

# In Vitro and In Silico Vibrational–Rotational Spectroscopic Characterization of the Next-Generation Refrigerant HFO-1123

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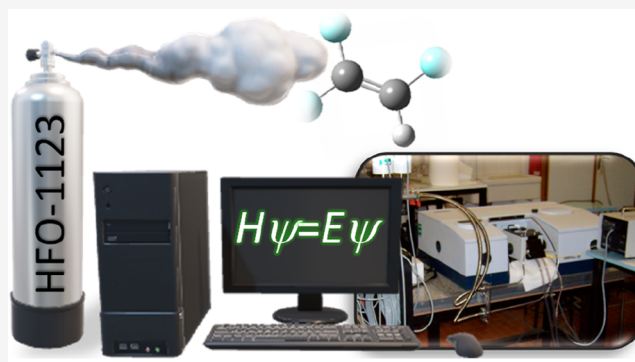
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**ABSTRACT:** Very short-lived substances have recently been proposed as replacements for hydrofluorocarbons (HFCs), in turn being used in place of ozone-depleting substances, in refrigerant applications. In this respect, hydro-fluoro-olefins (HFOs) are attracting particular interest because, due to their reduced global warming potential, they are supposed to be environmentally friendlier. Notwithstanding this feature, they represent a new class of compounds whose spectroscopic properties and reactivity need to be characterized to allow their atmospheric monitoring and to understand their environmental fate. In the present work, the structural, vibrational, and ro-vibrational properties of trifluoroethene (HFO-1123,  $F_2C = CHF$ ) are studied by state-of-the-art quantum chemical calculations. The equilibrium molecular structure has an expected error within 2 mÅ and  $0.2^\circ$  for bond lengths and angles, respectively. This represents the first step toward the computation of highly accurate rotational constants for both the ground and first excited fundamental vibrational levels, which reproduce the available experimental data well within 0.1%. Centrifugal distortion parameters and vibrational–rotational coupling terms are computed as well and used to solve some conflicting experimental results. Simulation of the vibrational transition frequencies and intensities beyond the double harmonic approximation and up to three quanta of vibrational excitation provides insights into the couplings ruling the vibrational dynamics and guides the characterization of the gas-phase infrared spectrum experimentally recorded in the range of 200–5000  $cm^{-1}$ . The full characterization of the IR features is completed with the experimental determination of the absorption cross sections over the 400–5000  $cm^{-1}$  region from which the radiative forcing and global warming potential of HFO-1123 are derived.



## 1. INTRODUCTION

Halogenated organic molecules, when released into the environment, influence atmospheric chemistry and, hence, Earth's climate by contributing to either ozone depletion or global warming. Due to this role as trace gas pollutants, over the years, halocarbons have been the subject of many experimental and theoretical investigations aimed on the one side at allowing their monitoring and on the other at understanding their atmospheric reactivity.<sup>1–3</sup> Halogenated hydrocarbons are in general widely used as blowing agents, propellants, refrigerants, and fire extinguishers, and halogenated olefins are also widely employed to produce synthetic polymers and copolymers characterized by a good mechanical, thermal, and chemical resistance. However, during the last three decades, a transition from chlorofluorocarbons (CFCs) to hydro-chlorofluorocarbons (HCFCs) and then to hydrofluorocarbons (HFCs) has taken place, driven by the need of mitigating stratospheric ozone depletion. In fact, the Montreal protocol that was enforced in 1987 has been phased out with a

few exceptions: the use of CFCs since 1996 due to their capacity to catalytically destroy stratospheric ozone molecules. This feature comes from the chlorine atom that, once in the stratosphere, acts as a catalyst in the ozone depleting reaction cycles. In addition, CFCs are also greenhouse gases because they strongly absorb the electromagnetic radiation within the atmospheric window located in the 8–12  $\mu m$  wavelength spectral region. Indeed, these molecules generally show very strong infrared (IR) absorption bands due to C–F and C–Cl stretching vibrations. To make things worse, CFCs generally have long atmospheric lifetimes that on the one side favor their

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accumulation in the atmosphere and on the other side contribute even more to their global warming activity in the long period. As replacement gases, HCFCs and HFCs with analogous chemical and physical properties have been proposed. Between the two, HFCs have appeared to be the more convenient option because, given the absence of chlorine atoms, they cannot contribute to ozone loss. Furthermore, these molecules present shorter atmospheric lifetimes than CFCs, because the C–H bond undergoes a reaction toward hydroxyl radicals in the troposphere.<sup>4</sup> Nevertheless, these gases still contribute to global climate changes as the presence of the C–F chromophore makes them greenhouse gases that absorb IR radiation around 9  $\mu\text{m}$ .

In 2016, the Kigali amendment to the Montreal protocol posed the onset of a new transition with the requirement to reduce HFCs emissions and to switch to the use of compounds characterized by a low global warming potential (GWP). Low-GWP molecules are currently being developed in order to meet the new regulation, and hydro-fluoro-olefins (HFOs) have appeared to be a good solution for applications in the field of refrigeration and air conditioning. In 2014, Asahi Glass Co., Ltd. employed trifluoroethene ( $\text{F}_2\text{C}=\text{CHF}$ ), known as HFO-1123, as the main component of refrigerant blends characterized by extremely low GWPs. As such, HFO-1123 is emerging as a new molecule in the refrigeration market, and it is expected to be a next-generation refrigerant for air conditioners.<sup>5</sup> From a chemical point of view, presenting unsaturated C=C bonds, HFOs have low chemical stabilities with a consequent shortening of the atmospheric lifetime that is, in turn, at the basis of the low GWP. This increased chemical reactivity however may cause self-decomposition reactions, and in fact, it has been reported that HFO-1123 undergoes a disproportionation reaction under high-temperature and high-pressure conditions.<sup>6,7</sup> As such, the thermodynamic properties of this molecule as well as its binary mixtures with other fluorinated refrigerants have been studied in recent years (see refs 8–11 and the references therein). Furthermore, HFO-1123 reactivity toward the OH radical has also been investigated.<sup>12</sup>

Among the different techniques available for environmental and process monitoring, one of the most effective and easy-to-use is IR spectroscopy whose successful application, however, depends on the availability of accurate spectroscopic data, above all the transition frequencies and intensities, for the species of interest.<sup>1,13,14</sup> While high-resolution spectroscopic investigations can be exploited to obtain the requested information to a very high accuracy, the necessary ro-vibrational analyses are complicated by the presence of resonances and very often by the spectral congestion due to the presence of low-lying vibrational states.<sup>15,16</sup> The determination of line intensities requires a careful line-by-line approach<sup>17,18</sup> that, being hampered by the overlap among different spectral features, can often be effectively exploited only for the lighter species. Vibrational assignments and band intensities, besides delivering the fundamental knowledge that is instrumental for the ro-vibrational analysis of high-resolution IR spectra, provide the description of vibrational couplings occurring within the molecule.<sup>19–21</sup> This data, coupled with sophisticated experimental methods capable of simplifying the spectral structure (like cooling cells<sup>22,23</sup> and supersonic jet expansion<sup>24,25</sup>) and with purposely tailored data-analysis software,<sup>26–28</sup> aids and speeds up the huge and time-consuming task of interpreting high-resolution spectra.

On their own, vibrational investigations are ever increasingly supported by quantum chemical (QC) calculations that work as a guide and a support for the assignment of the experimental signals. In fact, reliable predictions of vibrational transition frequencies and IR intensities can be obtained beyond the rigid-rotor-harmonic-oscillator model without the need of any empirical scaling factor and with a full account of anharmonic resonances with accuracies within 10  $\text{cm}^{-1}$  for fundamental frequencies and a few  $\text{km mol}^{-1}$  for IR intensities.<sup>29,30</sup> Concerning halocarbons, it has been demonstrated that such accuracy can be reached using the coupled cluster theory with singles, doubles, and a perturbative estimate of the triples, CCSD(T),<sup>31,32</sup> coupled to large basis sets.<sup>33–35</sup> An alternative and more cost-effective approach is represented by QM/QM' computational protocols involving CCSD(T)-based composite schemes for geometry and harmonic properties,<sup>36</sup> coupled with anharmonic effects described through density functional theory (DFT).<sup>37,38</sup>

The microwave spectra of HFO-1123 were investigated about 50 years ago, leading to the determination of the ground and some excited vibrational level rotational constants and to an effective structure,<sup>39</sup> which was later refined using microwave and electron diffraction data.<sup>40</sup> The rotational spectra were reanalyzed by obtaining centrifugal distortion terms some years later,<sup>41</sup> while more recently, improved rotational parameters have been determined for the ground and some low lying vibrational excited states together with the analysis of the ro-vibrational spectrum of the  $\nu_6$  band.<sup>42</sup> The vibrational spectrum of trifluoroethene was measured a long time ago,<sup>43</sup> while in the middle of the 70s, it was considered in a study devoted to explore the C–H stretching frequencies in a series of halogenated ethenes.<sup>44</sup> Some high-resolution investigations were carried out in more recent times with the aim of determining the molecular parameters in some excited vibrational levels, even though the observed irregularities in the spectra patterns, due to Coriolis and anharmonic interactions, were only recognized without any formal treatment of the couplings.<sup>45–47</sup> The vibrational properties of HFO-1123 were the subject of theoretical computations within the harmonic approximation,<sup>48,49</sup> and more recently, Jiang et al. compared the available experimental data on fundamental frequencies with the corresponding values computed by scaled ab initio calculations using the B3PW91 functional.<sup>50</sup>

As the characterization of the vibrational spectrum of trifluoroethene dates back to almost 70 years ago and an accurate modeling of its ro-vibrational spectroscopic properties has never been undertaken, the present work deals with a combined experimental and theoretical study of the vibrational and ro-vibrational properties of HFO-1123 in order to fill these gaps in view of the renewed interest in this species for applications as a refrigerant of the next generation. Experimentally, the vibrational quantum assignment of the gas-phase IR spectrum is carried out between 200 and 5000  $\text{cm}^{-1}$  and the determination of integrated absorption cross sections is performed over the 400–5000  $\text{cm}^{-1}$  range. The vibrational analysis is guided by state-of-the-art quantum chemical calculations involving hybrid CCSD(T)-based composite schemes and DFT anharmonic calculations. These are combined into hybrid force fields from which vibrational energies, transition strengths, rotational parameters, and coupling constants are determined.

