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An intercomparison of measured pressure-broadening, pressure shifting
parameters of carbon dioxide and their temperature dependence

Robert R. Gamache^{†a}, Julien Lamouroux^{a,b}, Valérie Blot-Lafon^c, Eldon Lopes^a

^a Department of Environmental, Earth, and Atmospheric Sciences
University of Massachusetts Lowell
One University Avenue
Lowell, MA 01854, USA

^b current address
Laboratoire Interuniversitaire des Systèmes Atmosphériques (LISA): CNRS UMR 7583,
Universités Paris Est Créteil et Paris Diderot, Institut Pierre-Simon Laplace. Université Paris Est
Créteil, 94010 Créteil Cedex, FRANCE

^c Department of Physics
Université de Rennes 1
F-35042 Rennes Cedex, FRANCE

[†] Corresponding author

Robert_Gamache@uml.edu

978-934-3904

Abstract:

An intercomparison of measured pressure-broadening and pressure-shifting coefficients for carbon dioxide absorption lines was done. The work focuses on collision systems where a significant number of data can be found ($\text{CO}_2\text{-N}_2$, $\text{CO}_2\text{-O}_2$, $\text{CO}_2\text{-air}$, and $\text{CO}_2\text{-CO}_2$) and yield information important to applications to Earth's atmosphere. The literature was searched for measured line shape parameter data for the collision systems mentioned above. Databases were created for each perturbing gas with the ro-vibrational transition as the key. Using these databases, intercomparisons of the measurements of half-widths, their temperature dependence, and line shifts were made. The data allow the investigations of trends in the data with respect to the vibrational and the rotational quantum numbers, various line shape models, and isotopologue effects.

The data were averaged and an estimated uncertainty determined. The averaged data sets are evaluated with respect to the need of the spectroscopic and remote sensing communities. In general many data points do not agree within the stated uncertainty estimate independent of the broadening species. Agreement between measurements of the temperature dependence of the half-width is in a poor state and the understanding of the temperature dependence of the line shift is in its infancy. The intercomparison data show more measurements are needed for the $\text{CO}_2\text{-O}_2$, $\text{CO}_2\text{-N}_2$ and $\text{CO}_2\text{-CO}_2$ systems. In addition to the half-width at 296 K, data are needed for the temperature dependence of the half-width and line shift.

Keywords: CO_2 , measured half-widths, line shifts, temperature dependence of half-width and line shift

INTRODUCTION

Carbon dioxide is a trace gas in the Earth's atmosphere with a concentration of 0.039445% [1]. Even with this seemingly low mixing ratio it is the second strongest absorber of infrared radiation in the Earth's atmosphere after water vapor [2]. CO₂, however, has a residence time of about 120 years in the atmosphere compared to water vapor which has a residence time of ~9 days. Because of its long lifetime, CO₂ is evenly mixed in the atmosphere. Due to these facts carbon dioxide has become the standard by which other greenhouse gases are gauged in global climate change. The mixing ratio of CO₂ is being monitored daily and records of direct measurement go back to 1957 [3, 4]. Proxy methods allow the records to be extended back hundreds of thousands of years [5]. The current concentration of 394 ppm is larger than it has been in the last 650 thousand years [5, 6] and probably much longer. Recent proxy data show an abrupt rise in the CO₂ concentration after the industrial revolution and this increase is thought to be of anthropogenic origin [7].

It is paramount that the role of CO₂ in global climate change is better understood. As such the international community is actively monitoring the sources and sinks of CO₂ globally. Measurements are being made from ground-based networks [8], from balloons [9, 10], and from a number of satellite platforms. NASA's AQUA satellite [11] has the Atmospheric Infrared Sounder (AIRS) [12] and the METOP-A satellite [13] has the Infrared Atmospheric Sounder Interferometer (IASI) [14] both measuring in the mid-infrared region. In 2009 the Greenhouse Gases Observing Satellite (GOSAT) [15] was launched by Japan to make measurements of carbon dioxide in the 1.6 μm and 2 μm region. The Jet Propulsion Laboratory attempted to launch the Orbiting Carbon

Observatory [16] in 2009 but the satellite failed to reach orbit. The importance of the mission [17] has prompted NASA to go forward with the launch of OCO-2 in the 2013-2014 time frame. The OCO-2 mission will require mixing ratios be retrieved with a precision of 0.3% placing strong demands on the spectroscopic community [18]. Miller *et al.* [18] state that the OCO remote sensing precision requirements for the Lorentz broadened half-width and line shift are 0.6% and $< 0.0002 \text{ cm}^{-1} \text{ atm}^{-1}$ respectively. This requirement is for air-CO₂ thus in this work the 0.6% precision level is applied to N₂-, O₂-, and air-CO₂ mixtures. While self-broadening of CO₂ in the Earth's atmosphere is negligible (Note, self-broadening is the dominant contribution in some planetary atmospheres.) the same precision requirement is used to be consistent with the other broadeners. They also point out that line shape models more accurate than the Voigt model are needed.

