (type I) and $\omega_i \approx \omega_j + \omega_k$ (type II), and then the magnitude of the term, using Martin's test⁸⁰ to estimate the deviation of the term from the variational energy of a model, ad hoc system. Currently, DVPT2 can be employed for both canonical and polar representations, the only difference lying in the definition of the χ matrix and the construction of the \mathbf{g} matrix in the second case. However, it can be demonstrated that the formalism described in [Section 2.2](#) can be easily extended to DPVT2 calculations since [eq 7](#) is valid for both resonant and non-resonant terms separately

$$\tilde{\mathbf{H}}_{\nu,\nu'}^{P,DVPT2} = \mathbf{P}_\nu^\dagger \tilde{\mathbf{H}}_{\nu,\nu'}^{C,DVPT2} \mathbf{P}_\nu \quad (15)$$

At this point, the transformation becomes similar to the VPT2 case presented above. An analogous procedure can be employed out to evaluate the χ^P and \mathbf{g} matrices.

In practice, subtle differences could be observed because of the numerical parameters and tests used to define the resonances. However, for an equivalent set of resonances, the results obtained through eqs 6 and 15 (χ^P and \mathbf{g} matrices being deprived of Fermi resonances) will converge.

2.3.2. Variational Correction in GVPT2. GVPT2 is built on top of DVPT2 by adding a final step to calculate the anharmonic energies as eigenvalues of a variational matrix, whose diagonal elements are the DVPT2 energies, discussed in the previous section, and the off-diagonal elements represent the corrective terms to Fermi resonances, complemented by Darling–Dennison interactions, evaluated over the basis of the canonical harmonic-oscillator wave functions.

On the premise that our reference will remain the canonical representation, which also fully fits the conditions of the application of the GVPT2 scheme at first, we will discuss the theory underlying the definition of the polar variational matrix and then how a full equivalence can be reached between the two representations.

The calculation of a specific resonant term can be carried out by applying a simple generalization of [eq 8](#) to the off-diagonal block coupling states differing in terms of principal quanta ($\nu \neq \nu'$), and it can be organized into two steps:

- Step 1: calculation of anharmonic energies and resonant terms in the canonical representation through the expressions reported in [ref 78](#) (see [Section S4](#) of the [Supporting Information](#) for more details);
- Step 2: combination of the canonical quantities evaluated in step 1 in order to obtain the polar resonant term of interest.

Through the symmetry relations between the anharmonic force constants of symmetric and linear tops, it is possible to prove that the corrective terms due to the Fermi resonances are equivalent in the two representations. Hence, the focus in the following will be on the couplings between states, collectively referred to as Darling–Dennison resonances or interactions.

In this context, particular importance is given to the resonances between states for which the condition $\nu = \nu'$ holds, whose interaction generates the off-diagonal elements of the blocks $\tilde{\mathbf{H}}_{\nu,\nu}^P$.

2.3.3. I-Type Doubling in the Polar Representation. I-type doubling terms involve overtones of a given degenerate mode or combination bands of two degenerate modes. Amat derived a general rule²⁴ for a priori identification of the non-vanishing off-diagonal terms

$$\begin{aligned} & \left\langle \psi_{\nu_s, l_s}^P \left| \tilde{\mathcal{H}} \right| \psi_{\nu_s, (l_s \pm 4)_k}^P \right\rangle \\ & = U_s^\pm \sqrt{(\nu_s \pm l_s + 4)(\nu_s + l_s \pm 2)(\nu_s - l_s \mp 2)(\nu_s \mp l_s)} \end{aligned} \quad (16)$$

$$\begin{aligned} & \left\langle \psi_{\nu_s, l_s}^P \left| \tilde{\mathcal{H}} \right| \psi_{\nu_s, (l_s \pm 2)_s (l_t \mp 2)_t}^P \right\rangle \\ & = R_{st}^\pm \sqrt{(\nu_s \pm l_s + 2)(\nu_t \mp l_t + 2)(\nu_s \mp l_s)(\nu_t \pm l_t)} \end{aligned} \quad (17)$$

$$\begin{aligned} & \langle \psi_{v_s, l_s}^P | \tilde{\mathcal{H}} | \psi_{v_s, (l_s \pm 2), (l_t \pm 2)}^P \rangle \\ & = S_{st}^{\pm} \sqrt{(v_s \pm l_s + 2)(v_t \pm l_t + 2)(v_s \mp l_s)(v_t \mp l_t)} \quad (18) \end{aligned}$$

where the elements defined in eq 16 contribute to the energy only if the order of the principal symmetry axis (n in C_n) is a multiple of 4 and those in eq 18 only if n is even. The expressions of the terms U_s^{\pm} , R_{st}^{\pm} , and S_{st}^{\pm} have been first derived by Grenier and Bresson^{83,84} and then re-derived in ref 71.

In the present derivation, there is no need of developing specific equations for computing the matrix elements defined in eqs 14–16 since they are off-diagonal elements of the diagonal blocks $\tilde{\mathbf{H}}_{2,2_s}^P$ and $\tilde{\mathbf{H}}_{1,1_t,1,1_t}^P$ (the full equations in terms of canonical quantities are reported in Section S5 of the Supporting Information). As a matter of fact, the calculation of l -type terms can be performed concurrently with the conversion of the anharmonic energies through eq 7.

2.3.4. Diagonalization and GVPT2 Energies. In the preceding sections, it has been shown that the blocks of the polar variational matrix ($\tilde{\mathbf{H}}^P$) can always be expressed in terms of their canonical counterpart. In order to understand the effects of such a connection on the GVPT2 energies, we will first consider a $\tilde{\mathbf{H}}^P$ matrix only containing the l -type terms as off-diagonal elements. In this context, the variational problem simplifies to diagonalizing blocks of the type $\tilde{\mathbf{H}}_{2,2_s}^P$ and $\tilde{\mathbf{H}}_{1,1_t,1,1_t}^P$, whose eigenvalues are equal to those of their canonical counterparts. Therefore, the inclusion of l -type doubling at the variational level implies the convergence of GVPT2 energies in the two representations.

This result can be easily generalized, given that an equivalent set of resonances is included in both representations, leading to the following identity:

$$\tilde{\mathbf{H}}^P = \mathbf{P}^{\dagger} \tilde{\mathbf{H}}^C \mathbf{P} \quad (19)$$

where \mathbf{P} is a block-diagonal matrix composed of all rotation matrices required for converting the different diagonal blocks $\tilde{\mathbf{H}}_{v_s, v_t}^C$, and it is itself unitary.

In analogy with the treatment of l -type doubling, the invariance of the eigenvalues of a matrix under unitary transformations can be exploited to state that canonical and polar GVPT2 energies converge to the same values. Let us remark that the l -type terms are present even in the absence of accidental resonances. Consequently, their inclusion is mandatory in order to reach the convergence of the GVPT2 energies. From a practical point of view, eq 19 prevents any ambiguity connected to the representation choice since the set of GVPT2 energies is unique.

2.4. Transition Moments and Intensities. Starting from the available literature (e.g., the study of Tarrago and co-workers⁸⁵ on the transition dipole moments of C_{3v} -symmetry systems), a general computational framework to calculate both IR intensities and Raman activities of molecular systems with non-Abelian symmetries has been devised and implemented in our platform.

Let us start from the band intensities and the associated transition moments of linear and symmetric tops taking into account that thanks to the transformation shown in eq 5, it is possible to use the formulas obtained for asymmetric tops (and reported in Section S6 of the Supporting Information) to obtain their counterparts in the polar representation.

If we look at the fundamental bands (the full equation is reported in Sections S6.1.1 and S6.2.1 of the Supporting Information for degenerate modes), terms of the form

$$-\frac{s_0}{8} \sum_{j=1}^N \sum_{k=1}^N f_{s,ijk} \mathbf{P}_j \left[\frac{1}{\omega_s + \omega_j} - \frac{S(1 - \delta_{sj})}{\omega_s - \omega_j} \right] \quad (20)$$

where \mathbf{P}_j collects the Cartesian components of the first derivative of the property \mathbf{P} with respect to the j th dimensionless normal coordinate, will present a singularity whenever $j = s_2$. For this reason, it is necessary to exclude those modes in the summation so that the degenerate modes are assumed to be resonant, and the “resonant” form (Section S6.2.1 in the Supporting Information) is used instead. For simplicity, only transitions from the ground state (noted 0) to a given final state f are considered, noted “ $0_j f$ ”.

The quantities of interest here for IR and Raman spectroscopies are the dipole strength and Raman activity, labeled in the canonical representation $D_{0_j f}^C$ and $S_{0_j f}^C$, respectively,

$$\begin{aligned} D_{0_j f}^C &= |\langle \boldsymbol{\mu} \rangle_{0_j f}^C|^2 \\ S_{0_j f}^C &= 45 \{a^2\}_{0_j f}^C + 7 \{\gamma^2\}_{0_j f}^C \quad (21) \end{aligned}$$

Here, the invariants $\{a^2\}_{0_j f}^C$ (isotropic) and $\{\gamma^2\}_{0_j f}^C$ (anisotropic) for the most general case of a complex tensor are defined as^{86,87}

$$\begin{aligned} \{a^2\}_{0_j f}^C &= \frac{1}{9} \sum_{\tau=x,y,z} \sum_{\eta=x,y,z} \langle \alpha_{\tau\tau} \rangle_{0_j f}^C \{ \langle \alpha_{\eta\eta} \rangle_{0_j f}^C \}^* \\ \{\gamma^2\}_{0_j f}^C &= \frac{1}{2} \sum_{\tau=x,y,z} \sum_{\eta=x,y,z} [3 \langle \alpha_{\tau\eta} \rangle_{0_j f}^C \{ \langle \alpha_{\tau\eta} \rangle_{0_j f}^C \}^* \\ &\quad - \langle \alpha_{\tau\tau} \rangle_{0_j f}^C \{ \langle \alpha_{\eta\eta} \rangle_{0_j f}^C \}^*] \quad (22) \end{aligned}$$

where τ and η run over the Cartesian axes, while $\langle \boldsymbol{\mu} \rangle_{0_j f}^C$ and $\langle \alpha_{\tau\eta} \rangle_{0_j f}^C$ represent respectively the transition integrals of the electric dipole and a component of the polarizability tensor between the canonical states $|\psi_0^C\rangle$ and $|\psi_f^C\rangle$. Let us anticipate that a closure relation having the same form as eq 8 also holds for dipole strengths and Raman activities due to the unitarity of the rotation matrices.

The theoretical framework currently used for the calculation of transition moments can be straightforwardly extended to the polar representation (for more details, see Appendix B). Thus, once the transition dipole moments and polarizabilities are converted through eq 60, the calculation of both IR and Raman intensities in the polar representation is possible. It is worth mentioning that even though the transition moments evaluated in this way are generally complex, the corresponding intensities are always real.

2.4.1. Infrared Intensities and Raman Activities at the VPT2 Level. For readability, the initial-state label will be dropped, so $D_{0_j f}$ will be simply written D_f . In analogy with energies, the dipole strengths and Raman activities for fundamental states are the same if they are degenerate so that their contribution to the anharmonic spectra is independent of the representation. This result simplifies considerably the whole conversion procedure since among states with up to two quanta, the only states potentially different with respect to the canonical representation are the sets $|\psi_{2_s}^P\rangle$ and $|\psi_{1,1_t}^P\rangle$.

Concerning degenerate overtones, the dipole strengths of the polar states are

$$D_{2_s, \pm 2_s}^P = \frac{1}{4}[D_{2_s}^C + D_{1_s, 1_{2_s}}^C - 2\langle \boldsymbol{\mu} \rangle_{2_s}^C \langle \boldsymbol{\mu} \rangle_{2_s}^C]$$

$$D_{2_s, 0_s}^P = \frac{1}{2}[D_{2_s}^C - D_{1_s, 1_{2_s}}^C + 2\langle \boldsymbol{\mu} \rangle_{2_s}^C \langle \boldsymbol{\mu} \rangle_{2_s}^C]$$
(23)

where $\langle \boldsymbol{\mu} \rangle_{2_s}^C$ and $\langle \boldsymbol{\mu} \rangle_{2_{s_2}}^C$ are respectively the vectors containing the Cartesian components of the transition dipole moments associated with the states $|\psi_{2_s}^C\rangle$ and $|\psi_{2_{s_2}}^C\rangle$, and $D_{2_s}^C$ is defined as

$$D_{2_s}^C = D_{2_{s_1}}^C + D_{2_{s_2}}^C + D_{1_s, 1_{2_s}}^C$$
(24)

The Raman activities can be expressed in a compact notation through the introduction of the variables $A_{r,s}^C$, $\Gamma_{r,s}^C$, and $S_{r,s}^C$

$$A_{r,s}^C = \frac{1}{9} \sum_{\tau=x,y,z} \sum_{\eta=x,y,z} \langle \alpha_{\tau\tau} \rangle_r^C \{ \langle \alpha_{\eta\eta} \rangle_s^C \}^*$$

$$\Gamma_{r,s}^C = \frac{1}{2} \sum_{\tau=x,y,z} \sum_{\eta=x,y,z} [3\langle \alpha_{\tau\eta} \rangle_r^C \{ \langle \alpha_{\eta\tau} \rangle_s^C \}^* - \langle \alpha_{\tau\tau} \rangle_r^C \{ \langle \alpha_{\eta\eta} \rangle_s^C \}^*]$$

$$S_{r,s}^C = 45A_{r,s}^C + 7\Gamma_{r,s}^C$$
(25)

which can be interpreted respectively as the “off-diagonal” terms of $\{a^2\}_r^C$, $\{\gamma^2\}_r^C$, and S_r^C .

As a consequence, the expressions of the Raman activities of the polar states of interest are

$$S_{2_s, \pm 2_s}^P = \frac{1}{4}[S_{2_s}^C + S_{1_s, 1_{2_s}}^C - 2S_{2_s, 2_{s_2}}^C]$$

$$S_{2_s, 0_s}^P = \frac{1}{2}[S_{2_s}^C - S_{1_s, 1_{2_s}}^C + 2S_{2_s, 2_{s_2}}^C]$$
(26)

where

$$S_{2_s}^C = S_{2_{s_1}}^C + S_{2_{s_2}}^C + S_{1_s, 1_{2_s}}^C$$
(27)

A similar analysis applied to the transition moments of binary combination bands involving degenerate modes yields the dipole strengths of the polar states

$$D_{1_s, 1_r, \pm 1_s, \pm 1_r}^P = \frac{1}{4}[D_{st}^C - 2\langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_1}}^C \langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_2}}^C + 2\langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_2}}^C \langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_1}}^C]$$

$$D_{1_s, 1_r, \pm 1_s, \mp 1_r}^P = \frac{1}{4}[D_{st}^C + 2\langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_1}}^C \langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_2}}^C - 2\langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_2}}^C \langle \boldsymbol{\mu} \rangle_{1_s, 1_{t_1}}^C]$$
(28)

where

$$D_{st}^C = D_{s_1, t_1}^C + D_{s_1, t_2}^C + D_{1_s, 2_{t_1}}^C + D_{1_s, 2_{t_2}}^C$$
(29)

Finally, the corresponding Raman activities are

$$S_{1_s, 1_r, \pm 1_s, \pm 1_r}^P = \frac{1}{4}[S_{st}^C - 2S_{s_1, 1_{t_1}; 1_{s_2}, 1_{t_2}}^C + 2S_{1_s, 1_{t_2}; 1_{s_2}, 1_{t_1}}^C]$$

$$S_{1_s, 1_r, \pm 1_s, \mp 1_r}^P = \frac{1}{4}[S_{st}^C + 2S_{s_1, 1_{t_1}; 1_{s_2}, 1_{t_2}}^C - 2S_{1_s, 1_{t_2}; 1_{s_2}, 1_{t_1}}^C]$$
(30)

where

$$S_{st}^C = S_{s_1, t_1}^C + S_{s_1, t_2}^C + S_{s_2, t_1}^C + S_{s_2, t_2}^C$$
(31)

From a comparison of eqs 25 and 26 with eqs 11a and 11b and eqs 28 and 30 with eqs 13a and 13b, it is possible to observe that the conversion of dipole strengths and Raman activities is ruled

by expressions similar to those employed for the anharmonic energies.

2.4.2. Introduction of the Variational Correction. It has been demonstrated that if the variational matrix $\tilde{\mathbf{H}}^P$ in the polar representation can be expressed by a rotation of the canonical one $\tilde{\mathbf{H}}^C$ (see eq 19), the anharmonic energies within the two representations converge to the same values. This result can be easily extended to vibrational intensities. Indeed, while the definition of variational states is dependent on the representation of the reference states, polar and canonical variational states are equivalent when projected onto the same basis. In this context, the canonical basis is chosen as a reference.

As a matter of fact, the variational states, and hence the intensities, do not depend on the representation. Let us remark that the equality of the variational states holds even when only l -type doubling terms are included as off-diagonal elements of the variational matrix, that is, even in the absence of accidental resonances.

In summary, the anharmonic spectrum is completely independent of the representation, with the only difference being the harmonic-state basis. Thus, the computational protocol currently employed for asymmetric tops can be straightforwardly extended to the treatment of symmetric/linear tops without any loss of accuracy.

2.5. Extension to Spherical Tops. In this section, we will show how the framework devised for symmetric and linear tops can be extended to spherical tops. For the sake of concision, the rotation-based formulation will be applied to systems presenting at most threefold degenerate vibrations. Actually, the eigenstates of the three-dimensional isotropic harmonic oscillator are again complex linear combinations of the canonical wave functions. This type of combinations can be extended to higher degeneracy orders, in which case, the following derivation can be straightforwardly adapted to systems exhibiting such characteristics.

Back to our application, the starting point remains the canonical representation since the specificity of the spherical top lies in the definition of the rotation matrices necessary for the transformation of the wave functions. While the canonical wave function is still given by eq 2, the so-called spherical wave function has the following form, where non-degenerate and doubly and triply degenerate modes are gathered in different terms

$$\psi_v^S(\mathbf{q}) = \prod_{m=1}^{N'} \varphi_{v_m}^C(q_m) \prod_{s=1}^{N''} \varphi_{v_s, l_s}^P(\rho_s, \theta_s) \prod_{s'=1}^{N'''} \varphi_{v_{s'}, k_{s'}, m_{s'}}^S(r_{s'}, \gamma_{s'}, \phi_{s'})$$
(32)

where $\varphi_{v_s, k_{s'}, m_{s'}}^S(r_{s'}, \gamma_{s'}, \phi_{s'})$ are the solutions of the three-dimensional isotropic harmonic oscillator Hamiltonian expressed with respect to the spherical coordinates stemming from a trio of degenerate modes, r_s , γ_s , and ϕ_s , and the quantum numbers v_s , $k_{s'}$, and $m_{s'}$ can assume the following values:

$$v_s = 0, 1, 2, \dots$$

$$k_{s'} = v_s, v_s - 2, v_s - 4, \dots, 0 \text{ or } 1$$

depending if v_s is even or odd

$$m_{s'} = -k_{s'}, -k_{s'} + 1, \dots, 0, \dots, k_{s'} - 1, k_{s'}$$
(33)

In analogy with symmetric and linear tops, the analysis of the spherical wave functions enables the definition of the rotation matrices \mathbf{Q} .

$$|\psi_\nu^S\rangle = \mathbf{Q}_\nu^T |\psi_\nu^C\rangle \quad (34)$$

where $|\psi_\nu^S\rangle$ contains the spherical states sharing the same principal angular number ν and $|\psi_\nu^C\rangle$ is the corresponding canonical counterpart. Let us stress that when the states $|\psi_\nu^S\rangle$ only involve non-degenerate and doubly degenerate modes, the rotation matrix is \mathbf{P}_ν .

Let us now analyze the effect of rotations in deeper detail, highlighting the analogies with symmetric and linear tops. The canonical expression of ZPVE, whose expression is reported in eq 9, can be equivalently expressed through the following formula:

$$\varepsilon_0^C = \langle \psi_0^C | \tilde{\mathcal{H}} | \psi_0^C \rangle \quad (35)$$

where ψ_0^C represents the harmonic ground-state wave function in the canonical representation. The main advantage of the rotation-based framework is that the specific properties of the rotor are collected in the definition of the wave function, more specifically through the use of rotation matrices to carry out the conversion procedure described above. As a result, the reference basis is always the canonical one, and the operators depending on the normal coordinates and their conjugate momenta are never subject to any modifications. Based on this, the expression of the spherical counterpart ε_0^S is

$$\varepsilon_0^S = \langle \psi_0^S | \tilde{\mathcal{H}} | \psi_0^S \rangle \quad (36)$$

where ψ_0^S is the harmonic ground-state wave function in the spherical representation. Since the ground-state wave function is independent of the representation (see Appendix B for more details), we can conclude that

$$\varepsilon_0^C = \varepsilon_0^S \quad (37)$$

Hence, the resonance-free ZPVE in the spherical representation can be still evaluated through the customary expression, which can be then used for systems presenting both doubly and triply degenerate modes without any restriction.

In the previous sections, all different types of bands involving up to two-quanta excitations characterizing symmetric and linear tops have been derived and analyzed separately. In general terms, this separation is not necessary since only a few rotation matrices are actually sufficient to build all the other ones. With the aim of treating all the states up to two quanta, the largest value that the quantum numbers ν_m , ν_s , and ν_t reported in eq 34 can assume is 2 so that the matrices \mathbf{P}_1 , \mathbf{P}_2 , \mathbf{Q}_1 , and \mathbf{Q}_2 allow to express any spherical state in terms of canonical ones (the extension to three-quanta states would only require additional matrices, \mathbf{P}_3 , and \mathbf{Q}_3).

In the same way as symmetric and linear tops, both transition energies and intensities for states only involving one excited quantum of a threefold degenerate vibration are still independent of the representation, and this is also true for the ZPVE. Concerning overtones and binary combination bands involving triply degenerate modes, a complete derivation of both energies and intensities in the spherical representation for states up to two quanta is reported in Section S7 of the Supporting Information.