Table 1. Equilibrium Structure of HFO-1123<sup>a</sup>

	CCSD(T)/CBS+CV	CCSD(T) <sup>b</sup>	$\Delta$ CBS <sup>c</sup>	$\Delta$ CV <sup>d</sup>	ChS <sup>e</sup>	B2PLYP <sup>f</sup>	rDSD <sup>g</sup>	rDSD+NL <sup>h</sup>	PW6 <sup>i</sup>	exp. <sup>j</sup>
C <sub>1</sub> =C <sub>2</sub>	1.3223	1.3221	-0.0043	-0.0024	1.3221	1.3224	1.3253	1.3229	1.3221	1.341(12)
C <sub>2</sub> -H <sub>3</sub>	1.0750	1.0745	-0.0012	-0.0011	1.0745	1.0750	1.0774	1.0749	1.0815	1.100(10)
C <sub>1</sub> -F <sub>4</sub>	1.3138	1.3137	-0.0030	-0.0017	1.3137	1.3203	1.3184	1.3143	1.3209	1.316(11)
C <sub>1</sub> -F <sub>5</sub>	1.3083	1.3082	-0.0036	-0.0017	1.3082	1.3147	1.3129	1.3089	1.3153	1.316(11)
C <sub>2</sub> -F <sub>6</sub>	1.3350	1.3350	-0.0024	-0.0018	1.3350	1.3417	1.3397	1.3356	1.3415	1.342(24)
$\angle$ C <sub>1</sub> C <sub>2</sub> H <sub>3</sub>	123.11	123.13	0.38	-0.04	123.13	123.21	123.09	123.09	123.43	124.0(17)
$\angle$ C <sub>2</sub> C <sub>1</sub> F <sub>4</sub>	122.87	122.85	0.11	0.00	122.85	122.96	122.91	122.91	123.06	123.1(15)
$\angle$ C <sub>2</sub> C <sub>1</sub> F <sub>5</sub>	125.13	125.14	-0.03	-0.02	125.14	125.21	125.18	125.18	125.20	124.0(6)
$\angle$ C <sub>1</sub> C <sub>2</sub> F <sub>6</sub>	120.22	120.26	-0.32	0.03	120.26	120.47	120.44	120.44	120.35	120.0(7)

<sup>a</sup>Bond lengths and angles in Å and deg, respectively. <sup>b</sup>f.c.-CCSD(T)/cc-pVTZ level of theory. <sup>c</sup>Correction due to the extrapolation to the CBS using MP2 theory with the cc-pVQZ and cc-pVTZ basis sets. <sup>d</sup>Correction due to core-correlation effects. <sup>e</sup>Best estimate according to the ChS, i.e., CCSD(T)+ $\Delta$ CBS+ $\Delta$ CV. <sup>f</sup>B2PLYP-D3/aug-cc-pVTZ. <sup>g</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>h</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ augmented through Nano-LEGO. CCF angles are not corrected due to the lack of parametrization. <sup>i</sup>PW6B95-D3/jul-cc-pVDZ. <sup>j</sup>From ref 40; figures in parentheses are uncertainties referred to the last significant digits.

## 2. EXPERIMENTAL DETAILS

To record the gas-phase IR spectra, two different Fourier-Transform IR (FTIR) spectrometers were used according to the spectral region. In the range between 200 and 400 cm<sup>-1</sup>, the spectra were recorded at a resolution of 1.0 and 0.5 cm<sup>-1</sup> using a Spectrum One (PerkinElmer) interferometer and a glass cell (optical path of 15 cm) equipped with KRS-5 windows. In the range of 400–5000 cm<sup>-1</sup>, the spectra were measured by a Bruker Vertex 70 spectrometer with a resolution ranging from 1.0 to 0.2 cm<sup>-1</sup> and using a double-walled, stainless-steel cell (optical path of 13.4 cm) equipped with KBr windows. Cross section measurements were performed at a resolution of 0.5 cm<sup>-1</sup>, keeping the cell temperature constant at 293.0 K ( $\pm$ 0.5 K). The pressure of the sample was varied in the range of 1.1–222.5 hPa, and for each experimental run, 128 interferograms were collected to maximize the signal-to-noise ratio. Following the methodologies established in previous works,<sup>33,51</sup> distortions due to finite-resolution effects<sup>52</sup> were minimized by mixing the sample with N<sub>2</sub> (SIAD, purity >99%) up to a total pressure of 101 kPa. Adsorption of the sample on the cell walls, checked by monitoring the pressure inside the cell and inspecting the absorption spectra, was found to be negligible over a period longer than that needed to obtain a spectrum. The gas sample of F<sub>2</sub>C=CHF (99% pure), supplied by Peninsular Chemical Research, Inc., was used without further purification.

## 3. COMPUTATIONAL DETAILS

The equilibrium structure of HFO-1123 was determined according to two different composite schemes relying on the CCSD(T) method. The first one exploits the additivity relation on the energy gradient used in the geometry optimization:<sup>53,54</sup>

$$\frac{dE_{\text{CBS+CV}}}{dx} = \frac{dE_{\text{HF-SCF}}^{\infty}}{dx} + \frac{dE_{\text{CCSD(T)}}^{\infty}}{dx} + \frac{d\Delta E(\text{CV})}{dx} \quad (1)$$

where  $\frac{dE_{\text{HF-SCF}}^{\infty}}{dx}$  and  $\frac{dE_{\text{CCSD(T)}}^{\infty}}{dx}$  are the energy gradients corresponding to the exp(- $Cn$ ) extrapolation to the complete basis set limit (CBS) at the Hartree-Fock self-consistent field level<sup>55</sup> and to the  $n^{-3}$  CBS extrapolation for the correlation contribution,<sup>56</sup> respectively. In these expressions,  $n$  represents the cardinal number of the Dunning's family of correlation consistent basis sets, and specifically, the HF-SCF extrapolation was carried out using the cc-pV $n$ Z with  $n = T, Q$ , and

$S$ ,<sup>57,58</sup> while  $n = T$  and  $Q$  were employed for the correlation contribution. The third term,  $\frac{d\Delta E(\text{CV})}{dx}$ , which accounts for core-valence correlation effects, was computed as the difference between all-electron (a.e.) and frozen core (f.c.) terms computed using the core-valence cc-pCVTZ basis set.<sup>59</sup> The second composite method relies on the cheap scheme (ChS), where the additivity relation is applied directly on the value of the geometrical parameter.<sup>60,61</sup> Briefly, it starts from the geometry optimized at the CCSD(T)/cc-pVTZ level and, on top of it, adds contributions accounting for the extrapolation to the CBS and core-valence correlations, evaluated according to the second-order Møller-Plesset (MP2) perturbation theory<sup>62</sup> in order to limit the computational cost. The best estimates of the harmonic vibrational frequencies were also obtained by applying the ChS, while for IR intensities and centrifugal distortion constants, the following expression was employed:<sup>63</sup>

$$I^{\text{best}} = I^{\text{CCSD(T)/VTZ}} + \Delta I^{\text{T-Q}} + \Delta I(\text{CV}) + \Delta I(\text{aug}) \quad (2)$$

where  $I^{\text{CCSD(T)/VTZ}}$  is the harmonic IR intensity (or the centrifugal distortion constant) obtained at the CCSD(T)/cc-pVTZ level;  $\Delta I^{\text{T-Q}}$  represents the correction due to the enlargement of the basis set obtained as the difference between the MP2 values obtained with the cc-pVQZ and cc-pVTZ basis;  $\Delta I(\text{aug})$  takes into account the effects of diffuse functions, which are particularly relevant for intensity calculations (it is calculated as the difference between MP2 values obtained with the aug-cc-pVTZ and cc-pVTZ basis sets);  $\Delta I(\text{CV})$  has its usual meaning. All the required harmonic force fields were obtained through analytic evaluation of the Hessian matrix.

On the basis of the recent literature,<sup>64,65</sup> full anharmonic calculations were carried out at the DFT level by adopting the double-hybrid B2PLYP<sup>66</sup> and rev-DSDPBEP86<sup>67</sup> functionals coupled to the aug-cc-pVTZ and jun-cc-pVTZ basis sets<sup>68</sup> as well as the PW6B95 functional<sup>69</sup> in conjunction with the jul-cc-pVDZ basis set.<sup>68</sup> All the functionals were corrected for dispersion correlation effects by means of the Grimme's DFT-D3 method<sup>70</sup> with the use of the Beck-Johnson damping,<sup>71</sup> Cubic, and semidiagonal quartic force constants, and up to the third-derivatives of the dipole moment were obtained by numerical differentiation of displaced analytic Hessians and electric dipole moment first derivatives along the normal coordinates, respectively. Vibrational corrections to rotational

constants, vibrational wavenumbers, and IR intensities beyond the double-harmonic approximation were obtained by applying vibrational perturbation theory to second-order (VPT2)<sup>72–74</sup> on the anharmonic force field. To improve the accuracy of the computed vibrational frequencies, a hybrid force field<sup>36,65</sup> was devised by mixing the harmonic vibrational wavenumbers evaluated in the frame of the ChS with the cubic- and quartic-force constants computed by means of double-hybrid functionals. In order to tackle the problem of resonances plaguing the VPT2 approach, generalized second-order vibrational perturbation theory (GVPT2) was adopted in which (near-) singular terms are removed from the perturbative summations (leading to the so-called deperturbed approach) and the energy levels coupled by the resonances are treated in a second step by a proper variational calculation of reduced dimensionality.<sup>75,76</sup> All DFT and MP2 calculations were performed with the Gaussian 16 quantum chemical package,<sup>77</sup> while for CCSD(T) computations, the CFOUR program<sup>78</sup> was employed. Spectroscopic parameters were evaluated according to the general VPT2 engine, implemented within the Gaussian software, and IR intensities evaluated up to three quanta of vibrational excitation.<sup>79</sup>

#### 4. RESULTS AND DISCUSSION

The F<sub>2</sub>C=CHF molecule is a planar near-prolate asymmetric rotor belonging to the C<sub>s</sub> symmetry point group. It has 12 normal modes of vibration classified in terms of symmetry species, as 9A' ⊕ 3A'', with A' vibrations producing A/B hybrid bands with different contributions of the components and A'' vibrations giving rise C-type band envelopes.

