

This discussion paper is/has been under review for the journal Atmospheric Measurement Techniques (AMT). Please refer to the corresponding final paper in AMT if available.

Rapid, optical measurement of the atmospheric pressure on a fast research aircraft using open-path TDLAS

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Received: 17 March 2014 – Accepted: 10 April 2014 – Published: 13 May 2014

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Published by Copernicus Publications on behalf of the European Geosciences Union.

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Because of the high travel speed, the complex flow dynamics around an aircraft and the complex dependency of the fluid dynamics on numerous airborne parameters, it is quite difficult to obtain accurate pressure values at a specific instrument location of an aircraft's fuselage. Complex simulations using computational fluid dynamics (CFD) models can in theory computationally "transfer" pressure values from one location to another. However, for long flight patterns, this process is inconvenient and cumbersome. Furthermore these CFD transfer models require a local experimental validation, which is rarely available.

In this paper, we describe an integrated approach for a spectroscopic, calibration-free, in-flight pressure determination in an open-path White cell on an aircraft fuselage using ambient, atmospheric water vapour as the "sensor species". The presented measurements are realized with the HAI (Hygrometer for Atmospheric Investigations) instrument, built for multiphase water detection via calibration-free TDLAS (tunable diode laser absorption spectroscopy). The pressure determination is based on raw data used for H₂O concentration measurement, but with a different post-flight evaluation method, and can therefore be conducted at deferred time intervals on any desired flight track.

The spectroscopic pressure is compared in-flight with the static ambient pressure of the aircraft avionic system and a micro-mechanical pressure sensor, located next to the open-path cell, over a pressure range from 150 hPa to 800 hPa, and a water vapour concentration range of more than three orders of magnitude. The correlation between the micro-mechanical pressure sensor measurements and the spectroscopic pressure measurements show an average deviation from linearity of only 0.14 % and a small offset of 9.5 hPa. For the spectroscopic pressure evaluation we derive measurement uncertainties under laboratory conditions of 3.2 % and 5.1 % during in flight operation on the HALO airplane. Under certain flight conditions we quantified for the first time stalling-induced, dynamic pressure deviations of up to 30 % (at 200 hPa) between the avionic sensor and the optical and mechanical pressure sensors integrated

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in HAI. Such severe local pressure deviations from the usually used avionic pressure are important to take into account for other airborne sensors employed on such fast flying platforms as the HALO aircraft.

1 Introduction

5 Open-path tunable diode laser absorption spectroscopy (op-TDLAS) is a promising approach for airborne in-situ gas analysis. Op-TDLAS does not require to take a gas sample into a closed measurement cell. Instead the gas is optically interrogated at its “natural” location (= in situ) in an open, atmospheric light path. This has the advantage of circumventing typical sampling problems like chemical reactions in the sampling line, delay and integration effects due to the gas transport, or sampling artefacts related to wall adsorption in the piping. This is particularly critical for strongly adsorbing gases like ammonia, HCl or water vapour. op-TDLAS in airborne scenarios thus allows measurements to be conducted directly outside the aircraft’s fuselage, e.g. using open-path cells such as those described in Zondlo et al. (2010), May (1998) or at least by hav-
10 ing a large part of the optical path outside the airplane’s boundary layer (Diskin et al., 2002). Hence, op-TDLAS is well suited for chemical analysis of reactive or strongly adsorbing gases like H₂O. However, absolute chemical species measurements – for example using TDLAS – also need accurate measurements of the physico-chemical boundary conditions, e.g. parameters like gas temperature and gas pressure. This is particularly true for airborne open-path gas analysis and especially for calibration-free
15 TDLAS techniques (Ebert and Wolfrum, 2000; Schulz et al., 2007) where for example no calibration to the flight speed is done and therefore pressure and temperature information is indispensable.

25 Pressure measurements represent a challenge in several ways, since they are needed for the final step in the evaluation of a volume concentration measurement (often applying the ideal gas law). The pressure might also be needed for the

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instrument can also still be calibrated/validated, e.g. after a campaign (Buchholz et al., 2013b), so it is not a choice for or against a principle.

The principle of non-calibrated, absolute dTDLAS is presented here very briefly (in reference to the above-mentioned literature), and the bulk of the effort is focused on the extraction of pressure values from the dTDLAS raw signals intended for species concentration measurements.

2.2 Non-calibrated, direct TDLAS (dTDLAS)

BASIC SETUP. The instrument (called “HAI” = Hygrometer for Atmospheric Investigations) used in this paper to extract absolute, high-speed pressure measurements from an open-path absorption signal is first of all designed for open-path water vapour concentration measurements aboard a specialised high-speed research aircraft (HALO: see www.halo.dlr.de), but the major functions for determining water vapour concentrations are secondarily for the pressure measurement. In the interest of a detailed description of the pressure measurement principles we refrain from explaining all details of the concentration detection procedure, the basic details of which have also been explained in detail elsewhere.

The basic setup of the HAI spectrometer is shown in Fig. 1 and can be visualised in two separate building blocks. Block (A) is the actual open-path measurement setup (Fig. 1, left) where the water molecules are detected. The entire data evaluation of the spectrometer is concentrated in building block (B), which comprises the main electronics unit of the spectrometer (Fig. 1, right). This spatial separation of the two building blocks allows the spectrometer optics to be adapted to the specific requirements of the individual application and is often advantageous when conditions at the measurement region itself are complex or critical (such as open-path measurements on the outer skin of an aircraft). Therefore the open-path cell of the HAI hygrometer has a direct fibre-coupling without any transfer optics and a built-in detector. This avoids exposing the rest of the spectrometer components to the large temperature (+50 to -70°C) and pressure (1000–150 hPa) fluctuations by installing them in the aircraft cabin.

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An absolute evaluation of dTDLAS raw signals to yield mixture fractions of the target species, which is for H₂O concentrations possible with an accuracy in the single digit percentage range (Buchholz et al., 2014), requires absolute gas temperature and gas pressure measurements inside the open-path cell. The temperature data needed for the laser signal evaluation are typically measured by a local T sensor at or near the open-path section. The pressure within the measurement region is usually derived from a pressure transmitter whose interface is located inside the optical unit of building block A and which senses the gas pressure. This is reliably done via a pressure pipe, as long as the frequency of the pressure variations over the flow time along the pipe length is negligible. In complex situations, like on high-speed airborne platforms, both values may be difficult to acquire due to dynamic effects. Therefore the open-path sensor has two temperature sensors and a built-in pressure sensor (described later) to provide, in combination with CFD models and several validations, reliable gas temperature and gas pressure values for the evaluation process.

EVALUATION. One of the major steps of TDLAS spectroscopy is laser current modulation, which is derived from a frequency generator signal, which is used to control a laser current driver. The laser chip temperature is stabilised using Peltier elements driven by a separate temperature control loop. The laser current modulation frequency is typically chosen in the range of 100 Hz–10 kHz, with a typical laser dependent current amplitude of 1–150 mA. The current modulation induces the desired, continuous, dynamic wavelength tuning (typically $< 3 \text{ cm}^{-1}$ for a distributed feedback laser (DFB) laser), as well as laser intensity or amplitude modulation. After transmitting the laser light through the measurement region the molecule-specific attenuation is captured and converted into a photo current using a detector, acquired with suitable data acquisition electronics, and saved to disk as raw data for further offline evaluation. In principle the HAI instrument contains two independent TDLAS spectrometers (working at 1.4 μm and 2.6 μm , respectively). For the spectroscopic pressure determination in this paper solely the 1.4 μm part with the H₂O transition (00–101, 110–211) at 1370 nm (7299.4 cm^{-1}) is used.

