A new photometric ozone reference in the Huggins bands: the absolute ozone absorption cross section at the 325 nm HeCd laser wavelength

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Abstract. The room temperature (294.09 K) absorption cross section of ozone at the 325 nm HeCd wavelength has been determined under careful consideration of possible biases. At the vacuum wavelength of 325.126 nm, thus in a region used by a variety of ozone remote sensing techniques, an absorption cross-section value of $\sigma = 16.470 \times 10^{-21}$ cm$^2$ was measured. The measurement provides the currently most accurate direct photometric absorption value of ozone in the UV with an expanded (coverage factor $k = 2$) standard uncertainty $u(\sigma) = 31 \times 10^{-24}$ cm$^2$, corresponding to a relative level of 2 ‰. The measurements are most compatible with a relative temperature coefficient $c_T = \sigma^{-1} \partial_T \sigma = 0.0031$ K$^{-1}$ at 294 K. The cross section and its uncertainty value were obtained using generalised linear regression with correlated uncertainties. It will serve as a reference for ozone absorption spectra required for the long-term remote sensing of atmospheric ozone in the Huggins bands. The comparison with commonly used absorption cross-section data sets for remote sensing reveals a possible bias of about 2 %. This could partly explain a 4 % discrepancy between UV and IR remote sensing data and indicates that further studies will be required to reach the accuracy goal of 1 % in atmospheric reference spectra.

1 Introduction

High-resolution reference data for ozone absorption in the UV are widely called for, as this region is used for remote and in situ measurement of atmospheric ozone concentrations, and new measurements are therefore under way in the framework of the ESA TROPOMI/Sentinel 5 precursor mission that aims at establishing an improved atmospheric spectroscopy database (SEOM-IAS). The demands for augmented quality of these atmospheric measurements have been increasing continuously over the last decades in order to fulfil the requirement of reliably detecting small atmospheric changes. This was highlighted in the last report of the “Absorption Cross-Sections of Ozone” (ACSO, http://igaco-o3.fmi.fi/ACSO) from the joint initiative of the International Ozone Commission (IO3C), the World Meteorological Organisation (WMO) and the Integrated Global Atmospheric Chemistry Observations (IGACO) O$_3$/UV subgroup, which was dedicated to studying, evaluating and recommending the most suitable cross-section data to be used in atmospheric ozone measurements (Orphal et al., 2016). Remote sensing of tropospheric ozone by joint retrieval of UV and IR satellite instruments is another emerging application (e.g. Cuesta et al., 2013) that strongly depends on unbiased UV spectroscopic data as most of the ozone resides in the stratosphere, but accurate knowledge of the ozone spectrum is also required for the retrieval of other, less abundant trace gases that absorb in spectral ranges where ozone acts as an interfering species.
Reference cross-section values with an uncertainty of 1 % or better at the 90 % confidence level have only recently become available at and around the Hg line position of 253.65 nm (Viallon et al., 2015). This wavelength is particularly important, because absorption at this position is currently used as an ozone standard via standard reference photometers (Hearn, 1961; Viallon et al., 2006). At other wavelengths, such SI-traceable data at a similar accuracy level are not available, and currently used absorption cross-section data in the atmospheric remote sensing of ozone (GSWCB, BDM, BP which stand for Gorshelev, Serdyuchenko, Weber, Chehade and Burrows Gorshelev et al., 2014; Serdyuchenko et al., 2014, Brion, Daumont and Malicet, Brion et al., 1993; Daumont et al., 1992; Malicet et al., 1995, and Bass and Paur, Bass and Paur, 1985; Paur and Bass, 1985) do not provide the same level of accuracy and traceability, which might lead to inconsistent and biased results.

However, the UV range between 302 and 340 nm in the Huggins bands of ozone is particularly interesting for ozone column measurements from the ground using Brewer and Dobson spectrophotometers or for ozone retrieval using differential optical absorption spectroscopy with ground-based or satellite instruments. The traceability of total column ozone including a comprehensive uncertainty budget is thus an important objective of the Joint Research Project ATMoz (traceability for ATMospheric total column Ozone) within the European Metrology Research Programme (EMRP). The retrieval of total column ozone from solar radiation measurements in the Huggins band requires cross sections with very low uncertainties and well-defined temperature coefficients to take into account the effective ozone temperature which varies depending on location and season.

In this article we present new measurements of the UV absorption cross section at the HeCd laser wavelength using the photometric method. Particular attention has been paid to the pressure measurement, the sample purity and the decomposition of ozone during the measurement process. This has led to an improvement of a factor of about 10 in the overall uncertainty of the measurement when compared to the reference of Hearn. The measurement thus provides a new reference in the spectral region that is most important for atmospheric remote sensing of ozone. An uncertainty budget following the ISO Guide to the Expression of Uncertainty in Measurement (GUM) is given and instrumental biases that might have affected earlier measurements are discussed in detail.

2 Experimental setup and methodology

2.1 Ozone production and handling

Ozone is produced from high purity oxygen gas (99.9995 %, Air Liquide, France) in a dedicated vacuum system that has been described elsewhere (Janssen et al., 2011). Here we briefly describe some key points (see Fig. 1). The system is made from Pyrex and equipped with all glass valves using PTFE fittings. The only metal parts are gas flasks, pressure gauges, the turbo-molecular pump and stainless steel parts that connect these components. Ozone is produced by electric discharge at LN2 temperatures in a 3 L reactor, to which copper electrodes are attached at the outside of the walls. After several evaporation and re-condensation cycles, the sample is transferred into a cold trap operating at 65 K, where it is further purified and then released into the absorption cell. The total volume of the cell, which can be closed off by an all-glass stopcock equipped with PTFE fittings, is 113 cm3.

Figure 1. Vacuum system for ozone sample preparation. AC: UV absorption cell; B1: Baratron 690 (10 Torr); B2: capacitive pressure gauge (1000 hPa); BS: gas buffer spiral; CT: cold trap operable at 65 K; DR: electric discharge reactor chamber (3 L); and TMP: turbo-molecular pump.

2.1.1 Sample pressure

Over the last 3 years, the capacitive 10 Torr pressure head (Baratron 690, MKS) of high accuracy (0.08 % nominal) is regularly calibrated at 1-year intervals by the French National Metrology Institute (LNE; last certificate no. P156207/1). Due to metal surfaces in the gauge and the stabilisation at +45 °C, slight ozone decomposition has been observed. In order to improve the stability during the pressure reading, a buffer gas technique has been employed (Janssen et al., 2011).

2.1.2 Sample temperature

Four thin film four-wire Pt100 sensors were distributed over the length of the absorption cell and attached to its outside. The signals were registered continuously by a Picotech (pt-104) data logger. The probes and data logger were calibrated right after the measurement series by an in-house comparison with a traceable standard platinum reference thermometer (SPRT-5626, Hart Scientific) coupled to a readout unit (1502A, Hart Scientific). The calibration uncertainty ($k = 2$) of 14 mK is smaller than observed temperature gradients.
2.2 Photometer setup

The absorption measurements are performed using a customized photometer, of which an overview is given in Fig. 2. As a light source, a HeCd laser (Kimmon) is used. It delivers around 12 mW of output power at the laser wavelength of 325 nm. The laser light passes through a chopper, which modulates the beam amplitude at a frequency of about 2 kHz. The beam is then widened and only a small portion is selected by a ~1 mm pinhole. A 30:70 beam splitter divides the beam and projects the reflected part on the reference detector. The transmitted beam is guided twice through the 30 cm long absorption cell using a flat mirror at the backside of the cell. Both the signal (I) and the reference intensities (I_r) are measured using cooled Si photovoltaic detectors (Newport/Oriel) with integrated transimpedance preamplifiers. The cell windows have a vertical inclination of 3° with respect to the optical axis in order to avoid light being reflected back and forth between the two cell windows to fall onto the detectors. The amplitude modulation of the beam intensity allows for phase-sensitive detection of the reference and absorption signals, which are measured by digital lock-in amplifiers (SRS 830). Their output signals are registered by a PC using a multi-purpose data acquisition card (NI PCI-6281).

