



HDO and H_2O total column retrievals from TROPOMI shortwave infrared measurements

Remco Scheepmaker¹, Joost aan de Brugh¹, Haili Hu¹, Tobias Borsdorff¹, Christian Frankenberg², Camille Risi³, Otto Hasekamp¹, Ilse Aben¹, and Jochen Landgraf¹

¹SRON Netherlands Institute for Space Research, Utrecht, The Netherlands

²Jet Propulsion Laboratory (JPL), California Institute of Technology, Pasadena, CA, USA

³Laboratoire de Météorologie Dynamique, Insitut Pierre Simon Laplace, CNRS, Paris, France

Correspondence to: J. Landgraf (J.Landgraf@sron.nl)

Abstract. The Tropospheric Monitoring Instrument (TROPOMI) onboard the European Space Agency Sentinel-5 Precursor mission is scheduled for launch in the last quarter of 2016. As part of its operational processing the mission will provide CH_4 and CO total columns using backscattered sunlight in the shortwave infrared band (2.3 μ m). By adapting the CO retrieval algorithm, we

- 5 have developed a non-scattering algorithm to retrieve total column HDO and H₂O from the same measurements under clear sky conditions. The isotopologue ratio HDO/H₂O is a powerful diagnostic in the efforts to improve our understanding of the hydrological cycle and its role in climate change, as it provides insight in the source and transport history of water vapour, nature's strongest greenhouse gas. Due to the weak reflectivity over water surfaces we need to restrict the retrieval
- 10 to cloud-free scenes over land. We exploit a novel two-band filter technique, using strong-vs-weak water or methane absorption bands, to pre-filter scenes with medium-to-high level clouds, cirrus or aerosol and to significantly reduce processing time. Scenes with cloud top heights ≤ 1 km or very low fractions of high-level clouds, or scenes with an aerosol layer above a high surface albedo are not filtered out. We use an ensemble of realistic measurement simulations for various conditions to
- 15 show the efficiency of the cloud filter and to quantify the performance of the retrieval. The single measurement precision in terms of δD is better than 15–25‰ for even the lowest surface albedo (2– 4‰ for high albedos), while a small bias remains possible of up to ~ 20‰ due to remaining aerosol or up to ~ 70‰ due to remaining cloud contamination. We also present an analysis of the sensitivity towards prior assumptions, which shows that the retrieval has a small but significant sensitivity
- 20 to the a priori assumption of the atmospheric trace gas profiles. Averaging multiple measurements over time and space, however, will reduce these errors, due to the quasi-random nature of the pro-





file uncertainties. The sensitivity of the retrieval with respect to instrumental parameters within the expected instrument performance is $< 3\%_o$, which represents only a small contribution to the overall error budget. Spectroscopic uncertainties of the water lines, however, can have a larger and more

- 25 systematic impact on the performance of the retrieval and warrant further reassessment of the water line parameters. With TROPOMI's high radiometric sensitivity, wide swath (resulting in daily global coverage) and efficient cloud filtering, in combination with a spatial resolution of $7 \times 7 \text{ km}^2$, we will greatly increase the amount of useful data on HDO, H₂O and their ratio HDO/H₂O. We showcase the overall performance of the retrieval algorithm and cloud filter with an accurate simulation
- 30 of TROPOMI measurements from a single overpass over parts of the USA and Mexico, based on MODIS satellite data and realistic conditions for the surface, atmosphere and chemistry (including isotopologues). This shows that TROPOMI will pave the way for new studies of the hydrological cycle, both globally and locally, on timescales of mere days and weeks instead of seasons and years, and will greatly extend the HDO/H₂O datasets from the SCIAMACHY and GOSAT missions.

35 1 Introduction

Water vapour, being the strongest natural greenhouse gas, plays a vital role in our understanding of climate change. It is part of a positive atmospheric feedback mechanism (Soden et al., 2005; Randall et al., 2007), and it plays a role in the mechanisms of cloud formation, of which the feedback mechanisms are still poorly understood (Boucher et al., 2013). A correct understanding of the many

40 interacting processes that control atmospheric humidity are crucial for General Circulation Models (GCMs) to come to accurate climate projections (Jouzel et al., 1987; Yoshimura et al., 2011; Risi et al., 2012a,b).

Measurements of stable water isotopologues, such as HDO, can be a unique diagnostic to improve our knowledge of the hydrological cycle (Dansgaard, 1964; Craig and Gordon, 1965). Different iso-

- 45 topologues have different equilibrium vapour pressures, which leads to a temperature dependent isotope fractionation whenever phase changes occur. The ratio HDO/H₂O of an air parcel is therefore dependent on the source region's location and temperature and the entire transport history of the air parcel, including all evaporation, condensation and mixing events. This makes measurements of the ratio HDO/H₂O a valuable benchmark for the evaluation and further development of GCMs
- 50 and explains why isotopologues have been used for decades in the fields of paleoclimatology, either using ice cores (Dansgaard et al., 1969; Jouzel et al., 1997) or speleothems (Lee et al., 2012), and hydrology in general (Mook, 2000; Aggarwal et al., 2005).

In the last decade there has been a rise in the application of water isotopologues to the atmospheric component of the hydrological cycle. This is directly related to improved remote-sensing

55 techniques that can accurately measure water vapour isotopologues from ground-based networks, such as the Total Carbon Column Observing Network (TCCON, Wunch et al., 2011) and the Net-





work for Detection of Atmospheric Composition Change (NDACC, formerly the Network for Detection of Stratospheric Change, Kurylo and Solomon, 1990; Schneider et al., 2016), as well as global measurements from space with instruments such as the Interferometric Monitor for Greenhouse

- 60 gases (IMG, Zakharov et al., 2004), the Thermal Emission Spectrometer (TES, Worden et al., 2007), the SCanning Imaging Absorption spectroMeter for Atmospheric CHartographY (SCIAMACHY, Frankenberg et al., 2009; Scheepmaker et al., 2015), the Infrared Atmospheric Sounding Interferometer (IASI, Herbin et al., 2009) and the Greenhouse gases Observing Satellite (GOSAT, Frankenberg et al., 2013; Boesch et al., 2013). These new techniques allow for more frequent and global
- 65 measurements of the ratio HDO/H₂O in water vapour, and show the clear potential to further our understanding of the atmospheric hydrological cycle through comparisons with GCMs (Frankenberg et al., 2009; Yoshimura et al., 2011; Risi et al., 2012a,b).

Here, we present an algorithm and performance analysis for new measurements of total column HDO and H_2O using the Tropospheric Monitoring Instrument (TROPOMI, Veefkind et al.,

- 70 2012), onboard the European Space Agency (ESA) Sentinel-5 Precursor (S5P) mission, scheduled for launch in Q4 2016. Like SCIAMACHY, TROPOMI will measure HDO and H_2O in a 2.3 μ m shortwave infrared (SWIR) band of backscattered sunlight, which provides a high sensitivity near the surface. TROPOMI, however, will have a higher spatial resolution with 7×7 km² ground pixels, better radiometric performance, a larger swath and shorter revisit time, resulting in daily global
- 75 coverage and many more measurements over cloud-free land pixels, while also demanding more efficient processing. With TROPOMI we have the opportunity to extend and improve the existing global HDO/H₂O time series, and study spatial and temporal gradients with higher spatial sampling and resolution.

In Sect. 2 we describe how we adapted TROPOMI's CO algorithm to retrieve HDO, H₂O and their respective averaging kernels and how we filter for cloudy scenes. We then describe the performance of the algorithm in Sect. 3, as tested on a series of synthetic measurements with systematically varying scattering layers. A sensitivity analysis of the various input parameters is presented in Sect. 4. The performance on a realistic scenario of measurements above North America is presented in Sect. 5. Finally, in Sect. 6 we discuss our results in the context of other studies and we formulate

85 our conclusions.

2 Retrieval algorithm description

Due to the large difference in atmospheric abundance between HDO and H_2O , the measurement sensitivity, reflected in the averaging kernels, is very different for HDO and H_2O . This makes the interpretation of their ratio very challenging under conditions of light scattering by clouds. We

90 therefore have to pre-filter for the most cloudy conditions, which at the same time reduces processing time. This cloud filter will be described in Sect. 2.3. After cloud filtering we use a non-scattering





retrieval algorithm, adapted from the SICOR algorithm that has already been developed as part of ESA's operational CO algorithm, which is described by Landgraf et al. (2016) in this issue. By using this heritage of TROPOMI's CO processing, we benefit from an algorithm optimised for speed,

95 while also leveraging already existing expertise and software. Sections 2.1 and 2.2 describe the specific implementation of the algorithm needed to retrieve both the $\rm H_2O$ and $\rm HDO$ total column densities.

