Dual-comb spectroscopy for high-temperature reaction kinetics

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Received 16 October 2019, revised 6 January 2020
Accepted for publication 22 January 2020
Published 5 March 2020

Abstract

In the current study, a quantum-cascade-laser-based dual-comb spectrometer (DCS) was used to paint a detailed picture of a 1.0 ms high-temperature reaction between propyne and oxygen. The DCS interfaced with a shock tube to provide pre-ignition conditions of 1225 K, 2.8 atm, and 2% p-C3H4/18% O2/Ar. The spectrometer consisted of two free-running, non-stabilized frequency combs each emitting at 179 wavelengths between 1174 and 1233 cm⁻¹. A free spectral range, \( f_r \), of 9.86 GHz and a difference in comb spacing, \( \Delta f_r \), of 5 MHz, enabled a theoretical time resolution of 0.2 μs but the data was time-integrated to 4 μs to improve SNR. The accuracy of the spectrometer was monitored using a suite of independent laser diagnostics and good agreement observed. Key challenges remain in the fitting of available high-temperature spectroscopic models to the observed spectra of a post-ignition environment.

Supplementary material for this article is available online

Keywords: propyne, dual-comb spectroscopy, reaction kinetics, time-resolved, laser absorption spectroscopy, shock tubes, high temperature

(Some figures may appear in colour only in the online journal)

1. Introduction

Energy systems involving high temperatures and fast reactions rely on absorption spectroscopy for fundamental research and field diagnostics. However, measurements at extreme conditions are fundamentally challenging due to short timescales and complex thermochemistry that are often present. For such environments, high-bandwidth measurement of a spectral surface (absorbance as a function of wavenumber and time) could provide sufficient information for speciation, thermometry, and knowledge of quantum state populations in non-equilibrium environments. However, many broadband measurement techniques, such as Fourier transform infrared spectrometers or rapid-tuning broad-scan external cavity quantum cascade lasers [1], are mechanically limited to low time-resolutions. Advanced broadband diagnostics being developed without such mechanical limitations involve super-continuum and frequency-comb absorption spectroscopy [2–9].

Two frequency combs can be combined to help interpret the signal of multiple wavelengths simultaneously, a technique called dual-comb spectroscopy (DCS). Notable studies involving DCS include: the first dual-comb field diagnostic monitoring greenhouse gases [2], dual-combs in a rapid compression machine [8], and DCS used to study protein dynamics [10].

It is of great interest to apply DCS to study high-temperature reaction kinetics. However, this application puts stringent constraints on a spectrometer’s capabilities. At high-temperatures, reactions occur over short timescales and often in vessels with short path lengths. Correspondingly, it is important to have microsecond time-resolution and a high power across all wavelengths (power-per-wavelength) to ensure sufficient signal to noise in a dynamic environment with
emission and vibration. Furthermore, it is desirable for measurements to be made in the mid-infrared between 3–20 μm with access to the prominent absorption features of hydrocarbons, such as CH$_2$ and CH$_3$ stretching modes in the 3 μm region, –CH = CH$_2$ bending modes near 11 μm, or CH bending modes near 15 μm [11]. To the authors’ knowledge, reaction kinetics using frequency-comb spectroscopy was first studied by Fleisher et al. and Bjork et al. involving measurements of a photolysis-induced reaction at 25 μs time resolution [12, 13]. These studies did not use DCS but used a single frequency comb and diffraction grating to interpret the signal. The use of two frequency combs, dual-comb spectroscopy, has been applied to high-temperature, non-reacting conditions in a rapid compression machine by Draper et al. at 704 μs time-resolution and also by Schroeder et al. in the exhaust of a gas turbine at high temperatures and 10–60 s time-resolution [4, 8]. Draper et al. and Schroeder et al. achieved exceptional spectral resolution with tens of thousands of wavelengths in the near-IR. A worthy extension of the work of Draper and Schroeder et al. would be to establish similar techniques in the mid-IR with improved time-resolution.