2.3 Sample purity and control measurements

In order to control and assess the purity of the ozone sample, a strict protocol of sample preparation, cell filling and pressure measurements has been followed, as described elsewhere (Janssen et al., 2011). After the measurement has been completed, the sample was re-condensed in an empty cell measurement, kept at a temperature of about 65 K. From the residual pressure, the mole fraction v_nc of non-condensable impurities, such as air, that might have entered the system through small leaks, or oxygen that originates from ozone decomposition, could be estimated. The small mole fraction v_c of condensable impurities that might be present in the current absorption cell has been estimated previously (Janssen et al., 2011). No attempt was made to repeat that quantification here. This was motivated by the fact that ozone decomposition rates in the absorption cell and the amount of non-condensable impurities after the experiment have not changed since. In the earlier study, the mole fractions of water, carbon dioxide, nitrous oxide and nitrate were measured and found to be \(-0.10(17), 0.07(7), 0.3(3)\) and \(-0.01(6)\) mmol mol\(^{-1}\) respectively. Moreover, an upper limit of all nitrogen containing impurities of \(1.3\) mmol mol\(^{-1}\) was found (Janssen et al., 2011, Table I).

2.4 Straight-line fit and data evaluation

The Beer–Lambert law implies a proportionality between the optical density \(\tau\) and the absorbers’ column density \(\xi = n \cdot L\):

\[
\tau = -\ln \left( \frac{I/I_0}{I_r/I_{r0}} \right) = \sigma \cdot (n \cdot L) = \sigma \xi,
\]

with the absorption cross-section \(\sigma\) being the proportionality constant. The \(I/I_0\) ratios designate intensities that are normalised for laser intensity fluctuations by means of a reference beam (\(I_r\)) and indices \(m\) and 0 indicate an ozone and an empty cell measurement respectively. In a plot of the optical density \(\tau\) vs. \(\xi\), the cross-section \(\sigma\) is obtained as the slope of this linear relation. Due to uncertainties in both variables, a standard least-squares fit is not appropriate. Because ozone column data are correlated (Bremser and Hässelbarth, 1998; Viallon et al., 2015), a weighted total least-squares (WTLS) fit with correlated uncertainties is required. The solution of the total least-squares problem ultimately goes back to Deming (1943), and there is now a wealth of literature on the York–Williamson algorithm which treats the straight-line adjustment with and without correlated uncertainties in \(x-y\) data pairs (York, 1966, 1968; Williamson, 1968; York et al., 2004; Reed, 2015, for example). The algorithm is frequently used in environmental, geochemical and isotope studies (e.g. York, 1968; Ludwig and Titterington, 1994; Cantrell, 2008; Wehr and Saleska, 2017). It seems, however, that fewer studies (e.g. Amir-Simkooei et al., 2014; Bremser and Hässelbarth, 1998; Bremser et al., 2007; Malengo and Pennecchi, 2013) are devoted to the problem of when the structure of the covariance matrix is more complex and when correlations exist between uncertainties in different values of \(x\) and/or \(y\). This type of question arises in chemometric or metrological applications when calibration lines need to be used or when instruments are to be compared.

In order to treat the latter problem, we use here an algorithm from Amir-Simkooei et al. (2014), which we implement using the Mathematica software (Wolfram Research, Inc., 2016). Our implementation provides the fit coefficients \(a\) and \(b\) of the straight-line function \(y = a x + b\), the associated standard uncertainties \(u(a)\) and \(u(b)\), Pearson’s correlation coefficient \(r(a, b)\) and the chi-squared value \(\chi^2\). The code has been tested on all benchmarks in the ISO technical specification (ISO, 2010, data given in Tables 4, 6, 10, 22 and 25 therein). These include the case of uncertainties in \(x\) and \(y\), and the two cases when there are covariances associated with the \(y\) values and when covariances are associated with both \(x\) and \(y\) values. Our results agreed within all digits indicated. We also note that our implementation further matched all example calculations given in the original publication of Amir-Simkooei et al. (2014). This comprises the classical data set from Pearson with York’s weights assuming no correlations in \(x-y\) data pairs (York et al., 2004), but agreement to all digits as given by Reed (2015) is also obtained when such a correlation is considered.
3 Analysis and uncertainty budget

3.1 Laser wavelength

We are not aware of direct interferometric measurements of the $4d^95s^2:2D_{3/2} \rightarrow 4d^95p^2P_{1/2}$ laser transition at 325 nm. Reported wavelength values are based on the analysis of emission spectra of the Cd$^+$ ion produced in electric discharges. Previous atmospheric studies (Lakkala et al., 2008; Lantz et al., 2002) using a HeCd laser mostly report an air wavelength of 325.029 nm. This number emanates probably from term energies reported in the handbook of basic atomic spectroscopic data (Sansonetti and Martin, 2005) that are ultimately based on one comprehensive study (Shenstone and Pittenger, 1949). Other databases, reference tables and handbooks (e.g. Reader et al., 1980; Haynes, 2015) recommend the slightly different value of 325.033 nm, which corresponds to the wavelength that Shenstone and Pittenger (1949) actually measured for this transition. While the 0.004 nm difference between the measured and the term energy derived transition energy is compatible with the measurement uncertainty of $\sim 0.1$ cm$^{-1}$, Burns and Adams (1956) confirmed the Shenstone and Pittenger (1949) measured value at a much lower degree of uncertainty ($< 0.01$ cm$^{-1}$). Indeed, the more accurate of the two measurements resulted in a vacuum wavelength $\lambda_{\text{vac}} = 325.126$ nm, which under standard conditions ($T = 15$ °C, $p = 101325$ Pa) and reasonable variation of the air molecular composition (RH = (50 ± 50) % and x(CO$_2$) = (0.4 ± 0.1) mmol mol$^{-1}$), corresponds to the air wavelength $\lambda_{\text{air}} = 325.033$ nm with all figures significant to the last digit (Cid-dor, 1996).

Further confidence into the claimed wavelength accuracy might be obtained by comparing measured and tabulated wavelengths of the well-studied HeNe laser. The Atomic Spectra Database gives an air wavelength of 632.8614 nm for the Ne I transition, corresponding to a vacuum wavelength $\lambda_{\text{HeNe}} = 632.9914$ nm (when standard conventions are applied: $T = 15$ °C, $p = 101325$ Pa, x(CO$_2$) = 0.33 mmol mol$^{-1}$, RH = 0). This result agrees to all digits with the reproducible line position of typical HeNe lasers (Mielenz et al., 1968).
3.2 Optical density

The stability of the laser fluctuation-corrected signal $I/I_0$ is shown in Fig. 3. The displayed curve is characteristic for our system and allows for a conservative uncertainty estimation, because curves at other measurement days gave values at a lower level. We chose integration times of about 30 s for the ozone and empty cell measurements, which were taken within a time span of about 2 min. For the ratio $(I/I_0)_m/(I/I_0)_h$ we infer a measurement uncertainty of

$$u(r) = 1.8 \times 10^{-4} \sqrt{(3 + \exp(r))}/2,$$  \hspace{1cm} (2)

where $r$ denotes the optical density of the absorption measurement. In deriving the above expression we assumed firstly that the measurements of the empty cell (0) and of the cell filled with ozone ($O_3$) are stochastically independent (white noise behaviour) during the whole measurement period that lasted for about 2 min, secondly that intensities of the empty cell and the reference beam contribute equally and thirdly that the relative uncertainty of the filled cell signal scales with $1/\sqrt{T_m} \sim \sqrt{\exp(r)}$. As the measurement signal is the ratio of two intensity ratios (thus the product/ratio of four measurement signals), the relative uncertainty must yield $\sqrt{2}$ times the 30 s level in Fig. 3 for $r = 0$. Note that the value of $2.1 \times 10^{-4}$ in Fig. 3 is conservative, because the spectrometer stability at other days was always better. We thus replaced that number with the moderately lower and more representative value of $1.8 \times 10^{-4}$. For optical densities between 0.025 and 0.32, as in this study, Eq. (2) implies standard uncertainties $u(r)$ between $2.6 \times 10^{-4}$ and $2.7 \times 10^{-4}$. This is only slightly higher than the residual scatter ($2.5 \times 10^{-4}$) of our measurements (see Fig. 5b in Sect. 4.1). The uncertainties correspond to relative values of $u_r(r)$ in the 0.08 to 1.0 % range.