Finally, it is worth mentioning that the equivalence of representations at the GVPT2 level is not affected by the presence of triply degenerate vibrations, with it being strictly related to the connection of the wave functions by unitary

matrices. Therefore, the GVPT2 results remain independent of the representation.

3. COMPUTATIONAL DETAILS

The theoretical framework presented in the previous sections has been implemented in a development version of the Gaussian package.⁸⁸ Most of the available electronic structure computations allowing analytic computation of second energy and first property derivatives have been employed. These include HF and second-order Møller–Plesset (MP2) wave-function methods together with different flavors of DFT including representative hybrid (B3LYP^{89–92}) and double-hybrid (B2PLYP^{93,94}) exchange–correlation functionals, with the inclusion of empirical dispersion contributions by means of Grimme’s D3 model with Becke–Johnson damping^{95,96} (hereafter noted B3D3 and B2D3, respectively). As the core of this work concerns the development, functionals with a well-documented reliability on the molecular systems chosen here for illustration purposes were selected. The reliability of the B2PLYP functional in the calculation of both harmonic and anharmonic frequencies has been demonstrated in the literature.^{97,98} The B3LYP functional has been used for the calculation of both harmonic and anharmonic frequencies only for linear systems, where B2PLYP results have also been shown. Concerning symmetric and spherical tops, the B3LYP functional has been only used for the calculation of the anharmonic corrections in the hybrid scheme. In this respect, it has been demonstrated that the quality of the harmonic frequencies is much more critical if compared with that of the corresponding anharmonic corrections.² The so-called calendar basis sets jun-cc-pVDZ and jun-cc-pVTZ⁹⁹ have been consistently employed (referred to in the following as JnDZ and JnTZ, respectively).

The anharmonic data required for the VPT2 calculation of frequencies and intensities have been obtained by finite differences of analytical force constants (full cubic and semi-diagonal quartic force constants) and first-order derivatives of the properties (full second and semi-diagonal third derivatives), employing a default displacement of $\delta Q_i = 0.01 \sqrt{\text{amu} \cdot \text{Å}}$ along each mass-weighted normal coordinate Q_i .^{42,100} From a practical point of view, the generation of the anharmonic force field and higher-order property derivatives is the most expensive step once the equilibrium structure has been found. Indeed, $2N$ frequency calculations are needed in addition to the one at the reference geometry, required to generate the displacement vectors. Being independent from one another, they can be run in parallel on separate machines, with the final constants built at the end of the process. Hence, in an optimal scenario, the computational cost can be reduced to roughly twice what is needed at the harmonic level. The VPT2 calculations themselves on systems of this size last a few minutes, independent of the scheme chosen.

For the treatment of resonances, the protocol detailed in ref 76 was used with the default parameters.

In addition to the basic formulation of VPT2, the DVPT2 and GVPT2 schemes are also used for the calculation of both anharmonic energies and intensities. The use of DVPT2 is necessary in the presence of Fermi resonances to avoid the unphysical results issuing from the standard VPT2 equations. In addition to recovering the discarded terms from DVPT2, the GVPT2 scheme allows a straightforward account of Darling–Dennison resonances, which are not explicitly considered within the purely perturbative approach. Besides improving the overall agreement with experimental energies, they can be critical to

Table 1. Comparison of Experimental and Computed Anharmonic Fundamental Wavenumbers (in cm^{-1}) for the Linear Molecules HCN, HNC, and C_2H_2 ^a

| | symm. | MP2//MP2 ^b | | B3D3//B3D3 ^b | | B2D3//B2D3 ^b | | exp. |
|-------------------------------------|------------|-----------------------|---------------------|-------------------------|---------------------|-------------------------|---------------------|------|
| | | ω | ν_{VPT2} | ω | ν_{VPT2} | ω | ν_{VPT2} | |
| HCN ^c | | | | | | | | |
| $ 1_1\rangle$ | Σ | 3459 | 3328 | 3440 | 3309 | 3455 | 3322 | 3312 |
| $ 1_2\rangle$ | | 2023 | 1989 | 2199 | 2172 | 2125 | 2094 | 2097 |
| $ 1_3, \pm 1_3\rangle$ | Π | 710 | 704 | 757 | 738 | 740 | 726 | 714 |
| MAE | | | 45 | | 34 | | 8 | |
| HNC ^d | | | | | | | | |
| $ 1_1\rangle$ | Σ | 3819 | 3658 | 3801 | 3632 | 3816 | 3650 | 3653 |
| $ 1_2\rangle$ | | 2019 | 1986 | 2103 | 2070 | 2060 | 2025 | 2029 |
| $ 1_3, \pm 1_3\rangle$ | Π | 492 | 479 | 471 | 436 | 470 | 445 | 477 |
| MAE | | | 17 | | 34 | | 13 | |
| C_2H_2 ^e | | | | | | | | |
| $ 1_1\rangle$ | Σ_g | 3525 | 3389 | 3512 | 3378 | 3524 | 3389 | 3372 |
| $ 1_2\rangle$ | | 1969 | 1930 | 2068 | 2036 | 2024 | 1988 | 1975 |
| $ 1_3\rangle$ | Σ_u | 3437 | 3312 | 3412 | 3287 | 3431 | 3305 | 3289 |
| $ 1_4, \pm 1_4\rangle$ | Π_g | 592 | 561 | 663 | 621 | 638 | 602 | 613 |
| $ 1_5, \pm 1_5\rangle$ | Π_g | 748 | 718 | 768 | 735 | 762 | 730 | 730 |
| MAE | | | 30 | | 16 | | 11 | |

^aMean absolute errors (MAEs) are also reported. The polar vibrational states are indicated as $|v_i, l_i\rangle$. ^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated through the JnTZ basis set. ^cExperimental values from ref 69. ^dExperimental values from ref 111. ^eExperimental values from ref 112.

Table 2. Comparison of Experimental and Computed Anharmonic VPT2, DVPT2, and GVPT2 Wavenumbers (in cm^{-1}) of CO_2 in the Polar Representation^a

| state | MP2 ^b | | | | B3D3 ^b | | | | B2D3 ^b | | | | exp. |
|----------------------------|------------------|---------------------|----------------------|----------------------|-------------------|---------------------|----------------------|----------------------|-------------------|---------------------|----------------------|----------------------|---------------------------|
| | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | |
| Fundamentals | | | | | | | | | | | | | |
| $ 1_1, \pm 1_1\rangle$ | 659 | 657 | 657 | 657 | 674 | 670 | 670 | 670 | 666 | 662 | 662 | 662 | 668 ^{c,d,e,f} |
| $ 1_2\rangle$ | 1326 | 1697 | 1309 | 1262 | 1369 | 1492 | 1349 | 1291 | 1341 | 1646 | 1321 | 1272 | 1285 ^{c,d,e,f,g} |
| $ 1_3\rangle$ | 2405 | 2367 | 2367 | 2367 | 2403 | 2356 | 2356 | 2356 | 2387 | 2342 | 2342 | 2342 | 2349 ^{c,d,e,f,g} |
| Overtones | | | | | | | | | | | | | |
| $ 2_1, \pm 2_1\rangle$ | 1319 | 1315 | 1315 | 1315 | 1347 | 1341 | 1341 | 1341 | 1332 | 1326 | 1326 | 1326 | 1336 ^{d,e,f} |
| $ 2_1, 0_1\rangle$ | 1319 | 933 | 1320 | 1368 | 1347 | 1203 | 1346 | 1403 | 1332 | 1005 | 1330 | 1379 | 1388 ^{d,e,f,g} |
| $ 2_2\rangle$ | 2652 | 3389 | 2614 | 2614 | 2739 | 2979 | 2692 | 2692 | 2682 | 3287 | 2635 | 2635 | 2548 ^g |
| $ 2_3\rangle$ | 4810 | 4714 | 4714 | 4714 | 4806 | 4688 | 4688 | 4688 | 4774 | 4660 | 4660 | 4660 | 4673 ^{f,g} |
| Combinations | | | | | | | | | | | | | |
| $ 1_1 1_2, \pm 1_1\rangle$ | 1986 | 2736 | 1960 | 1960 | 2043 | 2300 | 2013 | 2013 | 2007 | 2628 | 1978 | 1978 | 2077 ^{d,f} |
| $ 1_1 1_3, \pm 1_1\rangle$ | 3065 | 3013 | 3013 | 3013 | 3076 | 3014 | 3014 | 3014 | 3053 | 2992 | 2992 | 2992 | 3004 ^{f,g} |
| $ 1_2 1_3\rangle$ | 3731 | 4051 | 3664 | 3664 | 3772 | 3830 | 3686 | 3686 | 3728 | 3970 | 3644 | 3644 | 3613 ^{d,e,f,g} |
| MAE | | 291 | 43 | 38 | | 130 | 43 | 34 | | 244 | 36 | 29 | |

^aThe polar vibrational states are indicated as $|v_i, l_i\rangle$. ^bBasis set: JnTZ. ^cReference 113. ^dReference 114. ^eReference 115. ^fReference 116. ^gReference 117.

obtain correct band shapes. However, the induced transformation can result in a significant deviation from the harmonic oscillator-based description of the vibrational states preserved by VPT2, thus requiring some extra work for the band assignment. Further details concerning the different VPT2 schemes concerning both vibrational energies and intensities have been recently reported in ref 101.

The so-called hybrid force field scheme has also been employed.^{2,102–105} In this approach, harmonic and anharmonic contributions are treated at different levels of theory in view of their different contribution to the final VPT2 result. In particular, anharmonic contributions have been consistently computed with the JnDZ basis set and harmonic terms with JnTZ. In some cases, which will be explicitly mentioned in the discussion, the results at the CCSD(T) level in conjunction with

extended basis sets, already available in the literature, were employed for the latter. It should be noted that the higher-level harmonic frequencies are not simply added to anharmonic corrections but are also employed in the conversion of the anharmonic force constants and property derivatives, in the construction of the χ matrix, the definition of the resonant terms, and for the intensity. The JnTZ basis set has been employed for the calculation of anharmonic contributions of aromatic systems since it is well known that out-of-plane vibrations of these molecules are particularly sensitive to the basis-set dimension.^{106–109} An analogous remark applies to CO_2 .¹¹⁰

Table 3. Computed Anharmonic VPT2, DVPT2, and GVPT2 Wavenumbers (in cm^{-1}) of CO_2 in the Canonical Representation^a

| state | MP2 ^b | | | | B3D3 ^b | | | | B2D3 ^b | | | |
|------------------------|------------------|---------------------|----------------------|----------------------|-------------------|---------------------|----------------------|----------------------|-------------------|---------------------|----------------------|----------------------|
| | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} |
| Fundamentals | | | | | | | | | | | | |
| $ 1_{1a}\rangle$ | 659 | 657 | 657 | 657 | 674 | 670 | 670 | 670 | 666 | 662 | 662 | 662 |
| $ 1_{1b}\rangle$ | 659 | 657 | 657 | 657 | 674 | 670 | 670 | 670 | 666 | 662 | 662 | 662 |
| $ 1_2\rangle$ | 1326 | 1697 | 1309 | 1262 | 1369 | 1492 | 1349 | 1291 | 1341 | 1646 | 1321 | 1272 |
| $ 1_3\rangle$ | 2405 | 2367 | 2367 | 2367 | 2403 | 2356 | 2356 | 2356 | 2387 | 2342 | 2342 | 2342 |
| Overtones | | | | | | | | | | | | |
| $ 2_{1a}\rangle$ | 1319 | 1124 | 1318 | 1315 | 1347 | 1272 | 1344 | 1341 | 1332 | 1166 | 1328 | 1326 |
| $ 2_{1b}\rangle$ | 1319 | 1124 | 1318 | 1315 | 1347 | 1272 | 1344 | 1341 | 1332 | 1166 | 1328 | 1326 |
| $ 1_{1a}1_{1b}\rangle$ | 1319 | 1315 | 1315 | 1368 | 1347 | 1341 | 1341 | 1403 | 1332 | 1326 | 1326 | 1379 |
| $ 2_2\rangle$ | 2652 | 3389 | 2614 | 2614 | 2739 | 2979 | 2692 | 2692 | 2682 | 3287 | 2635 | 2635 |
| $ 2_3\rangle$ | 4810 | 4714 | 4714 | 4714 | 4806 | 4688 | 4688 | 4688 | 4774 | 4660 | 4660 | 4660 |
| Combinations | | | | | | | | | | | | |
| $ 1_{1a}1_2\rangle$ | 1986 | 2736 | 1960 | 1960 | 2043 | 2300 | 2013 | 2013 | 2007 | 2628 | 1978 | 1978 |
| $ 1_{1b}1_2\rangle$ | 1986 | 2736 | 1960 | 1960 | 2043 | 2300 | 2013 | 2013 | 2007 | 2628 | 1978 | 1978 |
| $ 1_{1a}1_3\rangle$ | 3065 | 3013 | 3013 | 3013 | 3076 | 3014 | 3014 | 3014 | 3053 | 2992 | 2992 | 2992 |
| $ 1_{1b}1_3\rangle$ | 3065 | 3013 | 3013 | 3013 | 3076 | 3014 | 3014 | 3014 | 3053 | 2992 | 2992 | 2992 |
| $ 1_21_3\rangle$ | 3731 | 4051 | 3664 | 3664 | 3772 | 3830 | 3686 | 3686 | 3728 | 3970 | 3644 | 3644 |

^aThe canonical vibrational states are indicated as $|\nu_i\nu_j\rangle$, and the subscripts “a” and “b” distinguish degenerate modes. ^bBasis set: JnTZ.

4. RESULTS AND DISCUSSION

The framework previously discussed has been validated through a series of applications to linear and symmetric tops, illustrative of representative non-Abelian symmetry groups.

4.1. Linear Molecules. First, a set of three- and four-atom molecules, including hydrogen cyanide (HCN), hydrogen isocyanide (HNC), and acetylene (C_2H_2), has been considered. The absence of Fermi and 1–1 Darling–Dennison resonances (between fundamental states) implies that the anharmonic fundamentals do not vary going from VPT2 to DVPT2 or GVPT2 schemes. The fundamental harmonic and anharmonic frequencies obtained with the hybrid force-field model described above are reported in Table 1.

As can be seen from Table 1, the computed fundamental energies are in good agreement with the experimental counterparts, with the largest discrepancy concerning the degenerate mode of HNC. At the B3D3 and B2D3 levels of theory, such an error is most likely due to the underestimation of the corresponding harmonic frequency, which is lower of the experimental value in both cases.

As an example of comparison between the polar and canonical representations, the vibrational frequencies of CO_2 for all states up to two quanta in the polar representation have been calculated and analyzed. The set of wavenumbers in the polar representation is reported in Table 2 and compared with reference experimental values.

For comparison purposes, the VPT2, DVPT2, and GVPT2 wavenumbers in the canonical representation are reported in Table 3.

As expected, the states involving at most one vibrational quantum in the degenerate bending mode have the same frequency irrespective of the chosen representation. Conversely, the frequencies of the first overtones related to the bending mode change between the representations, even though the energies converge to the same values when the GVPT2 model is applied.

By comparing Tables 2 and 3, it is straightforward to verify that the sum of the energies of the states $|\psi_{2,\pm 2,1}^P\rangle$ and $|\psi_{2,0,1}^P\rangle$

equals that of the states $|\psi_{2,1a}^C\rangle$, $|\psi_{2,1b}^C\rangle$, and $|\psi_{1_{1a}1_{1b}}^C\rangle$. Such an outcome is in full agreement with eq 8.

Due to the strong Fermi resonance between the overtone of the bending mode and the fundamental of the symmetric stretching ($2\omega_1 \approx \omega_2$), the CO_2 molecule has been widely used as a prototype in the study of resonances. Let us underline that only the state $|\psi_{2,0,1}^P\rangle$ is involved in the resonance since the coupling between the states $|\psi_{2,\pm 2,1}^P\rangle$ and $|\psi_{1_2}^C\rangle$ is symmetry-forbidden. In order to show this from the mathematical point of view, let us consider the general case of a Fermi resonance between the fundamental of a non-degenerate mode m and the overtone of a degenerate mode s . Both possible resonant terms, $\langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,s,0}^P \rangle$ and $\langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,\pm 2,s}^P \rangle$, can be conveniently expressed in terms of canonical interaction terms

$$\begin{aligned} \langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,s,0}^P \rangle &= -\frac{1}{\sqrt{2}} [\langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,s_1}^C \rangle + \langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,s_2}^C \rangle] \\ &= -\frac{f_{mss}^{(I)}}{\sqrt{2}} \\ \langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,\pm 2,s}^P \rangle &= \frac{1}{2} [\langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,s_1}^C \rangle - \langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,s_2}^C \rangle] \\ &\quad \pm \frac{i}{\sqrt{2}} \langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{1_{s_1}1_{s_2}}^C \rangle \\ &= \frac{1}{4} [2f_{mss}^{(III)} \pm if_{mss}^{(IV)}] \end{aligned} \quad (38)$$

where the expressions of the matrix elements associated to the Fermi resonances in the canonical representation⁷⁶ have been used in conjunction with the symmetry rules and the definition of the force constants $f_{mss}^{(\sigma)}$ ($\sigma = \text{I, III, IV}$) reported in Table A2 of ref 71. For linear molecules, the only non-vanishing force constant $f_{mss}^{(\sigma)}$ corresponds to $\sigma = \text{I}$ so that $\langle \psi_{1_m}^C | \tilde{\mathcal{H}} | \psi_{2,\pm 2,s}^P \rangle$ always vanishes for this kind of systems.

The results obtained at the B3D3/JnTZ level of theory have been employed in the calculation of both IR and Raman spectra,

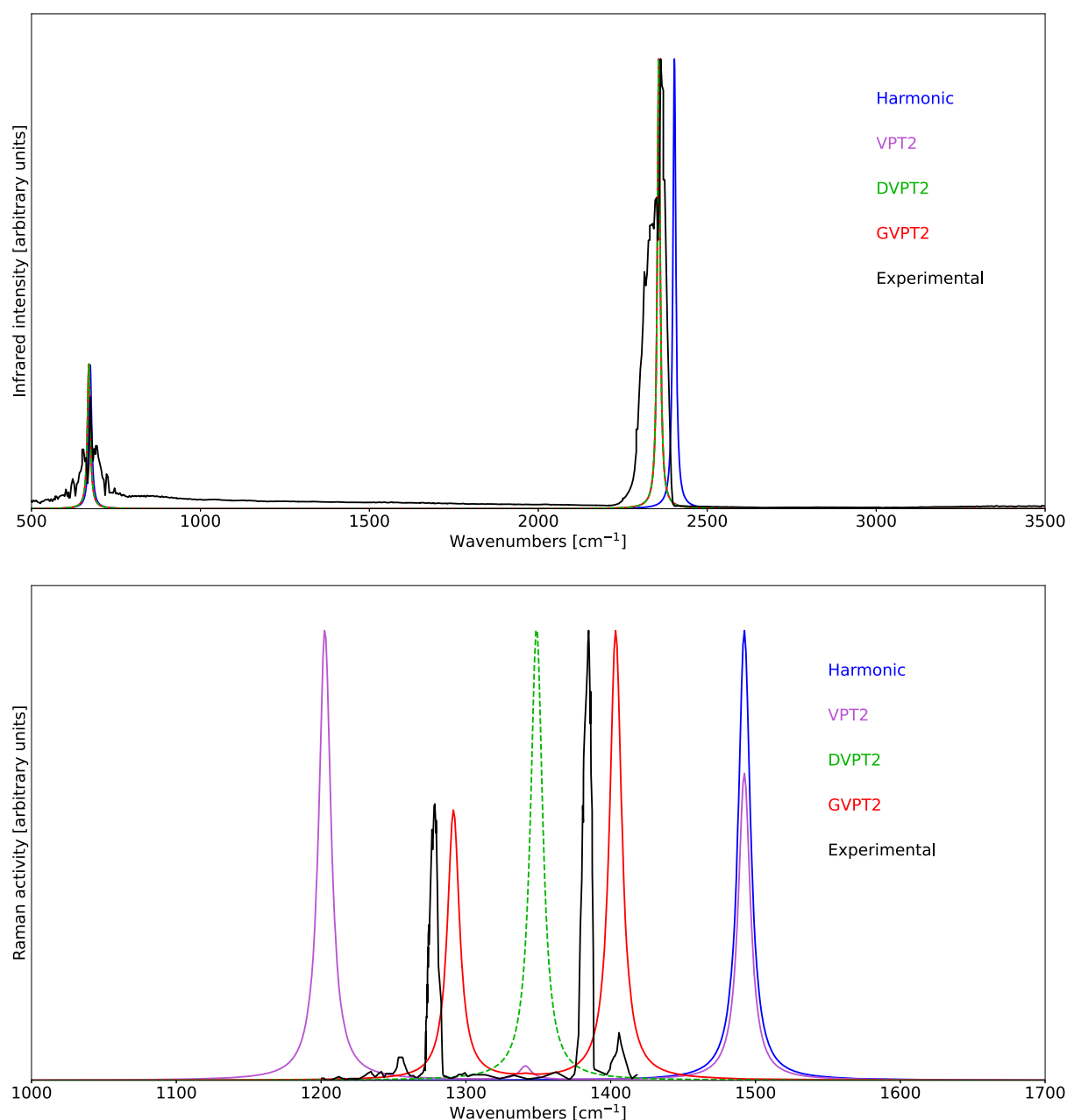


Figure 1. Comparison of the computed IR (top) and Raman (bottom) spectra of CO₂ at the B3D3/JnTZ level of theory with the experimental data. Spectral line shapes have been convoluted by Lorentzian distribution functions with HWHMs of 5 cm⁻¹. Experimental IR spectrum from the NIST Web Book.¹¹⁸ Experimental Raman spectrum from ref 119. All spectra are normalized by setting the intensity of their highest peak to unity.

with the harmonic and anharmonic results obtained within the VPT2, DVPT2, and GVPT2 schemes being compared with the experimental data in Figure 1. As expected, the best agreement between theoretical and experimental spectra is reached within the GVPT2 scheme. This is particularly evident in the Raman spectrum, in which the inclusion of the Fermi resonance discussed above at the variational level leads to an excellent reproduction of the relative intensities characterizing the Fermi diad present in the experimental spectrum.

