HAI –
A new, airborne, absolute, twin dual-channel, multi-phase TDLAS-hygrometer:
background – design – setup – first flight data

B. Buchholz1,2,5, A. Afchine1, A. Klein1, C. Schiller4, M. Krämer4, V. Ebert1,2,3
1 Physikalisch-Technische Bundesanstalt Braunschweig, Germany
2 Physikalisch Chemisches Institut, Universität Heidelberg, Germany
3 Center of Smart Interfaces, Technische Universität Darmstadt, Germany
4 Forschungszentrum Jülich, IEK-7, Germany
†deceased
Corresponding author: volker.ebert@ptb.de

Abstract

The novel Hygrometer for Atmospheric Investigations (HAI) realizes a unique concept for simultaneous
gas-phase and total (gas-phase + evaporated cloud particles) water measurements. It has been developed
and successfully deployed for the first time on the German HALO research aircraft. This new instrument
combines direct Tunable Diode Laser Absorption Spectroscopy (dTDLAS) with a first-principle evaluation
method to allow absolute water vapor measurements without any initial or repetitive sensor calibration
using a reference gas or a reference humidity generator. HAI contains two completely independent dual-
channel (closed-path, open-path) spectrometers, one at 1.4 µm and one at 2.6 µm, which allow together to
cover the entire atmospheric H2O range from 1 to 40 000 ppmv with a single instrument. Both spectrometers
comprise each a separate, wavelength-individual extractive, closed-path cell for total water (ice and gas-
phase) measurements. Additionally, both spectrometers couple light into a common, open-path-cell outside
of the aircraft fuselage for a direct, sampling-free and contactless determination of the gas-phase water con-
tent. This novel twin dual-channel setup allows for the first time multiple self-validation functions i.e. in
particular a reliable, direct, in-flight validation of the open-path channels. During the first field campaigns,
the in-flight deviations between the independent and calibration-free channels (i.e. closed-path to closed-
path and open-path to closed-path) were on average in the 2% range. Further, the fully autonomous HAI
hygrometer allows measurements up to 240 Hz with a minimal integration time of 1.4 ms. The best preci-
sion is achieved by the 1.4 µm closed-path cell at 3.8 Hz (0.18 ppmv) and by the 2.6 µm closed-path cell at
13 Hz (0.055 ppmv). The requirements, design, operation principle and first in-flight performance of the
hygrometer are described and discussed in this work.

1. Introduction

1.1. Atmospheric water vapor

Water vapor is in many ways one of the most important measurand for atmospheric investigations
(Ludlam, 1980; Möller et al., 2011; Ravishankara, 2012). Water vapor is the most important greenhouse gas
(Kiehl and Trenberth, 1997), and known as a key atmospheric coupling element of almost all microscopic
(e.g. droplets/ice crystals formation), macroscopic (e.g. clouds/precipitation) and global processes (e.g. hy-
drological cycle). It is strongly related to numerous highly relevant topics of atmospheric science and close-
ly related to “climate change” (Held and Soden, 2000; Houghton, 2009; Kiehl and Trenberth, 1997; Maycock
et al., 2011). Unsurprisingly, numerous water vapor studies have been carried out targeting its atmospheric
trends and variability (Lu and Takle, 2010; McCarthy et al., 2009; Ross and Elliott, 1996; Scherer et al., 2008; Trenberth et al., 2005; Xie et al., 2011), its influence on transport models (Kiemle et al., 2012; Schäfler et al., 2010), or its impact on radiation balance models (Lockwood, 1990; Ramanathan et al., 1989; Schneider, 1972). The high complexity of atmospheric water vapor is linked to its occurrence in all three phases which is unique among most atmospheric molecules. Water in the gas-phase is a very strong infrared absorber and significantly impacts atmospheric energy fluxes through latent heat transfers by the different phase transitions. Condensation to the liquid phase or freezing to solid particles leads to effective scattering of solar radiation, which directly raises links to the formation process of cirrus clouds (Krämer et al., 2009; Spichtinger et al., 2004). These few relationships show the complexity from a theoretical as well as a modeling point of view. Today, however, the quality (particularly accuracy and comparability) of atmospheric water measurements frequently limit a better understanding of key atmospheric processes (Krämer et al., 2009; Peter et al., 2006; Scherer et al., 2008; Sherwood et al., 2014). Despite the outlined importance and the large effort invested in the developments of hygrometers in recent years, water vapor remains a target molecule that is very difficult to measure accurately.

Several major issues exacerbate airborne water vapor measurements. Atmospheric water vapor encompasses a very large concentration range (3 – 40 000 ppmv). The high spatial variability of H₂O leads on fast aircraft (typically around 800 km/h cruising speed) to highly dynamic H₂O variations in airborne measurements of up to several 1000 ppmv/s in the gas-phase and several 10 000 ppmv/s for total water (gas-phase + evaporated ice/liquid phase). Airborne hygrometers thus require high time resolution, precision, and accuracy at the same time. Additionally, water vapor is very effectively absorbed from nearly any surface. This challenges in a highly complex manner not only the entire gas sampling system but also the calibration infrastructure which is typically required for most hygrometers. By waiving the entire calibration process, special laser-based hygrometers (Wolfrum et al., 2011) circumvent all calibration related issues efficiently which will be explained later in chapter 3.2. Last but not least, the measurement boundary condition such as gas pressure, gas temperature, gas velocity, instrument operation temperature changes or vibrations during airborne measurement are defined by the environment inside and outside of the aircraft itself causing a variety of complications.

1.2. Airborne Hygrometry

This brief compilation illustrates the complex challenges associated with developing airborne water vapor instruments, especially to measure at tropospheric and stratospheric atmospheric conditions. A high number of (mostly single-channel and single-H₂O-phase) hygrometers have been developed in the last decades with various advantages and drawbacks (Wiederhold, 1997). A non-exhaustive selection of instruments is described here: (Buck, 1985; Busen and Buck, 1995; Cerni, 1994; Desjardins et al., 1989; Diskin et al., 2002; Durry et al., 2008; Ebert et al., 2000b; Gurlit et al., 2005; Hansford et al., 2006; Helten et al., 1998; Hunsmann et al., 2008; Karpechko et al., 2014; Kley and Stone, 1978; May, 1998; Meyer et al., 2015; Ohtaki and Matsui, 1982; Roths and Busen, 1996; Salasama and Kostamo, 1986; Sargent et al., 2013; Schiff et al., 1994; Silver and Hovde, 1994b, 1994a; Thornberry et al., 2014; Webster et al., 2004; Zöger et al., 1999a, 1999b; Zondlo et al., 2010). Consequently, the question should be raised from the opposite point of view: What are the important and required properties to be covered and combined for a near-universal “Hygrometer for Atmospheric
Investigation” to serve as an innovative and cutting-edge tool to explore open and new scientific questions related to atmospheric water vapor?

2. General design targets for HAI (Hygrometer for Atmospheric Investigation)

2.1.1. Open-path or closed-path or both?

It is highly desirable for an airborne system to be constructed in a way that it operates on the ground in the same manner as in flight and that every change of environment and “boundary conditions” of the instrument is logged, supervised and impacts are minimized or even prevented by design. This allows extensive validation as well as laboratory comparisons with other instruments and avoids systematic, barely detectable deviations only occurring in flight. Instruments that are calibrated/validated on ground in different configurations such as (Diskin et al., 2002), or under different working conditions such as (Sargent et al., 2013) often fail to provide a clear, reliable, unbroken chain of calibrations from the in-flight measurement to a (metrological) laboratory standard, which is necessary to guarantee accuracy.

This notion related to water vapor leads directly into the everlasting discussion about sampling via open-path systems operation in the free flow versus closed-path systems extracting the air to be analyzed into the instrument. Open-path hygrometers offer numerous great benefits such as: the prevention of any sampling errors or uncertainties (caused by surface absorbing effects) as well as the high response time that is limited only by the transfer functions of optical or electrical components but not by the gas exchange rate. The latter circumvents the complicated and adulterant deconvolution of smoothing effects with time-response functions caused by a sampling system. On the other hand, the boundary conditions such as gas pressure and gas temperature as well as possible spatial inhomogeneity of both parameters are difficult to take accurately into account for an open-path system. Additionally, an airborne open-path sensor has to operate in harsh boundary conditions, i.e. over a large range of temperatures (-80 to 50 °C), pressures (70 to 1000 hPa), for large ram pressures (900 km/h gas velocity) and mechanical stress through accretion of ice or liquid water.

The major problem of all present open-path systems is their highly complex calibration, or even just validation, since realistic flight conditions, in particular the dynamics, are extremely difficult to realize in a lab with sufficient accuracy. A direct metrological link to test the in-flight performance, e.g. a dynamic calibration facility for open-path hygrometers, is therefore missing.

