

Scattering resonances in the rotational excitation of HDO by Ne and *normal*-H₂: theory and experiment[†]

Ricardo Manuel García-Vázquez,  ^a Astrid Bergeat,  ^{*a} Otoniel Denis-Alpizar,  ^b Alexandre Faure,  ^c Thierry Stoecklin  ^a and Sébastien B. Morales 

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The rotational excitation of a singly deuterated water molecule (HDO) by a heavy atom (Ne) and a light diatomic molecule (H₂) is investigated theoretically and experimentally in the near-threshold regime. Crossed-molecular-beam measurements with a variable crossing angle are compared to close-coupling calculations based on high-accuracy potential energy surfaces. The two lowest rotational transitions, $0_{00} \rightarrow 1_{01}$ and $0_{00} \rightarrow 1_{11}$, are probed in detail and a good agreement between theory and experiment is observed for both transitions in the case of HDO + Ne, where scattering resonances are however blurred out experimentally. In the case of HDO + H₂, the predicted theoretical overlapping resonances are faithfully reproduced by experiment for the $0_{00} \rightarrow 1_{11}$ transition, while the calculated strong signal for the $0_{00} \rightarrow 1_{01}$ transition is not detected. Future work is needed to reconcile this discrepancy.

1 Introduction

Water ranks as the third most abundant neutral molecule in the interstellar medium, behind H₂ and CO. Consequently, H₂O and its isotopologues, D₂O and HDO, have been observed in different environments and play a key role in the oxygen and deuterium chemistry of star- and planet-forming regions.^{1–3} Due to the water sublimation temperature, water is mainly frozen as ice in cold clouds and becomes abundant in the gas phase at higher temperatures, which defines the snowline.⁴ Similarly, water can be produced in the gaseous phase or on the surface of interstellar dust grains in the coldest regions.¹ Deuterated water (HDO), on the other hand, is formed mainly on dust grains in cold clouds prior to core

^aUniv. Bordeaux, CNRS, ISM, 33400 Talence, France. E-mail: astrid.bergeat@u-bordeaux.fr

^bGrupo de Investigación en Física Aplicada, Instituto de Ciencias Aplicadas, Facultad de Ingeniería, Universidad Autónoma de Chile, 7500912 Santiago, Chile

^cUniv. Grenoble Alpes, CNRS, IPAG, 38000 Grenoble, France

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collapse.^{1,5–7} As a result, the HDO/H₂O ratio varies significantly depending on the environment, rendering it a potent tracer of water chemistry and history.^{1,2,5,8} Recently, HDO was detected in a young proto-planetary disk, V883, in the molecular clouds of Orion, using the Atacama Large Millimeter/Submillimeter Array (ALMA).⁶ The HDO emission lines were observed at 225.89672 and 241.561550 GHz, which correspond to the $3_{12} \rightarrow 2_{21}$ and $2_{11} \rightarrow 2_{12}$ rotational transitions, respectively.

Accurate comprehension of how energy is exchanged between the kinetic energy of the gas and H₂O or HDO molecules is a key parameter for modelling astrophysical observations. It is essential to determine the effective cross-sections of inelastic collisions with the main interstellar neutral species, such as H₂ and He. They can be calculated theoretically over a wide range of collision energies. However, the most demanding test is the comparison of these calculations with experiments at the threshold of water's internal energy transitions. We conducted studies on inelastic collisions of HDO with H₂ within this framework, while those on the other isotopologues of water were already published.^{9–11} As the inelastic cross sections of HDO with helium will experimentally be too small,¹² we opted to simulate the study of isotopologues of water with atom, using neon.

Water is an asymmetric top molecule, and its rotational levels are conventionally labelled $j_{k_a k_c}$ where j is the rotational angular momentum quantum number (see Fig. 1). The pseudo-quantum numbers k_a and k_c are formally the projections of the rotational angular momentum vector along the principal inertia axes a and c , respectively. Considering the nuclear spins of H ($I = \frac{1}{2}$) and D ($I = 1$), the water molecules H₂O and D₂O have two nuclear spin modifications, labelled *ortho* and *para*. The sum $k_a + k_c$ for H₂O(D₂O) is even(odd) for *para* and odd(even) for *ortho* levels. It should be noted that the *para* and *ortho* levels do not interconvert in inelastic collisions. For HDO, this *ortho/para* modification does not apply and all HDO levels are collisionally connected. The HDO rotational

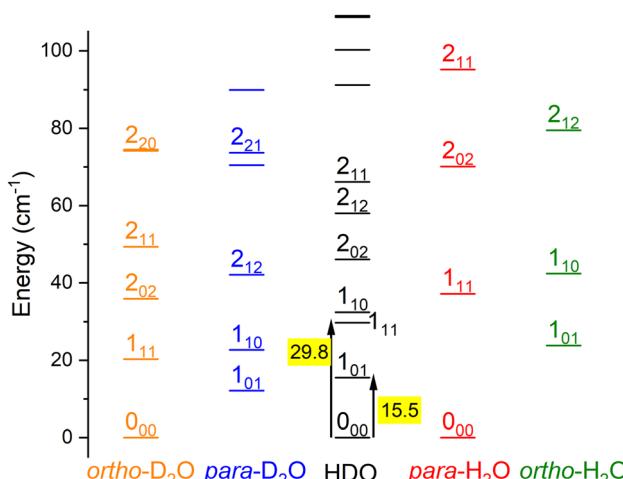


Fig. 1 Rotational energy levels of the water isotopologues, D₂O, HDO and H₂O, for $E < 100$ cm⁻¹. Levels are labelled by $j_{k_a k_c}$. Transition energies between the ground state 0₀₀ and levels 1₀₁ and 1₁₁ are given in cm⁻¹.



constants for the ground vibrational state were taken as $A = 23.41395 \text{ cm}^{-1}$, $B = 9.10340 \text{ cm}^{-1}$, and $C = 6.40628 \text{ cm}^{-1}$.¹³ The H_2 molecule also has two fermionic nuclei and the *para* and *ortho* rotational levels correspond to the even and odd values of its angular momentum j_{H_2} , respectively. *Normal*- H_2 is a mixture of *ortho*- and *para*- H_2 , considering there is a Boltzmann distribution of the rotational levels in the *ortho* or *para* forms at the surrounding temperature, but a spin temperature is maintained at room temperature, as the *ortho*-to-*para* conversion is extremely slow. The *ortho*-to-*para* ratio in *normal*- H_2 is thus 3 : 1.

Previous theoretical studies of the $\text{HDO}-\text{H}_2$ interactions are extensive.^{11,14} Faure *et al.*¹⁴ published the first rotational rate constants in the range 5–300 K. The calculations were recently improved¹¹ and radiative transfer calculations based on the new rate constants have shown that the HDO line at 225.9 GHz ($3_{12} \rightarrow 2_{21}$) observed by ALMA towards the V883 Ori proto-planetary disk⁶ is a weak maser.¹¹ It should be noticed that all the above studies rely on the full nine-dimensional potential energy surface (PES) computed by Valiron *et al.*³³ for the $\text{H}_2\text{O}-\text{H}_2$ system and adapted for all water isotopes.¹⁵ Theoretical calculations using this PES were compared to the only previous experimental study¹⁶ on the rotational inelastic collisions between HDO and *normal*- H_2 or *para*- H_2 . The differential cross sections (DCSs) were observed by Sarma *et al.*¹⁶ at collision energies of 580 cm^{-1} and 440 cm^{-1} . A good agreement was found between experiment and theory for the 0_{00} to 1_{11} , 2_{12} , 2_{21} and 2_{20} transitions. To the best of our knowledge, there are no previous theoretical calculations or experiments on the $\text{HDO}-\text{Ne}$ system.

The paper is organised as follows. Section 1 outlines the experimental setup and the procedure used. In Section 2, we describe the theoretical calculations for the $\text{HDO}-\text{Ne}$ system and summarise those for the $\text{HDO}-\text{H}_2$ system previously published.¹¹ Details on the PES for the $\text{HDO}-\text{Ne}$ system are given in Section 3. The theoretical integral cross-sections (ICSSs), as well as DCSs for different rotational transitions of HDO (0_{00} and 1_{01}) scattered by Ne , are presented in section 3. In this section, a comparison of experimental and calculated integral cross-sections (ICSSs) for the 0_{00} to 1_{01} or 1_{11} transitions at the thresholds for HDO scattered by Ne and *normal*- H_2 is also shown.