2.1 Forward model and averaging kernels

Throughout this paper we consider all water isotopologues present in the SWIR range, i.e. H¹⁶₂O,
H¹⁸₂O and HD¹⁶O, as separate absorbing species. For readability, however, we will simply write "H₂O" when in fact we refer to the main isotopologue H¹⁶₂O, and "HDO" when we refer to HD¹⁶O. To simulate the SWIR radiance measurement, we employ a non-scattering forward model F that simulates the reflected Earth radiance at its spectral sampling point λ_i by the spectral convolution of the simulated radiance at the top of the model atmosphere I^{TOA} with the instrument spectral 105 response function (ISRF) s_i,

$$F_i = s_i * I^{\text{TOA}} \,. \tag{1}$$

Here, we assume that sunlight is scattered only at the Earth surface into the satellite line of sight (LOS) and is attenuated by atmospheric absorption along its path. Using this approximation, the simulated radiance at wavelength λ is given by:

110
$$I^{\text{TOA}}(\lambda) = A_s(\lambda) \frac{\mu_0 F_0(\lambda)}{\pi} \exp\left(-\frac{1}{\tilde{\mu}} \tau_{\text{tot}}(\lambda)\right),$$
 (2)

where A_s is the Lambertian surface albedo, $\mu_0 = \cos(\Theta_0)$ with the solar zenith angle Θ_0 . For low solar zenith angles, μ_0 is corrected for the sphericity of the Earth according to Kasten and Young (1989). F_0 is the solar irradiance inferred from TROPOMI solar measurements (van Deelen et al., 2007; Landgraf et al., 2016) and

115
$$\tilde{\mu} = \frac{\mu_0 \mu_v}{\mu_0 + \mu_v}$$
 (3)

indicates the air mass factor with $\mu_v = \cos(\Theta_v)$ and viewing zenith angle Θ_v . The total optical thickness τ_{tot} is given by

$$\tau_{\rm tot}(\lambda) = \sum_{k} \int_{0}^{z_{\rm TOA}} \sigma_k(z,\lambda) \,\rho_k(z) \,dz \,, \tag{4}$$





where z indicates the altitude ranging from the surface z = 0 to the top of the model atmosphere 120 z_{TOA} . Index k represents the relevant absorbers: CO and CH₄ including all their isotopologues, and H₂O, H₂¹⁸O and HDO. $\rho_k(z)$ is the concentration of absorber k at altitude z and $\sigma_k(z, \lambda)$ are the corresponding wavelength- and altitude-dependent absorption cross sections.

The retrieval relies on a priori concentration profiles for CO and CH_4 from the TM5 chemistry model (Krol et al., 2005) and specific humidity profiles from the European Centre for Medium-

- 125 Range Weather Forecast (ECMWF, Dee et al., 2011). These profiles are interpolated to the higher resolution of the TROPOMI pixels, using a digital elevation model to account for variations in air mass due to orography. Specific humidity is converted into concentration profiles for the absorbers H₂O, H₂¹⁸O and HDO using their natural abundance ratios.
- In Fig. 1 we show a simulated transmission spectrum for the entire TROPOMI SWIR spectral range (2305–2385 nm). The top panel shows the total transmission, while the lower four panels show the individual transmissions of the main absorbing species (H₂O, HDO, CH₄ and CO). For the retrieval of the HDO and H₂O total column densities we chose the spectral window between 2354.0–2380.5 nm (indicated with the grey band), as a trade-off between inclusion of the strongest HDO absorption lines with only minor overlap with the strongest H₂O absorption lines. The smaller
- 135 spectral windows in respectively blue (H_2O) and red (CH_4) indicate the weak and strong absorption bands used for cloud filtering (see below). Although we additionally fit $H_2^{18}O$ to improve the fit quality of the other species, its absorption lines are weaker than those of HDO (not shown). An accurate retrieval of total column $H_2^{18}O$ in this spectral range is not yet feasible and therefore not part of the final retrieval product.
- 140 For the following analysis, we define two relative profiles:

$$\rho_k^{\rm rel} = \frac{\rho_k}{c_k} \tag{5}$$

and

$$\rho_{k,k'}^{\text{rel}} = \frac{\rho_{k'}}{\rho_k} \,, \tag{6}$$

where ρ_k^{rel} is the relative profile of absorber k with respect to the vertically integrated total column

$$145 \quad c_k = \int \rho_k(z) dz \,, \tag{7}$$

and $\rho_{k,k'}^{\text{rel}}$ is the relative profile of absorber k' with respect to absorber k. Assuming that the abundance of trace gas k changes by a scaling of the reference profile ρ_k^{rel} , the derivative of the total optical depth with respect to the trace gas column density is given by

$$\frac{\partial \tau_{\text{tot}}}{\partial c_k} = \frac{1}{c_k} \int \sigma_k(z,\lambda) \,\rho_k(z) \,dz \tag{8}$$







Fig. 1. Simulated spectral transmittance in the SWIR spectral range, showing the total transmittance (top panel) as well as the absorption features of the individual species (lower 4 panels). The simulation was performed assuming a solar zenith angle of 0° and a viewing zenith angle of 40° . The 2354.0–2380.5 nm retrieval window is indicated in grey. The coloured windows highlight the weak and strong absorption bands of H₂O (blue) and CH₄ (red), used for cloud filtering.

150 and thus

$$\frac{\partial I^{\text{TOA}}}{\partial c_k} = -\frac{I^{\text{TOA}}}{\tilde{\mu}c_k} \int \sigma_k(z)\rho_k(z)dz \,. \tag{9}$$

Corresponding expressions hold for the radiance derivative with respect to the trace gas concentration at a certain altitude level. Finally, the derivative of I^{TOA} with respect to surface albedo A_s is

155
$$\frac{\partial I^{\text{TOA}}}{\partial A_s} = \frac{\mu_0 F_0}{\pi} \exp\left(-\frac{1}{\tilde{\mu}}\tau_{\text{tot}}\right).$$
(10)





After the spectral convolution in equation (1), we have a linearised forward model,

$$\mathbf{F}(\mathbf{x}, \mathbf{b}) = \mathbf{F}(\mathbf{x}_0, \mathbf{b}) + \mathbf{K}\{\mathbf{x} - \mathbf{x}_0\} + \mathcal{O}(\mathbf{x}^2)$$
(11)

with de Jacobian $\mathbf{K} = \frac{\partial \mathbf{F}}{\partial \mathbf{x}}(\mathbf{x}_0, \mathbf{b})$. Here, we distinguish between the state vector \mathbf{x} that comprises in its components the parameters to be retrieved and forward model parameters \mathbf{b} describing parameters.

160 eters other than the state vector that influence the measurement. Equation (11) represents a Taylor expansion of the forward model around state vector \mathbf{x}_0 truncated to first order.

2.2 Inversion

To determine the column density of water vapour isotopologues from SWIR measurements, we adjust the state vector \mathbf{x} to fit the forward model to the measurement vector \mathbf{y} , with spectral residuals

165 e_y, by a least squares fitting approach. The state vector x includes the total columns of CO, CH₄, H₂O, H₂¹⁸O and HDO, two coefficients to describe the linear spectral dependence of the surfacae albedo A_s, and a spectral shift of the ISRF to adjust the spectral calibration of the TROPOMI instrument per retrieval. We apply the profile scaling approach as described by Borsdorff et al. (2014) employing a Gauss-Newton iteration scheme. The least squares minimisation problem

170
$$\hat{\mathbf{x}} = \min_{\mathbf{x}} ||\mathbf{S}_{\mathbf{y}}^{-1/2}(\mathbf{F}(\mathbf{x}) - \mathbf{y})||^2$$
(12)

is solved per iteration step with the solution

$$\hat{\mathbf{x}} = \mathbf{x}_0 + \mathbf{G}(\mathbf{y} - \mathbf{F}(\mathbf{x}_0)) \tag{13}$$

with the gain matrix

$$\mathbf{G} = (\mathbf{K}^T \mathbf{S}_{\mathbf{v}}^{-1} \mathbf{K})^{-1} \mathbf{K}^T \mathbf{S}_{\mathbf{v}}^{-1}$$
(14)

175 and the measurement covariance matrix S_y .

After convergence, the column averaging kernel can be calculated in a straight forward manner:

$$\mathbf{A}_{k,k'} = \frac{dc_{\text{ret},k}}{d\rho_{k'}} = \mathbf{g}_k \mathbf{K}_{k'}^{\text{prof}} \,. \tag{15}$$

- g_k is the row vector of the gain matrix G that belongs to the trace gas k and K^{prof}_{k'} is the forward
 model Jacobian with respect to the trace gas profile ρ_{k'}. The height dependence of the profile ρ_{k'} is omitted for a clear presentation. For the full mathematical proof, the reader is referred to Borsdorff et al. (2014). For k ≠ k', A_{k,k'} describes the interference of the retrieved column c_k with the real trace gas vertical distribution of another trace gas k'. For k = k', it is the standard column averaging kernel and we use the more simple notation A_k = A_{k,k}.
- 185 The relation between our retrieval product $c_{ret,k}$ (the retrieved total column of species k) and the true state (concentration profile ρ_k) of the atmosphere is given by

$$c_{\text{ret},k} = \mathbf{A}_k \rho_k + \sum_{k' \neq k} \mathbf{A}_{k,k'} \rho_{k'} + \mathbf{e}_x , \qquad (16)$$





where e_x is the error on the retrieved total column due to the forward model and measurement errors e_y. In the case of a single trace gas retrieval the interference terms A_{k,k'} do not exist. In such case,
the meaning of the remaining column averaging kernel A_k is related to the proper choice of the reference profile and the effective null-space of the regularisation (see the discussion in Borsdorff et al. (2014) and Wassmann et al. (2015)). If the chosen reference profile is correct, the equation is equal to a geometrical integration of ρ_k. In the case of multiple trace gas retrievals we need to assess Eq. 16 in more detail. Using Eq. 6 the above equation can be written as

195
$$c_{\operatorname{ret},k} = \left[\mathbf{A}_k + \sum_{k' \neq k} \rho_{k,k'}^{\operatorname{rel}} \mathbf{A}_{k,k'} \right] \rho_k + \mathbf{e}_x ,$$
 (17)

showing that the contribution of the interference kernel can be interpreted as an error term for every level of the averaging kernel. Because atmospheric humidity can show strong variability (in both time and space), and the variability of HDO is (to first order) strongly correlated to the variability of the main water isotopologue, we are particularly interested in possible interferences between H_2O

and HDO. We want to be certain that a measured variability in HDO is truly caused by variations in HDO, and not a result of the interferences between H₂O and HDO. Therefore, we need to test if the interferences are small for the cases $k = H_2O$ and k' = HDO and vice versa.