As a last example of linear molecules, dicyanoacetylene (C₄N₂, see Figure 2) is considered.

Dicyanoacetylene has been detected on the Titan moon of Saturn by IR spectroscopy, and it is used as a prototype for similar astrochemical systems, such as cyanopolynes.¹²⁰



Figure 2. Molecular structure of dicyanoacetylene.

The IR and Raman spectra of this system have been the object of several experimental works,^{121–128} while a theoretical analysis has been recently presented by Dargelos and Pouchan.¹²⁰ With the aim of showing the application of our computational framework to longer chain systems, the VPT2, DVPT2, and GVPT2 fundamentals at different levels of theory have been calculated and compared with their experimental counterparts in Table 4.

At both B3D3 and B2D3 levels, two Fermi resonances of the first type coupling the states $|1_4\rangle$ and $|2_5\rangle$, and $|1_3\rangle$ and $|2_7,0_7\rangle$ are

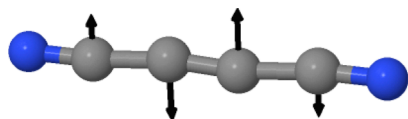
Table 4. Comparison of Experimental and Computed Anharmonic Fundamental VPT2, DVPT2, and GVPT2 Wavenumbers (in cm^{-1}) of Dicyanoacetylene^a

| | symm. | MP2//MP2 ^b | | | B3D3//B3D3 ^b | | | B2D3//B2D3 ^b | | | exp. | | | |
|------------------------|------------|-----------------------|---------------------|----------------------|-------------------------|----------|---------------------|-------------------------|----------------------|----------|------|---------------------|----------------------|----------------------|
| | | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} |
| $ 1_1\rangle$ | Σ_g | 2235 | 2174 | 2187 | 2178 | 2377 | 2607 | 2340 | 2315 | 2313 | 2236 | 2269 | 2254 | 2270 |
| $ 1_2\rangle$ | | 2031 | 1991 | 1991 | 1991 | 2219 | 2190 | 2190 | 2190 | 2144 | 2110 | 2110 | 2110 | 2123 |
| $ 1_3\rangle$ | | 600 | 606 | 606 | 606 | 617 | 634 | 611 | 617 | 610 | 623 | 604 | 610 | 606 |
| $ 1_4\rangle$ | Σ_u | 2152 | 2105 | 2105 | 2105 | 2342 | 2309 | 2309 | 2309 | 2261 | 2221 | 2221 | 2221 | 2245 |
| $ 1_5\rangle$ | | 1155 | 1149 | 1149 | 1149 | 1186 | 1186 | 1186 | 1186 | 1174 | 1171 | 1171 | 1171 | 1155 |
| $ 1_{6\pm 1_6}\rangle$ | Π_g | 504 | 422 | 422 | 422 | 559 | 330 | 330 | 330 | 537 | 375 | 375 | 375 | 505 |
| $ 1_{7\pm 1_7}\rangle$ | | 263 | 203 | 203 | 203 | 285 | 233 | 233 | 233 | 277 | 219 | 219 | 219 | 261 |
| $ 1_{8\pm 1_8}\rangle$ | Π_u | 478 | 463 | 463 | 463 | 510 | 475 | 475 | 475 | 497 | 469 | 469 | 469 | 472 |
| $ 1_{9\pm 1_9}\rangle$ | | 109 | 82 | 82 | 82 | 114 | 94 | 94 | 94 | 112 | 87 | 87 | 87 | 107 |
| MAE ^c | | | 58 | 57 | 58 | | 71 | 35 | 33 | | 21 | 15 | 17 | |

^aMean absolute errors (MAEs) are also reported. The polar vibrational states are indicated as $|v_i, l_i\rangle$. ^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set. ^cStates $|1_{6\pm 1_6}\rangle$ excluded.

detected, with only the former being present at the MP2 level. Within the GVPT2 scheme, such Fermi resonances have been included variationally, together with the proper l -type doubling terms, and 2–2 Darling–Dennison resonances, with the latter being present only at the B3D3 and B2D3 levels.

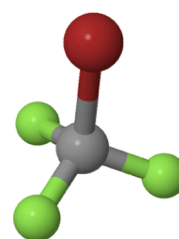
The best agreement between theoretical and experimental fundamentals is reached at the B2D3 level, despite an out-of-scale discrepancy detected for the degenerate states $|1_{6\pm 1_6}\rangle$ regardless of the electronic level of theory. The corresponding normal mode is depicted in Figure 3.

**Figure 3.** Graphical representation of normal mode 6 of dicyanoacetylene.

At the MP2 level, such a difference can be ascribed to an underestimation of the harmonic frequency (which is lower than the experimental value), whereas density functional calculations show an excessive anharmonic correction (229 and 158 cm^{-1} for B3D3 and B2D3, respectively). Interestingly, the MP2 harmonic energies of the $|1_{6\pm 1_6}\rangle$ fundamentals are very close to those of the experiment, a trend observed for all modes below 1000 cm^{-1} .

4.2. Symmetric Tops. Shifting to symmetric-top systems, we analyze a set of six molecules, namely, bromotrifluoromethane (CF_3Br), mono- and tri-deuterated methane (CH_3D and CHD_3 , respectively), the cyclopentadienyl anion (C_5H_5^-), benzene (C_6H_6), and pentaborane (B_5H_9), which are characterized by a principal axis of order ranging from 3 to 6 and belong to the molecular point groups C_{nv} and D_{nh} . As already pointed out, anharmonic corrections at the MP2 level are free from the issues of numerical integration, which is not always true for methods rooted into DFT. In this respect, MP2 is more suitable for validation purposes of the new rotation-based framework, significantly reducing the possibility of errors in the symmetry relations proposed by Amat and Henry, and characterizing the anharmonic force field of symmetric and linear tops. As a matter of fact, while MP2 corrections will be used for all symmetric tops studied in the following, the use of DFT will be limited to the C_{3v} systems.

Let us start from CF_3Br (see Figure 4), whose anharmonic IR spectrum has been simulated at the MP2/JnDZ level of theory

**Figure 4.** Molecular structure of CF_3Br and CH_3D .

with harmonic frequencies corrected by CCSD(T)/aug-cc-pVTZ-PP¹²⁹ calculations. In the former case, no Fermi resonances are present, while in the latter case, a single Fermi resonance of the second type, involving the non-degenerate fundamental $|1_1\rangle$ and combination $|1_2 1_3\rangle$, has been detected. In both cases, the only Darling–Dennison resonances detected correspond to the l -type doubling of type R (which is the only one present when the principal axis order is 3).

The set of anharmonic fundamental frequencies, together with the energy of state $|1_2 1_3\rangle$ (the only two-quanta state involved in a Fermi resonance within the CC//MP2 scheme), is compared with the experimental data in Table 5 where, as expected, the improvement of the results due to the use of coupled-cluster (CC) harmonic frequencies leads to excellent agreement.

The hybrid CC//MP2 results have also been used in the calculation of the anharmonic IR spectrum. Furthermore, in order to show the importance of anharmonic effects in the reproduction of both the position and intensity of the bands of the spectrum, the theoretical spectra have been evaluated by a stepwise inclusion of the anharmonic corrections. More specifically, the IR spectrum has been first evaluated at the purely harmonic level (HH), followed by the inclusion of the anharmonic corrections to the energies (AH) and finally to both energies and intensities (AA). A full comparison of theoretical and experimental spectra is reported in Figure 5.

As can be seen from the top panel of Figure 5, a remarkable improvement in the position of the bands is obtained by correcting the transition energies. Conversely, the bottom panel is characterized by a total absence of theoretical peaks unless the anharmonic contributions to the intensities are included. Indeed, in the spectral window corresponding to the bottom panel, only overtones and combination bands have been detected experimentally, while only fundamental bands have

Table 5. Comparison of Experimental and Computed Anharmonic VPT2, DVPT2, and GVPT2 Wavenumbers (in cm^{-1}) of CF_3Br^a

| state | symm. | MP2//MP2 ^b | | | | CC//MP2 ^c | | | | exp. ^{e,f,g} |
|------------------------|-------|-----------------------|---------------------|----------------------|----------------------|----------------------|---------------------|----------------------|----------------------|-----------------------|
| | | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω^d | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | |
| $ 1_1\rangle$ | A_1 | 1101 | 1080 | 1080 | 1080 | 1104 | 1082 | 1091 | 1086 | 1085 |
| $ 1_2\rangle$ | | 770 | 764 | 764 | 764 | 763 | 757 | 757 | 757 | 761 |
| $ 1_3\rangle$ | | 363 | 361 | 361 | 361 | 353 | 351 | 351 | 351 | 350 |
| $ 1_4, \pm 1_4\rangle$ | E | 1221 | 1196 | 1196 | 1196 | 1230 | 1206 | 1206 | 1206 | 1209 |
| $ 1_5, \pm 1_5\rangle$ | | 553 | 547 | 547 | 547 | 548 | 542 | 542 | 542 | 550 |
| $ 1_6, \pm 1_6\rangle$ | | 309 | 307 | 307 | 307 | 304 | 302 | 302 | 302 | 305 |
| $ 1_2 1_3\rangle$ | | 1132 | 1132 | 1132 | 1132 | 1116 | 1119 | 1110 | 1115 | 1120 |
| MAE | | | 7 | 7 | 7 | | 3 | 4 | 4 | |

^aThe polar vibrational states are indicated as $|v_i, l_i\rangle$. ^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set. ^cAnharmonic calculations performed at the MP2/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/aug-cc-pVTZ-PP level. ^dReference 130. ^eReference 131. ^fReference 132. ^gReference 133.

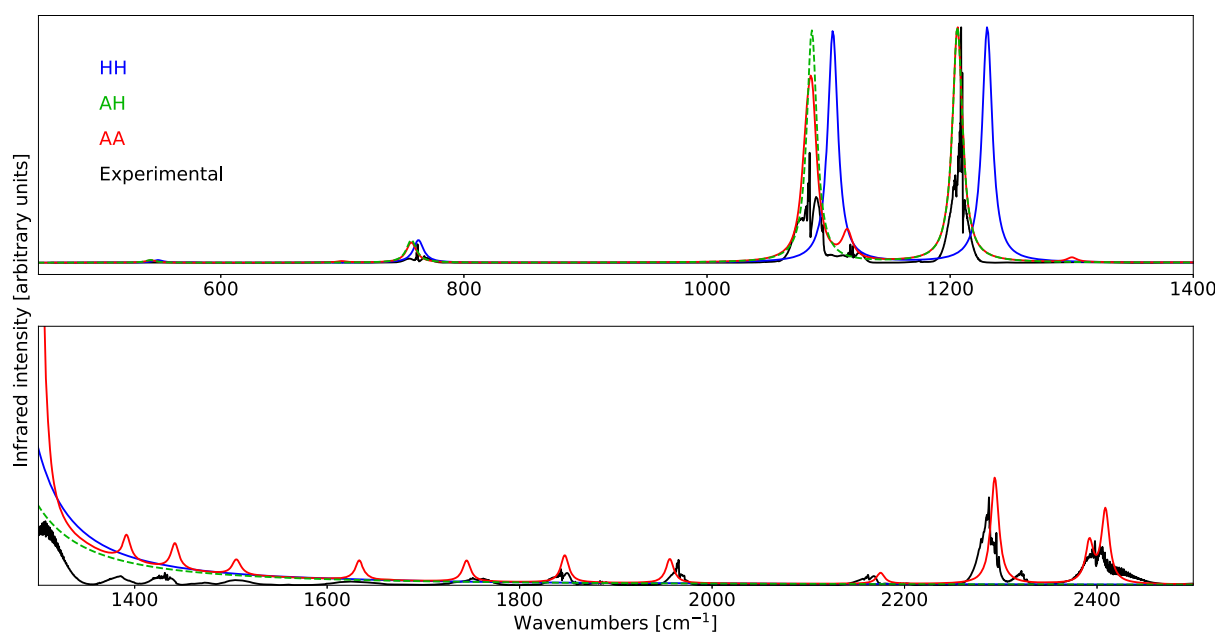


Figure 5. Comparison of the computed IR spectrum of CF_3Br at the hybrid CC//MP2 level of theory with the experimental data. The labels HH and AA indicate respectively the full harmonic and anharmonic spectra, while the label AH indicates the inclusion of anharmonic effects only for the correction of the energies. The spectral range has been segmented to highlight the structure in the region above 1400 cm^{-1} , with the panel below reporting the spectrum scaled by a factor 100 with respect to the above one. Spectral line shapes have been convoluted by Lorentzian distribution functions with HWHMs of 5 cm^{-1} . Experimental IR spectrum from ref 133. All spectra are normalized by setting the intensity of their highest peak to unity.

non-vanishing intensities within the harmonic oscillator model. The anharmonic corrections to the intensities allow not only to bypass this limit but also to produce a spectral profile in full agreement with the experimental one.

With the aim of showing the application of the new framework to the calculation of isotopomers belonging to different point groups, the CH_3D and CHD_3 molecules (symmetric tops) will be compared to the CH_2D_2 isotopomer (asymmetric top), whereas the spherical top isotopomers (CH_4 and CD_4) will be analyzed in a later section. In this context, four hybrid schemes have been used for computing the wavenumbers of the systems under consideration, namely, the MP2//MP2, B2D3//B3D3, CC//MP2, and CC//B3D3 models in conjunction with the JnDZ (B3D3 and MP2), JnTZ (B2D3), and cc-pVQZ (CC) basis sets.¹³⁴ A full comparison of all sets of theoretical data with the corresponding experimental counterparts is reported in Table 6.

Concerning CH_2D_2 , two Fermi resonances, namely, $\omega_2 \approx 2\omega_7$ and $\omega_8 \approx \omega_4 + \omega_9$, have been found at all levels of calculation, with an additional one ($\omega_1 \approx 2\omega_3$) detected at the B2D3//B3D3 level. Conversely, 1–1 and 2–2 Darling–Dennison resonances have not been identified. The resonance analysis carried out for CH_3D shows the presence of a Fermi resonance of type I ($\omega_1 \approx 2\omega_5$) and a 1–1 Darling–Dennison resonance ($\omega_1 \approx \omega_4$) at all the computational levels employed here. A second Fermi resonance of type I ($\omega_2 \approx 2\omega_6$) is also present at the MP2//MP2 level. Furthermore, a series of 2–2 Darling–Dennison interactions have been detected (including l -type terms). Finally, CHD_3 is consistently characterized by two Fermi resonances ($\omega_2 \approx 2\omega_3$ and $\omega_4 \approx \omega_3 + \omega_5$), with the addition of $\omega_4 \approx 2\omega_6$ at the CC//MP2 level. While 1–1 Darling–Dennison resonances have not been found, different 2–2 Darling–Dennison resonances have been included at the variational level. Let us remark that in agreement with the analysis performed by

Table 6. Comparison of Experimental and Computed Anharmonic Fundamental VPT2, DVPT2, and GVPT2 Wavenumbers (in cm^{-1}) of CH_2D_2 , CH_3D , and CHD_3 ^a

| symm. | MP2//MP2 ^b | | | B2D3//B3D3 ^c | | | CC//MP2 ^d | | | CC//B3D3 ^e | | | exp. ^f |
|------------------|-------------------------|---------------------|----------------------|-------------------------|---------------------|----------------------|----------------------|---------------------|----------------------|-----------------------|---------------------|----------------------|-------------------|
| | ω | ν_{VPT2} | ν_{DVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | |
| | CH_2D_2 | | | | | | | | | | | | |
| $ 1_1\rangle$ | 3148 | 3017 | 3017 | 3017 | 2986 | 2973 | 2994 | 2972 | 2950 | 2975 | 2961 | 2982 | 2975 |
| $ 1_2\rangle$ | *2268 | 2249 | 2197 | 2230 | 4174 | 2178 | 2153 | 2133 | 2446 | 2430 | 2169 | 2142 | 2203 |
| $ 1_3\rangle$ | 1483 | 1449 | 1449 | 1449 | 1445 | 1445 | 1445 | 1436 | 1436 | 1440 | 1440 | 1440 | 1435 |
| $ 1_4\rangle$ | 1060 | 1040 | 1040 | 1040 | 1058 | 1041 | 1041 | 1034 | 1034 | 1036 | 1036 | 1036 | 1033 |
| $ 1_5\rangle$ | 1373 | 1346 | 1346 | 1346 | 1364 | 1339 | 1339 | 1332 | 1332 | 1335 | 1335 | 1335 | 1331 |
| $ 1_6\rangle$ | 3208 | 3060 | 3060 | 3060 | 3163 | 3018 | 3018 | 2999 | 2999 | 3009 | 3009 | 3009 | 3012 |
| $ 1_7\rangle$ | 1122 | 1098 | 1098 | 1098 | 1122 | 1101 | 1101 | 1092 | 1092 | 1095 | 1095 | 1095 | 1091 |
| $ 1_8\rangle$ | *2376 | 2305 | 2287 | 2312 | *2342 | 2257 | 2241 | 2225 | 2285 | *2337 | 2290 | 2233 | 2235 |
| $ 1_9\rangle$ | 1272 | 1243 | 1243 | 1243 | 1273 | 1248 | 1248 | 1237 | 1237 | 1265 | 1240 | 1240 | 1236 |
| MAE ^g | 26 | 24 | 24 | 27 | 19 | 10 | 10 | 4 | 10 | 6 | 6 | 4 | 4 |
| | CH_3D | | | | | | | | | | | | |
| $ 1_1\rangle$ | 3113 | 3004 | 2977 | 2931 | 2984 | 2952 | 2995 | 2978 | 2962 | 2971 | 2934 | 2983 | 2970 |
| $ 1_2\rangle$ | 2320 | 2220 | 2237 | 2226 | 2291 | 2201 | 2201 | 2183 | 2183 | 2285 | 2192 | 2192 | 2220 |
| $ 1_3\rangle$ | 1347 | 1314 | 1314 | 1314 | 1347 | 1318 | 1318 | 1307 | 1307 | 1340 | 1310 | 1310 | 1307 |
| $ 1_4\rangle$ | 3208 | 3064 | 3064 | 3064 | 3163 | 3022 | 3024 | 3003 | 3003 | 3157 | 3013 | 3015 | 3017 |
| $ 1_5\rangle$ | 1522 | 1488 | 1488 | 1488 | 1513 | 1480 | 1480 | 1473 | 1473 | 1508 | 1475 | 1475 | 1471 |
| $ 1_6\rangle$ | 1195 | 1169 | 1169 | 1169 | 1194 | 1170 | 1170 | 1162 | 1162 | 1188 | 1164 | 1164 | 1161 |
| MAE | 19 | 14 | 14 | 20 | 8 | 9 | 10 | 4 | 10 | 3 | 8 | 4 | 4 |
| | CHD_3 | | | | | | | | | | | | |
| $ 1_1\rangle$ | 3179 | 3035 | 3035 | 3035 | 3138 | 2997 | 2997 | 2978 | 2978 | 3131 | 2987 | 2987 | 2992 |
| $ 1_2\rangle$ | *2220 | 2170 | 2157 | 2171 | *2201 | 2157 | 2143 | 2140 | 2142 | 2191 | 2132 | 2147 | 2143 |
| $ 1_3\rangle$ | 1031 | 1010 | 1010 | 1010 | 1031 | 1012 | 1012 | 1005 | 1005 | 1025 | 1007 | 1007 | 1003 |
| $ 1_4\rangle$ | *2376 | 2320 | 2288 | 2277 | *2342 | 2234 | 2252 | 2239 | 2202 | 2337 | 2212 | 2253 | 2251 |
| $ 1_5\rangle$ | 1332 | 1304 | 1304 | 1304 | 1326 | 1301 | 1301 | 1293 | 1293 | 1321 | 1295 | 1295 | 1291 |
| $ 1_6\rangle$ | 1062 | 1043 | 1043 | 1043 | 1060 | 1042 | 1042 | 1036 | 1036 | 1056 | 1037 | 1037 | 1036 |
| MAE | 23 | 18 | 18 | 16 | 8 | 6 | 5 | 4 | 11 | 8 | 3 | 3 | 3 |

^aMean absolute errors (MAEs) are also reported. The polar vibrational states are indicated as $|i_j\rangle$. The frequencies impacted by resonances are indicated with a “*”. ^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set. ^cAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the B2D3/JnTZ level. ^dAnharmonic calculations performed at the MP2/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/cc-pVQZ level. ^eAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/cc-pVQZ level. ^fReference 135. ^gState $|1_2\rangle$ excluded.