Closed-path systems, on the other side, are simply installed inside an air-conditioned cabin in a much more protected environment. Gas is sampled with a suitable inlet and led via a tubing system to an “internal” measurement chamber, such as an optical absorption cell or a “cavity”, e.g. for a dew point mirror hygrometer. On one hand, it is much easier to accurately control and maintain the physical boundary conditions of the sample gas, e.g. temperature, pressure, flow, etc. in the measurement volume. On the other hand, it is difficult to ensure and maintain a representative sampling process and to quantify and correct sampling related deviations. These maybe caused by adsorption and desorption effects, which occur on all surfaces of the sampling system, and have to be carefully minimized e.g. by heated (HAI =80 °C), electro-polished stainless steel sampling pipes, along with an instrument design ensuring high gas flows (HAI =100 liter/min) under all flight conditions.
Gas-phase H$_2$O measurements in clouds are often carried out via backward facing sampling inlets. But such inlets readily sample small liquid or solid water particles possibly causing systematically positive offsets. The measurement of total water, i.e. the sum of water in the gas-phase as well as in the condensed phase (ice/droplets), relies in additionally on an accurate distribution model of the particles entering the inlet system. Usually, the particle sampling characteristics are considered individually (Krämer and Afchine, 2004), and sampling corrections are done during further metrological analysis and therefore not part of an instrument description. For every H$_2$O measurement using airborne extractive (=closed-path) instruments on aircraft, it needs to be taken into account that the instruments response reflects contributions from the sensor element itself as well as the sampling system and its dynamic properties.

One indisputable, major advantage of typical extractive (closed-path) instruments is the possibility for careful tests outside of the aircraft, i.e. in a hangar or laboratory. But, to take full advantage of this, it is desirable for a sophisticated instrument to integrate supervising and monitoring functions in a way that a performance comparable to the laboratory can be ensured during any in-flight situation. This generates the great benefit of transferring the performance from laboratory to field, quite similar how a metrological transfer standard is typically used. This directly reinforces the question, how to assess the accuracy of airborne hygrometers on ground and in particular during flight operation.

2.1.2. State of the art instrument accuracy

In general, the highest measurement and preparation accuracy is realized by the validated primary standards of national metrology institutes such as PTB (Germany) or NIST (USA). The international, metrological water vapor scale is defined by traceable primary water vapor generators (Brewer et al., 2011). The mixing ratio range, required to cover the entire tropospheric and stratospheric range (3 - 40000 ppmv), is realized by a combination of generators based on different physical principles. Their uncertainty is in the order of 0.5% relative (Brewer et al., 2011; Buchholz et al., 2014a; Mackrodt, 2012). In other words, it is not possible to validate any hygrometer with a better accuracy due to the lack of a suitable accurate reference.

Comparing the available metrological accuracy to some results from field comparisons of airborne hygrometers demonstrates the large potential for improvement. For example, long-term (> 10 years) change studies of stratospheric H$_2$O (Oltmans and Hofmann, 1995; Rosenlof et al., 2001; Solomon et al., 2010) suffer from significant, difficult to quantify relative deviations between different instruments in the range of 50–100% (Fahey et al., 2014; Peter et al., 2006; Vömel et al., 2007) which recent studies such as (Rollins et al., 2014) confirm (±40–50% for < 3ppmv and ±20% for > 3ppmv). Radiosonde comparisons with polymer sensors and chilled dew point mirror hygrometer, covering the entire troposphere and lower stratosphere region such as (Miloshevich et al., 2006), show averaged overall agreements in the 10% range (but also, local deviations in the 30% range). These deviations are quite common for many airborne campaign results e.g. (Smit et al., 2014) and become even worse when focusing on the relative deviations in regions containing highly variable H$_2$O structures. Hence in 2007, an international comparison exercise, “AquaVIT” (Fahey and Gao, 2009), was organized to compare the world’s best airborne hygrometers under well-controlled, quasi-static, equivalent conditions to evaluate the accuracy under well controlled laboratory conditions, without the influence of any typical dynamic sampling effects. AquaVIT comprised 22 hygrometers (tunable diode laser spectrometers, TDL, dew or frost point mirror hygrometers, D/FPH, Lyman alpha fluorescence and absorp-
tion hygrometers, LAFH and other principles) from 17 international research groups. The instruments were categorized in well validated “core” instruments (APicT, FISH, FLASH, HWV, JLH, CFH, see (Fahey and Gao, 2009) for details) and “younger, less mature” non-core instruments. Even the core-hygrometers deviated in the important 1 to 150 ppmv WVMR range by up to ±10% from their common mean value. In other words, core instruments differed by up to 20% from each other, even under quasi-static conditions. Other, less representative and extensive comparisons such as (Hoff, 2009; Mangold and Wodca Team, 2003) yielded similar results.

2.1.3. Educated guess for accuracy deviations between state of the art hygrometers

The assessment of the required accuracy depends strongly on the purpose of the instrument and its data. In terms of commonly used climatologies or strongly averaged, coarse validation studies deviations can be acceptable, but as far as e.g. retrieval models for satellite data improve, such uncertainties and deviations can become critical. In many other cases such as the currently often discussed atmospheric super saturations (Peter et al., 2006), the instrument uncertainties prevent deeper investigations and therefore a better understanding. Reconsidering the entire situation and perceiving that after so much development effort over the past decades these deviations remain quite high, leads to the inevitable question of concealed, common impact factors. Contemplating the typical metrological efforts needed at national metrology institutes (NMI) to generate an accurately humidified gas stream (with a sub-percent uncertainty) suggests that the uncertainties generated by typical calibration processes under field conditions could be a major contribution to these hygrometers deviations found in AquaVIT and other studies. In particular, comparing the performance and strategies of lab-based, metrological and portable field calibration facilities (Friehe et al., 1986; Helten et al., 1998; Podolske et al., 2003; Smit et al., 2000; Smorgon et al., 2014; Zöger et al., 1999b) indicate three significant discrepancies: required time for calibration, frequency of calibration, and traceability of the humidity reference itself. Calibrations in low concentration ranges at NMIs take several hours up to days per individual humidity value. During airborne campaigns, however, calibrations often have to be realized (for practical reasons) in a short time, certainly less than a few hours for a large number of concentration steps often including several pressure levels, thereby taking the obvious risk that the instrument/reference is not fully stabilized or equilibrated. Ideally, the time between two calibrations should be shorter than the expected time required for a drift/change exceeding the boundaries of the instrument uncertainties. Some airborne instruments require for the same reason calibrations before and after each flight in order to interpolate between both calibrations (Zöger et al., 1999b). Some even work with in-flight calibrations (Kaufmann et al., 2016; Thornberry et al., 2013) sacrificing measurement time and shifting the accuracy issue to the necessary airborne H2O-source. Undoubtedly, many of these instruments have benefits e.g. in terms of precision, space, weight, prime cost etc., which justifies the calibration effort. However, vice versa, it is often condoned that the calibration process is hampered and turns out to be the major influence on the accuracy of such a sensor.

Lastly, it seems necessary to implement a traceable link to the metrological humidity scales to improve the overall accuracy of airborne hygrometry (Joint Committee for Guides in Metrology (JCGM), 2009). By realizing an unbroken chain of calibrations, it is possible to link the instrument performance and the metrologi-
The precision and response time are under certain circumstances reciprocally correlated to each other (Allan, 1966). Typical figures for response time of airborne hygrometers in the literature are 0.5 – 1 Hz (Petersen et al., 2010; Szakáll et al., 2004; Zöger et al., 1999b); some instruments deliver faster data 4 Hz (Weinstock et al., 1994) or special instruments up to 25 Hz (Zondlo et al., 2010). Typical precisions at 1 Hz are in 0.1-0.2 ppmv range (Sargent et al., 2013; Zöger et al., 1999b; Zondlo et al., 2010). While in the stratosphere (<10 ppm), the precision certainly can become a limiting factor, this is much less the case inside clouds or within the troposphere, where frequent, very strong spatial variations (up to 1000 ppmv per 100 m flightpath in the gas-phase, or up to 20000 ppmv per 100 m during total water phase) pose a larger problem. An instrument with a time response of just a few Hz causes significant under-sampling which can lead to strong aliasing effects at standard cruising speed of research aircraft (approx. 700-900 km/h). Important under such conditions is the instrument’s linearity and range to accurately cover the entire H₂O concentration range up to five magnitudes for total water vapor measurements without cut-offs or complex deconvolution functions.

2.1.5. Other requirements guiding the design
HAI was designed for the German HALO aircraft (HALO, 2016; Krautstrunk and Giez, 2012), a Gulfstream G500, similar to the American HIAPER (UCAR/NCAR, 2016) which is a Gulfstream G500. HALO offers broad spatial coverage (> 10 000 km), high altitudes (up to 15 km), large payloads of up to 3000 kg payload and a pressurized and air-conditioned cabin. Due to the high operation costs for aircraft and the high scientific demand, H₂O data have to be measured continuously without any interruptions. The instrument thus has to be highly reliable, robust, require low maintenance and conduct an entirely automatic and autonomous startup procedure. The restrictions in weight and space as associated with operation on aircraft result in the necessity for a compact and lightweight construction. The utterly complex and mandatory certification process (at least in Germany) enforces an instrument design freeze before a campaign; this results in very stiff constraints for improvements/repairs during a campaign.

3. H₂O Spectroscopy related to the the HAI instrument
HAI consists of four independent but interconnected spectrometers. For each individual channel an individual evaluation procedure is done following an identical, common spectroscopic method: calibration-free direct Tunable Diode Laser Absorption Spectroscopy (cal-free dTDLAS).
3.1. Direct Tunable diode laser absorption spectroscopy (dTDLAS)

The requirements for fast measurements and high chemical selectivity in combination with a robust and small system calls for a contact-less, spectroscopic (hence optical) measurement technique rather than contact sensing methods such as dew point mirror hygrometer (DPH) or capacitive polymer sensors. The latter (e.g. a “Humicap” (Salasamaa and Kostamo, 1986; Smit et al., 2000)) is quite frequently used in meteorological environments for weight, size, and cost reasons (Busen and Buck, 1995; Hansford et al., 2006; Wiederhold, 1997). By choice of the spectroscopic method, optical hygrometers can be set up to become quite immune to hydrophilic/hydrophobic substances (unavoidable in the vicinity of aircraft) as well as particles (dust, soot, ice, etc.) carried by the gas to be analyzed. These capabilities were e.g. extensively demonstrated via measurements inside of combustion processes in industrial power plants (Ebert et al., 2000a; Schlosser et al., 2002; Sun et al., 2013; Teichert et al., 2003).

