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Precision measurement of the fundamental vibrational frequencies of tritium-bearing hydrogen molecules: T₂, DT, HT

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High-resolution coherent Raman spectroscopic measurements of all three tritium-containing molecular hydrogen isotopologues T₂, DT and HT were performed to determine the ground electronic state fundamental Q-branch ($\nu = 0 \rightarrow 1$, $\Delta J = 0$) transition frequencies at accuracies of 0.0005 cm⁻¹. An over hundred-fold improvement in accuracy over previous experiments allows the comparison with the latest *ab initio* calculations in the framework of non-adiabatic perturbation theory including nonrelativistic, relativistic and QED contributions. Excellent agreement is found between experiment and theory, thus providing a verification of the validity of the NAPT-framework for these tritiated species. While the transition frequencies were corrected for ac-Stark shifts, the contributions of non-resonant background as well as quantum interference effects between resonant features in the nonlinear spectroscopy were quantitatively investigated, also leading to corrections to the transition frequencies. Methods of saturated CARS with the observation of Lamb dips, as well as the use of continuous-wave radiation for the Stokes frequency were explored, that might pave the way for future higher-accuracy CARS measurements.

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1 Introduction

Molecular hydrogen represents a benchmark molecule for testing quantum chemical theories. The comparison between precision spectroscopic measurements on the level structure of hydrogen and the results from calculations has made H₂ and its isotopologues a test bed for searches of fifth forces,¹ higher dimensions,² and physics beyond the Standard Model of physics.³ On the experimental side the measurement of the dissociation energy of H₂ has witnessed great improvements in the past decade^{4–7} now reaching a relative accuracy of 3×10^{-10} . Each isotopologue supports over 300 bound and long-lived rovibrational levels, that provide an extended playing field for performing fundamental tests, such as the ground tone vibrational splitting,⁸ the quadrupole overtone transitions of H₂^{9–11} and in the D₂ species,^{12,13} as well as in the mixed isotopologue HD.^{14–17}

On the theoretical side level energy calculations of the molecular hydrogen four-body system have undergone equally

great improvements, producing highly accurate level energies of H₂ and D₂¹⁸ as well as for HD.¹⁹ These methods were based on calculations of Born–Oppenheimer energies,²⁰ and separate approaches for adiabatic²¹ and non-adiabatic²² corrections, as well as computations of relativistic²³ and quantum electrodynamic effects.²⁴ This development has led to the comprehensive approach of nonadiabatic perturbation theory (NAPT),^{25,26} which now enables the rapid computation of all bound rovibrational states in the ground electronic manifold of the hydrogen isotopologues. In addition, and alongside, an even more refined and more accurate method involving direct 4-body calculations is being pursued, but this is currently limited to the computation of the binding energy of the lowest rovibrational level.^{27,28}

Most of the experimental precision spectroscopic studies were performed in the typical environment of molecular beams that can be easily applied to the stable, non-radioactive, hydrogen isotopologues H₂, HD and D₂. Tritium-containing isotopologues, HT, DT and T₂ have been much less frequently studied, because of limited access and handling difficulties due to safety requirements holding for radioactive species. The first spectroscopic studies on tritium bearing hydrogen molecules were performed by Dieke and Tomkins at Argonne National Laboratories recording emission spectra.^{29,30} Later, other methods were applied, such as spontaneous Raman spectroscopy^{31–33} and intracavity laser absorption, combined with an optophone.³⁴ A rationale for

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exploring tritium spectroscopy is that the heavier, tritiated molecular hydrogen species exhibit smaller non-adiabatic contributions to the level energies. High precision studies on these species may help to disentangle the effect of mass-dependent terms in the calculation of hydrogen level energies. Also, inclusion of tritium provides two additional heteronuclear species, HT and DT. These species provide information on the g/u mixing effects present in heteronuclear species. Such studies might help resolve the discrepancy observed in HD, between the experimental value for the dissociation energy³⁵ and the theoretical value.²⁸

The precise level structure of the tritium bearing hydrogen molecules is of relevance for the Karlsruhe Tritium Neutrino experiment (KATRIN) focusing on the β decay from T_2 .³⁶ The β -electron energy spectrum depends on the final state distribution of the $^3\text{HeT}^+$ daughter molecules,³⁷ requiring accurate level energies of $^3\text{HeT}^+$, but also on the isotopologue distribution, parent quantum state distribution, and the level structure of the parent states.^{38–40} The Raman spectroscopic techniques for molecular hydrogen isotopologues, including tritium-bearing species, were further developed to determine composition and concentration of the isotope mixtures for the KATRIN experiment.^{41–43}

Here we present a study of transition frequencies in the fundamental vibrational band of all tritiated species, T_2 , DT and HT, *via* Coherent Anti-Stokes Raman Scattering (CARS). After preliminary reports on the CARS detection,⁴⁴ on the spectroscopy of the T_2 isotopologue⁴⁵ and DT,⁴⁶ we now present a comprehensive report, including results on HT and further detailing the experimental methods. It is noted that remeasurements were performed for the T_2 species yielding improved statistics, following up on a previous set of data.⁴⁵ A comparison is made with the findings of theoretical calculations in the NAPT-framework.²⁶

2 Experiment

For designing an experiment on tritium-bearing hydrogen isotopologues aspects of obtainable experimental precision have to be matched to safety regulations with regard to radioactivity. The requirement of a legal upper limit of 1 GBq of tritium stored in the laboratory, corresponding in case of T_2 to a gas amount of 11.5 mbar cm^3 at room temperature, and the prohibition of pumping gas into an exhaust, sets constraints on the experimental method. Whereas in many previous precision studies on molecular hydrogen multi-step laser excitation was combined with ion detection^{4–6,47} this is not possible in case of tritium, in view of the $\sim 10^9$ fast electrons and ions being produced per second from the radioactive sample. In view of containment requirements molecular beam studies are ruled out as well.

A closed-cell environment with a fill of static gas is chosen as a safe solution. Spectroscopic detection *via* coherent Raman processes is more sensitive than direct linear infra-red absorption, while in both cases Doppler broadening scales linearly with the

vibrational energy, so imposing the same level of frequency uncertainty. These arguments led us to the choice of performing CARS experiments for the tritium-bearing hydrogen molecules. The CARS nonlinear optical process generates a coherent beam of light propagating into the far field where the Anti-Stokes signal can be collected and detected. In this section the handling of the tritium gas cell, the optical layout of the CARS setup, and the frequency calibration are detailed, while systematic effects are discussed.

2.1 Tritium gas samples

The sample cell for the present CARS measurements is specially built from well-proven materials for tritium gas containment, and has been described previously.⁴⁴ The cell with an inner volume of 4 cm^3 has input and output windows placed at a distance of 8 cm, so that focusing at $f = 20$ cm is possible without burning window surfaces and avoiding a too high intensity in the focus, in view of issues related to the ac-Stark effect (see below).

First, a high purity T_2 gas sample was prepared in the Tritium Laboratory Karlsruhe (TLK). The cell filled was filled to 2.5 mbar to be below the legal activity limit including a safety margin.

The heteronuclear molecules HT or DT can be generated by catalytic self-exchange reactions driven by ions⁴⁸ produced in the constant β -radiation intensity from tritium decay.

with X being H or D. The estimated HT and DT partial pressures are derived from equilibrium constants K_{eq} :

$$K_{\text{eq}} = \frac{[\text{XT}]^2}{[\text{X}_2][\text{T}_2]},$$

Values for K_{eq} are determined⁴⁹ at 2.58 for HT and 3.82 for DT at 300 K.