The major evaluation steps to obtain the H₂O absorber concentrations are as follows: the wavelength-dependent light intensity $I(\lambda)$ captured behind the measurement region of thickness L is described via the extended Lambert–Beer equation (Eq. 1), which can be written as

$$I(\lambda) = E(t) + I_0(\lambda) \cdot \text{Tr}(t) \cdot \exp[-S(T) \cdot g(\lambda - \lambda_0) \cdot N \cdot L] \quad (1)$$

with N being the number density of the molecular absorbers. The initial laser intensity $I(\lambda)$, the background emission $E(t)$ and the broadband transmission losses $\text{Tr}(t)$ are synchronously derived from the individual raw signals and absorption profiles. Molecular line data like absorption line strength $S(T)$ and its temperature dependence is either obtained from the HITRAN08 database (Rothman et al., 2009) or derived from our own dedicated measurements (Hunsmann et al., 2006). Other molecular spectral parameters, e.g. self or foreign broadening coefficients and their temperature dependence as well as pressure-induced line shift etc., of the normalised shape function $g(\lambda - \lambda_0)$ (centred at the wavelength λ_0) are also obtained from the same data sources.

The combination of the extended Lambert–Beer equation (Eq. 1) with the ideal gas law allows to derive the amount fraction c

$$c = \frac{k_B \cdot T}{S(T) \cdot L \cdot p} \int \ln \left(\frac{I(\nu) - E(t)}{I_0(\nu) \cdot \text{Tr}(t)} \right) \frac{d\nu}{dt} dt \quad (2)$$

The *amount fraction* c is in metrological units correctly specified as [$\text{mol mol}^{-1} = \text{mol absorber per mol gas}$] which is in the environmental community more frequently assigned as “volume fraction” e.g. in units of ppmv or Vol.-%. In Eq. (2) the dynamic tuning coefficient of the laser $\frac{d\nu}{dt}$ (which is a constant property of the individual laser) and fundamental physical constants such as the Boltzmann constant k_B are needed to solve this equation. The first is derived experimentally using the airy signal of the laser light passing through a planar, air-spaced etalon (Ebert and Wolfrum, 2000; Schlosser et al., 2002). From long-term $\frac{d\nu}{dt}$ measurements over several years we could verify, for

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the used laser type, a long-term stability of this basic laser property to better than 1 %, which is within the current uncertainties of the tuning characterisation.

For the pressure (p) and temperature (T) measurements it is important to note that no other hidden parameters generate unwanted additional degrees of freedom in the evaluation, so that no calibration, in particular, has to be conducted for the target species. This explains why we term this technique “calibration-free”. (Of course we have to calibrate our individual pressure and temperature sensors, but the response of the entire spectrometer with regard to the derived target gas concentration is not calibrated, in contrast to usual procedures, Muecke et al., 1994.)

2.3 Line broadening

The dTDLAS signal evaluation uses, via the spectral broadening parameters of the molecule, previous knowledge in order to model the precise shape of the molecular absorption signal contained in the raw signal. This modelling also connects the measured gas pressure and temperature with the spectral broadening parameters to the modelled line shape and line area. These links are therefore based on the dependence of the line shape, width and position of the used absorption line on changes in the physicochemical boundary conditions such as gas pressure and temperature as well as composition of the gas matrix. As soon as sufficient information on the boundary conditions in the measurement zone is available it is also possible to invert those relations and not only determine *absorber concentration* via laser absorption spectroscopy, i.e. dTDLAS, but also *gas temperature* (Teichert, 2003; Yang et al., 2011) or – as described here – also the *gas pressure*.

2.3.1 Line-broadening effects

The width of the absorption line, in our case the H₂O transition (000–101, 110–211) at 1370 nm (7299.4 cm⁻¹), is influenced by gas temperature, gas pressure and the matrix composition. For atmospheric applications the matrix changes can be ignored in most

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cases as the basic air composition remains constant. Regarding the individual broadening contributions, natural, Doppler and collisional broadening have to be considered. Natural line broadening is too small to be noted in tropospheric signals and thus can be neglected at these conditions. The influence of temperature-induced broadening by the Doppler-effect arises from the absorber movement because each molecule, depending on its velocity of movement towards the light, is penetrated in its inertial system by a small Doppler-shifted light frequency. Therefore, this velocity (and hence, the frequency) is Maxwell–Boltzmann-distributed and can be described by a Gaussian shape function with the width γ_{Doppler} as follows:

$$\gamma_{\text{Doppler}} = \left(\frac{v_0}{c}\right) \sqrt{\frac{2kT \ln 2}{m}} = 3.581 \times 10^{-7} v_0 \sqrt{\frac{T}{M}} \quad (3)$$

If the gas temperature is measured in the experiment with sufficient accuracy, Doppler broadening can be calculated directly and thus does not have to be fitted.

The collisional broadening, which mainly describes the pressure influence, i.e. the interaction between the absorber molecule and its collision partners, is to be discerned with regard to collisions between absorber species (leading to *self*-broadening) or “mixed” absorber-matrix-species collisions (leading to *foreign* broadening). This process is also strongly affected by the complexity (Peach, 1981), type (long/short-range forces) and strength of the interaction and the quantum state of both collision partners. Collisional broadening coefficients thus typically have to be determined experimentally. The commonly used line shape profile is the Lorentzian shape function. The pressure and temperature dependence of the collisional half width can be empirically expressed as

$$\gamma_{\text{Lorentz}} = \left(\gamma_{\text{L self}}^0 \cdot p_{\text{self}} + \gamma_{\text{L foreign}}^0 \cdot p_{\text{foreign}}\right) \frac{1}{p_0} \left(\frac{T_0}{T}\right)^n \quad (4)$$

The broadening coefficients γ ($\gamma_{\text{L self}}^0$ representing self-broadening, and $\gamma_{\text{L foreign}}^0$ for foreign broadening) are obtained from databases such as HITRAN (Rothman et al., 2009)

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about $0.1\text{--}0.02\text{ cm}^{-1}$ and for γ_{Doppler} 0.01 cm^{-1} in typical atmospheric conditions (200–1000 hPa). Strict minimization of the degrees of freedom of the fit is an issue, in particular for low signal-to-noise spectra, which can strongly influence the fit stability and reliability. Thus, with respect to the objective to measure pressure (and concentration) on fast airborne platforms, we did not use HoLS since they would only be interesting for the lowest targeted atmospheric pressure range, where the signal-to-noise ratio is, due to the associated low H_2O concentrations, also low.

2.4 Pressure determination using TDLAS

Many TDLAS studies on pressure broadening (e.g. Ortwein et al., 2010; Pustogov et al., 1994; Vorsa et al., 2005; Giesen et al., 1992) are available in the literature. These use in principle the inverse method for pressure determination via TDLAS (e.g. Brown et al., 2010).

Figure 2 shows – at constant temperature and humidity – the above-discussed Voigt line shape profiles for different pressures. The pressure-induced line broadening clearly offers the possibility to determine the total pressure from the measured line broadening. The apparent problem of the algebraically missing solution for the deconvoluted Voigt function can be solved by a precalculation of the Doppler component with the measured gas temperature as described in Eq. (3) and fitting the Voigt line shape to the measurement data by using the Voigt approximation (Eq. 5) to obtain the Lorentzian width γ_{Lorentz} .