Figure 2 shows an example of the column averaging kernels $A_{\rm H_2O}$ and $A_{\rm HDO}$, including the interference kernels multiplied with the relative profiles as in Eq. 17. The averaging kernel for

- 205 H_2O (A_{H_2O} , top left panel) shows that the retrieval is only sensitive to H_2O in the lower atmosphere. This is a result of strong pressure broadening of the H_2O absorption lines (Frankenberg et al., 2009). Since the HDO lines are weaker, the averaging kernel for HDO (A_{HDO} , lower left panel) is more uniform, showing only slightly lower sensitivity at high layers. The interference kernels show that above ~ 10 km variations of HDO have a minor impact on the retrieval of H_2O
- 210 $(\rho_{\text{H}_2\text{O},\text{HDO}}^{\text{rel}}\mathbf{A}_{\text{H}_2\text{O},\text{HDO}} \approx -0.02)$, top right panel in Fig. 2), and variations of H₂O have a small impact on the retrieval of HDO ($\rho_{\text{HDO},\text{H}_2\text{O}}^{\text{rel}}\mathbf{A}_{\text{HDO},\text{H}_2\text{O}} \approx -0.04 \pm 0.01$, lower right panel in Fig. 2). Since the column averaging kernels $\mathbf{A}_{\text{H}_2\text{O}}$ and \mathbf{A}_{HDO} are much larger, and the density profiles $\rho_{\text{H}_2\text{O}}$ and ρ_{HDO} will be very low higher in the atmosphere, the induced errors on the total columns due to this interference are practically negligible.
- 215 For a proper error characterisation of the retrieval product, we calculate the error covariance matrix $\mathbf{S_x}$ by

$$\mathbf{S}_{\mathbf{x}} = \mathbf{G}\mathbf{S}_{\mathbf{y}}\mathbf{G}^T \,. \tag{18}$$

This allows us to quantify the retrieval noise standard deviation σ_k of the individual column densities and a possible correlation between those.

For data interpretation, it is common to consider the relative abundance of HDO with respect to $H_2O: r = c_{ret,HDO}/c_{ret,H_2O}$, and to reference the ratio to the Vienna Standard Mean Ocean Water







Fig. 2. Top left: total column averaging kernel for H_2O . Lower left: total column averaging kernel for HDO. Top right: the sensitivity of total column H_2O to variations in HDO at different altitudes. Lower right: the sensitivity of total column HDO to variations in H_2O at different altitudes.

(VSMOW) ratio $r_s = 3.1153 \cdot 10^{-4}$:

$$\delta \mathbf{D} = \begin{bmatrix} \frac{r}{r_s} - 1 \end{bmatrix},\tag{19}$$

225

where δD is typically given in units of per mil. The first studies that measured δD in the atmosphere (as mentioned in the introduction) have shown that typical variations in δD over time or space are of the order of 50–100‰, which we therefore regard as the required accuracy for a useful product. The diagnostic tools of the individual columns can be used to derive the corresponding quantities for δD . For example, the standard deviation of the retrieval noise is given by

$$\sigma_{\delta \mathbf{D}} = \frac{r}{r_s} \sqrt{\frac{\sigma_{\rm H_2O}^2}{c_{\rm H_2O}^2} + \frac{\sigma_{\rm HDO}^2}{c_{\rm HDO}^2} - \frac{2\mathbf{S}_{\rm HDO,H_2O}}{c_{\rm H_2O}c_{\rm HDO}}}$$
(20)

and in a similar manner the column averaging kernel with respect to the $\rm H_2O$ and $\rm HDO$ abundance can be derived.





2.3 Two-band cloud filtering

Since the HDO and H₂O retrieval algorithm does not account for clouds nor any other scattering layers, we need to filter for clouds to avoid large retrieval inaccuracies. This filtering is achieved using the retrieved columns in a weak and strong absorption band of either CH₄ or H₂O. The bands used are indicated in Fig. 1. Elevated scattering layers not accounted for in the forward model generally cause a retrieval bias by scattering photons directly into the instrument. This optical path length shortening leads to negative biases in the retrieved total columns. The two-band cloud filter relies on the fact that a total column measurement using a strong absorption band is more strongly

- 240 affected by this "shielding bias" than the measurement using a weak absorption band. As a result, the relative difference in the retrieved total column between the weak and strong absorption band can be used to indicate the presence of clouds. Using a set of simulated measurements for varying cloudy conditions (as will be described in Sect. 3.1), we have tested that using a threshold < 6% for the relative difference in total column CH₄ between the weak and strong bands we filter for ground
- scenes that have a cloud fraction of more than 10–20% (cloud top height \geq 1 km). Scenes with low-level clouds (cloud top heights < 1 km) are not affected by this filter. Since low-level clouds above sea pass the filter, the retrieval allows for measurements over sea above these clouds due to their high albedo. The albedo of the sea surface itself is too low in the shortwave infrared for a meaningful retrieval above cloud-free sea pixels.
- As an example, Fig. 3 shows how the relative difference in retrieved methane absorption between the strong and weak absorption bands (shown in red in Fig. 1) changes for the scenarios with clouds (left panel), cirrus (middle panel) and aerosol (right panel). Scenes with strongest effects on the light path of the observed signal will also show the largest relative difference. We find that with a relative difference in methane absorption < 6% we effectively filter for clouds and cirrus, as well as for low
- 255 surface albedo scenes affected by aerosol. For example, not affected by the filter are scenes with a cloud top height ≤ 1 km or scenes with a low fraction of higher-level clouds (i.e. the contours with the two darkest shades of blue in the left panel of Fig. 3). A similar performance is achieved with a two-band water filter, using the weak and strong water bands as shown in blue in Fig. 1 and a threshold for the relative difference in water absorption in these bands of 8%. In the next sections the impact of this cloud filter on the retrievals of HDO and H₂O will be shown.

The two-band cloud filtering will be part of TROPOMI's operational methane pre-processing pipeline (Hu et al., 2016), and so synergies with the operational data processing can be used to reduce the processing time significanty, as we have estimated that on average 20% of all the measured ground scenes will pass the cloud filter above land, and 14% above sea.





Table 1. Overview of the generic scenarios used for the performance analysis. For the scenarios where the SZA was not variable, it was fixed at 50° . For the clouds scenario the surface albedo was fixed at 0.05.

Scenario	Variable X	Variable Y	
cloud free	surface albedo: 0.03-0.6	SZA: $0.0-70.0^{\circ}$	
clouds	cloud top height: 1-8 km	cloud fraction: 0.0-1.0	
cirrus	surface albedo: 0.03-0.6	$\tau_{\rm cir}$ (2300 nm): 0.0–1.0	
aerosol	surface albedo: 0.03-0.6	AOT (550 nm): 0.0-1.0	

265 3 Performance analysis for generic scenarios

To assess the performance of the retrieval algorithm, we applied the retrieval to simulated measurements for various generic scenarios. For each scenario, we systematically varied two variables such as surface albedo, solar zenith angle (SZA), cloud parameters (cloud top height, cloud fraction and cloud optical thickness (τ_{cld})) and aerosol optical thickness (AOT). An overview of the scenarios is given in Table 1.

3.1 Measurement simulations

270

The measurement simulations for the generic scenarios were created using the S-LINTRAN radiative transfer model (Schepers et al., 2014). The implementation of S-LINTRAN for TROPOMI simulations, including the instrument model, is described in detail in Landgraf et al. (2016), as the same simulations have been used to assess the performance of the CO retrieval algorithm. A summary of

the implementation is provided in the following two paragraphs.

The model is a scalar plane-parallel radiative transfer model that fully accounts for multiple elastic light scattering by clouds, cirrus, air molecules and the reflection of light by the Earth surface. The optical properties of water clouds are calculated using Mie theory with microphysical cloud

- 280 properties given in Table 2. For ice clouds the ray tracing model of Hess and Wiegner (1994); Hess et al. (1998) is employed assuming hexagonal, columnar ice crystals randomly oriented in space. Cirrus and water clouds are described by a cloud top and base height, and a cloud optical thickness. While cirrus fully cover the observed ground scene, water clouds can show partial cloud coverage by utilising the independent pixel approximation (Marshak et al., 1995) for the simulation.
- 285 Measurement noise was superimposed on the radiance spectra using the TROPOMI noise model (Tol et al., 2011). This assumed an observed ground scene of $7 \times 7 \text{ km}^2$ and a telescope aperture of $6 \cdot 10^{-6} \text{ m}^2$. The resulting signal to noise ratio is 120 in the continuum of the spectrum for a dark reference scene (surface albedo $A_s = 0.05$, viewing zenith angle VZA = 0° and solar zenith angle SZA = 70°).
- 290 The atmospheric model assumed the US standard atmosphere (1976) for the profiles of dry air





Table 2. Microphysical properties of water and ice clouds: n(r) represents the size distribution type, r_{eff} and v_{eff} are the effective radius and variance of the size distribution, m = n - ik is the refractive index. The ice cloud size distribution follows a power-law distribution as proposed by Heymsfield and Platt (1984).

	water clouds	ice clouds
n(r)	gamma	$(r/r_i)^{-3.85}$
r_{eff} [µm]	20	-
$v_{\rm eff}$	0.10	-
n	1.28	1.26
k	$4.7\cdot 10^{-4}$	$2.87\cdot 10^{-4}$

density, temperature, pressure, water and CO. The CH_4 profile is taken from the CAMELOT European background profile scenario (Levelt and Veefkind, 2009), interpolated to the same pressure grid and converted from mixing ratios to densities using the air densities from the US standard atmosphere. We separated the water profile into individual profiles for the three isotopic components

- 295 with absorption features in the TROPOMI SWIR range: $H_2^{16}O$, $H_2^{18}O$ and HDO. First, the water profile was scaled with the VSMOW abundance of the respective species. Additionally, a realistic altitude-dependent depletion of HDO and $H_2^{18}O$ was assumed. For HDO we assumed a linear decrease from $\delta D = -100 \%$ at the surface to $\delta D = -600 \%$ at 15 km, followed by a linear increase to $\delta D = -400 \%$ at the top of the atmosphere at an altitude of 48 km (Ehhalt, 1974; Ehhalt et al.,
- 300 2005; Schneider et al., 2010). We further assumed that the concentration of H₂¹⁸O is related to the concentration of HDO according to the empirically determined "global meteoric water line" (Craig, 1961)

$$\delta \mathbf{D} = 8 \cdot \delta^{18} \mathbf{O} + 10\%, \tag{21}$$

where δ^{18} O is defined in the same way as δ D (Eq. 19). All the atmospheric profiles used for the 305 measurement simulations are shown in Fig. 4.