Tunable diode laser absorption spectroscopy (TDLAS), especially in near infrared spectral range, is a powerful as well as versatile diagnostic technique and led to numerous applications in atmospheric hygrometry (Diskin et al., 2002; Fahey and Gao, 2009; Gurlit et al., 2005; May, 1998; Schiff et al., 1994; Thornberry et al., 2014). Advantageous properties of diode lasers are very high spectral resolution (providing excellent chemical selectivity), high power density, and continuous wavelength tuneability in combination with interesting technical features such as low cost, very low size/weight/power consumption, long life time, excellent beam quality, and optical fiber coupling to name just a few.

The typical setup and working principle of a TDLAS instrument has been frequently described in detail (Lackner, 2011; Schiff et al., 1994; Schulz et al., 2007; Werle, 1998). Therefore, only HAI’s design relevant topics are discussed. Important for an understanding of the novel HAI instrument is the classification of TDLAS instruments by their optical detection schemes in classical single- (Ebert et al., 2000b) or multi-path (Gurlit et al., 2005; Hunsmann et al., 2008; Lübken et al., 1999; May, 1998; McManus et al., 1995) beam setups. In general, longer path lengths - achieved by multi-path optics with a high number of reflections - provide better sensitivity. But higher reflection numbers are also more critical to align. Compared to most airborne TDLAS instruments, HAI has a relatively low number of reflections and thus relatively short path lengths. Further categorizing distinguishes between wavelength-modulation schemes like single modulation frequency = direct TDLAS or dTDLAS (Ebert and Wolfrum, 1994) or double modulation schemes like wavelength modulation spectroscopy WMS (Podolske and Loewenstein, 1993; Silver, 1992; Silver and Hovde, 1994a; Silver and Zondlo, 2006; Vance et al., 2011; Webster et al., 2004). WMS, often used for very compact sensors, provides on the first glance higher sensitivities by using lock-in technologies to efficiently filter noise. This, however, sacrifices the possibility of direct physics-based quality and reliability checks, since the actual measured WMS raw signal contains less spectral information than a dTDLAS raw signal. This aggravates or sometimes even prevents detailed signal analysis based on fundamental physical explanations. Using dTDLAS instead with a special, but less common, first principles evaluation procedure (Ebert and Wolfrum, 2000; Farooq et al., 2008; Mihalcea et al., 1997; Schulz et al., 2007) allows sophisticated evaluation, characterization, quality management and a holistic view on the physical principles behind the data. This circumstance can even be used to avoid typical calibration procedures with reference gas standards as demonstrated in (Buchholz et al., 2013b).
“Calibration-free” TDLAS: An explanation of the term

The term “calibration-free” is often used in different communities with dissimilar meanings. To distinguish, one should consider how calibration is defined by metrology (JCGM 2008, 2008): “calibration (…) in a first step, establishes a relation between the measured values of a quantity with measurement uncertainties provided by a measurement standard (…), in a second step, this information is used to establish a relation for obtaining a measurement result from an indication (of the device to be calibrated)”. In other words, an instrument with a deterministic relation between indication and measured quantity can be kind of “brute-force” calibrated, without knowing the physical details behind the measurement process. This is often used to compensate non-linearities, offsets, drifts or response changes over time as long as they are stable, predictable, or can be extrapolated. But again, the causes and physical explanation of this disadvantageous behavior is then neither known nor understood – it is only corrected for.

For HAI we use the term “calibration-free” to emphasize that HAI does not rely on such a correction process. This means that the hygrometer described in this paper is neither initially calibrated (i.e. like the factory calibration of a commercial instrument) nor is it regularly calibrated or adjusted by a repetitive comparison to a water vapor primary standard. Parameters like gas pressure and temperature that are used for the calculation of the water vapor content via our first principle spectro-physics model are of course measured with classically calibrated sensors. This is done from a practical point because a) primary standards for temperature and pressure are by themselves large facilities and b) the influence in the final uncertainty budget doesn’t justify a first principles approach for pressure and temperature too. Thus, calibration-free doesn’t mean that the all measurement parameters in the entire setup are based on principles. The whole idea behind traceability (JCGM 2008, 2008) is to use other units, higher in the hierarchy of the SI units to generate/analyze the target value. In Metrology, this is named “an unbroken chain of measurements”. To visualize, this means that the first and initial measurement result of the HAI instrument delivers the final concentration value. There is no subsequent “adjustment process” to “tune” an intermediate result towards a target value defined by a higher reference standard. The HAI evaluation directly delivers final values.

This requires that the entire measurement system has to be characterized in advance on such a level that e.g. all influences of the instrument temperature on the end-value has to characterized and the very first measurement value is already determined within its uncertainty limits. Therefore, since there was no calibration, it is termed calibration-free.

The uncertainty budget of HAI’s calibrated sub sensors e.g. for pressure and temperature also has to contain a p-T-sensor drift consideration between two calibration cycles of these p-T-sensors. Sufficient information about the required calibration interval and maximum sensor drifts have to be acquired from trustworthy sources such as the manufactures datasheet or in-house validations. These drift values have then to be considered in the frame of the entire uncertainty budget. E.g. one of the pressure sensors used in HAI (Omega PAA33X-V-1) has long-term stability of 0.5 mbar per year. The uncertainty budget of HAI (see below) clearly indicates that even if this was five times higher, it wouldn’t require a calibration cycle of less than a year.

The calibration-free approach used in HAI could certainly be enhanced by a calibration of HAI (such as (Muecke et al., 1994)) at any time (Buchholz et al., 2013b), even after a campaign, if this seems advantageous, since the requirements for a calibrated instrument are lower than for a calibration-free instrument.
For a limited time section between two calibration intervals we could therefore improve the accuracy of HAI at the expense of additional time and cost effort for the calibration itself. However, we also would then “downgrade” HAI by making its performance dependent on the availability of a calibration reference and we would also limit this performance “gain” to the time span between two calibrations. In addition, it is also a fact that a calibration only indicates the agreement between the reference and the sensor at the time of the calibration. Additional uncertainty contribution would have to be taken into account for drifts of the instrument in the time span between two calibration points. The many "ifs" related to calibration procedures and the high additional “costs” led us to design and realize an instrument which completely avoids such an process and which has a well-defined performance based on a physical first principles model.

3.2. Non-calibrated direct TDLAS (dTDLAS)

The spectroscopic principle of non-calibrated absolute dTDLAS is very briefly presented in the following section. For more detailed information regarding dTDLAS is referred to in the above-mentioned literature. The sketch in Figure 1 shows the schematics of a dTDLAS spectrometer with two independent channels. For low light intensities I(\(\lambda\)) in the mW-range, the transmitted light I(\(\lambda\)) can be described by the extended Lambert-Beer equation (Equation 1) including possible disturbances by background radiation E(t) or broadband transmission losses Tr(t).

Equation 1: 
\[ I(\lambda) = E(t) + I_0(\lambda) \cdot Tr(t) \cdot \exp[-S(T) \cdot g(\lambda - \lambda_0) \cdot N \cdot L] \]

By applying the ideal gas law Equation 1 can be used to retrieve the H2O volume mixing ratio c.

Equation 2: 
\[ c = -\frac{k_B T}{S(T) \cdot L \cdot p} \int ln \left( \frac{I(\nu)-E(t)}{I_0(\nu) \cdot Tr(t)} \right) \frac{d\nu}{dt} \, dt \]

The amount fraction c is in metrological units officially specified as [mol/mol = mol absorber per mol gas] which is in the environmental community better known as “volume fraction” e.g. in units of ppmv or Vol.-%. The term \(\frac{d\nu}{dt}\) is called the dynamic tuning coefficient of the used laser. It can be determined experimentally and is directly linked to the SI units (length) by using the Airy signal of the laser light passing through a planar, air-spaced etalon (Ebert and Wolfrum, 2000; Schlosser et al., 2002). Yet, unpublished, on-going, and long-term \(\frac{d\nu}{dt}\) measurements for the used 1.37 μm DFB laser type over several years indicate a long-term stability of its tuning characteristics better than 1%, which is within the current uncertainties of the tuning characterization. The variable t represents time, \(k_B\) is the Boltzmann constant and L is the optical path length. S(T) is the line strength of the selected molecular transition (see chapter below) and therefore a physical property of the molecule to be measured. The gas pressure (p) and gas temperature (T) can be accurately acquired in a closed-path cell; the quality of the respective measurements in the open-path cell are discussed in the following chapter describing the construction of the open-path sensor. Equation 2 also “explains” the term calibration-free quantitatively as there are no other “hidden” parameters used to derive the water vapour concentration which require a calibration.