**4.1. Equilibrium Geometry and Rotational Spectroscopic Parameters.** The HFO-1123 equilibrium structure is completely defined by nine structural parameters that are reported in Table 1 at different levels of theory with the labeling of the atoms as illustrated in Figure 1. More precisely,

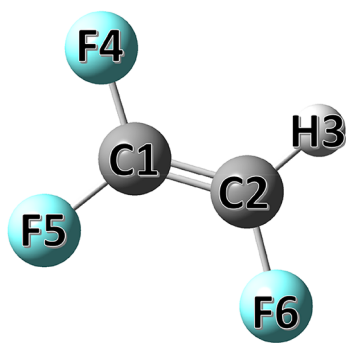


Figure 1. Molecular structure of HFO-1123 with atom labeling.

the table shows the results obtained from both the CCSD(T)/CBS+CV and the ChS and, for the latter, the f.c.-CCSD(T)/cc-pVTZ results are reported together with the CBS and CV corrections. As it can be seen from the results, the CCSD(T)/CBS+CV and the ChS yield the same predictions of the structural parameters with differences within 0.5 mÅ for bond lengths and 0.04° for bond angles. The expected accuracy for geometrical parameters obtained from the CCSD(T)/CBS+CV methods is 0.001–0.002 Å for distances and 0.05–0.1° for angles.<sup>36,53,54</sup> An inspection of the table reveals that the CCSD(T)/cc-pVTZ and the double-hybrid functionals yield predictions of the equilibrium geometry of similar quality with

differences, measured with respect to the CCSD(T)/CBS+CV structure, up to 0.007 Å and about 0.3° for distances and angles, respectively, while at the PW6B95, deviations increase slightly. The ΔCBS and ΔCV contributions are significant for the improvement of the CCSD(T)/cc-pVTZ accuracy. This is particularly true for bond lengths, systematically overestimated at the CCSD(T)/cc-pVTZ level (between 0.002 and 0.007 Å), for which both contributions are negative and range from –1 to –4 mÅ. Less systematic are the corrections for the bond angles for which the ΔCBS term ranges between –0.3° (∠ C<sub>1</sub>C<sub>2</sub>F<sub>6</sub>) and 0.4° (C<sub>1</sub>C<sub>2</sub>H<sub>3</sub>), while the ΔCV contribution is almost negligible. A comment is deserved concerning the comparison to the experimental structure derived by Mom et al.<sup>40</sup> When this geometry is compared against theoretical predictions, huge differences as large as 0.025 Å (C<sub>2</sub>–H<sub>3</sub>) and –1.1° (∠ C<sub>2</sub>C<sub>1</sub>F<sub>5</sub>) are noticed. In part, this is due to the fact that ref 40 reports an effective structure obtained from the fitting of the ground state rotational constants and, therefore, contaminated by the vibrational motion. Second and most importantly, because of the insufficient number of isotopic substitutions, limited to the main isotopologue and the deuterated species, the effective geometry was obtained by resorting to a number of approximations, like the equivalency of the C<sub>1</sub>–F<sub>4</sub> and C<sub>1</sub>–F<sub>5</sub> bond lengths, which is clearly not the case differing by about 6 mÅ. As mentioned above, the rev-DSDPBEP86 functional overestimates all the bond lengths. Hence, to improve its predictions the recently proposed Nano-LEGO<sup>80</sup> tool has been applied, and the results are also reported in Table 1. The Nano-LEGO augmented geometrical parameters are in an impressive agreement with those obtained by the most refined (and computationally demanding) composite schemes with deviations well within 0.001 Å from the CCSD(T)/CBS+CV bond lengths and a maximum difference of 0.2° for C<sub>1</sub>C<sub>2</sub>F<sub>6</sub>, which however, has not been corrected due to the lack of Nano-LEGO parametrization for this angle.

The spectroscopic parameters relevant for rotational spectroscopy, specifically equilibrium and ground state rotational constants as well as quartic centrifugal distortion terms, are summarized in Table 2, where theoretical predictions are compared against the experimental data in the *I'* representation of the Watson's A reduced Hamiltonian.<sup>81</sup> A well consolidated procedure was followed,<sup>30,38,80</sup> and the best estimates of ground state rotational constants have been obtained by augmenting those of the equilibrium configuration computed according to the CCSD(T)/CBS+CV composite scheme through rev-DSDPBEP86-D3/jun-cc-pVTZ vibrational corrections. The resulting parameters are in very good agreement with the corresponding experimental counterparts with discrepancies of 0.06%, thus confirming the accuracy of the CCSD(T)/CBS+CV equilibrium geometry as well as the reliability of the rev-DSDPBEP86 vibrational contributions. Analogous results (not reported) have been obtained using both the ChS and the Nano-LEGO augmented rev-DSDPBEP86 equilibrium rotational constants. The accuracy of the ground state rotational constants evaluated at the CCSD(T)/cc-pVTZ, B2LYP-D3/aug-cc-pVTZ, and rev-DSDPBEP86-D3/jun-cc-pVTZ levels of theory are similar with rev-DSDPBEP86 providing slightly better results and mirror that of the geometry: the three methods underestimate the experimental values and show deviations around –0.9%, –0.7%, and –0.6%, respectively. Upon inspection of Table 2, it can be observed that CCSD(T)/cc-pVTZ and the double-

Table 2. Rotational and Quartic Centrifugal Distortion Constants of HFO-1123 (MHz)

	best estimate <sup>a</sup>	CCSD(T) <sup>b</sup>	B2PLYP <sup>c</sup>	rev-DSD <sup>d</sup>	exp. <sup>e</sup>
$A_e$	10 723.27	10 626.84	10 650.43	10 660.92	
$B_e$	3887.73	3850.17	3855.48	3857.31	
$C_e$	2853.27	2826.22	2830.73	2832.47	
$A_0$	10 671.89	10 575.09	10 599.04	10 609.33	10 665.481287(51)
$B_0$	3874.60	3836.81	3842.35	3844.15	3872.406579(24)
$C_0$	2839.57	2812.40	2817.03	2818.74	2837.960953(29)
$\Delta_j \times 10^3$	0.7285	0.7075	0.7111	0.7099	0.731145(12)
$\Delta_{jK} \times 10^3$	7.8681	7.5942	7.6282	7.6187	7.671250(52)
$\Delta_K \times 10^3$	4.8572	4.6008	4.8030	4.7438	4.92912(16)
$\delta_j \times 10^3$	0.1811	0.1769	0.1775	0.1772	0.1831457(50)
$\delta_K \times 10^3$	4.8543	4.7171	4.7329	4.7288	4.83607(14)

<sup>a</sup>CCSD(T)/CBS+CV equilibrium rotational constants, rev-DSDPBEP86-D3/jun-cc-pVTZ vibrational corrections, and ChS quartic centrifugal distortion constants. <sup>b</sup>CCSD(T)/cc-pVTZ. <sup>c</sup>B2PLYP-D3/aug-cc-pVTZ. <sup>d</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>e</sup>From ref 42; values refer to the A-reduction Watson's Hamiltonian in the  $F$  representation. The figures in parentheses are uncertainties referred to the last significant digits.

Table 3. Theoretical Sextic Centrifugal Distortion Constants (A-Reduction Watson's Hamiltonian in the  $F$  Representation) of HFO-1123 (Hz) and Comparison to Experimental Values

	B2PLYP <sup>a</sup>	rev-DSD <sup>b</sup>	exp. <sup>c</sup>	exp. <sup>d</sup>
$\Phi_j \times 10^3$	0.214	0.207	0.038(20)	0.2010(29)
$\Phi_{jK}$	0.02064	0.02011	0.0027(105)	0.019977(56)
$\Phi_{Kj}$	-0.05331	-0.05313	0.0025(288)	-0.05529(18)
$\Phi_K$	0.07050	0.07076	0.033(21)	0.07534(20)
$\phi_j \times 10^3$	0.06858	0.06659	-0.02(18)	0.0765(15)
$\phi_{jK} \times 10^3$	10.04	9.77	14.4(63)	9.352(55)
$\phi_K$	0.11355	0.11277	-0.041(78)	0.11635(37)

<sup>a</sup>B2PLYP-D3/aug-cc-pVTZ. <sup>b</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>c</sup>From ref 41; figures in parentheses are uncertainties referred to the last significant digits. <sup>d</sup>From ref 42; figures in parentheses are uncertainties referred to the last significant digits.

Table 4. Rotational Constants (MHz) of the Excited Fundamental Vibrational Levels of HFO-1123<sup>a</sup>

vibrational level	$A^v$	$B^v$	$C^v$
$\nu_1 = 1$	10 654.68	3871.27	2836.66
$\nu_2 = 1$	10 646.88	3861.08	2831.09
$\nu_3 = 1$	10 647.42 (10 638.50)	3874.15 (3871.10)	2836.39 (2834.17)
$\nu_4 = 1$	10 655.46 (10 652.71) <sup>b</sup>	3864.76 (3863.89) <sup>b</sup>	2830.22 (2829.71) <sup>b</sup>
$\nu_5 = 1$	10 682.59 (10 651.20) <sup>c</sup>	3869.29 (3867.49) <sup>c</sup>	2837.08 (2833.86) <sup>c</sup>
$\nu_6 = 1$	10 647.03 (10 641.54) <sup>d</sup>	3869.98 (3867.84) <sup>d</sup>	2833.37 (2831.90) <sup>d</sup>
$\nu_7 = 1$	10 681.78	3878.58	2840.14
$\nu_8 = 1$	10 681.66 (10 673.41) <sup>e</sup>	3876.78 (3874.51) <sup>e</sup>	2837.35 (2835.74) <sup>e</sup>
$\nu_9 = 1$	10 637.11 (10 624.33) <sup>e</sup>	3873.82 (3870.68) <sup>e</sup>	2837.47 (2835.67) <sup>e</sup>
$\nu_{10} = 1$	10 671.26	3874.57	2842.42
$\nu_{11} = 1$	10 656.06	3872.38	2841.43
$\nu_{12} = 1$	10 699.53 (10 698.10) <sup>e</sup>	3880.83 (3879.12) <sup>e</sup>	2843.05 (2841.51) <sup>e</sup>

<sup>a</sup>Predicted values obtained by correcting CCSD(T)/CBS+CV equilibrium rotational constants through vibrational contributions at rev-DSDPBEP86-D3/jun-cc-pVTZ. Reported in parentheses are experimental values. <sup>b</sup>From ref 45. <sup>c</sup>From ref 46. <sup>d</sup>From ref 47. <sup>e</sup>From ref 42.

hybrid computations also systematically underestimate quartic centrifugal parameters with absolute deviations in the ranges of 1.1–6.7%, 0.6–3.2%, and 0.7% – 3.6% respectively, an accuracy coherent with previous benchmark studies.<sup>64,65,82</sup> The ChS results reproduce quartic centrifugal distortion parameters very well, the absolute percentage deviations being in the range of 0.3–2.6%, thus sensibly improving the CCSD(T)/cc-pVTZ outcomes in all cases but the  $\Delta_{jK}$  term.