For the preparation of DT, first D_2 and T_2 were injected at a 4:1 ratio into a large mixing vessel. After a short time DT is formed *via* radio-chemical equilibration. A sample was extracted and filled into the CARS cell up to a total pressure of 12.7 mbar. The partial pressure of DT is estimated to be about 3.8 mbar. At the time of the preparation of the HT cell, this accurate mixing procedure was not available due to constraints with other running experiments. Thus, HT preparation needed to be performed inside the CARS-cell. The 2.5 mbar T_2 sample was topped up with H_2 up to a total pressure of 11.5 mbar. This would lead to a theoretical partial pressure of 3.6 mbar. It should be noted that this procedure is less accurately controllable as tritium gas might have been partially back-diffused from the cell reducing the actual HT yield.

In view of the tritium radioactive half-life of 12.3 years these (partial) pressures effectively remained constant during the measurements performed in time windows of months. The safety-controlled and tritium-gas filled cell was transported to LaserLaB Amsterdam, where the CARS measurements were carried out.

2.2 Precision CARS setup

In the present study, the fundamental band ($\nu = 1 \leftarrow 0$) Q-branch transitions of all three tritiated species are measured by the CARS non-linear optical technique. The specific resonant four-wave mixing scheme of the CARS process is shown in Fig. 1 alongside with an overview spectrum of the Q-lines in DT. An anti-Stokes field is generated, at frequency $\omega_{AS} = 2\omega_P - \omega_S$, using input fields of a pump frequency ω_P and a Stokes frequency ω_S . The integrated anti-Stokes signal intensity I_{AS} is produced as a coherent light beam in the forward direction under phase-matching conditions and is proportional to:⁵⁰

$$I_{AS} \propto \int |\chi^{(3)}|^2 I_P^2(z, t) I_S(z, t) dz dt, \quad (1)$$

where $I_P(z, t)$ and $I_S(z, t)$ are the spatial and temporal evolution of pump and Stokes intensity and z is the optical path. The third order nonlinear susceptibility $\chi^{(3)}$ depends on the molecular number density and the differential Raman cross-section.⁵⁰

The optical layout of the CARS setup is shown in Fig. 2. The pump beam is obtained from a frequency-doubled injection-seeded Nd-YAG laser (Spectra-Physics GCR-330) delivering pulses at 532 nm, of 8 ns duration, which would correspond to a bandwidth of about 55 MHz in case of Fourier transform (FT) limited laser pulses. During the measurements, only low pulse energies were used, ranging from 30 to 500 μ J in the interaction range.

For the precision CARS measurements the Stokes beam is obtained from an injection-seeded travelling-wave pulsed dye amplifier (PDA) operating on DCM dye in methanol.⁵¹ The continuous-wave (cw) seed light for the PDA is obtained through an optical fiber from a ring-dye laser, covering the range from 590 nm to 650 nm. With 150 mW seed power and 50 mJ pumping from a second frequency-doubled injection-seeded Nd-YAG laser (Spectra-Physics Pro-250), the output energy reaches about 5 mJ at 633 nm (peak performance wavelength) at 6 ns pulse duration, which would correspond to a bandwidth of about a 74 MHz in case of FT-limited laser

Fig. 2 Optical layout for the CARS spectroscopy setup. The frame encloses the setups for frequency calibration of the pump and Stokes beams. See text for more details.

pulses; the actual width will be slightly larger in view of the chirp phenomena detected (see below). The pulse energy of the Stokes beam in the CARS experiment was kept unchanged for most part of the measurement campaigns at <20 μ J.

The pump and Stokes beams are combined on a longpass dichroic mirror and passed through a polarization beam splitter cube to ensure parallel polarizations. The two pulses are focused co-linearly into a gas cell with a $f = 20$ cm lens. The flat-top pump pulse beam waist is about 44 μ m, as measured by a CCD camera, for a 3 mm Rayleigh length. The Stokes pulse is estimated to be 1.7 times larger in diameter than the pump beam. Part of the merged beams is sampled out over a long distance for alignment purposes, with the angle mismatch estimated to be better than 2 mrad. The generated CARS signal (at $\lambda \sim 450$ –470 nm) travels along the same propagation axis of the pump and Stokes beams. The CARS-signal beam is collimated with a $f = 10$ cm lens, dispersed from the incident beams by a Pellin–Broca prism, and propagated over a 2 m separation path. The signal beam is then passed through an aperture and bandpass filter, and detected by photomultiplier tube (Philips XP-1911) mounted inside a dark box.

The measurements are performed with few-mbar pressure gas samples, and the spectra obtained are expected to be Doppler-limited. The expression for Doppler broadening (FWHM) of forward spontaneous Raman scattering is:⁵²

$$\Delta\nu = \frac{2\nu_0}{c} \left(\frac{2k_B T \ln 2}{m} \right)^{1/2} \quad (2)$$

where ν_0 is the fundamental vibrational frequency splitting, m is the molecular mass, k_B the Boltzmann constant, c the speed of light, and T the temperature. This results in a Doppler width at room temperature of 370 MHz for the heaviest species T_2 , and about 450 MHz and 630 MHz for DT and HT, respectively. CARS spectral profile simulations by Lucht and Farrow⁵² showed that the spectral width of the CARS resonances is about 1.2 times that of the spontaneous Raman scattering in the forward direction. These values for the width are to be further increased by contributions of the laser bandwidth of both lasers, some additional broadening caused by the timing jitter in the temporal overlap of the two pulses (within 3.5 ns), and by ac-Stark broadening.

Fig. 1 Overview recording of the $X^1\Sigma_g^+$ ($\nu = 0 \rightarrow 1$) Q(0–6) lines in DT using a combination of an injection-seeded Nd:YAG laser (pump) and a grating-based pulsed-dye laser (Stokes). The inset shows an energy diagram for the CARS four-wave mixing scheme, with the generated Anti-Stokes radiation detected as signal.

2.3 Frequency calibration

The frequency calibration of the Stokes beam was performed in two stages. As shown in Fig. 2, part of the cw seed-radiation was split off to perform I_2 -saturation spectroscopy for absolute calibration against well-calibrated I_2 -standards,⁵³ and another part for measuring transmission markers from an etalon (FSR = 150.67(2) MHz) stabilized by a HeNe laser (Thorlabs HRS-015). The resulting uncertainty in the absolute frequency calibration of the cw seed laser amounts to 2 MHz.

In a second step the chirp-induced pulse-cw frequency offset was measured by techniques previously employed for frequency calibration of a PDA-system.^{51,54,55} The frequency of the cw-seed was shifted by +700 MHz by an acousto-optic modulator (AOM) in a double-pass configuration and the beatnote between the shifted cw-seeding and Stokes pulses was recorded using a fast photodetector and a 4-GHz bandwidth oscilloscope. The setup for producing a beat-note signal is displayed in Fig. 3a. Following Hannemann *et al.*⁵⁶ for the frequency chirp analysis, the time-dependent phase in the duration of the pulse is extracted from the beat-note signal, to yield a time-dependent frequency offset curve as plotted in Fig. 3b. The reported average frequency offset values are obtained by time-integrating the frequency offset over the pulse duration, weighted by the instantaneous intensity.