3.3 Temperature

A temperature gradient along the cell of about 100 mK was observed. In the absence of more accurate data, we determined the cell temperature as the average of the minimal and maximal temperature during a measurement ($\sim 30$ s). Taking into account the uncertainty of the calibration (7 mK), the standard uncertainty was determined as

$$u^2(T) = (T_{\text{max}} - T_{\text{min}})^2/12 + (7 \text{ mK})^2.$$ \hspace{1cm} (3)

3.4 Pressure

The capacitive pressure sensor is regularly calibrated at LNE. The calibration determines the measurement uncertainty from the scatter of repeated readings and the standard uncertainty of the LNE working standard. In the relevant 10 Torr range, reading errors were shown to be negligible. However, there is a small pressure rise observed during the measurement, which is likely due to some ozone decomposi-

$$u(p) = 0.05 \text{ Pa} + 10^{-3} \cdot p.$$ \hspace{1cm} (4)

The laboratory temperature has an effect on offset and span of our sensor. While the offset is always readjusted, we need to consider the manufacturer-specified span temperature coefficient of $2 \times 10^{-5}$ K$^{-1}$. We are always within $\pm 2.5 \text{ K}$ of the calibration temperature, which adds an uncertainty of $5 \times 10^{-5}$ to the span. Because this is at least 20 times smaller than the calibration uncertainty, we can simply ignore it here. The dominant pressure uncertainties in Eq. (4) are of type B and do not lessen over repeated measurements. This is confirmed by the long-term drift between our sensor and the LNE working standard, which shows a characteristic pattern that evolves slowly over the calibration period of 3 years.

3.5 Optical path length

The path length was determined from the window thickness and calliper measurements of the outer cell dimensions combined with the observation of the entrance and exit positions on the two cell windows. A HeNe laser beam has temporarily been superposed to the UV beams and the cell centre axis in order to determine the different inclination angles. The procedure is described in detail in Appendix A. Altogether, seventeen different measurements contribute to the determination of the cell geometry and the orientation of the two beams with respect to the cell. All of these are included in the uncertainty budget of the optical path length $L = l_1 + l_2 = (596,654 \pm 0.243) \text{ mm}$ (see Table 1), which is obtained as the sum of the individual lengths on the round trip through the cell. We only list and discuss the four factors that contribute most. The remaining non-listed quantities add less than 1 %.

The most important (95 %) contribution to the uncertainty is from the window thickness. The manufacturer-specified tolerance, which we verified on other windows of the same production batch, is $\pm 0.1 \text{ mm}$. We therefore deduce a standard uncertainty $u(d) = 100 \mu\text{m} / \sqrt{3} = 57.7 \mu\text{m}$. Superposing transparent millimetre paper on the cell windows allowed determination of the coordinates $(x_1, y_1, x_2, y_2)$; see Appendix A) where the laser beams passed through the cell windows. A standard uncertainty of $u(y_1) = u(y_2) = u(x_1) = u(x_2) = 1 \text{ mm} / \sqrt{3} = 577 \mu\text{m}$ is estimated for these measurements. They are listed at second and third position in Table 1. Since the cell is passed through by the laser beam 2 times, resulting in two beams with separate optical path lengths, these two contributions need to be accounted for twice. Due to the window inclination, the length is more sensitive to the vertical coordinate.

The forth most important contribution to the uncertainty is due to the shortest distance $L$ between the inclined, but not exactly parallel windows. It was measured using a cal-

www.atmos-meas-tech.net/11/1707/2018/ Atmos. Meas. Tech., 11, 1707–1723, 2018
that was compared to gauge block combinations with overall lengths of 290 and 300 mm. As the resolution of the calliper is 10 µm, the comparison with the gauge blocks always gave perfect agreement. The uncertainty of the length measurement was therefore obtained as the quadratic sum of two contributions: the standard uncertainty related to the calliper resolution (5 µm/√3) and the error of the mean of eight measurements at two different days, which was found to be 9.8 µm.

The finite dimensions of the laser beam was also taken into account and found to be negligible compared to other factors. The beam divergence was estimated using the divergence angle $\alpha_d \sim 5 \times 10^{-4}$, determined from beam profile measurements before the entrance and after the exit of the cell. To first approximation, its effect ($\sim \sec(\alpha_d) - 1 \approx \alpha_d^2/2 \approx 10^{-7}$) on the path length is negligible. The finite diameter was explored through numerical simulations of parallel displacements of our beam centre. We shifted the centre by 1.5 mm (the beam diameter is between 2.6 and 3.3 mm) in all directions and found that the two displacements lead to changes of the length by less than 2 parts in $10^6$.

### 3.6 Sample purity

The sample purity is characterised by the mole fractions of condensable ($\nu_c$) and non-condensable gases ($\nu_{nc}$). In a previous study, the mole fractions of CO$_2$, H$_2$O, N$_2$O, and NO$_3$ were determined and an upper limit for the sum of all oxides and hydrogen oxides of nitrogen, of $\nu_{nc} < 1.3$ mmol mol$^{-1}$ was found. Despite the fact that the observed mole fractions of the directly measurable quantities were all within one standard uncertainty or close to 0, we assume a rectangular probability distribution function (pdf) with bounds at 0 and 1.3 mmol mol$^{-1}$. We thus obtain $\nu_{HxCyOz} = 0.65$ mmol mol$^{-1}$ with a standard uncertainty of 0.38 mmol mol$^{-1}$. When combining this result with the observations on CO$_2$ and H$_2$O, we obtain $\nu_c = 0.62$ mmol mol$^{-1}$ with a standard uncertainty $u(\nu_c) = 0.42$ mmol mol$^{-1}$.

The mole fraction of non-condensables was determined from measurements of the residual pressure after condensation of the cell content as

$$\nu_{nc,max} = \gamma \cdot \frac{p_{res}}{p}, \quad (5)$$

where $p_{res}$ is the residual gas pressure, and $\gamma = 7.24 \pm 0.44$ is a factor which takes into account the volume ratios and temperature gradients between the absorption cell, the volume where the residual pressure measurements are made and the cold finger where ozone is frozen back. The uncertainty of $\gamma$ comprises the reproducibility of test measurements (1.9 %) and varying levels of LN$_2$ that change the effective volume of the cold finger (5.8 %). Residual pressure measurements are impacted by the thermal transpiration effect (Daudé et al., 2014) caused by the heating of the gauge (45°C). It can be taken into account by assuming that the actual pressure is somewhere between the indicated value and the maximum of 4.2 % induced by thermal transpiration. This leads to a $+2.1 \ %$ correction of the pressure reading with an associated standard uncertainty of the residual pressure measurement of 1.2 %.

The value in Eq. (5) is a limiting value, as it was obtained only after the measurement and it is likely that the non-condensables, in addition to entering into the measurement cell during sample admission, also get in especially when the re-condensation of ozone takes place and the residual pressures are measured. In the absence of further information, we simply assume a rectangular probability distribution 0 $\leq$ $\nu_{nc} \leq \nu_{nc,max}$ for each measurement. $\nu_{nc,max}$ varies between 1.3 and 4.8 mmol mol$^{-1}$ with an average of 2.9 mmol mol$^{-1}$. This amounts to a typical value of $\nu_{nc} = 1.4$ mmol mol$^{-1}$ with a standard uncertainty of $u(\nu_{nc}) = 0.84$ mmol mol$^{-1}$. We note that the dominating source of uncertainty comes from the unknown origin of the residual pressure and not from individual measurements.