3.3. Absorption line selection

Suitable absorption lines have to be selected for a specific application by several criteria (Wagner et al., 2012; Wunderle et al., 2006). Besides a line strength maximization to ensure high sensitivity, other im-
important parameters have to be taken into account. For atmospheric measurements, the cross sensitivity to other gases such as CO₂ needs to be minimized. This ensures a better control of the fitting process due to the fewer degrees of freedom. Similarly, the line should be isolated from other lines to simplify the retrieval of the baseline function. For the open-path measurements, it is highly important to minimize temperature dependence of the line strength in order to minimize the influence of gas temperature uncertainties. Lastly, sometimes the primary constraint is the availability of suitable laser diodes and additional accessories such as fibers, optic components, lenses or detectors. As the certification for airborne instruments nearly prevents improvements/repairs during a campaign, all components need to be very reliable. For HAI, we selected two specific water lines at 2596 nm and 1370 nm. The latter has been used by us and others before (Buchholz et al., 2012, 2013b; Ebert, 2006; Ebert et al., 2004; Fahey and Gao, 2009; Hovde et al., 2001; Hunsmann et al., 2008; May, 1998; Seidel et al., 2012; Witzel et al., 2012; Wunderle et al., 2008) and improved spectral parameters were available (Hunsmann et al., 2006). The 2.6 µm laser is not fiber-coupled, doesn’t have an optical isolator, is less stable in terms of temperature fluctuations and has a lower beam quality, but accesses a factor of 20 stronger line and thus ultimately promises 20 x higher sensitivity. Both lines are shown (Figure 2) as a simulation based on the HITRAN (Rothman et al., 2009) database. Assuming an optical resolution of 5.10⁻⁴ OD, the hygrometer is expected to provide for a 1.4 µm closed-path cell (assuming a 1.5 m path length) under atmospheric conditions a water vapor concentration range from 10 to 40 000 ppmv and for the 2.6 µm closed-path channel from 0.5 to 5000 ppmv. The lower limit will finally be defined by the capability of minimizing and compensating the effects of parasitic H₂O offsets (Buchholz and Ebert, 2014b) and their related uncertainties. For the 2.6 µm channel, the upper limit depends strongly on the retrieval quality of the baseline. This means e.g. under lower pressure conditions (<500 hPa) evaluations up to 40 000 ppmv (similar to 1.4 µm) can be performed. At higher pressures and higher concentrations the absorption line is strongly saturated and such broadened that the baseline cannot be retrieved with a sufficiently low uncertainty. The upper limit in general, i.e. also for the 1.4 µm channels, is defined by the dew point related to the instrument temperature; 40 000 ppmv is equal to 100 %RH at 28.5 °C and 1013 hPa.

4. **HAI configuration**

HAI can be installed on an aircraft in several different ways and configurations. This paper focuses on the deployment during the first HALO (Krautstrunk and Giez, 2012) campaigns (TACTS (Engel et al., 2013) and ESMVal (Schlager, 2014)) which targeted multiphase H₂O measurements. Figure 3 (right) shows the fuselage of HALO with the two mountings belonging to HAI. In the very front is a trace gas inlet (TGI), where HAI uses the second opening from the top to sample air in flight. The sampling height is approx. 25 cm above the aircraft skin. Next to the TGI is the HAI open-path cell (OPc), which is a fiber-coupled, White-type three-mirror configuration mounted on two pylons. These mirrors fold the 4.8 m laser light beam shown in the middle (yellow). Both the TGI and the OPc are installed on HALO specific aperture plates. Figure 3 (left) depicts the cabin installations with the golden HAI main instrument in the middle. The black insulated ½” pipes on the right side of the rack are the inlet pipes guiding the air flow sampled by the TGI to the HAI main unit. These electro-polished stainless steel pipes are heated (70 °C) to evaporate
ic particles and droplets in the gas stream at a flow rate of approximately 100 volume liter/min. The op-
path sensor is not visible in this photo since it is on the left side of the rack.

4.1. HAI main unit

HAI’s main unit comprises all laser/electronic/control etc. modules except for some minor electronics parts
of the OPc, which are directly mounted cabin sidewise on the OPc to minimize noise and electromagnetic-
irradiation: These components on the OPc comprise low noise preamplifiers for detectors (InGaAs respec-
tively InGaAsExt), five converters for PT-100 temperature sensors, mirror heating controllers (20 Watt per
mirror), and two gas pressure sensors.

Many of the modules in the HAI main unit contain innovative developments, which are or will be pub-
lished in subsequent papers to prevent this overview-paper from being overloaded with long lists of tech-
nical details. The important functions/modules to understand the HAI concept are as follows:

a. Automatic line identification and spectral locking to compensate temperature and electrical influences as
well as drifts

b. Internal mechanism (Buchholz and Ebert, 2014b) to minimize and compensate effects from parasitic
absorption which usually leads to variable offset problems in spectrometers like HAI

c. Gas handling system which tempers the air to be analyzed to the temperature of the instrument and
therefore avoids the risk of temperature inhomogeneity even at high gas flows as well provides detailed
information about temperature, pressure, and gas flow in and through the system

d. Several individual sensors and electronics to collect in total more than 120 system parameters about the
HAI system’s health – the so called “housekeeping data”

e. Sequence control system for steering, controlling, and exception treatment of HAI with all its submod-
ules

4.2. HAI’s closed-path cells (CPc)

Core parts of the HAI main unit are the extractive i.e. “closed-path” optical cells (Figure 4) which are inte-
grated in individual 1.4 µm and 2.6 µm optics modules. The 1.4 µm laser is a DFB fiber-coupled diode laser
at 1370 nm from NEL, the 2.6 µm diode laser is a DFB diode laser from Nanoplus, which is collimated with
a ZnSe aspheric lens designed by our group. Both optics modules are completely independent. Both closed-
path cells use a new, compact, high numerical aperture miniature White type (White, 1976) design that
folds the laser beam (total approx. 1.5 m) with three mirrors at approx. 7.5 cm distance. The 1.4 µm cell is an
improved, lens-less fiber-coupled version of the White cell described in (Kühnreich et al., 2013). It is com-
bined with a new vacuum tight fiber-feed-through (Buchholz and Ebert, 2014a), which reaches a very low
leakage rate of 1.9·10⁻⁶ hPaL/s and prevents parasitic absorption offsets due to ambient air. Three mirrors,
one fiber, and one detector are the only optical parts inside the cell. This reduced amount of optics efficient-
ly minimizes fringing effects down to a long-term 10⁻⁴ optical density signal-to-noise (ODsn) level. No read-
justment of the cells was needed and exercised in the approximately 400 flight hours which has been real-
ized over more than three years up to now. The temperature is measured via a 3 mm² accurate (0.3 K) PT100
sensor for slow temperature changes (to₂₅ approx. 2.5sec) and in addition with a thermocouple (type T,
0.5 mm diameter) for faster gas temperature changes (to₂₅ approx. 0.5sec). Due to the above mentioned gas
handling system and the thermal mass of the whole instrument (approx. 31 kg) temperature fluctuations are
measured to be below 0.1 K during aircraft operation. Each cell has a volume of approx. 300 cm³ and is
(depending on the installing scenario) typically flushed during flight with a gas flow of 20 – 50 vol.liter/min
per cell. This leads to an approximate exchange time of 0.7 sec by using a “bulk flow” estimation. The
2.6 µm closed-path cell is by its design quite similar to the 1.4 µm. The most significant difference is the
free-space beam guidance into the cell. Fibers at 2.6 µm are quite critical to handle and suitable fiber beam
splitter etc. weren’t available when HAI was developed. Therefore, the 2.6 µm laser, the reference cell for
spectral stabilization, beam splitters, mirrors, and fiber coupling section for the open-path cell are free-space
coupled without fibers and enclosed in a specially designed gas-tight 2.6 µm optics module box. This mod-
ule is internally dried by a proprietary purging cycle containing a gas drying system based on the same
idea as for removing parasitic offsets in fiber-coupled hygrometer (Buchholz and Ebert, 2014b). The 2.6 µm
closed-path cell is directly coupled to this laser module. Before reaching the cell, the laser beam is therefore
“free-space” guided in a water vapor free environment (in typical operation <0.5 ppmv) for approx. 19 cm.
This efficiently avoids parasitic effects in the beam path. The optical-path ratio between closed-path cell and
“free-space” area of 7.9 (1.5m/0.19m) yields in the close-path 2.6 µm cell an offset of less than 0.063 ppmv
(<0.5 ppmv/7.9) to be corrected. The total uncertainty of the offset determination (realized via the so called
2PS-method (Buchholz and Ebert, 2014b)) amounts to less than 0.4 ppmv which also defines the lower
measurement limit of HAI’s 2.6 µm channel.