Of the spectroscopic parameters, it is the line shape parameters that are the least well known and while the shift can affect retrievals [19-22] the half-width has a stronger effect in retrievals from atmospheric spectra [23-26] reaching 1:1, error in the half-width to error in the retrieval, at surface pressures. This work presents an intercomparison of the measured half-widths and line shifts of carbon dioxide important for terrestrial applications. This study was undertaken to assess the state of knowledge of measurements of the line shape parameters for CO₂ and to address if the needs of the remote sensing community can be satisfied. These data allow an understanding of the influence of vibration, J dependence of the broadening, shifting, and temperature dependence, different collision partners, different line shape models, and broadening for different isotopologues. The intercomparison will allow a sense of the uncertainty in the

parameters. These data will point to gaps in the measurement record and suggest measurements that are crucially missing in order to complete a selected set of data. These data were also used to identify measurements that could be used to test the theoretical models.

Section 2 discusses the collection of the experimental results and the creation of the databases. Section 3 reports the analysis of the data including the dependence of the collision-induced parameters on vibration, rotation, temperature, broadening partner, and line shape models. Intercomparisons of the data are also discussed in this section. In section 4, conclusions and recommendations are made.

2.0 THE EXPERIMENTAL RESULTS

This work considers the collision-broadened half-widths and pressure induced line shifts of carbon dioxide for terrestrial applications. A search of the literature was made for line shape parameters for the $\text{CO}_2\text{-N}_2$, $\text{CO}_2\text{-O}_2$, $\text{CO}_2\text{-air}$, $\text{CO}_2\text{-CO}_2$ collision systems. The result was 87 papers from 1968 to 2012 [22, 27-112]. Most of the work focused on measurements of the pressure-broadened half-width with 33, 13, 24, and 48 papers, respectively, reporting data on N_2 -, O_2 -, air-, and self-broadening of CO_2 . The first measurements of the pressure-induced line shift for CO_2 were reported in 1998. The number of papers reporting measurements of the pressure-induced line shift drops to 9, 3, 11, and 12 for the perturbors respectively. Considering the temperature dependence of the half-width the number of papers drops further with 9, 3, 4, and 4 for N_2 -, O_2 -, air-, and self-broadening of CO_2 . The temperature dependence of the line shift has only

recently been studied with 2 papers each for the air- and self-collision systems. A small number of papers (8) report measurements on the lesser isotopologues of CO₂ (¹⁶O¹³C¹⁶O, ¹⁶O¹²C¹⁸O, and ¹⁶O¹³C¹⁸O). The data also allow the vibrational dependence of these line shape parameters to be considered with measurements available for 17 (CO₂-N₂), 8 (CO₂-O₂), 17 (CO₂-air), and 37 (CO₂-CO₂) vibrational bands. Tables 1-4 present a summary of the data for the CO₂-N₂, CO₂-O₂, CO₂-air, CO₂-CO₂ collision systems respectively. These tables list the maximum J'' value, the number of data for the half-width, n , the line shift, n_s , and the temperature dependence of the half-width, n_n , the minimum and maximum values of γ , δ , and n as a function of isotopologue and vibrational band. The notation of the vibrational quantum numbers in the Table is that of Toth *et al.* [113].

Approximate position of Tables 1-4

The data in Tables 1-4 show that half-widths change by a factor of 2.01, 1.8, 2.2, and 6.4 for N₂-, O₂-, air- and self-broadening respectively. As an example, the air-broadened value can be compared with that for water where the half-widths change by a factor of 44 [114]. As a function of J'', the half-widths increase at very low J'' values and then as J'' increases the half-widths decrease. The line shifts are mostly negative in the database files. For example the CO₂-N₂ collision system has a few positive values for transitions to low energy vibrational states. One of these values is much larger than the others and may be erroneous. Similar problems may arise looking at the maximum values of the temperature exponent for N₂-, O₂-, and self-broadening. In the study by

Predoi-Cross *et al.* [68] the air-broadening n values ranged from roughly 0.65 to 0.89. Theoretical calculations by Gamache *et al.* [115-117] for the 4 broadeners of this study show $n < 0.8$. Caution should be exercised when using values greater than 1.0.

Tables 1-4 lists the vibrational bands that have been studied for the different CO₂ collision systems. The table shows that the maximum number of ν_1 , ν_2 , and ν_3 quanta exchanged in the ro-vibrational transitions that have been measured are small; $\Delta \nu_{1\max}$, $\Delta \nu_{2\max}$, and $\Delta \nu_{3\max}$ are 2, 4, and 3 for N₂-broadening; 2, 4, and 1 for O₂-broadening; 2, 4, and 3 for air-broadening; and 3, 4, and 3 for self-broadening.

To account for the various collisional effects observed for CO₂ at different experimental conditions a number of different line shape models have been used to reduce the measured spectra. For a review of the various models see the work of Hartmann *et al.* [118] and references therein. Table 5 gives the line shape models used to fit the spectra and an index labeling the model in the database. The results from these models are discussed below.

Approximate position of Table 5

These data have been organized into databases for each perturbing gas with the values corrected, when necessary, to 296 K. The format is similar to that of the line shape parameter databases for water [114, 119] or ozone [120] with modifications for the quantum numbers and an index for the line shape model used in the reduction of the measured data.