2 Experiments

2.1 Experimental set-up

Inelastic collisions down to low collision energies can be studied under single-collision conditions with the crossed-molecular-beam experiment in Bordeaux.¹⁷ The angle between the two molecular beams generated in two different chambers is variable. This allows the collision energy between the molecules to vary:¹⁸ the lowest value is for the lowest angle when the reactant beams have almost the same velocity. The molecules are then probed by a time-of-flight mass spectrometer (TOF-MS) following resonance-enhanced multiphoton ionization (REMPI).

The HOD beam was formed by expanding water vapour seeded in neon as a carrier gas through an Even–Lavie valve at 320 K from a high-pressure reservoir (11 bar), which contained a mixture with equal amounts of H_2O and D_2O maintained at a temperature of 45 °C. The carrier gas is neon to ensure better cooling of the water molecules.⁹ Supersonic expansion by a second Even–Lavie pulsed



valve at 300 K produced a pure Ne molecular beam. However, the two beams do not have exactly the same velocities due to the different valve temperatures and geometry of the two chambers: hence, the supersonic expansion efficiencies differ slightly.¹⁸ The H₂ molecular beam was also generated in the second chamber, with the valve cryo-cooled at a setpoint of 50 K, allowing decreasing the beam velocity and rotationally cooling down the H₂. As there is no *ortho*-to-*para* conversion, the *normal*-H₂ beam remains a mixture of *para*- and *ortho*-H₂ with the room temperature ratio, *i.e.*, 1 : 3. The *normal*-H₂ beam is thus composed of 25% H₂ ($j_{\text{H}_2} = 0$) and 75% H₂ ($j_{\text{H}_2} = 1$).⁹ The beam properties were inferred from measurements at the crossing point and 393.3 mm downstream, as explained previously,^{19,20} and are reported in Table 1.

The HOD rotational populations in the supersonic beam were measured using (2 + 1) REMPI at *ca.* 248 nm, *via* the C¹B₁ Rydberg state.²¹ Laser pulse energies of 2–5.5 mJ per pulse at a repetition rate of 10 Hz were generated by doubling the output of a dye laser operating with Coumarin 500 dye in ethanol, pumped by the third harmonic (355 nm) of a Nd:YAG laser. The laser beam was directed perpendicular to the collision plane and a wavemeter was used to monitor the wavelength. The output of the frequency-doubled dye laser was monitored with a photodiode to select data acquired with a laser pulse energy at $\pm 20\%$ of the mean.

2.2 Excitation functions: acquisition and data treatment

A typical REMPI spectrum of the HDO beam is presented in Fig. 2. A simulation of the experimental spectrum performed with pGopher²² software and the spectroscopic data of Yang *et al.*²¹ leads to the conclusion that HOD is mainly in the ground electronic, vibrational and rotational state (98.8%) and only 1.2% of the molecules are in the first excited rotational state, 1₀₁, which corresponds to a rotational temperature of 5 K. In some experiments, a slightly higher ratio of excited HDO was found (<5%).

The REMPI signal of HOD (1₁₁ or 1₀₁) is then acquired as a function of the angle, θ , between the two beams. To minimize background contributions from any rotationally excited molecules in the water beam or the main chamber, the H₂ or Ne beam was triggered at 5 Hz, with signals averaged in alternating pulse mode and recorded as two intensities: signal and background. The experimental ICSs in arbitrary units, σ , were then obtained from the subtraction of these averaged

Table 1 Characteristics of the molecular beams used in the determination of the integral cross-sections

Beam	Velocity ^a	Velocity spread ^a HWHM	Pulse duration ^b HWHM	Angular spread ^c HWHM
HOD in Ne at 320 K	$828 \pm 10 \text{ m s}^{-1}$	16 m s^{-1}	$26 \mu\text{s}$	0.9°
Ne at 300 K	$801 \pm 9 \text{ m s}^{-1}$	26 m s^{-1}	$15 \mu\text{s}$	1.2°
<i>n</i> -H ₂ at 50 K	$1174 \pm 13 \text{ m s}^{-1}$	34 m s^{-1}	$7 \mu\text{s}$	1.4°

^a Beam velocity peak values and half width at half maximum (HWHM) of the velocity spread deduced from temporal profiles at the crossing point and 393.3 mm downstream. ^b Pulse duration (HWHM) at the crossing point. ^c Angular divergence (HWHM).



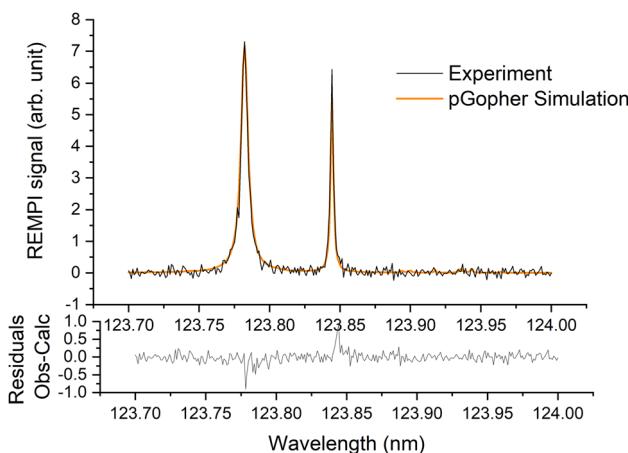


Fig. 2 Experimental (in black) and simulated (in orange) (2 + 1) REMPI spectra of the $\text{C}^1\text{B}_1, v' = 0 \leftarrow \text{X}^1\text{A}_1, v = 0$ transition of HDO in the supersonic beam. See Yang *et al.*²¹ for more details.

REMPI signal intensities over 5000 to 10 000 data. To take into account the density-to-flux transformation under our work conditions,²⁰ the values were divided by the relative velocity of the water and H_2 or Ne beams and the mean interaction time between the two beam pulses, considering a forward angular distribution of scattered water molecules as demonstrated later by the theory for the $\text{HOD} + \text{Ne}$ system. This conversion is described in detail in ref. 17 and 20. In

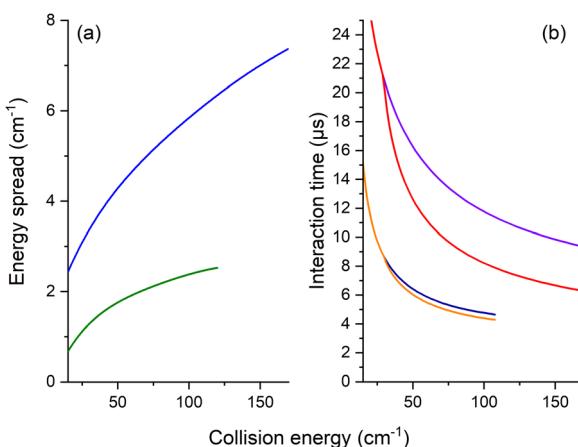


Fig. 3 Experimental collision-energy spread (a) and interaction time (b) as a function of the collision energy. (a) Half-width at 1/e of the collision-energy distribution assuming velocities and collision angular spreads as Gaussian functions for the two systems $\text{HOD} + \text{Ne}$ (in blue) and *normal-H₂* (in green). (b) Interaction time (right panel) considering an isotropic distribution of the scattered HDO (in violet, by Ne and in navy blue, by H_2) and a forward distribution (in red, by Ne and in orange, by H_2); the threshold is at 29.8 cm^{-1} for the $1_{11} \leftarrow 0_{00}$ rotational transition.

Fig. 3, the mean interaction time corrections are presented *versus* the collision energy. We note that the angular distribution (forward or isotropic) was found to have an almost negligible impact in the case of HOD + *normal*-H₂, contrary to the HDO + Ne system.

Moreover, for comparison with the experiment, the theoretical values were convoluted with the experimental collision-energy spread presented in Fig. 3. The poor collision-energy spread for the HDO + Ne system is due to the reduced mass, which is 9.746 a.m.u. in this case compared to 1.822 a.m.u. for the HDO + H₂ system. The convolution with the experimental collision-energy spread was computed from the interpolated calculated cross-sections. A short sensibility test was carried out, by changing each experimental parameter used to calculate the density-to-flux transformation and the experimental collision-energy spread. The energy resolution depends mainly on the beam velocity spreads and the pulse duration of the Ne or H₂ beam. The product of the relative velocity and the interaction time was essentially sensitive to the beam angular spreads, the beam velocities and the pulse duration of the Ne or H₂ beam. The influences of the other experimental characteristics remain within the statistic experimental error bars. Also reported on the experimental ICSs are the energy uncertainties due to the uncertainties in the beam velocities and the beam angle. These collision-energy uncertainties will just shift the excitation functions on the collision-energy scale.