4.3. HAI’s open-path cell (OPc)

HAI’s open-path cell (Buchholz et al., 2013a), installed in the HALO’s fuselage, uses a similar optical White
cell concept like the closed-path cells (White, 1976) in HAI’s main unit. Contrary to the closed-path cells,
both wavelengths (2.6 µm and 1.4 µm) are simultaneously coupled into a single set of White arranged mir-
rors via two independent glass fibers for 1.4 and 2.6 µm. The cell (made of 2” diameter copper mirrors) has a
mirror base distance of approximately 15 cm (2x more than the closed-path cells) and is set to an optical
path length of 4.2 m. The optical measurement volume is located approximately 23 cm above the aircraft
skin. The air temperature within the open-path cell is measured via two surface-mounted platinum PT100
sensors. These temperature sensors provide, together with the HALO’s temperature measurement (Giez,
2012) and our CFD simulation (to be discussed in a subsequent publication, some preliminary data in at-
tachment of (Buchholz, 2014)), an estimated (±5 K) temperature in the actual measurement volume of the
open-path cell. Improved temperature field simulations will be realized in the future as soon as a full CFD
model of the aircraft and HAI is accessible. To prevent a dew or frost buildup on the mirror surface, all
three mirrors are individually heated with approximately 20 W each. This raises the mirror temperature,
defined by core temperature measurements inside the copper mirrors, typically 15 – 20 K above the ambient
temperature. The gas pressure measurements are done by two commercial piezo pressure transmitters.
Their individual ports are located on either side of the open-path cell. An in-flight pressure sensor valida-
tion has been realized recently via the first optical airborne pressure measurements (Buchholz et al., 2014b).
This calibration-free approach exploits the pressure dependence of the collisional broadening of the water
absorption line.
4.4. TDLAS based multi-phase H₂O measurements

By simultaneously performing closed-path and open-path water measurements at high-speed, HAI is the first TDLAS hygrometer allowing a calibration-free, synchronous multi-channel, multi-phase water detection. The schematic of the HAI configuration is shown in Figure 4:

Both independent closed-path cells are connected via heated inlet lines (visible in Figure 3 left) to the forward facing TGI sampling in clouds gas-phase water vapor and ice particles or H₂O droplets together. Droplets and ice particles are evaporated in heated inlet pipes, so that both closed-path cells analyze the so called total water content, i.e. the sum of the interstitial gas-phase water plus the evaporated condensed phase water. This evaporation step is necessary since the dTDLAS signal is “blind” for direct absorption from condensed water phases. The shifted broad band spectra of condensed/ice phase water is thus suppressed and dTDLAS is only sensitive to the narrow spectral structure of water vapor. The combination of the narrow absorption line width with the narrow diode laser tuning range allows a very effective discrimination of the broad spectra of ice or liquid water. Depending on the final usage, the total water content has to be further corrected for the particle sampling efficiency of the inlet system impacted by travelling speed, inlet orientation and other parameters causing an enhancement of certain particle size fractions (over-representation of large particles due to their momentum). A detailed sampling characterization and sampling analysis of these HALO inlet (TGI systems) can be found in literature (Krämer and Afchine, 2004).

The open-path cell, directly located on the fuselage of HALO, measures the pure interstitial gas-phase water vapor of the air flowing through the cell. By combining the total water and the gas-phase water measurement, i.e. closed-path and open-path signals, it is possible to derive from the difference of both HAI signals the pure condensed water phase, i.e. the ice/droplet phase. To our knowledge, HAI is the first airborne instrument that can measure all water phases with the same detection principle, the same evaluation strategy, and with two channels, each spectroscopically and temporally linked by one single laser. When independent devices have been used previously to measure these water products, even with similar TDLAS instruments like in (Vance et al., 2015), this always led to the discussion on how much of the deviations are caused by atmospheric effects and how many are linked to discrepancies between the participating instruments caused by different characteristics, response times or nonlinearities of the individual instruments. Indeed, the calibration or even validation of all other open-path hygrometers cited above is based on laboratory measurement or in-flight comparisons with other hygrometers with different properties making it nearly impossible to justify more than statistical (=averaged) statements. Especially in mid-tropospheric regions, small-scale spatial variations of water vapor content lead at typical HALO cruising speeds to local, temporal gradients of up to 1000 ppmv/s in the gas-phase and several 10 000 ppmv/s in the total water phase respectively. HAI’s effective average time of just 1.3 milliseconds at an output data rate of 120 independent measurements per second allows for the first time a detailed investigation of such structures as well as a very high temporal resolution, point-by-point comparison of all four spectrometer signals of HAI. Pairwise comparisons of HAI’s four channels allow therefore various possibilities for self-validation of the in-flight signals. Further, it is possible to avoid averaged statements and facilitate discrimination/analysis of effects such as sampling artefacts, hysteresis, vibrations, oscillations etc.
4.5. Self-validation

The new self-validating capabilities are a consequence of the unique 2x2 multi-channel configuration, which enables six individual validation pathways, as illustrated in Figure 5. Under clear-sky (= cloud-free) conditions, the independent open-path and closed-path measurements should each deliver the same value. Important to note is that both the 1.4 µm and the 2.6 µm spectrometer channels are evaluated without common parameters in calibration-free (= first principles) mode. This means in detail: two different lasers, two different absorption paths, two different absorption lines, different line parameters, different electronics, different gas pressure/temperature sensors, etc. Nevertheless, both spectrometer channels have to agree within their uncertainty if the first principle evaluation is applicable. Due to the large overlapping H₂O concentration range of approximately 50 to 5000 ppmv, all disturbances can be clearly identified as long as they affect both spectrometers differently. Outside of clouds (clear-sky) – i.e. in ice particle/droplet-free air – all four channels spectrometer channels have to agree. This in particular allows to validate (or even if necessary to calibrate) the closed-path cells in a laboratory environment and to use them as an airborne transfer standard to validate or even calibrate in turn the open-path sensor in flight. This is a unique property of HAI, since each open- and closed-path spectrometer is coupled by one single laser and all four spectrometers employ the same evaluation procedure. Hence, HAI is called a 2x2 spectrometer.

5. Results

5.1. Signal assessment

Figure 6 shows - at the same instant of time - all seven raw optical signals from HAI’s measurement channels at approx. 15000 ppmv and 900 hPa. For visualization, the signals of each laser are scaled to the same peak maximum, since transmission changes are irrelevant according to the equation for concentration retrieval (Equation 2). The time axis (240 Hz laser repetition rate) is already converted to the wavelength axis using the dynamic tuning rate. The triangular diode laser current modulation yields to a triangular modulation of the beam intensity and a similar shape in the wavelength space. The tuning range of the 2.6 µm laser is set to be broader in order to allow the evaluation of high absorbance spectra when the peak absorption is in saturation in the absorption line center. The Lorentzian line wings still carry enough information to retrieve the concentration. The absorbance of the open-path signal (e.g. the 1.4 µm signal in purple) is higher than the one from the closed-path cell, due to the threefold longer optical path length in the open-path cell.

The 1.4 µm reference cell signal (blue) is used for spectral stabilization and acquired from a low pressure (approx. 20 hPa) fiber-coupled reference cell. The signal for retrieving and compensating parasitic absorption is shown in green and the associated evaluation is described in (Buchholz and Ebert, 2014b). Accordingly, the three 2.6 µm signals are shown. The parasitic absorption compensation of the 2.6 µm system is due to the “free-space” signal in the sealed laser module (see description above) at least as important as the one for the 1.4 µm spectrometer. This compensation treatment is done by using an ultra-low pressure (approx. 0.5 hPa) pure water reference cell which provides information for both, the spectral stabilization and the parasitic absorption correction. The latter approach is described as “two pressure separation method (2pS)” in (Buchholz and Ebert, 2014b).
The exact gas pressure or gas matrix in the reference cells is irrelevant as we only use the cell for line locking but not for concentration calibration as other TDLAS instruments do. The requirement for such a cell is that it contains a sufficiently high water vapor content to ensure a sufficiently high S/N H₂O absorption signal. The temperatures of both reference cells are accurately measured (PT100 sensor) to monitor the gas pressure increase in the reference cell caused by the temperature raise. This pressure raise induces a measureable, significant shift of the absorption line position which needs to be corrected to achieve a very high line stabilization quality. The retrieved line position thus is corrected for this shift based in the measured temperature and the known pressure shift coefficient of the absorption line.

Figure 7 and Figure 8 show four typical absorption signals in flight with a pre-average of 50 raw measurements recorded at 240 Hz. This leads to 4.8 measurements per second. In-flight situations similar to that, only approx. 1/6 of the whole (up- and down-ramp) spectral scan contains significant contributions of the water vapor absorption line (= the pressure broadened target absorption line). The “rest” of the wavelength scan is used to retrieve the baseline, the laser-out region and for situations where gas pressure and water vapor concentrations are much higher (e.g. low flights in warm areas, especially in clouds). This small section with the narrow absorption line can thus be interpreted as the “effective” water measurement interval, which could then be interpreted as an effective time resolution (= after averaging) of 70 ms for each reading (1/240 * 1/6 * 50).

Figure 7 allows the simultaneous comparison of both closed-path cells. Gas pressure and temperature are slightly different due to the different piping and installation placement inside of HAI. Both H₂O concentrations agree (∆ = 1.6%) within the combined uncertainties (7.9%) of both channels (4.3% respectively 5.9%). Each graph in Figure 7 shows in the top the measured data (green dots) with the fitted Voigt profile (black line). The measurement data are shown as absorbance (ODₑ = -ln(I/I₀)) values, i.e. the detector (I) signal divided by the so called “base line” (I₀). The baseline resembles the absorption spectrum in the absence of the molecular target absorption. It is retrieved from the raw spectrum (I) by applying a synchronous polynomial fit of the baseline and a Voigt profile for the absorption line. The residuals between measured data and the model function are depicted below. The remaining structures visible in the residual are optical interference fringes, caused by light scattered from imperfect mirrors, fiber surface, detector arrangement, and wall backscattering. The frequency of a fringe can be linked to the optical distances between the interfering surfaces. So far, these baseline structures as well as the cell alignment remained stable and the mirrors have never required any cleaning during the last four years.

“Stability” always has to be related to the right timing context: Calibration-free dTDLAS are usually operated at relatively moderate OD resolution levels (of about >10⁻³). Otherwise impacts which affect the long-term stability cannot be controlled or assessed well enough. However, in the context of “short term” precision statements the residual structures only have to be stable in the order of minutes instead of years. The entire fringe structures displayed in the pictures above are phase shifted by temperature due to mechanical expansion of the cell structure. Though, the thermal mass of the cells prevents fast temperature changes, which leads to very stable “short-term” fringes down to the 10⁻⁴ OD levels.

