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No detection of methane on Mars based on early ExoMars Trace Gas Orbiter observations

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The detection of methane on Mars has been interpreted as indicating that geochemical or biotic activities could persist on Mars today¹. A number of different measurements of methane show evidence for transient, locally elevated methane concentrations and seasonal variations in background methane concentrations²⁻⁵. These measurements, however, are difficult to reconcile with current understanding of the chemistry and physics of the Martian atmosphere^{6, 7}, which predicts the methane life time of several centuries resulting in its even, well mixed distribution^{1, 6, 8}. Here we report highly sensitive attempts to detect atmospheric methane by the ACS and NOMAD instruments onboard the ESA-Roscosmos ExoMars Trace Gas Orbiter (TGO) from April to August 2018. We do not detect methane during the time period of our measurements and over a range of latitudes in both hemispheres. Our upper limit for methane of ~0.05 ppbv is 10-100 times lower than previously reported positive detections^{2, 4}. We suggest that reconciliation between the present findings and the background methane concentrations found in Gale crater⁴ would require an unknown process that can rapidly remove or sequester methane from the lower atmosphere before it spreads globally.

The first positive detections of methane on Mars were published in 2004 from the analysis of 1999 ground-based spectroscopic observations⁹, and from the Planetary Fourier Spectrometer (PFS) instrument on board ESA's Mars Express orbiter¹⁰. Mixing ratios of methane of ~10 ppbv were reported. This stirred up excitement in the scientific community but both observations were at the limit of sensitivity. Another set of ground-based echelle-spectroscopy observations reported a plume of methane developed over 60 northern summer sols² in 2003, reaching a peak value of 45±10 ppbv. No or little methane (≤7–8 ppbv) was detected before and after this event^{2, 11}. One more ground-based detection of 10 ppbv was reported in 2005¹². Starting from October 2012 the Tunable Laser Spectrometer (TLS) of the Sample Analysis at Mars (SAM) instrument onboard NASA's Curiosity rover (MSL, Mars Science Laboratory) performed local samplings of Mars's atmosphere in Gale crater. The readings first remained below 2-3 ppbv, yet were followed by a number of positive detections in 2013-

2017, the most notable of 9 ppbv in January 2014³. This result was questioned in ref.¹³ on the basis of potential rover self-contamination, an argument later rejected by the TLS team who showed such hypothesis was excluded based on a number of supporting evidences⁴. One of TLS detections of 5.8 ppbv in 2013⁴ has been independently confirmed by PFS target observation of the Gale surroundings resulting in 15.5 ppbv value⁵. More sensitive TLS samplings led to the discovery of a seasonally varying “background level” ranging between 0.24 and 0.65 ppbv⁴.

In the oxidizing martian atmosphere methane is slowly destroyed by UV photolysis and reactions with OH and O(¹D). Based on our current understanding of Mars photochemistry, it should have a lifetime of 250-300 years^{1, 6, 8}. Therefore, its detection, even in small quantities, requires a sustained replenishment. Methane on Mars has attracted much interest because on Earth, most of the atmospheric methane has a biological origin. Thus the Martian atmospheric methane might hint at active or extant microbial life or at the existence of organic matter. However, methane can also be formed abiotically, by low-temperature chemical reactions (e.g., CO₂ hydrogenation) or magmatic processes^{14, 15}.

Given its potential implications for exobiology or geochemistry, highly sensitive measurements of atmospheric methane and other trace species were identified as the primary science goal of the TGO mission¹⁶. The 2-hour circular orbit of the TGO satellite was designed for detecting trace gases using solar occultations, a technique in which the spacecraft instruments observe the atmospheric absorption spectrum of sunlight during sunsets and sunrises¹⁷. Solar occultations provide very high sensitivity for trace gas concentration measurements because: (1) the Sun’s brightness results in very high signal to noise ratio (SNR) spectra; and (2) the atmospheric optical path length in occultation viewing geometry is up to 10x longer than that achieved when observing the planet’s surface. Two instrument suites on board TGO were designed to perform such measurements: ACS (the Atmospheric Chemistry Suite)¹⁸ and NOMAD (Nadir and Occultation for MArS Discovery)¹⁹. Both ACS and NOMAD cover the 3.3 μm spectral range that includes the strongest fundamental absorption bands for hydrocarbons such as CH₄, in particular the ν₃ asymmetric stretching band on which all the

previous detections were made. TGO started its science operations in April 2018, with the first
80 occultation taking place on April 21st. From June until August 2018 a planetary encircling dust storm
has reduced the transparency of the atmosphere (see companion paper²⁰), while high northern polar
latitudes remained suitable for sensitive soundings. A map of measurements by the most sensitive
channels of ACS and NOMAD in the CH₄ range is shown in Figure 1.