A systematic study of frequency deviations between the cw-seed frequency and the PDA frequency was performed for various settings of the PDA. Typical results for measured frequency offsets in the PDA, running on DCM in methanol, are displayed in Fig. 4. The chirp-offset is found to vary from 23 MHz at 610 nm (Stokes wavelength for T_2) to 5 MHz at 650 nm (Stokes wavelength for HT) as shown in Fig. 4a, while also a dependence on the intensity was observed as shown in Fig. 4b. In addition, the effect of frequency chirp over the spatial beam profile of the Stokes beam was measured, and was observed to vary within 2 MHz. During the CARS measurements

Fig. 3 Setup for the frequency chirp measurement of the output of the PDA laser system. The cw-seed laser is shifted by double-passing an AOM at 350 MHz modulation frequency. The shifted cw and pulsed laser outputs are coupled through a single mode 2×2 fused fibre coupler for imposing spatial overlap. The beat note in (a) is detected on a fast photodetector (Thorlabs DET025AFC) and stored in a Tektronix TDS7404 oscilloscope. (b) The instantaneous frequency offset, $V_{\text{offset}} = \nu_{\text{pulse}} - \nu_{\text{cw}}$, is obtained from the phase evolution of the 700 MHz carrier signal.⁵⁶

Fig. 4 Result of frequency chirp measurements of the PDA laser system. (a) Pulse-cw frequency offset Stokes beam from PDA (DCM in Methanol) at different wavelengths with 50 mJ pump energy. (b) Frequency offset of the PDA at 633 nm for different pump energies. (c) Daily variation of frequency offset of the PDA at 633 nm.

the chirp analysis was performed after co-propagating the output of the pulsed laser with that of the AOM-shifted cw-laser output in a single mode fiber, so that this spatial effect was averaged out. Systematic chirp-analysis measurements were performed for the laser settings during every measurement day. The frequency calibration of the PDA-laser was corrected for the chirp results during the CARS precision measurements, for which an estimated uncertainty of 5 MHz represents the standard deviation over different measurement days, as shown in Fig. 4c.

The frequency of the 532 nm pump pulse, obtained from the injection-seeded Nd:YAG laser, was directly measured by a High Finesse WSU-30 (Toptica) wavemeter. The absolute calibration of the wavemeter was verified during each measurement run by measurements of I_2 -hyperfine transitions in the range 600 to 650 nm, as well as measurement of a cw Ti:Sa laser at 718 nm locked to a frequency comb laser.⁵ These measurements result in a value for the typical drift of the wavemeter, determined at 0.3 MHz h^{-1} . In addition the effect of measuring cw laser *vs.* pulsed laser radiation was verified by measuring the frequency of the output of the cw ring laser seed frequency *vs.* the output of the PDA. As a result of these repeated procedures the uncertainty of the frequency of the pump pulse is estimated at 6 MHz. In a recent study a similar model wavemeter (WSU-2) was used in measurements of several $^{40}\text{Ca}^+$ ion transitions and the calibrations methods led to an absolute accuracy of 5 MHz.⁵⁷

2.4 ac-Stark shift

The ac-Stark effects occurring in CARS measurements of the hydrogen molecule have been investigated in detail.^{58–60} In our current setup, the ac-Stark shift is caused by both Stokes and pump pulses and can be approximated in terms of pulse energies E_P , E_S , areas at the beam waist A_P , A_S and pulse durations (FWHM) τ_P , τ_S , and the ac-Stark shift can be expressed as:

$$\delta\nu = \kappa_P \frac{E_P}{A_P \tau_P} + \kappa_S \frac{E_S}{A_S \tau_S}.$$

The constant κ , for both P and S depends on the difference of the ac-Stark shift in upper and lower states *via*:

$$\kappa = -\frac{2\pi}{hc} \left((\alpha_{\nu'} - \alpha_{\nu''}) + \frac{2J(J+1) - 3m_f^2}{3(2J+3)(2J-1)} (\gamma_{\nu'} - \gamma_{\nu''}) \right) \quad (3)$$

where $\alpha_{\nu'}$ and $\gamma_{\nu'}$ are the dynamic (frequency-dependent) isotropic polarizabilities and anisotropic polarizabilities at frequencies ν' and ν'' , respectively. The zero field transition frequency can be obtained by performing measurements at different Stokes and pump energies and extrapolation to zero. Since the frequencies used in CARS are far from the dipole-allowed transitions in molecular hydrogen, the polarizability term only weakly depends on the frequency in the optical range⁶⁰ and the expression can be approximated as (with $\kappa_P = \kappa_S = \kappa$):

$$\delta\nu = \kappa \left(\frac{E_P}{A_P \tau_P} + \frac{E_S}{A_S \tau_S} \right).$$

As the effective area and pulse width of the pump and Stokes beams are affected by spatial and temporal overlap between the two pulses, the value and the reproducibility of the shift coefficient in terms of pulse energy for pump and Stokes were measured extensively on the DT Q(1) transition,⁴⁶ yielding $\frac{\kappa_P}{A_P \tau_P} = -0.0064(7) \text{ cm}^{-1} \text{ mJ}^{-1}$ and $\frac{\kappa_S}{A_S \tau_S} = -0.0006(2) \text{ cm}^{-1} \text{ mJ}^{-1}$. To find a balance between minimal ac-Stark shift and sufficient signal strength, the Stokes pulse energy was kept below 10 μJ for the T_2 and DT measurements. In such conditions, the uncertainty of the ac-Stark shift from the Stokes beam, upon extrapolation to zero intensity, is estimated to be less than 1 MHz. Due to the weaker signal strengths obtained in the HT experiments, the Stokes energy was raised to 20 μJ , while the pump energy was varied from 90 to 500 μJ . Here the pump energy was not increased further in view of saturation and asymmetry phenomena observed in the spectra, as discussed below.

2.5 Pressure shift

Collisional shift coefficients of Raman transitions in stable molecular hydrogen isotopologues are well studied in the large pressure range from tens of millibars to a few bars of total pressure.^{61–63} For the reported shift coefficients the presently measured CARS lines are estimated to undergo shifts below ~ 1 MHz. In addition, the collisional shift of D_2 was investigated in the present study by pressure-dependent CARS measurements of pure D_2 in an identical gas cell. The shift coefficient is found to equal $0.06 \text{ MHz mbar}^{-1}$ under room temperature conditions.

For the T_2 measurements, the sample contains 93.4% of T_2 .⁴⁴ The collisional shift is assumed to be similar to the self-collisional shift coefficient of H_2 and D_2 yielding a shift of about 0.3 MHz. From the reported result of HD,⁶³ the pressure shift in this heteronuclear species was found to be twice that of H_2 and D_2 , which was ascribed to the presence of a small permanent dipole moment and the lack of a selection rule prohibiting *para-ortho* conversion during rotationally inelastic

collisions.⁶⁴ The DT and HT samples inevitably contain some amounts of D_2 and H_2 , respectively. We conclude that based on previous studies,^{61–63} and on present data for D_2 , the collisional shift in the present CARS experiments is well below 1 MHz.

Some of the CARS measurements on D_2 were performed from the DT gas sample cell, hence under conditions of a low-density plasma environment, undergoing constant production of about 10^9 electrons per second at an average kinetic energy of 5 keV and the same rate of ion production from the primary beta decays. In addition, secondary electrons and ions are produced from ionization of the neutral gas. The transition frequencies of the D_2 CARS signals were not found to deviate from the signals from a different gas cell filled with 4 mbar of pure D_2 , without addition of radioactive species. We conclude that there occurs no measurable plasma shift on the transition frequencies as reported in Table 2.

2.6 Other effects

There is underlying hyperfine structure in the T_2 , DT and HT transitions, where the hyperfine splitting spans a range of less than a MHz. This hyperfine structure is similar to those in the stable molecular hydrogen species such as HD, which was shown to affect vibrational resonance lineshapes in the high-resolution investigations of Diouf *et al.*⁶⁵ However, in view of the resolution of the present experiment this does not affect the determination of the transition frequency positions (hyperfine center-of-gravity). Due to its closed-shell nature, the ground electronic state transitions in molecular hydrogen are insensitive to DC Stark shifts. Using the static polarizabilities from ref. 66, we estimate⁶⁰ an upper limit of well below 1 Hz for a DC field amplitude of 1 V cm^{-1} . The Stark effect due to the plasma generated from the β -decay might be expected to be higher, but this plasma effect was not observed in the D_2 measurements discussed in the previous section. Likewise, the Zeeman shifts of the hyperfine components are also very low, with estimated shifts using constants from ref. 67 of well below 1 Hz due to the earth's magnetic field. In addition, for the Q transitions with the same upper J' and lower J'' rotational quantum numbers and almost identical hyperfine constants for the $\nu = 0$ and $\nu = 1$ states, the shifts in the transition frequency effectively cancel out. We thus neglect the contributions of the hyperfine structure, DC Stark and Zeeman effects from the uncertainty budget.