Efforts to extend DCS into the 3–20 μm region have involved distributed feedback generation DFG [14], optical parametric oscillators (OPO) [15–17], and quantum-cascade-laser (QCL) approaches [6, 7, 18–21]. Obtaining coherency between two frequency combs is an enabling factor for DCS and DFG approaches can achieve this readily. However, DFG methods generally have limited power-per-wavelength, which limits application in many environments [19]. Alternatively, OPO-based combs can obtain a higher power-per-wavelength while also providing access to the mid-infrared. However, a disadvantage of OPO combs involves the fact that coherency between separate combs is more challenging to obtain. For this reason, dispersion was used when Fleisher et al. and Bjork et al. first studied reaction kinetics with a mid-infrared OPO comb [12, 13]. An extension of the work of Fleisher and Bjork et al. would involve the use of two coherent, mid-IR frequency combs configured as a dual-comb spectrometer.

Prominently, a class of OPO combs operating in degeneracy, known also as divide-by-two subharmonic OPOs, have successfully demonstrated coherency and dual-comb operation [17, 22, 23]. The OPO-based DCS of Muraviev et al. [17] offered remarkable spectral coverage (3.1–5.5 μm with 350 000 spectral data points), although with a minimum time-resolution of only 7 ms. While only a matter of time, OPO-based DCS has yet to demonstrate both dual-comb and microsecond operation in the mid-infrared.

Advances in the field of quantum-cascade-laser (QCL) frequency combs have led to the development of mid-IR, solid-state spectrometers that have attracted considerable attention in the recent literature [7, 10, 18, 20, 24–27]. In particular, the recent advent of QCL frequency combs provides an alternative to DFG and OPO combs when a high power-per-wavelength and fast time resolution are needed [18, 20, 21]. QCL combs can offer over 1 mW per wavelength and microsecond time-resolution, with the tradeoff of having a larger comb spacing (~0.3 cm$^{-1}$) and less spectral range than alternative systems. To date, these QCL-based systems have performed well in proof-of-concept demonstrations, e.g. to observe protein dynamics and to study the spectrum of Bromomethane and Freon 134a [10, 20].

In this study, we demonstrate a mid-infrared QCL-based dual-comb spectrometer (DCS) capable of microsecond-resolved measurements of energetic gas phase reactions. Specifically, an aggressive chemical reaction, propyne oxidation, was studied using a pressure-driven shock tube equipped with two QCL frequency combs and a suite of independent validation diagnostics. This study details the dual-comb spectrometer employed, provides a demonstrative propyne oxidation measurement, and concludes with an assessment of system accuracy.

2. Background

2.1. Laser systems

An overview of the experimental setup is provided in figure 1, illustrating the placement of frequency combs and other validation diagnostics on a shock tube with a bore of 14.13 cm. The shock tube was used to generate the high-temperature conditions and was described by Strand et al. [1] previously. The shock tube interfaced with a dual-comb (QCL-based) spectrometer, and two additional interband cascade lasers (ICL) to study water and propyne. A side-wall pressure transducer was also used to monitor the pressure during the reaction.

2.1.1. Dual-comb spectrometer. The DCS developed for the current study relies on two free-running QCL-based frequency combs (from IRsweep AG), each spanning about 60 cm$^{-1}$. These frequency combs rely on the principles of four-wave-mixing (FWM) and are Fabry–Pérot-based QCL combs. The native emission power of each comb was >500 mW resulting in >2 mW per comb tooth on average. Each QCL comb was slightly detuned from another to yield two different repetition frequencies, $f_1$ and $f_2$ that were both at approximately 0.3 cm$^{-1}$ or 9.86 GHz. The difference in repetition rates,
\[ \Delta f_{\text{rep}} = f_1 - f_2, \]  was typically around 5 MHz. This slight difference resulted in the creation of a set of beating frequencies between each detuned pair of spectral lines. This set of beat-notes constitutes an additional frequency comb in the radio-frequency (RF) domain. Unlike the mid-infrared combs that generated the beat-notes, this RF comb is at much slower frequencies and its intensity vs. time can be measured on a detector.