In the following subsections, we characterise the retrieval performance for the generic scenarios considering separately the retrieval statistical errors (i.e. the single measurement noise) σ_{H_2O} , σ_{HDO} and $\sigma_{\delta D}$ (neglecting the small HDO–H₂O cross-correlation term for $\sigma_{\delta D}$) and the biases in the total columns c_{ret,H_2O} , $c_{ret,HDO}$ and their ratio δD . The retrieval statistical error estimate for δD is

310 given by Eq. 20. For the bias in δD , which we refer to as " $\Delta \delta D$ ", we first determine $\delta D_{retrieval}$ by removing the noise on the retrieved total columns HDO and H¹⁶₂O using linear error propagation for the particular noise realisation. We need to compare $\delta D_{retrieval}$ with δD_{model} , where δD_{model} is δD





of the "true" model atmosphere:

$$\delta \mathbf{D}_{\text{model}} = \frac{c_{\text{true,HDO}}}{c_{\text{true,H_2O}}} \frac{1}{r_s} - 1.$$
(22)

315 Finally, the retrieval bias on δD is defined as:

 $\Delta \delta \mathbf{D} = \delta \mathbf{D}_{\text{retrieval}} - \delta \mathbf{D}_{\text{model}} \,. \tag{23}$

3.2 Cloud-free conditions

In Fig. 5 we show the simulated cloud-free retrieval bias for the total column H_2O (left panel), total column HDO (middle panel) and their ratio ($\Delta\delta D$, right panel), as a function of surface albedo and

320 SZA (no clouds or aerosol present). The figure shows that the retrieval performs very well for the majority of the scenes, with $\Delta\delta D$ less than 0.8%. Only for the lowest surface albedos (0.03–0.05) the bias in δD increases to a few per mil, due to slightly more negative bias in H₂O compared to HDO.

The corresponding statistical error estimates are shown in Fig. 6. σ_{H_2O} reaches maximum values 325 of 1.6–2.0% for the lowest surface albedos and highest SZAs and σ_{HDO} is about a factor 2 larger due to the weaker HDO absorption features. Combined, it results in values for $\sigma_{\delta D}$ of the order of 15–25‰ for the lowest surface albedos. For high surface albedo regions such as deserts (surface albedo ~ 0.3 in the SWIR) typical values for $\sigma_{\delta D}$ are 2–4‰. This is roughly an order of magnitude better than what is achieved with SCIAMACHY (Scheepmaker et al., 2015).

330 3.3 Clouds and cirrus

As the retrieval algorithm does not account for scattering, any clouds, cirrus and aerosol present in the observed scene will lead to biases in the retrieval of the total columns HDO and H₂O. We have tested the performance of the retrieval under cloudy conditions with a scenario assuming a cloud with an optical thickness of $\tau_{cld} = 5$, with varying cloud top heights (between 1–8 km in steps of

- 335 1 km) and varying cloud fractions (between 0.0–1.0 in steps of 0.1). The same scenario was used to demonstrate the two-band cloud filter in the left panel of Fig. 3. Due to differences in their retrieval sensitivities, the observed bias is stronger for H₂O than for HDO, leading to significant biases in their ratio, increasing with both cloud fraction and cloud top height (although not shown, we find that $\Delta\delta D$ can reach values > 900% for clouds above 7 km with 100% cloud coverage). Similarly,
- 340 by simulating scenarios with varying surface albedos and a uniform cirrus or aerosol layer with varying optical thickness (see Table 1), we find that this bias increases with the optical thickness of the layer and with lower surface albedos, as both lead to a lower contribution of photons from below the scattering layer reaching the instrument. As described in Sect. 2.3, the two-band cloud filtering technique will be used to pre-filter the scenes most affected by this shielding bias. We find that, after





applying the two-band methane filter to scenes affected by clouds and cirrus, $\Delta \delta D \lesssim 70\%$ and $\sigma_{\delta D} = 10-20\%$ (for scenes with a low surface albedo of $A_s = 0.05$).

3.4 Aerosol

An aerosol layer will typically have a lower optical thickness than clouds, and occurs lower in the atmosphere, leading to a different impact on our non-scattering retrieval. Our aerosol scenario

- 350 assumes a uniform layer of a sulphate-type aerosol in the boundary layer between 0–2 km. Figure 7 shows how this induces a bias in the total columns H₂O, HDO and their ratio, as a function of aerosol optical thickness and surface albedo. We see that for very low surface albedos, direct reflection off the aerosol layer leads to path length shortening and a corresponding negative (shielding) bias for the total column H₂O. This effect is weaker for HDO due its more uniform averaging kernel. For
- 355 higher surface albedos, however, we see that the bias becomes positive, likely due to an increased amount of light scattering in the boundary layer. The contribution of photons from the brighter surface increases and a fraction of these photons undergo multiple scattering events between the aerosol layer and the surface, enhancing the path length. The net effect on δD is that its bias due to aerosol is highest for the lowest surface albedos and highest AOT (right panel in Fig. 7). If we
- 360 take the two-band cloud filter into account (right panel in Fig. 3) to filter the lowest surface albedos affected by aerosol, we are left with $\Delta\delta D \lesssim 20\%$ due to boundary layer aerosol with AOT = 1.0 (at 550 nm). The statistical error (not shown) does not depend significantly on AOT, but varies primarily with surface albedo, reaching similar peak values as in the cloud free scenario ($\sigma_{\delta D} \approx 20\%$).

3.5 Summary of the general performance

- 365 In summary, we can conclude that the retrieval performs well under cloud-free conditions. The bias $\Delta\delta D$ will be less than 2‰, even for the lowest surface albedos, and the statistical errors vary from 2–4‰ for high albedos to 15–25‰ for the lowest albedos. Under conditions with clouds, cirrus or aerosol the retrieval performs less well and we generally find a positive bias in δD . To restrict this bias we need strict filtering against clouds and aerosol by applying the two-band cloud filter either
- 370 to methane or water (which additionally leads to a great reduction in the computational effort). Applying a two-band methane threshold of 6%, we restrict the bias in δD to $\Delta \delta D < 70\%$ for all simulated measurements. Averaging multiple single measurements over time and space will further reduce the statistical error and will improve the accuracy to better than the maximum 70%, bringing the measurements within the requirements to study typical temporal and spatial gradients (which are
- 375 of the order of 50–100%*c*).





Table 3. Summary of the sensitivity to the meteorological input and instrument parameters, expressed as the mean difference in δD between the perturbed and default retrievals for the 45 scenes with the three lowest surface albedos (0.03, 0.05 and 0.075).

Prior parameter	Systematic error in δD [%]		
Temp -0.5 K	-6.9	±	0.74
Temp $+0.5 \text{ K}$	+7.0	±	0.11
Temp −1 K	-14	±	0.74
Temp +1 K	+14	\pm	0.17
Pressure $\times 0.99$	-4.5	±	0.74
Pressure $\times 1.01$	+4.5	\pm	0.54
Rad. offset $+0.1\%$	-0.054	±	0.11
Rad. offset $+0.5\%$	-0.20	\pm	0.16
ISRF FWHM -1%	+0.36	±	0.85
ISRF FWHM $+1\%$	-0.33	±	0.84

4 Sensitivity to prior assumptions

Similarly to what was done for the CO TROPOMI retrievals (Landgraf et al., 2016), we have tested the sensitivity of the H₂O and HDO total column retrievals to the prior assumptions, including the impact on δD. These so-called forward modelling errors were tested on the cloud-free scenario
(with varying SZA and surface albedo) using the same measurement simulation as described in Sect. 3. A perturbation in one of the input assumptions was introduced, after which the retrievals were performed and compared with the default retrievals without the perturbation. The impact of the perturbation is expressed as a systematic error and standard deviation, where we define the systematic error as the mean difference in δD between the perturbed and default retrievals for the 45 scenes with the three lowest surface albedos (0.03, 0.05 and 0.075). The results are summarised in

000

Table 3.

The retrieval uses a priori temperature profiles and surface pressures from the ECMWF. To test the impact of uncertainties in the temperature profile, we have varied this profile by ± 0.5 K and ± 1 K. This is primarily affecting the retrieved total column H₂O, while the total column HDO is not very

390 sensitive to temperature variations. A perturbation of +1 K (-1 K) leads to a decrease (increase) in the retrieved H₂O column of 1.8%, inducing a systematic error in δ D of +14‰ (-14‰). This error is constant for all surface albedos and SZAs and scales linearly with the size of the temperature perturbation.

The atmospheric pressure profile is derived from the surface pressure. To test the impact on





395 inaccuracies in the ECMWF surface pressure, we applied a perturbation of $\pm 1\%$. This leads to systematic errors of about 0.5% in H₂O and 0.13% in HDO (with reversed sign), together inducing errors of about 4.5% in δD .