The sextic centrifugal distortion constants computed at different levels of theory are reported in Table 3 together with those determined experimentally in ref 41 and more recently by Tamassia et al.<sup>42</sup> within the  $F$  representation of the Watson's  $A$  reduced Hamiltonian.<sup>81</sup> The inspection of this table reveals huge discrepancies between the two sets of

experimental values, even though sextic centrifugal distortion parameters determined in ref 41 suffer from large uncertainties and, in practice, cannot be considered well determined. On the other side, according to the recent literature on the subject,<sup>83,84</sup> the computed sextic centrifugal distortion constants are expected to have an average accuracy of around 10%, which is indeed fully matched by the more recent experimental values<sup>42</sup> with which the average agreement is around 5% for both the functionals, and the maximum discrepancies, reported for the  $\phi_j$  term, amount to 10% and 13% at the B2PLYP and rev-DSDPBEP86-D3 levels, respectively. Therefore, the present theoretical outcomes confirm the reliability of the ground state rotational centrifugal distortion terms measured by Tamassia et al.<sup>42</sup>

Before moving to the characterization of the vibrational properties of HFO-1123, the attention is briefly moved to the ro-vibrational spectroscopic parameters. In particular, Table 4 shows the best estimates of the rotational constants for the excited fundamental vibrational levels obtained by correcting the best estimated equilibrium rotational constants through vibrational contributions evaluated at the rev-DSDPBEP86-D3/jun-cc-pVTZ. It should be noted that, according to the VPT2 framework, the vibrational dependence of the rotational constants is ruled by the  $\alpha_k^\beta$  vibration–rotation interaction constants that, for the sake of completeness, are reported in Table 5. The rotational constants of  $\nu_3 = 1$ ,  $\nu_4 = 1$ ,  $\nu_5 = 1$ ,  $\nu_6 =$

**Table 5.**  $\alpha_k^\beta$  Vibrational Interaction Constants (MHz) of HFO-1123 Evaluated at the rev-DSDPBEP86-D3/jun-cc-pVTZ Level of Theory

normal mode	<i>a</i>	<i>b</i>	<i>c</i>
1	17.326	3.217	2.859
2	25.120	13.421	8.431
3	24.577	0.352	3.129
4	16.551	9.740	9.309
5	−10.596	5.198	2.431
6	24.963	4.528	6.157
7	−9.784	−4.101	−0.635
8	−9.648	−2.295	2.164
9	34.886	0.689	2.041
10	0.734	−0.070	−2.911
11	15.952	2.129	−1.924
12	−27.518	−6.346	−3.534

1,  $\nu_8 = 1$ ,  $\nu_9 = 1$ , and  $\nu_{12} = 1$  vibrational levels evaluated theoretically can be compared with those determined experimentally,<sup>42,45–47</sup> which are listed as well in Table 4 within the parentheses. For these, a very good agreement is noted between the predicted and measured values with errors in all cases well within 0.1% with the only exception of the *A* rotational constant of the  $\nu_9 = 1$  and  $\nu_5 = 1$  levels. While the former deviates by only 0.12% from the experimental counterpart, the latter show a larger discrepancy, around 0.3%. It is worthwhile to note, however, that the experimental value may be affected by spurious Coriolis and anharmonic interactions. It can be concluded that the rotational constants reported in Table 4 almost retain the same accuracy as that reached for the ground state rotational constants, thus highlighting the reliability of the computed  $\alpha_k^\beta$  constants, and hence, they can be employed as a guide for future high-resolution investigations on this molecule. For this purpose, the most relevant Coriolis coupling constants computed at the rev-DSDPBEP86-D3/jun-cc-pVTZ are also given in Table 6.

**4.2. Harmonic Vibrational Frequencies and IR Intensities.** Harmonic wavenumbers and IR intensities calculated within the double-harmonic approximation at different levels of theory are summarized in Tables 7 and 8, respectively. In addition to the best-estimated values from the ChS method, these tables also list its different contributions as well as predictions from the B2PLYP-D3, rev-DSDPBEP86-D3, and PW6B95-D3 density functionals. Concerning wavenumbers, corrections due to the CBS extrapolation are always negative, and they range between  $-1$  and  $-7$   $\text{cm}^{-1}$  with the exception of  $\omega_3$ ,  $\omega_4$ , and  $\omega_5$  for which the contributions are  $-20$ ,  $-16$ , and  $-12$   $\text{cm}^{-1}$ , respectively. Diffuse functions also provide a negative contribution between  $-2.7$   $\text{cm}^{-1}$  ( $\omega_6$ ) and

$-19.7$   $\text{cm}^{-1}$  ( $\omega_3$ ) and usually are of a larger magnitude than the  $\Delta\text{CBS}$  term. Conversely, the core–valence correlation yields positive corrections, generally of smaller magnitude, being  $6$   $\text{cm}^{-1}$  at most in the case of the  $\omega_2$  normal mode. Summed up, the different contributions cause a blue-shift of the CCSD(T)/cc-pVTZ harmonic wavenumbers for all the vibrational normal modes, which reaches  $-36.6$  and  $-30.1$   $\text{cm}^{-1}$  for the  $\omega_3$  and  $\omega_4$  vibrations, respectively. Concerning the intensities, all three corrections can be either positive or negative and, in absolute terms, they are generally small. In particular, core–valence contributions are always lower than  $0.7$   $\text{km mol}^{-1}$ , and in general, they are almost negligible. Likewise, the  $\Delta I(\text{aug})$  term is always lower than  $0.8$   $\text{km mol}^{-1}$  with exceptions being given by  $\omega_4$  and  $\omega_5$  for which it amounts to  $8.1$  and  $5.1$   $\text{km mol}^{-1}$ , respectively, and  $\omega_1$  for which it is  $1.6$   $\text{km mol}^{-1}$ . The latter is small in absolute value but, in relative terms, it represents 14.1% of the overall ChS harmonic IR intensity for this normal mode. The largest CBS correction, for the  $\omega_4$  vibration, is around  $9$   $\text{km mol}^{-1}$ , which corresponds to only 5% of the overall intensity. Conversely, the CBS correction for  $\omega_1$  amounts to  $1.9$   $\text{km mol}^{-1}$ , which accounts for almost 17% of the ChS IR intensity.

When the ChS best estimates are taken as the reference, it is seen that the double-hybrid functionals yield, on the whole, similar predictions with a mean absolute deviation (MAD) from ChS harmonic frequencies of  $6.7$  and  $4.0$   $\text{cm}^{-1}$  at the B2PLYP-D3/aug-cc-pVTZ and rev-DSDPBEP86-D3/jun-cc-pVTZ levels of theory, respectively. For the former, the maximum deviations, around  $14$   $\text{cm}^{-1}$  but in opposite directions, are observed for  $\omega_4$  and  $\omega_{11}$ , which is also responsible for the largest deviation ( $18$   $\text{cm}^{-1}$ ) observed among the rev-DSDPBEP86-D3 harmonic frequencies. This functional shows a better agreement to the ChS harmonic intensities than B2PLYP-D3. Indeed, the former presents a MAD of c.a.  $2$   $\text{km mol}^{-1}$  and maximum discrepancies in the range of  $-2.6$ – $10$   $\text{km mol}^{-1}$ , while the latter reproduces the best estimates with a MAD of  $5$   $\text{km mol}^{-1}$  and deviations between  $-14.5$  and  $20.8$   $\text{km mol}^{-1}$  for the intensities of normal modes 3 and 5, respectively. As expected, the hybrid PW6B95-D3 functional shows slightly larger deviations for harmonic wavenumbers with a MAD of around  $13$   $\text{cm}^{-1}$  and maximum deviation of about  $35$   $\text{cm}^{-1}$  for  $\omega_2$ . On the other hand, it well reproduces ChS harmonic intensities with the MAD being  $3.6$   $\text{km mol}^{-1}$ .

**4.3. Anharmonic Force Field and Characterization of the Gas-Phase Spectra.** Fundamental vibrational frequencies calculated at different levels of theory are presented in Table 9, where they are also compared to the wavenumbers measured experimentally, while theoretical IR intensities beyond the double-harmonic approximation are collected in Table 10. Following the outcomes obtained for harmonic frequencies and intensities, in addition to full anharmonic computations at the rev-DSDPBEP86-D3/jun-cc-pVTZ and PW6B95-D3/jul-cc-pVDZ, a hybrid force field, termed ChS:rDSD, has been worked out. This has harmonic frequencies estimated by the ChS method mixed with cubic and semidiagonal quartic force constants evaluated at the rev-DSDPBEP86-D3 level, while hybrid anharmonic intensities have been obtained from ChS and rev-DSDPBEP86-D3 contributions, according to eq 2. According to the available literature, the adopted hybrid force field is expected to predict fundamental transition frequencies with an average accuracy within  $5$   $\text{cm}^{-1}$ , a maximum error of  $10$   $\text{cm}^{-1}$ , and anharmonic IR intensities with an error of a few

**Table 6. Relevant Coriolis Coupling Constants of HFO-1123 Evaluated at the rev-DSDPBEP86-D3/jun-cc-pVTZ Level of Theory**

A-type Coriolis			B-type Coriolis			C-type Coriolis		
mode k	mode l	$ \zeta_{kl}^a $	mode k	mode l	$ \zeta_{kl}^b $	mode k	mode l	$ \zeta_{kl}^c $
1	10	0.967	1	10	0.106	1	2	0.286
1	11	0.147	2	10	0.457	1	3	0.645
1	12	0.106	2	11	0.739	1	4	0.610
2	11	0.413	2	12	0.162	1	5	0.278
3	10	0.139	3	10	0.632	1	6	0.178
3	11	0.713	3	12	0.137	1	9	0.117
3	12	0.135	4	10	0.564	2	3	0.549
4	11	0.347	4	11	0.572	2	4	0.401
4	12	0.152	4	12	0.401	2	5	0.460
5	11	0.361	5	10	0.168	2	6	0.246
5	12	0.466	5	11	0.121	2	7	0.366
6	10	0.165	5	12	0.696	2	8	0.191
6	12	0.302	6	10	0.124	2	9	0.107
7	11	0.118	6	12	0.339	3	4	0.325
7	12	0.459	7	11	0.248	3	7	0.310
8	11	0.165	8	11	0.106	3	8	0.230
8	12	0.377	8	12	0.122	3	9	0.110
9	12	0.533	9	10	0.127	4	6	0.188
			9	11	0.182	4	7	0.247
			9	12	0.423	4	8	0.453
						4	9	0.236
						5	6	0.457
						5	7	0.332
						5	9	0.586
						6	7	0.161
						6	8	0.147
						6	9	0.394
						7	8	0.518
						8	9	0.556