2.7 Frequency uncertainty in the CARS measurements

Table 1 lists all the uncertainty contributions to the line center determination in the present CARS study. For T_2 Q($J = 1-5$) and DT Q($J = 0-5$), the total uncertainty is about 12 MHz. For the T_2 Q(0) and DT Q(6) lines the uncertainty is estimated at 20 MHz, due to interference effects, which will be discussed in the last section. For HT, the lowest pulse energy used in the measurement was about 2.5 times larger than that for T_2 and DT measurements, thus resulting in a larger uncertainty from the ac-Stark effects. In addition, the Doppler width for the HT lines is 1.4 times larger than that for DT. Such differences are also found in results from different measurement days. For HT, the total uncertainty is about 16 MHz for the Q($J = 0-3$) lines.

Table 1 Uncertainty budget for the CARS measurements. Systematic and statistical contributions to the frequency uncertainties in the fundamental vibrational Raman Q($J < 6$) transitions, excluding T_2 Q(0). Values in units of 10^{-4} cm^{-1} and representing 1σ

Contribution	T_2 /DT	HT
Pump (ω_p) calibration	2	2
Stokes (ω_s) cw calibration	0.7	0.7
Stokes chirp correction	1.7	1.7
ac-Stark analysis	2	3.3
Collisional shift	0.3	0.3
Plasma shift	0.3	0.3
CARS interference shift	1	1
Statistics	2.3	3
Combined (1σ)	4.2	5.3

3 Results and discussions

Representative high-resolution CARS spectra of the ($\nu = 0 \rightarrow 1$) Q(1) transition for tritiated molecular hydrogen species T_2 , DT, and HT are displayed in Fig. 5. The line widths of the Q(1) transitions exhibit the expected Doppler widths with the broadest width for HT (lightest) and narrowest for T_2 (heaviest). While the Q($J = 0-5$) transitions in T_2 were previously reported in ref. 45, these lines have been remeasured in this study and the Q-branch extended up to $J = 7$. Moreover, the ($\nu = 0 \rightarrow 1$) S($J = 0-3$) transitions, which are much weaker than the Q lines, were also investigated in the present work. The DT ($\nu = 0 \rightarrow 1$) Q($J = 0-5$) transitions were recently reported in ref. 46 and extended here up to $J = 7$. The high-resolution measurements of the HT Q-branch transitions are presented here for the first time. The transition frequencies of the Q-branch lines in the fundamental band splitting ($\nu = 0 \rightarrow 1$) of all tritiated species are listed in Table 2.

3.1 HT, DT and T_2 transitions

The HT measurements include Q transitions from $J = 0-3$, where the signal strength involving higher J states is limited due to decreasing population. The HT partial pressure was somewhat lower than the targeted equilibrium partial pressure of 3.6 mbar due to issues in the HT gas filling procedure. In addition, the lower transmission of the bandpass filter in the spectral region of the HT Q-branch lines, led to a weaker observed signal strength. To compensate for the weaker signals

Table 2 Fundamental vibrational ($\nu = 0 \rightarrow 1$) splittings of the Q(J) transitions in T_2 , DT, and HT. The weaker S($J = 0-3$) transitions in T_2 are also listed. The measured values appear in the second column while the theoretical values are listed in the third column, with uncertainties indicated within parentheses in units of the last digit in the indicated value. The last column represents the difference ($\omega_{\text{exp}} - \omega_{\text{calc}}$) with the combined uncertainty from experiment and calculation indicated. Experimental values include corrections from the interference phenomena affecting the CARS lineshape. Lines indicated with asterisks (*) include uncertainty contributions from line profile asymmetry, while those with uncertainties of $5 \times 10^{-3} \text{ cm}^{-1}$ are dominated by ac-Stark broadening. All values in units of cm^{-1}

Line	Experiment	Calculation	Difference
T_2			
Q(0)	2464.50394 (67)*	2464.50415 (6)	-0.00021 (67)
Q(1)	2463.34817 (42)	2463.34836 (6)	-0.00019 (42)
Q(2)	2461.03917 (42)	2461.03917 (6)	+0.00000 (42)
Q(3)	2457.58135 (42)	2457.58137 (6)	-0.00002 (42)
Q(4)	2452.98233 (42)	2452.98211 (6)	+0.00022 (42)
Q(5)	2447.25061 (42)	2447.25085 (6)	-0.00025 (42)
Q(6)	2440.397 (5)	2440.39934 (6)	-0.002 (5)
Q(7)	2432.442 (5)	2432.44152 (6)	+0.000 (5)
S(0)	2581.114 (5)	2581.10522 (6)	+0.009 (5)
S(1)	2657.281 (5)	2657.28290 (6)	-0.002 (5)
S(2)	2731.716 (5)	2731.71084 (6)	+0.006 (5)
S(3)	2804.164 (5)	2804.16421 (6)	-0.000 (5)
DT			
Q(0)	2743.34160 (42)	2743.34174 (11)	-0.00014 (43)
Q(1)	2741.73204 (39)	2741.73210 (11)	-0.00006 (40)
Q(2)	2738.51662 (42)	2738.51697 (11)	-0.00035 (43)
Q(3)	2733.70479 (42)	2733.70466 (11)	+0.00013 (43)
Q(4)	2727.30745 (42)	2727.30755 (11)	-0.00010 (43)
Q(5)	2719.34221 (42)	2719.34202 (11)	+0.00019 (43)
Q(6)	2709.82835 (67)*	2709.82832 (11)	+0.00003 (68)
Q(7)	2698.787 (5)	2698.79048 (11)	-0.003 (5)
HT			
Q(0)	3434.81248 (53)	3434.81333 (44)	-0.00085 (69)
Q(1)	3431.57509 (53)	3431.57553 (44)	-0.00044 (69)
Q(2)	3425.11265 (53)	3425.11324 (44)	-0.00059 (69)
Q(3)	3415.45258 (53)	3415.45298 (44)	-0.00040 (69)

Fig. 5 High resolution CARS spectra of the Q(1) lines in the fundamental vibrational band ($\nu = 0 \rightarrow 1$) of T_2 , DT, and HT.

higher pulse energies were employed to obtain sufficient SNR. This resulted, however, in a larger ac-Stark uncertainty. These factors as well as the larger Doppler width lead to a larger total uncertainty in the HT transition energies than those of the DT and T_2 measurements. The present Q-branch transition energies are 6.4 cm^{-1} higher than the values reported by Edwards *et al.*³² These discrepancies were already pointed out

in the subsequent studies of Chuang and Zare³⁴ and Veirs and Rosenblatt.³³ The present HT Q-branch measurement results are consistent with and improve the measurement uncertainty of Veirs and Rosenblatt³³ by a factor of 200. Chuang and Zare³⁴ performed absorption measurements of weakly-allowed dipole P- and R-branch transitions in the (1, 0) fundamental band, as well as the (4, 0) and (5, 0) overtone bands of HT. When converting the transition frequencies of that study³⁴ into fitted molecular constants, good agreement is found with the present results, which are more accurate by seventeen-fold.