3.0 Intercomparison of the data

3.1 Intercomparison of the collision-induced parameters

The database files for each collision system were taken and sorted to group transitions with the same quantum numbers; $v_1'v_2'v_3'J' \leftarrow v_1''v_2''v_3''J''$. These data appear in groups of 2-9 points. The half-width data were compared to each other (intercomparison) by determining the average percent difference (APD),

$$APD = Ave \left\{ \sum_i \sum_{j \neq i} 100 \times \left(\gamma_i - \gamma_j \right) / \gamma_i \right\}, \text{ and average absolute percent difference (AAPD),}$$

$$AAPD = Ave \left\{ \sum_i \sum_{j \neq i} Abs \left[100 \times \left(\gamma_i - \gamma_j \right) / \gamma_i \right] \right\}, \text{ by comparing points I and J where I=1 to}$$

N, J=I+1 to N and I≠J, where N is the number of intercomparison points. For the line shifts the APD and AAPD were calculated however due to comparing small numbers many of the values are large. As an alternative the average absolute ratio,

$$AR = Ave \left\{ \sum_i \sum_{j \neq i} \frac{\delta_i}{\delta_j} \right\}, \text{ of the data were taken. Table 6 lists the number of points in the}$$

intercomparison, n_i , the number of transitions with n_i , the APD, AAPD, and AR. This table clearly shows the lack of data for the CO₂-O₂ system. In general the APDs range from ~0.2 to 3% and the AAPDs from ~2 to 6%. The ARs vary considerably from 0.65 for the oxygen-induced shift to 2.27 for air-induced shift with $n_i=3$. These differences are more apparent in the plots. Figures 1-2 and 3-4 show example plots for the half-widths and line shifts for CO₂-N₂ (Figs. 1-2) and CO₂-CO₂ (Figs. 3-4) collision systems respectively.

Approximate position of Table 6

Approximate position of Figs 1-4

3.2 Effects of rotational transition

There are a number of measured data sets, each for a particular vibrational band, that contain a sufficient number of transitions to address the effect of the rotational transition on the line shape parameters. Figure 5a-c present the measured half-width, line shift, and temperature dependence of the half-width as a function of m , where $m = -J''$, $J''+1$ for P- and R-branch transitions, respectively for the $30012 \leftarrow 00001$ band of CO_2 in collision with N_2 ; note measurements go to $|m| \approx$ mid sixties. In general the data show the half-width decreases as $|m|$ increases. This structure is true for the other perturbors studied here. A similar structure is observed for the line shift with the shift becoming more negative as $|m|$ increases. The temperature dependence of the half-width decreases with some oscillation as seen in the plot. Note, this structure is evident even considering the scatter in the data. Recent theoretical work for the $\text{CO}_2\text{-N}_2$, $\text{CO}_2\text{-O}_2$, $\text{CO}_2\text{-air}$, and $\text{CO}_2\text{-CO}_2$ systems [115-117], which demonstrates excellent agreement with measured half-widths, line shifts, and temperature dependence of the line shift, shows similar plots with calculations to $|m|=120$. Complete sets of measurement plots are available on the web site of one of the authors (faculty.uml.edu/Robert_Gamache).

Approximate position of Fig. 5

3.3 Effects of vibration

The number of vibrational transitions observed varies with the particular perturber, see Tables 1-4 for details. There are no experimental studies of the vibrational dependence of the half-width that have been both systematic and comprehensive. A good number of papers have been published addressing the vibrational dependence of the half-width within confined spectral regions. It is difficult to draw any firm conclusions from these studies. A number of papers [22, 35, 36, 48, 52-55, 64, 66, 107, 108] report that the vibrational dependence of the half-width is very small; “within the error bars.” Several studies report a measurable vibrational dependence in CO₂ self-broadened half-widths [64, 71]. There are also studies that contradict one another; for example Young and Chapman [106] report half-widths in the 9.4 μm band are 5% smaller than those in the 10.4 μm band whereas Moskalenko and Zotov [121] state half-widths in the 9.4 μm band are 5% larger than those in the 10.4 μm band.

In a recent theoretical work on the vibrational dependence of half-widths and line shifts of CO₂ transitions, Gamache and Lamouroux [122] noted that in the above analyses most of the comparisons are for vibrational bands within the same polyad or between bands that have roughly the same number of vibrational quanta exchanged. In their study they began with the expression for the vibrational dependence of γ and δ developed by Gamache and Hartmann [123] and modified the expressions for a study of the vibrational dependence of γ and δ of CO₂ transitions giving,

$$\gamma \left[(v'_1, v'_2, v'_3) f \leftarrow (v''_1, v''_2, v''_3) i \right] = I_{f \leftarrow i}^\gamma + A_{f \leftarrow i} \text{abs} \left(c_1 |\Delta v_1| + c_2 |\Delta v_2| + c_3 |\Delta v_3| \right)^{p_\gamma} \quad (1a)$$

$$\delta \left[\left(v_1', v_2', v_3' \right) f \leftarrow \left(v_1'', v_2'', v_3'' \right) i \right] = I_{f \leftarrow i}^{\delta} + B_{f \leftarrow i} \text{abs} \left(c_1 |\Delta v_1| + c_2 |\Delta v_2| + c_3 |\Delta v_3| \right)^{p_{\delta}} \quad (1b)$$

where γ and δ are the half-width and line shift for the ro-vibrational transition