**Table 7. Approximate Description and Harmonic Wavenumbers ( $\text{cm}^{-1}$ ) Obtained at Different Levels of Theory for the HFO-1123 Vibrational Normal Modes**

sym.	mode	approximate description	$\omega^{\text{CCSD(T)}^a}$	$\Delta\omega[\text{CBS(T,Q)}]^b$	$\Delta\omega(\text{CV})^c$	$\Delta\omega(\text{aug})^d$	$\omega^{\text{ChS}^e}$	$\omega^{\text{B2f}^f}$	$\omega^{\text{rDSDg}}$	$\omega^{\text{PW6h}^h}$
A'	$\omega_1$	CH stretch	3272.0	-0.3	5.1	-4.5	3272.3	3279.9	3277.4	3296.9
	$\omega_2$	CC stretch	1836.3	-6.7	6.0	-11.3	1824.3	1827.9	1837.1	1871.2
	$\omega_3$	CH bend + CF <sub>2</sub> asym. stretch	1402.4	-20.4	3.5	-19.7	1365.8	1373.1	1384.2	1365.2
	$\omega_4$	CF <sub>2</sub> asym. stretch + CH bend	1301.5	-16.4	3.3	-16.9	1271.4	1274.1	1286.2	1270.3
	$\omega_5$	CF stretch (CHF) + CH bend	1196.1	-12.3	3.2	-13.9	1173.1	1175.4	1185.1	1180.4
	$\omega_6$	CF <sub>2</sub> sym. stretch + CH bend	944.1	-3.1	2.8	-6.3	937.5	941.9	945.8	944.9
	$\omega_7$	CHF rock + CF <sub>2</sub> scissor	628.8	-2.7	2.0	-5.0	623.2	626.6	628.9	622.4
	$\omega_8$	CF <sub>2</sub> scissor	488.9	-1.8	1.6	-4.8	484.0	486.5	488.1	480.9
	$\omega_9$	CF <sub>2</sub> rock	232.9	-1.5	0.9	-2.7	229.6	233.1	233.1	225.2
A''	$\omega_{10}$	CHF wag	771.2	-7.0	4.1	-16.3	752.0	778.7	783.1	783.0
	$\omega_{11}$	CF <sub>2</sub> wag	564.6	-2.6	3.3	-11.6	553.7	579.4	583.4	586.7
	$\omega_{12}$	C=C torsion	310.3	-1.8	1.0	-4.4	305.0	310.1	313.0	310.8

<sup>a</sup>f.c.-CCSD(T)/cc-pVTZ. <sup>b</sup>CBS contribution based on MP2 computations with cc-pVTZ and cc-pVTZ basis sets. <sup>c</sup>Correction due to core-correlation effects. <sup>d</sup>Contribution from diffuse functions. <sup>e</sup>Best estimate according to ChS, i.e., CCSD(T)+ $\Delta[\text{CBS(T,Q)}]+\Delta(\text{CV})+\Delta(\text{aug})$ . <sup>f</sup>B2PLYP-D3/aug-cc-pVTZ. <sup>g</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>h</sup>PW6B95-D3/jul-cc-pVDZ.

km mol<sup>-1</sup>; hence, ChS:rDSD simulations have been used to guide and support the analysis of the observed spectra.<sup>30,36,38,64,65</sup>

Survey gas-phase IR spectra of HFO-1123 in the ranges of 200–400 cm<sup>-1</sup> and 400–5000 cm<sup>-1</sup> are reported in Figure 2 in which some relevant absorptions are also labeled. The former region is characterized by the absorptions of the lowest lying fundamentals: the  $\nu_9$  vibration, corresponding to the CF<sub>2</sub>

rocking is located at 232.9 cm<sup>-1</sup>, and the  $\nu_{12}$  normal mode, due to the C=C torsion, measured at 302.7 cm<sup>-1</sup> in agreement with previous measurements.<sup>43</sup> Both of them perfectly match the theoretical predictions of the ChS:rDSD hybrid force field that places the  $\nu_9$  and  $\nu_{12}$  transitions at 232 and 304 cm<sup>-1</sup>, respectively. In addition to the fundamentals, the  $2\nu_{12}-\nu_{12}$  hot band has been observed, as detailed in Table 11 that shows the full list of assigned vibrational transitions and

**Table 8. Harmonic Intensities ( $\text{km mol}^{-1}$ ) Obtained at Different Levels of Theory for the HFO-1123 Vibrational Normal Modes**

sym.	mode	$I^{\text{CCSD(T)}^a}$	$\Delta I[\text{CBS(T,Q)}]^b$	$\Delta I(\text{CV})^c$	$\Delta I(\text{aug})^d$	$I^{\text{ChS}^e}$	$I^{\text{B2}^f}$	$I^{\text{rDSD}^g}$	$I^{\text{PW6}^h}$
A'	$\omega_1$	7.76	1.90	0.13	1.60	11.38	10.49	9.91	11.44
	$\omega_2$	66.47	1.17	0.14	0.79	68.58	77.97	76.03	79.56
	$\omega_3$	121.93	-2.75	0.63	-0.10	119.71	105.19	117.09	136.21
	$\omega_4$	179.22	9.48	-0.39	8.10	196.41	205.94	197.56	200.12
	$\omega_5$	131.68	3.63	-0.05	5.11	140.38	161.13	150.40	145.04
	$\omega_6$	55.98	0.82	0.12	0.09	57.01	58.57	57.84	60.37
	$\omega_7$	3.50	0.18	-0.02	0.25	3.90	3.36	3.57	3.51
	$\omega_8$	2.12	-0.05	-0.01	0.05	2.11	1.84	1.91	2.09
	$\omega_9$	4.11	-0.15	-0.01	-0.11	3.85	3.94	3.97	3.97
A''	$\omega_{10}$	30.64	-0.44	0.21	0.05	30.46	34.11	33.18	33.43
	$\omega_{11}$	0.44	-0.33	-0.06	-0.19	0.00	0.00	0.01	0.00
	$\omega_{12}$	4.24	0.13	0.03	-0.09	4.31	4.02	3.98	4.42

<sup>a</sup>f.c.-CCSD(T)/cc-pVTZ. <sup>b</sup>CBS contribution based on MP2 computations with cc-pVTZ and cc-pVTZ basis sets. <sup>c</sup>Correction due to core-correlation effects. <sup>d</sup>Contribution from diffuse functions. <sup>e</sup>Best estimate according to ChS, i.e., CCSD(T)+ $\Delta[\text{CBS(T,Q)}]+\Delta(\text{CV})+\Delta(\text{aug})$ . <sup>f</sup>B2PLYP-D3/aug-cc-pVTZ. <sup>g</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>h</sup>PW6B95-D3/jul-cc-pVDZ.

**Table 9. Experimental and Theoretical Fundamental Wavenumbers ( $\text{cm}^{-1}$ ) of the HFO-1123 Normal Modes of Vibration**

symmetry	mode	exp.	ChS:rDSD <sup>a</sup>	rDSD <sup>b</sup>	PW6 <sup>c</sup>
A'	$\nu_1$	3163.65	3174 (3148) <sup>d</sup>	3176	3187
	$\nu_2$	1787.42	1794	1796	1825
	$\nu_3$	1360.8	1357	1357	1336
	$\nu_4$	1264.78	1261	1259	1244
	$\nu_5$	1172.41	1171 (1165) <sup>d</sup>	1172	1167
	$\nu_6$	929.5	929	931	931
	$\nu_7$	623.83	624	624	620
	$\nu_8$	484.93	485	484	477
	$\nu_9$	232.9	232	233	228
A''	$\nu_{10}$	750.59	749	764	770
	$\nu_{11}$	553.76	554	573	576
	$\nu_{12}$	305.1	304	307	306
ME <sup>e</sup>			-0.4	-3.9	-3.1
MAE <sup>f</sup>			2.6	5.5	14.6
max. neg. <sup>g</sup>			-10.4	-19.2	-37.9
max. pos. <sup>h</sup>			3.8	5.3	20.8

<sup>a</sup>Hybrid anharmonic wavenumbers obtained by mixing ChS harmonic properties with cubic and semidiagonal quartic force constants at the rev-DSDPBEP86-D3/jun-cc-pVTZ level. <sup>b</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>c</sup>PW6B95-D3/jul-cc-pVDZ. <sup>d</sup>Deperturbed value within parentheses. <sup>e</sup>Mean error. <sup>f</sup>Mean absolute error. <sup>g</sup>Maximum negative error. <sup>h</sup>Maximum positive error.

where the frequencies measured experimentally are compared with those from the ChS:rDSD predictions. The comparison, in terms of differences between the theoretical and experimental values as a function of increasing wavenumbers, is illustrated in Figure 3 in which the data are reported grouped according to the transition type (i.e., fundamentals, overtones, two- and three-quanta combinations, and hot bands). As it can be seen, the almost completeness of the transitions assigned experimentally are reproduced within  $10 \text{ cm}^{-1}$ , and only four bands fall outside this range. While deviations appear randomly scattered around zero, the only dependence that can be inferred to some extent is that most experimental values are underestimated below c.a.  $2000 \text{ cm}^{-1}$ , a trend that reverses moving toward higher frequencies. At room temperature, the populations of the  $\nu_9 = 1$  and  $\nu_{12} = 1$  levels are about 32% and

**Table 10. Theoretical Anharmonic IR Intensities ( $\text{km mol}^{-1}$ ) of the HFO-1123 Fundamental Transitions**

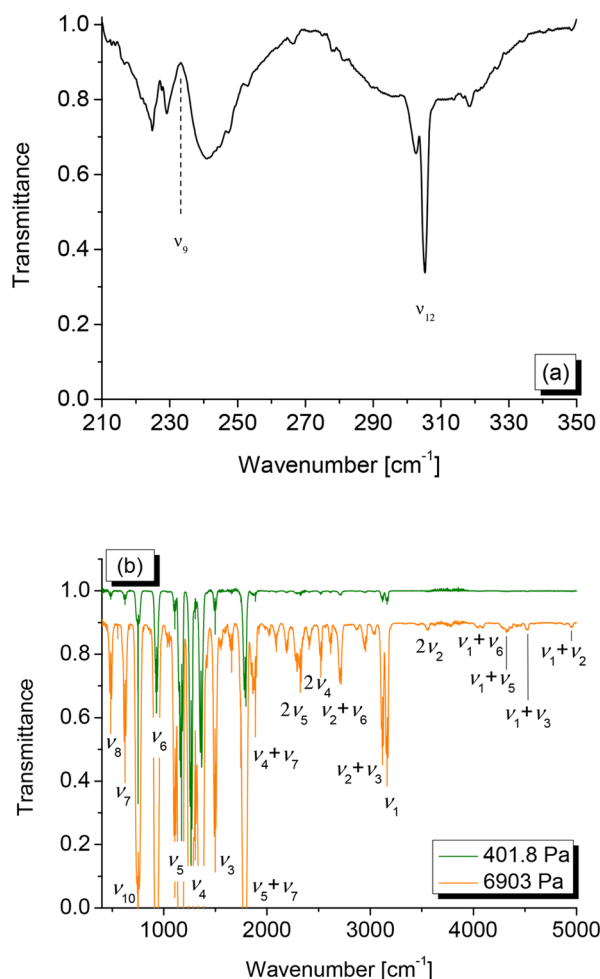
symmetry	mode	ChS:rDSD <sup>a</sup>	rDSD <sup>b</sup>	PW6 <sup>c</sup>
A'	$\nu_1$	6.63	5.16	6.33
	$\nu_2$	62.42	69.88	60.49
	$\nu_3$	113.34	110.72	128.16
	$\nu_4$	175.67	176.82	171.39
	$\nu_5$	51.36	61.38	54.09
	$\nu_6$	54.94	55.77	58.00
	$\nu_7$	3.19	2.87	2.39
	$\nu_8$	2.05	1.86	2.02
	$\nu_9$	3.86	3.98	3.98
A''	$\nu_{10}$	29.27	31.98	31.46
	$\nu_{11}$	0.00	0.01	0.01
	$\nu_{12}$	4.37	4.05	4.32

<sup>a</sup>Hybrid intensities obtained from ChS harmonic intensities and rev-DSDPBEP86-D3/jun-cc-pVTZ anharmonic contributions. <sup>b</sup>rev-DSDPBEP86-D3/jun-cc-pVTZ. <sup>c</sup>PW6B95-D3/jul-cc-pVDZ.