Measurements of the DT Q-branch were reported in our recent work⁴⁶ and are extended here to include Q(6) and Q(7) lines. The uncertainty for the much weaker Q(7) line is substantially larger than those for $J < 5$, caused mainly by the asymmetric line profile, which is discussed below. A comparison of the present DT Q-branch transition frequencies with the $Q(J = 0-4)$ values reported by Edwards *et al.*³² shows differences of some 0.15 cm^{-1} , hence much larger than their claimed uncertainty of 0.005 cm^{-1} .³² Veirs and Rosenblatt³³ already had cast doubt on the accuracy claimed by Edwards *et al.*³² given the discrepancies in the analogous HT and T_2 measurements. The DT Q-branch measurements presented here are consistent with those of Veirs and Rosenblatt³³ and improve the measurement uncertainty by a factor of more than 200 for the $Q(J = 2-7)$ transitions.

The present T_2 $Q(J = 0-5)$ values were obtained after implementing improvements in the setup used in ref. 45. Comparing the T_2 results listed in Table 2 with the values reported in ref. 45, reveals an offset of $< 0.001 \text{ cm}^{-1}$ for the Q(0), Q(1), and Q(3) transitions while showing consistent values for Q(2), Q(4) and Q(5). We attribute these discrepancies to several difficulties in the measurements of T_2 in the previous study. For example, the cw-seed instability of the pump laser for both CARS laser beams resulted in substantial time jitter leading to poor SNR, especially for weaker transitions. To compensate for such signal loss, the Stokes and the pump pulse energies used during the previous experiment were much higher than in the present study. The cross-sectional areas of both laser beams are now more accurately determined using two methods, a CCD camera beam profiler system and knife-edge method, leading to consistent values. Compared to the estimates of laser intensities in the previous study,⁴⁵ these could be off by a factor of three, which is important in the ac-Stark extrapolation studies. A set of measurements, carried out during a day, is based on an extrapolation relying only on relative intensities, assuming little alignment drift, and should lead to robust results. This becomes an issue when combining results of different days where any misalignment may lead to a slightly different overlap of the pump and Stokes laser beams, yielding a different effective area for production of CARS signal. Moreover, in the previous study the ac-Stark analysis for all transitions except Q(1) was performed by altering the pump beam intensity only, and thus the ac-Stark shift contribution due to differences in the Stokes beam overlap could be another source for the offset. These shortcomings were repaired in the present study leading to more reliable results.

The frequency calibration of the Stokes laser is also improved in the present study. Some transitions exhibit a more than 0.15 cm^{-1} separation from the calibration line, represented by a particular I_2 hyperfine component. Since such frequency span covers more than 30 markers of the reference etalon, length stabilization is now ensured over the whole recording time by locking to a stabilized HeNe-laser. Improved frequency calibration of the free spectral range (FSR = $150.67(2) \text{ MHz}$) of the reference cavity is also applied. The issues discussed here could have led to an underestimate of the uncertainties in the previous study.⁴⁵ In addition, the present measurements of T_2 lines in the improved setup allowed for recordings at lower pulse energies than in ref. 45, resulting in a reduced uncertainty of ac-Stark shift described in the previous section. These improvements have reduced systematic shifts, while at the same time have achieved better sensitivity and enhanced SNR of the recordings. These in turn enabled extending the measurements to weaker Q(6) and Q(7) transitions, as well as the T_2 S-branch with the transition frequencies listed in Table 2. The T_2 $Q(J = 0-5)$ results obtained in our new measurements represent a nearly 200-fold improvement compared with results from spontaneous Raman spectroscopy by Veirs and Rosenblatt.³³

3.2 Benchmark measurements on D_2

CARS measurements on pure D_2 were performed in an identical gas cell focusing on studies of systematic effects, such as the validation of pressure shift effects. This allowed for the comparison of D_2 Q(0), Q(1) and Q(2) of the fundamental vibrational splitting with previous precision studies using molecular beams and a deep-UV REMPI scheme.^{8,68}

In addition, quasi-cw CARS measurements were performed on D_2 Q(1) and Q(2) lines, using a cw Stokes beam, in combination with a pulsed pump beam. A quasi-cw CARS spectrum of the D_2 Q(2) transition is shown in Fig. 6. Such experiments bear the advantage that frequency chirp effects in the Stokes beam are absent. With 80 mW of cw Stokes radiation and a $20 \mu\text{m}$ beam waist, it was possible to detect the CARS signal using a pump energy of as low as $300 \mu\text{J}$ in a cell containing 8 mbar of D_2 gas. The accuracy of these quasi-cw CARS measurements is estimated to be 15 MHz, limited by the ac-Stark extrapolation.

Fig. 6 Quasi-cw CARS spectra of the Q(2) line in the fundamental band of D_2 recorded at 8 mbar using continuous wave Stokes radiation at 80 mW and a 8 ns pump pulse with 0.47 mJ pulse energy.

Results of a number of test experiments performed on D_2 are included in the rightmost panel of Fig. 7. The results of these studies using pulsed pump and Stokes radiation for $Q(J=0-2)$, as well as using pulsed pump and cw Stokes beams for $Q(1)$ and $Q(2)$ are in good agreement with the precision molecular beam experiments in ref. 8 and 68. These test experiments on D_2 verify the assessment of pressure shifts in the present CARS study. In addition, the use of chirp-free Stokes radiation in the quasi-cw CARS measurements on D_2 validates the chirp correction procedures implemented for the majority of data recorded with the PDA-laser system.

3.3 Comparison with calculations

Ab initio calculations, in the framework of NAPT, on level energies in the ground electronic state, are described in ref. 26 and 46. for all tritiated species. Rovibrational level energies and transition frequencies for all molecular hydrogen isotopologues in the ground electronic state can now be accessed through a publicly available program.⁶⁹ The differences between experimental results and calculated values, expressed as $\omega_{\text{exp}} - \omega_{\text{calc}}$, are plotted in Fig. 7 and listed in Table 2.

In Fig. 7, the experimental values are represented by data points with error bars attached, while a yellow-shaded region represents the uncertainty of the calculation. Very good agreement between measurement and theory is found for all tritiated species, as can be read from Fig. 7. For the tritium-bearing species, the major contribution to the theoretical uncertainty derives from the non-relativistic energy, $E^{(2)}$, which is about $\sim 6 \times 10^{-5} \text{ cm}^{-1}$ for the heaviest T_2 species, and about $\sim 4 \times 10^{-4} \text{ cm}^{-1}$ for HT, owing to its lighter mass.⁴⁶ Hence, the results from the calculations are more accurate than from experiments for T_2 and DT, while for the case of HT a similar accuracy is obtained, due to the less accurate $E^{(2)}$ term.

For the stable species H_2 , HD and D_2 , the $E^{(2)}$ terms have been calculated directly in a 4-particle variational approach (*i.e.* without invoking the Born-Oppenheimer approximation and corrections, hence outside the NAPT-framework), achieving

10^{-7} cm^{-1} uncertainty levels.²⁶ Therefore, the frequencies for the D_2 species are extremely accurate ($1.2 \times 10^{-5} \text{ cm}^{-1}$),²⁶ as represented by the narrow yellow strip in the rightmost panel of Fig. 7.

A similar calculation of improved accuracy might in future be performed for the tritiated species, thereby reducing the total uncertainty to the 10^{-5} cm^{-1} level, limited by higher-order $E^{(5)}$ and $E^{(6)}$ terms. However, the current theoretical uncertainty of the NAPT-framework is sufficient for a comparison with experimental values, with uncertainties on the order of a few 10^{-4} cm^{-1} for the tritiated molecular hydrogen species.

4 Lineshape analysis

For most lines observed at good signal-to-noise ratios presented in this study, the lineshapes were found to be symmetric. These resonances were fitted with Voigt profiles, dominated by a Gaussian component with Doppler and instrumental contributions. At sufficiently high intensities for strong transitions, saturation dips are observed and were in fact exploited for T_2 spectroscopy in ref. 45 for the ac-Stark extrapolation. We discuss this saturation effect in this section and its consequence for extracting the line position from the profile. For the weaker transitions such as those in the S-branch of T_2 , some asymmetry in the lineshape is observed. We discuss several contributions that result in the asymmetric profiles including the ac-Stark effect, non-resonant background and interference effects occurring in the non-linear optical CARS process.