The combs emitted at 179 different wavelengths from 1174 to 1233 cm\(^{-1}\) providing access to high-temperature absorption features for alkynes and water in this region [28, 29]. FTIR measurements of the corresponding emission from each comb are presented in figure 2. The narrow emission of each comb tooth is below the 0.3 cm\(^{-1}\) spectral resolution of the FTIR and this under-sampling manifests itself as the broadened peaks seen in figure 2. However, the FTIR measurements in figure 2 very effectively illustrate the emission range of the QCL’s and the spectral power distribution that can be expected from the spectrometer.

The emission from the pair of QCL combs was coaligned and split to yield two combined beams. The first (reference) beam was attenuated and focused onto a reference HgCdTe (MCT) detector, while the second (sample) beam was attenuated and transmitted through wedged, anti-reflection-coated ZnSe windows in the shock tube. After traversing the shock tube, this beam was directed through a long-pass filter and focused on another MCT detector. The superimposed wavelengths of light from the two frequency combs heterodyned on each detector and generated a comb in the RF domain spaced at \( \Delta f_{\text{rep}} \). The detectors had a bandwidth of 1 GHz and were digitally sampled at 2.0 GS s\(^{-1}\) to record a sequence of interferograms spaced at a period of \( 1/\Delta f_{\text{rep}} \) (0.2 μs). The shortest achievable time resolution of this configuration is determined by the period of a single interferogram and was therefore 0.2 μs. However, the interferograms were then averaged over a time-interval of 4 μs to improve signal-to-noise ratio. In the time domain, a representative signal from the DCS is shown in figure 3, which demonstrates a typical interferogram pattern observed.

![Figure 2](image1.png)

**Figure 2.** FTIR measurements of each frequency comb. While emission of each comb line is well below the spectral resolution of the FTIR (0.3 cm\(^{-1}\)), these FTIR measurements provide a clear picture of the wavenumber-dependent power distribution of the system with high power near 1185 and 1220 cm\(^{-1}\) and low power near 1205 cm\(^{-1}\).

\[ \text{SNR is improved further through time-integration to 4 μs.} \]

Figure 4 presents the result of a Fourier transformation of the time-domain data to assess the frequency content of the DCS signal. As can be seen in figure 4, the slight detuning between the two frequency combs manifests itself as an additional frequency comb in the RF domain. The beating, or heterodyning, between DCS comb teeth creates the multi-heterodyne pattern in figure 4. The discrete and equally spaced peaks confirm that the system is generating a successful multi-heterodyne signal and provides insights about the QCL combs that generated the pattern. Prominently, the RF frequency comb in figure 4 has a similar shape as the power profiles from the FTIR measurements presented in figure 2.

The power in each RF beat-note is proportional to the product of the electric fields of the two contributing laser modes. Consequently, changes to the intensities of the measured beat-notes directly correlates to attenuation of light in the optical domain. To calculate absorbance using the DCS system, the attenuation at each optical wavelength was calculated, adjusted by a pre-experimental background measurement, and normalized by an unattenuated signal. The absorbance of the mixture at each wavelength was then determined using the Beer–Lambert relation.

Notably, the spectral separation of the superimposed laser lines contributing to an individual beat-note varied from 50 to 900 MHz; however, an average frequency for each superimposed pair of comb teeth was used for absorbance. Accordingly, the optical resolution of each emission line was limited to 900 MHz (0.03 cm\(^{-1}\)), and overall spectroscopic sampling.
resolution governed by the comb spacing of 0.3 cm\(^{-1}\). To ensure quality measurements, the system was calibrated before and after each experiment against the well-known spectrum of polyethylene. Ultimately, while no active wavelength stabilization method was used during an experiment, a comparison of the pre- and post-shock calibration indicated drift of <100 kHz. Recent studies have indicated that it is possible to reduce this drift to Hz-level performance through full-phase stabilization of QCL-based dual-combs [26]. While full-phase stabilization not employed in the current study, such advancements are very promising.