### 3.7 Temperature dependence

For small variability, the temperature dependence of the absorption cross-section $\sigma$ in the vicinity of some reference value $\sigma_0$ at $T = T_0$ is given by
\[
\frac{\sigma(T)}{\sigma_0} - 1 = c_T (T - T_0),
\]
where \(c_T\) is the normalised linear temperature coefficient. As shown further below, our measurements are most compatible with \(c_T = 0.0031 \text{ K}^{-1}\). This is close to data in the literature: 0.0033 K\(^{-1}\) (Serdyuchenko et al., 2014), 0.0042 K\(^{-1}\) (Malicet et al., 1995), and 0.0039 K\(^{-1}\) (Paur and Bass, 1985) at 325.03 nm and 294 K. Since the absorption cross section in the Huggins band is strongly wavelength and temperature dependent, we prefer using \(c_T = 0.0031 \text{ K}^{-1}\) as other values might be biased by small wavelength shifts. For the uncertainty estimate we assume a rectangular pdf with 0.0011 K\(^{-1}\) half width to obtain \(u(c_T) = 0.00064 \text{ K}^{-1}\).

### 3.8 Uncertainty budget for a single measurement

The uncertainty of a cross-section measurement is obtained from the Beer–Lambert law (Eq. 1), taking into account that the ozone column density \(\xi_i\) is given by

\[
\xi_i = (n \cdot L_i) = (1 - \nu_c - \nu_{\text{nc},i}) L \frac{p_i}{k_B T_i},
\]
where the different quantities have their previously defined meanings. However, as will become clear later, it is useful to define an adjusted ozone column density

\[
x_i = \xi_i (1 + c_T (T_i - T_0)) = (1 + c_T (T_i - T_0)) (1 - \nu_c - \nu_{\text{nc},i}) L \frac{p_i}{k_B T_i},
\]
where the slight temperature dependency of the absorption cross section is incorporated into the coordinate axis (see Sect. 4.1). Quantities with added index \(i\) vary between runs and must be determined for each individual ozone absorption measurement, while others, such as the path length \(L\), always remain the same. These constants necessarily introduce a correlation between different values of \(x_i\). For an individual measurement, where we ignore correlations and the temperature dependency of the cross section, simple error propagation rules yield the following equation for the relative uncertainty of the cross section

\[
u_r^2(\sigma) = \frac{u_r^2(\tau_i) + u_r^2(\xi_i)}{(1 - u(\nu_c) - u(\nu_{\text{nc},i}))^2}.
\]

The different contributions are summarised in Table 2 and a total relative standard uncertainty of \(u_r(\sigma) = 2.3 \times 10^{-3}\) is obtained for an individual measurement. This, for the moment, ignores the uncertainty caused by repeating measurements at slightly different temperatures, taken into account in the full analysis presented later in Sect. 4. The most prominent contributions (> 1 %\(\_\text{o}\)) are due to the measurement of the optical density and the pressure. Repeated measurements will allow improvement in measurement uncertainty, provided that correlations in the pressure and other data contributing to the ozone column density are taken into account.

### 3.9 Correlations between realisations of the ozone column density

Equation (8) provides also the basis for the evaluation of measurement correlations. Constants in that equation clearly introduce a correlation between different values of \(x_i\), but individual realisations of temperature, pressure and the mole fraction of non-condensable impurities are also not strictly independent from one run to another, because their measurements rely on the same calibrations and sensors. The correlation coefficients \(r_{ij} = u^2(x_i, x_j)/u^2(x_i)u^2(x_j)\) between two measurements \(i\) and \(j\) of the ozone column \(x\) can be calculated from Eq. (8), using correlations between the independent measurement quantities and a generalised error propagation rule. Details of the procedure are presented in Appendix B. We obtain

\[
u_r^2(x_i, x_j) = \frac{u_r^2(k_B) + u_r^2(L) + (T_i - T_0)(T_j - T_0)u^2(c_T) + (1 - c_T(T_i - T_0))(1 - c_T(T_j - T_0))}{(1 - c_T(T_i - T_0))(1 - c_T(T_j - T_0))} + \frac{u^2(p_i, p_j)}{u^2(T_i, T_j)} + \frac{u^2(T_i, T_j)}{u^2(T_i, T_j)},
\]

where \(k_B\), \(L\), \(\nu_c\) and \(c_T\) are the independent quantities common to all determinations and where the variables \(p_i\), \(T_i\) and \(\nu_{\text{nc},i}\) are newly determined in each run. The similarity with Eq. (9) is apparent. Indeed with the exception of the term for the optical density, we immediately recover Eq. (9) by setting \(i = j\) and \(u(c_T) = 0\). There is no uncertainty \(u(T_0)\) associated with the arbitrarily chosen reference temperature \(T_0\), which explains the absence of a corresponding term. The calculation of the different terms for \(i \neq j\) is detailed in the remainder of this section.

The Boltzmann constant and the absorption path length contribute via their absolute or relative standard uncertainties to \(r_{ij}\). We obtain (see Tables 2 and 3)

\[
u_r(x_i)u_r(x_j) r_{ij} \big|_{k_B, L} = 1.672 \times 10^{-7}.
\]

Similarly, the contribution to the correlation coefficient through temperature variation of the absorption cross section is

\[
u_r(x_i)u_r(x_j) r_{ij} \big|_{c_T} = 4.033 \times 10^{-7} K^{-2} (T_i - T_0)(T_j - T_0) / (1 - c_T(T_i - T_0))(1 - c_T(T_j - T_0)).
\]
Table 2. Uncertainty budget of a single absorption cross-section measurement at average pressure

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Unit</th>
<th>Probability distribution</th>
<th>Typical or recommended value</th>
<th>Rel. standard uncertainty $u_r(X)$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Length $L$</td>
<td>mm</td>
<td>rect.</td>
<td>596.654</td>
<td>$4.1 \times 10^{-4}$</td>
</tr>
<tr>
<td>Mole fraction complement of non-condensable impurities $(1 - \nu_{nc})$</td>
<td>1</td>
<td>rect.</td>
<td>$1 - 1.4 \times 10^{-3}$</td>
<td>$8.2 \times 10^{-4}$</td>
</tr>
<tr>
<td>Mole fraction complement of condensable impurities $(1 - \nu_{c})$</td>
<td>1</td>
<td>rect.</td>
<td>$1 - 6.2 \times 10^{-4}$</td>
<td>$4.2 \times 10^{-4}$</td>
</tr>
<tr>
<td>Temperature $T$</td>
<td>K</td>
<td>gauss.</td>
<td>294.09</td>
<td>$5.8 \times 10^{-4}$</td>
</tr>
<tr>
<td>Pressure $p$</td>
<td>hPa</td>
<td>gauss.</td>
<td>7.6</td>
<td>$1.1 \times 10^{-3}$</td>
</tr>
<tr>
<td>Opt. density $\tau$</td>
<td>1</td>
<td>gauss.</td>
<td>0.18</td>
<td>$1.6 \times 10^{-3}$</td>
</tr>
<tr>
<td>Temperature dependence of cross section $c_T$</td>
<td>$K^{-1}$</td>
<td>rect.</td>
<td>0.0031</td>
<td>$2.1 \times 10^{-1}$</td>
</tr>
<tr>
<td>Boltzmann constant $k_B$</td>
<td>$J K^{-1}$</td>
<td>–</td>
<td>$1.38064852 \times 10^{-23}$</td>
<td>$5.7 \times 10^{-7}$</td>
</tr>
<tr>
<td>Cross-section $\sigma$</td>
<td>$cm^2$</td>
<td>–</td>
<td>$16.47 \times 10^{-21}$</td>
<td>$2.3 \times 10^{-3}$</td>
</tr>
</tbody>
</table>

Table 3. Contributions to correlation coefficients between different realisations $x_i$ and $x_j$ of the ozone column ($i \neq j$).