4.1 Saturation CARS spectroscopy

The accuracy of the current CARS measurements is mainly limited by the Doppler-broadened linewidth. In view of testing future improvements of QED-calculations beyond the NAPT-framework experimental uncertainties at the level of $< 10^{-5} \text{ cm}^{-1}$ would be desired. Such reduced measurement uncertainties might be achievable *via* saturated CARS spectroscopy, where a narrow

Fig. 7 Comparison of the experimental and calculated fundamental vibrational splittings of all tritiated species. The error bars on the data points indicate the measurement uncertainty while the (yellow) shaded regions indicate the uncertainty of the calculations. The rightmost trace shows results of CARS spectra for D_2 , measured in a molecular beam expansion (red triangle), from a quasi-cw CARS measurement (green square), and measurements with pulsed pump and Stokes laser beams (blue circle) as used for the tritiated species. The D_2 points are slightly displaced horizontally for clarity.

Doppler-free Lamb dip is detected. Saturated CARS profiles were explored by Lucht and Farrow⁷⁰ who proposed a theoretical description *via* a density matrix formalism. In that study, a mechanism for the saturation effect was invoked either *via* strong population pumping or by spectral interference between different velocity classes. The population hole-burning effect in saturation spectroscopy gives rise to typical Lamb dips at the resonance positions. The spectral interference effect, discussed in detail below, results in the cancellation of the real part of the third order susceptibility of a particular velocity class $\vec{u}$ with that of the opposite direction $-\vec{u}$. Such cancellation is also most pronounced at the resonance position, which then also contributes to saturation of the CARS signal. In a different approach Owyong and Esherick⁷¹ have demonstrated saturation in stimulated Raman loss spectroscopy of D₂ in a fashion of typical pump-probe spectroscopies, using a three and four-beam counterpropagating configuration. The observed saturation dip exhibited a width of 110 MHz, limited by laser bandwidth.⁷¹ This suggests that with a narrower laser bandwidth and improved frequency calibration procedures, sub-MHz accuracies could potentially be achieved for tritiated species.

We performed systematic studies of saturation CARS on the D₂ Q(2) and T₂ Q(1) transitions. The D₂ Q(2) measurements were performed using a gas cell containing 4 mbar of pure D₂. A clear saturation feature appears when the pump and Stokes intensity product is $I_P I_S \sim 5 \text{ GW}^2 \text{ cm}^{-4}$. Fig. 8 shows recordings at various pump pulse energies. As shown, the width of the Lamb dip in D₂ is about 300 MHz, while the Doppler-limited profile is more than 1.2 GHz wide, with width contributions from the instrument function and power broadening. The spectra were fitted with two Gaussian function components, with opposite amplitude signs to represent the saturation Lamb dip and the Doppler-broadened profile. It turned out that the linear ac-Stark extrapolation of the Lamb dip feature and that of the Doppler-broadened envelope do not produce consistent values for the zero-field transition frequency. The Lamb-dip positions plotted against intensity shows a slope that is a factor of five smaller than expected from the unsaturated Doppler-broadened profile.

We suspect that this stems from (1) the spatial intensity distribution in the sample volume, which is difficult to integrate over, especially for the Lamb dips exhibiting a different nonlinear response to intensity than that of the Doppler-broadened envelope; and (2) the occurrence of CARS cross-interference effect⁷² on saturation.

Saturated spectra of the T₂ Q(1) transition, recorded at similar alignment and intensity conditions as for D₂ Q(2), are shown in Fig. 8. The T₂ Q(1) saturation dip is observed to be shifted stronger at the higher-frequency flank with respect to the center of the Doppler-broadened profile at increasing pump energy. The resulting asymmetric profile appears to be qualitatively different from the case of D₂, where the Lamb dip center coincides with the Doppler envelope center at lower intensities. Trivikram *et al.*⁷³ studied a related phenomenon on the electronic transitions of H₂, where the line shift no longer follows a linear trend at sufficiently high intensities as the transitions exhibit skewed line profiles. This presents a serious difficulty in the ac-Stark extrapolation, as high intensity measurements cannot be used in a linear extrapolation to the ac-Stark free value. The intensity values reported in the previous T₂ study⁴⁵ was probably overestimated thus staying in the linear regime for the ac-Stark shift. In the present study, the line positions of the T₂ Lamb dip positions in Fig. 8(d)–(f) already showed a much smaller power-dependence slope compared to measurements for unsaturated profiles at lower intensity.

A more accurate treatment of these phenomena needs to address the asymmetric ac-Stark broadening due to spatial and temporal intensity variations, such as those discussed in ref. 73 and 74, which is not straightforward in the case of CARS. In principle the Lamb-dips should produce more accurate results in determining transition frequencies, however, more refined studies are required to obtain a quantitative understanding of the saturation effect in CARS and to extract the most accurate line positions from the narrower resonance features.

4.2 Asymmetric profile of weak transitions

The CARS measurements were extended to transitions involving higher rotational states up to $J = 7$ for T₂ and DT and also

Fig. 8 Observed spectral profiles for CARS signals exhibiting saturation Lamb dips for two lines recorded at different pump pulse energies. Specific conditions: D₂ Q(2) with Stokes pulse energy 0.74 mJ and pump energy: (a) 0.58 mJ; (b) 1.2 mJ; (c) 1.7 mJ; and T₂ Q(1) with Stokes pulse energy 1 mJ and pump energy: (d) 1.5 mJ; (e) 1.75 mJ; (f) 2 mJ.

to the weaker S-branch of T_2 , with $\Delta J = +2$. The lineshapes for these weaker lines were observed to give rise to asymmetric profiles. These effects are partly caused by the required high intensity in the measurements to compensate for the weakness of transition probabilities, either from having a smaller Raman cross-section or a lower population. The weak transitions are barely detectable with 1 mJ pump and Stokes energy and show different degrees of asymmetry in their spectral profile. Nearly all these weak transitions show different tailing behavior in the flanks below and above the resonance and, despite the higher pulse energy used, no saturation effect at resonance is apparent. As a consequence the accuracy for which the $Q(J = 6, 7)$ transition frequencies in T_2 and DT, as well as T_2 $S(J = 0-3)$ lines can be determined is much lower than for the strong CARS lines (the transition frequencies of these weak lines are included Table 2). We discuss three possible effects contributing to the asymmetry: (1) spatial beam profile and time evolution of pulses; (2) ac-Stark splitting of the magnetic sublevels, and; (3) resonant cross interference between lines and interference with the non-resonant background.⁵⁰

The first two effects were discussed by Moosmüller *et al.*,⁵⁹ studying the spectral profile from coherent Stokes Raman scattering (CSRS) of N_2 exhibiting the effects of spatial and temporal variation of intensity. Similarly as for the CARS profiles investigated here, molecules at different positions in the interaction volume and throughout the time evolution of tightly focused pump and Stokes pulses will have varying contributions to the overall spectrum, because of the spatial and temporal variation of the intensities. It is expected that signal contributions near the focal point, which is highly red-shifted by ac-Stark effect in this case, are the strongest.

Apart from the ac-Stark shift due to the spatial and temporal intensity distribution, the ac-Stark induced splitting of the m_J sublevels, which are unresolved in the measurements, will lead to an asymmetry of the spectral profile. From eqn (3), the second term refers to the ac-Stark splitting for different m_J , where higher $|m_J|$ values experience less shift and overall produce a broadening of the spectrum. Such phenomena of ac-Stark splittings in CARS were quantitatively investigated previously.^{60,75} Here we address such effects phenomenologically in the line fitting of asymmetric lines (see below).