This is a crucial point compared to WMS systems, where fringe levels have to be controlled down to the 10⁻⁶ OD level to achieve the sensitivities advantages WMS is known for. Our calibration-free dTDLAS evalua-
tion is designed to require a stable baseline only in the $10^4$ OD range, which makes it inherently robust and improves the stability of the entire system. In other words, dTDLAS systems like HAI are usually not as sensitive to misalignments, dirt, or optical interfering structures as WMS systems but require higher absorbance (= higher line strength or longer path length) to allow an implementation of our calibration-free approach.

Figure 8 shows a simultaneous comparison between the open-path and the closed-path channel. The outside gas temperature in the open-path cell is approx. 52 K lower than in the closed-path cell. The gas pressure in the closed-path cell is much higher than in the open-path cell, due to the ram pressure guiding the air through the TGI and piping. The measured water vapor concentrations are slightly shifted in time due to the gas transport and the sampling delay in the pipes. The 1σ residual is a factor of six higher in the open-path than in the closed-path sensor. This mainly results from the high wind speed through the open-path cell (approx. 800 km/h), vibrations, and the thermal mass of the closed-path cell as mentioned above.

It is important to mention that for typical diode laser instruments (direct as well as WMS) a change of gas temperature, pressure, and concentration over a large range can cause large systematic offsets, which frequently require at least a three dimensional calibration procedure (including pressure, temperature, and concentration dependence) or the integration of complex assumptions (Duffin et al., 2007; Goldenstein et al., 2014; Rieker et al., 2009) to correct the data via a simulation model. WMS closed-path systems usually avoid this problem by always keeping gas temperature and pressure as close and constant as possible to the calibrated level which is obviously not a viable approach for an open-path system.

5.2. Precision

One common figure of merit to quantify the short term optical precision of a spectrometer is the 1σ noise level in the residuum. The idea is based on the assumption that the residual is governed by random noise which limits the precision of the system. The Signal-to-Noise Ratio (SNR) is then defined by the signal as the ODepeak value and the noise by the statistical standard deviation of the residual and thus equals to ODnoise.

This definition yields for the 1.4 µm (2.6 µm) closed-path cells at 4.8 Hz time resolution a SNR of 54 (204).

The corresponding precision is then $100/54$ ppmv = 1.9 ppmv (at 1.4 µm) respectively $100/204$ ppmv = 0.49 ppmv (at 2.6 µm). This determination is quite consistent with Figure 8 where the precision of the 1.4 µm closed-path can be estimated to 1.3 ppmv in the same way. However, as the fringe structure of the closed-path cell is, as discussed earlier, quite stable it is better to determine the instrument precision via the Allan variance (Allan, 1966). Figure 9 shows the Allan plot for both closed-path cells. The measurement was done at 255 ppmv by measuring gas from a big vessel. Under these conditions, the precision at 4.8 Hz is $0.22$ ppmv (for the 1.4 µm closed-path cell) respectively $0.065$ ppmv (2.6 µm closed-path cell). Both values are approximately a factor of 8 lower than the one derived from the single scan analysis, which confirms the stability thesis for these kinds of fringes. Vice versa we can derive an actual (= unstable, random) 1σ residual OD noise level of 2.3E-5 for the 1.4 µm closed-path and 1.7E-4 ppmv for the 2.6 µm closed-path cell. The best precision is achieved at 3.9 Hz (0.18 ppmv) for the 1.4 µm cell and at 13 Hz (0.055 ppmv) for the 2.6 µm cell. Normalizing this with respect to time resolution and optical path length yields a 1σ precision of 187 ppbv-m-Hz for the 1.4 µm and 31 ppbv-m-Hz for the 2.6 µm.
Comparing that precision ratio of 6 between the 1.4 μm and the 2.6 μm to the ratio in line strength of 20 (see line selection chapter 3.3) confirms the statement mentioned above that drawbacks of the longer wavelength technology significantly constrain the practically achievable maximum gain.

For the open-path sensor, we have not yet been able to calculate an Allan variance so far, since we haven’t acquired a data set in flight where the atmospheric water was constant enough not to dominate the Allan variance via natural H₂O fluctuations. However, using the short term single spectrum evaluation method in Figure 8 yield for the 1.4 μm path a SNR of 145, which leads to a precision of 2.1 ppmv (= 4.6 ppmv·m·Hz⁻¹).

5.3. Uncertainty consideration

One of the major benefits of the dTDLAS evaluation done with HAI is the high level of control over the fitting/evaluation process. In particular, combining it with the complete storage of all raw data allows a dedicated and highly flexible post-flight analysis or post-flight processing. Raw data storage for HAI not only covers all raw spectra but also approximately 120 measured parameters depicting the complete status of the HAI instrument. This “housekeeping data” for example, encompasses 15 temperature measurements that enable diagnostic statements e.g. about temperature inhomogeneity, electronic drift, or sensor malfunctions. Housekeeping data like these facilitate assignments of a measurement uncertainty closer to the one in a metrological sense (Joint Committee for Guides in Metrology (JCGM), 2008). If measurements are done in harsh environments or under rapidly changing conditions, typical concepts of metrological uncertainty assessments fall short since most of their concepts are developed for laboratory applications under quasi-static conditions. But even in this case the housekeeping parameters provide information for a trustful uncertainty consideration. The uncertainty calculations below are based on a physical model (Equation 2) and performed with an approach independent from the actual measurement. Depending on the individual science community, these uncertainties, errors, reproducibility, or misreading are often determined by comparing the readings of an instrument to reference instrument. This is contrary to a metrological uncertainty and furthermore a non-independent approach; we call that therefore simply deviation, standard deviation etc.

According to the Equation 2 for retrieving the final concentration, the individual contributions to the total uncertainty budget can be assessed as followed for both closed-path cells. The optical path length was primarily determined with a Zemax based ray tracing simulation. In addition, it was calculated by the mechanical mirror distance and compared to a test measurement with a known amount of CH₄ gas in the cell. These measurements lead to an uncertainty of the optical path length of 15 mm (1.1%). The uncertainty for the used H₂O absorption line strength is 3.5% [44]. The temperature sensors (PT100 and thermocouple type E as described above) are calibrated against metrological transfer standards (Rosemount platinum resistance thermometer type 162CE (PR), uncertainty 1.5 mK). Due to the well-known fact that temperature measurements in moving gases suffer from many issues, we use a conservative estimation of 1K (0.3%).

The manufacture of the pressure sensors (Omega PAA33X) advertises a resolution of 0.02 hPa and a long-term accuracy of 0.5 hPa. We use 1 hPa as a conservative uncertainty estimation. A general statement for the laser tuning uncertainty is difficult in a metrological sense, since it depends on local effects in the spectra, pressure range, concentration level and number of absorption lines fitted. From our experience, we assume that deviations related to the fitting process in total (tuning + fit process) are in the range of below 1.5% for situations similar to Figure 6 (It has to be kept in mind, that our cal-free dTDLAS systems are usually de-
signed to work in the > 10³ OD level, where a clear signal assessment is feasible. As a conservative approximation we use 2% for the fit and 1% for the tuning. This yields to a total uncertainty of 4.3% for the 1.4 µm closed-path cell.

The largest relative influence to this total uncertainty budget results from the line strength (66%), followed by the fitting uncertainty (21%), as well as the optical path length (7%). Additionally, the offset uncertainty is defined in a calibration-free dTDLAS system by the capability of minimizing and quantifying parasitic offset effects. It is ±3 ppmv for the 1.4 µm closed-path cell of HAI, which uses the same parasitic prevention treatment as SEALDH-II (Buchholz and Ebert, 2014b). Thus, the total uncertainty of the 1.4 µm closed-path system is 4.3% ±3 ppmv. Similar considerations lead to an uncertainty of 5.9% ±0.4 ppmv for the 2.6 µm closed-path cell.

Related to the open-path cell, similar uncertainty statements are more difficult. First of all, a full CFD model of the boundary layer around HALO is currently missing, which would allow a retrieval of the gas temperature in the open-path cell. The first CFD test runs of HAI’s open-path sensor and its immediate surroundings have been realized (A. Afchine (FZJ), published in (Klostermann, 2011)), and led to a first temperature uncertainty estimate of ±7 K. The full CFD model could also provide pressure field calculations. However, in this case, we developed a method on how to retrieve the pressure directly from the dTDLAS signal (Buchholz et al., 2014b) and use that for a validation of the built-in micromechanical pressure sensor. The pressure dependent uncertainty is in the range of 3-5%. The uncertainty of the fitting process reveals a much higher complexity. It depends on all impacts on the spectral signal itself such as: background radiation on the detector, misalignment due to the distortion of the HALO fuselage, forming of an ice layer on the mirrors, high optical dense cloud transects, eddies around the aircraft etc. From all the data gathered and compared during the HALO missions TACTS and ESMVal, we couldn’t find any deviation larger than ±5% in clear sky conditions between the open-path and the closed-path sensor which couldn’t be clearly linked to explainable effects. Besides of the availability of an entire CFD model in the future, HAI will be validated by sampling the pure water vapor gas-phase with its closed-path cells (connected to backward facing TGI measurement) in future campaigns. These datasets will sustainably prove if the ±5% deviation is valid under all conditions, since all four channels of HAI (Figure 5) should permanently measure then the same atmospheric value. Compared to the deviations, revealed in the AquaVIT (Fahey et al., 2014) comparison of state of the art, airborne hygrometer in 2007, this 5% is fourfold smaller than the span (20%) of those instruments there. Additionally, one has to keep in mind that this AquaVIT comparison was done under quasi-static laboratory conditions. Therefore, this in-flight 5% deviation between the open-path and the closed-path sensor shows the great performance of HAI, even if a general metrological uncertainty is difficult to be given at this time.