The partial pressures in the equation for the Lorentzian width Eq. (4) can be replaced by the water concentration combined with the total gas pressure. Thereby, the total gas pressure p becomes separable (Eq. 6)

$$\gamma_{\text{Lorentz}} = \left(\gamma_{\text{L H}_2\text{O self}}^0 \cdot C_{\text{H}_2\text{O}} + \gamma_{\text{L air foreign}}^0 \cdot (1 - C_{\text{H}_2\text{O}}) \right) \cdot \frac{p}{p_0} \cdot \left(\frac{T_0}{T} \right)^n \quad (6)$$

and can thus be isolated Eq. (7)

$$\rho = \frac{\gamma_{\text{Lorentz}} \cdot p_0}{\left(\gamma_{\text{L H}_2\text{O self}}^0 \cdot C_{\text{H}_2\text{O}} + \gamma_{\text{L air foreign}}^0 \cdot (1 - C_{\text{H}_2\text{O}}) \right) \cdot \left(\frac{T_0}{T} \right)^n} \quad (7)$$

Equation 7 contributes to the following discussion: (A) the Lorentzian width γ_{Lorentz} can be determined (as described above) using the fitting process during absorption profile evaluation. Again, the dynamic tuning coefficient of the laser $\frac{d\nu}{dt}$ is needed to transfer the raw scan from time space to wavenumber space. (B) The foreign broadening coefficient ($\gamma_{\text{L air foreign}}$) as well as the self-broadening coefficients ($\gamma_{\text{L H}_2\text{O self}}$), are molecule-specific parameters and can be measured in an independent lab experiment, or taken from HITRAN[13], but with relatively large uncertainties of 5 % and even 10 % for foreign and self-broadening, respectively. Since the foreign broadening coefficient has a significantly larger influence we recently re-measured it with improved uncertainty (HWHM $0.1077 \text{ cm atm}^{-1} \pm 2.5 \%$) at the PTB national humidity standard (Heinonen, 2002). (C) The influence of the concentration $C_{\text{H}_2\text{O}}$ on the optical pressure measurement has to be critically discussed, because conversely, for calibration-free concentration evaluation (Eq. 2), the absolute gas pressure is required. For typical atmospheric water vapour concentrations ranging from single-digit ppmv to approximately 45 000 ppmv (at -70 to $+40^\circ\text{C}$), the relative influence of self-broadening is strongly suppressed under most atmospheric sections (Fig. 3, left) and certainly well below 10 %. Figure 3 (right) shows for better visualisation a typical smoothed atmospheric water vapour profile. The pressure on the x axis is directly correlated to the flight height and, finally, with the altitude above the ground. Hence, even when the pressure is unknown, an iterative curve-fitting process can be done by neglecting the self-broadening ($C_{\text{H}_2\text{O}} = 0$) at the first iteration. Using this pressure value, the first calibration-free $C_{\text{H}_2\text{O}}$ evaluation iteration can be executed, to obtain the maximum 10%-deviated $C_{\text{H}_2\text{O}}$ values (Eq. 2). These values can be used in the second iteration to reduce the maximum total gas pressure deviation to 1 % by neglecting all other deviation contributions.

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An alternative way which avoids the iterative process can be described as follows: Eq. (2) can be written as: $\rho \cdot c_{\text{H}_2\text{O}} = \frac{k_{\text{B}} \cdot T}{S(T) \cdot L} \cdot \int \ln \left(\frac{I(\nu) - E(t)}{I_0(\nu) \cdot \text{Tr}(t)} \right) \frac{d\nu}{dt} dt$, where the line area $\int \ln \left(\frac{I(\nu) - E(t)}{I_0(\nu) \cdot \text{Tr}(t)} \right) \frac{d\nu}{dt} dt$ can be evaluated directly from the fitting process.

$$\text{Equation (6) can thus be rewritten as: } \rho = \frac{\gamma_{\text{Lorentz}}^0 \cdot \rho_0 \cdot \left(\frac{T}{T_0} \right)^n - (\gamma_{\text{L}}^0 \text{H}_2\text{O self} + \gamma_{\text{L}}^0 \text{air foreign}) \cdot \rho \cdot c_{\text{H}_2\text{O}}}{\gamma_{\text{L}}^0 \text{air foreign}}$$

From a practical point of view the fitting process can be better controlled and monitored in the iterative procedure, since the fitting process of noisier absorption profiles from field situations measured at gas velocities of up to 900 km h^{-1} is less robust as it also has to correct for various optical disturbances.

Combining all broadening influences and applying them to typical atmospheric conditions (described by pressure, temperature and concentration profiles vs. height), the simulated profiles shown in Fig. 4 for the four selected height levels (marked “1” to “4” in Figs. 3 and 4) can be obtained. The individual pressure, temperature and self-broadening contributions to the line width are graphically illustrated in Fig. 5 for a mid-value and two extremes, which show the maximum range of the contribution, that can be found under typical atmospheric variations. These contributions are considered later in detail when estimating the uncertainty contributions to the optical pressure determination.

Crucial in this process and similar to calibration-free concentration evaluation (Sect. 2.2) is the design of the optical pressure determination procedure in a way that avoids any pressure calibration for the optical signal. An accurate pressure calibration at 900 km h^{-1} air speed would indeed be quite difficult to achieve, especially because a calibration under static conditions is very difficult to transfer to the highly dynamic, high-speed flow situation on a flying aircraft. The calibration-free optical pressure determination is therefore an elegant method to provide an independent way of validating other, built-in, classical, inflight pressure sensors in the measurement region.

Generally it can be stated that water vapour is certainly a suboptimal sensor species for pressure determination owing to its very high atmospheric variability (more than

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four orders of magnitude) and its very low signal levels in and above the tropopause (compare Fig. 3). Other more suitable molecules like CO_2 , CH_4 etc. show negligible self-broadening effects and much weaker changes in the SNR (signal-to-noise ratio), e.g. the atmospheric CO_2 fluctuations are much smaller (only double-digit percentage range at a basis concentration of about 400 ppmv). However, the primary goal of the HAI open-path sensor (Buchholz et al., 2013) is the realisation of a calibration-free, multi-phase water vapour measurement, for which the built-in micro-mechanical pressure transmitter has to be validated to stabilise and accelerate HAI's fitting process. Thus an optical, water-based pressure determination is more an additional benefit of the HAI instrument, which comes at very little extra cost, but which significantly improves HAI's performance and robustness. With knowledge of the correct and validated total gas pressure, a spectrally stabilised spectrometer such as the HAI can precalculate pressure-dependent spectral parameters such as the pressure shift of the absorption line position as well as the line broadening and provide them as predetermined parameters, thereby significantly minimizing the degrees of freedom and thus the stability and robustness as well as the accuracy of the spectral fitting process. Ultimately the degrees of freedom for the absorption line could be reduced to one final value, namely the line area. This procedure may be executed only when the pressure transmitter data adequately matches with the spectroscopically "sensed" pressure.

2.5 Uncertainty consideration

For measurements under harsh conditions and with complex flow dynamics (e.g. on the aircraft fuselage), it is difficult to assign a measurement uncertainty in the metrological sense (Joint Committee for Guides in Metrology (JCGM), 2008). Nevertheless, the uncertainty is estimated for the purpose of classifying the accuracy of the final measurands (and always stated as 1 sigma values).