Another well-known artifact in CARS spectra is the spectral interference in the third order susceptibility $\chi^{(3)}$. The main contribution to the third order non-resonant susceptibility at 532 nm for the pump and 683 nm for the Stokes wavelengths is from the electronic polarizability, with only a smaller contribution due to molecular vibration and rotation.⁷⁶ In eqn (1), the third-order susceptibility $\chi^{(3)} = \chi_{NR} + \chi_R$ comprises of a resonant and non-resonant part giving rise to interference effects in the CARS signal, $I_{AS} \propto |\chi_{NR} + \chi_R|^2$ (where the superscript⁽³⁾ is dropped). The real and positive χ_{NR} , from far off-resonance contributions, will interfere with the real part of χ_R and distort the spectrum resulting in a dispersive-like signal.

In the following, we first discuss a numerical study to simulate the effects of the spectral interference effect on the lineshape that eventually shifts the peak position of the

resulting profiles. Such shifts are included in the frequency determinations as listed in Table 2. Thereafter, a fitting procedure is described using an asymmetric profile applied to the experimental spectra to assess shifts in the peak position. In the latter, it remains difficult to decouple the separate contributions of the ac-Stark effect and spectral interference, giving at best an estimate of possible systematic frequency shifts due to the asymmetric profile. For these cases the uncertainties in the frequency values as listed in Table 2 were appropriately enlarged.

4.3 Lineshape simulation for interference effects

Following the analyses in ref. 77 and 78 simulations were performed to account for the interference effects in the CARS lineshape that is proportional to the square of the third-order nonlinear susceptibility $|\chi|^2$:

$$|\chi|^2 = \left| \chi_{NR} + \sum_J a_0(J) \int_{-\infty}^{\infty} \frac{\frac{1}{\sqrt{\pi}u} e^{-v_z^2/u^2} dv_z}{\omega_0(J) - (\omega_P - \omega_S) - i\Gamma - \frac{1}{2\pi}(k_P - k_S)v_z} \right|^2, \quad (4)$$

where $\omega_0(J)$ is the resonance frequency of a particular transition, Γ is the relaxation rate, $(k_P - k_S)v_z/(2\pi c)$ is the Doppler shift, and u is the most probable speed $\sqrt{2k_B T/M}$ for temperature T . The integration over the Maxwell-Boltzmann distribution for all velocities v_z along the CARS beam propagation direction can be performed and the equation recast using a Faddeeva function $w(\zeta)$

$$|\chi|^2 = \left| \chi_{NR} - i \sum_J a_0(J) \frac{\sqrt{\pi}c}{\omega_R u} w\left(\frac{-\omega_0 + \omega_R + i\Gamma}{\omega_R \frac{u}{c}}\right) \right|^2 \quad (5)$$

where $\omega_R = (\omega_P - \omega_S)$. The $(v = 1, J)$ states are extremely long-lived, hence the Lorentzian width Γ is set to zero in the present low-pressure and Doppler-limited case. The $a_0(J)$ coefficients for the Q-branch transitions,

$$a_0(J) = \frac{Np(J)}{h} \left(\alpha_0^1(J)^2 + \frac{4}{45} b_J^J \gamma_0^1(J)^2 \right), \quad (6)$$

depend on the population number density $Np(J)$ for the rotational ground states J and b_J^J is the Placzek-Teller coefficient. The isotropic α_0^1 and anisotropic γ_0^1 polarizabilities between $v = 1$ and $v = 0$ are estimated from that of H_2 , HD and D_2 from ref. 79, since no values for the tritiated species are available. For $J = 0$, $\alpha_0^1(0) = 0.61$ a.u. and $\gamma_0^1(0) = 0.50$ a.u. and these values do not vary significantly for other J states.

There are two contributions to the interference effect in CARS. Firstly, the summation over all populated states gives rise to cross-interferences χ_R between all lines in the $v = 0 \rightarrow 1$ fundamental band, and secondly a nonresonant term χ_{NR} that represents the contribution from the far-lying electronic resonances. The χ_{NR} value is estimated from the H_2 value of $6.7 \times 10^{-18} \text{ cm}^3 \text{ erg}^{-1}$ amagat given in ref. 80. For the 2.5 mbar T_2 sample at 298 K, this value converts to $1.5 \times 10^{-20} \text{ cm}^3 \text{ erg}^{-1}$

or 1.5×10^{-20} esu, while for the DT sample with 12.7 mbar total pressure a value of 7.6×10^{-20} esu is estimated. The presence of D_2 in the DT sample results in additional cross-resonance contribution from the D_2 Q-branch (Raman shift at $\sim 2991 \text{ cm}^{-1}$) on the DT Q-branch (Raman shift at $\sim 2743 \text{ cm}^{-1}$) which is estimated to be 1.8×10^{-20} esu.

Fig. 9 illustrates the effect of CARS interference on the T_2 Q(0) line, for which the dominant interference contribution is from the neighboring T_2 Q(1) transition. Since Q(1) is the most intense and the nearest lying resonance, it produces the strongest effect as follows from eqn (5). The real part $\text{Re}[\chi]$ is plotted in Fig. 9b showing the spectral region that spans T_2 Q(0), with the grey horizontal line indicating zero amplitude. The dashed blue dispersive line belongs to Q(0) while the dashed red line below zero represents the contribution of the Q(1) line, yielding the solid blue dispersive line as the sum. At the Q(0) resonance position, the nonresonant contribution χ_{NR} is more than a hundred times smaller than the resonant χ_R contribution from Q(1). It is noted that this holds for the present specific case, given the rather low pressures in the sample yielding a small χ_R . The square of the real part of χ is plotted in Fig. 9c, with the dashed curve indicating the pure contribution from Q(0) and the solid curve from the summation. Fig. 9d, shows both contributions of the real $\text{Re}[\chi]$ and imaginary parts $\text{Im}[\chi]$ of the third-order susceptibility as well as the square $|\chi|^2$. Finally, Fig. 9e shows the line profile of the T_2 Q(0) line. The solid line includes the cross-interference effect, while the dashed curve represents the case when the effect of interference is neglected. This numerical example demonstrates how the large CARS susceptibility contribution from T_2 Q(1), due to the factor of 7 higher population in $J = 1$ compared to that of $J = 0$ (under room-temperature conditions), results in a large interference with the nearby Q(0) transition leading to a +13 MHz shift in Q(0). The positive sign of the shift in T_2 Q(0) is due to the negative contribution of the real part of the χ susceptibility from the Q(1) transition (see Fig. 9b).

Systematic shifts of the line centers due to spectral interference were simulated for all T_2 Q-branch transitions as a function of temperature, and results are shown in Fig. 10. The Q(0) line is blue-shifted due to cross-interference with other Q($J \neq 0$) lines. For increasing temperature there are two effects that enlarge the cross-interference shift on the Q(0) line: the population ratio between $J = 1$ and $J = 0$ grows and approaches 9:1 while also the larger Doppler width leads to a stronger interference experienced by Q(0). The sharp drop in the peak position offset of the Q(4) and Q(5) lines below 200 K is due to the drastic decrease in population of the $J = 4, 5$ states. At around 300 K where the measurements were performed, systematic shifts of the line centers for the T_2 Q($J = 1-5$) lines were found to be less than 3 MHz.

The Doppler-limited spectra for the DT lines were also simulated and depicted in Fig. 11. For the strong DT transitions Q($J = 0-5$), the asymmetry in the line profiles is not apparent, and the interference-induced shifts in the line positions are found to be below 3 MHz for Q($J = 0-4$), while it is -8 MHz for Q(5). However, for the weaker Q(6) and Q(7) lines, the interference effect is more visible as shown in the inset of Fig. 11. Based on eqn (5) and (6) the effects of the cross-interferences between the lines as well as the effects of the interference of the non-resonant background are quantitatively evaluated. A shift of -24 MHz is observed for Q(6) and a much higher shift of -83 MHz for the weaker Q(7) line. It is worth noting that for the DT Q(6) and Q(7) lines, the non-resonant susceptibility contribution χ_{NR} is an important contribution that is comparable to the cross-interference contributions, as shown in the insets of Fig. 11. The χ_{NR} for the DT sample is higher than in the case of T_2 mainly caused by the higher total pressure in the DT gas cell. Similar simulations were also performed on the HT Q($J = 0-3$) lines, with the resulting shifts in peak positions due to CARS interference were found to be less than 3 MHz. The line positions reported in Table 2 include corrections to these estimated CARS interference effects.