23% of the vibrational ground state; hence, a number of hot bands has to be expected in addition to cold transitions.

The spectral portion between  $400$  and  $1000 \text{ cm}^{-1}$  features a number of absorptions mainly related to the fundamental transitions  $\nu_8$ ,  $\nu_7$ ,  $\nu_{10}$ , and  $\nu_6$  and their hot bands involving, as expected, the  $\nu_9 = 1$  and  $\nu_{12} = 1$  excited levels. Among these, the most prominent ones are  $\nu_6$ , the  $\text{CF}_2$  symmetric stretching, and  $\nu_{10}$ , the CHF wagging, with a predicted intensity of  $54.9$  and  $29.3 \text{ km mol}^{-1}$  and located at  $929.5$  and  $750.6 \text{ cm}^{-1}$ , respectively, in perfect agreement with quantum chemical predictions ( $929$  and  $749 \text{ cm}^{-1}$ ). The CHF rocking and  $\text{CF}_2$  scissoring motions correspond to the  $\nu_7$  and  $\nu_8$  normal modes, respectively, that have similar intensities and give rise to the bands observed at  $623.8$  and  $484.9 \text{ cm}^{-1}$ . In between these two bands, the very weak  $\nu_{11}$  feature, stemming from the  $\text{CF}_2$  wagging motion, can be detected at  $553.8 \text{ cm}^{-1}$  only in the spectra recorded at the higher pressures, again in perfect accord with the ChS:rDSD predictions ( $554 \text{ cm}^{-1}$ ) that significantly improves the results obtained from the rev-DSDPBEP86 and PW6B95 functionals. The fundamental transitions  $\nu_5$ ,  $\nu_4$ , and  $\nu_3$ , associated, respectively, to the C–F stretching of the CHF group, the  $\text{CF}_2$  asymmetric stretching, and the CH in-plane bending motion cover the  $1000$ – $1700$





**Figure 2.** Gas-phase spectrum of HFO-1123. (a) Spectral region between 210 and 350  $\text{cm}^{-1}$  (resolution =  $0.5 \text{ cm}^{-1}$ , optical path = 15 cm, temperature = 293 K, sample pressure = 48.2 hPa). (b) Spectral region between 400 and 5000  $\text{cm}^{-1}$  (resolution =  $0.5 \text{ cm}^{-1}$ , optical path = 13.4 cm, temperature = 298 K): the upper trace (green) corresponds to a sample pressure of 401.8 Pa, and the lower trace (orange, displaced downward for clarity), corresponds to a sample pressure of 69.03 hPa.

$\text{cm}^{-1}$  spectral portion. The  $\nu_5$  and  $\nu_4$  modes give rise to two A/B hybrid bands with a prevalence of A-type character located at 1172.4 and 1264.8  $\text{cm}^{-1}$  and with a predicted intensity of 51.4 and 175.7  $\text{km mol}^{-1}$ , respectively. The  $\nu_5 = 1$  level is also involved in a type II Fermi interaction with the  $\nu_6 = \nu_9 = 1$  level, whose corresponding combination band is observed on the low-frequency side of  $\nu_5$  at 1155.1  $\text{cm}^{-1}$ . According to the ChS:rDSD hybrid force field, this two-level system can be described by the following interaction matrix,

$$\begin{array}{c|cc} & |v_6 = \nu_9 = 1\rangle & |v_5 = 1\rangle \\ \hline |v_6 = \nu_9 = 1\rangle & 1160.0 & -8.6 \\ |v_5 = 1\rangle & -8.6 & 1164.5 \end{array}$$

where diagonal elements are deperturbed and the off-diagonal coupling element is the cubic force constant  $\phi_{569}/\sqrt{8}$ . The eigenvalues of this matrix are 1153 and 1171  $\text{cm}^{-1}$  in optimal agreement with the measured transition frequencies. The  $\nu_3$  vibration appears at 1360.8  $\text{cm}^{-1}$  with a B-type band envelope and an intensity (according to the ChS:rDSD computations) of 113.3  $\text{km mol}^{-1}$ . The remaining relevant absorptions in this

spectral region include the first overtones of  $\nu_{11}$  and  $\nu_{10}$  at 1106.8 and 1498.1  $\text{cm}^{-1}$ , respectively, and the  $\nu_{10} + \nu_{11}$  combination measured at 1304.1  $\text{cm}^{-1}$ , all showing an A/B-type envelope with prevailing A-character. When one focuses on the portion between 1700 and 3500  $\text{cm}^{-1}$ , the strongest features come from the  $\nu_2$  fundamental, corresponding to the stretching of the C=C double bond and measured at 1787.4  $\text{cm}^{-1}$  and  $\nu_1$  centered at 3163.7  $\text{cm}^{-1}$  associated with the C-H stretching. The latter is involved in a quite strong Fermi resonance of type II with the nearby  $\nu_2 + \nu_3$  combination appearing at 3119.5  $\text{cm}^{-1}$  as an A/B-type band of intensity comparable to that of  $\nu_1$ . Using deperturbed frequencies obtained at the ChS:rDSD level, this resonance can be modeled by the following interaction Hamiltonian

$$\begin{array}{c|cc} & |v_2 = v_3 = 1\rangle & |v_1 = 1\rangle \\ \hline |v_2 = v_3 = 1\rangle & 3143.0 & 28.7 \\ |v_1 = 1\rangle & 28.7 & 3148.1 \end{array}$$

with eigenvalues 3116 and 3173  $\text{cm}^{-1}$ , and the perturbed levels are almost a 1:1 mixture of the unperturbed ones. In addition, the region presents a large number of weaker features mainly due to two quanta combinations bands, as detailed in Table 11. The last part of the MIR spectrum, in the range of 3500–5000  $\text{cm}^{-1}$ , shows a number of weak absorption features attributable to binary and ternary combination vibrations with an intensity within  $0.5 \text{ km mol}^{-1}$ . Among these, the most important ones come from the first overtone of  $\nu_2$  (3560  $\text{cm}^{-1}$ ) and the binary combinations involving the  $\nu_1$  vibration, namely,  $\nu_1 + \nu_7$  (3784.9  $\text{cm}^{-1}$ ),  $\nu_1 + \nu_6$  (4087.9  $\text{cm}^{-1}$ ),  $\nu_1 + \nu_5$  (4330.5  $\text{cm}^{-1}$ ),  $\nu_1 + \nu_4$  (4419.6  $\text{cm}^{-1}$ ),  $\nu_1 + \nu_3$  (4523.7  $\text{cm}^{-1}$ ), and  $\nu_1 + \nu_2$  (4951.0  $\text{cm}^{-1}$ ). These combination levels of the type  $\nu_1 = \nu_j = 1$  are expected to inherit the Fermi resonance  $\nu_1/\nu_2 + \nu_3$  and, hence, to be involved to various extents in 1–2 interactions with the corresponding three quanta combination levels  $\nu_2 = \nu_3 = \nu_j = 1$  that in the present investigation have been treated by manually setting up the proper interaction matrices on the basis of the computed quantities, resulting in an optimal agreement with the experiment. In particular for  $\nu_2 = \nu_3 = \nu_5 = 1$ , the resonant systems involving  $\nu_1/\nu_2 + \nu_3$  and  $\nu_5/\nu_6 + \nu_9$  intersect, giving rise to a four-level system that using the spectroscopic parameters evaluated from the ChS:rDSD hybrid force field has been described by the following interaction Hamiltonian:

$$\begin{array}{c|cccc} & |v_2 = v_3 = v_6 = v_9 = 1\rangle & |v_2 = v_3 = v_5 = 1\rangle & |v_1 = v_6 = v_9 = 1\rangle & |v_1 = v_5 = 1\rangle \\ \hline |v_2 = v_3 = v_6 = v_9 = 1\rangle & 4290.4 & -8.6 & 28.7 & 0 \\ |v_2 = v_3 = v_5 = 1\rangle & -8.6 & 4299.4 & 0 & 28.7 \\ |v_1 = v_6 = v_9 = 1\rangle & 28.7 & 0 & 4305.4 & -8.6 \\ |v_1 = v_5 = 1\rangle & 0 & 28.7 & -8.6 & 4308.5 \end{array}$$

Upon diagonalization, the following eigenvalues and eigenvectors are obtained:

$$\begin{array}{c|cccc} & |v_2 = v_3 = v_6 = v_9 = 1\rangle & |v_2 = v_3 = v_5 = 1\rangle & |v_1 = v_6 = v_9 = 1\rangle & |v_1 = v_5 = 1\rangle \\ \hline & 4262.4 & 4280.8 & 4321.2 & 4339.3 \\ |v_2 = v_3 = v_6 = v_9 = 1\rangle & -0.6 & -0.4 & 0.5 & 0.4 \\ |v_2 = v_3 = v_5 = 1\rangle & 0.4 & -0.6 & -0.3 & 0.5 \\ |v_1 = v_6 = v_9 = 1\rangle & -0.5 & -0.4 & -0.6 & -0.4 \\ |v_1 = v_5 = 1\rangle & -0.4 & 0.5 & -0.5 & 0.6 \end{array}$$

From these, it can be observed that the perturbed levels are better described as a superposition of the unperturbed ones. That said, when the labels corresponding to the most important component are retained, it can be concluded that the four quanta combination is predicted at 4262  $\text{cm}^{-1}$ , the  $\nu_2 + \nu_3 + \nu_5$  and  $\nu_1 + \nu_6 + \nu_9$  bands move to 4281 and 4321  $\text{cm}^{-1}$ , respectively, and the  $\nu_1 + \nu_5$  combination is expected around 4339  $\text{cm}^{-1}$ . Among these, only the  $\nu_2 + \nu_3 + \nu_5$  and  $\nu_1 + \nu_5$  transitions are expected to produce weak but observable absorptions, and indeed, the perturbed transition frequencies

Table 11. Vibrational Assignment of HFO-1123 and Comparison to Theoretical Wavenumbers ( $\text{cm}^{-1}$ )