Table 7. Comparison of Extrapolated and Computed Anharmonic VPT2 and DVPT2 χ^p and g Diagonal Elements (in cm^{-1}) of CH_2D_2 , CH_3D , and CHD_3 ^a

| | MP2//MP2 ^b | | B2D3//B3D3 ^c | | CC//MP2 ^d | | CC//B3D3 ^e | | CC. ^f | exp. ^g |
|------------------------------------|-----------------------|----------------------|-------------------------|----------------------|----------------------|----------------------|-----------------------|----------------------|------------------|-------------------|
| | ν_{VPT2} | ν_{DVPT2} | ν_{VPT2} | ν_{DVPT2} | ν_{VPT2} | ν_{DVPT2} | ν_{VPT2} | ν_{DVPT2} | | |
| CH₂D₂ | | | | | | | | | | |
| χ_{11}^c | -28.1 | -28.1 | -27.3 | -27.3 | -30.7 | -30.7 | -27.9 | -27.9 | -27.3 | -26.4 |
| χ_{22}^c | -14.4 | -14.4 | -14.1 | -14.1 | -15.7 | -15.7 | -14.4 | -14.4 | -14.1 | -13.5 |
| χ_{33}^c | -7.5 | -7.5 | -7.2 | -0.5 | -8.6 | -1.3 | -7.3 | -0.5 | -6.7 | -9.1 |
| χ_{44}^c | -5.2 | -5.2 | -5.0 | -5.0 | -6.0 | -6.0 | -5.0 | -5.0 | -4.5 | -7.1 |
| χ_{55}^c | -2.0 | -2.0 | -1.9 | -1.9 | -2.2 | -2.2 | -2.0 | -2.0 | -2.2 | -3.6 |
| χ_{66}^c | -32.2 | -32.2 | -31.5 | -31.5 | -35.2 | -35.2 | -32.1 | -32.1 | -31.6 | -32.7 |
| χ_{77}^c | -28.2 | -2.3 | -999.2 | -1.7 | -144.7 | -2.4 | -132.3 | -1.7 | -1.9 | 8.9 |
| χ_{88}^c | -18.8 | -18.8 | -18.5 | -18.5 | -20.5 | -20.5 | -18.9 | -18.9 | -18.6 | -19.3 |
| χ_{99}^c | -5.2 | -5.2 | -4.4 | -4.4 | -5.8 | -5.8 | -4.5 | -4.5 | -4.4 | -13.4 |
| MAE ^h | 3.4 | 0.5 | 111.0 | 0.8 | 17.6 | 2.2 | 14.8 | 1.0 | | |
| MAE ⁱ | 6.0 | 3.1 | 114.0 | 3.9 | 19.4 | 4.4 | 17.7 | 3.9 | | |
| CH₃D | | | | | | | | | | |
| χ_{11}^p | -17.8 | -17.8 | -17.2 | -17.2 | -19.3 | -19.3 | -17.6 | -17.6 | -17.2 | -15.9 |
| χ_{22}^p | -31.5 | -31.5 | -31.4 | -31.4 | -34.6 | -34.6 | -31.9 | -31.9 | -31.2 | -31.0 |
| χ_{33}^p | -7.5 | -7.5 | -6.9 | -6.9 | -8.2 | -8.2 | -7.0 | -7.0 | -6.8 | -20.3 |
| χ_{44}^p | -32.1 | -32.1 | -31.5 | -31.5 | -35.1 | -35.1 | -32.2 | -32.2 | -31.6 | -32.6 |
| χ_{55}^p | -8.4 | -1.7 | -9.4 | -1.5 | -10.5 | -1.8 | -9.8 | -1.5 | -1.9 | -19.3 |
| χ_{66}^p | 1.5 | -2.8 | 0.8 | 0.8 | 0.5 | 0.5 | 1.0 | 1.0 | 0.3 | -6.1 |
| g_{44} | 12.6 | 12.6 | 12.7 | 12.7 | 13.8 | 13.8 | 12.9 | 12.9 | 12.6 | 13.3 |
| g_{55} | 6.9 | 0.2 | 7.8 | -0.1 | 8.7 | 0.4 | 8.2 | -0.1 | 0.4 | 15.9 |
| g_{66} | -1.0 | 3.3 | -0.1 | -0.1 | 0.2 | 0.2 | -0.3 | -0.3 | 0.7 | 3.3 |
| MAE ^h | 2.0 | 0.9 | 1.9 | 0.3 | 3.2 | 1.4 | 2.2 | 0.5 | | |
| MAE ⁱ | 5.3 | 5.9 | 5.0 | 6.8 | 5.2 | 7.1 | 4.9 | 6.7 | | |
| CHD₃ | | | | | | | | | | |
| χ_{11}^p | -60.4 | -60.4 | -59.7 | -59.7 | -66.3 | -66.3 | -60.6 | -60.6 | -59.2 | -59.5 |
| χ_{22}^p | -9.1 | -9.1 | -8.8 | -8.8 | -9.8 | -9.8 | -9.1 | -9.1 | -8.8 | -8.1 |
| χ_{33}^p | -7.4 | -0.8 | -7.5 | -0.4 | -8.5 | -0.8 | -7.6 | -0.4 | -7.2 | -18.1 |
| χ_{44}^p | -18.9 | -18.9 | -18.6 | -18.6 | -20.6 | -20.6 | -19.0 | -19.0 | -18.6 | -19.3 |
| χ_{55}^p | -5.0 | -5.0 | -4.4 | -4.4 | -5.7 | -5.7 | -4.5 | -4.5 | -4.6 | -7.4 |
| χ_{66}^p | -5.9 | -5.9 | -6.0 | -6.0 | -6.9 | -2.2 | -6.0 | -6.0 | -5.5 | -12.4 |
| g_{44} | 8.2 | 8.2 | 8.3 | 8.3 | 9.0 | 9.0 | 8.4 | 8.4 | 8.3 | 8.7 |
| g_{55} | 5.7 | 5.7 | 5.5 | 5.5 | 6.2 | 6.2 | 5.5 | 5.5 | 5.4 | 6.7 |
| g_{66} | 3.3 | 3.3 | 3.3 | 3.3 | 4.0 | -0.7 | 3.4 | 3.4 | 3.3 | 7.7 |
| MAE ^h | 0.4 | 1.1 | 0.2 | 1.0 | 1.8 | 3.0 | 0.4 | 1.1 | | |
| MAE ⁱ | 3.1 | 3.8 | 3.1 | 3.9 | 3.5 | 5.4 | 3.1 | 3.9 | | |

^aMean absolute errors (MAEs) are also reported. ^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set. ^cAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the B2D3/JnTZ level. ^dAnharmonic calculations performed at the MP2/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/cc-pVQZ level. ^eAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/cc-pVQZ level. ^fTheoretical values obtained from CCSD(T)/cc-pVTZ anharmonic force constants in conjunction with harmonic frequencies at the CCSD(T)/cc-pVQZ level (see Tables V and VI of ref 134 for more details) and reported with one decimal place. ^gValues from ref 136 and reported with one decimal place. ^hComputed with respect to the reference theoretical (CC) values. ⁱComputed with respect to the experimental data.

Lee and co-workers,¹³⁴ the state $|1_1\rangle$ of CH_3D is strongly coupled with $|2_{7,0_7}\rangle$ (but not with $|2_{7,\pm 2_7}\rangle$ since the coupling element vanishes based on eq 38) so that at the GVPT2 level, the states with energies (see Table 6) 2931, 2995, 2978, and 2983 cm^{-1} for MP2//MP2, B2D3//B3D3, CC//MP2, and CC//B3D3, respectively, and those with energies 3017, 2911, 2890, and 2899 cm^{-1} are basically equal mixtures of $|1_1\rangle$ and $|2_{7,0_7}\rangle$.

As expected, the results closer to those of the experiment are those based on CC harmonic frequencies, although the B2D3//B3D3 scheme leads to quite satisfactory results. Besides, the results presented here are in good agreement with those reported in ref 134, as confirmed by a comparison of the χ^p and g

matrices, for which only the diagonal elements are reported in Table 7 for readability.

The next case studies are two planar aromatic systems, namely, the cyclopentadienyl anion and benzene (see Figures 6 and 7), with the former being noted Cp^- in the following.

As already anticipated, the presence of out-of-plane vibrations makes the use of a triple- ζ basis set mandatory for the calculation of the anharmonic force field so that the MP2/JnTZ level of theory has been employed for both systems. The assignment of the normal vibrations of the cyclopentadienyl anion has been recently revisited by Bencze and co-workers¹³⁷ on the basis of previous studies.¹³⁸ As outlined in ref 137, the cyclopentadienyl anion cannot exist without a counter cation under normal

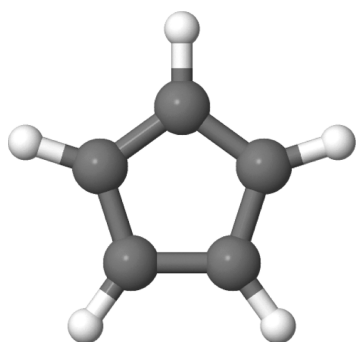


Figure 6. Molecular structure of the cyclopentadienyl anion.

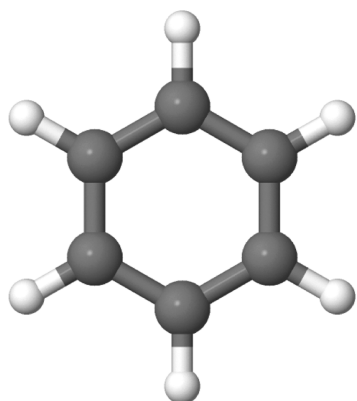


Figure 7. Molecular structure of benzene.

experimental conditions of IR and Raman spectroscopies. By means of a detailed analysis, the authors showed that the structural parameters of the Cp^- ring in the solid-state CpK should be close to those of the hypothetical free anion employed in the QC computations. Furthermore, the experimental frequencies of CpLi and CpNa in tetrahydrofuran (THF) were found to be very close to those of solid CpK. Experimental

and theoretical anharmonic fundamental frequencies of Cp^- are reported in Table 8.

The results reported in Table 8 show that the VPT2 results are in good agreement with the experiment and that inclusion of the variational correction improves the agreement. The largest discrepancies concern the $|1_{12}\rangle$ and $|1_{13,\pm 1_{13}}\rangle$ states. There are, of course, systematic shifts related to the difference between the structure of the cyclopentadienyl anion in the experimental complexes and that of the free anion, with the lowest frequencies being, as usual, the most sensitive to environmental effects.

The next molecule studied is benzene (see Figure 7), a D_{6h} symmetric-top system, which has been extensively studied by both IR and Raman spectroscopies.^{30,140–143}

The anharmonic calculations have been carried out at the MP2/JnTZ level or coupling the MP2/JnTZ anharmonic contributions to harmonic frequencies evaluated at the CCSD-(T)/ANO4321' level.¹⁴⁴ The VPT2, DVPT2, and GVPT2 fundamentals of benzene computed at both levels of theory are compared with experimental data in Table 9.

Concerning the GVPT2 scheme, Fermi resonances of type II ($\omega_i \approx \omega_j + \omega_k$) as well as 2–2 DD resonances, were found, with the latter including l -type doubling of types R and S. On the other hand, 1–1 DD resonances have not been identified. The use of the hybrid scheme leads to a remarkable improvement of the fundamentals, and this is particularly true for the state $|1_{14}\rangle$, with the corresponding MP2/JnTZ harmonic frequency being clearly overestimated. On top of this, the inclusion of the variational correction further improves the agreement with the experimental data for all the transitions involved in resonances. The IR spectrum evaluated by means of the CC//MP2 hybrid scheme is compared with that of the experiment in Figure 8.

All spectra show a very strong band between 650 and 700 cm^{-1} corresponding to the fundamental transition to $|1_{12}\rangle$, associated to the out-of-plane bending vibration sketched in Figure 9.

Since such a state is not involved in any resonance, its position is independent of the adopted VPT2 scheme.

Table 8. Comparison of Experimental and Computed Anharmonic Fundamental VPT2, DVPT2, and GVPT2 Wavenumbers (cm^{-1}) of the Cyclopentadienyl Anion^a

| state | symm. | MP2 ^b | | | | exp. | | |
|-----------------------------|---------|------------------|---------------------|----------------------|----------------------|-------------------|-------------------|------------------|
| | | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | CpLi ^c | CpNa ^d | CpK ^e |
| $ 1_1\rangle$ | A'_1 | 3218 | 3087 | 3087 | 3087 | 3104 | 3090 | 3088 |
| $ 1_2\rangle$ | | 1140 | 1118 | 1118 | 1118 | 1114 | 1114 | 1119 |
| $ 1_3\rangle$ | A'_2 | 1262 | 1237 | 1237 | 1237 | | | 1260 |
| $ 1_{4,\pm 1_4}\rangle$ | A''_2 | 663 | 652 | 652 | 652 | 710 | 722 | 686 |
| $ 1_{5,\pm 1_5}\rangle$ | E'_1 | 3196 | 3066 | 3066 | 3068 | 3082 | 3067 | 3061 |
| $ 1_{6,\pm 1_6}\rangle$ | | 1448 | 1415 | 1415 | 1415 | 1433 | | 1440 |
| $ 1_{7,\pm 1_7}\rangle$ | | 1014 | 996 | 996 | 996 | 1006 | 998 | 1008 |
| $ 1_{8,\pm 1_8}\rangle$ | E'_2 | 3171 | 3043 | 3043 | 3042 | 3080 | 3060 | 3068 |
| $ 1_{9,\pm 1_9}\rangle$ | | 1420 | 1612 | 1382 | 1375 | 1346 | 1342 | 1370 |
| $ 1_{10,\pm 1_{10}}\rangle$ | | 1061 | 1043 | 1043 | 1043 | 1067 | 1062 | 1070 |
| $ 1_{11,\pm 1_{11}}\rangle$ | | 830 | 819 | 819 | 819 | 854 | 848 | 854 |
| $ 1_{12}\rangle$ | E''_1 | 638 | 638 | 638 | 638 | 759 | 730 | 719 |
| $ 1_{13,\pm 1_{13}}\rangle$ | E''_2 | 780 | 787 | 787 | 787 | 735 | 722 | 686 |
| $ 1_{14,\pm 1_{14}}\rangle$ | | 616 | 612 | 612 | 612 | | | 600 |
| MAE ^f | | | 40 | 23 | 22 | | | |

^aMean absolute errors (MAEs) are also reported. The polar vibrational states are indicated as $|v_i, l_i\rangle$. ^bBasis set: JnTZ. ^cSolution of CpLi dissolved in THF (ref 139). ^dSolution of CpNa dissolved in THF (ref 139). ^eSolid CpK (refs 137 and 139). ^fMAE computed with respect to the experimental values of CpK, excluding the states $|1_{13,\pm 1_{13}}\rangle$.

Table 9. Comparison of Experimental and Computed Anharmonic Fundamental VPT2, DVPT2, and GVPT2 Wavenumbers (in cm^{-1}) of Benzene^a

| state | symm. | MP2 ^b | | | | CC//MP2 ^b | | | | exp. ^c |
|-----------------------------|----------|------------------|---------------------|----------------------|----------------------|----------------------|---------------------|----------------------|----------------------|-------------------|
| | | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | |
| $ 1_1\rangle$ | A_{1g} | 3235 | 3091 | 3107 | 3101 | 3210 | 3072 | 3072 | 3072 | 3074 |
| $ 1_2\rangle$ | | 1011 | 997 | 997 | 997 | 1003 | 988 | 988 | 988 | 993 |
| $ 1_3\rangle$ | A_{2g} | 1371 | 1338 | 1350 | 1346 | 1380 | 1348 | 1359 | 1355 | (1350) |
| $ 1_4\rangle$ | B_{2g} | 996 | 1019 | 1019 | 1019 | 1009 | 1030 | 1030 | 1030 | (990) |
| $ 1_5\rangle$ | | 710 | 709 | 709 | 709 | 709 | 708 | 708 | 708 | (707) |
| $ 1_{6,\pm 1_6}\rangle$ | E_{1g} | 864 | 858 | 858 | 858 | 865 | 858 | 858 | 858 | 847 |
| $ 1_{7,\pm 1_7}\rangle$ | E_{2g} | 3207 | 3084 | 3084 | 3084 | 3183 | 3061 | 3061 | 3061 | 3057 |
| $ 1_{8,\pm 1_8}\rangle$ | | 1636 | 1601 | 1601 | 1601 | 1637 | 1600 | 1600 | 1600 | 1601 |
| $ 1_{9,\pm 1_9}\rangle$ | | 1195 | 1180 | 1180 | 1180 | 1194 | 1179 | 1179 | 1179 | 1178 |
| $ 1_{10,\pm 1_{10}}\rangle$ | | 606 | 603 | 603 | 603 | 611 | 609 | 609 | 609 | 608 |
| $ 1_{11}\rangle$ | A_{2u} | 691 | 683 | 683 | 683 | 687 | 678 | 678 | 678 | 674 |
| $ 1_{12}\rangle$ | B_{1u} | 3195 | 3106 | 3072 | 3026 | 3173 | 3117 | 3047 | 3047 | (3057) |
| $ 1_{13}\rangle$ | | 1019 | 1014 | 1014 | 1014 | 1020 | 1015 | 1015 | 1015 | (1010) |
| $ 1_{14}\rangle$ | B_{2u} | 1460 | 1418 | 1418 | 1418 | 1326 | 1288 | 1288 | 1288 | 1309 |
| $ 1_{15}\rangle$ | | 1169 | 1156 | 1156 | 1156 | 1163 | 1149 | 1149 | 1149 | 1150 |
| $ 1_{16,\pm 1_{17}}\rangle$ | E_{1u} | 3224 | 3111 | 3111 | 3111 | 3200 | 3089 | 3073 | 3045 | 3047 |
| $ 1_{17,\pm 1_{17}}\rangle$ | | 1503 | 1476 | 1476 | 1476 | 1509 | 1481 | 1481 | 1481 | 1484 |
| $ 1_{18,\pm 1_{18}}\rangle$ | | 1060 | 1040 | 1040 | 1040 | 1056 | 1035 | 1035 | 1035 | 1038 |
| $ 1_{19,\pm 1_{19}}\rangle$ | E_{2u} | 977 | 978 | 978 | 978 | 985 | 985 | 985 | 985 | 976 |
| $ 1_{20,\pm 1_{20}}\rangle$ | | 404 | 401 | 401 | 401 | 406 | 403 | 403 | 403 | 398 |
| MAE | | | 18 | 17 | 17 | | 11 | 8 | 7 | |

^aThe polar vibrational states are indicated as $|l_i, l_j\rangle$. ^bBasis set: JnTZ. ^cReference 30, the values in parentheses have not been observed directly but have been deduced from combination bands.

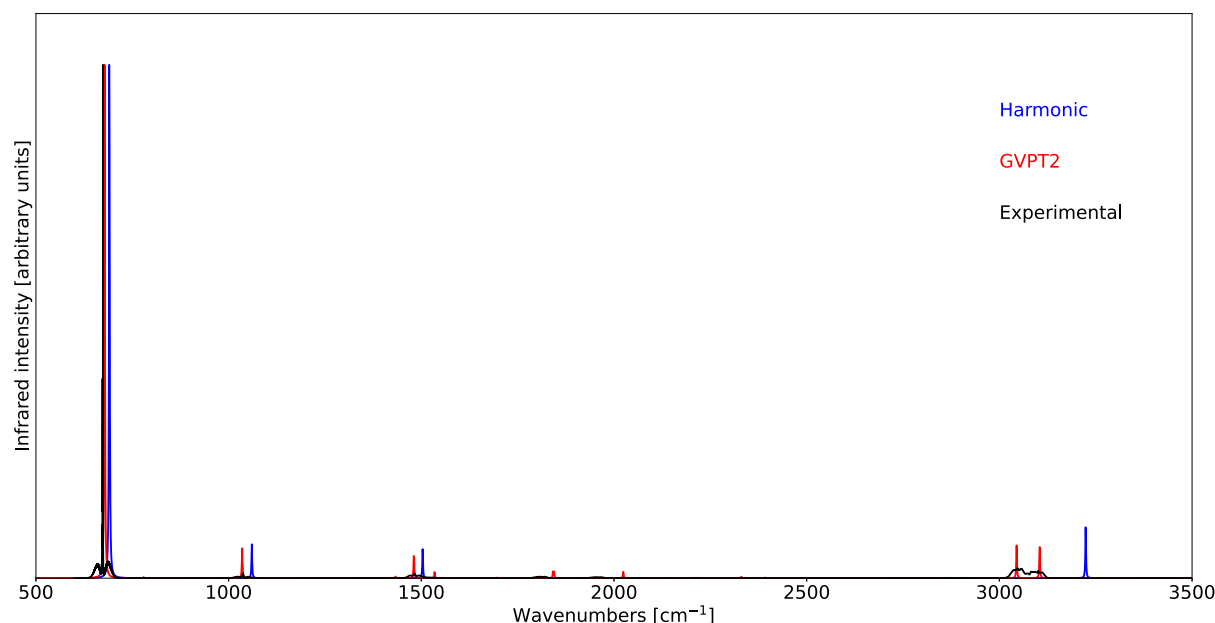


Figure 8. Comparison of the computed harmonic and anharmonic (GVPT2) IR spectra of benzene at the MP2/JnTZ level of theory with those of the experiment. Spectral line shapes have been convoluted by Lorentzian distribution functions with HWHMs of 1 cm^{-1} . The experimental IR spectrum is from refs 145 and 146. All spectra are normalized by setting the intensity of their highest peak to unity.

Finally, the simulation of the IR spectrum of pentaborane (see Figure 10), a C_{4v} system, has been carried out.

Among the systems considered in the present study, pentaborane is the only one showing all types of l -type doubling within the GVPT2 scheme since the order of its principal axis is a multiple of 4. The simulation has been performed by employing the same hybrid scheme as for the other systems, namely, by coupling MP2/JnDZ anharmonic contributions to MP2/JnTZ

harmonic frequencies. A full comparison of the theoretical and experimental fundamentals is reported in Table 10.

At the DVPT2 level, several Fermi resonances (most of type II) have been detected, which have been successively included at the variational level within the GVPT2 scheme, together with the proper l -type doubling terms and the other identified 2–2 Darling–Dennison resonances. Most of the frequencies are qualitatively correct and consistent with those reported in a

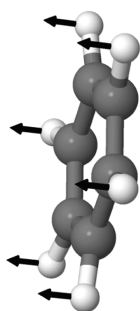


Figure 9. Vibration associated to the strongest band of the IR spectrum of benzene.