2.1.2. Supporting laser diagnostics. In addition to the DCS, two separate laser diagnostic systems, shown in figure 1, were employed simultaneously to monitor the reaction. A scanned-wavelength diagnostic for water was used to measure a water absorption transition at 4029.5 cm\(^{-1}\) using a wavelength-tuned interband cascade laser (ICL) with a scan repetition rate of 5 kHz [30, 31]. Secondly, a fixed-wavelength laser diagnostic for propyne detection was used at 2999.55 cm\(^{-1}\) using an additional ICL. Each ICL was coupled with a HgCdTe (MCT) detector that was sampled at 5 MHz.

2.2. Propyne oxidation

Propyne oxidation was selected as the reaction of interest due to a blended broadband absorption feature of propyne and wealth of narrow absorption transitions of water between 1174–1233 cm\(^{-1}\). The absorption feature of propyne is derived from the P-branch of the second harmonic of propyne’s \(-\text{CH}\) in-plane vibrational bending mode [11, 28]. At high temperatures, nearly the entire P-branch of the propyne absorption feature is accessible using the DCS system. The absorption features of water are derived from the P-branch of an in-plane symmetric bending mode [11]. During oxidation, the mixture’s spectrum evolved in time as the broadband spectrum of propyne transformed into the finely featured spectrum of water.

Figure 5. DCS data from 1215 to 1225 cm\(^{-1}\) of propyne oxidation (2% p-C\(_3\)H\(_4\), T\(_0\) = 1225 K, P\(_0\) = 2.8 atm) illustrating the arrival of the incident and reflected shock before time-zero. The broadband absorption feature of propyne is visible from 0 to 0.6 ms and can be seen transforming into a finely featured spectrum (water) after 0.8 ms. The DCS data demonstrates good SNR during the passage of each shock wave, occurrence of a combustion reaction, and temperature/pressure increase of roughly 2500 K/60x over only 1 ms.

Figure 6. DCS measurements of propyne oxidation with time-zero conditions of 2% p-C\(_3\)H\(_4\)/18% O\(_2\) in Ar, 1225 K, and 2.8 atm. All data are aligned in time and share the same x-axis. (a) Measured and simulated thermodynamic conditions for the reaction. (b) Dual-comb-measured absorbance spectrum evolving in time at 4 μs time resolution (see Visualization 1). Two regions with high SNR are delineated in dashed white areas. All simulations shown in (a) used the USC Mech II kinetic mechanism and assumed constant volume [29].

3. Experimental results

3.1. Overview

Data from all diagnostics are presented in figures 5–7 for an example shock tube experiment. The pressure-driven, stainless steel shock tube shown in figure 1 was used to propagate a shock wave at Mach 2.2 into a test gas mixture with an initial molar composition of 2% p-C\(_3\)H\(_4\)/18% O\(_2\)/80% Ar, a temperature of 298 K, and a pressure of 75 Torr. Figure 5
Figure 7. DCS measurements of propyne oxidation with time-zero conditions of 2% p-C₃H₄/18% O₂ in Ar, 1225 K, and 2.8 atm. Speciation measurements at a 95% confidence interval and simulated results for propyne and water. All simulations used the USC Mech II kinetic mechanism and assumed constant volume [29]. Agreement at a 95% confidence interval is observed between the DCS measurements and supporting laser measurements. While overall agreement exists between the DCS measurements and USC Mech II, key differences are visible at early times and just after ignition.

presents a sample of the DCS data from this reaction where the broadband propyne absorption feature can be seen at early times and the finely featured spectrum of water observed after 0.75 ms. The full dataset of the DCS absorbance measurements is presented in figure 6(b). The lowest pressure and absorbance measurements in figures 5 and 6(a)–(b), located before −75 μs, indicate this pre-shock region. Upon the arrival of the first shock wave, there is a step change in pressure and absorbance of the test gas, seen in figures 5 and 6(a)–(b) at −75 μs, as the mixture is heated and pressurized to 720 K and 0.7 atm. At this time, the tail of the propyne absorption feature first appears in figures 5 and 6(b) near 1225–1230 cm⁻¹ and the apex of the feature rests just beyond the domain limit of 1233 cm⁻¹.