$\left(v_1', v_2', v_3' \right) f \leftarrow \left(v_1'', v_2'', v_3'' \right) i$, $I_{f \leftarrow i}^{\gamma}$ and $I_{f \leftarrow i}^{\delta}$ are the intercepts and $A_{f \leftarrow i}$ and $B_{f \leftarrow i}$ are the

slopes both with units of $\text{cm}^{-1} \text{ atm}^{-1}$, and $\left(c_1 |\Delta v_1| + c_2 |\Delta v_2| + c_3 |\Delta v_3| \right)^{p_{\gamma}}$ and

$\left(c_1 |\Delta v_1| + c_2 |\Delta v_2| + c_3 |\Delta v_3| \right)^{p_{\delta}}$ are the *Quantum Coordinates* of the lines describing the

vibrational dependence of the half-widths and line shifts respectively. These expressions

fit the calculated data very well ($< 5 \times 10^{-4} \text{ cm}^{-1} \text{ atm}^{-1}$ for γ and $< 1.5 \times 10^{-3} \text{ cm}^{-1} \text{ atm}^{-1}$ for δ)

and allow prediction of the line shape parameters for any ro-vibrational transition of CO_2

[124]. They also explain why the studies within a polyad should not show appreciable

vibrational dependence.

Studying the vibrational dependence of the half-width and line shift from the

measurement data allows changes in vibrational quanta up to $\Delta v_1 = 3$, $\Delta v_2 = 6$, $\Delta v_3 = 3$ (see

Tables 1-4 for details). Here, plots of γ or δ versus the appropriate *Quantum Coordinate*

were made for each collision system, $\text{CO}_2\text{-X}$ with $\text{X} = \text{N}_2$, O_2 , air, and CO_2 . Figures 6 and

7 show examples for the P18 line of CO_2 broadened by air and the R20 line of CO_2 self-

broadened. Plotted are the half-widths and line shifts with the error bars versus the

quantum coordinate. Also shown in the plots are the predicted fits of the data from Ref.

[124] as the solid black line. What is clear is that there is considerable scatter in the

measured values. Many of the data points do not overlap within the quoted errors. The

prediction formulas give reasonable descriptions of the measurements. Complete sets of

plots for all collision systems are available on the web site of one of the authors (faculty.uml.edu/Robert_Gamache).

Approximate position of Figs 6-7

3.4 Line shape models

In an effort to reduce the fit residuals when determining the parameters that describe the spectrum many line shape models have been used. Table 5 shows that, for CO₂ as the radiating molecule, eleven line shape models have been employed when fitting spectra. These models, which are more advanced than the Lorentz (or Voigt) model, contain additional fitting parameters, some up to six parameters [125], and always reduce the fit residual. However, it has been shown [126, 127] that for some of these models some of the parameters giving the improved fits are unphysical. Figures 1 and 8 show the measured half-widths deduced using different line shape model versus line number. The plot symbols for the different line shape models are given in Table 5. Also shown in the plot is the predicted value [124] as the solid black triangle. The plots show small differences between the half-widths reduced using different line shape models. However, looking at many plots (available online at faculty.uml.edu/Robert_Gamache) the scatter in the data and uncertainties it is difficult to make definitive statements about the effect of the model on the derived half-widths. Luckily, there are a number of studies that applied two line shape models to a number of measured spectra and allow a statement. For example, Refs. [56, 67, 105] show that their speed-dependent model gives results a few percent higher than the Voigt model, which was also mentioned by Hartmann *et al.* [118].

Note however, there are plots that show one model's results higher than another model in one plot and lower in another.

Approximate position of Fig. 8

3.5 Isotopologue effects

A limited number of half-width measurements are available for several of the isotopologues of the CO₂ molecule; ¹²C¹⁶O₂, ¹³C¹⁶O₂, ¹⁶O¹²C¹⁸O, and ¹⁶O¹³C¹⁸O. A good number of intercomparisons are available for the pair ¹²C¹⁶O₂ and ¹³C¹⁶O₂, there are also a limited number of comparisons for the pairs ¹²C¹⁶O₂ and ¹⁶O¹²C¹⁸O and ¹²C¹⁶O₂ and ¹⁶O¹³C¹⁸O. Theory [128, 129] provides a guide to the effects of the mass dependence of the half-width. Assuming all collisions are resonant and a single intermolecular potential term of the form R^{-q} , the temperature dependence of the half-width in cm⁻¹/molecule is

$$\gamma(T) \propto T^{-\frac{(q-3)}{2(q-1)}} \quad (2)$$

Because the temperature dependence is essentially due to the mean relative speed $v_r(T)$ in the intermolecular collision, and since $v_r(T)$ is proportional to $\sqrt{T/\mu}$ the following relations hold

$$\gamma(T) \propto v_r(T)^{-\frac{(q-3)}{(q-1)}} \quad (3)$$

and

$$\gamma(T) \propto \mu^{-\frac{(q-3)}{2(q-1)}} \quad (4)$$

where μ is the reduced mass. Note all of the collision systems considered here have the quadrupole-quadrupole interaction as the leading term in the intermolecular potential, i.e. $q=5$. Thus, knowing the broadening of $^{12}\text{C}^{16}\text{O}_2$ by a broadener, the broadening of other isotopologues can be approximated by

$$\gamma(^a\text{C}^b\text{O}_2 - X) = \gamma(^{12}\text{C}^{16}\text{O}_2 - X) \left(\frac{\mu(^a\text{C}^b\text{O}_2 - X)}{\mu(^{12}\text{C}^{16}\text{O}_2 - X)} \right)^{-\alpha} \quad (5)$$

where $\alpha=0.25$.