3 Theoretical calculations

3.1 HOD + Ne potential energy surface

The coordinates employed for describing the potential energy surface (PES) of the HOD-Ne complex are shown in Fig. 4. R connects the centre of mass of the HOD molecule with Ne, θ is the angle between the z -axis and R , and φ is the azimuthal angle. The interatomic distances and bending angle of the HOD molecule are set to the vibrationally averaged values in the ground state ($r_{\text{OH}} = 0.9727 \text{ \AA}$, $r_{\text{OD}} = 0.9684 \text{ \AA}$ and $\gamma = 104.34^\circ$).¹⁵ The *ab initio* calculations were performed at the explicitly correlated coupled-cluster level of theory, including single, double and perturbative triple excitations (CCSD(T)-F12),²³ together with the cc-pVTZ-F12 and cc-pVQZ-F12 basis sets.²⁴ The energies at the completed basis set (CBS) limit were determined using the two-point extrapolation formula:²⁵

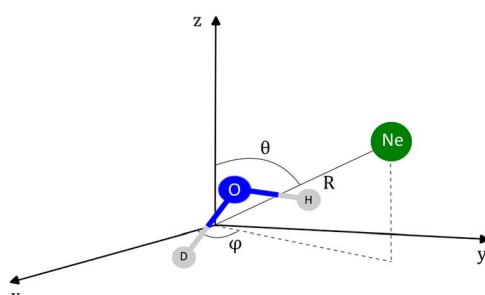


Fig. 4 Coordinates employed in the description of the HOD + Ne system. The origin of the coordinate system corresponds to the center of mass of the HDO molecule.



$$\left(E_{\text{CBS}} = \frac{E_X X^3 - E_Y Y^3}{X^3 - Y^3} \right),$$

where $X = 4$ and $Y = 3$ are the cardinality of the basis set. All the *ab initio* energies were computed with the MOLPRO package.²⁶

A grid of 6175 energies at the CBS limit was computed for the system. The grid includes 25 R values from 1.5 Å to 10.0 Å, with θ and φ varying from 0° to 180° in steps of 10° and 15°, respectively. It should be noted that for certain angular configurations, *ab initio* calculations could not produce energies for a few low R values due to the proximity of Ne and the atoms within the triatomic molecule. Consequently, local extrapolation was utilized.

The grid of energies was fitted to an analytical function, like the one used for H₂S + He²⁷ and H₂O/D₂O + Ne,²⁸ with the angular part expressed as the product of normalized associated Legendre functions $\bar{P}_l^m(\cos\theta)$ and cosine functions:

$$V(R, \theta, \varphi) = \sum_{l=0}^{10} \sum_{m=0}^{\min(l,4)} v_{lm}(R) \bar{P}_l^m(\cos\theta) \cos(m\varphi)$$

The $v_{lm}(R)$ coefficients were computed from a least-squares procedure and fitted using the reproducing kernel Hilbert space (RKHS) method,²⁹ using:

$$v_{lm}(R) = \sum_{k=1}^{N_R} \alpha_k^{lm} q^{2.5}(R, R_k)$$

where N_R is the number of R values of the grid, and $q^{2.5}$ is the one-dimensional kernel:²⁹

$$q^{2.5}(R, R_k) = \frac{2}{21R_{>}^6} - \frac{R_{<}}{14R_{>}^7}$$

with $R_{<}$ and $R_{>}$ defining the lower and larger values, respectively, between R and R_k . The α_k^{lm} coefficients were computed by solving the linear equation system $\mathbf{q}(R_k, R_k) \alpha^{lm} = \mathbf{v}_{lm}(R_k)$, where k and k' label different radial configurations of the grid. The van der Waals $\sim R^{-6}$ long-range behaviour is ensured using the $q^{2.5}(R, R_k)$ kernel.

3.2 HOD + Ne dynamical calculations

The Newmat code³⁰ was used to carry out the dynamical calculations of the HOD + Ne system. As in our previous work on H₂O/D₂O + Ne,²⁸ the rotational eigenfunctions and eigenenergies were determined by diagonalizing the rigid rotor Hamiltonian of an asymmetric top molecule:

$$\mathbf{H}_{\text{rot}} = A_x \mathbf{j}_x^2 + A_y \mathbf{j}_y^2 + A_z \mathbf{j}_z^2$$

In the usual symmetrised basis set:

$$|jkm, \varepsilon\rangle = [2(1 + \delta_{k0})]^{-\frac{1}{2}} (|jkm\rangle + (-1)^\varepsilon |jk - m\rangle)$$



where $|jkm\rangle$ is a symmetric top wave function and $A_x = A$, $A_y = C$ and $A_z = B$.¹⁵ These wave functions were used to perform close-coupling calculations, with the calculated eigenenergies being substituted for experimental ones.³¹

The basis set includes 7 values of HDO rotational angular momentum ($0 \leq j \leq 6$), which leads to energies up to 872.77 cm^{-1} . We carried out scattering calculations within the collision-energy range of $[15.5, 315.5] \text{ cm}^{-1}$, utilizing 0.1 cm^{-1} steps for energies below 115.5 cm^{-1} and 1 cm^{-1} steps above. The log derivative method³² was used to propagate the radial wave functions for $3.0 \leq R \leq 20a_0$ (a_0 is the Bohr radius). The relative convergence of the cross sections as a function of the total angular momentum J was imposed to be better than 10^{-3} , leading to a maximum value of $J = 66$ for the maximum energy value investigated. The reduced mass of the HDO-Ne system is $9.747\,406 \text{ a.m.u.}$

3.3 HDO + H₂ dynamical calculations

Scattering quantum close-coupling calculations of the rotational inelastic cross sections were previously published by Faure *et al.*¹¹ to provide rate constants over a large kinetic temperature range (5–300 K). For this study, the same V08 PES of Valiron *et al.*³³ was used, but more refined scattering calculations were performed. The calculations were conducted at the quantum close-coupling level with the MOLSCAT program.³⁴ The rigid-rotor PES for HDO-H₂ was derived from the full-dimensional (9D) PES of H₂O-H₂ determined at the CCSD(T) level of theory and further refined with higher-accuracy explicitly correlated CCSD(T)-R12 calculations.³³ To provide precise descriptions of the rotational thresholds and resonances, calculations were conducted up to total energies of 220 cm^{-1} with a grid increment of 0.25 cm^{-1} . Cross sections for *normal*-H₂ were obtained through calculations for the ground *para*-($j_{\text{H}_2} = 0$) and *ortho*-($j_{\text{H}_2} = 1$) states of H₂ and an *ortho*-to-*para* ratio of 3 was employed. The highest HDO rotational angular momentum in the basis set was $j_{\text{HDO}} = 5$, while two rotational levels of H₂ were included for the *para*-($j_{\text{H}_2} = 0, 2$) and *ortho*-($j_{\text{H}_2} = 1, 3$) symmetries of H₂. In the calculations of Faure *et al.*,¹¹ total energies of up to 2000 cm^{-1} (with a less-fine energy grid) were considered, but the level $j_{\text{H}_2} = 3$ of *ortho*-H₂ was neglected. The inclusion of this level was found to be important for a very detailed description of resonances in this work. The maximum value of the total angular momentum J used in the calculations was $J = 37$ at the highest collisional energy. The value of the H₂ rotational constant was $B_0 = 59.322 \text{ cm}^{-1}$. Finally, the reduced mass of the system HDO-H₂ was calculated to be 1.82252 a.m.u.

4 Results and discussion

4.1 HDO + Ne PES

The accuracy of the fitting procedures was assessed by measuring the root-mean-square deviation (RMSD). The RMSD for negative energies was determined to be $8.5 \cdot 10^{-4} \text{ cm}^{-1}$. Subsequently, the RMSD values obtained for energy intervals of $[0, 1000] \text{ cm}^{-1}$ and $[1000, 10\,000] \text{ cm}^{-1}$ were $7.7 \cdot 10^{-2} \text{ cm}^{-1}$ and 25.6 cm^{-1} correspondingly.