<table>
<thead>
<tr>
<th>Quantity ($y$)</th>
<th>First-order contribution</th>
<th>Sensitivity coefficient $(\partial \rho_{ij} / \partial x_i) / (x_i x_j)$</th>
<th>$r_{ij}$</th>
<th>$u_r(X)$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Boltzmann constant $k_B$</td>
<td>$u^2(k_B)$</td>
<td>$3.25 \times 10^{-13}$</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Length $L$</td>
<td>$u^2(L)$</td>
<td>$1.67 \times 10^{-7}$</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Temperature coefficient $c_T$</td>
<td>$(T_i - T_0)(T_j - T_0)u^2(c_T)$</td>
<td>$-4.79 \times 10^{-9}$</td>
<td>$-5.00 \times 10^{-7}$</td>
<td>$4.23 \times 10^{-7}$</td>
</tr>
<tr>
<td>Mole fraction $\nu_{c}$ of condensable impurities</td>
<td>$u^2(\nu_{c})$</td>
<td>$1.76 \times 10^{-7}$</td>
<td>$1.76 \times 10^{-7}$</td>
<td>$1.76 \times 10^{-7}$</td>
</tr>
<tr>
<td>Mole fraction $\nu_{nc,i}$ of non-condensables</td>
<td>$u(\nu_{nc,i}, \nu_{nc,j})$</td>
<td>$2.26 \times 10^{-7}$</td>
<td>$5.01 \times 10^{-8}$</td>
<td>$6.36 \times 10^{-7}$</td>
</tr>
<tr>
<td>Pressure $p_i$</td>
<td>$u(p_i, p_j)(p_i p_j)^{-1}$</td>
<td>$4.70 \times 10^{-8}$</td>
<td>$1.90 \times 10^{-8}$</td>
<td>$2.89 \times 10^{-7}$</td>
</tr>
<tr>
<td>Temperature $T_i$</td>
<td>$u(T_i, T_j)(T_i T_j)^{-1}$</td>
<td>$5.00 \times 10^{-11}$</td>
<td>$5.00 \times 10^{-11}$</td>
<td>$5.00 \times 10^{-11}$</td>
</tr>
</tbody>
</table>

Modified ozone column density $x_i$ 0.76 | 1.00 |

As discussed in Sect. 3.8, individual measurements of non-condensable impurities ($\nu_{nc,i}$) are essentially fully correlated, which is due to the fact that it is not known whether the small amounts of residual gases have already been present during the measurement or were added only afterwards. We thus have $u^2(\nu_{nc,i}, \nu_{nc,j}) = u(\nu_{nc,i})u(\nu_{nc,j})$ and, if we add the constant contribution from the condensables $u^2(\nu_{c}) = 1.764 \times 10^{-7}$, we obtain

$$u_t(x_i)u_t(x_j) \Big|_{\nu_{nc}, \nu_{c}} = \frac{0.5772u(\nu_{nc,i})u(\nu_{nc,j}) + 1.764 \times 10^{-7}}{(1 - \nu_{c} - \nu_{nc,i})(1 - \nu_{c} - \nu_{nc,j})}. \tag{13}$$

Temperature measurements are assumed to be non-correlated except for the contribution due to sensor calibration (7 mK):

$$u_t(x_i)u_t(x_j) \Big|_{T} = \frac{49 \times 10^{-6} K^2(1 - c_T T_0)^2}{(1 - c_T(T_i - T_0))(1 - c_T(T_j - T_0)) T_i T_j}. \tag{14}$$

The uncertainty of the pressure measurement is essentially limited by the calibration. Repeated measurements at the same pressure will thus be fully correlated. Less is known about the correlation of measurements at different pressures. As pointed out by Viallon et al. (2015), assuming a high degree of correlation does not alter the derived value of the absorption cross section, but leads to a conservative uncertainty estimate. Therefore we assume full correlation $u^2(p_i, p_j) = u(p_i)u(p_j)$:

$$u_t(x_i)u_t(x_j) \Big|_{p} = u_t(p_i)u_t(p_j). \tag{15}$$

The range of values for the different contributions is indicated in Table 3. Taking all parts together one gets hold of the correlation coefficient $r_{ij}$. Evidently, $r_{ij} = 1$ for $i = j$, but we still find an average value of $r_{ij} = 0.94$ for $i \neq j$, indicating a very strong correlation between different measurements of the ozone column density. Most of this is due to the foreign gas contamination and the optical path length.

4 Analysis and results

Figure 4 shows the results of 27 individual measurements and an unweighted linear fit to the data. The measurements span the range of $\tau$ between 0.025 and 0.32, corresponding to ozone columns from 0.15 to $1.95 \times 10^{19} \text{cm}^{-2}$. A high coefficient of determination ($r^2 = 0.999977$) attests to the excellent linearity between optical densities $\tau$ and ozone columns $\xi$. Before the cross-section value can be derived, the impact
of temperature and the choice of the fitting model need to be examined.

4.1 Preliminary analysis and choice of fitting function

As discussed previously, the data were obtained for temperatures varying slightly in the range between 293.17 and 295.37 K. This leads to some scatter due to the temperature dependence of the absorption cross section. Using a local linear dependency on temperature, the optical density will be given by

$$\tau(\xi, T) = \sigma(T)\xi = \sigma_0[1 + c_T(T - T_0)]\xi = \sigma_0 x(T),$$  

(16)

where we defined the new variable $x = [1 + c_T(T - T_0)]\xi$ (see Eq. 8). In order to determine the cross section at the average temperature $T_0$, we can now plot $\tau$ vs. $x$, which directly yields the cross section $\sigma_0$ as the slope term. While we allow for an offset $a$ in the linear fit that serves as an additional control, we also need to explore the possibility of non-linearities in our measurement chain, possibly caused by a saturation of the detectors or by other effects in the electronic acquisition and amplification modules. This can be accomplished by including a quadratic term $(b\xi^2)$ in the fit, leading to the following fitting function:

$$\tau(\xi) = a + \sigma_0[1 + c_T(T - T_0)]\xi + b\xi^2.$$

(17)

Figure 5 shows the residuals of fitting this function for different scenarios. In the lower panel relatively large residuals with a reduced standard deviation (SD) of $S_\varepsilon = S\cdot\sqrt{27/24} = 5.0 \times 10^{-4}$ and prominent features at $\xi \sim 1.7 \times 10^{10}$ cm$^{-2}$ are observed, when we assume $b = c_T = 0$. Interestingly, the most variable temperature conditions (between $-0.92$ and $+1.28$ K with respect to the average) prevailed during measurements at these column densities. When a first-order correction $c_T = 0.0001$ K$^{-1}$ for the temperature is taken into account, the largest residual features disappear (as shown in the middle panel of Fig. 5) and the spread of residuals (max–min) is reduced by a factor of 2.0. Correspondingly, the reduced SD of the temperature-corrected residuals in the middle panel of Fig. 5 of $S_\varepsilon = 0.0001$ K$^{-1}$ is only half of that in the lower panel. It has to be noted that this number is only slightly lower than the standard uncertainty of the optical density $\tau$, derived in Sect. 3.2.

Allowing for a quadratic term $+b\xi^2$ in the fit affects residuals (shown on the top panel of Fig. 5) only marginally, diminishing the reduced SD just by 3% to yield $S_\varepsilon = 2.4 \times 10^{-4}$. At the same time, the quadratic term introduces a strong anti-correlation between fit parameters ($r(\sigma_0, b) = -0.98$). This indicates that, while the effect of temperature on the fit is well significant, quadratic terms are not. Our restriction to a straight-line fit is thus well justified in what follows. We also fix the temperature coefficient to our best fit value of $c_T = 0.0001$ K$^{-1}$, because the value is consistent with previously observed data (see Sect. 3.7) and because reasonable changes to this parameter do not modify our result significantly (see Sect. 4.2).