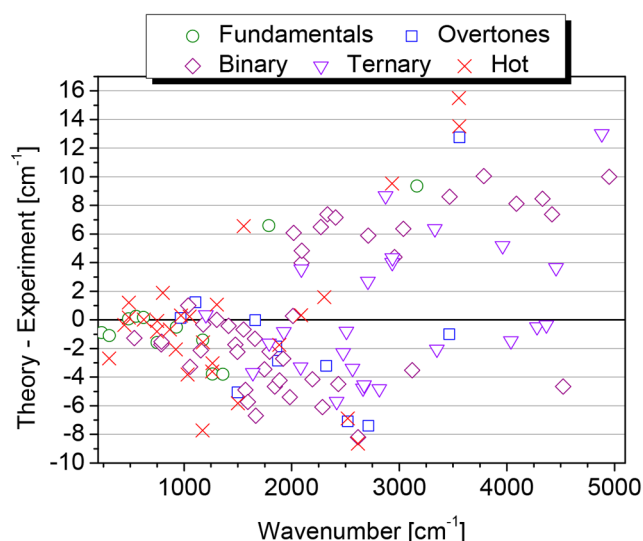
assignment	exp.	ChS:rDSD	assignment	exp.	ChS:rDSD	assignment	exp.	ChS:rDSD
$\nu_9$	232.9	232	$\nu_2 - \nu_{12}$	1492.74	1491	$\nu_3 + \nu_6 + \nu_9$	2509.80	2509
$2\nu_{12} - \nu_{12}$	302.69	300	$\nu_4 + \nu_9$	1495.24	1493	$2\nu_4 + \nu_9 - \nu_9$	2520.89	2514
$\nu_{12}$	305.1	304	$2\nu_{10}$	1498.07	1493	$2\nu_4$	2522.09	2515
$\nu_{10} - \nu_{12}$	445.36	445	$\nu_5 + \nu_{11} - \nu_9$	1498.83	1493	$\nu_4 + \nu_{10} + \nu_{11}$	2565.40	2562
$\nu_8$	484.93	485	$\nu_6 + \nu_7$	1552.68	1552	$\nu_3 + \nu_4 + \nu_9 - \nu_9$	2615.64	2607
$\nu_9 + \nu_9 - \nu_9$	485.84	486	$\nu_2 - \nu_9$	1555.45	1562	$\nu_3 + \nu_4$	2617.20	2609
$\nu_8 + 2\nu_9 - 2\nu_9$	486.80	488	$\nu_4 + \nu_{12}$	1568.89	1564	$\nu_3 + \nu_{10} + \nu_{11}$	2662.77	2658
$\nu_9 + \nu_{12}$	537.28	536	$\nu_3 + \nu_9$	1592.73	1587	$\nu_4 + \nu_6 + \nu_8$	2672.54	2668
$\nu_{11}$	553.76	554	$\nu_5 + 2\nu_9$	1639.7	1636	$\nu_5 + \nu_6 + \nu_7$	2709.30	2712
$\nu_7$	623.83	624	$\nu_5 + \nu_8$	1657.32	1656	$\nu_2 + \nu_6$	2711.11	2717
$\nu_{11} + \nu_{12} - \nu_9$	625.97	626	$3\nu_{11}$	1661.02	1661	$2\nu_3$	2713.40	2706
$\nu_9 + \nu_{10} - \nu_9$	747.82	747	$\nu_3 + \nu_{12}$	1665.7	1659	$\nu_4 + \nu_6 + \nu_7$	2813.82	2809
$\nu_{10} + \nu_9 - \nu_9$	749.08	749	$\nu_4 + \nu_8$	1746.46	1743	$\nu_2 + 2\nu_{11}$	2874.33	2883
$\nu_{10}$	750.59	749	$\nu_2$	1787.42	1794	$\nu_1 - \nu_9$	2931.47	2941
$\nu_9 + \nu_{11}$	787.72	786	$\nu_8 + \nu_{10} + \nu_{11}$	1789.62	1788	$2\nu_5 + \nu_7$	2934.00	2938
$\nu_8 + \nu_{12}$	789.52	788	$\nu_5 + \nu_7$	1795.22	1793	$\nu_2 + \nu_6 + \nu_9$	2936.66	2941
$2\nu_{11} - \nu_{12}$	802.12	804	$\nu_3 + \nu_8$	1843.65	1839	$\nu_2 + \nu_5$	2954.62	2959
$\nu_5 - \nu_{12}$	867.67	867	$2\nu_6$	1856.86	1855	$\nu_2 + \nu_4$	3038.64	3045
$\nu_6 + \nu_9 - \nu_9$	923.06	921	$3\nu_7$	1871.86	1869	$\nu_2 + \nu_3$	3119.52	3116
$\nu_6$	929.51	929	$\nu_4 + \nu_6 - \nu_{12}$	1885.75	1884	$\nu_1$	3163.65	3173
$2\nu_8$	969.86	970	$\nu_4 + \nu_7$	1887.23	1883	$\nu_2 + \nu_6 + \nu_7$	3329.64	3336
$2\nu_8 + \nu_9 - \nu_9$	971.70	972	$\nu_5 + \nu_{10}$	1920.71	1918	$\nu_4 + \nu_5 + \nu_6$	3349.05	3347
$\nu_4 - \nu_9$	1032.82	1029	$\nu_7 + \nu_{10} + \nu_{11}$	1926.83	1926	$3\nu_5$	3465.01	3464
$\nu_8 + \nu_{11}$	1038.00	1039	$\nu_3 + \nu_7$	1982.40	1977	$\nu_1 + \nu_{12}$	3468.40	3477
$\nu_{10} + \nu_{12} + \nu_9 - \nu_9$	1050.78	1051	$\nu_4 + \nu_{10}$	2010.73	2011	$2\nu_2 + \nu_{12} - \nu_{12}$	3553.50	3569
$\nu_{10} + \nu_{12}$	1054.28	1051	$\nu_2 + \nu_9$	2018.93	2025	$2\nu_2 + \nu_9 - \nu_9$	3557.47	3571
$2\nu_{11}$	1106.78	1108	$2\nu_6 + \nu_9$	2083.30	2080	$2\nu_2$	3560.25	3573
$\nu_6 + \nu_9$	1155.12	1153	$2\nu_5 - \nu_{12}$	2086.68	2087	$2\nu_3 + \nu_6$	3630.04	3622
$\nu_5 + \nu_{12} - \nu_{12}$	1170.73	1163	$\nu_5 + \nu_7 + \nu_{12}$	2089.45	2093	$\nu_1 + \nu_7$	3784.95	3795
$\nu_5$	1172.41	1171	$\nu_2 + \nu_{12}$	2091.02	2095	$\nu_2 + \nu_4 + \nu_6$	3960.83	3966
$\nu_6 + \nu_{11} - \nu_{12}$	1175.58	1174	$\nu_5 + \nu_6$	2093.18	2098	$\nu_2 + \nu_3 + \nu_6$	4038.46	4037
$\nu_7 + \nu_{11}$	1178.32	1178	$\nu_4 + \nu_6$	2192.15	2188	$\nu_1 + \nu_6$	4087.88	4096
$2\nu_8 + \nu_9$	1203.64	1204	$\nu_2 + \nu_8$	2270.51	2277	$\nu_2 + \nu_3 + \nu_5$	4281.48	4281
$\nu_4 + \nu_8 - \nu_8$	1261.62	1258	$\nu_3 + \nu_6$	2285.10	2279	$\nu_1 + \nu_5$	4330.54	4339
$\nu_4 + \nu_{12} - \nu_{12}$	1264.03	1261	$\nu_3 + \nu_4 - \nu_{12}$	2303.42	2305	$\nu_2 + \nu_3 + \nu_4$	4366.35	4366
$\nu_4$	1264.78	1261	$2\nu_5$	2322.22	2319	$\nu_1 + \nu_4$	4419.63	4427
$\nu_8 + \nu_{10} + \nu_{12} - \nu_9$	1302.94	1304	$\nu_2 + \nu_{11}$	2331.63	2339	$\nu_2 + 2\nu_3$	4457.36	4461
$\nu_{10} + \nu_{11}$	1304	1304	$\nu_2 + \nu_7$	2406.85	2414	$\nu_1 + \nu_3$	4523.66	4519
$\nu_3$	1360.80	1357	$\nu_4 + \nu_6 + \nu_9$	2417.70	2412	$\nu_2 + 2\nu_3$	4884	4897
$\nu_6 + \nu_8$	1413.42	1413	$\nu_4 + \nu_5$	2433.49	2429	$\nu_1 + \nu_2$	4951.00	4961
$\nu_5 + \nu_{12}$	1475.74	1474	$2\nu_6 + \nu_7$	2479.30	2477			

well reconcile with the bands observed at 4280.8 and 4330.5  $\text{cm}^{-1}$ . In passing, it should be noted that the band here assigned as  $2\nu_2$  was originally attributed to a three quanta combination in ref 43; however, according to the guidance of quantum chemical calculations, the overtone presents the more suited choice with the strongest transition expected at around 3560  $\text{cm}^{-1}$ .

**4.4. Absorption Cross Sections, Anharmonic Dipole Moment Surface, and Global Warming Potential.** The experimental investigation has also involved the determination of integrated absorption cross sections (i.e., IR band intensities) in the 400–5000  $\text{cm}^{-1}$  region. While the reader is referred to the available literature for a detailed description of the procedure,<sup>21,24,34</sup> here, only the main steps are summarized. The method is based on the least-squares fitting the point-by-point absorbance value,  $A(\tilde{\nu})$ , measured at each wavenumber,  $\tilde{\nu}$ , against the sample concentration. The slope of the straight line so obtained at each wavenumber,  $\sigma(\tilde{\nu})$ ,

provides the absorbance cross section per molecule ( $\text{cm}^2 \text{molecule}^{-1}$ ) with the statistical uncertainty obtained in a similar way. In passing, it should be noted that the procedure yields IR intensities in very good agreement with those determined by the line-by-line analysis of high-resolution spectra (see, for example, refs 51, 85, and 86). The integrated absorption cross sections obtained from the analysis of the medium resolution IR spectra of HFO-1123 are collected in Table 12, together with the corresponding theoretical predictions at the ChS:rDSD level of theory, while the cross section spectrum is detailed in Figure 4.