[Figure 1]

85 When staring at the solar disk outside the atmosphere, the SNR for ACS MIR (mid-IR) channel
reaches 10,000 (one detector line, 2 s integration time, 2.5 km vertical sampling rate), and for NOMAD
Solar Occultation (SO) channel SNR reaches ~2000 (one spectrum, 48 ms integration time, 1 km
sampling rate). During a solar occultation, the trace gas detection sensitivity increases as the line of
sight progressively samples closer to the surface, thereby intersecting a larger air mass. At the same
90 time, sensitivity suffers from the increasing presence of dust and clouds, which can drastically reduce
the intensity of light reaching the instrument. The atmospheric aerosol loading has previously been
shown to have a negative effect on the retrieval accuracy at lower altitudes¹⁸. The optimum sensitivity
is thus achieved at the lowest altitude where the atmosphere is still transparent enough, typically
between 5 to 25 km corresponding to an atmospheric transmission of 0.2 to 0.5. Figure 2 shows
95 examples of spectra acquired at an altitude close to the optimal one. No methane absorptions are
apparent, while we accurately measure the faint H₂O lines within the range, which, at very low water
content, have an absorption depth comparable to a 1 ppbv CH₄ absorption (panel c). Corresponding
profiles of H₂O (see Methods), are characterized by an unprecedented accuracy compared to previous
profiling^{20, 21}.

100 [Figure 2]

Based on the noise level, CH₄ absorption integrated over the line of sight can be tentatively
fitted along with absorption of CO₂ while taking into account the instrument spectral resolution (see
Methods). This way, an estimation of methane detection limits, converted into volume mixing ratios,
for the full data set acquired by ACS and NOMAD was made. Figure 3 illustrates the detection limits

105 for the ensemble of observations performed by both instruments. The gradual increase in upper limits
observed after the onset of the Planetary dust event (indicated by a light grey bar) is a direct
consequence of dust forcing detections to occur progressively above 30 km, that is above the
theoretically optimal altitude (detection-wise) usually found between 15 to 25 km. A few profiles,
measured in cleaner northern conditions, were able to achieve the most precise detection limits of
110 0.012 ppbv down to an altitude of ~3 km (cf. Figure 3).

[Figure 3]

This non-detection of methane by TGO and its associated upper limits are in contradiction
with the 0.41 ppbv background levels measured *in situ* by Curiosity at the same season⁴ in previous
years. As discussed above, TGO is able to detect concentrations at least ten times lower than 0.4 ppbv.
115 In fact, a simple comparison of the theoretical sensitivity of the solar occultation method with the TLS
instrument method shows that TGO should be more sensitive than what can be achieved with the TLS,
even when the measurements are performed using the TLS enrichment mode (see Methods).

Is it possible that the factor of ten difference between the MSL measurements and the TGO
upper limits could result from spatial variations in the methane mixing ratios? MSL measurements
120 were obtained at the bottom of Gale Crater near the equator, while the best TGO measurements were
achieved in the near-polar latitudes and a few kilometres above the surface. However, It is difficult to
understand why the Martian atmosphere would permit such a spatial differentiation of
concentrations. On Mars, the daytime atmospheric boundary layer is characterized by intense
convective motions, which mix any trace gas such as methane efficiently on a daily basis from the
125 surface up to the top of the convective boundary layer, usually 6 to 10 kilometres high. From there
the global wind circulation transports trace gases horizontally^{6, 22} and vertically around the planet.
Global uniform mixing of methane occurs on a scale of 2 to 3 months^{6, 8, 23}. Even in the extremely
unlikely case where Gale Crater would constitute the sole source of methane on Mars (note that Gale
Crater and surrounding areas along the Martian dichotomy host geological features where methane
130 could be released¹³), MSL measurements still remain in disagreement with the detection limits derived