First, the measurement by itself should be evaluated under laboratory conditions, followed by estimates of the additional influences when the measurements are performed on an aircraft. To estimate the total uncertainty of the "optical" pressure, which

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is determined via Eq. (7), we take into account the error in the self-broadening coefficient ($\gamma_{L, \text{H}_2\text{O self}}^0$), which is taken from the HITRAN08 database (Rothman et al., 2009) with 10 % uncertainty (1σ). The temperature coefficient (n) was measured by our group within 5 % (Hunsmann et al., 2006), and the air broadening coefficient ($\gamma_{L, \text{air foreign}}$) is obtained from unpublished measurements at the PTB with an uncertainty of 2.5 %.

For the uncertainty calculation we assume a water vapour mixture fraction of 2000 ppmv, which is determined with an uncertainty of 5 %, and a gas temperature of 273 K determined with 1 K uncertainty, which realistically can be achieved under quasi-static conditions. Finally, the collisional width ($\gamma_{L, \text{Lorentz}}$), was set to 0.12 cm^{-1} FWHM (equivalent to 560 hPa) with an uncertainty of 2 %, which includes a conservative uncertainty estimate for the fitting process and the uncertainty of the dynamic tuning, which has to be measured to evaluate the laser signals. All these contributions result under laboratory conditions in a total uncertainty in the optical pressure determination of 3.2 %. The largest relative contribution to the total uncertainty is caused by the uncertainty of the air foreign broadening coefficient (58 % relative), followed by the measurement of the collisional width $\gamma_{L, \text{Lorentz}}$ (38 % relative).

To transfer this uncertainty estimate to the inflight situation on the aircraft's fuselage, it is necessary to adapt the uncertainties of the measurands to the inflight situation. The gas temperature uncertainty is then assumed to be 7 K, which is derived as a quite conservative estimate from a CFD simulation of the pressure and temperature field within the open-path cell. The uncertainty for the width determination ($\gamma_{L, \text{Lorentz}}$) is doubled in light of the harsh environments in the high-speed flow plus broadband light scattering or radiative detector offsets caused by the sun. In total this results in an overall uncertainty of 5.1 % for the optical pressure determination, with the largest contribution coming from the width measurement $\gamma_{L, \text{Lorentz}}$ (61 %) followed by the uncertainty of the air broadening coefficient (23 %) and the temperature measurement (14 %). This estimate is realistic for absorption profiles such as in Fig. 7 (discussed later in detail), where the SNR is quite high. Clearly, during flight tracks with very strong optical disturbances or

in general at very low H_2O concentrations the width uncertainty increases owing to the reduction of the SNR.

3 Installing the open-path cell on HALO

The “HAI” multiphase hygrometer was successfully tested on the HALO research aircraft during the first scientific HALO mission in the second and third quarter of 2012. The aim of this science campaign, called TACTS (Transport and Composition in the UT/LMS), was to study the seasonality in the UTLS (Engel et al., 2013). Additionally, HAI was part of the scientific payload during the ESMVal (Earth System Model Validation) (Schlager, 2014) campaign.

The highest achievable flight level of the HALO aircraft determines the practically relevant pressure range for sensor validation, which is approximately 150 to 1000 hPa. The installation schematics (depicted in Fig. 6, top, together with a photograph of the HAI open-path cell, bottom) show the open-path sensor after being installed on the HALO fuselage. The White type (White, 1976) open-path cell has a mirror base distance of approximately 15 cm, which yields an optical path length of 4.2 m (indicated in red between the pylons in Fig. 6). The main electronics part and the HAI lasers are housed in the cabin. The Hai control unit and the open-path cell are connected via glass fibres. The air temperature within the HAI open path is measured via two platinum PT100 sensors. Certification requirements made it almost impossible to develop an isokinetic temperature sensor which protrudes in the gas flow. Hence, simulation models will be developed by the Forschungszentrum Jülich to estimate and correct the boundary layer effects acting on the surface-mounted PT100 sensors. The resulting temperature offsets (and corrections) are not as critical for the optical pressure measurement and hardly affect the pressure validation discussed in this paper in view of the uncertainty budgets mentioned above.

The Hai open-path sensor also includes its own pressure sensor based on a commercial piezo pressure transmitter (New Port Omega). This sensor is installed behind

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a rectangular interface in the open-path cell. Furthermore there is a second pressure “signal” via HALO’s static ambient avionics pressure (Giez, 2012) which is calculated based on a multidimensional pressure determination at the aircraft’s noseboom (Giez, 2012). Thus there are in total three independent pressure values which can be compared: the optical TDLAS and the piezo sensor within HAI and the HALO avionic system.

The high complexity of the HAI multiphase hygrometer does not allow a comprehensive description in this paper. Therefore, only the components relevant to the spectroscopic pressure measurement have been explained (Fig. 1). A full-length description of the HAI instrument is planned and will also be focusing on multi-phase H₂O measurements.

4 Results

4.1 Signal evaluation

A typical TDLAS absorption signal profile after baseline, offset and transmission correction is shown in Fig. 7 (top) for a water vapour concentration of approximately 1900 ppm at a temperature of 1 °C and 440 hPa pressure. Here, the DFB laser is scanned over the H₂O transition with a repetition rate of 240 Hz. The detector signal after the absorption path in the open-path HAI cell is digitised at 480 ksamples s⁻¹ and 14-bit resolution. Before further evaluation fifty individual raw absorption profiles are pre-averaged resulting in an effective time resolution of 210 ms. This averaged signal is fitted to the above-described Voigt profile to obtain the effective collisional contribution γ_{Lorentz} of the total line width (Eq. 6), which is then used to determine the pressure, as described above. Figure 7 (bottom) shows the residual between measurement and model as determined by the fitting process. (It has to be noted that – during this measurements – the gas velocity through the open-path White cell is approximately 900 kmh⁻¹.) The typical Allan variance evaluation of system stability is not possible for HAI owing to the

for evaluation of the HAI data. This was the main reason for developing an integrated, optical, open-path pressure monitoring system and to compare it with other means of dynamic pressure measurements.

In the following we compare the pressure signals in two typical situations: *disturbed* and *undisturbed*. Figure 8 shows a typical case of *undisturbed* superposition of the *three* pressure signals, captured during a descent of the HALO aircraft. This descent covers variations of a factor of four in pressure and a factor of 100 in H₂O concentration. The MMP transmitter (black line) is located (see rectangular pressure transmitter interface in Fig. 6) close to the optical measurement volume defined by the open light path in the HAI White cell. The static ambient avionic pressure (SAP, red line) is derived from HALO's avionic system, measured at the noseboom in front of the airplane and directly processed via HALO's main data acquisition to provide the static ambient pressure. Finally, there is the spectroscopy-based TDLAS pressure (TP, blue line) which is discussed in this paper.

At first glance all three sensors pretty much coincide. Looking closer, the sensors show consistent and systematic discrepancies, with the SAP being always high, HAI's MMP being always low (averaged difference SAP – MMP = 28 hPa) and the optical TP being all the time in between. These deviations have a simple physical/fluid-dynamical explanation: since the air masses are accelerated in between the HAI pylons (see Fig. 6), they experience a dynamical pressure reduction relative to the ambient pressure, so that MMP (black) and TP (blue) which are “sampling” between the pylons have to be lower than the static ambient pressure (SAP, red) measured by the avionic system.

Comparing, furthermore, the two HAI pressure sensors, it has to be taken into account that the pressure measured directly by the MMP internal pressure transmitter in the open-path cell (black) has to be lower, owing to the dynamic pressure reduction caused by the airflow parallel to the surface of the MMP's pressure transmitter.