Fig. 9 Simulation of the CARS spectrum of the Q(0) line of T_2 , focusing on the effect of CARS spectral cross-interference. The real and imaginary contributions of the Q(0) line are drawn in subfigures (a)–(d). See text for extensive discussion. (e) The line profile of the Q(0) line with the cross-interference effect is shown as the solid curve, while the dashed curve represents the case when the effect of interference is neglected. The cross-interference causes a shift of +13 MHz of the Q(0) peak position.

Fig. 10 Simulated shifts of peak positions of the T_2 Q-branch transitions due to spectral cross-interference as a function of temperature.

Fig. 11 Simulation of CARS spectra of DT displaying relative intensities for Q(0)–Q(7) lines at room temperature. Zoomed-in spectra of Q(6) and Q(7) are displayed showing the resonant and non-resonant interference effects on these weak lines; (a) the blue line represents a symmetric line profile without considering other contributions in the susceptibility; (b) the red line represents the cross-interference from nearby line contributions in the fundamental band Q-branch susceptibility; (c) the black line represents the inclusion of both interference effects: estimated electronic nonresonant susceptibility χ_{NR} and off-resonance fundamental band Q-branch susceptibility. The shift in peak position of Q(6) and Q(7) from (a) and (c) is about -24 MHz and -83 MHz, respectively.

4.4 Lineshape fitting

A phenomenological approach is adopted in fitting asymmetric profiles to experimental data from the combined effects of the ac-Stark and spectral interference phenomena, especially for the weak transitions. While a full quantitative assessment that would separate the contributions of these effects is not possible at present, the lineshape study enables us to better estimate the contribution to the systematic uncertainty resulting from the line profile asymmetry.

For the strong transitions with symmetric line profiles such as Q transitions from $J = 0-5$ in T_2 , DT and HT, a symmetric Gaussian or Voigt fitting function is used. However, despite the T_2 Q(0) and the DT Q(6) lines appearing to be symmetric, the simulations suggest that interference effects may potentially shift the line center. To estimate a systematic uncertainty contribution due to a choice of the line profile in the fitting

Fig. 12 Comparison of lineshape fitting of the DT Q(6) transition using the asymmetric profile of eqn (7) (red) and a symmetric Gaussian function (blue). Fitting with the Faddeeva profile, as in eqn (7), gives a better fit especially in the wing of the spectrum. The difference of central frequency between the two fitting functions is 20 MHz.

of the T_2 Q(0) and DT Q(6) lines, a Faddeeva function $w(\zeta)$ is employed:

$$I_{AS} = y_0 + \left| \chi_{NR} - i a_0 w \left(\frac{-\omega_0 + \omega_R + i\Gamma}{\Delta\omega} \right) \right|^2, \quad (7)$$

where $\Delta\omega$ is the Gaussian width and Γ is the Lorentzian width, while y_0 sets the experimental baseline. For T_2 Q(0) and DT Q(6), a 16 MHz discrepancy is observed from the fitting result between a simple symmetric Gaussian profile and the Faddeeva profile in eqn (7). This systematic offset is included in the uncertainty budget of these two lines as listed in Table 2.

For other weak lines such as the T_2 Q(6) and S(1) transitions shown in Fig. 13, the poor SNR makes it more difficult to separate the effects of the dispersive feature due to spectral interference and the ac-Stark splitting and broadening, leading to larger uncertainties in the line position determination. Note that when compared to the DT Q(6) line in Fig. 12, the analogous T_2 Q(6) line in Fig. 13 exhibits a more pronounced asymmetry. This is due to the *ortho-para* spin statistics leading to a lower population for $J = 6$ in homonuclear T_2 species, which is not present in DT. On the other hand, despite the T_2 S(1) line being of comparable strength to T_2 Q(6) (Fig. 13), the S(1) line at 2657 cm^{-1} exhibits a more symmetric profile. This can be explained from competing contributions of opposite signs: the cross-interference χ_R from the T_2 Q-branch at 2464 cm^{-1} having a negative sign and the non-resonance χ_{NR} background having a positive sign.

In addition to the cross-interference induced shifts in the peak positions (included in Table 2), the transition frequencies of the weaker lines are corrected for the ac-Stark shift, using the slope of low-intensity ac-Stark measurements in stronger transitions (e.g. Q(1) transition of the respective species). However, the dominant contribution to the uncertainties of $5 \times 10^{-3} \text{ cm}^{-1}$ for the weak lines derives from the large fitting

Fig. 13 Line fitting of the T_2 Q(6) line and the weaker S(1) transition, both with pronounced asymmetries, using a symmetric Gaussian fit (solid – blue) and asymmetric Faddeeva profile (dashed – red). Below the spectra the residuals of different fitting functions are plotted.

uncertainty in the peak positions given the significant ac-Stark broadening.

5 Conclusion and outlook

In summary, we have accurately determined transition frequencies of the fundamental vibrational band ($\nu = 0 \rightarrow 1$) of HT, DT and T_2 . For most Q-branch transitions with ($J = 0-5$), uncertainties below 0.0005 cm^{-1} are achieved, which represent more than a hundred-fold improvement over previous studies. For the weak Q($J = 6, 7$) transitions in T_2 and DT and the S-branch of T_2 , the high pulse intensities required for the measurements cause significant ac-Stark broadening. The transition frequencies of these weak lines are thus reported with a much higher uncertainty of 0.005 cm^{-1} . All measured transition energies are in very good agreement with the latest *ab initio* calculations based on NAPT theoretical framework.^{25,69} This work demonstrates that measurements on tritiated species can be achieved at high accuracy, to within a factor of three compared to the accuracy for the non-radioactive species. This doubles the number of benchmark candidates for testing the most accurate quantum chemical calculations of the energy structure of molecular hydrogen.

Due to interference effects with the non-resonant background in CARS spectra, the resulting asymmetric line profiles in weak transitions lead to a higher uncertainty in the line center determination. Cross-interference between resonances can be a limiting factor for the spectroscopic accuracy of CARS

measurements; it is shown here that this phenomenon can be quantitatively approached and corrections to transition frequencies calculated.

Future improvements in experiments on tritium-bearing molecular hydrogen would require sub-Doppler techniques, to enable improved comparison with ever advancing calculations. This could be achieved by exploiting Lamb dip spectroscopy with hybrid pulsed-cw CARS, using a pulsed pump beam and a continuous wave Stokes beam, as demonstrated here for D_2 . From these test studies it follows that an improved understanding of line shape profiles, in particular of the ac-Stark effect on Lamb-dips, is required to effectively achieve higher accuracy on the transition frequencies. Improvements might be achieved by using higher cw-Stokes powers, allowing measurements to be performed at even lower pump pulse energies, thereby reducing ac-Stark effects induced by the pump laser. With the narrower cw laser bandwidth and improved frequency calibration procedures, sub-MHz accuracies could potentially be achieved. Ultimately the application of novel cavity-enhanced cw-spectroscopies such as the NICE-OHMS technique on the heteronuclear species HT and DT, with the possibility of Doppler-free saturated absorption, may result in accuracy improvements by orders of magnitude,¹⁶ provided that those techniques can be combined with safe handling of radioactive species.

Conflicts of interest

There are no conflicts to declare.

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