from TGO measurements. Indeed, considering only the background concentration of 0.41 ppbv we assume that Gale is uniformly and constantly filled with CH₄ up to its lowest rim (at ~2 km) and that a mixing timescale of 1 sol⁻¹ (ref.⁴) is the typical time for air to leave the crater. The Gale emission would lead CH₄ to accumulate globally over one Martian year at a level of ~2 pptv. This implies that such a background emission from Gale crater could only have been going on for at most 24 Martian years (or 135 44 Earth years) before the detection limits reported here would have been reached. Taking into account the ppbv spikes of CH₄ concentration reported by MSL, this 24 years timeframe would be even more shortened. This is unrealistic. To maintain a level of methane ten times higher than elsewhere, Gale Crater should not only be the unique source, it should also preserve its air mass from exchanging 140 with the global atmosphere. Interestingly, mesoscale model simulations have shown that the depth of the boundary layer in Gale crater is significantly lowered²⁴ due to the crater size and depth. Even if this would tend to maintain methane locally, the same simulation^{24, 25} shows that the slope winds on the side of the crater and the induced updraft above the rims is so intense that methane should be efficiently injected into the atmosphere at 10 km altitude. In no case can we consider Gale to be an 145 isolated crater (see Methods for additional information).

To reconcile the lack of CH₄ detection in the TGO data and the positive CH₄ detection at the surface by Curiosity, one must invoke a mechanism able to fully eradicate methane in the lower atmosphere at a rate ~1000 times faster than that predicted by the conventional chemistry. However, existing conventional models not only describe very well the chemistry of methane on Earth but also 150 reproduce satisfactorily on Mars species that are sensitive to the oxidizing capacity of the atmosphere, such as hydrogen peroxide²⁶, ozone²⁷, and carbon monoxide²⁸. Unless a mechanism is discovered that can rapidly destroy or sequester methane and is compatible with our wide quantitative understanding of Mars photochemistry, all the methane detections reported to date appear inconsistent with present TGO measurements.

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Figure captions

Figure 1. Map of ExoMars TGO methane measurements by ACS (stars) and NOMAD (circles) used in this study. The symbols, corresponding to example measurements considered in this study are enlarged. The colour scale denotes L_s (the areocentric Solar Longitude). Gale crater (the Curiosity rover location) is marked by a bold, black square.

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Figure 2. TGO spectra encompassing the spectral range with multiple methane R-branch features.

Panels **a, b**: Example of spectra obtained by NOMAD SO in two different ranges. The measurements are plotted together with synthetic models of CH₄ and of water vapour absorption. Panel **c**: similar results obtained by ACS MIR before the dust storm. The ACS spectrum includes the same methane feature of methane as in Panel **b** (3048.2 cm⁻¹), and two stronger isolated features, allowing to constrain the methane content below few tens of pptv. Measured spectra show 1-σ instrument noise. The tangent altitudes above areoid are indicated.

Figure 3. Upper limits for CH₄ (95% confidence limit) obtained by TGO (ACS and NOMAD) compared to seasonally variable background methane as measured by SAM-TLS on Curiosity. The colour scale gives the latitude of the TGO sampling. The TGO dataset has been filtered to retain only the most precise upper limits below a threshold of 0.15 ppbv, encompassing values down to 0.012 ppbv. The gradual increase in upper limits is associated with the onset of the Planetary dust event (indicated by a light grey bar).

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Author contributions

O.K., A.C.V., F.M. conceived the study, collected inputs and wrote the paper. A.A.F. calibrated the ACS data and analysed the profiles assisted with A.T., K.S.O., J.A. The ACS dataset was prepared by A.T. assisted by L.B., J.A.P., and Y.S.I. A.T, A.G., N.I., A.S., and A.P. have designed ACS observations. D.B. analysed ACS CO₂ data. F.M. has derived the methane detection limits from ACS. I.R.T. analysed the solar occultation data and provided transmittances from NOMAD SO. J.T.E. and L.T. with S.A. derived the NOMAD methane detection limits. J.T.E. and S.R. provided and analysed the a priori knowledge and initial GCM fields, the latter were provided by L.N. and F.D. C.D. and Y.W. were involved in UVIS calibration and data pipeline. B.R., B.B., C.Q., and E.N. designed the NOMAD observations helped by J.M. for the UVIS channel. L.H., S.B., and R.C. are responsible for the uplink and downlink of telemetry and science data, and first conversion of those to scientific data. M.R.P., G.B. and J.J.L.M. provided support in the selection of the NOMAD observations based on scientific interest. F.F., F.L., F.D., provided critical inputs regarding the model chemistry and circulation, assisted with J.L.B., L.N., S.V., and G.E. C.W., L.Z. and H.S. coordinated observations of the various instruments on TGO. All authors contributed to the preparation of the manuscript.