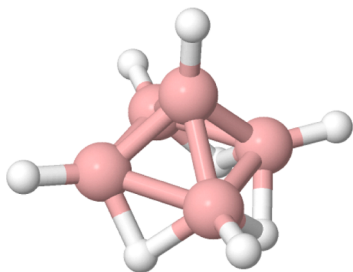


Figure 10. Molecular structure of pentaborane.

Table 10. Comparison of Experimental and Computed (VPT2, DVPT2, and GVPT2) Anharmonic Fundamental Wavenumbers (in cm^{-1}) of Pentaborane

| state | symm. | MP2 ^a | | | | Exp. |
|------------------------------|-------|------------------|---------------------|----------------------|----------------------|-------------------|
| | | ω | ν_{VPT2} | ν_{DVPT2} | ν_{GVPT2} | |
| $ 1_1\rangle$ | A_1 | 2773 | 2675 | 2675 | 2677 | 2628 ^b |
| $ 1_2\rangle$ | | 2755 | 2657 | 2657 | 2651 | 2610 ^b |
| $ 1_3\rangle$ | | 2026 | 1865 | 1852 | 1852 | 1844 ^b |
| $ 1_4\rangle$ | | 1167 | 1125 | 1125 | 1125 | 1126 ^b |
| $ 1_5\rangle$ | | 1011 | 981 | 981 | 981 | 985 ^c |
| $ 1_6\rangle$ | | 825 | 801 | 801 | 801 | 799 ^b |
| $ 1_7\rangle$ | | 725 | 709 | 709 | 709 | 701 ^b |
| $ 1_8\rangle$ | A_2 | 1523 | 1361 | 1373 | 1372 | 1450 ^c |
| $ 1_9\rangle$ | | 883 | 849 | 849 | 849 | |
| $ 1_{10}\rangle$ | B_1 | 1987 | 1721 | 1839 | 1787 | 1870 ^d |
| $ 1_{11}\rangle$ | | 1038 | 1010 | 1010 | 1010 | 1036 ^b |
| $ 1_{12}\rangle$ | | 784 | 752 | 752 | 752 | 741 ^b |
| $ 1_{13}\rangle$ | | 620 | 604 | 604 | 604 | 599 ^b |
| $ 1_{14}\rangle$ | B_2 | 2742 | 2640 | 2640 | 2639 | 2610 ^b |
| $ 1_{15}\rangle$ | | 1718 | 1554 | 1568 | 1565 | 1500 ^c |
| $ 1_{16}\rangle$ | | 812 | 792 | 792 | 792 | 785 ^b |
| $ 1_{17}\rangle$ | | 729 | 706 | 706 | 705 | |
| $ 1_{18}\rangle$ | | 491 | 473 | 473 | 473 | 470 ^d |
| $ 1_{19}, \pm 1_{19}\rangle$ | E | 2751 | 2649 | 2649 | 2649 | 2610 ^b |
| $ 1_{20}, \pm 1_{20}\rangle$ | | 1988 | 1817 | 1851 | 1812 | 1634 ^b |
| $ 1_{21}, \pm 1_{21}\rangle$ | | 1585 | 1467 | 1440 | 1436 | 1410 ^b |
| $ 1_{22}, \pm 1_{22}\rangle$ | | 1086 | 1101 | 1047 | 1041 | 1035 ^b |
| $ 1_{23}, \pm 1_{23}\rangle$ | | 958 | 927 | 927 | 927 | 918 ^b |
| $ 1_{24}, \pm 1_{24}\rangle$ | | 907 | 886 | 886 | 886 | 890 ^b |
| $ 1_{26}, \pm 1_{26}\rangle$ | | 649 | 624 | 624 | 624 | 618 ^b |
| $ 1_{27}, \pm 1_{27}\rangle$ | | 595 | 576 | 576 | 576 | 569 ^b |
| MAE | | | 30 | 21 | 22 | |

^aMean absolute error (MAE) does not include state $|1_{20}, \pm 1_{20}\rangle$.

^bReference 147. ^cReference 148. ^dReference 149.

recent study performed by Maillard and co-workers¹⁵⁰ for several B–H systems (including, for instance, the out-of-scale discrepancy characterizing the states $|1_{20}, \pm 1_{20}\rangle$). For the sake of completeness, the IR spectrum of B_5H_9 is reported in Figure 11, where the theoretical data obtained by means of the VPT2, DVPT2, and GVPT2 schemes are compared.

While DVPT2 and GVPT2 spectra show a common pattern, the atypical behavior of the VPT2 spectrum is due to the strong Fermi resonances between the fundamentals $|1_{22}, \pm 1_{22}\rangle$ and the combination bands $|1_{18}1_{27}, \pm 1_{27}\rangle$, which lead to a huge value for the transition dipole moment of the $|1_{22}, \pm 1_{22}\rangle$ states so that the corresponding band is the only one clearly visible in the theoretical spectrum. This problem is fixed within the DVPT2 scheme through the elimination of the resonant term, and the result is further refined at the GVPT2 level by the successive variational treatment.

4.3. Paving the Route to Spherical Tops. With the aim of showing the extension of our computational framework to spherical tops, we now analyze a series of systems of both tetrahedral and octahedral symmetries, including, for instance, tetraphosphorus (P_4), methane (CH_4) and its fully deuterated isotopologue (CD_4), and sulfur hexafluoride (SF_6), whose structures are sketched in Figure 12.

In the case of linear and symmetric tops, our computational protocol performs by default a full check of all the symmetry relations present between the anharmonic force constants.⁷¹ Such a procedure has not yet been implemented for tetrahedral XY_4 -^{136,151} and octahedral XY_6 -like¹⁵² systems so that some slight (and generally negligible) discrepancies can be detected between anharmonic force constants which should be in principle identical. With the aim of limiting this issue as much as possible, the pruned (99,590) grid employed before for DFT calculations (which are in principle the most sensitive to this problem) will be replaced with a larger one (175,974 and 250,974 for first-row atoms and atoms in the second and later rows, respectively).

4.3.1. Tetrahedral Molecules. The fundamental frequencies of the P_4 molecule obtained through the MP2//MP2 and B2D3//B3D3 hybrid schemes are compared with their experimental counterparts in Table 11.

The P_4 system does not show any Fermi or 1–1 Darling–Dennison resonance so that the values of the fundamentals do not vary going from VPT2 to DVPT2 or GVPT2 schemes. As can be seen from Table 11, the frequencies of the states $|1_1\rangle$ and $|1_2, \pm 1_2\rangle$ are significantly improved within the B2D3//B3D3 scheme, reaching values within 1 cm^{-1} from the experimental counterparts. Concerning the triply degenerate states $|1_1, 1_1, \pm 1_1\rangle$ or $|0_1\rangle$, the best estimate is reached by the MP2//MP2 scheme, even though in both cases, the value of the frequency is not as accurate as the previous ones. The origin of such a discrepancy can be traced back to the experimental conditions at which the gas-phase spectrum has been recorded,¹⁵³ as pointed out by Persson and co-workers¹⁵⁴ in their very detailed vibrational analysis of P_4 . For the sake of completeness and as a consistency check of our calculations, the fundamentals calculated within the MP2//MP2 scheme have been compared with the MP2 results of ref 154, showing good agreement in terms of both harmonic frequencies and anharmonic corrections (see Table III of ref 154 for more details).

A strategy often used to deal with systems presenting degeneracies is that of modifying slightly one or more masses/coordinates in order to lower the symmetry^{134,140} and then

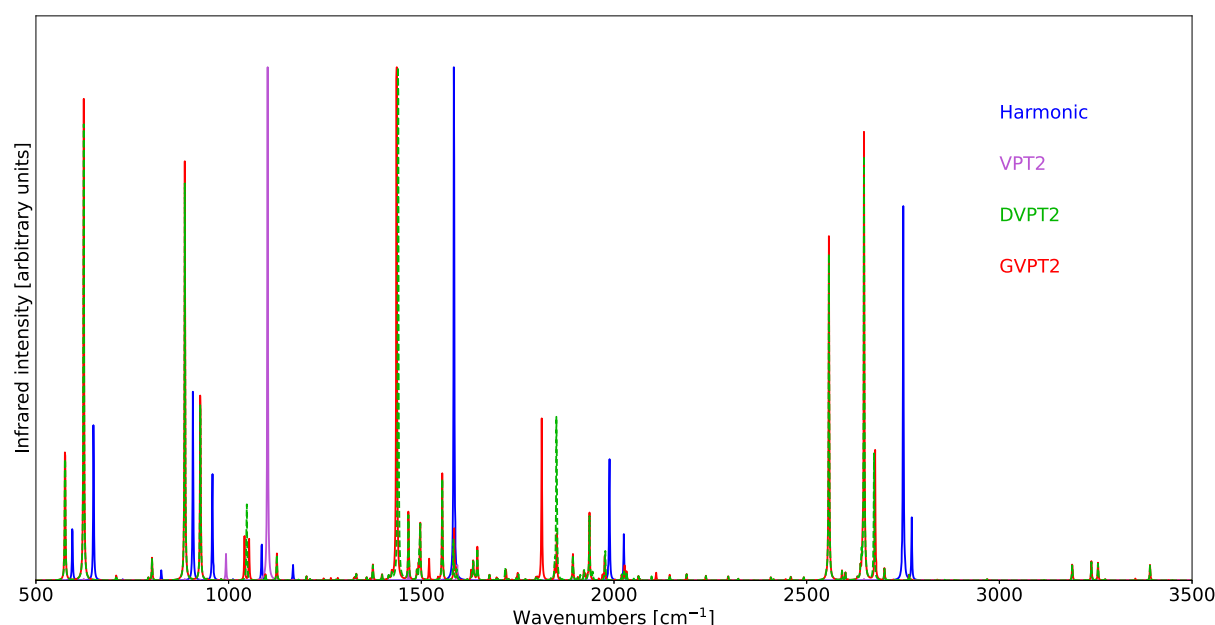


Figure 11. Comparison of the computed harmonic, VPT2, DVPT2, and GVPT2 IR spectra of pentaborane. Spectral line shapes have been convoluted by Lorentzian distribution functions with HWHMs of 1 cm^{-1} . All the spectra are normalized by setting the intensity of their highest peak to unity.

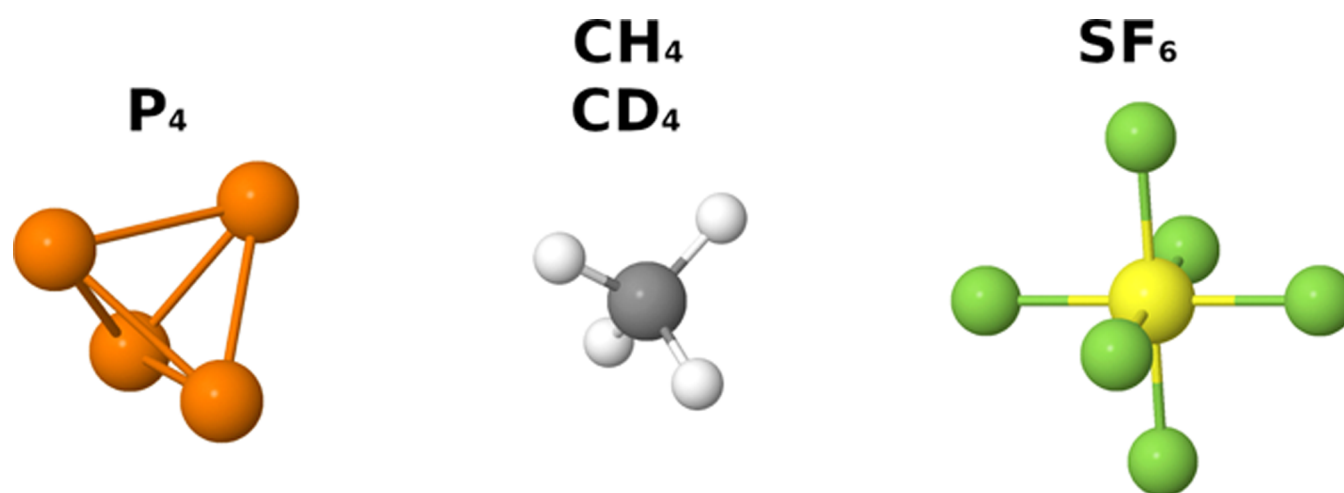


Figure 12. Molecular geometries of methane, tetraphosphorus, and sulfur hexafluoride.

Table 11. Comparison of Experimental and Computed Anharmonic Fundamental VPT2 Wavenumbers (in cm^{-1}) of Tetraphosphorus^a

| | symm. | MP2//MP2 ^b | | B2D3//B3D3 ^c | | exp. ^d |
|---|-------|-----------------------|---------------------|-------------------------|---------------------|-------------------|
| | | ω | ν_{VPT2} | ω | ν_{VPT2} | |
| $ 1_1\rangle$ | A_1 | 613 | 607 | 606 | 601 | 600 |
| $ 1_2, \pm 1_2\rangle$ | E | 365 | 363 | 363 | 361 | 361 |
| $ 1_3, 1_3, \pm 1_3 \text{ or } 0_3\rangle$ | T_2 | 463 | 459 | 458 | 454 | 467 |
| MAE ^e | | | 6 | | 5 | |

^aMean absolute errors (MAEs) are also reported. The spherical and polar vibrational states are indicated respectively as $|v_j, k_j, m_j\rangle$ and $|v_i, l_i\rangle$.
^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set.
^cAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the B2D3/JnTZ level.
^dReference 153.

employ a theoretical model able to treat only lower- or non-degenerate modes. This procedure has been applied to P_4 in order to perform a consistency check of our new implementation. More specifically, the T_d symmetry of this system has been gradually reduced without any geometry modification by slightly increasing (by 0.1%) the masses of a single and then a couple of phosphorous atoms, thus obtaining a symmetric (C_{3v} point group) or an asymmetric (C_{2v} point group) top, respectively. A comparison of the VPT2 fundamentals of the lower-symmetry systems with those of the fully symmetric molecule is reported in Figure 13.

As expected, triply degenerate states $|1_3, 1_3, \pm 1_3 \text{ or } 0_3\rangle$ are no more present, being replaced with a non-degenerate state of A_1 symmetry and a couple of doubly degenerate states belonging to the irreducible representation E in the C_{3v} system and with three non-degenerate states with irreducible representations A_1 , B_1 , and B_2 in the C_{2v} structure. This is clearly visible since the blue bars (C_{3v}) corresponding to the states $|1_3, 1_3, \pm 1_3\rangle$ are equal to each other but different from that of $|1_3, 1_3, 1_0\rangle$, while the

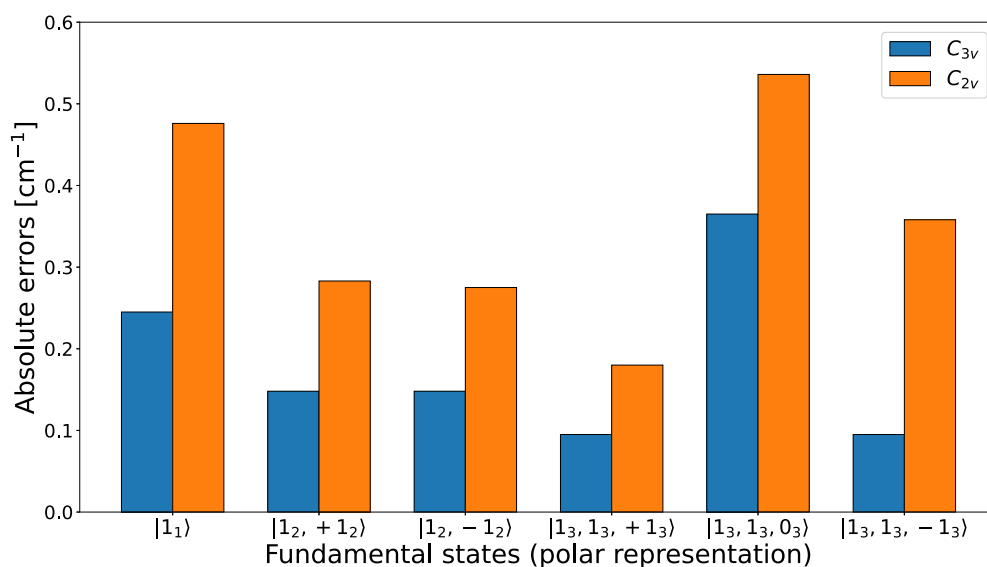


Figure 13. Errors between anharmonic fundamentals of P_4 and the corresponding counterparts of the C_{3v} and C_{2v} symmetry-broken geometries obtained through the MP2//MP2 hybrid scheme.

Table 12. Comparison of Experimental and Computed Anharmonic Fundamental VPT2 Wavenumbers (in cm^{-1}) of CH_4 and CD_4^a

| symm. | MP2//MP2 ^b | | B2D3//B3D3 ^c | | CC//MP2 ^d | | CC//B3D3 ^e | | exp. | |
|---|-----------------------|---------------------|-------------------------|---------------------|----------------------|---------------------|-----------------------|---------------------|------|------|
| | ω | ν_{VPT2} | ω | ν_{VPT2} | ω | ν_{VPT2} | ω | ν_{VPT2} | | |
| CH_4^f | | | | | | | | | | |
| 1 ₁ ⟩ | A ₁ | 3073 | 2945 | 3051 | 2928 | 3036 | 2902 | 3036 | 2911 | 2921 |
| 1 ₂ , ±1 ₂ ⟩ | E | 1586 | 1551 | 1575 | 1543 | 1570 | 1534 | 1570 | 1538 | 1532 |
| 1 ₃ , 1 ₃ , ±1 ₃ or 0 ₃ ⟩ | T ₂ | 3209 | 3067 | 3163 | 3027 | 3157 | 3007 | 3157 | 3017 | 3022 |
| 1 ₄ , 1 ₄ , ±1 ₄ or 0 ₄ ⟩ | | 1352 | 1319 | 1353 | 1323 | 1345 | 1311 | 1345 | 1316 | 1308 |
| MAE | | | 28 | | 10 | | 10 | | 7 | |
| CD_4^f | | | | | | | | | | |
| 1 ₁ ⟩ | A ₁ | 2173 | 2124 | 2158 | 2113 | 2148 | 2098 | 2148 | 2102 | 2124 |
| 1 ₂ , ±1 ₂ ⟩ | E | 1122 | 1103 | 1114 | 1097 | 1111 | 1092 | 1111 | 1094 | 1093 |
| 1 ₃ , 1 ₃ , ±1 ₃ or 0 ₃ ⟩ | T ₂ | 2376 | 2295 | 2342 | 2264 | 2337 | 2252 | 2337 | 2258 | 2260 |
| 1 ₄ , 1 ₄ , ±1 ₄ or 0 ₄ ⟩ | | 1022 | 1003 | 1023 | 1006 | 1017 | 998 | 1017 | 1000 | 1001 |
| MAE | | | 17 | | 6 | | 10 | | 7 | |

^aMean absolute errors (MAEs) are also reported. The spherical and polar vibrational states are indicated respectively as $|l_{\nu}, k_{\nu}, m_{\nu}\rangle$ and $|l_{\nu}, l_{\nu}\rangle$.

^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set.

^cAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the B2D3/JnTZ level.

^dAnharmonic calculations performed at the MP2/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/cc-pVQZ level.

^eAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the CCSD(T)/cc-pVQZ level.

^fReference 135.

corresponding orange bars (C_{2v}) are all different. The couple of states $|1_{2,\pm 1_2}\rangle$ belonging to the irreducible representation E are still present in the C_{3v} system, while they are replaced with two non-degenerate states with representations A_1 and A_2 in the C_{2v} system. Again, the blue bars of the states $|1_{2,\pm 1_2}\rangle$ are equal, while the corresponding orange ones are different, following the symmetry breaking.

As a further test, CH_4 and CD_4 are considered. In addition to the MP2//MP2 and B2D3//B3D3 schemes employed for P_4 , two hybrid calculations have been performed at the MP2/JnDZ and B3D3/JnDZ levels in conjunction with a set of harmonic frequencies at the CCSD(T)/cc-pVQZ level.¹³⁴ The VPT2 wavenumbers are compared with the experimental data in Table 12.

Both systems do not show any Fermi or 1–1 Darling–Dennison resonance, except for CD_4 within the CC//MP2

hybrid scheme, where a Fermi resonance of type I ($\omega_1 \approx 2\omega_4$) has been detected, leading the vibrational frequency of $|1_1\rangle$ shifts from 2064 to 2099 cm^{-1} between the DVPT2 and GVPT2 levels. The presence of 2–2 Darling–Dennison resonances has been detected in the GVPT2 model, and the resonances between states possessing the same principal quanta have been properly included. Concerning CH_4 , the best agreement between theory and experiments is reached with the CC//B3D3 hybrid scheme (MAE = 7 cm^{-1}), even if all sets of VPT2 theoretical wavenumbers are close to those of the experiment, with the only exception being the MP2//MP2 scheme (MAE = 28 cm^{-1}), for which larger discrepancies are observed.