The global minimum of the system is located in a deuterium-bonded configuration that is linear (-65.15 cm^{-1} , at $R = 3.17 \text{ \AA}$, $\theta = 73^\circ$, $\varphi = 0^\circ$). From a comparison with the minima found for H₂O and D₂O systems in our previous



Table 2 Equilibrium energies, De , and geometries (variables as defined in the text and Fig. 4) of the Ne–HOD, Ne–H₂O and Ne–D₂O systems. The values for Ne–HOD were obtained in this work, while the values for the two other systems were taken from our previous work²⁸

System	De (cm ⁻¹)	Re (Å)	θ_e (°)	φ_e (°)
HOD + Ne	65.15	3.18	73.50°	0.00°
H ₂ O + Ne	65.29	3.20	73.00°	0.00°
D ₂ O + Ne	65.29 ^a	3.24	71.50°	0.00°

^a The D₂O + Ne PES was obtained from the H₂O + Ne PES taking into account the shift of the centre of mass (see ref. 28 for more details).

work²⁸ (see Table 2), we notice that these three stationary points are very close in geometrical configuration and well depth. Nevertheless, it should be noted that despite the similarities observed among the three minima in the PES, the most significant changes in the dynamics are determined by other features such as the shape of the barriers and the anisotropy.

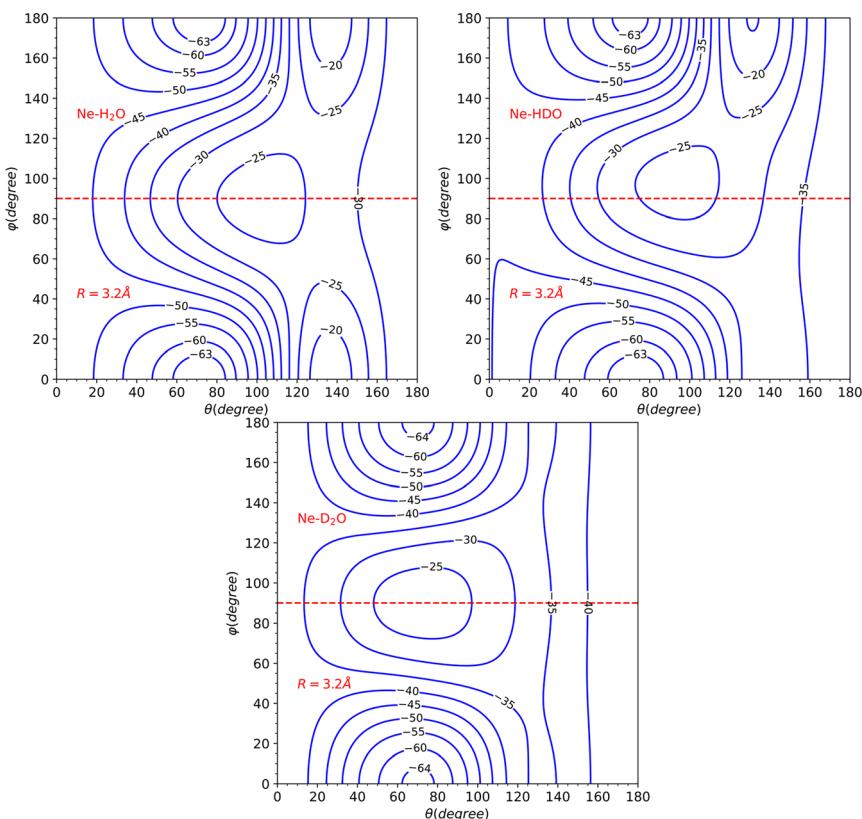


Fig. 5 Contour plots for the H₂O–Ne, HOD–Ne and D₂O–Ne PESs for $R = 3.2$ Å. A red dashed line is placed at $\varphi = 90^\circ$ to mark the symmetry of the H₂O/D₂O–Ne systems and the break in the symmetry of the HOD–Ne system. The labels of the contour lines correspond to the energy in cm⁻¹.



The topological variations amongst the systems can be more effectively visualized through Fig. 5. This displays a contour plot of the three systems, with $R = 3.2 \text{ \AA}$. To aid discussion of PES changes resulting from the H/D interchange, a red dashed line at $\varphi = 90^\circ$ has been included. From the plot of HDO, which corresponds to the top-right plot of Fig. 5, it can be observed that the contour lines behave similarly to those of H_2O , for the interval of $90^\circ \leq \varphi \leq 180^\circ$, which corresponds to the Ne atom approaching the H atom. The latter half of the interval for $0^\circ \leq \varphi \leq 90^\circ$, however, displays more variations compared to the D_2O system, implying that the system's dynamics will bear more similarities to that of H_2O .

This hypothesis receives further support by comparing the $v_{lm}(R)$ coefficients (refer to Fig. 6). For the H_2O and HDO system, the $l = 2, m = 0$ partial wave exhibits the only marked difference, while for D_2O , the $l = 1, m = 0$ partial wave differs significantly, being repulsive instead of attractive as seen for the water and singly-deuterated water systems. We should note that only even m partial waves are presented, since they are the only ones observed in the H_2O and D_2O systems.

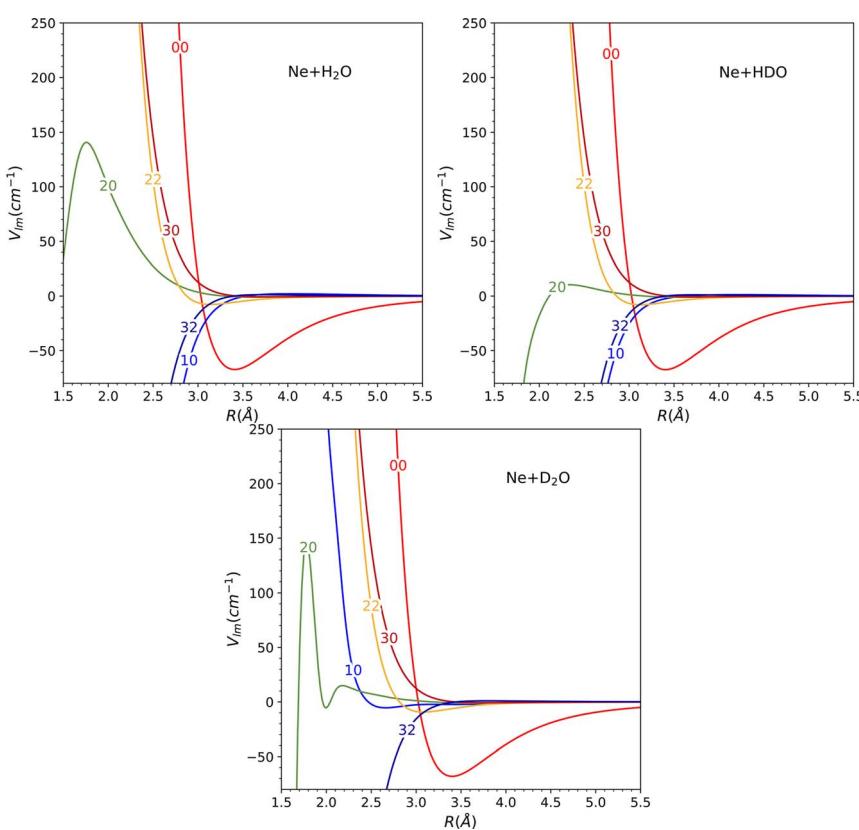


Fig. 6 $v_{lm}(R)$ expansion coefficients for the $\text{H}_2\text{O}-\text{Ne}$, $\text{HDO}-\text{Ne}$, and $\text{D}_2\text{O}-\text{Ne}$ PESs. Labels in the curves are associated with lm partial waves. Only even m values are allowed for H_2O and D_2O due to the C_{2v} symmetry of these two molecules, while for HDO, even though odd m values are now allowed, just the even m values are represented to simplify the comparison.



This is due to the fact that the molecule exhibits C_{2v} symmetry in these two cases. However, the inclusion of the odd m coefficients in the HDO PES expansion, which were not depicted in the plot to facilitate comparison, is likely to result in significant variations in the dynamics.

4.2 HDO + Ne inelastic collisions

The theoretical calculations focus on exciting the fundamental rotational level 0_{00} and the related transitions for exciting and relaxing from the first excited rotational state, 1_{01} . This is due to the experimental probing of only the first two excited rotational levels, 1_{01} and 1_{11} .

4.2.1 The rotational transition 0_{00} to 1_{11} . Calculated state-to-state cross sections for exciting the fundamental rotational level 0_{00} to the excited rotational states are illustrated in Fig. 7. The transition $1_{11} \leftarrow 0_{00}$ exhibits the largest cross-section, followed by the excitation $2_{02} \leftarrow 0_{00}$. It is noteworthy that all the water isotopes display a similar ordering for these largest cross-sections, providing evidence of similarities between the PESs. However, the symmetry between *para* and *ortho* in H_2O or D_2O systems has been broken, allowing previously forbidden transitions to occur. A few of these transitions, like $1_{01} \leftarrow 0_{00}$ and $2_{12} \leftarrow 0_{00}$, have significant cross sections. The energy threshold appears to have a small impact on the ordering of cross sections based on magnitude.