4.2 Linear regression

After having established the fitting function, the data are evaluated using the weighted total least-squares algorithm with correlated $x-\tau$ data. Table 4 summarises the results of the analysis.
The cross-section $\sigma_0 = 1.6470 \times 10^{-20}$ cm$^2$ with a relative standard uncertainty $u(\sigma) = 9.3 \times 10^{-4}$ ($k = 1$) is obtained. The small offset within the uncertainty range indicates that the data comply with our hypothesis of a straight-line passing through the origin, and thus that our measurement follows the Beer–Lambert law. The $\chi^2$ value falls within the 10 and 90 % quantiles of the cumulative $\chi^2$ distribution, which also indicates that the straight-line hypothesis does not need to be rejected and that the uncertainty analysis is compatible with our data.

The importance of considering covariances in this type of photometric absorption measurements (Bremser et al., 2007) is once more emphasised by comparing our results with numbers obtained when these covariances are omitted. Ignoring covariances firstly leads to an unrealistically small value of $\chi^2$ (12.6 instead of 20.8) and, secondly, underestimates $u(\sigma)$ by 34 %. However, the absolute value of $\sigma$ is remarkably robust against ignoring the covariance terms (it changes only by 0.011 % if these are ignored). This finding is in line with the discussion of Viallon et al. (2015), where an effectively constant correlation coefficient $r = r_{ij}$ for all $x_i$-$x_j$ pairs ($i \neq j$) was assumed. But we suspect that this might not generally be true. In particular if $r_{ij}$ strongly varies as a function of $i$ and $j$, we expect that the value of the cross section changes as well upon considering covariances. A possible scenario would be a measurement where different pressure sensors are utilised in different pressure ranges, possibly leading to little correlation between low- and high-pressure values, while maintaining a high correlation coefficient between measurements using the same gauge. We also note that the result is de facto independent of our choice of $C_T$. Using one of the highest values reported in the literature so far ($C_T = 0.0042$ K$^{-1}$ (Paur and Bass, 1985) instead of $C_T = 0.0031$ K$^{-1}$), the derived cross-section value changes by less than 1 part in 10$^5$ and the uncertainty estimate is not at all affected.

### Table 4. Linear fit statistics including standard uncertainty ($k = 1$) values.

<table>
<thead>
<tr>
<th>Quantity</th>
<th>Unit</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Degrees of freedom $\nu$</td>
<td>1</td>
<td>25</td>
</tr>
<tr>
<td>Slope $\sigma$</td>
<td>$10^{-20}$ cm$^2$</td>
<td>1.64704</td>
</tr>
<tr>
<td>Offset $\tau_0$</td>
<td>1</td>
<td>6.684 $\times$ 10$^{-6}$</td>
</tr>
<tr>
<td>Slope uncertainty $u(\sigma)$</td>
<td>$10^{-23}$ cm$^2$</td>
<td>1.530</td>
</tr>
<tr>
<td>Offset uncertainty $u(\tau_0)$</td>
<td>1</td>
<td>1.033 $\times$ 10$^{-4}$</td>
</tr>
<tr>
<td>Pearson’s coefficient $r(\sigma, \tau_0)$</td>
<td>1</td>
<td>$-0.4092$</td>
</tr>
<tr>
<td>Chi-squared $\chi^2$</td>
<td>1</td>
<td>20.8469</td>
</tr>
</tbody>
</table>

Figure 6. Comparison of high-resolution ozone absorption data around the 325 nm HeCd laser line (position indicated by vertical bar). The vacuum wavelength is used on the ordinate scale. Data are from the same sources as in Table 5. Straight lines between measurement points were inserted for visual guidance. Individual uncertainty bars have been omitted from the graph. The uncertainties of this work are smaller than the symbol size. Uncertainty assignments for other spectra are given in Table 5. As a reference, the black vertical bar indicates the $\pm 1$ % relative uncertainty range at $1.647 \times 10^{-20}$ cm$^2$.

## 5 Discussion

### 5.1 Comparison with laboratory data

Table 5 compares our result with previously published high-resolution data or cross sections commonly used for atmospheric retrieval. For convenience, the Table shows not only an analysis of the original data; it also includes the recent parametrisation of cross-section data from BP, BDM and GSWCB, and their uncertainties provided by Weber et al. (2016). Their investigation agrees well (better than 0.2 %) with our analysis of the original data when cross sections are smoothed. Interestingly, all of the literature data only insignificantly deviate from our reference measurement. Except for BP, the literature data sets agree well with each other at 325 nm, but they show values about 2 % higher than our measurement, independent of whether they were calibrated at the Hearn (1961) value (VOPB) or not (GSWCB, BDM). This is different from the situation around the top of the Hartley band. In that region Viallon et al. (2015) observed that data can be divided into two distinct groups: one where values are scaled to the absolute absorption cross section of Hearn, including the old Bremen (VOPB) data, and one where the absolute scale is determined independently, such as their own measurements, the new Bremen (GSWCB) and the Reims (BDM) data, the former group giving values about 2 to 3 % higher than the latter. Obviously this is not the case at the HeCd laser wavelength (see Table 5 and Fig. 6).

The negative offset of the BP data must mostly be explained by a wavelength bias (see Fig. 6). Early evaluations...
Table 5. Comparison of absolute high-resolution absorption cross-section data of ozone at 325.126 nm in vacuum (325.033 nm in air). Smoothed data were obtained from applying a Savitzky–Golay filter of order 2 over the range of 0.1 nm.

<table>
<thead>
<tr>
<th>Data set</th>
<th>Temperature</th>
<th>Cross-section $\sigma$ (10$^{-21}$ cm$^2$)</th>
<th>Rel. standard uncertainty$^c$ $a$($\sigma$) (%)</th>
<th>Relative deviation$^d$ from paper</th>
<th>Relative deviation$^d$ from this work (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>BP (1985)</td>
<td>294.1</td>
<td>16.335</td>
<td>2.3</td>
<td>2.31</td>
<td>-0.8/0.7...2.5</td>
</tr>
<tr>
<td>BDM (1995)</td>
<td>294.1</td>
<td>16.864</td>
<td>2–4</td>
<td>1.74</td>
<td>2.4</td>
</tr>
<tr>
<td>VOPB (2001)</td>
<td>294.1</td>
<td>16.819</td>
<td>4–7</td>
<td>2.1</td>
<td>2.1</td>
</tr>
<tr>
<td>GSWCB(2014)</td>
<td>294.1</td>
<td>16.740</td>
<td>1.1–3</td>
<td>1.65</td>
<td>1.6</td>
</tr>
<tr>
<td>This work</td>
<td>294.09</td>
<td>16.470</td>
<td>–</td>
<td>0.093</td>
<td></td>
</tr>
</tbody>
</table>

$^a$ Data were obtained from the ACSO website: http://igaco-oc.fmi.ck/ACSO/cross_sections.html. References are BP: Bass and Paur (1985), Paur and Bass (1985); BDM: Daumont et al. (1992), Malicet et al., 1995; VOPB: Voigt et al. (2001); and GSWCB: Gershevich et al. (2014), Seryadychenko et al. (2014)

$^b$ The temperature dependence of the literature data has been taken into account using a quadratic parametrisation $\sigma(t) = a_0 + a_1 T + a_2 T^2$, where $T = T - 273.15 \, K$. BP and GSWCB provided corresponding coefficients $a_0$, $c_1$ and $c_2$. For BDM and VOPB, these were obtained from a quadratic fit to cross sections given at fixed temperatures. $^c$ The uncertainty estimation of BDM contains the effect of wavelength shifts, not considered by BP, GSWCB and VOPB. $^d$ Based on smoothed data. BP cross sections suffer from wavelength bias: neg. value uncorrected; range of pos. values after correction (see text).