After 75 μs, the reflected shock wave arrives (defining time-zero) and provides another pressure increase that is visible in figure 6(a). At time-zero, chemistry is assumed to be momentarily frozen and the mixture condition changed to 1225 K and 2.8 atm. Also at time-zero, the system undergoes a rapid redistribution of the state populations, which shifts the crest of the propyne feature seen in figure 6(b) to approximately 1220 cm⁻¹. Soon after, oxidative pyrolysis begins and propyne is slowly consumed for the subsequent 0.75 ms. During the pyrolysis process, propyne was observed on the separate propyne-specific diagnostic and showed agreement to the overall decomposition rate registered by the DCS, shown on the left of figure 7.

After a residence time of approximately 0.75–0.9 ms, exponential growth of the radical pool takes hold and the mixture ignites. Formation of water during ignition was observed by both the DCS and the scanned-wavelength water diagnostic. In figure 6(b), water formation is represented by a brush of fine absorption lines just before 0.9 ms. These absorption features arise coincident with H₂O formation observed using the ancillary water diagnostic, which is shown on the right of figure 7.

The current study also includes a video in order to demonstrate the time evolution of the measured spectrum, pressure, and water mole fraction. This video is attached in the supplemental material as Visualization 1.

3.2. Speciation

Quantitative speciation measurements were performed using the DCS and the supporting laser diagnostics and compared to a kinetic model (USC Mech II [32]), all shown in figure 7. For DCS measurement of propyne and water mole fractions, a vectorized Beer–Lambert system was formulated as presented in equation (1). The system included: W a weighting parameter matrix, K a cross-section matrix (cm⁻¹ atm⁻¹), x a mole fraction vector, and b (cm⁻¹ atm⁻¹) a vector of the pressure- and path-length-normalized absorbance at each DCS wavelength as a function of time. The formulation of the Beer–Lambert system is discussed in detail in supporting literature [33]:

Minimize: ∥W(Kx − b)∥₂². (1)

The diagonal of the weighting matrix was set as the wavenumber- and time-dependent SNR of the DCS system (see figure A1). The SNR of each DCS wavenumber was defined as the absorbance at a given time divided by a measurement of the standard deviation at each wavenumber. The standard deviation at each DCS wavenumber was calculated from a statistical analysis of the DCS noise, which was determined by measuring the absorbance of a non-absorbing medium. Propyne and water were the only species included in the cross-section matrix, K. Expected intermediate species that form in trace amounts, but potentially detectable quantities, include: allene, acetylene, methane, ethylene, and formaldehyde. Of these potentially interfering species, only methane and acetylene absorb in this region, although acetylene absorbs very weakly.
Temperature-dependent absorption cross-sections were determined for propyne through a series of additional non-oxidative 2% p-CH$_2$H$_4$/98% Ar experiments ranging from 1100 to 1400 K. Water cross-sections were collected from a series of fuel-lean hydrogen oxidation experiments (2%–4% H$_2$/2%–4% O$_2$ at 1670–2030 K) and methane-oxidation experiments (4.5% CH$_4$/9% O$_2$/Ar near 2900 K) to generate water in the post-ignition environments [32]. These independent experiments provided the absorption cross-sections that were not available through existing experimental or theoretical databases. Cross-section correlations across all 179 wavelengths were assumed to have a linear temperature dependence. Acetylene and methane cross-section measurements were also collected due to their potential to absorb and interfere in this spectral region. From the additional cross-section measurements, acetylene was undetectable, but methane’s cross-section was non-negligible. Equation (1) was solved both with and without methane in order to assess whether it was detectable as an intermediate. Including methane in the absorption cross-section matrix did not yield statistically significant methane measurements but did contribute approximately 5% additional uncertainty to the propyne mole fraction measurements between 0.5 and 1 ms. This additional uncertainty was accounted for in the 95% confidence interval included with the data in figure 7.