To compare the theoretical data with the experimental results shown in Fig. 8, we applied the experimental collision-energy spread to the theoretical cross-sections. Moreover, the experimental ICSs were scaled by one factor so that the area in the $[30\text{--}165]\text{ cm}^{-1}$ range is the same as for the theoretical ICS calculated. It should be noted that in the HDO beam, more than 95% of the population was in the ground rotational state. Therefore, when the 1_{11} level was probed, the main contribution originated from the $1_{11} \leftarrow 0_{00}$ transition. The experiment successfully reproduced the transition threshold at 29.75 cm^{-1} , but the calculated resonance features were blurred out in the experimental ICSs due to the energy

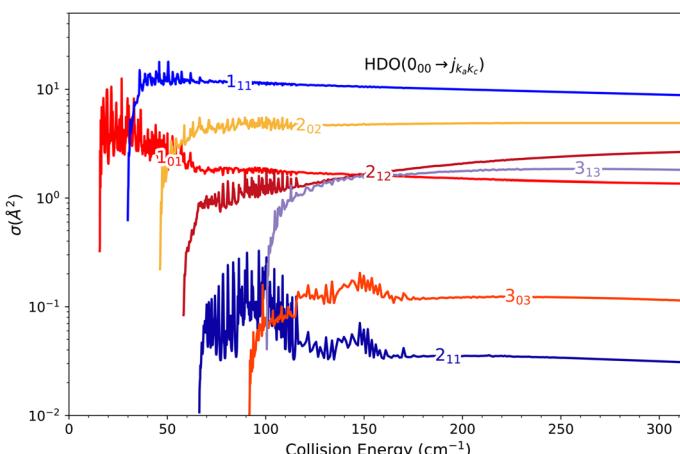


Fig. 7 Cross sections for the excitation of the fundamental rotational level of the HDO molecule by collision with Ne. The curves on the log-linear plot correspond to transitions from the 0_{00} level to the first eight rotational excited states.



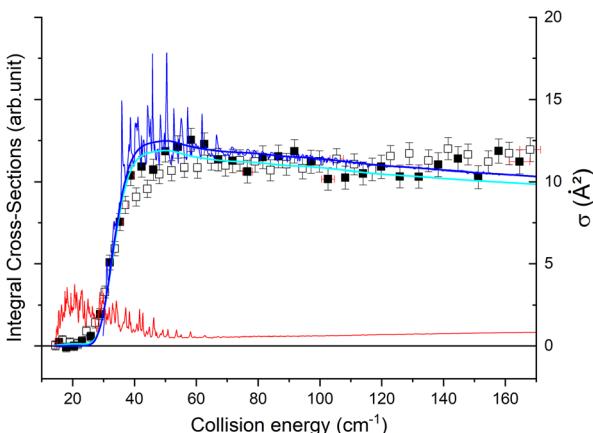


Fig. 8 Experimental and theoretical cross-sections for the $\text{HDO} + \text{Ne} \rightarrow \text{HDO (1}_{11}\text{)} + \text{Ne}$ inelastic collisions. Squares are experimental ICSSs for the 1_{11} rotational level, with error bars corresponding to statistical uncertainties at 95% of the confidence interval (white or black squares are the means of around 7000 or 9200 data, respectively). Shown as solid lines, the total theoretical contributions are convoluted to the experimental collision-energy spread, with partial contributions of the $1_{11} \leftarrow 0_{00}$ transition (as a thin blue line) and the $1_{11} \leftarrow 1_{01}$ transition (as a thin red line), assuming an initial rotational population from the 1_{01} state of 0% (as a bold blue line) or 5% (as a bold cyan line).

spread. To verify the reproducibility, two sets of experimental data were recorded at different periods of the year: the disparity in values was, however, mainly due to fluctuations in the laser power (as well as humidity in the laboratory). This might also explain why the experimental values exceed the theoretical cross-sections above 120 cm^{-1} (as the experimental acquisition was always carried out from the highest beam angle to the lowest angle).

4.2.2 The differential cross sections related to the rotational transition from 0_{00} to 1_{11} . The calculations of the differential cross sections (DCSs) for the inelastic collision between Ne and HDO have been performed with the usual formula:

$$\frac{d\sigma}{d\hat{R}}(j'\tau' \leftarrow j\tau | \hat{R}) = \frac{k_{j\tau}}{k_{j\tau}(2j+1)} \sum_{m_j=-j}^j \sum_{m_{j'}=-j'}^{j'} \left| f(j'\tau' \leftarrow j\tau | \hat{R}) \right|^2$$

where $\tau = k_a - k_{c'} \hat{R} = (\theta, \varphi)$ and $k_{j\tau} = \sqrt{2\mu(E - E_{j\tau})}$, while (j', τ') and (j, τ) define the final and initial level of the HDO molecule. $f(j'\tau' \leftarrow j\tau | \hat{R})$ is the scattering amplitude, which, in the case of the collision between a rigid asymmetric top and an atom, is defined as:

$$f(j'\tau' \leftarrow j\tau | \hat{R}) = \sqrt{\frac{\pi}{k_{j\tau} k_{j'\tau'}}} \sum_{J, M_J, M_{J'}} i^{l-l'+1} (-1)^{j+j'-l-l'} (2J+1) \sqrt{2l+1} \times \begin{pmatrix} j & l & J \\ m_j & 0 & -M \end{pmatrix} \\ \times \begin{pmatrix} j' & l' & J \\ m_{j'} & m_{l'} & -M \end{pmatrix} T_{j\tau l, j'\tau' l'} {}^J Y_{l' m_{l'}}(\hat{R})$$

where $T_{j\tau l, j'\tau' l'}$ are the transition matrix elements at a given collision energy and $Y_{l' m_{l'}}(\hat{R})$ is a spherical harmonic function.

In the present paragraph dedicated to the DCS associated with the $1_{11} \leftarrow 0_{00}$ transition, we extend the comparison between the HDO, H₂O and D₂O systems. The collision-energy ranges used to calculate the DCS were [37.5, 67.0] cm⁻¹, [30.0, 60.0] cm⁻¹ and [21.0, 60.0] cm⁻¹ for H₂O, HDO and D₂O, respectively. The choice of the energy interval allows a near-threshold study for each system.

Fig. 9 displays the computed DCSs. For the entire energy interval under consideration, the DCSs are observed to be strongly forward peaked for collisions with deuterated water (HDO) and H₂O, while significant back-scattering contributions are noted for collisions with D₂O. The uniqueness of the latter system stems from the $v_{10}(R)$ potential term, which is the only relevant $v_{lm}(R)$ term for this transition. As seen in Fig. 6, it is entirely attractive and primarily contributes to the forward-scattering direction for both H₂O and HDO. For D₂O, however, this interaction is predominantly repulsive, with a minor potential well of around 5 cm⁻¹ depth, the repulsion being a significant contributor to the backward-scattering phenomenon.

4.2.3 The rotational transition 0_{00} to 1_{01} . Theoretical state-to-state cross sections for the excitation and relaxation of the first excited rotational level 1_{01} are displayed in Fig. 10. The transitions $2_{12} \leftarrow 1_{01}$ and $1_{10} \leftarrow 1_{01}$ currently have the largest related cross sections, being approximately equal in magnitude. It is interesting to note that, despite the loss of *para/ortho* symmetry, the most significant cross sections for exciting the 0_{00} and 1_{01} levels correspond to those with similar symmetry in H₂O or D₂O molecules.²⁸ This proximity in symmetry may be linked to the similarities of the H₂O-Ne and HDO-Ne PESs, implying that partial waves with even values of m dominate the system's dynamics.