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Methods

ACS instrument and measurements

ACS¹⁸ consists of three infrared channels featuring high accuracy, a high resolving power, and
305 broad spectral coverage (0.7 to 17 μm). The mid-infrared (MIR) channel is a high dispersion echelle
spectrometer dedicated to solar occultation measurements in the 2.3-4.5 μm range. MIR has been
conceived to accomplish the most sensitive measurements of Martian trace gases, while
simultaneously profiling more abundant compounds such as CO₂, H₂O, and their isotopologues. ACS
MIR is a crossed dispersion spectrometer which measures spectra dispersed onto a cryogenic 512×640
310 CdHgTe infrared array. For each acquired frame, MIR measures ≥ 20 adjacent diffraction orders,
covering an instantaneous spectral range of 0.15 to 0.3 μm wide. To achieve the full spectral coverage,
a secondary dispersion grating can be rotated to one out of 12 distinct positions (see¹⁷ for a tabulated
description of all the grating positions). Together with two other channels, the near-infrared (NIR) and
the thermal infrared Fourier transform (TIRVIM) spectrometers, ACS continuously covers the spectral
315 range between 0.7 and 17 μm . NIR and TIRVIM channels are used to observe, both in solar occultation
and in nadir, water vapour H₂O, carbon monoxide CO, and other gases including molecular oxygen O₂
in a fundamental state. The broad spectral range acquired enables the characterization of the key
meteorological parameters, including dust and water ice cloud column opacities. In addition, the
temperature profile of the atmosphere can be retrieved from the 15- μm CO₂ band sensed by TIRVIM
320 in nadir.

In this work, we used ACS MIR data obtained with the secondary grating tuned to position
#12. In this range, MIR acquires frames containing 20 adjacent and partially overlapping diffraction
orders (from #172 to #192) from 3.09 μm to 3.45 μm (Figure 4). The instrument point-spread-function
(PSF) can be assimilated to a Gaussian function associated with a spectral resolving power of
325 $\lambda/\Delta\lambda > 30,000$ where $\Delta\lambda$ is taken as the full-width at half-maximum (FWHM) of the Gaussian, that is
 $\sim 2.36 \times \sigma$ (with σ , the standard deviation). ACS MIR was operated in the so-called “high-sensitivity”
mode, where 200 frames obtained from consecutive 6 ms integration frames are stacked together on-
board. One full measurement lasts 2.1 seconds. Depending on orbital parameters a profile of the
atmosphere from 0 to 200 km is measured within 3 to 6 minutes. The uppermost part of the

330 occultation corresponds to the clear sun observations, averaged to obtain a reference spectrum I_{sun} .
A dark signal I_{dark} (the sum of detector dark current and of the surrounding thermal background
emission) is estimated from the dark part of the occultation where the sun is fully obscured by the
solid body of Mars and is refined using dedicated observations of the open space. As the thermal
environment inside the instrument is slightly changing during the occultation session due to internal
335 and solar heating, I_{sun} and I_{dark} are also time-dependent. To account for the gradual sub-pixel drift of
the image during the occultation, we extrapolate the trend of each pixel measured during the clear
sun observation throughout the occultation. The methane spectral range around 3030 cm^{-1} is free of
any strong gaseous absorptions, and the atmosphere above 100 km can be considered clear from any
absorptions features, increasing the accuracy of the extrapolation. Each line of the detector within
340 each stripe represents a transmittance spectrum at a certain altitude and at a wavelength range
corresponding to the displayed diffraction. The resulting frame of transmittances undergoes a
projective transformation to correct the artefactual curved appearance of the stripes of each
diffraction order and to make it look more horizontal and thereby maintain spectral connectivity
between adjacent orders (see Extended Data Figure 1). A first-order approximation of the pixel-to-
345 wavelength calibration is established using solar lines and is then refined using the strong absorption
lines of CO_2 or H_2O .

NOMAD instrument and measurements

NOMAD¹⁹ includes three spectroscopic channels, operating from the ultraviolet (UV) and
visible range to $4.3\text{ }\mu\text{m}$. The channel most sensitive to trace gases is the SO (Solar Occultation)
350 spectrometer providing a spectral resolving power of $\lambda/\Delta\lambda\approx 20,000$ in the spectral range of $2.3\text{-}4.3\text{ }\mu\text{m}$.
Within this range, NOMAD SO acquires 10 separate wavelength sub-ranges to profile a variety of
atmospheric species. The two other channels of NOMAD are the UVIS (Ultraviolet and visible
spectrometer; $200\text{-}650\text{ nm}$), and LNO (Limb, Nadir, Occultation) spectrometers that can be operated
both in solar occultation and in nadir. NOMAD provides vertical profiling information for atmospheric
355 constituents at unprecedented spatial and temporal resolution. Indeed, in solar occultation, the

vertical resolution is less than 1 km for SO and UVIS, with a sampling rate of 1 s (one measurement every 1 km), and occultations range from the surface to 200 km altitude. NOMAD also provides mapping of several constituents (aerosols/dust/clouds, and O₃, H₂O, HDO, CO, and other trace gases) in nadir mode with an instantaneous footprint of 0.5 x 17 km² (LNO spectrometer) and 5 km² (UVIS spectrometer) respectively, with a repetition rate of 30 Martian days.