Notably, all 179 wavelengths were used in the optimization process. Each wavelength made a valuable contribution to the final mole fraction solution, however, with the specific contribution commensurate to the wavelength’s SNR. By employing this weighted optimization procedure, the mole fractions of propyne and water were found to be in fair agreement with the validation diagnostics and USC Mech. II (see figure 7), with fluctuations in the mole fraction of water roughly correlating to the pressure shown in figure 6(a). In the DCS-based mole fraction measurements at early times (0–0.2 ms), we observe a convex decay profile that appears higher and at odds to the concave predictions of the model and ancillary propyne diagnostic. This difference after time-zero is still being investigated as to the source.

### 3.3. Uncertainty quantification

Uncertainty was propagated for the DCS and the supporting laser diagnostics to assess whether the results agree within a 95% confidence interval. All uncertainties are included with each measurement in figure 7. The single-wavelength propyne diagnostic uncertainty is presented in light gray bars at 0.35 and 0.68 ms, the scanned-wavelength water diagnostic uncertainty as light blue bars at 0.7, 1, and 1.4 ms, and the DCS propyne (black) and DCS water (dark blue) uncertainty visible at all other times where ± bars are indicated.

DCS uncertainty was largely governed by the high-temperature absorption cross-section measurements used for speciation. These measurements were collected through the non-oxygenated propyne experiments and the water-generating experiments described previously. Linear temperature-dependent cross-section correlations were determined at each of the 179 wavelengths along with each correlation’s uncertainty to a 95% confidence interval. Equation (2) was used to compute the cross-section (σ) uncertainty at wavelength λ (u$_{\lambda,\sigma,95\%}$) and involved: the residual squared error for the linear fit at wavelength λ (RSS$_{\lambda}$), sample number (n), and the temperature for each cross-section measurement (T$_i$). This method for uncertainty propagation was described previously in [33] and derived from [34]:

\[
U_{\lambda,\sigma,95\%} = \text{RSS}_\lambda \left( \frac{1}{n} \right) \left( \frac{(T - \frac{1}{n} \sum_{i=1}^{n} T_i)^2}{\sum_{i=1}^{n} T_i^2 - \left( \frac{\sum_{i=1}^{n} T_i}{n} \right)^2} \right) \left( \frac{1}{2} \right).
\]

(2)

Uncertainty associated with the pressure- and path-length-normalized absorbance (u$_{\theta}$) was estimated by propagating error due to path length, pressure, and attenuated and un-attenuated laser intensities through the Beer–Lambert relation. Notably, the attenuated intensity was found to be the dominant contributor. Equation (3) was then used in the determination of the uncertainty in the mole fraction of species i at time t during speciation.

\[
u_i(t) = \sqrt{\sum_{j=1}^{M} \sum_{k=1}^{N} \left( \frac{\partial x_i}{\partial \sigma_{j,k}} \right)^2 u_{\sigma_{j,k}}^2 + \sum_{j=1}^{M} \left( \frac{\partial x_i}{\partial b_j} \right)^2 u_{b_j}^2}.
\]

(3)

The terms (\( \frac{\partial x_i}{\partial \sigma} \), (\( \frac{\partial x_i}{\partial b} \)) are the derivatives of the mole fraction of species $x_i$ with respect to the cross-section correlation, σ, and pressure- and path-length-normalized absorbance, b. Notably, the uncertainties in the final measurements were calculated assuming the potential presence of methane. In these calculations, methane was added to the matrix $K$ from equation (1) to assess how the measurements could be affected with a third absorber present. The propyne and water species time-histories were unaffected by this addition, however, their corresponding uncertainties were affected. The largest influence was to the propyne mole fraction uncertainty, which increased by about 5% in the 0.2 ms proceeding ignition. No statistically significant methane was observed to form and therefore the results with no methane are shown in figure 7 but with the larger uncertainties included. Future studies may also consider numerically perturbing the temperature profile used in the determination of the temperature-dependent cross-sections. However, the uncertainty here was largely dominated by the uncertainty in the absorption cross-section measurements and the absorbance. Therefore, adding these additional terms will yield only minor contributions to the overall uncertainty reported.