The retrieval algorithm requires a reflectance spectrum, acquired by dividing the radiance spectrum measured from the Earth's surface by the irradiance spectrum measured directly from the Sun.

- 400 Differences in the radiometric offset between these spectra could induce spectral features in the reflectance spectrum, leading to systematic errors. The TROPOMI instrument requirement for the radiometric offset on the radiance is 0.1% of the continuum level. We have tested the impact of an offset on the radiance of 0.1% and 0.5% of the maximum value in the retrieval window. However, the retrieval fits for an offset in the reflectance spectra, which partly mitigates the effects of an offset
- 405 in the radiance or irradiance. The systematic errors due to uncertainties in the radiometric offset are therefore very small (errors in δD less than 0.5%).

For the default retrieval and measurement simulations we have assumed a Gaussian slit function (ISRF) with a FWHM of 0.25 nm. We have tested the impact of perturbing this FWHM by $\pm 1\%$ and find that the induced systematic errors are strongly dependent on surface albedo and SZA. The

410 largest errors in δD reach $\pm 3\%$ and are found for high albedos and low SZAs. The mean systematic error for the lowest albedos is $0.36 \pm 0.85\%$.

In summary (also see Table 3), we find that the retrieval algorithm is most sensitive to uncertainties in the a priori temperature profiles, followed by the pressure profiles. The sensitivity to uncertainties in the instrument parameters is about an order of magnitude smaller. The uncertainties in the input

415 profiles are expected to be mostly quasi-random in nature, which means their impact on the error in δD will diminish when taking averages in time and space.

More structural systematic errors (i.e. those that will not diminish by averaging) are potentially caused by uncertainties in the water spectroscopy. Recent studies have shown that spectroscopic uncertainties of water can have a large impact on total column retrievals of CO (Galli et al., 2012), CH_4

- 420 (Frankenberg et al., 2008; Schneising et al., 2009), H₂O (Schrijver et al., 2009) and the HDO/H₂O ratio (Scheepmaker et al., 2013). As a test of the possible impact of uncertainties in the water line parameters, we have repeated the retrievals of the simulated clear-sky scenario after replacing the line parameters of the water isotopologues. For the simulated spectra the parameters from HITRAN (Rothman et al., 2009) were used. We then performed the retrievals using the water line parameters are a second second
- 425 ters from Scheepmaker et al. (2013). Table 4 shows the induced systematic errors for replacing a single isotopologue at a time, and for replacing all modelled water isotopologues simultaneously. It shows that the retrieval of HDO and H_2O can be very sensitive to spectroscopic uncertainties, especially in the ratio HDO/H₂O, since HDO and $H_2^{16}O$ can show sensitivities with opposite sign, which strengthen each other when taking the ratio (as can be seen from replacing only the $H_2^{16}O$
- 430 parameters). The differences in spectroscopy between HITRAN and Scheepmaker et al. (2013) can lead to differences in δD of up to 128%. Although we find that the differences do not depend on





Table 4. Sensitivity to a change of water line parameters from HITRAN2008 to the parameters from Scheepmaker et al. (2013).

Isotopologue	Systematic Error				
	${ m H}_{2}^{16}{ m O}$ [%]	HDO [%]	CH ₄ [%]	δD [‰]	
$\mathrm{H}_{2}^{16}\mathrm{O}$	$+8.0\pm0.55$	-4.6 ± 0.50	$+1.3\pm0.35$	-97 ± 1.5	
HDO	-0.069 ± 0.036	-4.2 ± 0.043	-0.040 ± 0.037	-34 ± 0.29	
$\mathrm{H}_{2}^{18}\mathrm{O}$	$+0.011 \pm 0.017$	-0.11 ± 0.0090	-0.029 ± 0.017	-0.97 ± 0.21	
All	$+7.9\pm0.53$	-8.7 ± 0.50	$+1.3\pm0.37$	-128 ± 1.5	

surface albedo or SZA, we cannot exclude a dependency on the total amount of water vapour, that might lead to seasonal and latitudinal biases. Similar to the retrieval of CO (Galli et al., 2012), the HDO/H_2O retrieval will very likely benefit from a reassessment of the spectroscopic line parameters

435 of water, a study which is currently on-going (Loos et al., 2015). Regardless of such reassessments, validation studies will be needed to verify spectroscopy and to define corrections that might mitigate spectroscopy related biases.

5 Performance analysis for a realistic scenario

To show the capabilities of the TROPOMI H₂O and HDO total column retrievals, we have sim-440 ulated an ensemble of measurements that reflect a realistic scenario as accurately as possible. In Sect. 5.1 we describe the input data and measurement simulations in more detail. In Sect. 5.2 we discuss the results of retrieving the simulated measurements, in terms of retrieval bias, precision, and effectiveness of the cloud filtering.

5.1 Measurement simulations

- 445 The simulated measurement ensemble covers a region over the South-West part of the US and the North-East part of Mexico as TROPOMI would observe it on August 4th 2009. Figures 8 and 9 show the region in terms of various input fields. This region comprises a clear gradient in the relative abundance of HDO with respect to H_2O due to the transport of humid air from coastal regions inland. We have combined data from the MODIS Aqua satellite (clouds, land/water coverage, sur-
- 450 face albedo, aerosol) with data from ETOPO5 (elevation), ECMWF (surface pressure, temperature profiles, specific humidity), TM5 (CO and CH_4 profiles) and LMDZiso (HDO and $H_2^{18}O$ profiles) to simulate 27405 TROPOMI measurements on a grid of 135 by 203 ground pixels. This ensemble represents roughly what TROPOMI will observe in five minutes with a daily revisiting cycle.

The viewing and solar geometry and ground pixel size were adapted from MODIS Aqua granule 455 2009216 (19h45m UT), where the MODIS information was spatially resampled on a pixel size of





 $10 \times 10 \text{ km}^2$ at sub-satellite point. The pixel distortion towards the outer swath was adopted from the MODIS observation. The surface reflection was estimated from the MODIS MCD43C4 data product at 2105–2155 nm for the same period (Strahler et al., 1999) in combination with the surface elevation from ETOPO5 (NOAA, 1988). Furthermore, the MODIS MYD06 cloud product (Platnick

- 460 et al., 2015) was used to estimate cloud cover and cloud top height for the individual TROPOMI ground pixels. Only clouds with top height above 100 meters were used. We derived cirrus optical thicknesses from the MODIS cirrus reflectance product employing the algorithm by Dessler and Yang (2003). For all pixels, the cirrus was located between 9–10 km. For the aerosol optical thickness (at 550 nm) we used the MODIS MYD08_M3 global monthly mean product (Platnick,
- 465 2015), resampled to the above mentioned granule with a pixel size of $10 \times 10 \text{ km}^2$ at sub-satellite point. For some pixels with missing aerosol data the optical thickness was set to 0.1. We assumed three different aerosol types: "oceanic" above water, "dust" above land and "urban" above all land regions with AOT > 0.23. The corresponding model fields are depicted in Fig. 8.

The distribution of atmospheric trace gases was estimated using TM5 chemistry model simula tions (Krol et al., 2005), which yields the CO and CH₄ abundances. Moreover, we used data from ECMWF (Dee et al., 2011) for the atmospheric pressure, temperature and humidity profiles. For realistic HDO/H₂O ratios we derived δD profiles from LMDZiso model simulations (Risi et al.,

2010) and for the δ^{18} O profiles we assumed correlation to δ D according the global meteoric water line (also see Eq. 21, Craig, 1961). Figure 9 shows the resulting total columns for the most impor-

475 tant species for our ensemble. Based on this input and the TROPOMI instrument model as described in Sect. 3.1, we simulated for each individual pixel the TROPOMI SWIR observations using the S-LINTRAN radiative transfer model.

5.2 Results

Using the simulated measurement ensemble we retrieved the water vapour abundances using the
SICOR retrieval algorithm including the retrieval of the two methane and water bands used for cloud filtering. In Fig. 10 the results are shown in terms of the retrieved bias in total column H₂O, HDO and δD. We also show the relative difference in the weak vs strong water bands that was used for cloud filtering. The cloud filter panel (lower-left panel in Fig. 10) shows that the algorithm retrieved some pixels above the Gulf of Mexico, even though these pixels did not contain low-level clouds. In

- 485 reality such pixels will be removed by pre-filtering for very low albedo regions. Once the two-band cloud filter threshold is applied (keeping only pixels with a relative difference < 8% using the water bands), practically all ocean pixels are removed, as well as all the lands pixels affected by clouds, resulting in 54.5% of the pixels remaining for further study. Using the two methane bands as a cloud filter with a threshold of 6% resulted in slightly less strict filtering (60.7% of the pixels remaining),</p>
- 490 as certain pixels with low and optical thin clouds in the west above Mexico and in the east above Alabama were not removed (not shown). The few rejected pixels in the centre and south of Texas





495

show that the cloud filter effectively removed high isolated clouds with low optical thickness (cf. the lower two panels in Fig. 8 with Fig. 10), but left pixels with clouds < 1 km intact (as is preferred). The large group of pixels in the north-east of the ensemble were rejected based on the presence of high and optically thick clouds, or the presence of aerosol above low surface albedo regions.