The results for CD_4 present a trend similar to that of CH_4 , with the MP2//MP2 scheme showing again the worst agreement with the experiment, although it is able to match exactly the experimental value of the fundamental band

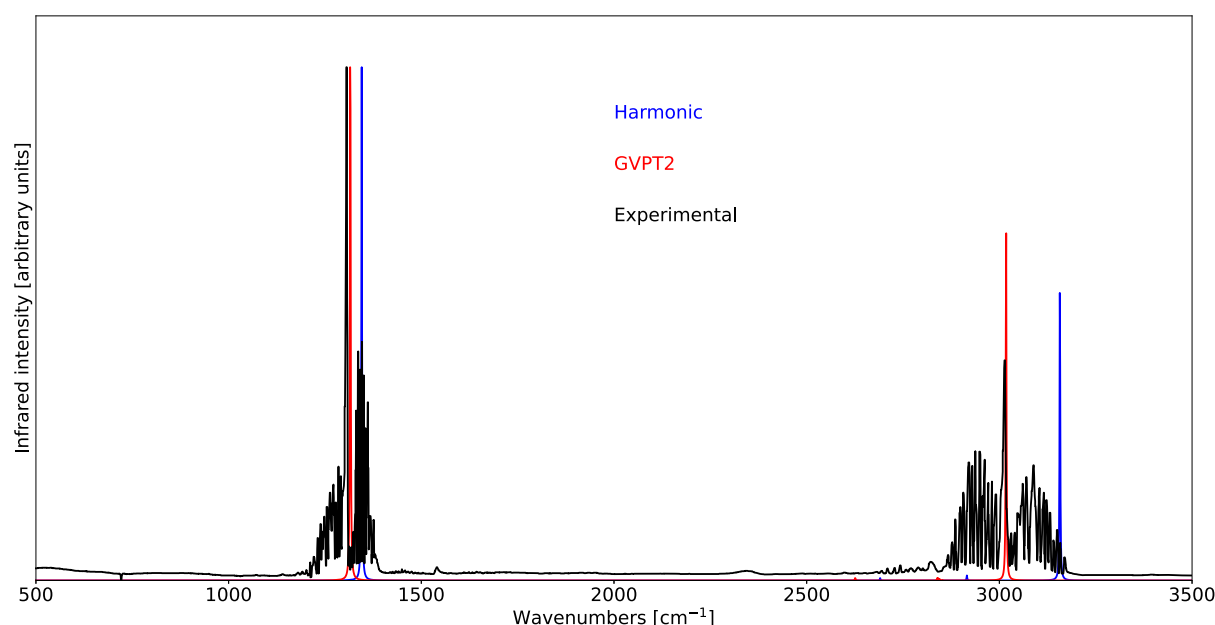


Figure 14. Comparison of the computed harmonic and GVPT2 IR spectra of methane at the CC//B3D3 level of theory with the experimental data. Spectral line shapes have been convoluted by Lorentzian distribution functions with HWHMs of 1 cm^{-1} . The experimental IR spectrum is from ref 145. All spectra are normalized by setting the intensity of their highest peak to unity.

associated to the state $|1_1\rangle$. The hybrid schemes based on CC harmonic frequencies (particularly CC//B3D3) are quite close to the experimental data, even though in this case, the best agreement is reached by the B2D3//B3D3 scheme (MAE = 6 cm^{-1}).

The anharmonic frequencies and intensities at the CC//B3D3 level have been employed in the simulation of the IR spectrum, which is compared with the harmonic and experimental ones in Figure 14.

As expected, the inclusion of anharmonic effects improves the agreement between theory and experiments, especially for the band around 3000 cm^{-1} , corresponding to the excitation of the triply degenerate fundamental $|1_3, 1_3, \pm 1_3\text{ or }0_3\rangle$.

4.3.2. Octahedral molecules. Finally, sulfur hexafluoride has been chosen as a test case for the O_h point group. The anharmonic calculations have been performed at the MP2/JnDZ and B3D3/JnDZ levels of theory, in conjunction with harmonic frequencies at the MP2/JnTZ and B2D3/JnTZ levels, respectively. A comparison between the VPT2 fundamental wavenumbers and their experimental counterparts is reported in Table 13.

This system does not show any Fermi or 1–1 Darling–Dennison resonance at the levels of theory employed here, while 2–2 Darling–Dennison resonances have been identified and included at the GVPT2 level. The best agreement with the experimental fundamentals is obtained at the MP2//MP2 level (MAE = 11 cm^{-1}), while the B2D3//B3D3 scheme is in this case characterized by a significant underestimation of all harmonic wavenumbers. For this reason, the anharmonic results obtained at the MP2//MP2 level have been employed in the simulation of the IR spectrum. Both theoretical harmonic and GVPT2 spectra are compared with the experimental one in Figure 15.

Similar to CH_4 , the inclusion of anharmonic contributions leads to a spectral profile closer to the experimental one, especially for the position of the band around 1000 cm^{-1} ,

Table 13. Comparison of Experimental and Computed Anharmonic Fundamental VPT2 Wavenumbers (in cm^{-1}) of Sulfur Hexafluoride^a

| | symm. | MP2//MP2 ^b | | B2D3//B3D3 ^c | | exp. ^d |
|---|----------|-----------------------|---------------------|-------------------------|---------------------|-------------------|
| | | ω | ν_{VPT2} | ω | ν_{VPT2} | |
| $ 1_1\rangle$ | A_{1g} | 771 | 762 | 748 | 740 | 775 |
| $ 1_2, \pm 1_2\rangle$ | E_g | 646 | 638 | 629 | 622 | 643 |
| $ 1_3, 1_3, \pm 1_3\text{ or }0_3\rangle$ | T_{1u} | 953 | 937 | 928 | 915 | 948 |
| $ 1_4, 1_4, \pm 1_4\text{ or }0_4\rangle$ | | 606 | 600 | 592 | 586 | 615 |
| $ 1_5, 1_5, \pm 1_5\text{ or }0_5\rangle$ | T_{2g} | 515 | 511 | 502 | 498 | 524 |
| $ 1_6, 1_6, \pm 1_6\text{ or }0_6\rangle$ | T_{2u} | 342 | 339 | 334 | 330 | 348 |
| MAE | | | 11 | | 27 | |

^aMean absolute errors (MAEs) are also reported. The spherical and polar vibrational states are indicated respectively as $|l_j, k_j, m_j\rangle$ and $|l_i, l_i\rangle$.

^bAnharmonic calculations performed with the JnDZ basis set based on a set of harmonic frequencies evaluated with the JnTZ basis set.

^cAnharmonic calculations performed at the B3D3/JnDZ level based on a set of harmonic frequencies evaluated at the B2D3/JnTZ level.

^dReference 155.

corresponding to the triply degenerate fundamentals $|1_3, 1_3, \pm 1_3\text{ or }0_3\rangle$.

5. CONCLUSIONS

In this work, we have shown how the canonical representation used for the development of VPT2 equations of asymmetric tops can be extended to linear and symmetric tops, followed by a series of a posteriori transformations, to give results identical to those obtained with the polar representation, thus offering the possibility of an ease of choice of the most convenient form for any application in vibro-rotational and vibrational spectroscopies. Such a strategy offers a number of advantages with respect to previous, ad hoc procedures. The first aspect concerns the ease of implementation since the new approach does not require any heavy modification of the codes already supporting VPT2 for asymmetric tops. The second aspect is the simplicity

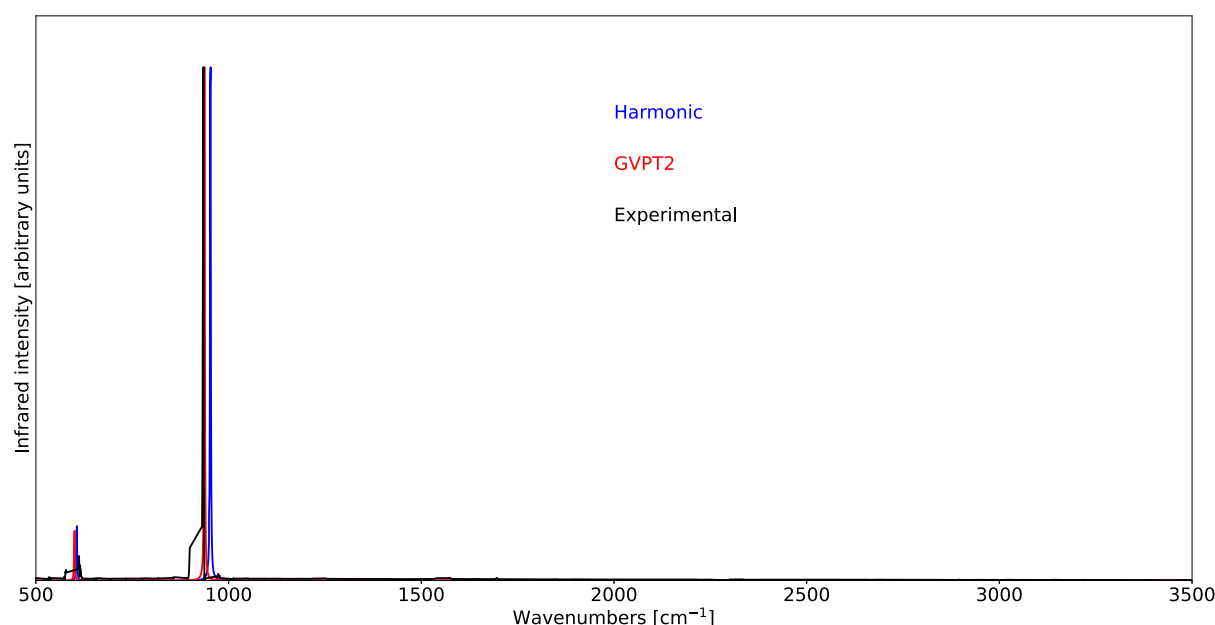


Figure 15. Comparison of the computed harmonic and GVPT2 IR spectra of sulfur hexafluoride obtained through the MP2//MP2 hybrid scheme of theory with the experimental data. Spectral line shapes have been convoluted by Lorentzian distribution functions with HWHMs of 1 cm^{-1} . The experimental IR spectrum is from ref 145. All spectra are normalized by setting the intensity of their highest peak to unity.

of the extension to spherical tops. Once the transformation matrix between the representations is known, it is possible to derive the necessary equations for any quantity of interest, which can be coded in small, specialized routines. However, the most important advantage is the availability of general equations for the intensities of all vibrational spectroscopies without the need of resorting to complex numbers. Actually, to the best of our knowledge, this is the first completely general implementation of intensities in the framework of the double-perturbation theory.

The results show that we dispose now of a general and robust implementation of GVPT2 for Abelian and non-Abelian groups allowing the effective treatment of medium- to large-sized molecules for all electronic structure methods for which analytical Hessians and first derivatives of properties are available. Hybrid methods in which harmonic and anharmonic contributions are treated at different levels can further extend the range of applications of the general platform. Studies in condensed phases can also be performed by means of mixed discrete-continuum models in which the solute and, possibly, some molecules of its cybotactic region are embedded in a polarizable continuum mimicking bulk solvent effects. Also in this case, the availability of analytical Hessians and first derivatives of properties allows an effective GVPT2 treatment.

Of course, all the intrinsic problems of a low-order perturbative treatment based on Cartesian normal modes are still present, especially concerning large-amplitude motions. Besides, the harmonic-oscillator wave functions do not always provide a suitable basis for the representation of vibrations, regardless of both the method used for the inclusion of anharmonic effects and the level of electronic theory employed. However, semi-rigid molecules can be routinely analyzed with remarkable results, largely sufficient for interpretation and assignment tasks. Extension to flexible systems can be pursued by coupling reduced-dimensionality treatments of large-amplitude motions to GVPT2 for small-amplitude motions. In this connection, use of curvilinear in place of rectilinear coordinates is an appealing option. While work in this and

related connections is underway in our laboratory, we think that already the present implementation offers a number of interesting perspectives for the study of molecular systems of current scientific and technological interest.

A.1 Degeneracies in the Perturbative Treatment

One of the obvious issues in the VPT2 expansion based on the canonical representation is the risk of singularity arising from the degeneracy of the vibrational modes and the related harmonic states. Let us consider the calculation of the energy ϵ_R of a vibrational state $|\nu_R\rangle$ (the superscript “C” has been dropped for clarity). Following the Rayleigh–Schrödinger perturbation theory, the following relations are obtained:

$$\epsilon_R = \epsilon_R^{(0)} + \epsilon_R^{(1)} + \epsilon_R^{(2)}$$

with

$$\epsilon_R^{(0)} = \langle \nu_R | \hat{H}^{(0)} | \nu_R \rangle = \sum_{i=1}^N \left(\frac{1}{2} + \nu_{R,i} \right) \omega_i \quad (39a)$$

$$\epsilon_R^{(1)} = \langle \nu_R | \hat{H}^{(1)} | \nu_R \rangle = 0 \quad (39b)$$

$$\epsilon_R^{(2)} = \langle \nu_R | \hat{H}^{(2)} | \nu_R \rangle + \sum_{S \neq R} \frac{\langle \nu_R | \hat{H}^{(1)} | \nu_S \rangle \langle \nu_S | \hat{H}^{(1)} | \nu_R \rangle}{\epsilon_R^{(0)} - \epsilon_S^{(0)}} \quad (39c)$$

where the vectorial form $|\nu_R\rangle = |\nu_{R,1} \dots \nu_{R,i} \dots \nu_{R,N}\rangle$ was preferred to represent the harmonic state $|\psi_R^{(0)}\rangle$.

The problem lies in the second term of the right-hand side of eq 39c since if $|\nu_R\rangle$ and $|\nu_S\rangle$ are degenerate, the denominator would be null. For the sake of simplicity but without loss of generality, let us consider that both states involve the same set of degenerate modes, $\{s_1, s_2\}$. In order to have $\epsilon_R^{(0)} = \epsilon_S^{(0)}$, the total number of quanta $\nu_s = \nu_{s_1} + \nu_{s_2}$ must be kept constant, and the number of quanta involving non-degenerate modes cannot vary. Mathematically, this translates in an even number of bosonic

ladder operators involving the degenerate modes. By construction, $\hat{\mathcal{H}}^{(1)}$ is defined as

$$\begin{aligned}\mathcal{H}^{(1)} &= \frac{1}{6} \sum_{i=1}^N \sum_{j=1}^N \sum_{k=1}^N \frac{\partial^3 V}{\partial q_i \partial q_j \partial q_k} q_i q_j q_k \\ &= \frac{1}{6} \sum_{i=1}^N \sum_{j=1}^N \sum_{k=1}^N f_{ijk} q_i q_j q_k\end{aligned}$$

which implies that three creation/annihilation operations are carried out. Hence, $\langle \mathbf{v}_S | \hat{\mathcal{H}}^{(1)} | \mathbf{v}_R \rangle = 0$ if $|\mathbf{v}_R\rangle$ and $|\mathbf{v}_S\rangle$ are degenerate. In other words, the representation matrix of $\hat{\mathcal{H}}^{(1)}$ over the degenerate state basis is null so that the canonical representation can be safely applied to treat systems with degenerate modes.

B.1 Calculation of Properties in the Polar Representation

In the following, we will describe the theoretical framework underlying the calculation of molecular properties (including transition ones) in the polar representation. Let us point out that even if the derivation reported below focuses on symmetric and linear tops, it still holds for systems presenting threefold degenerate vibrations.

Let us consider the operator $\hat{\Theta}$ describing a molecular property depending on normal coordinates or their conjugate momenta. Following previous works,^{34,47,98} a generic element of the representation matrix of $\hat{\Theta}$ over the canonical states can be written as

$$\langle \Theta \rangle_{R,S}^C = \frac{\langle \psi_R^C | \hat{\Theta} | \psi_S^C \rangle}{\sqrt{\langle \psi_R^C | \psi_R^C \rangle \langle \psi_S^C | \psi_S^C \rangle}} \quad (40)$$

The perturbative expansion of both $\hat{\Theta}$ and canonical states enables to write a generic element $\langle \Theta \rangle_{R,S}^C$ as

$$\langle \Theta \rangle_{R,S}^C = \langle \Theta \rangle_{R,S}^{C(0)} + \langle \Theta \rangle_{R,S}^{C(1)} + \langle \Theta \rangle_{R,S}^{C(2)} \quad (41)$$

where

$$\begin{aligned}\langle \Theta \rangle_{R,S}^{C(0)} &= \langle \psi_R^{C(0)} | \hat{\Theta}^{(0)} | \psi_S^{C(0)} \rangle \\ \langle \Theta \rangle_{R,S}^{C(1)} &= \langle \psi_R^{C(0)} | \hat{\Theta}^{(1)} | \psi_S^{C(0)} \rangle + \langle \psi_R^{C(1)} | \hat{\Theta}^{(0)} | \psi_S^{C(0)} \rangle \\ &\quad + \langle \psi_R^{C(0)} | \hat{\Theta}^{(0)} | \psi_S^{C(1)} \rangle \\ \langle \Theta \rangle_{R,S}^{C(2)} &= \langle \psi_R^{C(0)} | \hat{\Theta}^{(2)} | \psi_S^{C(0)} \rangle + \langle \psi_R^{C(2)} | \hat{\Theta}^{(0)} | \psi_S^{C(0)} \rangle \\ &\quad + \langle \psi_R^{C(0)} | \hat{\Theta}^{(0)} | \psi_S^{C(2)} \rangle \\ &\quad + \langle \psi_R^{C(1)} | \hat{\Theta}^{(1)} | \psi_S^{C(0)} \rangle + \langle \psi_R^{C(0)} | \hat{\Theta}^{(1)} | \psi_S^{C(1)} \rangle \\ &\quad + \langle \psi_R^{C(1)} | \hat{\Theta}^{(0)} | \psi_S^{C(1)} \rangle \\ &\quad - \frac{\langle \psi_R^{C(0)} | \hat{\Theta}^{(0)} | \psi_S^{C(0)} \rangle}{2} [\langle \psi_R^{C(1)} | \psi_R^{C(1)} \rangle + \langle \psi_S^{C(1)} | \psi_S^{C(1)} \rangle]\end{aligned} \quad (42)$$

Similar expressions stand for the polar representation. Since we already know that polar and canonical harmonic wave functions are linked through the customary expression

$$|\psi^P\rangle = \mathbf{P}^T |\psi^C\rangle \quad (43)$$

we will focus on the first- and second-order corrections to the wave functions.

B.1.1 Perturbative Corrections to the Polar States. B.1.1.1 First-Order Correction to the Wave Function. In the canonical representation, the first-order correction $|\psi_R^{C(1)}\rangle$ to the R th state is given by

$$\begin{aligned}|\psi_R^{C(1)}\rangle &= \sum_{S \neq R} \frac{\langle \psi_S^{C(0)} | \hat{\mathcal{H}}^{(1)} | \psi_R^{C(0)} \rangle}{\epsilon_R^{(0)} - \epsilon_S^{(0)}} |\psi_S^{C(0)}\rangle \\ &= \sum_{S \neq R} \frac{H_{SR}^{(1)}}{\epsilon_R^{(0)} - \epsilon_S^{(0)}} |\psi_S^{C(0)}\rangle\end{aligned} \quad (44)$$

The diagonal elements of the first-order Hamiltonian over the harmonic states are null as well as the off-diagonal ones when two degenerate states are considered. As a matter of fact, we can define the matrix $\bar{\mathbf{H}}^C$ as

$$\bar{H}_{SR}^C = \begin{cases} 0 & \text{if } R \text{ and } S \text{ degenerate} \\ \frac{H_{SR}^{(1)}}{\epsilon_R^{(0)} - \epsilon_S^{(0)}} & \text{otherwise} \end{cases} \quad (45)$$

so that eq 44 can be rewritten as

$$|\psi_R^{C(1)}\rangle = \sum_{S \neq R} \bar{H}_{SR}^C |\psi_S^{C(0)}\rangle \quad (46)$$

By defining $|\psi^C\rangle$ and $|\psi^{C(1)}\rangle$, the vectors collecting respectively the harmonic states and the corresponding first-order corrections in the canonical representation, eq 46 can be recast in the matrix form

$$|\psi^{C(1)}\rangle = \{\bar{\mathbf{H}}^C\}^T |\psi^C\rangle \quad (47)$$

and a similar expression holds in the polar representation

$$|\psi^{P(1)}\rangle = \{\bar{\mathbf{H}}^P\}^T |\psi^P\rangle \quad (48)$$

The diagonal blocks of $\bar{\mathbf{H}}^P$ and $\bar{\mathbf{H}}^C$ are null, and the energy difference $\epsilon_i^{(0)} - \epsilon_j^{(0)}$ reported above is constant within an off-diagonal block. As a matter of fact, we obtain the following identity:

$$\bar{\mathbf{H}}^P = \mathbf{P}^\dagger \bar{\mathbf{H}}^C \mathbf{P} \quad (49)$$

By substituting eqs 43 and 49 into eq 48, we obtain that

$$|\psi^{P(1)}\rangle = \mathbf{P}^T |\psi^{C(1)}\rangle \quad (50)$$

B.1.1.2 Second-Order Correction to the Wave Function. In the canonical representation, the second-order correction to the wave function is given by

$$\begin{aligned}|\psi_R^{C(2)}\rangle &= \sum_{S \neq R} \sum_{T \neq R} \frac{H_{ST}^{C(1)} H_{TR}^{C(1)}}{(\epsilon_R^{(0)} - \epsilon_S^{(0)})(\epsilon_R^{(0)} - \epsilon_T^{(0)})} \\ &\quad + \sum_{S \neq R} \frac{H_{SR}^{C(2)}}{\epsilon_R^{(0)} - \epsilon_S^{(0)}} |\psi_S^{C(0)}\rangle\end{aligned} \quad (51)$$

Let us define $\bar{\mathbf{H}}^C$ as

$$\bar{H}_{SR}^C = \begin{cases} 0 & \text{if } R \text{ and } S \text{ degenerate} \\ \sum_{T \neq R} \frac{H_{ST}^{C(1)} H_{TR}^{C(1)}}{\epsilon_R^{(0)} - \epsilon_T^{(0)}} = \sum_{T \neq R} H_{ST}^{C(1)} \bar{H}_{TR}^{C(1)} & \end{cases} \quad (52)$$

Equation 51 can be rewritten as follows:

$$|\psi_R^{C(2)}\rangle = \sum_{S \neq R} \frac{\bar{H}_{SR}^C + H_{SR}^{C(2)}}{\epsilon_R^{(0)} - \epsilon_S^{(0)}} |\psi_S^{C(0)}\rangle \quad (53)$$

Due to the presence of $H_{SR}^{C(2)}$ in eq 53, the terms for which the harmonic states $|\psi_R^{C(0)}\rangle$ and $|\psi_S^{C(0)}\rangle$ are degenerate must be excluded from the summation. From this, an analysis analogous to the one reported for the first-order correction leads to the following identity:

$$|\psi^{P(2)}\rangle = \mathbf{P}^T |\psi^{C(2)}\rangle \quad (54)$$

B.2.1 Molecular Properties: From the Canonical to the Polar Representation. Let us now consider the general expression for the conversion of a molecular property at the second order of perturbation. An element of the representation matrix of the operator $\hat{\Theta}$ over the perturbed polar states is given by an expression similar to eq 41

$$\langle \Theta \rangle_{R,S}^P = \langle \Theta \rangle_{R,S}^{P(0)} + \langle \Theta \rangle_{R,S}^{P(1)} + \langle \Theta \rangle_{R,S}^{P(2)} \quad (55)$$

where $\Theta^{P(0)}$, $\Theta^{P(1)}$, and $\Theta^{P(2)}$ are given by expressions similar to eq 42 that, recast in the matrix form, have the following expressions in the polar representation:

$$\begin{aligned} \langle \Theta \rangle^{P(0)} &= \langle \psi^{P(0)} | \hat{\Theta}^{(0)} | \psi^{P(0)} \rangle \\ \langle \Theta \rangle^{P(1)} &= \langle \psi^{P(0)} | \hat{\Theta}^{(1)} | \psi^{P(0)} \rangle + \langle \psi^{P(1)} | \hat{\Theta}^{(0)} | \psi^{P(0)} \rangle \\ &\quad + \langle \psi^{P(0)} | \hat{\Theta}^{(0)} | \psi^{P(1)} \rangle \\ \langle \Theta \rangle^{P(2)} &= \langle \psi^{P(0)} | \hat{\Theta}^{(2)} | \psi^{P(0)} \rangle + \langle \psi^{P(2)} | \hat{\Theta}^{(0)} | \psi^{P(0)} \rangle \\ &\quad + \langle \psi^{P(0)} | \hat{\Theta}^{(0)} | \psi^{P(2)} \rangle \\ &\quad + \langle \psi^{P(1)} | \hat{\Theta}^{(1)} | \psi^{P(0)} \rangle + \langle \psi^{P(0)} | \hat{\Theta}^{(1)} | \psi^{P(1)} \rangle \\ &\quad + \langle \psi^{P(1)} | \hat{\Theta}^{(0)} | \psi^{P(1)} \rangle \\ &\quad - \frac{\langle \psi^{P(0)} | \hat{\Theta}^{(0)} | \psi^{P(0)} \rangle}{2} [\langle \psi^{P(1)} | \psi^{P(1)} \rangle + \langle \psi^{P(1)} | \psi^{P(1)} \rangle]_N \end{aligned} \quad (56)$$

where the subscript N indicates a compact matrix notation for the normalization term.