This method of uncertainty propagation was also used for the fixed-wavelength propyne diagnostic presented as light gray in figure 7. This diagnostic used only one wavelength, rather than 179 for the DCS. Correspondingly, the uncertainty is much larger for the single-wavelength diagnostic due to a very large sensitivity from the term (\( \frac{\partial x_i}{\partial \theta} \)). This observation confirms the utility of using multiple wavelengths as a means of decreasing this sensitivity and, therefore, the overall uncertainty in mole fraction measurements. In contrast, for the DCS diagnostic, improvements in the reference cross-sections will yield a largest reduction in uncertainty.
Uncertainties associated to the scanned-wavelength measurements were determined using the RSS of the Voigt fit to the absorbance trace recorded and detailed in the literature [31]. In consideration of the uncertainties presented, the DCS and supporting diagnostic measurements of both propyne and water agree within a 95% confidence interval. However, before 0.1 ms and immediately after ignition a subtle but observable disagreement exists between the DCS and the supporting diagnostics that is still under investigation.

4. Time resolution and spectral accuracy

4.1. Time resolution

Figures 8(a) and (b) present wavenumber slices of the DCS measurement at 100 and 4 μs time resolution, respectively, to illustrate the DCS data as a function of the averaging time. Low SNR regions between 1200–1210 cm\(^{-1}\) and beyond 1230 cm\(^{-1}\) exhibit high noise yet contain valuable data even at 4 μs time resolution. The blended-spectrum of the P-branch of the propyne can be seen very clearly at early times for both 100 and 4 μs cases. Likewise, the consumption of propyne and the final spectrum of water can be made out from the last two frames of figures 8(a) and (b). We conclude that the 4 μs resolution data are of sufficient quality to make out the spectrum and its evolution. Although noise increases with shorter time resolution, and there are regions with very low SNR, this data demonstrates the opportunity to DCS to effectively capture reaction kinetics.

4.2. Spectral accuracy

Spectral accuracy of the spectrometer was tested both using an independent laser diagnostic for propyne and two available high-temperature spectroscopic models for water. The spectrum of high-temperature propyne showed good agreement to independent spectral measurements, confirming the spectral accuracy of the system. However, an interesting disagreement was found between the measured spectrum and available spectroscopic models for high-temperature water [29, 35]. At this time, no high-temperature models exist for propyne that can be used for comparison.

4.2.1. Independent spectral measurements | propyne. A rapidly-tunable external-cavity QCL was used to scan over the same portion of the spectral region as the DCS. A full description of the scanned QCL laser methodology is provided in the literature [1]. Non-oxygenated mixtures of 2% p-C\(_3\)H\(_4\)/Ar were used to compare each diagnostic. Experiments using the scanned-QCL were limited to temperatures below 1130 K, where propyne was thermally stable for the 2 ms scan time. A common condition of 1120 K and 3 atm was tested for both systems and the results are shown in figure 9. The DCS measurement in figure 9 was time-averaged over the same 2 ms as the scan from the scanned-QCL. Good agreement is visible between the DCS and the QCL spectra with the only discrepancies in the two measurements corresponding to low SNR regions of the DCS. The scanned-QCL’s spectral resolution is limited to 0.3 cm\(^{-1}\), roughly equal to the DCS point spacing. Accordingly, the scanned-QCL was only used on the broadband spectrum of propyne and spectroscopic models used to investigate the spectrum of water. This rapidly-tunable external-cavity QCL could not be used as a broadband method to study the propyne oxidation reaction because the laser system’s time resolution, of 2 ms, is greater than the time-scale of the 1 ms reaction.