5.2 Atmospheric implications

The discrepancy at the HeCd laser wavelength indicates a 2 % room temperature bias in current atmospheric reference spectra used by a variety of remote sensing platforms and techniques (Brewer, Dobson, lidar, Umkehr, SBUV, TOMS, OMI, SCIAMACHY and GOME(-2); see Orphal et al., 2016, for example). If that bias applies to a larger wavelength region (~310–340 nm) and to most of the atmospheric temperature range, actual retrievals in this spectral region systematically underestimate atmospheric ozone by about 2 %. Although identical in magnitude this tentative bias in the Huggins bands is different from the ongoing discussion of whether the reference absorption cross section of Hearn (1961) at the Hg line position of 253.65 nm in the Hartley band should be reduced by about 2 % (Viallon et al., 2015; Orphal et al., 2016), because both the GSWCB and the BDM data are already compatible with the lower value at 253.65 nm, and only the BP data set, which is no longer recommended for atmospheric retrieval (Orphal et al., 2016), would be affected by the revision of the absorption cross section at 253.65 nm.

There is a long-standing consistency problem of atmospheric ozone derived from remote sensing in UV and IR spectral regions (e.g. Barbe et al., 2013; Janssen et al., 2016). Both laboratory (Picquet-Varrault et al., 2005; Gratien et al., 2010; Guinet et al., 2010) and atmospheric (Kagawa et al., 2007; Viatte et al., 2011) studies imply that using recommended spectroscopic data in the UV (BP/BDM) and IR (HITRAN2012 Rothman et al., 2013) leads to results that disagree by about 4 to 5 %, with ozone abundances inferred from IR measurements being higher. Thus when comparing measured (msd) and database (db) IR intensities ($I$) with UV cross sections ($\sigma$) through the ratio

$$\bar{R} = (I/\sigma)_{\text{msd}}/(I/\sigma)_{\text{db}} = (\sigma_{\text{db}}/\sigma_{\text{msd}})(I_{\text{msd}}/I_{\text{db}}),$$

the above studies indicate a value $\bar{R}$ around 1.04 or 1.05, whereas consistent data require $\bar{R} = 1$. Note that the factor...
\(\frac{I}{\sigma}\) in this equation is the ratio of two absorption signals, where factors except for the molecular parameters, such as the concentration or light path geometry, ideally cancel. Our new absorption section at 325 nm suggests that the currently used values of \(\sigma_{db}\) might be too high by about 2% \(\left(\frac{\sigma_{db}/\sigma_{msd} - 1}{\sigma_{msd}}\right) \sim 1.02\) which brings IR and UV results to within 2 or 3% if this bias is taken into account. The remaining discrepancy is already close to most measurement uncertainties, but also agrees remarkably well with the value \(I_{msd}/I_{db} = 2.5\%\) observed by Guinet et al. (2010), who investigated 15 intense lines at 8.8 µm in the \(v_1\) fundamental. This situation thus is similar to the spectral conditions in the atmospheric UV–IR comparison of Viatte et al. (2011) using Brewer and FTS instruments, their IR analysis being based on the \(v_1-v_3^\prime\) region at 9.6 µm. The atmospheric comparison of Kagawa et al. (2007) between concentrations from TOMS (UV) and from ground-based FTS (IR) does not directly depend on the intensities in the \(v_1\) fundamental. But the fact that most atmospherically relevant ozone vibrational intensities in HITRAN directly depend on transition moments of the \(v_1\) and \(v_3\) fundamentals implies that IR intensities in the 3 µm region should be corrected by the same amount (Rothman et al., 2005; Fland et al., 2003), implying that the discrepancy observed in their study is resolved at the same time.

The new ozone cross section at the HeCd laser wavelength thus not only provides the first reference value with sub-percent accuracy for ozone spectra in the Huggins bands; it also supplies independent evidence for a shared contribution of IR and UV biases to the UV–IR consistency problem of atmospheric ozone. This is the first evidence directly based on a measurement in the Huggins band, i.e. in the same UV band that is actually utilised for the atmospheric (Kagawa et al., 2007; Viatte et al., 2011) and laboratory (Picquet-Varrault et al., 2005; Gratien et al., 2010) inter-comparisons. A previous laboratory study (Guinet et al., 2010) dependent on UV measurements in the Hartley band. Further systematic temperature- and wavelength-dependent studies with high accuracy will be required to work out the possible bias in currently used atmospheric reference spectra (BDM, GSWCB and forthcoming data) and confirm our assertion with respect to the share of bias between UV and IR data in the spectroscopic databases.

6 Conclusions

Using a HeCd laser spectrophotometer we have obtained the currently most accurate measurement of an ozone absorption cross section in the Huggins bands, and in the spectral region used by a variety of remote sensing techniques and platforms. The cross-section \(\sigma = (16.470 \pm 0.031) \times 10^{-21}\) cm\(^2\) was found at \(\lambda_{vac} = 325.126\) nm, and a full uncertainty budget in accordance with the guide to expression of uncertainty in measurements (GUM) has been presented. The expanded \((k = 2)\) relative uncertainty is at the 2% level and thus significantly more accurate than previous measurements and well below the current target of 1% for atmospheric applications. This high-accuracy level has been made possible by the use of a special ozone production and handling system and an elaborate analysis of the light path in a cell with slightly non-parallel windows. The measurement, together with a recent study at several wavelengths (244–257 nm) in the Hartley band (Viallon et al., 2015), demonstrates that a sub-percent accuracy can now well be achieved in laboratory ozone absorption investigations and shows that the accuracy of atmospheric measurements can be improved significantly.

Our new reference value suggests that absorption spectra currently used for atmospheric remote sensing of ozone possibly need to be revised towards lower values in the Huggins bands by about 2%. Such a revision would likely impact most ozone retrievals in the UV and would also reduce the \(~4\%\) UV–IR discrepancy reported in atmospheric and laboratory studies by a factor of 2. The remaining 2 to 3% need to be attributed to a bias in the IR data, which is compatible with a previous independent IR study. The often cited target uncertainty of 1% has obviously not yet been reached in atmospheric reference spectra. This implies that further studies are required. The possible bias in the atmospheric reference spectra is likely wavelength dependent because atmospheric reference spectra need to be acquired in spectral slices to be combined to cover the entire range from the UV to the NIR, which is a consequence of the 7 orders of magnitude in absorption between the Hartley and the Wulf bands. One would thus ideally make high-accuracy measurements at regular wavelength intervals (10 or 20 nm or so) in order to investigate this wavelength dependence. Unfortunately, this is not highly feasible due to the need for suitable laser sources at all these different wavelengths. In our next step, we propose extending the current measurements to selected UV and VIS wavelengths (particularly around 254, 325 and 633 nm, for example) using both gas and tuneable lasers, and including the whole temperature range down to 190 K. Thus, relevant reference points or even small regions for actual or new atmospheric reference spectra can be obtained. These can be used to calibrate existing and future cross-section data, assess their accuracy, identify wavelength shifts and assure traceability in limited wavelength regions.

Data availability. Raw data displayed in Fig. 4 are available from the corresponding author on request.
Appendix A: Path length in cell with non-parallel windows

Here we describe how the absorption path length in a cell with inclined (but not necessarily parallel) windows is obtained. The full analytic expression was derived using an algebraic software package (Mathematica) and our uncertainty analysis was based on this exact solution. Its analytical form is too clumsy to be fully reproduced here. We prefer to give the closed analytic solution for a cell with parallel windows, together with the first-order correction for slightly non-parallel windows.