The other three panels in Fig. 10 show the remaining biases in total column H_2O , HDO and δD after cloud and ocean filtering. Both H_2O and HDO show a positive bias of a few percent above the higher surface albedo regions in the west and a small negative bias over the lower surface albedo regions in the states of Louisiana and Mississippi show that even the

- solution albedo contrast caused by the Mississippi and Red River basins can be observed in the H₂O and HDO bias maps. We also see that the biases are slightly larger for H₂O, compared to HDO. The cause for these biases is aerosol, as the aerosol bias shows the same patterns as a function of surface albedo and AOT, and the effect is slightly larger for H₂O, as discussed in Sect. 3.4 and shown in Fig. 7. Combined into a ratio, the lower-right panel in Fig. 10 show that the retrieval bias in δD
- 505 is slightly negative above the highest albedos and increases to a positive bias with a maximum of $\sim 20\%$ above the lowest albedo regions. Areas at high altitudes usually have a lower humidity and therefore a lower δD compared to areas at lower altitudes. This gradient is visible in the bottom right panel in Fig. 9. Furthermore, areas at higher altitude generally have a higher surface albedo. Because the retrieval bias in δD is negative for high surface albedos and positive for low surface
- state albedos, the altitude gradient in δD is overestimated by the retrieval. This will likely be the case for all scenarios with a gradient between higher elevated (or drier) areas and lower elevated (or more humid) areas.

Figure 11 shows the same maps in terms of single measurement precision error (1σ) . As expected, the dominant factor to determine the precision error is surface albedo. The error of the strong ab-

- 515 sorbers H_2O and CH_4 is 0.05–0.15% for the highest surface albedos, and increases to 0.35% for the lowest surface albedos. The precision errors of the weak absorber HDO are larger, reaching 0.50% for the lowest surface albedos. This translates into precision errors in δD of at most 5‰ above the lowest albedos.
- This realistic scenario demonstrates the capabilities of TROPOMI and the SICOR algorithm to 520 retrieve accurate patterns in total column H₂O, HDO and δ D above land from a single overpass. After the two-band cloud filter effectively removed all measurements above water and high clouds, a small bias remains due to aerosol, which correlates with surface albedo. The bias is smaller (and of opposite sign) compared to the temporal or spatial gradients in δ D expected for typical science cases (e.g., as observed by SCIAMACHY in Yoshimura et al., 2011; Lee et al., 2012; Risi et al.,
- 525 2012a; Okazaki et al., 2015). This ensures the ability to detect and study patterns in δD on much smaller timescales and at higher spatial resolution compared to previous satellite missions, but it should also be kept in mind to be careful when using the data over regions with strong gradients in surface albedo.





6 Discussion and conclusions

- 530 We have presented an algorithm and performance analysis for the retrieval of total column H₂O and HDO from TROPOMI measurements onboard the Sentinel-5 Precursor mission. By adapting ESA's operational CO algorithm (Landgraf et al., 2016), we developed a relatively simple approach that is fast but relies on strict filtering for clouds, cirrus and aerosol using a two-band methane or water retrieval. The ratio HDO/H₂O will be a useful scientific product in the fields of hydrology and climate research, with the potential to improve our understanding of the processes controlling
 - atmospheric humidity and transport.

The first studies in these directions that used a similar type of column-averaged satellite product were using SCIAMACHY data (Frankenberg et al., 2009; Yoshimura et al., 2011; Lee et al., 2012; Risi et al., 2012b,a; Scheepmaker et al., 2015). These studies showed that the typical seasonal or

- spatial gradients in δD are about 50–100‰. The measurement precision and accuracy needs to be higher than this in order to contribute significantly to the science. For SCIAMACHY, this implied either taking monthly averages or binning to a spatial resolution of at least $1 \times 1^{\circ}$ in order to bring the statistical error down to about 20‰ (the single measurement precision being ~ 115‰, Scheepmaker et al., 2015). The newer GOSAT measurements show an improvement in precision by a factor
- 545 of about two, compared to SCIAMACHY (Frankenberg et al., 2013; Boesch et al., 2013). Both SCIAMACHY and GOSAT products show a negative bias of about 30–70‰ compared to groundbased Fourier-transform spectroscopy (FTS) networks.

Our analysis has shown that TROPOMI is expected to deliver a much better performance than SCIAMACHY and GOSAT in terms of δD in only a single overpass. The single measurement noise

- 550 will be better than 15-25% for even the lowest surface albedos, while at the same time the spatial resolution of 7×7 km² is much higher than SCIAMACHY's 120×30 km² and provides a better coverage than GOSAT's sparse spatial sampling. Even though we still need to filter for clouds, due to this higher spatial resolution TROPOMI will observe many more useful scenes in between clouds compared to SCIAMACHY or GOSAT. This allows for new opportunities of studying the
- 555 hydrological cycle on timescales of mere days or weeks instead of seasons or years, or over longer periods if a high spatial resolution is desired.

Mainly due to the presence of low-level aerosol in the atmosphere, the cloud-filtered TROPOMI measurements of total column HDO and H_2O are not expected to be completely bias free. Changes to the light paths of the reflected photons due to any scattering particles remaining after filtering are

560 not accounted for in the retrieval algorithm, and lead to biases of a few percent in total column HDO and H₂O, and up to $\sim 20\%$ in δ D, depending on surface albedo, as shown by our simulated scenario of measurements above the USA and Mexico.

After launch and commissioning of the instrument in Q4 2016, validation using ground-based FTS data from the TCCON and NDACC networks is needed to test the performance of the algorithm on real measurements. Ideally, the HDO/H₂O products from these networks should first be intercom-





pared, both using the results from the on-going reassessment of the water spectroscopy (Loos et al., 2015), and for a range of atmospheric conditions and geographical locations. Any possible differences due to either spectroscopy or location (e.g. as found by Scheepmaker et al., 2015) need to be understood before the next generation of HDO and H_2O global retrievals from space can be ex-

570 ploited to come to a better understanding of the atmospheric hydrological cycle and the role it plays in our changing climate.

Acknowledgements. LMDZiso simulations used the computing resources of IDRIS under the allocation 0292 made by GENCI.





References

- 575 Aggarwal, P. K., Gat, J. R., and Froehlich, K. F.: Isotopes in the Water Cycle: Past, present and future of a developing science, Springer, Dordrecht, The Netherlands, 2005.
 - Boesch, H., Deutscher, N. M., Warneke, T., Byckling, K., Cogan, A. J., Griffith, D. W. T., Notholt, J., Parker, R. J., and Wang, Z.: HDO/H₂O ratio retrievals from GOSAT, Atmos. Meas. Tech., 6, 599–612, doi:10.5194/amt-6-599-2013, 2013.
- 580 Borsdorff, T., Hasekamp, O. P., Wassmann, A., and Landgraf, J.: Insights into Tikhonov regularization: application to trace gas column retrieval and the efficient calculation of total column averaging kernels, Contrib. Atmos. Phys., 7, 523–535, doi:10.5194/amt-7-523-2014, 2014.
 - Boucher, O., Randall, D., Artaxo, P., Bretherton, C., Feingold, G., Forster, P., Kerminen, V.-M., Kondo, Y., Liao, H., Lohmann, U., Rasch, P., Satheesh, S., Sherwood, S., Stevens, B., and Zhang, X.: Clouds and
- 585 Aerosols. In: Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change [Stocker, T.F., D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley (eds.)], Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2013.
- Craig, H.: Isotopic Variations in Meteoric Waters, Science, 133, 1702–1703, 590 doi:10.1126/science.133.3465.1702, 1961.
 - Craig, H. and Gordon, L.: Deuterium and oxygen 18 variations in the ocean and marine atmosphere, in: Stable Isotopes in Oceanographic Studies and Paleotemperatures, Spoleto, Italy, edited by Tongiogi, E., pp. 9–130, V. Lishi e F., Pisa, Italy, 1965.

Dansgaard, W.: Stable isotopes in precipitation, Tellus, 16, 436-468, 1964.

- 595 Dansgaard, W., Johnsen, S. J., Møller, J., and Langway, C. C.: One Thousand Centuries of Climatic Record from Camp Century on the Greenland Ice Sheet, Science, 166, 377–380, doi:10.1126/science.166.3903.377, 1969.
 - Dee, D. P., Uppala, S. M., Simmons, A. J., Berrisford, P., Poli, P., Kobayashi, S., Andrae, U., Balmaseda, M. A., Balsamo, G., Bauer, P., Bechtold, P., Beljaars, A. C. M., van de Berg, L., Bidlot, J., Bormann, N.,
- 600 Delsol, C., Dragani, R., Fuentes, M., Geer, A. J., Haimberger, L., Healy, S. B., Hersbach, H., Hólm, E. V., Isaksen, L., Kållberg, P., Köhler, M., Matricardi, M., McNally, A. P., Monge-Sanz, B. M., Morcrette, J.-J., Park, B.-K., Peubey, C., de Rosnay, P., Tavolato, C., Thépaut, J.-N., and Vitart, F.: The ERA-Interim reanalysis: configuration and performance of the data assimilation system, Q. J. Roy. Meteor. Soc., 137, 553–597, doi:10.1002/qj.828, 2011.
- 605 Dessler, A. E. and Yang, P.: The Distribution of Tropical Thin Cirrus Clouds Inferred from Terra MODIS Data, J. Climate, 16, 1241–1247, 10.1175/1520–0442, 2003.
 - Ehhalt, D. H.: Vertical profiles of HTO, HDO, and H2O in the troposphere, NCAR Tech. Note NCAR-TN-STR-100, 1974.
- Ehhalt, D. H., Rohrer, F., and Fried, A.: Vertical profiles of HDO/H2O in the troposphere, J. Geophys. Res.,110, 13 301, doi:10.1029/2004JD005569, 2005.
- Frankenberg, C., Bergamaschi, P., Butz, A., Houweling, S., Meirink, J. F., Notholt, J., Petersen, A. K., Schrijver, H., Warneke, T., and Aben, I.: Tropical methane emissions: A revised view from SCIAMACHY onboard ENVISAT, Geophys. Res. Lett., 35, 15811, doi:10.1029/2008GL034300, 2008.