The calculated isotopologue scaling factor (ratios of the reduced masses raised to the power $-\alpha$) are listed in Table 7 for N_2 , O_2 , air, and $^{12}\text{C}^{16}\text{O}_2$ as the collision partners. For air the reduced mass was determined using the average molecular weight of dry air, 28.96 g/mol [130]. The table indicates that the half-widths for the other heavier isotopologues will be slightly smaller than those of $^{12}\text{C}^{16}\text{O}_2$, of order less than 1%. Looking at the intercomparison plots there are many where the heavier isotopologue half-widths are $\sim 1\%$ smaller, however there is much scatter in the measurement database and the relationship between isotopologues is not always the same. Figure 9 shows the P 34 line of the $00011 \leftarrow 10001$ (top panel) and $00011 \leftarrow 10002$ (bottom panel) bands. Looking at the data for these two bands [22] and comparing the half-width for the $^{13}\text{C}^{16}\text{O}_2$ isotopologue to that of the $^{12}\text{C}^{16}\text{O}_2$ isotopologue one finds the $^{13}\text{C}^{16}\text{O}_2$ value slightly larger for the $00011 \leftarrow 10001$ band and slightly smaller for the $00011 \leftarrow 10002$ band.

While there are some exceptions this observation generally holds in the data [22], indicating the possibility of a vibrational effect. Looking at the other plots for isotopologues (available at faculty.uml.edu/Robert_Gamache) it is difficult to draw any conclusions due to the scatter in the data. Often there are data points above and below the lesser isotopologue half-widths. The same observation is made for the line shift intercomparisons with respect to the isotopologues.

Approximate position of Fig. 9

3.6 Temperature dependence of the half-width

As discussed above, the number of papers that report the temperature dependence of the half-width is relatively small. The temperature dependence of the half-width is expressed using the power law model, see section 3.5, and is generally expressed as

$$\gamma(T) = \gamma(T_0) \left[\frac{T_0}{T} \right]^n \quad (6)$$

where n is the temperature exponent. The use of the power law model has been called into question for certain radiator-perturber collision pairs and for large temperature ranges [131, 132] with important considerations for CO₂ as the radiating molecule [115-117]. Note, the n data from Ref. [37] for CO₂-N₂ are unusually large (some values are greater than 2) well outside the range of other measurements or theoretical calculations.

From the measured data there are no intercomparisons of n for the CO₂-N₂ and CO₂-O₂ collision systems, there are 4 intercomparisons for the CO₂-air collision system, and for the CO₂-CO₂ collision system 112 plots were made, however the intercomparisons are of the data from Ref. [68] where two different line shape models, Voigt and speed dependent Voigt, were used. Thus, the intercomparisons are of the line shape models and no conclusion can be drawn from the data as the Voigt model gives n values sometimes a few percent lower and sometimes a few percent higher than the speed dependent model. Considering J'' and J' and ignoring vibrational dependence, which is expected to be small (see section 3.3), more intercomparisons are possible. Figure 10 shows measured temperature exponents with the corresponding error bars for the R20 transition for CO₂-CO₂ versus the *Quantum Coordinate*. As before, the plot symbols refer to the line shape model used to reduce the spectra. Note, for the *Quantum Coordinate* around 0.5 there are 4 data points. These are from Ref. [68] for the 30012 $\leftarrow$ 00001 and 30013 $\leftarrow$ 00001 bands using the Voigt and a speed dependent Voigt model. Finally, Fig. 11 shows the temperature exponent versus m , where m equals $-J''$ for P-, J'' for Q-, and $J''+1$ for R-branch transitions. In black are the recent measurements of Predoi-Cross *et al.* [68] for the 30012 $\leftarrow$ 00001 and 30013 $\leftarrow$ 00001 bands. The plot symbols reflect the line shape model used to reduce the data. The scatter in the data is quite noticeable. It appears the comment in the HITRAN article [2] discussing improvements needed in the line shape parameters of ozone, “This especially concerns the temperature exponents where the experimental measurements rarely agree with each other or with the theoretical calculations,” also applies to CO₂. This observation is not particularly surprising since it has been shown [120] that the uncertainty in the temperature exponent n is related to the

uncertainty in the half-width and the temperature range used in the measurements. The uncertainties in n in the databases appear to be underestimated, they are generally 1- (or 2-) sigma of the fit and do not include systematic errors or the temperature range of the measurements.