4.2.2. Spectral modeling | water. Additional measurements were taken to study the spectrum of water and compare against available high-temperature spectroscopic models. Hydrogen oxidation experiments were used to generate a post-ignition condition of 4% H\(_2\)O, \(T = 1700–1800\) K, \(P = 3–4\) atm, which was studied using the DCS. Hydrogen oxidation provided a controlled method to produce water that could be measured using the DCS and compared to two spectroscopic models: the HITEMP model and a recent model put forth by Polyansky \textit{et al} and part of the ExoMol database [36, 37]. Publically available fortran scripts, ExoCross, were used to
execute the ExoMol model in this study [38]. Conditions used for the spectroscopic simulation relied on modeled predictions of the post-ignition temperature using USC Mech II, the water mole fraction measured by the supporting water diagnostic, and pressure measured using a side-wall transducer. A resulting comparison between the DCS measurement and HITEMP modeled spectrum of high-temperature water is provided in figure 10. The DCS measurements use the post-ignition spectrum of hydrogen oxidation averaged from 400 to 600 μs. It is evident that the measured spectrum under-samples the predicted spectrum and there is poor agreement between the measured and simulated spectra. This under-sampling is due to a comb-spacing that is larger than the size of the spectral features of water at these conditions. Even when considering the under-sampling rate, the difference between the modeled and measured spectrum cannot be reconciled at this time.

It is of great interest to the authors to fit high-temperature spectroscopic models to spectra measured using the DCS. This is in contrast to the method employed in the current study that involved the use a library of measured spectra and absorption cross-sections for speciation. While also very effective, an improved methodology would rely upon fitting spectroscopic models to the DCS data for reduced uncertainty in speciation and to potentially enable thermometry. However, across several species that absorb in the region (propyne, methane, and water), there is a lack of high-temperature spectroscopic data to compare to, or to found models upon, in the 1200 cm$^{-1}$ region. Many studies have begun to address this absence of data already [1, 28, 37, 39–45]. This discrepancy suggests an opportunity for future research as measurements push to shorter timescales and higher temperatures, expanding the utility of models and DCS for use in high-temperature reaction kinetics.

5. Conclusion

A clear tradeoff exists in molecular DCS between spectral and time resolution, with the former investigated thoroughly. Despite the fewer comb teeth used in this study, the mid-infrared dual-comb system demonstrated was capable of resolving propyne oxidation kinetics at 4 μs time resolution. The time resolution and quality of the multi-heterodyne signal showed merit in the high-enthalpy test environment despite having several regions with low SNR. The DCS was validated against a suite of independent laser diagnostics with statistically significant agreement demonstrated between laser systems. The current study presented speciation of propyne and water by comparing a measured absorption surface to a library of experimentally measured, high-temperature absorption cross-sections. Agreement was also observed between the modeled and measured mole fractions of propyne and water, although slight discrepancies did exist at early times and just after ignition, a phenomenon warranting further investigation.

While agreement was obtained between this DCS and a supporting laser diagnostic to measure the broadband spectrum of propyne, a notable disagreement exists between our measured spectra of water and existing high-temperature spectroscopic models. Therefore, whether QCL-based DCS has the ability to measure the finely featured spectra of molecules such as water remains an open question. Towards this end, future studies are needed to make additional measurements of molecules such as water or methane with finely featured spectra in this spectral region. It is also important to make use of ever-improving high-temperature spectroscopic models and cross-section databases, which are also still quite under-developed at these temperatures and in this region.

The case study shown here indicates that a clear opportunity exists for DCS to greatly increase the information content of the traditional laser diagnostics used in high-enthalpy and reacting test environments. As mid-infrared dual-comb spectroscopy emerges, the study of high-enthalpy reacting environments offers a wide canvas of applications on which DCS can make a lasting impact.

Acknowledgment

We acknowledge Wey-Wey Su, Adam Susa, and Séné Cassady for their contributions to the project.

Funding

This material is based upon work supported by, or in part by, the U. S. Army Research Laboratory and the U. S. Army Research Office under contract/grant number W911NF-17-1-0420 and the Air Force Office of Scientific Research, AFOSR Grant No, FA9550-16-1-0195, with Dr Chiping Li as contract monitor.

Conflict of interest

No conflict of interest is reported for this work.

Appendix A

Figure A1 presents the wavenumber- and time-dependent SNR of the system during the propyne oxidation reaction. These
SNR values were used in the formulation of the diagonal weighting matrix used in the speciation calculations.

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