Frankenberg, C., Yoshimura, K., Warneke, T., Aben, I., Butz, A., Deutscher, N., Griffith, D., Hase,
F., Notholt, J., Schneider, M., Schrijver, H., and Röckmann, T.: Dynamic Processes Governing Lower-Tropospheric HDO/H2O Ratios as Observed from Space and Ground, Science, 325, 1374–1377, doi:10.1126/science.1173791, 2009.

Frankenberg, C., Wunch, D., Toon, G., Risi, C., Scheepmaker, R., Lee, J.-E., Wennberg, P., and Worden, J.: Water vapor isotopologue retrievals from high-resolution GOSAT shortwave infrared spectra, Atmo-

- 620 spheric Measurement Techniques, 6, 263–274, doi:10.5194/amt-6-263-2013, http://www.atmos-meas-tech. net/6/263/2013/, 2013.
- Galli, A., Butz, A., Scheepmaker, R. A., Hasekamp, O., Landgraf, J., Tol, P., Wunch, D., Deutscher, N. M., Toon, G. C., Wennberg, P. O., Griffith, D. W. T., and Aben, I.: CH₄, CO, and H₂O spectroscopy for the Sentinel-5 Precursor mission: an assessment with the Total Carbon Column Observing Network measurements, Atmos. Meas. Tech., 5, 1387–1398, doi:10.5194/amt-5-1387-2012, 2012.
- Herbin, H., Hurtmans, D., Clerbaux, C., Clarisse, L., and Coheur, P.-F.: H216O and HDO measurements with IASI/MetOp, Atmos. Chem. Phys., 9, 9433, 2009.
 - Hess, M. and Wiegner, M.: COP: a data library of optical properties of hexagonal ice crystals, ao, 33, 7740– 7746, 1994.
- 630 Hess, M., Koelemeijer, R. B. A., and Stammes, P.: Scattering matrices of imperfect hexagonal ice crystals, jqrst, 60, 301–308, doi:10.1016/S0022-4073(98)00007-7, 1998.

Heymsfield, A. J. and Platt, C. M. R.: A Parameterization of the Particle Size Spectrum of Ice Clouds in Terms of the Ambient Temperature and the Ice Water Content., jas, 41, 846–855, doi:10.1175/1520-0469(1984)041<0846:APOTPS>2.0.CO;2, 1984.

635 Hu, H., Hasekamp, O., and Landgraf, J.: Bla, Atmos. Meas. Tech., p. to be submitted, 2016.

Jouzel, J., Russell, G. L., Suozzo, R. J., Koster, R. D., White, J. W. C., and Broecker, W. S.: Simulations of the HDO and H₂¹⁸O atmospheric cycles using the NASA/GISS general circulation model: The seasonal cycle for present-day conditions, J. Geophys. Res., 92, 14739–14760, doi:10.1029/JD092iD12p14739, 1987.

Jouzel, J., Alley, R. B., Cuffey, K. M., Dansgaard, W., Grootes, P., Hoffmann, G., Johnsen, S. J., Koster, R. D.,

640 Peel, D., Shuman, C. A., Stievenard, M., Stuiver, M., and White, J.: Validity of the temperature reconstruction from water isotopes in ice cores, J. Geophys. Res., 102, 26471–26488, doi:10.1029/97JC01283, 1997.

- 645 Krol, M., Houweling, S., Bregman, B., van den Broek, M., Segers, A., van Velthoven, P., Peters, W., Dentener, F., and Bergamaschi, P.: The two-way nested global chemistry-transport zoom model TM5: algorithm and applications, Atmos. Chem. Phys., 5, 417–432, doi:10.5194/acp-5-417-2005, 2005.
- Kurylo, M. and Solomon, S.: Network for the Detection of Stratospheric Change: A Status and Implementation Report, NASA, Upper Atmosphere Research Program and NOAA Climate and Global Change Program,
 Washington DC, 1990.
 - Landgraf, J., aan de Brugh, J., Scheepmaker, R., Borsdorff, T., Houweling, S., Aben, I., Hu, H., Hasekamp, O., and Butz, A.: Carbon Monoxide total column retrievals from TROPOMI shortwave infrared measurements, Atmos. Meas. Tech., p. to be submitted, 2016.

Kasten, F. and Young, T.: Revised optical air mass tables and approximation formula, Appl. Opt., 28, 4735–4738, 1989.





665

- Lee, J.-E., Risi, C., Fung, I., Worden, J., Scheepmaker, R. A., Lintner, B., and Frankenberg, C.: Asian monsoon hydrometeorology from TES and SCIAMACHY water vapor isotope measurements and LMDZ simulations: Implications for speleothem climate record interpretation, J. Geophys. Res., 117, D15112, doi:10.1029/2011JD017133, 2012.
 - Levelt, P. and Veefkind, P.: ESA CAMELOT study: Challenges in future operational missions for GMES atmospheric monitoring, sentinel 4 and 5, in: EGU General Assembly Conference Abstracts, edited by
- 660 Arabelos, D. N. and Tscherning, C. C., vol. 11 of EGU General Assembly Conference Abstracts, p. 8911, 2009.
 - Loos, J., Birk, M., Wagner, G., Mondelain, D., Kassi, S., Vasilchenko, S., Campargue, A., Hase, F., Dufour, G., Eremenko, M., Orphal, J., Cuesta, J., Bigazzi, A., Perrin, A., Daumont, L., and Zehner, C.: Spectroscopic database for TROPOMI/Sentinel-5 Precursor, in: ESA ATMOS 2015 Conference Proceedings (ESA SP-735), 2015.
 - Marshak, A., Davis, A., Wiscombe, W., and Titov, G.: The verisimilitude of the independent pixel approximation used in cloud remote sensing, rse, 52, 71–78, doi:10.1016/0034-4257(95)00016-T, 1995.
 - Mook, W. G.: Environmental Isotopes in the Hydrological Cycle. Principles and Applications, IAEA and UN-ESCO, http://www-naweb.iaea.org/napc/ih/IHS_resources_publication_hydroCycle_en.html, 2000.
- 670 NOAA, 1988: Data Announcement 88-MGG-02, Digital relief of the Surface of the Earth, 1988. Okazaki, A., Satoh, Y., Tremoy, G., Vimeux, F., Scheepmaker, R., and Yoshimura, K.: Interannual variability of isotopic composition in water vapor over western Africa and its relationship to ENSO, Atmos. Chem. Phys., 15, 3193–3204, doi:10.5194/acp-15-3193-2015, 2015.

Platnick, S., King, M. D., G., M. K., Wind, G., N., A., B., M., T., A. G., Z., Z., A., H. P., B., R., and J., R.:

675 MODIS Cloud Optical Properties: User Guide for the Collection 6 Level-2 MOD06/MYD06 Product and Associated Level-3 Datasets, Report, NASA MODIS Adaptive Processing System. Goddard Space Flight Center, USA, http://modis-atmos.gsfc.nasa.gov/_docs/C6MOD060PUserGuide.pdf, 2015.

Platnick, S. e. a.: MODIS Atmosphere L3 Monthly Product, doi:10.5067/MODIS/MOD08_M3.006, 2015.

- Randall, D., Wood, R., Bony, S., Colman, R., Fichefet, T., Fyfe, J., Kattsov, V., Pitman, A., Shukla, J., Srini-vasan, J., Stouffer, R., Sumi, A., and Taylor, K.: Climate Models and Their Evaluation. In: Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change [Solomon, S., D. Qin, M. Manning, Z. Chen, M. Marquis, K.B. Averyt, M.Tignor and H.L. Miller (eds.)], Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 2007.
- 685 Risi, C., Bony, S., Vimeux, F., and Jouzel, J.: Water-stable isotopes in the LMDZ4 general circulation model: Model evaluation for present-day and past climates and applications to climatic interpretations of tropical isotopic records, J. Geophys. Res.-Atmos., 115, D12 118, doi:10.1029/2009JD013255, 2010.
 - Risi, C., Noone, D., Worden, J., Frankenberg, C., Stiller, G., Kiefer, M., Funke, B., Walker, K., Bernath, P., Schneider, M., Bony, S., Lee, J., Brown, D., and Sturm, C.: Process-evaluation of tropospheric humidity
- 690 simulated by general circulation models using water vapor isotopic observations: 2. Using isotopic diagnostics to understand the mid and upper tropospheric moist bias in the tropics and subtropics, J. Geophys. Res.-Atmos., 117, D05304, doi:10.1029/2011JD016623, 2012a.

Risi, C., Noone, D., Worden, J., Frankenberg, C., Stiller, G., Kiefer, M., Funke, B., Walker, K., Bernath, P.,





Schneider, M., Wunch, D., Sherlock, V., Deutscher, N., Griffith, D., Wennberg, P. O., Strong, K., Smale, D.,
Mahieu, E., Barthlott, S., Hase, F., García, O., Notholt, J., Warneke, T., Toon, G., Sayres, D., Bony, S., Lee,
J., Brown, D., Uemura, R., and Sturm, C.: Process-evaluation of tropospheric humidity simulated by general
circulation models using water vapor isotopologues: 1. Comparison between models and observations, J.
Geophys. Res.-Atmos., 117, D05303, doi:10.1029/2011JD016621, 2012b.

Rothman, L. S., Gordon, I. E., Barbe, A., Benner, D. C., Bernath, P. F., Birk, M., Boudon, V., Brown, L. R.,
Campargue, A., Champion, J.-P., Chance, K., Coudert, L. H., Dana, V., Devi, V. M., Fally, S., Flaud, J.M., Gamache, R. R., Goldman, A., Jacquemart, D., Kleiner, I., Lacome, N., Lafferty, W. J., Mandin, J.-Y.,
Massie, S. T., Mikhailenko, S. N., Miller, C. E., Moazzen-Ahmadi, N., Naumenko, O. V., Nikitin, A. V.,
Orphal, J., Perevalov, V. I., Perrin, A., Predoi-Cross, A., Rinsland, C. P., Rotger, M., Šimečková, M., Smith,
M. A. H., Sung, K., Tashkun, S. A., Tennyson, J., Toth, R. A., Vandaele, A. C., and Vander Auwera, J.:

705 The HITRAN 2008 molecular spectroscopic database, J. Quant. Spectrosc. Radiat. Transfer, 110, 533–572, doi:10.1016/j.jqsrt.2009.02.013, 2009.