5.4. First airborne measurements with HAI on the HALO aircraft

Figure 10 shows typical HAI measurements during the first HALO science mission “TACTS/ESMVal”. The chosen flight profile shows 15 min of a slow ascent from the lower to the upper troposphere. Depicted in Figure 10 are: the H₂O concentration measured with the 1.4 µm CPc (black), the 1.4 µm OPc (magenta), and the 2.6 µm CPc (red) channel of HAI (left axis) as well as the ambient gas pressure (right y-axis, in blue), which is directly connected to the flight level (height). The green dotted line shows the water vapor satura-
tion mixing ratio (SMR) calculated from the static ambient temperature and pressure measured by the HALO avionic system (Giez, 2012). The SMR indicates that HALO flew in the early part of the flight (left) in clear sky conditions (no clouds) and in the second part of the flight though cirrus clouds. All Figure 10 data were plotted with 1 Hz temporal resolution. With increasing height, the H$_2$O concentration declines about 2.5 magnitudes from approximately 2000 ppmv to 75 ppmv, in combination with simultaneous relative pressure variation from approx. 175 to 450 hPa ($\Delta p=275$ hPa). The H$_2$O concentrations, measured with HAI’s two independent closed-path spectrometer channels, fit nicely to each other and show only a small average relative deviation of 1.9% (bottom graph in red), which is entirely covered by the uncertainty range of each spectrometer (4.3% resp. 5.9%). This result is consistent with other measurements, where the 2.6 $\mu$m path was persistently on average 2% higher than the 1.4 $\mu$m channel. This could be easily explained with the 2.6 $\mu$m line strength from the HITRAN database being 2% too small. The 1.9% deviation also needs to be related to the fact that both hygrometer channels are evaluated completely independently, without any calibration of the instrument and most importantly under real-world flight conditions. Despite these demanding conditions HAI’s two closed-path channels generate a relative deviation which is just 1/10 of the above mentioned 20% in AquaVIT (Fahey et al., 2014), which demonstrates the excellent accuracy of the HAI spectrometers.

The purple signal in Figure 10 shows the measurement of the 1.4 $\mu$m open-path sensor (1.4 $\mu$m OPr). Compared to the 1.4 $\mu$m closed-path cell (1.4$\mu$m CPC), the 1.4 $\mu$m OPr shows a relative deviation of $\pm$5% peak-to-peak or 2% on average. Even though this performance seems good in comparison with the AquaVIT results, a more suitable literature reference would be one with in-flight open-path measurements rather than quasi static laboratory comparisons. However, there is none like that and therefore a “state of the art” statement is quite difficult. This holds especially when we consider the fact that the “uncertainties” or “errors” stated for other the AquaVIT participants were claimed to be on such a low level that they did not cover or explain the deviations observed. This is strongly correlated to different and non-standardized definitions, procedures on how to calculate sensor accuracy and other performance figures yielding often a systematic underestimation of uncertainties compared to metrological procedures such as (JCGM 2008, 2008; Joint Committee for Guides in Metrology (JCGM), 2009).

Though, a short incomprehensive overview on the performance of previous and current open-path instruments should be given here to compare HAI with their claimed accuracy level:

a) A calibrated, 1 m direct absorption, open-path sensor at 1.4$\mu$m was developed in a feasibility study for the Strato 2C research aircraft and described to cover a concentration range of 2-2000 ppmv with 100 ppbv precision at 1 Hz. It was claimed to have an accuracy of 5%, but this hygrometer never flew (Roths and Busen, 1996).

b) A calibrated open-path sensor with an optical path length of 375 cm, folded in a Herriot cell, and a time resolution of 25 Hz has a precision of 80 ppbv. This instrument is designed for the American HIAPER aircraft, a Gulfstream 500 similar to the HALO aircraft. Typical deviations to other instruments were in the range between 2% and 10% (Zondlo et al., 2010).
c) A calibrated, Herriott cell based open-path hygrometer with a total path length of 11 m, a measurement frequency up to 10 Hz and a precision of 50 ppbv at 0.5 Hz. This instrument claims to have an uncertainty of 5% to 10% (May, 1998).

d) An innovative concept was deployed by (Diskin et al., 2002). The spectrometer’s (2f, calibrated) hardware is attached from the inside of the cabin to a window. The laser beam hits a retro reflector mounted on the outer engines under then airfoil. The total absorption path is 28.5 m; the measurements frequency is 1 Hz; a 1σ error is claimed to be 3.7% (Podolske et al., 2003). Measurements published in (Diskin et al., 2002) shows deviations in the same order.

In summary, HAI’s 2% deviation appears to be better or at least equal to typical statements in publications but is based for the first time on a metrologically evaluated, calibration-free approach.

To further evaluate the performance of HAI’s open-path channels we discuss some of the TACTS data. The dark purple line shows the data after smoothing them with a 60 second sliding window algorithm. Quite interesting are the two dents at 13:41 and 13:44 in the relative deviation, which means the closed-path measurement was higher than the open-path measurement. Both of them happened after the water vapor concentration dropped sharply. Hence, the dents might be explained by desorption processes inside of the TGI, its piping and the measurement cell. This idea is backed by the 1.4 µm to 2.6 µm closed-path comparison (red). The same behavior is also visible at 13:44, but here the 2.6 µm closed-path cell added less desorption water to the gas steam than the 1.4 µm one. Another influence leading to that deviation could be a temperature measurement on HALO affected by fast changes of humidity. The total temperature influence in Equation 2 is at this temperature and pressure approximately 0.6%/K caused by the temperature influence via the ideal gas law, the line strength temperature dependence and the Gaussian width. Other major influences can be excluded since all relevant housekeeping data clearly indicate that HAI operating as expected during the entire time period. The final distinction of small local effects like that can be done by acquiring more flight data in the troposphere for a better understanding of the sampling system. This is planned to be realized in a future, more mid-troposphere oriented campaign.

This case study and analysis of HAI’s capabilities and performance demonstrates the unique opportunities offered by HAI, which allow a deep analysis to identify and eventually classify different measurement sections of a flight in order to distinguish instrumental artefacts from real atmospheric situations.

On the second part of the flight shown in Figure 10, the aircraft flew through ice clouds (cirrus, temperature < -40 °C). The evaporation of the ice crystals captured by the gas inlet leads to a clear enhancement of the amount of water vapor in the closed-path cells. The relative comparison shows a ± 5% noisy structure, which is caused by the low time resolution of just 1 Hz. H2O gradients in this particular part of the flight are in the range of several 1000 ppmv/s. Due to SSD space limitations in the first campaign, only parts of the flights where sampled with the highest time resolution of 240 Hz; other parts the data were online pre-averaged. Like in the clear sky region, the average deviation between both closed-path cells during cirrus cloud transects remains at 2%. To further check the accuracy of the open-path channel, we use two methods: The first is based on the physical argument that the gas-phase measurement always has to be lower than the total water measurement, which consists of gas-phase plus evaporated ice/liquid-phase. This requirement is entirely fulfilled. The second check compares the absolute gas-phase measurement with the
saturation mixing ratio (SMR). In case of a fully equilibrated cloud, the gas-phase has to be saturated at temperature T, meaning that the relative humidity is \( rH = 100\% \), from which we can derive the SMR with the help of the water vapor partial pressure curve. This is a weaker check, since it is a well-known fact that super saturation can occur during strong air updrafts.

The transition region at the edge of the cloud is also very interesting, where the closed-path cell sees an enhancement while being still below the SMR. This means that the fringe of the cloud was not saturated but had sublimating ice particles in it.

The data between 13:40 and 13:48 of Figure 10 right are plotted in Figure 10 left in a different way: The x-axis shows the water vapor concentration measured by the 2.6 \( \mu m \) closed-path cell; the y-axis shows the corresponding relative deviation between the 1.4 \( \mu m \) closed-path cell (in black; the 1.4 \( \mu m \) open-path cell in purple respectively) and the 2.6 \( \mu m \) closed-path cell. The \( \pm 10\% \) range of the y-axis is chosen to show HAI’s independent in-flight performance in the perspective of the results of the cited laboratory comparison AquavIT. As explained above, there is a small offset between both instrument channels. This is not surprising as all HAI channels are evaluated completely independent in the described calibration-free approach. The deviations between the 1.4 \( \mu m \) and 2.6 \( \mu m \) closed-path cells of about 2% have consistently been seen in other measurements too, but are within the metrological uncertainty of the evaluation. This systematic deviation is most likely caused by an inaccurate line strength parameter in the HITRAN (Rothman et al., 2009) database used for this evaluation. The very tiny concentration dependency (0.18% per 1000 ppmv) of the relative deviation between the 1.4 \( \mu m \) and the 2.6 \( \mu m \) close-path cell reflects the larger complexity of evaluating the 2.6 \( \mu m \) spectra at very high optical densities (OD) – e.g. approx. OD = 16 at 8000 ppmv.

As a remark: An evaluation of the 2.6 \( \mu m \) close-path cell at 8000 ppmv is from a spectroscopic point of view similar to the evaluation of a hypothetical 160,000 ppmv concentration in the 1.4 \( \mu m \) close-path cell. Comparing this 160 000 ppmv with HAI’s (1.4 \( \mu m \) close-path cell) calculated offset uncertainty of 3 ppmv (laboratory validations show real fluctuation of just around 0.6 ppmv) demonstrates the impressive linearity which can be achieved with our dTDLAS systems like HAI over a range of 5 to 6 orders of magnitude.