As to the greenhouse effect exerted by this molecule, the most important spectral regions are, as expected, those featuring the strong absorptions due to the C–F stretching vibrations. In particular, the spectral region between 1000 and 1445  $\text{cm}^{-1}$  accounts for almost 50% of the total IR absorption strength measured over the 400–5000  $\text{cm}^{-1}$  range. As previously pointed out, it hosts the absorptions stemming



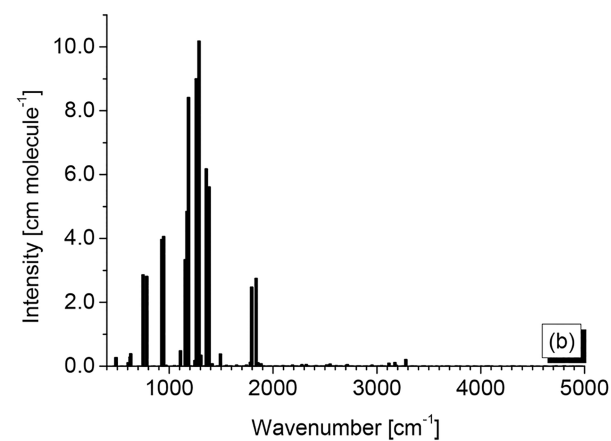
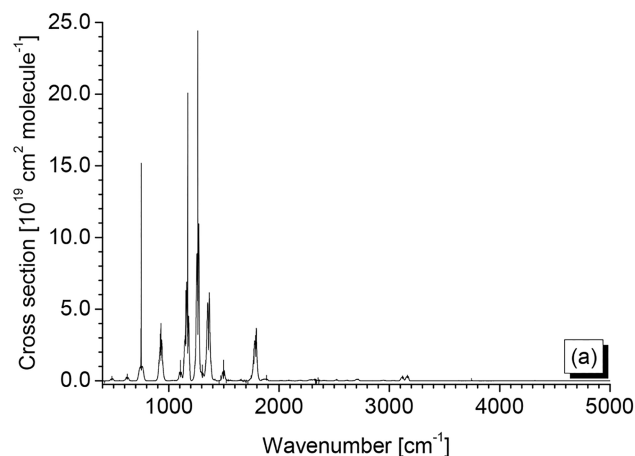
**Figure 3.** Differences between computed and observed transition frequencies as a function of the wavenumbers and type of transition for the assigned bands of HFO-1123.

**Table 12. Experimental and Theoretical Integrated Absorption Cross Sections ( $\text{km mol}^{-1}$ ) of HFO-1123**

spectral range ( $\text{cm}^{-1}$ )	main absorption	exp. <sup>a</sup>	ChS:rDSD <sup>b</sup>
440–530	$\nu_8$	2.1(1)	2.07
530–570	$\nu_9 + \nu_{12}, \nu_{11}$	0.068(3)	0.01
570–670	$\nu_7$	3.7(2)	3.81
670–820	$\nu_{10}, \nu_8 + \nu_{12}$	27(2)	29.66
820–1000	$\nu_6, 2\nu_8$	52(1)	54.96
1000–1205	$\nu_8 + \nu_{11}, \nu_{10} + \nu_{12}, 2\nu_{11}, \nu_6 + \nu_9, \nu_5, \nu_7 + \nu_{11}$	129(3)	142.37
1205–1445	$\nu_4, \nu_{10} + \nu_{11}, \nu_3, \nu_6 + \nu_8$	279(3)	313.55
1705–1950	$\nu_4 + \nu_8, \nu_2, \nu_5 + \nu_7, \nu_3 + \nu_8, 2\nu_6, \nu_4 + \nu_7, \nu_5 + \nu_{10}$	63(2)	68.97
1950–2240	$\nu_3 + \nu_7, \nu_2 + \nu_9, \nu_2 + \nu_{12}, \nu_4 + \nu_6$	1.92(8)	1.88
2460–2640	$2\nu_4, \nu_3 + \nu_4$	1.62(2)	2.03
2640–2800	$\nu_2 + \nu_6$	2.24(2)	2.79
2985–3250	$\nu_2 + \nu_4, \nu_2 + \nu_3, \nu_1$	8.5(3)	11.18
3415–3580	$2\nu_2$	0.34(4)	0.33
3980–4130	$\nu_1 + \nu_6$	0.34(3)	0.33
4150–4665	$\nu_1 + \nu_5, \nu_1 + \nu_4, \nu_1 + \nu_3$	1.08(3)	1.15
4665–5000	$\nu_1 + \nu_2$	0.23(2)	0.20
MAE <sup>c</sup>			3.97

<sup>a</sup>Values in parentheses are standard errors in the units of the last significant digits. <sup>b</sup>Harmonic frequencies and intensities from ChS, anharmonic contributions at the rev-DSD-PBEP86-D3/jun-cc-pVTZ level. <sup>c</sup>Mean absolute error.

from the  $\nu_3$  and  $\nu_4$  normal modes involving the asymmetric stretching of the  $\text{CF}_2$  moiety and predicted with an anharmonic intensity of 119.7 and 196.4  $\text{km mol}^{-1}$ , respectively. Next, the C–F stretching motion of the CHF group ( $\nu_5$ ) is located within the 1000–1205  $\text{cm}^{-1}$  spectral range within the atmospheric window, which accounts for about 23% of the overall integrated band intensity. In particular, according to ChS:rDSD anharmonic computations, it has an intensity of 75.5  $\text{km mol}^{-1}$ , almost half with respect to the harmonic value (140.4  $\text{km mol}^{-1}$ ). The reason for this weakening accounting for both mechanical and electrical anharmonicity has to be sought in the aforementioned Fermi



**Figure 4.** (a) Experimental cross-section spectrum of HFO-1123 in the 400–5000  $\text{cm}^{-1}$  spectral region. (b) Theoretical stick spectrum obtained at the hybrid ChS:rDSD level of theory over the same spectral range.

type II interaction with  $\nu_6 + \nu_9$  that borrows intensity from the fundamental, and indeed, its strength is predicted to be 57.9  $\text{km mol}^{-1}$ . The ranges 820–1000 and 1705–1950  $\text{cm}^{-1}$  each contain about the 10% of the total measured intensity. The former, still within the atmospheric window, presents  $\nu_6$ , the symmetric  $\text{CF}_2$  stretching, as the only relevant contribution, while the latter is dominated by the  $\nu_2$  C=C stretching vibration whose IR intensity is predicted to be 59.3  $\text{km mol}^{-1}$ . Then, the spectral intervals 670–820  $\text{cm}^{-1}$  (27  $\text{km mol}^{-1}$ ) and 2985–3250  $\text{cm}^{-1}$  (8.5  $\text{km mol}^{-1}$ ) contribute to about 5% and 1% of the overall integrated band intensity, respectively. Their measured intensities are mainly due to the  $\nu_{10}$  vibration, whose computed intensity is 29  $\text{km mol}^{-1}$  and the  $\nu_1$  and  $\nu_2 + \nu_3$  bands. In the latter case, as well, the combination vibration steals intensity from the C–H stretching vibration, thus reaching an absorption cross section of about 4  $\text{km mol}^{-1}$ , while the  $\nu_1$  fundamental loses about 50% of its strength (from 11.4 to 6.3  $\text{km mol}^{-1}$ ) upon introduction of anharmonic effects. The remaining spectral portions account for less than 1% of the total integrated absorption cross section measured over the whole 400–5000  $\text{cm}^{-1}$  range. Overall, a good accord between the measured intensities and the corresponding theoretical counterparts is noted for the different integration ranges, the mean absolute error of around 4  $\text{km mol}^{-1}$  being coherent with the expected accuracy.<sup>63–65</sup>

Finally, when one utilizes the measured cross section spectrum and the narrowband model of ref 87, the HFO-1123 radiative forcing (RF) is estimated to be  $0.10 \text{ W m}^{-2} \text{ ppbv}^{-1}$ . Since this approach might not be well suited for short-lived molecules like HFO-1123, the value determined should be taken as an estimate of the actual RF, although the data computed in this way are in good agreement with those obtained using a more sophisticated model on a similar halogenated olefin.<sup>88</sup> To the best of our knowledge, there is no data about the atmospheric lifetime of HFO-1123, which however is required in order to evaluate the global warming potential index. On the basis of the available literature,<sup>89</sup> an atmospheric lifetime of 10–30 days appears as a reasonable upper limit guess, leading to an estimated GWP between 2.0 and 5.3 (taking  $\text{CO}_2$  as the reference) on a time horizon of 100 years.

## 5. CONCLUSIONS

Several research efforts have been devoted to the study of the infrared spectroscopic behavior of both HFCs and HCFCs with the aim to assess the environmental impacts of these compounds. Indeed, it is important for radiative transfer calculations and atmospheric models to include data about their absorptions, and furthermore, accurate characterizations of their spectral properties constitute the basic requirement for monitoring their concentration in the atmosphere or work places. The present work aimed at investigating the structural and rotational/vibrational spectroscopic behavior of HFO-1123, one of the next generation refrigerants proposed as a replacement for HFCs, which has a supposedly shorter atmospheric lifetime and hence a lower impact on global warming. In particular, an accurate equilibrium geometry has been first worked out by applying post-Hartree–Fock composite methods. On the one side, this has been the first step toward the detailed modeling of its spectroscopic features, particularly those of relevance for rotational spectroscopy. On the other side, it represents the most accurate equilibrium geometry presented up to date for this species. The accuracy reached in the geometry has been mirrored in the accuracy in the computed ground state- and excited fundamental vibrational level- rotational constants of HFO-1123, which showed a remarkable agreement with the available experimental data with deviations well within 0.1%. Quartic and sextic centrifugal distortion parameters have been evaluated as well, addressing the dichotomy existing on the experimental data determined for the latter terms. Together with the  $\alpha_k^l$  vibration–rotation constants and the  $\zeta_{kl}^a$  Coriolis coupling terms, the computed data can be used to drive further high-resolution vibrational–rotational investigations. Attention has then been moved to the integrated experimental–theoretical investigation of HFO-1123 vibrational spectra. From the theoretical point of view, accurate harmonic frequencies and IR intensities have been obtained using the ChS method, and subsequently, these have been combined with anharmonic contributions evaluated using the rev-DSDPBEP86 functional that has been shown to provide excellent performances for the computation of vibrational transition frequencies and intensities. The anharmonic spectra simulated in this way have been used to assist the characterization of the experimental gas-phase spectra. To this end, the observed absorption features have been identified as fundamentals, overtones, a combination, and hot bands up to three quanta of vibrational excitations, and the analysis resulted in the assignment of almost 120 vibrational transitions

in the 200–5000  $\text{cm}^{-1}$  spectral range. A number of 1–2 Fermi resonances have been recognized and consistently treated, thus explaining the observed spectral pattern and intensity alterations. Furthermore, the integrated absorption cross sections have been experimentally determined for the spectral region between 400 and 5000  $\text{cm}^{-1}$ , reporting also in this case a very good agreement with the theoretical predictions that have furnished a quantitative description of the overall band shape over the whole spectral region investigated in the present work. Finally, the obtained spectroscopic data have been employed to obtain an estimate of the HFO-1123 radiative forcing. From this, an upper limit of the global warming potential has been determined, which is of interest in view of the use of HFO-1123 for refrigeration applications.

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