Scheepmaker, R. A., Frankenberg, C., Galli, A., Butz, A., Schrijver, H., Deutscher, N. M., Wunch, D., Warneke, T., Fally, S., and Aben, I.: Improved water vapour spectroscopy in the 4174-4300 cm⁻¹ region and its impact on SCIAMACHY HDO/H₂O measurements, Atmos. Meas. Tech., 6, 879–894, doi:10.5194/amt-6-879-2013, 2013.

- 2013.
 Scheepmaker, R. A., Frankenberg, C., Deutscher, N. M., Schneider, M., Barthlott, S., Blumenstock, T., Garcia, O. E., Hase, F., Jones, N., Mahieu, E., Notholt, J., Velazco, V., Landgraf, J., and Aben, I.: Validation of SCIAMACHY HDO/H2O measurements using the TCCON and NDACC-MUSICA networks, Atmos. Meas. Tech., 8, 1799–1818, doi:10.5194/amt-8-1799-2015, 2015.
- 715 Schepers, D., aan de Brugh, J. M. J., Hahne, P., Butz, A., Hasekamp, O., and Landgraf, J.: {LINTRAN} v2.0: A linearised vector radiative transfer model for efficient simulation of satellite-born nadir-viewing reflection measurements of cloudy atmospheres, J. Quant. Spectrosc. Radiat. Transfer, 149, 347–359, doi:10.1016/j.jgsrt.2014.08.019, 2014.
- Schneider, M., Toon, G. C., Blavier, J.-F., Hase, F., Leblanc, T., and Gutman, S.: H₂O and δD profiles
 remotely-sensed from ground in different spectral infrared regions, Atmos. Meas. Tech., 3, 1599–1613, doi:10.5194/amt-3-1599-2010, 2010.
 - Schneider, M., Wiegele, A., Barthlott, S., González, Y., Christner, E., Dyroff, C., García, O. E., Hase, F., Blumenstock, T., Sepúlveda, E., Mengistu Tsidu, G., Takele Kenea, S., Rodríguez, S., and Andrey, J.: A framework for accurate, long-term, global and high resolution observations of tropospheric H2O-δD pairs –

725 a MUSICA review, Atmos. Meas. Tech. Disc., 2016, 1–47, doi:10.5194/amt-2015-330, 2016.

- Schneising, O., Buchwitz, M., Burrows, J. P., Bovensmann, H., Bergamaschi, P., and Peters, W.: Three years of greenhouse gas column-averaged dry air mole fractions retrieved from satellite - Part 2: Methane, Atmos. Chem. Phys., 9, 443–465, 2009.
- Schrijver, H., Gloudemans, A. M. S., Frankenberg, C., and Aben, I.: Water vapour total columns from SCIAMACHY spectra in the 2.36 μm window, Atmos. Meas. Tech., 2, 561–571, 2009.
 - Soden, B. J., Jackson, D. L., Ramaswamy, V., Schwarzkopf, M. D., and Huang, X.: The Radiative Signature of Upper Tropospheric Moistening, Science, 310, 841, doi:10.1126/science.1115602, 2005.

Strahler, A. P., Lucht, W., Schaaf, C. B., Tsang, T., Gao, F., Li, X., Muller, J.-P., Lewis, P., and Barnsley,





 J.: MODIS BRDF Albedo Product: Algorithm Theoretical Basis Document Version 5.0, Report, NASA
 MODIS Adaptive Processing System. Goddard Space Flight Center, USA, http://modis.gsfc.nasa.gov/data/ atbd/atbd_mod09.pdf, 1999.

Tol, P., Landgraf, J., and Aben, I.: Instrument noise model for the Sentinel 5 SWIR bands, Tech. rep., SRON Netherlands Institute for Space Research, Utrecht, The Netherlands, 2011.

- van Deelen, R., Hasekamp, O. P., and Landgraf, J.: Accurate modeling of spectral fine-structure in Earth
 radiance spectra measured with the Global Ozone Monitoring Experiment, Appl. Opt., 46, 243–252, doi:10.1364/AO.46.000243, 2007.
 - Veefkind, J. P., Aben, I., McMullan, K., Förster, H., de Vries, J., Otter, G., Claas, J., Eskes, H. J., de Haan, J. F., Kleipool, Q., van Weele, M., Hasekamp, O., Hoogeveen, R., Landgraf, J., Snel, R., Tol, P., Ingmann, P., P., V., Kruizinga, P. B., Vink, R., Visser, H., and Levelt, P. F.: TROPOMI on the ESA Sentinel-5 Precursor:
- 745 A GMES mission for global observations of the atmospheric composition for climate, air quality and ozone layer applications, Remote Sens. Environ., 120, 70–83, doi:doi:10.1016/j.rse.2011.09.027, 2012.
 - Wassmann, A., Borsdorff, T., aan de Brugh, J. M. J., Hasekamp, O. P., Aben, I., and Landgraf, J.: The direct fitting approach for total ozone column retrievals: a sensitivity study on GOME-2/MetOp-A measurements, Atmos. Meas. Tech., 8, 4429–4451, doi:10.5194/amt-8-4429-2015, 2015.
- 750 Worden, J., Noone, D., Bowman, K., Beer, R., Eldering, A., Fisher, B., Gunson, M., Goldman, A., Herman, R., Kulawik, S. S., Lampel, M., Osterman, G., Rinsland, C., Rodgers, C., Sander, S., Shephard, M., Webster, C. R., and Worden, H.: Importance of rain evaporation and continental convection in the tropical water cycle, Nature, 445, 528–532, doi:10.1038/nature05508, 2007.

Wunch, D., Toon, G. C., Blavier, J.-F. L., Washenfelder, R. A., Notholt, J., Connor, B. J., Griffith, D. W. T.,

755 Sherlock, V., and Wennberg, P. O.: The Total Carbon Column Observing Network, Philos. T. R. Soc. A., 369, 2087–2112, doi:10.1098/rsta.2010.0240, 2011.

Yoshimura, K., Frankenberg, C., Lee, J., Kanamitsu, M., Worden, J., and Röckmann, T.: Comparison of an isotopic atmospheric general circulation model with new quasi-global satellite measurements of water vapor isotopologues, J. Geophys. Res.-Atmos., 116, D19118, doi:10.1029/2011JD016035, 2011.

760 Zakharov, V. I., Imasu, R., Gribanov, K. G., Hoffmann, G., and Jouzel, J.: Latitudinal distribution of the deuterium to hydrogen ratio in the atmospheric water vapor retrieved from IMG/ADEOS data, Geophys. Res. Lett., 31, 12 104, doi:10.1029/2004GL019433, 2004.







Fig. 3. Two-band CH₄ filter results for clouds (top), cirrus (middle) and aerosol (bottom). Plotted is the relative difference in total column CH₄ retrieved from the weak and strong bands: CH₄ (weak - strong) / strong (%). The cloud scenario assumed a cloud optical thickness of $\tau_{cld} = 5$ and a variable cloud-top-height (x-axis) and cloud fraction (y-axis). The cirrus scenario assumed a cloud fraction of 100% for a layer between 9 and 10 km and a variable surface albedo (x-axis) and cirrus optical thickness (y-axis). The aerosol scenario assumed a sulphate-type aerosol in the boundary layer between 0–2 km and a variable surface albedo (x-axis) and aerosol optical thickness (y-axis).







Fig. 4. Atmospheric profiles for the number densities of the absorbers (bottom axis, normalised to the surface value) and temperature (top axis) used as input for the model atmosphere.







Fig. 5. Cloud free rerieval bias as a function of surface albedo and SZA for the total columns of H_2O (top), HDO (middle) and the HDO/ H_2O ratio (bottom).







Fig. 6. Cloud free statistical error estimates (single measurement noise) as a function of surface albedo and SZA for the total columns of H_2O (top), HDO (middle) and the HDO/ H_2O ratio (bottom).







Fig. 7. Rerieval bias for an aerosol layer between 0-2 km as a function of surface albedo and AOT for the total columns of $H_2O(\%)$ (top), HDO (%) (middle) and the HDO/ H_2O ratio (%) (bottom). Applying the two-band methane cloud filter with a threshold of 6% to this scenario would result in filtering of the scenes with surface albedo < 0.1.







Fig. 8. A selection of the input for the realistic scenario simulation. Top: SWIR surface albedo. Second: Aerosol optical thickness at 550 nm. Third: Cloud optical thickness. Bottom: Cloud top height.







Fig. 9. Input total columns for the most important absorbing species for the realistic scenario simulation. Top: H_2O . Second: HDO. Third: CH_4 . Bottom: the resulting total column HDO/ H_2O ratio expressed in δD .







Fig. 10. Retrieval biases for water in the realistic scenario simulation. Except for the bottom left panel, the results are cloud filtered using a weak-vs-strong water band threshold of 8%. Top: $\mathrm{H_2O}$ bias. Second: HDO bias. Third: relative difference in the weak-vs-strong water bands used for cloud filtering. Bottom: bias in the derived HDO/H₂O ratio. 34







Fig. 11. Retrieved, cloud-filtered, single measurement precision of H_2O (top), HDO (second), CH_4 (third) and of the derived HDO/ H_2O ratio (bottom). 35