The obviously higher fluctuations of the TP are caused by a combination of several factors:

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1. Real pressure fluctuations which cannot be detected by the slow and integrating MMP pressure transmitter.
2. Fitting noise has to be considered which shows up since quite fine broadening effects have to be extracted at rather high accuracy to be used for pressure determination. This noise (9 hPa (1σ) at 220 hPa) is also a consequence of the very harsh measurement conditions, such as the high gas velocity of up to 900 km h^{-1} which also causes very fast optical transmission fluctuations. In addition the signal is impeded by fluctuating background radiance (caused by solar radiation) impinging on the open detector (see Eq. 1).
3. Further, one has to consider imperfect separation of the self-broadening effects, owing to the strong (100×) and fast H_2O concentration fluctuation (see Fig. 8, green line and right-hand scale).
4. Finally, the strong H_2O absorbance variations also lead to strong variations in the signal-to-noise ratio and hence to limitations in the fitting process.

Despite these imperfections, Fig. 8 nicely illustrates the mutual validation of all three sensors, as the small differences (averaged difference SAP-MMP = 28 hPa, averaged difference SAP-TP = 13 hPa; see also Figs. 9 and 10) can be attributed to the different measurement locations and sensor installations.

This result therefore not only justifies water vapour as a suitable target species for optical pressure determination. It also allows the validation of the optical pressure technique permanently – without any additional hardware effort (!) – while the standard evaluation of the HAI water vapour signals is maintained at any given time.

The *disturbed* intercomparison case (shown in Fig. 9) removes the apparent redundancy of the three pressure signals and explains why a “local” pressure measurement at the HAI pylons is needed, for example to recognise fluid-dynamically critical situations, which may lead to significant systematic deviations in the pressure as well as the H_2O concentration measurement derived from the HAI open-path cell.

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Similar to the figures above, Fig. 9 shows the three measured pressure values in the same colour code. On the right-hand side (at $t > 11:02$ LT) a steep descent from 200 to 800 hPa occurs, followed by four, fast, repetitive, small-scale, ascent-descent movements with a pressure amplitude of only 50 hPa (see inset Fig. 9 on the right). As in the case of the previous figure all sensors nicely overlap with average deviation over the whole time of only within one percent (9 hPa at about 830 hPa).

The left-hand side of Fig. 9 ($t = 10:32$ to $11:02$) shows, in contrast to Fig. 8, on two occasions a significant discrepancy between the SAP and both HAI pressure sensors. These strong, several-minutes-long, block-shaped pressure deviations with a relative magnitude of 30 % to SAP ($\Delta p =$ approximately 70 hPa) are not visible by the SAP. However, MMP and TP agree in this situation quite well with regard to the amplitude as well as the phase of this pressure excursion (to better than 5 %). If this deviation would be measured by only MMP it would be very difficult to exclude any malfunction, e.g. due to ice accumulation in the MMP interface. The excellent match between MMP and TP, however, falsifies this assumption and asks for another explanation than a malfunction of the MMP sensor.

This explanation can be found looking at additional aircraft parameters which influence the flow field around HALO. Also shown in Fig. 9 are the travelling speed of HALO (= Mach number, orange line and scale on the left) and the angle between the atmospheric fluid velocity vector and the wing surface (= angle of attack, green line and scale). As depicted the strong pressure excursions correlate in an almost “binary” way with the crossing of a minimum Mach number or a maximum angle of attack (dashed lines).

Obviously the flow around the HAI sensor is obstructed or stalled at a certain angle of attack/speed of the aircraft, so that the pressure in between the HAI pylons changes abruptly. With a fully detailed model of the flow around the aircraft and its external component parts, these pressure effects could also be simulated and superimposed on the avionic pressure signal. But these calculations would have to be recomputed for every flight parameter set (flight speed, HALO orientation), thus involving intensive

computation over several hours of flight, which makes this a slow and unattractive approach. Therefore, the possibility of validating the pressure transmitter in an open-path cell with the same raw data for final humidity evaluation allows a much easier clarification and verification of malfunctions (if there are any), thus allowing easy but advanced data quality management.

Finally (see Fig. 10) we compare the HAI pressure sensors by directly correlating the data from Fig. 9. The pressure detected by the MMP, the internal pressure transmitter in the open-path cell, and the TP, the spectroscopically determined pressure, show an excellent agreement which falls very close to the 1 : 1 bisector line. A linear fit is made to elucidate the correlation quantitatively and yields a very small slope deviation of only 0.14 % from the ideal bisector slope. This demonstrates nicely the good agreement and the linearity of the two sensors. The correlation of both sensors also falls well within the $\pm 5\%$ corridor defined by the overall TP sensor uncertainty of 5.1 %. The correlation plot also shows nicely that most of the individual data points over a wide pressure – and hence H_2O concentration – range match well. Only at quite low pressures, where the SNR dropped significantly, a weakened correlation, caused by the dynamic changes in the flight pattern at the high flight levels, appears. Figure 10 also clearly shows the transition regions resulting from the pressure drop at the edges as a dynamic effect, and the higher noise level caused by the low water vapour concentration (approximately 150 ppmv (Fig. 8)). Even fine details can be found: the offset of 9.5 hPa extracted from the slope fit can be interpreted as a consequence of the air mass movement towards the rectangular pressure transmitter interface, which leads – as mentioned above – to a local pressure reduction. Perspectively, this offset will be an interesting parameter to be checked when an entire CFD model becomes available. Overall Figs. 8–10 nicely allow a mutual validation of the two HAI pressure sensors with each other and during suitable conditions also of the TP and the MMP with the avionic pressure data within a few percent.

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5 Conclusion and outlook

Highly accurate pressure measurements on research aircraft are quite difficult to realise because of the complex flow dynamics and their dependency on several other airborne flight parameters. To solve many flight issues, an accurate knowledge of the static ambient pressure is required. Static pressure sensors are typically available with a high level of accuracy. However, the determination of the exact pressure at a specific location on the fuselage of an aircraft requires computationally intensive simulations (using CFD models) for every flight parameter set.

In this paper, we presented an integrated approach to a purely spectroscopic, calibration-free pressure determination using water vapour as the sensor species and HAI as the sensing instrument. The HAI instrument is initially designed for multiphase water vapour detection and serves here as an additional TDLAS-based gas pressure sensor in the light path region of the HAI open-path White cell on the aircraft fuselage. The optical pressure measurement was designed to be calibration-free and is purely based on the knowledge of the relevant spectral parameters of the sensor molecule describing the line broadening. The correction process, which has to be executed to obtain accurate pressure values, has been described and takes self-broadening and temperature dependence of the broadening parameters into account. For the spectroscopic pressure detection using HAI we determine – despite the high atmospheric water vapour variability – a pressure uncertainty of 3.2 % for laboratory conditions and 5.1 % under flight operating conditions in the field.

The spectroscopic pressure determination was compared in the range of 150–800 hPa with (a) a HAI-internal micro-mechanical pressure sensor and (b) the static ambient pressure measurements of the aircraft avionics. The linearity deviation between MMP and TP was 0.14 %. A small dynamically induced pressure offset of 9.5 hPa was found.