4.0 CONCLUSIONS AND RECOMMENDATIONS

A literature search of measured half-widths, line shifts, temperature dependence of the half-widths and line shifts for CO₂ in collision with N₂, O₂, air, and CO₂ has led to the collection of 7087 data points that have been stored in databases. Using these databases intercomparison of the data was done to determine trends in the data, rotational-, vibrational-, isotopologue-, and temperature-dependence. In general many measurements do not agree within the stated uncertainty estimate. Considering measurements made in the past decade the data points still do not agree within the quoted error. The uncertainties reported seem to be one standard deviation of the fit of the line rather than the combined uncertainty accounting for all statistical and systematic uncertainties. Miller *et al.* [18] state “The multispectrum technique can determine spectrum line half-widths with an uncertainty better than 0.1%, but the systematic uncertainties have always limited the absolute accuracy of half-widths determined from experimental spectra to 1–2%.” It is likely that the reported uncertainties often underestimate the actual uncertainties because systematic contributions to measurement uncertainty are not considered. In the future, researchers should report the true error of the measurements.

The needs of the spectroscopic community, especially the OCO-2 mission, place a heavy burden on measurement. While Table 6 reports averages for the intercomparisons, the data can be studied line by line. For air-broadening only 35 of the transitions out of the 224 intercomparisons have the average absolute difference less than or equal to 0.6%. For N₂-, O₂-, and self-broadening there are 21, 4, and 67 transitions out of the 233, 39, and 536 intercomparisons respectively that have an AAD $\leq$ 0.6%. In the air-broadened data none of the 35 transitions are in the spectral range of importance to OCO-2. An additional 43 transitions have an APDs between 0.6 and 1 and 137 have APDs between 1 and 5 for air- broadening. Thus there is a need for more measurements of CO₂-air mixtures.

To aid theoretical calculations measurements need to be made for single perturbing gases (N₂, O₂, CO₂) and not mixtures (air). Looking at the databases there is a shortage of half-width measurements for the CO₂-O₂ system allowing only 36 2-point and 3 3-point intercomparisons. More high-accuracy measurements of half-widths for the CO₂-N₂ and CO₂-CO₂ systems would also be useful. While the line shifts are less important to remote sensing they are very useful to gauge the theory. Measurements of the temperature dependence of the half-width are in a poor state as demonstrated by the intercomparisons. More measurements should be done considering larger temperature ranges. Gamache *et al.* [120] have shown that the uncertainty in the temperature exponent is proportional to the ratio of the uncertainty in the half-width over the half-width divided by the natural log of T_{max}/T_{min} (see Appendix A of Ref. [120]). Thus the larger the temperature range the smaller the uncertainty in n .

Measurements of the temperature dependence of the line shift are still in their infancy. In early works by Varanasi and Chudamani [133] and by Grossmann and Browell [134, 135] a model similar to that used for the half-width given by

$$\delta(T) = \delta(T_0) \left[\frac{T_0}{T} \right]^{n'} \quad (7)$$

was used. Note, all parameters have their usual meaning. However, as pointed out by Gamache and Rothman [136], and later by Smith *et al.* [137], this model does not allow the shift to change sign. Frost [138] proposed a model that allows for a change of sign in the line shift modified here to give the shift at temperature T from the shift at a reference temperature T_0 (note the correction of Eq. (5) of Ref. [137])

$$\delta(T) = \delta(T_0) \left[\frac{T_0}{T} \right]^{n'} \left[\frac{1 + A \ln(T)}{1 + A \ln(T_0)} \right] \quad (8)$$

where A is a fitted constant which is negative (see Frost for details). Gamache and Rothman [136] proposed a different model that allows for the sign of the shift to change

$$\delta(T) = \delta(T_0) S(T') \left[\frac{T_0}{T} \right]^{n'} \quad (9)$$

where $S(T')$ changes from -1 to 1 at T' . The last two models require fits to obtain the additional parameters. The work of Smith *et al.* [137, 139-141] suggest their data can be fit by a linear model

$$\delta(T) = \delta^0(T_0) + \delta'(T - T_0) \quad (10)$$

where the sign of δ' depends on the choice of T_0 . Frost states that their expression has no physical significance; it simply gives a reasonable fit to the data. The same is true for Eqs. (9) and (10). More work on a theoretical expression is needed.

Researchers should try to measure the half-width and line shift and the temperature dependence of these parameters. It has been found that adjusting the potential surface in CRB calculations by fitting γ , δ , and n simultaneously places strong constraints on the parameters describing the surface. What has been found is that only a unique set of parameters can reproduce all three line shape parameters [115-117]. It would be interesting to see how the addition of the temperature dependence of the line shift will affect the fits.

For new measurements to be useful more effort must be made to determine the true uncertainty of the measurements. An excellent discussion of the determination of uncertainties in line shape measurements can be found Ref. [142]. The emphasis should be on the accuracy of the measurement rather than the number of transitions measured.

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al. [68] for the 30012 $\leftarrow$ 00001 and 30013 $\leftarrow$ 00001 bands. The plot symbols reflect the line shape model used to reduce the data.

Table 1 Summary of measurement data for the CO₂-N₂ collision system.