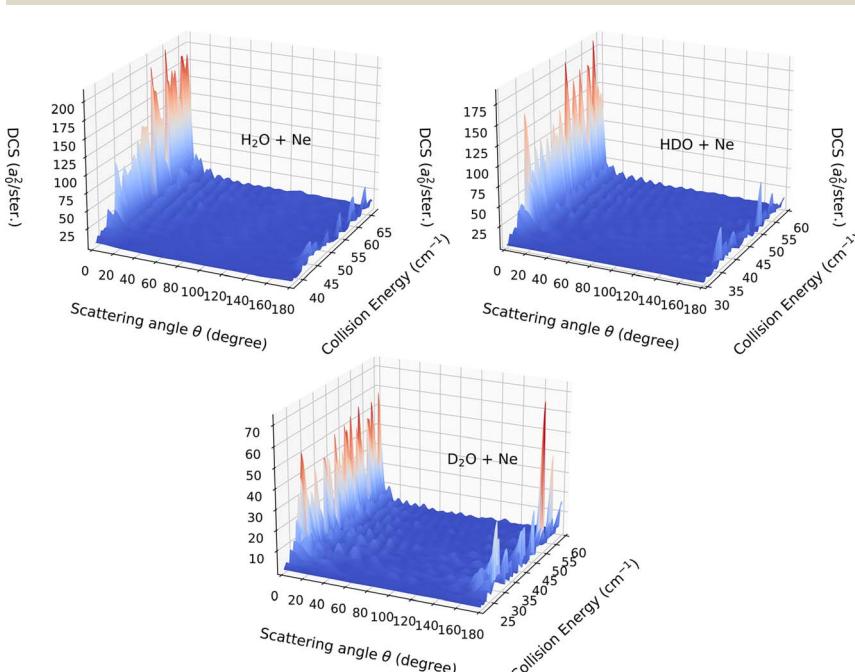


Fig. 9 Differential cross section for the $(1_{11} \leftarrow 0_{00})$ transitions induced by the collision of H₂O, HOD or D₂O with Ne.



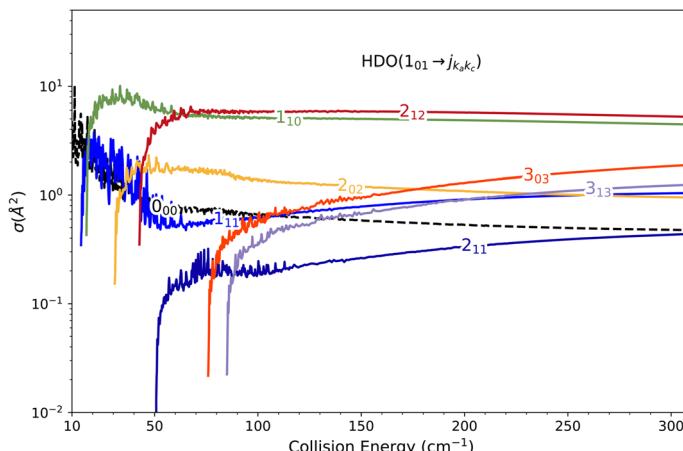


Fig. 10 Cross sections for the relaxation (black dashed line) and the excitation (solid lines) of the 1_{01} rotational level of the HDO molecule by collision with Ne. Excitation cross sections correspond to transitions to the seven closest in energy excited states.

Direct comparison with the experiment is not possible as the HDO beam may contain water in its first excited rotational state 1_{01} . To obtain the effective ICSs, it is thus necessary to combine the state-to-state calculated cross-sections. The HDO (1_{01}) formation is due to the $1_{01} \leftarrow 0_{00}$ transition, weighted by 1 or 0.95, depending on the ground HDO rotational level population. When there is 5% rotationally excited HDO in the water beam, HDO (1_{01}) also disappears due to the transitions from this 1_{01} level to the fundamental level or all the excited levels up to 2_{12} . The sum of these cross-sections is therefore subtracted with a weight of 0.05 from the HDO (1_{01}) production cross-section. The effective ICSs are then convolved with the experimental energy dispersion. The two computed excitation functions shown in Fig. 11 thus correspond, respectively, to the two extreme scenarios: the presence of 0% or 5% HDO in state 1_{01} in the water beam. Although the statistical uncertainties at 95% confidence remain high, due to the low HDO (1_{01}) intensity detected, the two undulations due to resonances predicted by theory can be seen. To verify the ratio of effective cross-sections between the 1_{01} to the 1_{11} states, a study at 44 cm^{-1} was carried out. Some of the experimental data used are presented in the ESI file.† REMPI intensities were acquired from both levels 1_{01} and 1_{11} while maintaining perfectly identical experimental conditions, with the exception of MCP amplification and the wavelength of the ionizing laser (its power having varied very little between the 2 wavelengths). The ratio of 1_{01} to 1_{11} population variations due to HDO + Ne collisions is roughly 0.2, with a relative uncertainty of 50%. This was deduced from pGopher simulations with the spectroscopic data from Yang *et al.*²¹ The ratio of the effective theoretical cross-sections convolved with the experimental energy dispersion at this collision energy is found to be in the range of 0.196 to 0.248 (depending on the population in state 1_{01} in the beam).

4.3 HDO + H₂ inelastic collisions

The state-to-state cross sections for the experimental energy range are displayed in Fig. 12. The HDO + *normal*-H₂ cross sections are similar in magnitude to those

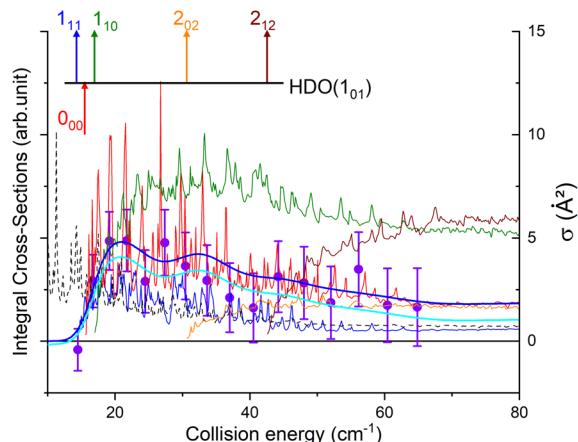


Fig. 11 Experimental and theoretical cross-sections for the $\text{HDO} + \text{Ne} \rightarrow \text{HDO} (1_{01}) + \text{Ne}$ inelastic collisions. Circles are experimental ICSs for the 1_{01} rotational level, with error bars corresponding to statistical uncertainties at 95% of the confidence interval (a circle represents the mean of around 9200 data). Shown as solid lines, the effective theoretical contributions are convoluted to the experimental collision-energy spread assuming an initial rotational population from the 1_{01} state of 0% (as a bold blue line) or 5% (as a bold cyan line), with partial contributions of the $1_{01} \leftarrow 0_{00}$ transition (as a red line), the $1_{01} \rightarrow 0_{00}$ transition (as a black dashed line), the $1_{11} \leftarrow 1_{01}$ transition (as a blue line), the $1_{10} \leftarrow 1_{01}$ transition (as a green line), the $2_{02} \leftarrow 1_{01}$ transition (as an orange line) and the $2_{12} \leftarrow 1_{01}$ transition (as a wine-colored line). The respective thresholds of the transitions are indicated by arrows.

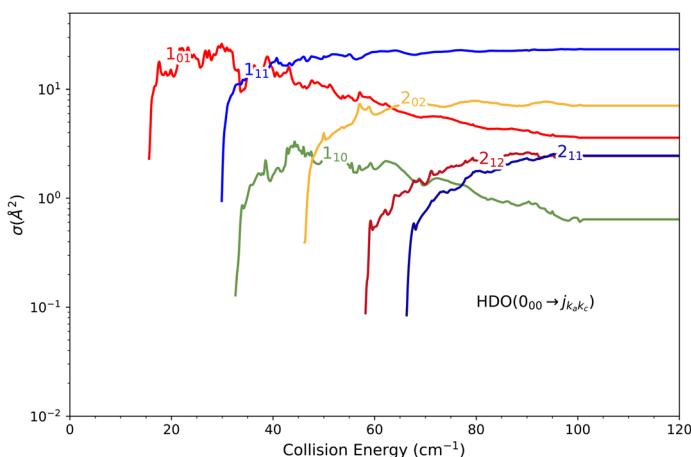


Fig. 12 Cross sections for the excitation of the fundamental 0_{00} rotational level of the HDO molecule by collision with *normal*-H₂. The curves on the log-linear plot correspond to transitions to the first six rotational excited states.

of the HDO + Ne system, except for the $1_{01} \leftarrow 0_{00}$ transition, which is approximately 5 times stronger at the peak when averaged over resonances. Moreover, for the HDO + *normal*-H₂ system, sharp resonances are not observed, in contrast to the HDO + Ne system. These differences can be rationalized by the fact that the

well for HDO–Ne is 65 cm^{-1} deep, whereas for HDO–H₂ it is 237 cm^{-1} . In the *normal*-H₂ system, it is also relevant to consider the angular momentum (and quadrupole) of H₂ ($j_{\text{H}_2} = 1$). Consequently, more scattering resonances may be found when H₂ is the collider, and their overlap broadens the excitation function in the HDO + *normal*-H₂ system. In the case of HDS + H₂,³⁵ the $1_{01} \leftarrow 0_{00}$ transition has an effective maximum cross-section whose magnitude is close to that of the $1_{11} \leftarrow 0_{00}$ transition, as observed here for HDO + H₂.