By combining eqs 43, 50 and 54 with eq 56, the following identity is obtained:

$$\langle \Theta \rangle^P = \mathbf{P}^\dagger \langle \Theta \rangle^C \mathbf{P} \quad (57)$$

where $\langle \Theta \rangle^P$ and $\langle \Theta \rangle^C$ are respectively the representation matrices of the operator $\hat{\Theta}$ over the perturbed polar and canonical states. Consequently, the conversion of molecular properties at the anharmonic level is ruled by expressions similar to those used for the transformation of the contact-transformed Hamiltonian.

B.2.1.1 Transition Properties from the Ground State. Finally, let us consider the ground state as the initial state. It is worth mentioning that the matrix \mathbf{P} only mixes polar and canonical states that are degenerate at the harmonic level so that the perturbed wave function associated to the ground state is independent of the representation

$$|\psi_0^P\rangle = |\psi_0^C\rangle \quad (58)$$

As a result, the row vector $\langle \Theta \rangle_{0,S}^P$, containing the values of the transition property (e.g., a component of the dipole moment or polarizability) from the ground state to a set of states sharing the

set of principal quantum numbers ν_S , is related to its canonical counterpart $\langle \Theta \rangle_{0,S}^C$ as follows:

$$\langle \Theta \rangle_{0,S}^P = \langle \Theta \rangle_{0,S}^C \mathbf{P} \quad (59)$$

or equivalently,

$$\langle \Theta \rangle_{0,S}^P = \mathbf{P}^T \langle \Theta \rangle_{0,S}^C \quad (60)$$

where in eq 60, $\langle \Theta \rangle_{0,S}^P$ and $\langle \Theta \rangle_{0,S}^C$ are column vectors.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.jctc.1c00240>.

Rotation matrices, transition energies and intensities, and ZPVE for systems containing doubly or triply degenerate vibrations, χ and \mathbf{g} matrices, and l -type doubling coefficients in terms of the canonical representation, and expression for the transition properties accounting for degeneracies (PDF)

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Notes

The authors declare no competing financial interest.

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■ REFERENCES

- Barone, V. The Virtual Multifrequency Spectrometer: a New Paradigm for Spectroscopy. *Wiley Interdiscip. Rev.: Comput. Mol. Sci.* **2016**, *6*, 86–110.
- Puzzarini, C.; Bloino, J.; Tasinato, N.; Barone, V. Accuracy and Interpretability: The Devil and the Holy Grail. New Routes across Old Boundaries in Computational Spectroscopy. *Chem. Rev.* **2019**, *119*, 8131–8191.
- Tennyson, J. Perspective: Accurate ro-vibrational calculations on small molecules. *J. Chem. Phys.* **2016**, *145*, 120901.
- Puzzarini, C.; Stanton, J. F.; Gauss, J. Quantum-chemical calculation of spectroscopic parameters for rotational spectroscopy. *Int. Rev. Phys. Chem.* **2010**, *29*, 273–367.
- Császár, A. G.; Fábri, C.; Szidarovszky, T.; Mátyus, E.; Furtenbacher, T.; Czako, G. The fourth age of quantum chemistry: molecules in motion. *Phys. Chem. Chem. Phys.* **2012**, *14*, 1085–1106.
- Lipparini, F.; Mennucci, B. Perspective: Polarizable continuum models for quantum-mechanical descriptions. *J. Chem. Phys.* **2016**, *144*, 160901.

- (7) Barone, V.; Biczysko, M.; Puzzarini, C. Quantum Chemistry Meets Spectroscopy for Astrochemistry: Increasing Complexity toward Prebiotic Molecules. *Acc. Chem. Res.* **2015**, *48*, 1413–1422.
- (8) Hochlaf, M. Advances in spectroscopy and dynamics of small and medium sized molecules and clusters. *Phys. Chem. Chem. Phys.* **2017**, *19*, 21236–21261.
- (9) Császár, A. G. Anharmonic molecular force fields. *Wiley Interdiscip. Rev.: Comput. Mol. Sci.* **2011**, *2*, 273–289.
- (10) Baiardi, A.; Bloino, J.; Barone, V. General formulation of vibronic spectroscopy in internal coordinates. *J. Chem. Phys.* **2016**, *144*, 084114.
- (11) Nielsen, H. H. The Vibration-Rotation Energies of Molecules. *Rev. Mod. Phys.* **1951**, *23*, 90–136.
- (12) Mills, I. M. In *Molecular Spectroscopy: Modern Research*; Rao, K. N., Mathews, C. W., Eds.; Academic Press: New York, 1972; chapter 3.2, pp 115–140.
- (13) Whitehead, R. J.; Handy, N. C. Variational calculation of vibration-rotation energy levels for triatomic molecules. *J. Mol. Spectrosc.* **1975**, *55*, 356–373.
- (14) Bowman, J. M. Self-consistent field energies and wavefunctions for coupled oscillators. *J. Chem. Phys.* **1978**, *68*, 608–610.
- (15) Dunn, K. M.; Boggs, J. E.; Pulay, P. Vibrational energy levels of hydrogen cyanide. *J. Chem. Phys.* **1986**, *85*, 5838–5846.
- (16) Bowman, J. M. The self-consistent-field approach to polyatomic vibrations. *Acc. Chem. Res.* **1986**, *19*, 202–208.
- (17) Jung, J. O.; Gerber, R. B. Vibrational wave functions and spectroscopy of $(\text{H}_2\text{O})_n$, $n=2,3,4,5$: Vibrational self-consistent field with correlation corrections. *J. Chem. Phys.* **1996**, *105*, 10332–10348.
- (18) Carter, S.; Culik, S. J.; Bowman, J. M. Vibrational self-consistent field method for many-mode systems: A new approach and application to the vibrations of CO adsorbed on Cu(100). *J. Chem. Phys.* **1997**, *107*, 10458–10469.
- (19) Christiansen, O. Møller-Plesset perturbation theory for vibrational wave functions. *J. Chem. Phys.* **2003**, *119*, 5773–5781.
- (20) Christiansen, O. Vibrational coupled cluster theory. *J. Chem. Phys.* **2004**, *120*, 2149–2159.
- (21) Cassam-Chenaï, P.; Liévin, J. The VMFCI method: A flexible tool for solving the molecular vibration problem. *J. Comput. Chem.* **2006**, *27*, 627–640.
- (22) Christiansen, O. Vibrational structure theory: new vibrational wave function methods for calculation of anharmonic vibrational energies and vibrational contributions to molecular properties. *Phys. Chem. Chem. Phys.* **2007**, *9*, 2942–2953.
- (23) Christiansen, O. Selected new developments in vibrational structure theory: potential construction and vibrational wave function calculations. *Phys. Chem. Chem. Phys.* **2012**, *14*, 6672–6687.
- (24) Papoušek, D.; Aliev, M. R. *Molecular Vibrational-Rotational Spectra*; Elsevier Scientific Publishing Company, 1982.
- (25) Califano, S. *Vibrational States*; John Wiley & Sons, 1976.
- (26) Truhlar, D. G.; Olson, R. W.; Jeannotte, A. C.; Overend, J. Anharmonic force constants of polyatomic molecules. Test of the procedure for deducing a force field from the vibration-rotation spectrum. *J. Am. Chem. Soc.* **1976**, *98*, 2373–2379.
- (27) Isaacson, A. D.; Truhlar, D. G.; Scanlon, K.; Overend, J. Tests of approximation schemes for vibrational energy levels and partition functions for triatomics: H_2O and SO_2 . *J. Chem. Phys.* **1981**, *75*, 3017–3024.
- (28) Isaacson, A. D.; Hung, S. C. Use of second-order perturbation theory for the vibrational energy levels and partition functions at a saddle point. *J. Chem. Phys.* **1994**, *101*, 3928–3935.
- (29) Clabo, D. A., Jr.; Allen, W. D.; Remington, R. B.; Yamaguchi, Y.; Schaefer, H. F., III A systematic study of molecular vibrational anharmonicity and vibration-rotation interaction by self-consistent-field higher-derivative methods. Asymmetric top molecules. *Chem. Phys.* **1988**, *123*, 187–239.
- (30) Maslen, P. E.; Handy, N. C.; Amos, R. D.; Jayatilaka, D. Higher analytic derivatives. IV. Anharmonic effects in the benzene spectrum. *J. Chem. Phys.* **1992**, *97*, 4233–4254.
- (31) Zhang, Q.; Day, P. N.; Truhlar, D. G. The accuracy of second order perturbation theory for multiply excited vibrational energy levels and partition functions for a symmetric top molecular ion. *J. Chem. Phys.* **1993**, *98*, 4948–4958.
- (32) Barone, V. Vibrational zero-point energies and thermodynamic functions beyond the harmonic approximation. *J. Chem. Phys.* **2004**, *120*, 3059–3065.
- (33) Bloino, J.; Biczysko, M.; Crescenzi, O.; Barone, V. Integrated computational approach to vibrationally resolved electronic spectra: Anisole as a test case. *J. Chem. Phys.* **2008**, *128*, 244105.
- (34) Vázquez, J.; Stanton, J. F. Simple(r) algebraic equation for transition moments of fundamental transitions in vibrational second-order perturbation theory. *Mol. Phys.* **2006**, *104*, 377–388.
- (35) Vázquez, J.; Stanton, J. F. Treatment of Fermi resonance effects on transition moments in vibrational perturbation theory. *Mol. Phys.* **2007**, *105*, 101–109.
- (36) Van Vleck, J. H. On σ -Type Doubling and Electron Spin in the Spectra of Diatomic Molecules. *Phys. Rev.* **1929**, *33*, 467–506.
- (37) Kemble, E. C. *The Fundamental Principles of Quantum Mechanics*; Dover Publications: New York, 2005.
- (38) Messiah, A.; Schiff, L. I. *Quantum Mechanics*; McGraw-Hill College, 1968; Vol. 643.
- (39) Jensen, P.; Bunker, P. R. *Computational Molecular Spectroscopy*; John Wiley and Sons Ltd: Chichester, UK, 2000.
- (40) Willetts, A.; Handy, N. C.; Green, W. H., Jr.; Jayatilaka, D. Anharmonic Corrections to Vibrational Transition Intensities. *J. Phys. Chem.* **1990**, *94*, 5608–5616.
- (41) Ruden, T. A.; Taylor, P. R.; Helgaker, T. Automated calculation of fundamental frequencies: Application to AlH_3 using the coupled-cluster singles-and-doubles with perturbative triples method. *J. Chem. Phys.* **2003**, *119*, 1951–1960.
- (42) Barone, V. Anharmonic vibrational properties by a fully automated second-order perturbative approach. *J. Chem. Phys.* **2005**, *122*, 014108.
- (43) Carbonnière, P.; Dargelos, A.; Pouchan, C. The VCI-P code: an iterative variation-perturbation scheme for efficient computations of anharmonic vibrational levels and IR intensities of polyatomic molecules. *Theor. Chem. Acc.* **2010**, *125*, 543–554.
- (44) Krasnoshchekov, S. V.; Isayeva, E. V.; Stepanov, N. F. Numerical-Analytic Implementation of the Higher-Order Canonical Van Vleck Perturbation Theory for the Interpretation of Medium-Sized Molecule Vibrational Spectra. *J. Phys. Chem. A* **2012**, *116*, 3691–3709.
- (45) Gaw, F.; Willetts, A.; Handy, N.; Green, W. Chapter SPECTRO - a program for derivation of spectroscopic constants from provided quartic force fields and cubic dipole fields. In *Advances in Molecular Vibrations and Collision Dynamics*; Bowman, J. M., Ed.; JAI Press, 1992; Vol. 1, pp 186–195.
- (46) Dressler, S.; Thiel, W. Anharmonic force fields from density functional theory. *Chem. Phys. Lett.* **1997**, *273*, 71–78.
- (47) Barone, V.; Bloino, J.; Guido, C. A.; Lipparini, F. A fully automated implementation of VPT2 Infrared intensities. *Chem. Phys. Lett.* **2010**, *496*, 157–161.
- (48) Bloino, J.; Biczysko, M.; Barone, V. General Perturbative Approach for Spectroscopy, Thermodynamics, and Kinetics: Methodological Background and Benchmark Studies. *J. Chem. Theory Comput.* **2012**, *8*, 1015–1036.
- (49) Hermes, M. R.; Hirata, S. Second-order many-body perturbation expansions of vibrational Dyson self-energies. *J. Chem. Phys.* **2013**, *139*, 034111.
- (50) Werner, H.-J.; Knowles, P. J.; Knizia, G.; Manby, F. R.; Schütz, M.; Celani, P.; Korona, T.; Lindh, R.; Mitrushenkov, A.; Rauhut, G.; Shamasundar, K. R.; Adler, T. B.; Amos, R. D.; Bernhardsson, A.; Berning, A.; Cooper, D. L.; Deegan, M. J. O.; Dobbyn, F. A. J.; Eckert, Goll, E.; Hampel, C.; Hesselmann, A.; Hetzer, G.; Hrenar, T.; Jansen, G.; Köppl, C.; Liu, Y.; Lloyd, A. W.; Mata, R. A.; May, A. J.; McNicholas, S. J.; Meyer, W.; Mura, M. E.; Nicklass, A.; O'Neill, D. P.; Palmieri, P.; Peng, D.; Pflüger, R. K.; Pitzer, Reiher, M.; Shiozaki, T.; Stoll, H.; Stone, A. J.; Tarroni, R.; Thorsteinsson, T.; Wang, M. *MOLPRO*, version 2012.1, a package of ab initio programs, 2012.
- (51) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Petersson, G.

- A.; Nakatsuji, H.; Li, X.; Caricato, M.; Marenich, A. V.; Bloino, J.; Janesko, B. G.; Gomperts, R.; Mennucci, B.; Hratchian, H. P.; Ortiz, J. V.; Izmaylov, A. F.; Sonnenberg, J. L.; Williams-Young, D.; Ding, F.; Lipparini, F.; Egidi, F.; Goings, J.; Peng, B.; Petrone, A.; Henderson, T.; Ranasinghe, D.; Zakrzewski, V. G.; Gao, J.; Rega, N.; Zheng, G.; Liang, W.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Throssell, K.; Montgomery, J. A., Jr.; Peralta, J. E.; Ogliaro, F.; Bearpark, M. J.; Heyd, J. J.; Brothers, E. N.; Kudin, K. N.; Staroverov, V. N.; Keith, T. A.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A. P.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Millam, J. M.; Klene, M.; Adamo, C.; Cammi, R.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Farkas, O.; Foresman, J. B.; Fox, D. J. *Gaussian 16*, Revision A.03; Gaussian Inc: Wallingford CT, 2016.
- (52) Quade, C. R. Internal coordinate formulation for the vibration-rotation energies of polyatomic molecules. *J. Chem. Phys.* **1976**, *64*, 2783–2795.
- (53) Isaacson, A. D. Including Anharmonicity in the Calculation of Rate Constants. 1. The HCN/HNC Isomerization Reaction†. *J. Phys. Chem. A* **2006**, *110*, 379–388.
- (54) Baiardi, A.; Bloino, J.; Barone, V. Simulation of Vibronic Spectra of Flexible Systems: Hybrid DVR-Harmonic Approaches. *J. Chem. Theory Comput.* **2017**, *13*, 2804–2822.
- (55) Richter, F.; Carbonnière, P. Vibrational treatment of the formic acid double minimum case in valence coordinates. *J. Chem. Phys.* **2018**, *148*, 064303.
- (56) Wang, D.; Shi, Q.; Zhu, Q.-S. An ab initio quartic force field of PH₃. *J. Chem. Phys.* **2000**, *112*, 9624–9631.
- (57) Bunker, P. R.; Kraemer, W. P.; Špirko, V. An ab initio investigation of the potential function and rotation-vibration energies of NH₃. *Can. J. Phys.* **1984**, *62*, 1801–1805.
- (58) Mattioda, A. L.; Ricca, A.; Tucker, J.; Bauschlicher, C. W., Jr; Allamandola, L. J. Far-infrared spectroscopy of neutral coronene, ovalene, and dicyronylene. *Astron. J.* **2009**, *137*, 4054.
- (59) Zhang, Z.; Li, B.; Shen, Z.; Ren, Y.; Bian, W. Efficient quantum calculation of the vibrational states of acetylene. *Chem. Phys.* **2012**, *400*, 1–7.
- (60) Tamsamani, M. A.; Herman, M. The vibrational energy levels in acetylene 12C₂H₂: Towards a regular pattern at higher energies. *J. Chem. Phys.* **1995**, *102*, 6371–6384.
- (61) Henry, L.; Amat, G. The quartic anharmonic potential function of polyatomic molecules. *J. Mol. Spectrosc.* **1965**, *15*, 168–179.
- (62) Simmonett, A. C.; Schaefer, H. F., III; Allen, W. D. Enthalpy of formation and anharmonic force field of diacetylene. *J. Chem. Phys.* **2009**, *130*, 044301.
- (63) McNaughton, D.; Bruget, D. N. The high-resolution infrared spectrum of triacetylene. *J. Mol. Spectrosc.* **1991**, *150*, 620–634.
- (64) McNaughton, D.; Bruget, D. N. The high-resolution infrared spectrum of diacetylene and structures of diacetylene, triacetylene and dicyanoacetylene. *J. Mol. Struct.* **1992**, *273*, 11–25.
- (65) Shindo, F.; Bénilan, Y.; Chaquin, P.; Guillemin, J.-C.; Jolly, A.; Raulin, F. IR spectrum of C₈H₂: integrated band intensities and some observational implications. *J. Mol. Spectrosc.* **2001**, *210*, 191–195.
- (66) Gronowski, M.; Kolos, R. Isomers of cyanodiacetylene: Theoretical structures and IR spectra. *Chem. Phys. Lett.* **2006**, *428*, 245–248.
- (67) Janoschek, R. Novel carbon suboxides and subsulfides (C₅O₂, C₅S₂, C₄O₂ and C₂S₂): assignment of UV and IR spectra by quantum chemical calculations. *J. Mol. Struct.: THEOCHEM* **1991**, *232*, 147–154.
- (68) Wang, H.; Szczepanski, J.; Cooke, A.; Brucat, P.; Vala, M. Vibrational absorption spectra of C_nS (n = 2, 6) and C_nS₂ (n = 7, 9, 11, 13, 15) linear carbon-sulfur clusters. *Int. J. Quantum Chem.* **2005**, *102*, 806–819.
- (69) Mellau, G. C. Complete experimental rovibrational eigenenergies of HCN up to 6880 cm⁻¹ above the ground state. *J. Chem. Phys.* **2011**, *134*, 234303.
- (70) Schulze, G.; Koja, O.; Winnewisser, B. P.; Winnewisser, M. High resolution FIR spectra of DCNO and HCNO. *J. Mol. Struct.* **2000**, *517–518*, 307–325.
- (71) Piccardo, M.; Bloino, J.; Barone, V. Generalized vibrational perturbation theory for rovibrational energies of linear, symmetric and asymmetric tops: Theory, approximations, and automated approaches to deal with medium-to-large molecular systems. *Int. J. Quantum Chem.* **2015**, *115*, 948–982.
- (72) Pliva, J. Anharmonic constants for degenerate modes of symmetric top molecules. *J. Mol. Spectrosc.* **1990**, *139*, 278–285.
- (73) Franke, P. R.; Stanton, J. F.; Douberly, G. E. How to VPT2: Accurate and Intuitive Simulations of CH Stretching Infrared Spectra Using VPT2+ K with Large Effective Hamiltonian Resonance Treatments. *J. Phys. Chem. A* **2021**, *125*, 1301–1324.
- (74) Willetts, A.; Handy, N. C. The anharmonic constants for a symmetric top. *Chem. Phys. Lett.* **1995**, *235*, 286–290.
- (75) Bloino, J. VPT2 Route to Near-Infrared Spectroscopy: The Role of Mechanical and Electrical Anharmonicity. *J. Phys. Chem. A* **2015**, *119*, 5269–5287.
- (76) Bloino, J.; Baiardi, A.; Biczysko, M. Aiming at an accurate prediction of vibrational and electronic spectra for medium-to-large molecules: An overview. *Int. J. Quantum Chem.* **2016**, *116*, 1543–1574.
- (77) Schuurman, M. S.; Allen, W. D.; von Ragué Schleyer, P.; Schaefer, H. F., III The highly anharmonic BH₃ potential energy surface characterized in the *ab initio* limit. *J. Chem. Phys.* **2005**, *122*, 104302.
- (78) Rosnik, A. M.; Polik, W. F. VPT2+K spectroscopic constants and matrix elements of the transformed vibrational Hamiltonian of a polyatomic molecule with resonances using Van Vleck perturbation theory. *Mol. Phys.* **2014**, *112*, 261–300.
- (79) Krasnoshchekov, S. V.; Isayeva, E. V.; Stepanov, N. F. Criteria for first- and second-order vibrational resonances and correct evaluation of the Darling-Dennison resonance coefficients using the canonical Van Vleck perturbation theory. *J. Chem. Phys.* **2014**, *141*, 234114.
- (80) Martin, J. M. L.; Lee, T. J.; Taylor, P. R.; François, J. P. The anharmonic force field of ethylene, C₂H₄, by means of accurate ab initio calculations. *J. Chem. Phys.* **1995**, *103*, 2589–2602.
- (81) Bloino, J.; Barone, V. A second-order perturbation theory route to vibrational averages and transition properties of molecules: General formulation and application to infrared and vibrational circular dichroism spectroscopies. *J. Chem. Phys.* **2012**, *136*, 124108.
- (82) Darling, B. T.; Dennison, D. M. The Water Vapor Molecule. *Phys. Rev.* **1940**, *57*, 128–139.
- (83) Grenier-Besson, M. L. Résonances et dédoublements rotationnels du type l dans les molécules à symétrie axiale. *J. Phys. Radium* **1960**, *21*, 555–565.
- (84) Grenier-Besson, M. L. Résonances et dédoublements vibrationnels du type l dans les molécules à symétrie axiale. *J. Phys.* **1964**, *25*, 757–762.
- (85) Tarrago, G.; Ulenikov, O. N.; Poussigues, G. Dipole moment matrix for vibration-rotation transitions in C_{3v} molecules. *J. Phys.* **1984**, *45*, 1429–1447.
- (86) Long, D. A. *The Raman Effect: A Unified Treatment of the Theory of Raman Scattering by Molecules*; John Wiley & Sons Ltd, 2002.
- (87) Egidi, F.; Bloino, J.; Cappelli, C.; Barone, V. A robust and effective time-independent route to the calculation of resonance raman spectra of large molecules in condensed phases with the inclusion of duschinsky, herzberg-teller, anharmonic, and environmental effects. *J. Chem. Theory Comput.* **2014**, *10*, 346–363.
- (88) Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Scalmani, G.; Barone, V.; Petersson, G. A.; Nakatsuji, H.; Li, X.; Caricato, M.; Marenich, A. V.; Bloino, J.; Janesko, B. G.; Gomperts, R.; Mennucci, B.; Hratchian, H. P.; Ortiz, J. V.; Izmaylov, A. F.; Sonnenberg, J. L.; Williams-Young, D.; Ding, F.; Lipparini, F.; Egidi, F.; Goings, J.; Peng, B.; Petrone, A.; Henderson, T.; Ranasinghe, D.; Zakrzewski, V. G.; Gao, J.; Rega, N.; Zheng, G.; Liang, W.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Vreven, T.; Throssell, K.; Montgomery, J. A., Jr.; Peralta, J. E.; Ogliaro, F.; Bearpark, M. J.; Heyd, J. J.; Brothers, E. N.; Kudin, K. N.; Staroverov, V. N.; Keith,