Occasional dynamic deviations from the static ambient avionic pressure confirm the need for permanent CFD simulations under all flight conditions for validating the

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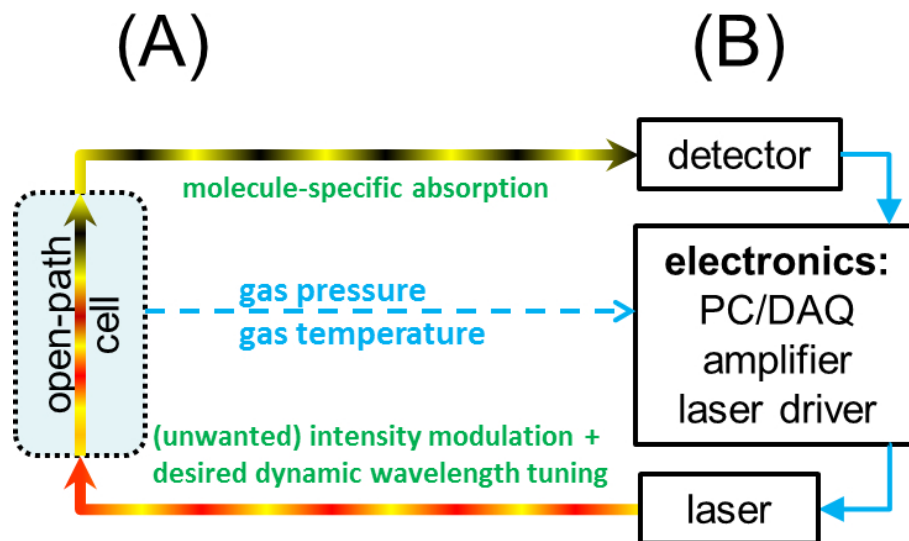


Fig. 1. TDLAS principle and basic setup: the laser current modulation at typically 100 Hz–10 kHz initiates a rapid wavelength tuning in combination with a strong intensity modulation of the diode laser radiation. The molecule-specific light loss caused by the absorbers within the open-path cell is captured for each individual wavelength scan by a photo detector, digitised, and saved as raw data for subsequent data analysis.

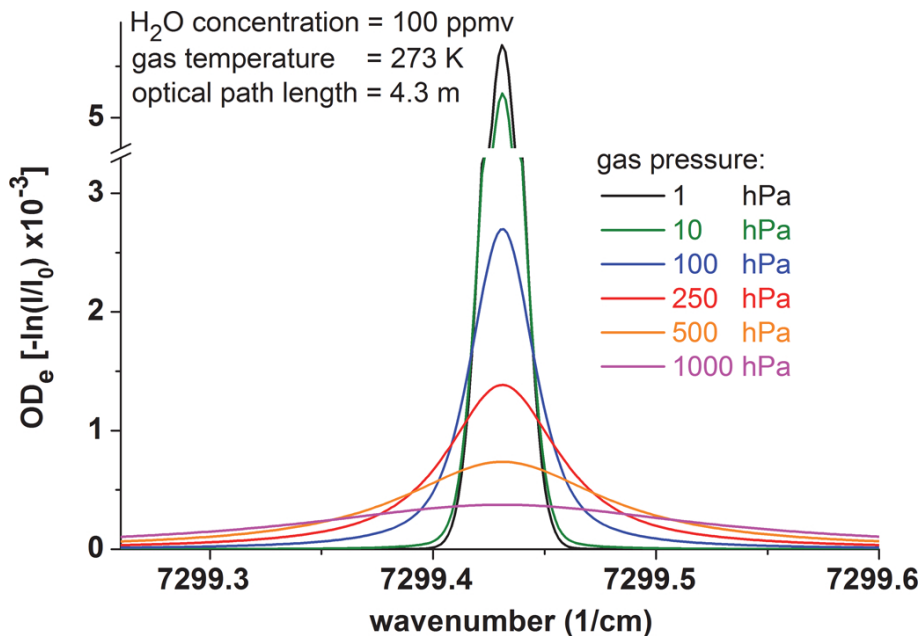


Fig. 2. Simulated absorption profiles of the used (000–101, 110–211) water vapour transition at 1370 nm (7299.4 cm^{-1}) at 100 ppmv water vapour for the atmospheric pressure range between ground level and the mid-stratosphere. The line width is mainly influenced by the collisional, foreign broadening by air, inducing a Lorentzian line shape for gas pressures above 100 hPa. At the lowest pressures of 1 and 10 hPa the line shape (Gaussian) is mainly influenced by Doppler broadening.

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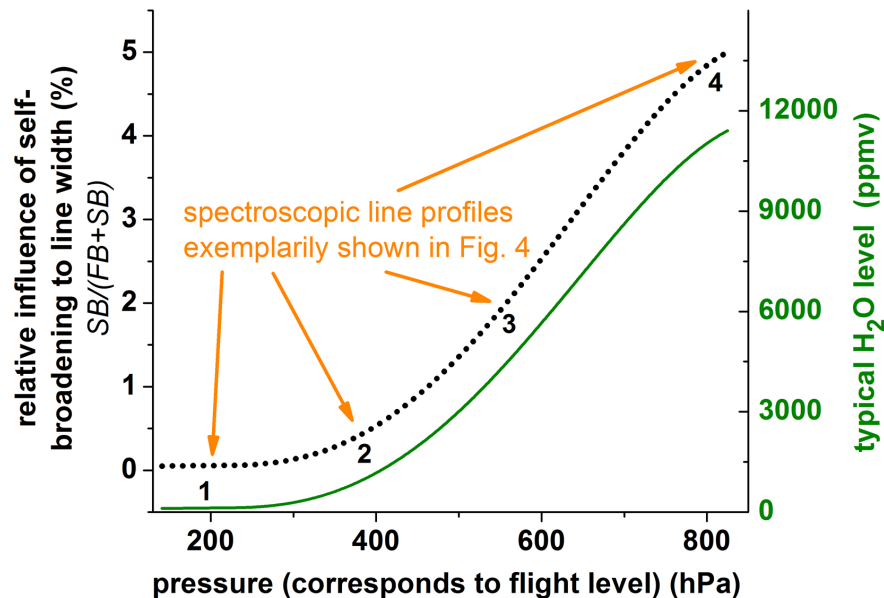


Fig. 3. Typical atmospheric vertical water vapour concentration profile (green solid line) vs. total pressure. The total pressure is usually the preferred indicator for flight height. Also shown (black broken line) is the height dependence of the relative contribution (in percent) of self-broadening to the total line width for the used water transition (1370 nm). This shows that self-broadening correction is absolutely essential for precise spectroscopic pressure measurement as the self-broadening can contribute up to 5% to the total width. Exemplary absorption profiles at height levels indicated by numbers 1 to 4 are depicted in Fig. 4.

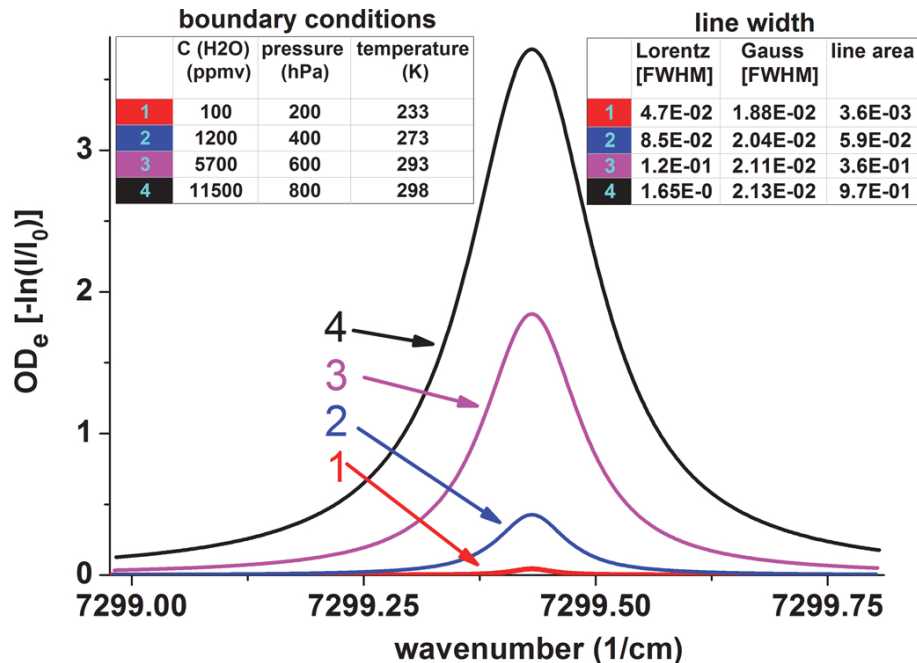


Fig. 4. Simulated atmospheric H₂O absorption profiles at positions 1 to 4 in Fig. 3 illustrating the effect of (details see text) temperature and foreign- and self-broadening on the used water transition (1370 nm) for a typical vertical atmospheric height profile. Parameter sets (T , p , C) of the individual lines and the resulting line widths are also shown. All line shapes are based on the same Voigt profile approximation, which is also used to pursue the fitting process.