Experimentally, a weak signal from the 1_{01} rotational level is clearly visible when probing the water beam, indicating the presence of HDO in its first excited rotational state, mainly on the border of the water gas pulse (see the ESI file).† Nonetheless, no increase is detectable when colliding with *normal*-H₂. This leads us to the conclusion that the cross-section for the $1_{01} \leftarrow 0_{00}$ transition should be lower than about 7 \AA^2 (our limit for the HDO + Ne system), which is a factor of 2–3 lower than the theoretical value. In Fig. 13, a simulation of the excitation function has been carried out, taking into account the experimental dispersion of the energy and the population and depopulation of the 1_{01} rotational state (theoretical state-to-state cross-sections are reported in Fig. 14), as carried out for the HDO + Ne system. The simulation also considered two scenarios: initial 1_{01} state populations of 0% and 5% in the beam. The maximum in the cross-section for the $1_{01} \leftarrow 0_{00}$ transition is still of the same order of magnitude as the $1_{11} \leftarrow 0_{00}$ cross-sections, in disagreement with our lack of experimental signal. In contrast, the agreement between theory and experiment is excellent when probing the 1_{11} state (Fig. 13). The transition threshold is replicated accurately and the ICS oscillations

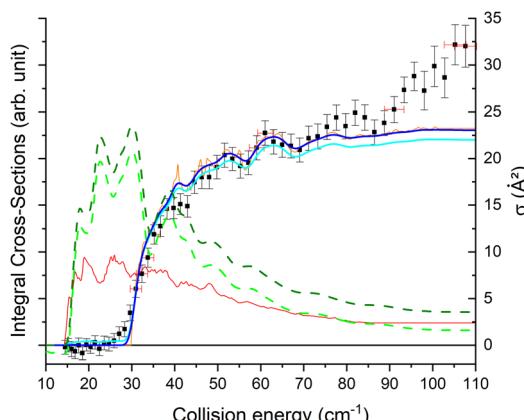


Fig. 13 Experimental and theoretical cross-sections for the HDO + *normal*-H₂ → HDO (1_{01} or 1_{11}) + H₂ inelastic collisions. Squares are experimental ICSs for the 1_{11} rotational level, with error bars corresponding to statistical uncertainties at 95% of the confidence interval (a square represents the mean of around 5000 data). Shown as solid lines, the total theoretical contributions are convoluted to the experimental collision-energy spread assuming an initial rotational population from the 1_{01} state of 5% (as a bold cyan line) or 0% (as a bold blue line), with partial contributions of the $1_{11} \leftarrow 0_{00}$ transition (as an orange line) and the $1_{11} \leftarrow 1_{01}$ transition (as a red line). Shown as dashed lines, the simulated total theoretical contributions are convoluted to the experimental collision-energy spread for the 1_{01} rotational level, assuming an initial rotational population from the 1_{01} state of 5% (in green) or 0% (in olive), as done for the HDO + Ne system.



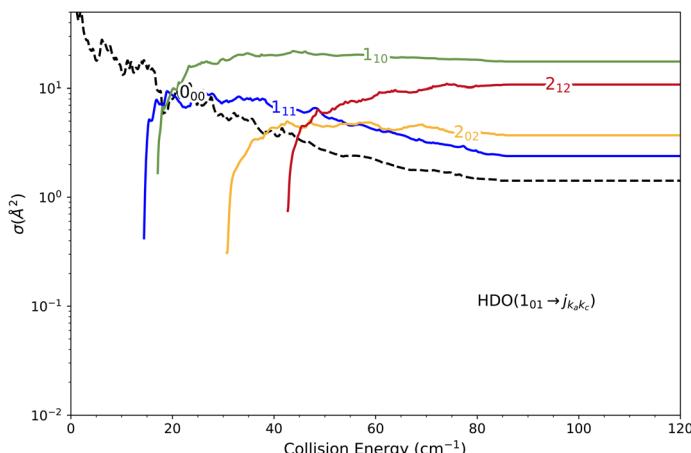


Fig. 14 Cross sections for the excitation (as solid lines) and relaxation (as a dashed line) of the first excited rotational level of the HDO molecule by collision with *normal*-H₂. The curves correspond to transitions from the 1₀₁ level to the six closest in energy rotational states.

caused by overlapping resonances are in perfect concordance up to about 80 cm⁻¹. For comparison, the experimental data have been normalised with a single rescaling value corresponding to the cross-section at 51.6 cm⁻¹. It also indicates that the water beam contains less than 5% HDO in the first excited 1₀₁ state: there is no signal below 25 cm⁻¹ collision energy. The excellent agreement between theory and experiment for the 1₁₁ \leftarrow 0₀₀ ICSs makes the disagreement for the 1₀₁ \leftarrow 0₀₀ even more puzzling. Moreover, for collision energies above 80 cm⁻¹, it can be noticed that the experimental 1₁₁ \leftarrow 0₀₀ cross-section is significantly larger than the theoretical prediction. In this collision-energy regime, a large resonance is, however, not expected. More experimental acquisitions are needed to reduce the error bars. In conclusion, in both cases, additional work is clearly necessary to reconcile these discrepancies.

5 Conclusions

Scattering resonances in the rotational excitation of HDO by Ne and *normal*-H₂ were studied through a joint theoretical and experimental approach. For the 1₁₁ \leftarrow 0₀₀ transition of HDO + Ne collisions, a good agreement was found for the overall shape of the cross-sections and the near-threshold rise, but the large experimental collision-energy spread due to the reduced mass of the system did not allow us to see the calculated resonances. In the case of the 1₀₁ \leftarrow 0₀₀ transition, even if the experimental signal was in the noise, the small undulations due to resonances were nicely reproduced by the calculations. Moreover, this system afforded the supplementary opportunity to theoretically compare the collisions of Ne with the three water isotopes, H₂O, D₂O, and HDO. We discovered that HDO and H₂O produce very similar outcomes, whereas D₂O differs from the other two. From this perspective, we demonstrated that straightforward reasoning, hinging on the symmetry of the potential terms $v_{lm}(R)$, facilitated the comprehension of

the scattering outcomes. In the case of HDO + *normal*-H₂, an excellent agreement was found for the $1_{11} \leftarrow 0_{00}$ cross-section between the theoretical and experimental results. However, for the $1_{01} \leftarrow 0_{00}$ cross-section, theory predicted an effective maximum cross-section whose magnitude is close to that of the $1_{11} \leftarrow 0_{00}$ transition, although no signal was experimentally detected. Thus, additional work is clearly necessary to reconcile this discrepancy.

Author contributions

All the authors have participated in the investigation, validation and writing. RMGV, AB, ODA, AF and TS carried out the formal analysis of the theoretical and experimental data. AB, ODA, AF and TS acquired the funding.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 E. F. van Dishoeck, *et al.*, *Astron. Astrophys.*, 2021, **648**, A24, DOI: [10.1051/0004-6361/202039084](https://doi.org/10.1051/0004-6361/202039084).
- 2 J. L. Neill, S. Wang, E. A. Bergin, N. R. Crockett, C. Favre, R. Plume and G. J. Melnick, *Astrophys. J.*, 2013, **770**, 142, DOI: [10.1088/0004-637X/770/2/142](https://doi.org/10.1088/0004-637X/770/2/142).
- 3 K. Furuya, E. F. van Dishoeck and Y. Aikawa, *Astron. Astrophys.*, 2016, **586**, A127, DOI: [10.1051/0004-6361/201527579](https://doi.org/10.1051/0004-6361/201527579).
- 4 I. Kamp, *et al.*, *Faraday Discuss.*, 2023, **245**, 112, DOI: [10.1039/d3fd00013c](https://doi.org/10.1039/d3fd00013c).
- 5 A. Andreu, A. Coutens, F. Cruz-Sáenz de Miera, N. Houry, J. K. Jørgensen, A. Kóspál and D. Harsono, *Astron. Astrophys.*, 2023, **677**, L17, DOI: [10.1051/0004-6361/202347484](https://doi.org/10.1051/0004-6361/202347484).
- 6 J. J. Tobin, M. L. R. van't Hoff, M. Leemker, E. F. van Dishoeck, T. Panque-Carreño, K. Furuya, D. Harsono, M. V. Persson, L. I. Cleeves, P. D. Sheehan and L. Cieza, *Nature*, 2023, **615**, 227, DOI: [10.1038/s41586-022-05676-z](https://doi.org/10.1038/s41586-022-05676-z).
- 7 M. Sewiło, *et al.*, *Astrophys. J.*, 2022, **933**, 64, DOI: [10.3847/1538-4357/ac6de1](https://doi.org/10.3847/1538-4357/ac6de1).
- 8 L. Yang, F. J. Ciesla and C. M. O. 'D. Alexander, *Icarus*, 2013, **226**, 256, DOI: [10.1016/j.icarus.2013.05.027](https://doi.org/10.1016/j.icarus.2013.05.027).
- 9 (a) A. Bergeat, A. Faure, L. Wiesenfeld, C. Miossec, S. B. Morales and C. Naulin, *Molecules*, 2022, **27**, 7535, DOI: [10.3390/molecules27217535](https://doi.org/10.3390/molecules27217535); (b) A. Bergeat,