The general situation with arbitrarily inclined window plates is illustrated in Fig. A1. In the laboratory system, we define the $z$ axis along the centres of two parallel plates of thickness $d$ and radius $R$, measured between the outer surfaces of the windows. The first centre is located at $z = 0$, and the second at $z = L_0$, $x_1$ and $y_1$ coordinates respectively designate axes in the vertical and horizontal directions in the laboratory frame ($x_2$ and $y_2$ are similarly defined at the origin of the second window). Because windows are assumed to be spherical, two Euler angles suffice to define the window inclination: $\beta_1$ for rotation around the $y_1$ axis, and $\alpha_1$ for rotation around the newly obtained $x_1'$ axis. The passage of the light beam is defined by the coordinates on the entrance ($x_1', \ y_1') = (a_1, b_1)$) and exit window ($x_2', \ y_2') = (a_2, b_2)$) surfaces. We define $\alpha = (\alpha_1 + \alpha_2)/2$ and $\beta = (\beta_1 + \beta_2)/2$ to be the average inclination angles, and $\Delta y = b_2 - b_1$ and $\Delta x = a_2 - a_1$ as the changes of the horizontal and vertical displacements of the window coordinates between the beams’ exit and entrance. We can also characterise the degree of non-parallelism by introducing the angle differences $\Delta \alpha = (\alpha_2 - \alpha_1)/2$ and $\Delta \beta = (\beta_2 - \beta_1)/2$.

Let us first note that the window centre distance $L_0$ can be obtained from the shortest distance $L$ between the two inclined plates, measured with a calliper where the two outside jaws are oriented along the $y_1$ and $y_2$ axes:

$$L_0 = \frac{1}{\cos(\alpha - \Delta \alpha)} \times \left( L \cos^2 \alpha + L \sin(\alpha) \sin(\alpha - 2\Delta \alpha) + R \sin(2\Delta \alpha) \right).$$

(A1)

In deriving this formula we made the convenient but non-restricting assumption that $0 \leq \alpha_1 \leq \alpha_2 < \pi/2$. Assuming that windows are parallel ($\Delta \alpha = \Delta \beta = 0$), the length of a single pass is given by

$$l^{(0)} = l_p^{(0)} A(\Delta x, \Delta y),$$

(A2)

where

$$l_p^{(0)} = \frac{L}{\cos \alpha} \left( 1 - \frac{2d}{L \cos \beta} \right)$$

is the length of the beam propagating parallel to the $z$ axis and where the correction term

$$A(\Delta x, \Delta y) =$$

(A3)

is the length of the beam propagating parallel to the $z$ axis and where the correction term

$$l = l^{(0)} + l_p^{(1)} \Delta \alpha + l_p \Delta \beta + O(\Delta \alpha^k \Delta \beta^m),$$

$$k, m \geq 0 \land k + m = 2.$$  

(A4)

As a matter of fact, the agreement between this approximation and the exact solution is better than 2 nm for a single pass in our configuration. For extreme conditions with $\alpha = 5^\circ$, $\beta = 0$ or $\alpha = 0$, $\beta = 5^\circ$ and $\Delta \alpha$ and $\Delta \beta$ in the $0.5^\circ$ range, where the beam passes through the 30 cm cell within 5 mm of the centre, we find that the linear approximation for one pass always agrees with the full analytic solution by better than 12 µm, which is close to the calliper resolution. Let us introduce some quantities for deriving the coefficients in Eq. (A4):

$$\Delta x = x + \Delta x/2, \quad \Delta y = y + \Delta y/2,$$

$$\Delta = \sqrt{\Delta x^2 + \Delta y^2}.$$  

(A5)
\[ B = 3 \sin \alpha + \sin 3\alpha - 4 \frac{R + \gamma}{L} \cos \alpha. \]  
(A6)

Here, the average horizontal \( \tau \) and vertical \( \gamma \) beam displacements are introduced. Using these abbreviations and the definition of \( A \) in Eq. (A3), the partial derivatives for the first-order corrections in \( \Delta \beta \) and \( \Delta \alpha \) are given as

\[
\begin{align*}
\iota_{1\alpha}^{(1)} &= \frac{L}{2A} \left[ B \left( 1 + \frac{2d}{\cos \beta \cos \alpha} \right) \left( \frac{\Delta x}{L} \sin \beta \frac{\Delta y}{L} \tan \alpha \right) \right. \\
&\quad \left. - \frac{1}{\cos^2 \alpha} - \frac{2d}{L \cos \beta} \left( \frac{\Delta x}{L} \right)^2 \right] + 4 \cos \alpha \left( 1 - \frac{2d}{L \cos \beta} \right) \\
&\quad \times \left( \frac{\Delta y}{L} - \tan \alpha \right) \left( \frac{\tau}{L} \sin \beta + \frac{\gamma}{L} \tan \alpha \right), \quad \text{(A7)}
\end{align*}
\]

\[
\begin{align*}
\iota_{1\beta}^{(1)} &= \frac{2 \tau \cos \alpha}{A} \left\{ \cos \beta \left[ 1 - \frac{2d}{L} \left( \frac{\Delta x}{\cos \alpha \cos \beta} \right)^2 \right] \right. \\
&\quad \left. - \frac{\tau}{L} \sin \beta \right\}. \quad \text{(A8)}
\end{align*}
\]

Appendix B: Correlation terms

The correlation coefficients \( r_{ij} = u^2(x_i, x_j)/(u(x_i)u(x_j)) \) can be obtained from a generalised uncertainty propagation rule:

\[
u^2(x_i(y), x_j(y)) = \sum_{k=1}^{n} \sum_{r=1}^{m} \left( \frac{\partial x_i}{\partial y_{k,r}} \right) \left( \frac{\partial x_j}{\partial y_{k,r}} \right) u^2(y_{k,r}) \\
+ \sum_{l,k=1}^{n} \sum_{r=1}^{m} \sum_{s=1}^{m} \left( \frac{\partial x_i}{\partial y_{k,r}} \right) \left( \frac{\partial x_j}{\partial y_{l,s}} \right) u^2(y_{k,r}, y_{l,s}). \quad \text{(B1)}
\]

Summation indices \((k,l)\) and \((r,s)\) respectively go over the number \( n \) of observables \( y \) in Eq. (8) and the number \( m \) of different measurements. By setting \( i = j \) and identifying covariance terms \( u^2(x_i) \) by variances \( u^2(x_i) \), we recover the familiar propagation rule for standard uncertainties with contributions from both variance and covariance terms. Equation (B1) considerably simplifies when cross-correlation terms vanish. In our case, variables \( k_B, \nu_c, L, c_T \) and \( T_0 \) are common to all realisations and stochastically independent of all other quantities. Their covariance terms thus disappear completely. Due to temperature \( (T) \), pressure \( (p) \) and residual gas \( (v_{nc}) \) measurements being independent of each other, covariances between \( T \) and \( p \), between \( T \) and \( v_{nc} \) and between \( p \) and \( v_{nc} \) also mutually vanish, and so do the variance terms of these variables, because their sensitivity coefficients are necessarily 0 for \( i \neq j \). One thus finds

\[
u^2(x_i, x_j) \quad \frac{x_i}{x_j} \quad \frac{x_i}{x_j} = u^2(k_B) + u^2(L) \\
+ \frac{(T_i - T_0)(T_{j} - T_0)}{(1 - c_T(T_i - T_0))(1 - c_T(T_j - T_0))} u^2(c_T) \\
+ \frac{u^2(\nu_c) + u^2(v_{nc,i}, v_{nc,j})}{(1 - \nu_c - v_{nc,i})(1 - \nu_c - v_{nc,j})} \frac{u^2(p_i, p_j)}{p_i p_j} \\
+ \frac{(1 - c_T T_0)^2}{(1 - c_T(T_i - T_0))(1 - c_T(T_j - T_0))} \frac{u^2(T_i, T_j)}{T_i T_j}. \quad \text{(10)}
\]

Note that \( u(T_0) = 0 \) due to \( T_0 \) being an arbitrary constant and that we have introduced the normalisation factor \( x_i x_j \), which expresses the covariances on the same footing as (the squared) relative uncertainties.
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Competing interests. The authors declare that they have no conflict of interest.

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