CO ₂ -N ₂ Band	J _{max}	n _v	n _s	n _n	$\gamma_{\min}^*$	$\gamma_{\max}^*$	$\delta_{\min}^*$	$\delta_{\max}^*$	n _{min}	n _{max}
00011←00001	82	268	133	21	58.50	96.96	-3.72	-0.09	0.720	0.860
00011←10001	48	266	210		60.79	97.00	-4.32	0.90		
00031←00001	36	23	26		71.10	96.30	-9.50	-5.00		
11101←00001	60	37			74.18	106.48				
11101←10002	38	19			66.51	83.75				
20012←00001	46	17	13	2	61.10	111.97	-7.00	-3.40	1.059	1.130
30012←00001	50	102	51		55.73	97.29	-8.49	-2.73		
30013←00001	50	106	49	2	60.71	99.49	-8.52	-3.53	0.740	0.800
00011←10002	52	227	218		67.11	98.40	-4.90	1.73		
01111←01101	52	63		7	69.03	95.00			0.860	0.860
10001←00001	50	93	92		63.50	105.10	-5.50	4.30		
10001←01101	40	9			62.45	83.36				
01101←00001	56	27		16	59.55	91.32			0.608	1.132
10002←00001	46	93	90		62.10	99.60	-5.60	1.40		
10011←00001	48	9			68.70	95.80				
20031←00001	20	1			70.00	70.00				
02211←02201	36	3		3	65.63	84.55			0.860	0.860

* in units of cm⁻¹atm⁻¹

Table 2 Summary of measurement data for the CO₂-O₂ collision system.

CO ₂ -O ₂ Band	J _{max}	n _i	n _s	n _n	$\gamma_{\min}^*$	$\gamma_{\max}^*$	$\delta_{\min}^*$	$\delta_{\max}^*$	n _{min}	n _{max}
00011←00001	82	21		9	51.30	83.80			0.820	0.820
11101←00001	60	31			64.16	93.79				
11101←10002	26	18			56.34	73.98				
20012←00001	46	13	13		54.70	73.00	-7.10	-2.90		
30012←00001	56	72	61		53.13	86.99	-14.44	-2.98		
30013←00001	56	76	56	2	53.20	86.43	-16.36	-1.24	0.660	0.720
00011←10002	30			4					0.930	1.090
10011←00001	48	9			57.50	85.10				

* in units of cm⁻¹atm⁻¹

Table 3 Summary of measurement data for the CO₂-air collision system.

CO ₂ -air Band	$J_{\max}$	n_i	n_s	n_n	$\gamma_{\min}^*$	$\gamma_{\max}^*$	$\delta_{\min}^*$	$\delta_{\max}^*$	$n_{\min}$	$n_{\max}$
00011 $\leftarrow$ 00001	82	71	63		57.00	92.80	-6.40	-1.10		
00011 $\leftarrow$ 10001	48	226	218		62.80	98.33	-4.69	0.70		
00031 $\leftarrow$ 00001	48	30	41		66.40	94.50	-12.10	-3.86		
20012 $\leftarrow$ 00001	64	60	53		60.91	95.20	-7.60	-2.08		
20013 $\leftarrow$ 00001	62	58	61		63.20	94.40	-7.41	-2.32		
30012 $\leftarrow$ 00001	56	290	263	55	49.05	95.75	-15.21	-0.62	0.646	0.884
30013 $\leftarrow$ 00001	56	284	260	56	63.14	96.42	-12.16	-2.40	0.673	0.876
00011 $\leftarrow$ 10002	52	226	220		62.40	93.21	-4.20	1.83		
10001 $\leftarrow$ 00001	50	93	93		63.10	92.60	-3.10	1.70		
31113 $\leftarrow$ 01101	31	49	39		67.25	89.47	-10.28	-0.02		
20011 $\leftarrow$ 00001	60	51	50		62.40	95.50	-8.08	-2.27		
21112 $\leftarrow$ 01101	47	52	42		66.20	94.20	-6.36	-2.61		
21113 $\leftarrow$ 01101	45	37	60		66.70	92.20	-6.71	-2.67		
21111 $\leftarrow$ 01101	44	48	50		67.30	90.90	-6.05	-2.81		
10002 $\leftarrow$ 00001	46	93	91		64.40	106.40	-3.50	1.30		
10011 $\leftarrow$ 00001	50	6			66.70	77.50				
20031 $\leftarrow$ 00001	20	1			80.00	80.00				

* in units of cm⁻¹atm⁻¹

Table 4 Summary of measurement data for the CO₂-CO₂ collision system.