These first detailed close-ups in HAI’s multi-phase measurements show the potential to investigate many interesting questions in the following campaigns focusing on cirrus clouds which are strongly linked to a very reliable accuracy study for the open-path cell in clouds. To really investigate and fully understand all effects, HAI needs more data in cirrus, mixed-phase and other clouds in a multi-phase configuration as well as a validation campaign in which closed- and open-path cells measure the pure gas-phase via a backward facing measurements. The latter is necessary to distinguish between sampling-effects and optical/spectral-effects in and outside of clouds. This knowledge, especially in combination with the mentioned CFD model, will then allow reevaluating HAI’s raw data which are always saved during flights and extent statements even for this short cirrus transect.

5.5. Performance summary

Figure 11 shows an overview of HAI’s current performance features. The 2.6 \( \mu m \) open-path channel is not mentioned due to low light transmission in the 2.6 \( \mu m \) fiber. This problem is likely to be related to the novel ZBLAN fiber used here which prevented the 2.6 \( \mu m \) open-path to excel the 1.4 \( \mu m \) open-path performance
during the first flight campaign. It is expected for future campaigns that the optical transmission and thus the 2.6 \( \mu \text{m} \) open-path channel performance can be improved significantly.

6. Conclusion and outlook

The novel Hygrometer for Atmospheric Investigation, HAI, realizes in a unique concept a simultaneous gas-phase and total water measurement. Based on calibration-free direct tuneable diode laser absorption spectroscopy, HAI provides a variety of unique feature as summarized in Figure 11 such as accurate (< 4%), precise (0.065 ppmv at 4.8 Hz, channel depended) and very fast (down to 1.3 ms) measurements with metrologically defined uncertainties (4.3%, channel dependent). HAI contains four measurement channels, grouped as two completely independent dual-channel spectrometers, one at 1.4 \( \mu \text{m} \) and one at 2.6 \( \mu \text{m} \) to cover the entire H\textsubscript{2}O concentration range of the aircraft accessible atmosphere. Each spectrometer feeds light in a wavelength-individual extractive, closed-path cell with an optical absorption path length of 1.5 m for total water measurements. Additionally, both spectrometers couple their light in a common open-path cell (optical path of 4.8 m) located outside of the aircraft fuselage, for a sampling-free and contactless determination of the gas-phase water content. These four spectroscopic channels plus three additional supplementary spectroscopic channels allow multiple self-validation strategies inside and outside of clouds and therefore solve the current lack of an integrated approach to validate open-path sensors in flight. HAI’s complex control software minimizes maintenance at ground and ensures almost entirely autonomous operation. In addition, instrument health is permanently supervised by permanent storage of more than 120 housekeeping data. This enables a novel, holistic quality management and a sophisticated signal cross check, which guarantees a high trust level of the final H\textsubscript{2}O values.

HAI was operated for the first time during the TACTS/ESMVal flight campaign for more than 120 operation hours without any malfunction. The entirely independent, never calibrated first principles evaluation of the closed-path spectrometer channels yield in-flight deviations of only 1.9% over a large concentration (75 to 2000 ppmv) and pressure range (175 to 450 hPa). The deviation between the open-path and the closed-path measurements in the same flight segment was on average just 2% with a short-term deviation between +3% and -5%. Despite measuring with a single evaluation concept, over a very broad range of conditions (i.e. temperatures (-70°C to 30°C), gas speeds (cm/s vs 100 m/s), optical disturbances (no background light vs. sunlight, clean mirrors vs. dirty scratched mirrors outside), HAI provided a high trust level of the data over extensive science missions. Laboratory evaluations demonstrated the lowest achievable precision of 0.18 ppmv (at 3.8 Hz) for the 1.4 \( \mu \text{m} \) closed-path cell and 0.055 ppmv (at 13 Hz) for the 2.6 \( \mu \text{m} \) closed-path cell.

In conclusion, HAI proved during its first deployment a novel, highly complex and demanding set of capabilities. This will enable in the future a much more accurate and stringent evaluation of atmospheric multiphase water vapor data inside and outside of clouds and foster in further HALO missions the investigation of new scientific questions in the atmospheric water cycle. HAI can serve in the future as a major, powerful tool for cutting-edge atmospheric water vapor measurements.

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7. References


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Figures:

Figure 1: Simplified schematic of a standard dTDLAS setup with several spectroscopic channels: HAI combines two lasers (one at 1.4 µm for high, one at 2.6 µm for low water vapor concentrations respectively), with two locations cells (1x open- and 2x closed-path), and supplementary spectroscopic channels (parasitic water detector (Buchholz and Ebert, 2014b) and reference cell for spectral stabilization). HAI has in total seven spectroscopic channels.
Figure 2: HITRAN 2008 line strength plot (Rothman et al., 2009) for CO$_2$ (green) and H$_2$O (green). The turquoise spectrum in the back shows the scaled (factor 39) CO$_2$ line strengths for visualization of the stratospheric situation with a low water vapor content (10 ppmv) and standard CO$_2$ (390 ppmv) level.

Figure 3: Photos of HALO’s cabin layout with installed payload for the TACTS/ESMVal campaign (left). Photo of HALO’s fuselage showing the trace gas inlet (TGI) for HAI’s closed-path cells (for total water detection) as well as HAI’s open-path cell (for gas-phase water detection; yellow marked are the open-path laser beams).

Figure 4: Schematic of HAI’s working principle in the multi-phase configuration. By combining selective open-path gas-phase measurements with total water measurements in closed-path, extractive cells (ice, droplets are evaporated by...
heated inlet lines before reaching the cells), it is possible to derive the ice water content from the difference between closed-path and open-path readings.

Figure 5: HAI’s self-validation possibilities under clear-sky conditions (black) and measurement redundancies under all atmospheric conditions (red). For details see text.

Figure 6: dTDLAS raw signals (spectra) of all seven spectroscopic channels permanently acquired and saved by HAI. The x-axis is already converted to the wavelength axis via the individual dynamic tuning rate of each laser. Each spectrum contains approx. 1700 16-bit values. (PA: parasitic absorption, sStab: spectral stabilization)
Figure 7: Typical pre-processed absorption signals (after baseline, offset and transmission correction) for both closed-path spectrometer channels (left 1.4 μm right: 2.6 μm) during flight. The laser modulation frequency was 240 Hz. 50 individual raw scans are pre-averaged yielding 4.8 Hz O2 measurements per second with 69 ms total integration time. This corresponds to a spatial resolution of 15 m at 800 km/h cruising speed. Without averaging, HAI achieves a maximum time resolution of 1.3 ms, respectively a spatial resolution of 30 cm at 800 km/h cruising speed. (It should be noted that the four vertical axes have different scales).

Figure 8: Comparison of pre-processed TDLAS scans (similar to Figure 7) of HAI’s 1.4 μm closed-path (left) and the 1.4 μm open-path cell (right) during a HALO flight. Despite the entirely different measurement conditions such as wind speed (cm/s to 100 m/s) or temperature (+28 °C to -23 °C), both channels are evaluated calibration-free with the exact same methods and model.
Figure 9: Allan variance plots for both closed-path spectrometer channels. The measurements were done by analysing gas from a big buffer vessel with a H$_2$O concentration of 255 ppmv. The optimal precision at 4.8 Hz derived from this measurement is 0.22 ppmv for the 1.4 µm closed-path cell and 0.065 ppmv for the 2.6 µm closed-path cell. The best i.e. highest precision of 0.18 ppmv is achieved at 3.8 Hz for the 1.4 µm cell and 0.055 ppmv at 13 Hz for the 2.6 µm cell.

Figure 10: HAI measurement on board of HALO during the TACTS/ESMVal campaign. Left: The flight profile shows a 15 min section of a slow ascent from the lower troposphere to the upper troposphere. The first part (under clear sky condition) can be used to perform an absolute in-flight accuracy comparison of HAI’s measurement channels; the second part demonstrates a multi-phase H$_2$O measurement. For further explanations of all signals and visible effects see text.

Right: The scatterplot displays on the x-axis the water vapor concentration measured by the 2.6 µm closed-path cell and on the y-axis the relative deviation of the 1.4 µm closed-path cell (and the 1.4 µm open-path cell respectively) from the 2.6 µm closed-path cell measurements. This scatterplot contains all data from 13:40 to 13:48 shown in the left graph.
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Figure 11: Performance overview of HAI:

0) lower limit defined by offset uncertainty, upper limit by instrument temperature or spectroscopic effects respectively; 1) calculated (!) uncertainty based on calibration-free evaluation model; 2) calculated, based on the accuracy level of the independent parasitic absorption minimization and determination (Buchholz and Ebert, 2014b); 3) general statements for typical airborne application; 3+4+5+6+7) statements based on several metrological method validations which are not part of this paper; 8) precision derived from single absorption signal as described in paper; 9) precision derived from Allan variance approach; 10) depending on broadening of the absorption line the max time resolution variates between 1 and 2 ms; 11) HAI reports 120 independent measurements per second; 12) the effective time resolution is limited for the closed-path cell by the gas flow rate (exchange rate) and for the open-path cell by a minimum average number of two raw spectra to save SSD space; 13) the Allan variance is not a feasible approach for retrieving reliable in-flight statements; *) this open-path figures are inferred from closed-path measurements, a direct metrological validation not viable; **) figures are based on a variety of studies which will be published in individual, coherent publications.