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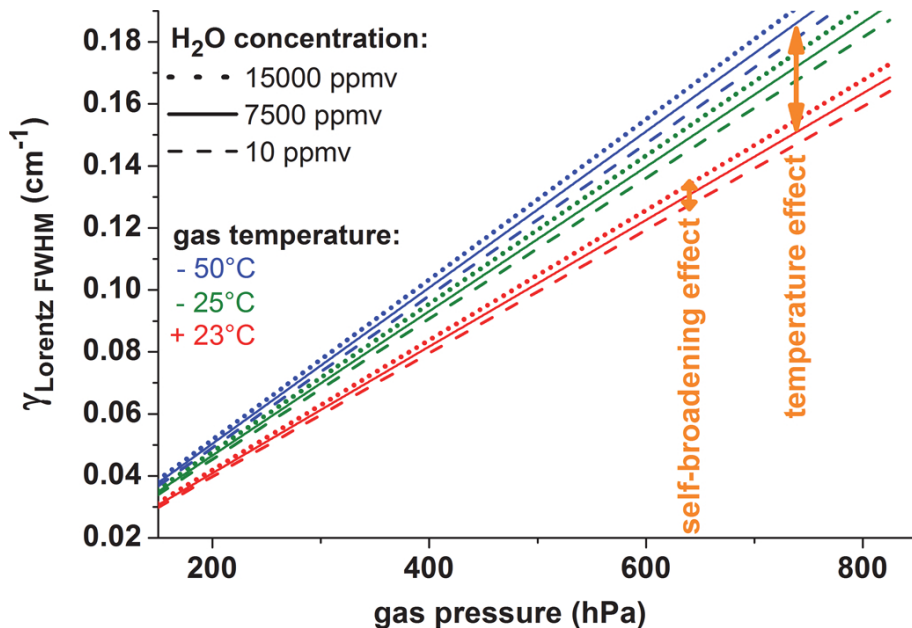


Fig. 5. Collisional line widths γ_{Lorentz} induced by foreign broadening (of the used 1370 nm transition) as a function of flight height (scaled as total pressure) for three atmospheric temperatures (-50°C , -25°C , $+25^\circ\text{C}$) indicating the strong pressure and temperature dependence of the collisional width. In addition the influence of self-broadening on the line width is shown for high, mid and low H_2O concentrations (dashed lines). A clear hierarchy becomes visible with the largest impact from pressure, then temperature, then matrix composition. Thus the line width can be used for optical pressure determination if the latter two can be compensated, i.e. if concentration (self-broadening) and temperature effects are corrected for.

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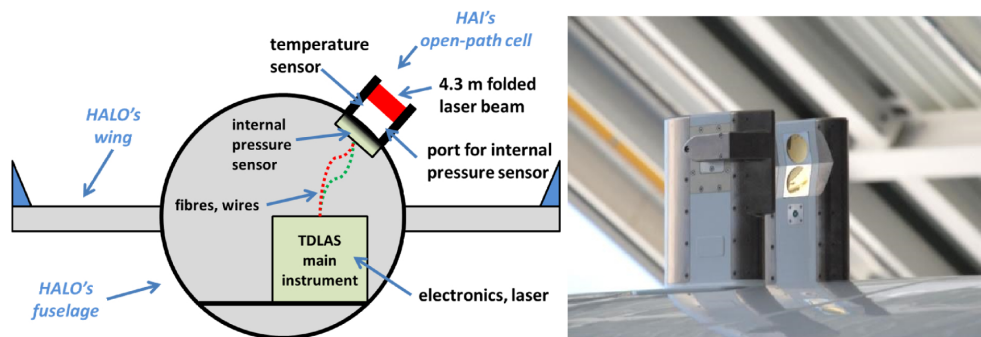


Fig. 6. Left: schematics of the HAI open-path sensor installed on the HALO research aircraft fuselage. The open-path cell itself comprises two platinum PT100 sensors for air temperature measurements for use in data evaluation. Our HAI pressure measurement interface port allows comparison between the “spectroscopically determined” pressure and a standard pressure transmitter. Additionally, several pressure sensors are installed in the noseboom of the HALO aircraft. Right: photo of the installed, open-path TDLAS sensor. At the rear pylon two adjacent, gold-coated, circular mirrors of the White cell are visible as well as one air temperature sensor below the mirrors.

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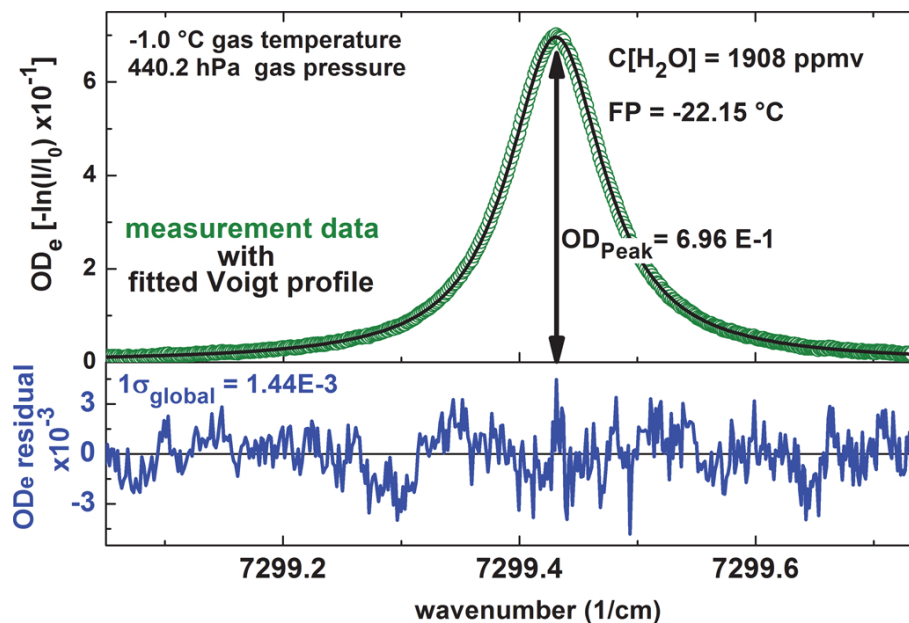


Fig. 7. Typical, pre-processed, in-flight line profile measurement after baseline-, offset- and transmission-correction. The DFB diode laser scans across the H_2O transition as shown with 240 Hz repetition rate; 50 individual raw scans are pre-averaged and the resulting average scan is then fitted, resulting in 210 ms time resolution. The global residual variance $1\sigma_{global}$ of the fit is defined as standard deviation of the residual over shown entire fit range. It has to be kept in mind that the gas flow is passing through the open-path cell at approximately 900 km h^{-1} (more details see text).