- T. A.; Kobayashi, R.; Normand, J.; Raghavachari, K.; Rendell, A. P.; Burant, J. C.; Iyengar, S. S.; Tomasi, J.; Cossi, M.; Millam, J. M.; Klene, M.; Adamo, C.; Cammi, R.; Ochterski, J. W.; Martin, R. L.; Morokuma, K.; Farkas, O.; Foresman, J. B.; Fox, D. J. *Gaussian Development Version, Revision J.08*; Gaussian, Inc.: Wallingford CT, 2020.
- (89) Becke, A. D. Density-functional thermochemistry. III. The role of exact exchange. *J. Chem. Phys.* **1993**, *98*, 5648–5652.
- (90) Lee, C.; Yang, W.; Parr, R. G. Development of the Colle-Salvetti correlation-energy formula into a functional of the electron density. *Phys. Rev. B: Condens. Matter Mater. Phys.* **1988**, *37*, 785–789.
- (91) Vosko, S. H.; Wilk, L.; Nusair, M. Accurate spin-dependent electron liquid correlation energies for local spin density calculations: a critical analysis. *Can. J. Phys.* **1980**, *58*, 1200–1211.
- (92) Stephens, P. J.; Devlin, F. J.; Chabalowski, C. F.; Frisch, M. J. Ab Initio Calculation of Vibrational Absorption and Circular Dichroism Spectra Using Density Functional Force Fields. *J. Phys. Chem.* **1994**, *98*, 11623–11627.
- (93) Grimme, S. Semiempirical hybrid density functional with perturbative second-order correlation. *J. Chem. Phys.* **2006**, *124*, 034108.
- (94) Dierksen, M.; Grimme, S. A theoretical study of the chiroptical properties of molecules with isotopically engendered chirality. *J. Chem. Phys.* **2006**, *124*, 174301.
- (95) Grimme, S.; Antony, J.; Ehrlich, S.; Krieg, H. A consistent and accurate ab initio parametrization of density functional dispersion correction (DFT-D) for the 94 elements H-Pu. *J. Chem. Phys.* **2010**, *132*, 154104.
- (96) Grimme, S.; Ehrlich, S.; Goerigk, L. Effect of the damping function in dispersion corrected density functional theory. *J. Comput. Chem.* **2011**, *32*, 1456–1465.
- (97) Biczysko, M.; Panek, P.; Scalmani, G.; Bloino, J.; Barone, V. Harmonic and Anharmonic Vibrational Frequency Calculations with the Double-Hybrid B2PLYP Method: Analytic Second Derivatives and Benchmark Studies. *J. Chem. Theory Comput.* **2010**, *6*, 2115–2125.
- (98) Barone, V.; Biczysko, M.; Bloino, J. Fully anharmonic IR and Raman spectra of medium-size molecular systems: accuracy and interpretation. *Phys. Chem. Chem. Phys.* **2014**, *16*, 1759–1787.
- (99) Papajak, E.; Zheng, J.; Xu, X.; Leverentz, H. R.; Truhlar, D. G. Perspectives on basis sets beautiful: Seasonal plantings of diffuse basis functions. *J. Chem. Theory Comput.* **2011**, *7*, 3027–3034.
- (100) Schneider, W.; Thiel, W. Anharmonic force fields from analytic second derivatives: Method and application to methyl bromide. *Chem. Phys. Lett.* **1989**, *157*, 367–373.
- (101) Yang, Q.; Mendolicchio, M.; Barone, V.; Bloino, J. Accuracy and Reliability in the Simulation of Vibrational Spectra: A Comprehensive Benchmark of Energies and Intensities Issuing from Generalized Vibrational Perturbation Theory to Second Order (GVPT2). *Front. Astron. Space Sci.* **2021**, *8*, 77.
- (102) Begue, D.; Carbonniere, P.; Pouchan, C. Calculations of Vibrational Energy Levels by Using a Hybrid ab Initio and DFT Quartic Force Field: Application to Acetonitrile. *J. Phys. Chem. A* **2005**, *109*, 4611–4616.
- (103) Carbonniere, P.; Lucca, T.; Pouchan, C.; Rega, N.; Barone, V. Vibrational computations beyond the harmonic approximation: Performances of the B3LYP density functional for semirigid molecules. *J. Comput. Chem.* **2005**, *26*, 384–388.
- (104) Puzzarini, C.; Barone, V. Toward spectroscopic accuracy for organic free radicals: Molecular structure, vibrational spectrum, and magnetic properties of F₂NO. *J. Chem. Phys.* **2008**, *129*, 084306.
- (105) Biczysko, M.; Bloino, J.; Puzzarini, C. Computational challenges in Astrochemistry. *Wiley Interdiscip. Rev.: Comput. Mol. Sci.* **2018**, *8*, No. e1349.
- (106) Fortenberry, R. C.; Lee, T. J.; Layfield, J. P. Communication: The failure of correlation to describe carbon-carbon bonding in out-of-plane bends. *J. Chem. Phys.* **2017**, *147*, 221101.
- (107) Fortenberry, R. C.; Novak, C. M.; Layfield, J. P.; Matito, E.; Lee, T. J. Overcoming the Failure of Correlation for Out-of-Plane Motions in a Simple Aromatic: Rovibrational Quantum Chemical Analysis of c-C₃H₂. *J. Chem. Theory Comput.* **2018**, *14*, 2155–2164.
- (108) Westbrook, B. R.; Del Rio, W. A.; Lee, T. J.; Fortenberry, R. C. Overcoming the out-of-plane bending issue in an aromatic hydrocarbon: the anharmonic vibrational frequencies of c-(CH)₃H. *Phys. Chem. Chem. Phys.* **2020**, *22*, 12951–12958.
- (109) Lee, T. J.; Fortenberry, R. C. The unsolved issue with out-of-plane bending frequencies for C=C multiply bonded systems. *Spectrochim. Acta, Part A* **2021**, *248*, 119148.
- (110) Cheeseman, J. R.; Frisch, M. J. Basis Set Dependence of Vibrational Raman and Raman Optical Activity Intensities. *J. Chem. Theory Comput.* **2011**, *7*, 3323–3334.
- (111) Lee, T. J.; Dateo, C. E.; Gazdy, B.; Bowman, J. M. Accurate quartic force fields and vibrational frequencies for hydrogen cyanide and hydrogen isocyanide. *J. Phys. Chem.* **1993**, *97*, 8937–8943.
- (112) Martin, J. M. L.; Lee, T. J.; Taylor, P. R. A purely ab initio spectroscopic quality quartic force field for acetylene. *J. Chem. Phys.* **1998**, *108*, 676–691.
- (113) Rauhut, G.; Knizia, G.; Werner, H.-J. Accurate calculation of vibrational frequencies using explicitly correlated coupled-cluster theory. *J. Chem. Phys.* **2009**, *130*, 054105.
- (114) Adel, A.; Dennison, D. M. The infrared spectrum of carbon dioxide. Part I. *Phys. Rev.* **1933**, *43*, 716.
- (115) Suzuki, I. General anharmonic force constants of carbon dioxide. *J. Mol. Spectrosc.* **1968**, *25*, 479–500.
- (116) Miller, C. E.; Brown, L. R. Near infrared spectroscopy of carbon dioxide I. 16O12C16O line positions. *J. Mol. Spectrosc.* **2004**, *228*, 329–354.
- (117) Requena, A.; Bastida, A.; Zúñiga, J. Curvilinear Jacobi and Radau normal coordinates for linear triatomic molecules. Application to CO₂. *Chem. Phys.* **1993**, *175*, 255–264.
- (118) NIST Mass Spec Data Center; Stein, S.E. Chapter Infrared Spectra. In *NIST Chemistry WebBook, NIST Standard Reference Database Number 69*; Linstrom, P. J., Mallard, W. G., Eds.; National Institute of Standards and Technology: Gaithersburg MD, 20899, 2015. (retrieved September 20, 2014). <http://webbook.nist.gov>.
- (119) Howard-Lock, H. E.; Stoicheff, B. P. Raman intensity measurements of the Fermi diad $\nu_1, 2\nu_2$ in 12CO₂ and 13CO₂. *J. Mol. Spectrosc.* **1971**, *37*, 321–326.
- (120) Dargelos, A.; Pouchan, C. Ab Initio Modeling of the IR Spectra of Dicyanoacetylene in the Region 100–4800 cm⁻¹. *J. Phys. Chem. A* **2016**, *120*, 6270–6273.
- (121) Miller, F. A.; Hannan, R. B., Jr The infrared and Raman spectra of dicyanoacetylene. *J. Chem. Phys.* **1953**, *21*, 110–114.
- (122) Miller, F. A.; Hannan, R. B., Jr; Cousins, L. R. Infrared and Raman Spectra of Dicyanoacetylene. II. *J. Chem. Phys.* **1955**, *23*, 2127–2129.
- (123) Winther, F.; Ketelsen, M.; Guarnieri, A. The infrared and Raman spectrum of dicyanoacetylene. The ν_9 fundamental. *J. Mol. Struct.* **1994**, *320*, 65–73.
- (124) Winther, F.; Schönhoff, M. The Fundamental Vibrations of NC–CC–CN (Dicyanoacetylene). *J. Mol. Spectrosc.* **1997**, *186*, 54–65.
- (125) Khelifi, M.; Paillous, P.; Bruston, P.; Guillemin, J. C.; Bénilan, Y.; Daoudi, A.; Raulin, F. Gas infrared spectra, assignments, and absolute IR band intensities of C₄N₂ in the 250–3500 cm⁻¹ region: implications for Titan's stratosphere. *Spectrochim. Acta, Part A* **1997**, *53*, 707–712.
- (126) Fayt, A.; Vigouroux, C.; Winther, F. Analysis of the ν_9 band complex of dicyanoacetylene and application of a theory of relative intensities to all subbands. *J. Mol. Spectrosc.* **2004**, *224*, 114–130.
- (127) Jolly, A.; Cottini, V.; Fayt, A.; Manceron, L.; Kwabia-Tchana, F.; Benilan, Y.; Guillemin, J.-C.; Nixon, C.; Irwin, P. Gas phase dicyanoacetylene (C₄N₂) on Titan: New experimental and theoretical spectroscopy results applied to Cassini CIRS data. *Icarus* **2015**, *248*, 340–346.
- (128) Couturier-Tamburelli, I.; Gudipati, M. S.; Lignell, A.; Jacovi, R.; Piétri, N. Spectroscopic studies of non-volatile residue formed by photochemistry of solid C₄N₂: A model of condensed aerosol formation on Titan. *Icarus* **2014**, *234*, 81–90.
- (129) Peterson, K. A.; Figgen, D.; Goll, E.; Stoll, H.; Dolg, M. Systematically convergent basis sets with relativistic pseudopotentials.

II. Small-core pseudopotentials and correlation consistent basis sets for the post-d group 16–18 elements. *J. Chem. Phys.* **2003**, *119*, 11113–11123.

(130) Carnimeo, I.; Puzzarini, C.; Tasinato, N.; Stoppa, P.; Charmet, A. P.; Biczysko, M.; Cappelli, C.; Barone, V. Anharmonic theoretical simulations of infrared spectra of halogenated organic compounds. *J. Chem. Phys.* **2013**, *139*, 074310.

(131) Drage, E. A.; Jaksch, D.; Smith, K. M.; McPheat, R. A.; Vasekova, E.; Mason, N. J. FTIR spectroscopy and estimation of the global warming potential of CF₃Br and C₂F₄. *J. Quant. Spectrosc. Radiat. Transfer* **2006**, *98*, 44–56.

(132) Martins Filho, H. P.; Guadagnini, P. H. Infrared vibrational intensities and polar tensors of CF₃Br and CF₃I. *J. Mol. Struct.: THEOCHEM* **1999**, *464*, 171–182.

(133) Charmet, A. P.; Tasinato, N.; Stoppa, P.; Baldacci, A.; Giorgianni, S. Jet-cooled diode laser spectrum and FTIR integrated band intensities of CF₃Br: rovibrational analysis of $2\nu_3$ and $\nu_2+\nu_3$ bands near 9 μm and cross-section measurements in the 450–2500 cm^{-1} region. *Mol. Phys.* **2008**, *106*, 1171–1179.

(134) Lee, T. J.; Martin, J. M. L.; Taylor, P. R. An accurate ab initio quartic force field and vibrational frequencies for CH₄ and isotopomers. *J. Chem. Phys.* **1995**, *102*, 254–261.

(135) Albert, S.; Bauerecker, S.; Boudon, V.; Brown, L. R.; Champion, J.-P.; Loëte, M.; Nikitin, A.; Quack, M. Global analysis of the high resolution infrared spectrum of methane 12CH₄ in the region from 0 to 4800 cm^{-1} . *Chem. Phys.* **2009**, *356*, 131–146.

(136) Gray, D. L.; Robiette, A. G. The anharmonic force field and equilibrium structure of methane. *Mol. Phys.* **1979**, *37*, 1901–1920.

(137) Bencze, É.; Lokshin, B. V.; Mink, J.; Herrmann, W. A.; Kühn, F. E. Vibrational spectra and structure of the cyclopentadienyl-anion (Cp⁻), the pentamethylcyclopentadienyl-anion (Cp^{*-}) and of alkali metal cyclopentadienyls CpM and Cp^{*}M (M= Li, Na, K). *J. Organomet. Chem.* **2001**, *627*, 55–66.

(138) Nakamoto, K. *Infrared and Raman Spectra of Inorganic and Coordination Compounds. Part B: Application in Coordination, Organometallic, and Bioinorganic Chemistry*, 5th ed.; Wiley: New York, 1997; p 286.

(139) Garbuzova, I. A.; Garkusha, O. G.; Lokshin, B. V.; Borisov, G. K.; Morozova, T. S. Vibrational spectra of alkali metal cyclopentadienides. *J. Organomet. Chem.* **1985**, *279*, 327–335.

(140) Handy, N. C.; Willetts, A. Anharmonic constants for benzene. *Spectrochim. Acta, Part A* **1997**, *53*, 1169–1177.

(141) Miani, A.; Cané, E.; Palmieri, P.; Trombetti, A.; Handy, N. C. Experimental and theoretical anharmonicity for benzene using density functional theory. *J. Chem. Phys.* **2000**, *112*, 248–259.

(142) Goodman, L.; Ozkabak, A. G.; Thakur, S. N. A benchmark vibrational potential surface: ground-state benzene. *J. Phys. Chem.* **1991**, *95*, 9044–9058.

(143) Henry, B. R.; Siebrand, W. Anharmonicity in polyatomic molecules. The CH-stretching overtone spectrum of benzene. *J. Chem. Phys.* **1968**, *49*, 5369–5376.

(144) Martin, J. M. L.; Taylor, P. R.; Lee, T. J. The harmonic frequencies of benzene. A case for atomic natural orbital basis sets. *Chem. Phys. Lett.* **1997**, *275*, 414–422.

(145) Gordon, I. E.; Rothman, L. S.; Hill, C.; Kochanov, R. V.; Tan, Y.; Bernath, P. F.; Birk, M.; Boudon, V.; Campargue, A.; Chance, K. V.; et al. The HITRAN2016 molecular spectroscopic database. *J. Quant. Spectrosc. Radiat. Transfer* **2017**, *203*, 3–69.

(146) Rinsland, C. P.; Devi, V. M.; Blake, T. A.; Sams, R. L.; Sharpe, S.; Chiou, L. Quantitative measurement of integrated band intensities of benzene vapor in the mid-infrared at 278, 298, and 323 K. *J. Quant. Spectrosc. Radiat. Transfer* **2008**, *109*, 2511–2522.

(147) Kalasinsky, V. F. Raman spectra and vibrational dephasing of pentaborane (9). *J. Phys. Chem.* **1979**, *83*, 3239–3243.

(148) Hrostowski, H. J.; Pimentel, G. C. The infrared spectra of stable pentaborane and deuterated pentaborane. *J. Am. Chem. Soc.* **1954**, *76*, 998–1003.

(149) Taylor, W. J.; Beckett, C.; Tung, J.; Holden, R.; Johnston, H. Raman and infra-red spectra of pentaborane. *Phys. Rev.* **1950**, *79*, 234.

(150) Maillard, R.; Sethio, D.; Hagemann, H.; Lawson Daku, L. M. Accurate Computational Thermodynamics Using Anharmonic Density Functional Theory Calculations: The Case Study of B–H Species. *ACS Omega* **2019**, *4*, 8786–8794.

(151) Raynes, W. T.; Lazzeretti, P.; Zanasi, R.; Sadlej, A. J.; Fowler, P. W. Calculations of the force field of the methane molecule. *Mol. Phys.* **1987**, *60*, 509–525.

(152) Hodgkinson, D. P.; Heenan, R. K.; Hoy, A. R.; Robiette, A. G. Vibrational anharmonicity in octahedral XY₆ molecules: Theoretical expressions for the spectroscopic constants. *Mol. Phys.* **1983**, *48*, 193–208.

(153) Edwards, H. G. M. Vibrational Raman spectrum and force constants of phosphorus, P₄. *J. Mol. Struct.* **1993**, *295*, 95–100.

(154) Persson, B. J.; Taylor, P. R.; Lee, T. J. Ab initio geometry, quartic force field, and vibrational frequencies for P₄. *J. Chem. Phys.* **1997**, *107*, 5051–5057.

(155) McDowell, R. S.; Krohn, B. J.; Flicker, H.; Vasquez, M. C. Vibrational levels and anharmonicity in SF₆-I. Vibrational band analysis. *Spectrochim. Acta, Part A* **1986**, *42*, 351–369.