S. B. Morales, C. Naulin, A. Faure and L. Wiesenfeld, *Phys. Rev. Lett.*, 2020, **125**, 143402, DOI: [10.1103/PhysRevLett.125.143402](https://doi.org/10.1103/PhysRevLett.125.143402).

10 A. Bergeat, A. Faure, S. B. Morales, A. Moudens and C. Naulin, *J. Phys. Chem. A*, 2020, **124**, 259, DOI: [10.1021/acs.jpca.9b04753](https://doi.org/10.1021/acs.jpca.9b04753).

11 A. Faure, M. Źołtowski, L. Wiesenfeld, F. Lique and A. Bergeat, *Mon. Not. R. Astron. Soc.*, 2024, **527**, 3087, DOI: [10.1093/mnras/stad3037](https://doi.org/10.1093/mnras/stad3037).

12 S. Green, *Astrophys. J., Suppl. Ser.*, 1989, **70**, 813, DOI: [10.1086/191358](https://doi.org/10.1086/191358).

13 H. M. Pickett, R. L. Poynter, E. A. Cohen, M. L. Delitsky, J. C. Pearson and H. S. P. Müller, *J. Quant. Spectrosc. Radiat. Transf.*, 1998, **60**, 883, DOI: [10.1016/S0022-4073\(98\)00091-0](https://doi.org/10.1016/S0022-4073(98)00091-0).

14 A. Faure, L. Wiesenfeld, Y. Scribano and C. Ceccarelli, *Mon. Not. R. Astron. Soc.*, 2012, **420**, 699, DOI: [10.1111/j.1365-2966.2011.20081.x](https://doi.org/10.1111/j.1365-2966.2011.20081.x).

15 L. Wiesenfeld, Y. Scribano and A. Faure, *Phys. Chem. Chem. Phys.*, 2011, **13**, 8230, DOI: [10.1039/c0cp02591g](https://doi.org/10.1039/c0cp02591g).

16 G. Sarma, C.-H. Yang, A. Kumar Saha, D. H. Parker and L. Wiesenfeld, *J. Chem. Phys.*, 2013, **138**, 024314, DOI: [10.1063/1.4772600](https://doi.org/10.1063/1.4772600).

17 (a) S. Chefdeville, T. Stoecklin, A. Bergeat, K. M. Hickson, C. Naulin and M. Costes, *Phys. Rev. Lett.*, 2012, **109**, 023201, DOI: [10.1103/PhysRevLett.109.023201](https://doi.org/10.1103/PhysRevLett.109.023201); (b) S. Chefdeville, T. Stoecklin, C. Naulin, P. Jankowski, K. Szalewicz, A. Faure, M. Costes and A. Bergeat, *Astrophys. J. Lett.*, 2015, **799**, L9, DOI: [10.1088/2041-8205/799/1/L9](https://doi.org/10.1088/2041-8205/799/1/L9).

18 M. Lara, F. Dayou, J.-M. Launay, A. Bergeat, K. M. Hickson, C. Naulin and M. Costes, *Phys. Chem. Chem. Phys.*, 2011, **13**, 8127, DOI: [10.1039/C0CP02705G](https://doi.org/10.1039/C0CP02705G).

19 A. Bergeat, S. B. Morales, C. Naulin, J. Klos and F. Lique, *Front. Chem.*, 2019, **7**, 164, DOI: [10.3389/fchem.2019.00164](https://doi.org/10.3389/fchem.2019.00164).

20 C. Naulin and A. Bergeat, *Cold Chemistry: Molecular Scattering and Reactivity Near Absolute Zero*, ed. O. Dulieu and A. Osterwalder, The Royal Society of Chemistry, 2017, ch. 3, p. 92, DOI: [10.1039/9781782626800](https://doi.org/10.1039/9781782626800).

21 C.-H. Yang, G. Sarma, J. J. ter Meulen, D. H. Parker and C. M. Western, *Phys. Chem. Chem. Phys.*, 2010, **12**, 13983, DOI: [10.1039/c0cp00946f](https://doi.org/10.1039/c0cp00946f).

22 C. M. Western, *J. Quant. Spectrosc. Radiat. Transf.*, 2017, **186**, 221, DOI: [10.1016/j.jqsrt.2016.04.010](https://doi.org/10.1016/j.jqsrt.2016.04.010).

23 G. Knizia, T. B. Adler and H.-J. Werner, *J. Chem. Phys.*, 2009, **130**, 054104, DOI: [10.1063/1.3054300](https://doi.org/10.1063/1.3054300).

24 K. A. Peterson, T. B. Adler and H. J. Werner, *J. Chem. Phys.*, 2008, **128**, 084102, DOI: [10.1063/1.2831537](https://doi.org/10.1063/1.2831537).

25 A. Halkier, W. Klopper, T. Helgaker, P. Jorgensen and P. R. Taylor, *J. Chem. Phys.*, 1999, **111**, 9157, DOI: [10.1063/1.479830](https://doi.org/10.1063/1.479830).

26 H.-J. Werner, P. J. Knowles, G. Knizia, F. R. Manby and M. Schütz, *Wiley Interdiscip. Rev. Comput. Mol. Sci.*, 2012, **2**, 242, DOI: [10.1002/wcms.82](https://doi.org/10.1002/wcms.82).

27 O. Denis-Alpizar and T. Stoecklin, *Astron. Astrophys.*, 2020, **638**, A31, DOI: [10.1051/0004-6361/202037821](https://doi.org/10.1051/0004-6361/202037821).

28 R. M. García-Vázquez, O. Denis-Alpizar and T. Stoecklin, *J. Phys. Chem. A*, 2023, **127**, 4838, DOI: [10.1021/acs.jpca.3c02086](https://doi.org/10.1021/acs.jpca.3c02086).

29 T. S. Ho and H. Rabitz, *J. Chem. Phys.*, 1996, **104**, 2584, DOI: [10.1063/1.470984](https://doi.org/10.1063/1.470984).

30 T. Stoecklin, O. Denis-Alpizar, A. Clergerie, P. Halvick, A. Faure and Y. Scribano, *J. Phys. Chem. A*, 2019, **123**, 5704, DOI: [10.1021/acs.jpca.9b04052](https://doi.org/10.1021/acs.jpca.9b04052).



31 B. A. Voronin, J. Tennyson, R. N. Tolchenov, A. A. Lugovskoy and S. N. Yurchenko, *Mon. Not. R. Astron. Soc.*, 2010, **402**, 492, DOI: [10.1111/j.1365-2966.2009.15904.x](https://doi.org/10.1111/j.1365-2966.2009.15904.x).

32 D. E. Manolopoulos, *J. Chem. Phys.*, 1986, **85**, 6425, DOI: [10.1063/1.451472](https://doi.org/10.1063/1.451472).

33 P. Valiron, M. Wernli, A. Faure, L. Wiesenfeld, C. Rist, S. Kedžuch and J. Noga, *J. Chem. Phys.*, 2008, **129**, 134306, DOI: [10.1063/1.2988314](https://doi.org/10.1063/1.2988314).

34 J. M. Hutson and S. Green, *Astrophysics Source Code Library*, 2012, record ascl: 1206.004.

35 P. J. Dagdigian, *Mon. Not. R. Astron. Soc.*, 2022, **511**, 3440, DOI: [10.1093/mnras/stac216](https://doi.org/10.1093/mnras/stac216).