CO ₂ -CO ₂ Band	J _{max}	n _i	n _s	n _n	$\gamma_{\min}^*$	$\gamma_{\max}^*$	$\delta_{\min}^*$	$\delta_{\max}^*$	n _{min}	n _{max}
00011←00001	82	125		25	53.19	126.46			0.891	2.158
00011←10001	60	213	114	14	64.00	127.20	-4.80	-0.60	0.580	0.820
00031←00001	56	69	28		78.00	126.70	-13.30	-4.75		
01121←00001	38	47	25		79.40	116.70	-8.71	-4.31		
10011←10002	32	34			82.50	113.50				
11101←00001	12	6			107.00	134.30				
11101←10002	40	20			75.81	104.64				
20012←00001	74	160	48	15	64.80	127.00	-11.60	-2.36	0.48	0.801
20013←00001	72	161	66		67.70	126.40	-9.61	-2.81		
21102←01111	59	8			57.80	74.00				
30012←00001	62	382	270	112	56.48	135.70	-20.53	-1.90	0.428	0.871
31112←01111	16	1			103.94	103.94				
30013←00001	62	354	232	112	34.88	134.41	-12.68	-2.00	0.429	0.823
00011←10002	62	159	122	19	63.60	126.70	-3.40	-0.70	0.600	1.090
01111←01101	48	42			65.34	124.53				
10001←00001	50	23		12	49.93	123.38			0.706	0.990
10001←01101	40	9			99.32	120.69				
01101←00001	56	23		10	73.40	123.05			0.187	0.743
31112←01101	48	129	121		69.91	147.34	-11.55	0.30		
40013←10002	30	24	17		86.91	116.91	-14.24	-2.90		
31113←01101	46	122	111		76.77	125.09	-9.89	-0.72		
40014←10002	32	18	11		70.77	141.94	-14.07	-2.63		
20011←00001	72	65	59		66.30	127.10	-11.77	-2.55		
21112←01101	62	58	43		68.90	111.60	-9.48	-4.13		
21113←01101	59	69	62		66.80	123.30	-9.27	-2.64		
21111←01101	57	73	54		69.50	121.70	-8.80	-2.54		
30014←00001	48	47	44		72.60	120.60	-10.77	-2.82		
30011←00001	44	34	31		76.00	112.30	-9.80	-2.76		
01131←01101	37	50			86.90	116.30				
10002←00001	32	7			85.80	109.90				
10011←00001	70	2		2	130.00	171.00			0.654	0.695
10012←00001	42	19			78.37	126.28				
12211←00001	46	19			157.87	223.97				
20031←00001	30	14			63.00	106.98				
11102←01101	48	20	20		80.07	111.73	-11.36	-11.36		
02211←02201	15	1	1		102.60	102.60	-3.44	-3.44		
12201←01101	46	105			75.40	126.50				

* in units of cm⁻¹atm⁻¹

Table 5 Database Index and corresponding line shape model.

Index	Model	Plot symbol	CO ₂ -N ₂	CO ₂ -O ₂	CO ₂ -air	CO ₂ -CO ₂
1	Voigt	+	X	X	X	X
2	Galatry	*	X	X	X	X
3	Speed dependent Voigt	○			X	
4	Rautian	×			X	X
5	Speed dependent Nelkin-Ghatak	□			X	
6	Speed dependent Rautian	△				
7	Lorentz (pure)	◇	X	X		X
8	Speed dependent Lorentz	☆				X
9	Modified Voigt (includes line-mixing)	▲	X			X
10	Speed dependent - See Ref. [27]	■	X	X	X	X
11	Speed dependent - See Ref. [55]	●			X	X

Table 6 Intercomparison of the half-width and line shift data. See text for details.

CO ₂ -N ₂	γ			δ	
n_i	n data	APD	AAPD	n data	AR
2	68	2.79	5.52	15	1.17
3	38	0.46	5.42	51	1.25
4	16	0.82	4.25	73	1.22
5	68	0.38	1.78		
6	40	0.68	2.34		
7	2	2.52	5.84		
8	1	1.54	2.40		
CO ₂ -O ₂	γ			δ	
2	36	0.23	5.15	5	0.65
3	3	1.70	2.07		
CO ₂ -air	γ			δ	
2	21	-1.80	2.74	15	1.02
3	15	-1.53	1.98	14	2.27
4	22	0.30	1.67	21	1.03
5	134	-0.17	1.06	159	1.18
6	18	-0.56	2.82	4	1.24
7	10	-2.38	2.92		
8	2	2.91	5.84		
9	2	1.81	5.69		
CO ₂ -CO ₂	γ			δ	
2	234	0.28	2.97	216	0.96
3	150	-1.67	4.00	4	0.99
4	21	-0.28	3.38	74	0.99
5	27	-1.62	4.25	29	1.03
6	55	-0.20	6.01	5	1.54
7	18	1.12	3.73		
8	29	2.65	5.03		

Table 7 Isotopologue scaling factors for the half-width and line shift.

Broadening species	$^{13}\text{C}^{16}\text{O}_2/^{12}\text{C}^{16}\text{O}_2$	$^{16}\text{O}^{12}\text{C}^{18}\text{O}/^{12}\text{C}^{16}\text{O}_2$	$^{16}\text{O}^{13}\text{C}^{18}\text{O}/^{12}\text{C}^{16}\text{O}_2$
N ₂	0.9978	0.9957	0.9937
O ₂	0.9976	0.9954	0.9932
air	0.9978	0.9956	0.9936
CO ₂	0.9972	0.9945	0.9919

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Figure 1

Figure 2

Figure 3

Figure 4

Figure 5

Figure 6

Figure 7

Figure 8

Figure 9

Figure 10

Figure 11