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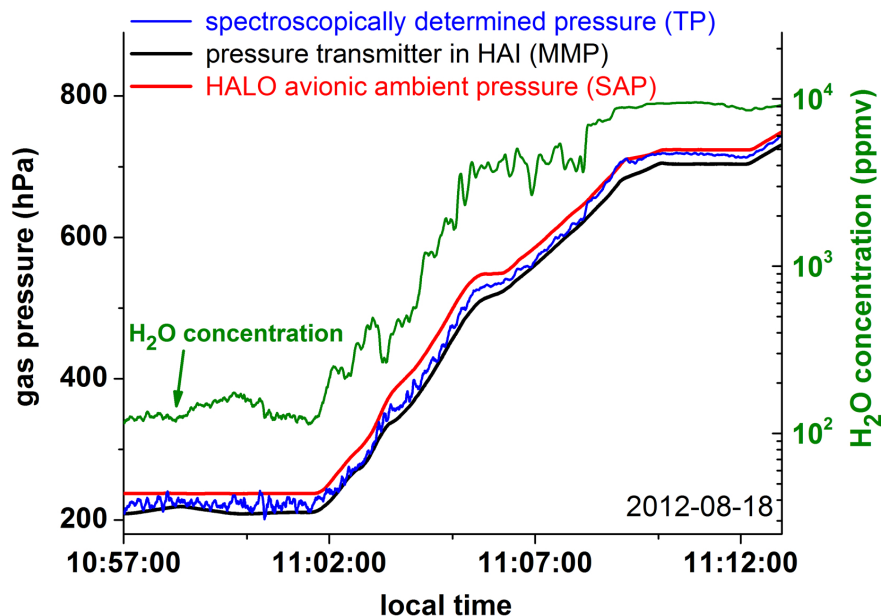


Fig. 8. Superposition of all three pressure measurands (bottom traces, left scale) as well as the measured water vapour profile (top trace and right scale) vs. time during an aircraft descent from 200 hPa to 800 hPa. Despite the micro-mechanical HAI pressure transmitter (MMP, black) being placed very close to the open-path measurement volume, clear discrepancies to the other pressure sensors are visible. These are attributed to the high-speed flow around the HAI sensor and especially towards the rectangular pressure transmitter interface (see Fig. 6), which can cause complex dynamic effects depending on the HAI orientation in the flow. The red curve shows static ambient pressure measurements (SAP) from HALO’s avionic sensors (Giez, 2012), located “in front” of Halo in the noseboom. The typical large atmospheric H₂O concentration variations (log scale!) are visible, indicating the necessity for detailed corrections of the optical pressure measurement (explained in detail in text).

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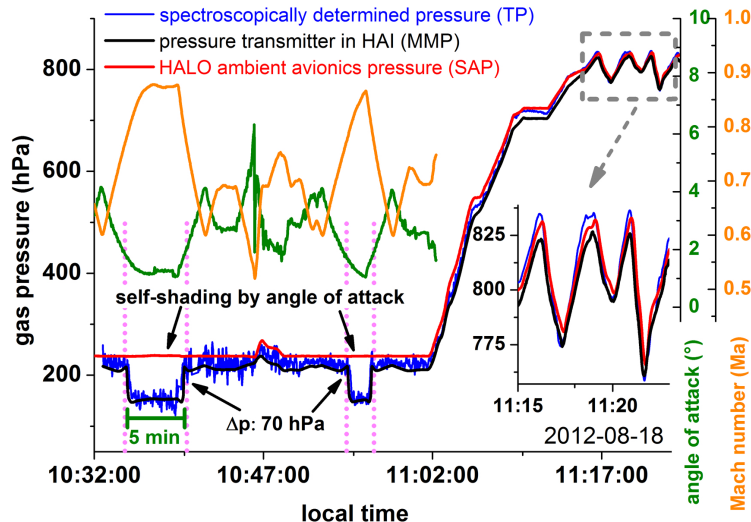


Fig. 9. Intercomparison of static ambient avionic pressure (SAP, red), micro-mechanical HAI pressure (MMP, black) and spectroscopic TDLAS pressure (TP blue, all left pressure scale) during certain HALO high-altitude flight manoeuvres and a subsequent descent. A clear incidence of HALO's aerodynamic influence of the effective pressure within the open-path volume of HAI, leading to a strong deviation between the SAP and the HAI pressures (MMP/TP), can be seen on the left between 10:30 and 11:00. The block-shaped pressure deviations of 70 hPa (= 30% relative Δp) are caused by self-shading of HAI during a HALO manoeuvre caused by a change in the angle of attack (green line, right scale in green) and velocity (orange line and scale on right) of the aircraft. (The resulting constraints in using HALO's avionic pressure measurement are discussed in the text.) At lower flight heights and less critical manoeuvres ($t = 11:20$), all three pressure signals agree within 1% (9 hPa at about 830 hPa) (see inset on the right).

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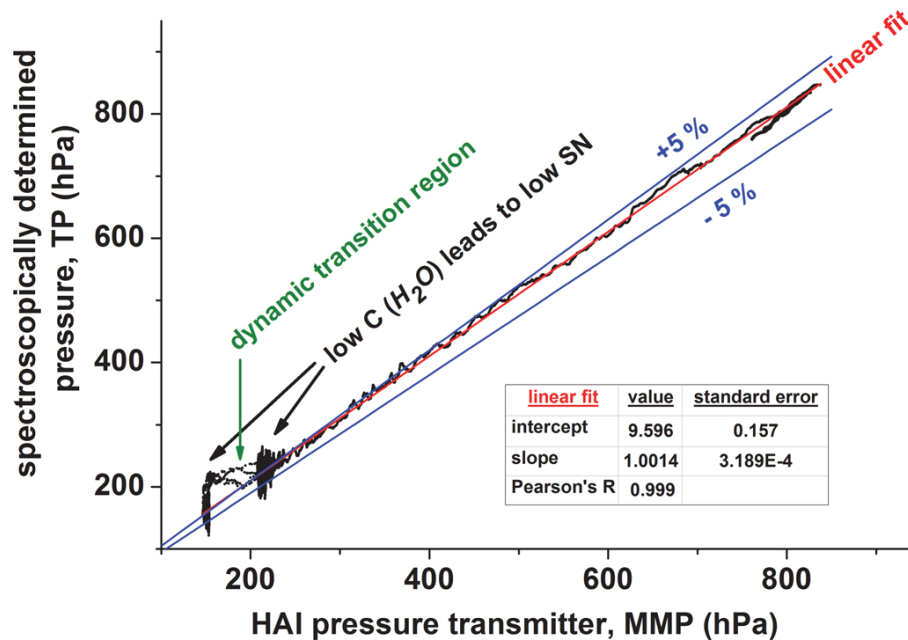


Fig. 10. Correlation plot between HAI's micro-mechanical pressure transmitter (MMP) and the spectroscopic TDLAS pressure measurement (TP). Overall, the correlation can be well represented with a linear fit (red line) with a slope of 1.0014 and $R = 0.9992$. The complex flow dynamic effects (e.g. Fig. 8) have some influence on the correlation, particularly at lower pressure, and can lead to self-shading effects (green-marked, see Fig. 9) in the dynamic transition region. The higher noise level at low pressures results from the strong H_2O decrease at higher flight heights.

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