For this work, we analysed SO channel data measured between 21 April and 1 August. SO measures 4 spectral bins in each of 5 or 6 diffraction orders per second in solar occultation mode, among which a series of specific diffraction orders were chosen (order 133 to 136 spanning 2990 to 3080 cm⁻¹ spectral range) where methane features are located. To increase the sensitivity of the NOMAD SO measurements, we accumulate all the spectral measurements in each occultation from the 4 detector bins into 3 km vertical bins. The transmittance calibration and error calculation as described by Trompet et al.²⁹ is adapted to consider this accumulation. By accumulating multiple measurements we improve the SNR, by increasing the 48 ms integration time for a single detector bin to an effective integration time of 144 ms for all illuminated detector bins. By accumulating multiple measurements, we effectively increase the typical 48 ms integration time of a single measurement to an average of 500 ms integration, thereby increasing the SNR as compared to the previous estimation³⁰.

Calculation of the methane detection limits

Attempting to detect methane we have used the ACS data from diffraction orders #180 (which contains the Q-branch of the fundamental ν_3 asymmetric stretching band of CH₄), and #182 (which contains the strongest lines among the P- and R-branches) as shown in Extended Data Figure 1, and described in the main text. Five detector lines were stacked together to form a spectrum to be fitted. We then applied a method to retrieve vertical profiles of trace gas vmr as developed for Mars Express solar occultation^{21, 31}. First, a more accurate wavelength-to-pixel dependence is established using the most intense H₂O absorption lines and is then propagated elsewhere in the occultation in particular where the water lines are too faint. Finally, the vmr profiles of the trace constituents are retrieved by

fitting the retrieved occultation portion with a pre-computed look-up synthetic model in a three-parameter space (H_2O , CH_4 vmr, and aerosol extinction) using Rodgers' regression with additional Tikhonov regularization³². The synthetic spectra were computed with spectral line parameters from HITRAN 2016³³ corrected to account for the CO_2 atmosphere, as described in²¹. Temperature and pressure profiles were extracted from the Mars Climate Database³⁴. Aerosol extinction is assumed to have a grey behaviour within a diffraction order. The retrieved profiles of H_2O and the attempts to retrieve CH_4 for two selected occultations are shown in Extended Data Figure 2. The H_2O profiles, even though obtained from faint lines corresponding to very dry conditions, are nevertheless characterized by a greater accuracy compared to previous H_2O profilings²¹. However, this formal retrieval shows no trace of methane. Sometimes very small methane abundances of ~ 20 pptv in the R-branch (3058 cm^{-1}) were retrieved above the $1\text{-}\sigma$ level. However, this result was not confirmed, in any case, by a corresponding result in the Q-branch (3018 cm^{-1}). We conclude that these constitute false positive detections, perhaps due to a residual fixed pattern noise in the spectrum. The effect of the atmospheric aerosol loading on the retrieval accuracy is illustrated in the right panel where the "profile" of transmittance noise is shown along with the optical slant density, as measured along the line of sight.

As no methane was detected within the analysed dataset, we selected a faster approach to estimate robust upper limits of CH_4 vmr for the considered dataset covering the April to August period of 2018. The sensitivity in solar occultation is produced by the number of species molecules observed along the LOS. We assume that no methane can be retrieved within the $1\text{-}\sigma$ error bars.

Near the centre of the detector, the best SNR are achieved at a level of $\geq 10,000$. This very low noise level implies that the main source of errors is systematic. This is the fixed pattern noise. The part of the systematic error originating from the detector dark current non-uniformity (DCNU) is not fully cancelled out when divided by solar reference I_{sun} because of the non-linearity of the detector pixels. This noise appears as a pattern with a relative level of $\leq 10^{-3}$ exhibiting a pixel-to-pixel correlation remanent on consecutive spectra. In Figure 2, the fixed pattern noise is responsible for

most of the visible irregularities, while the random noise (error bars shown by hairlines) only becomes visible near the edges of the detector where the signal is weaker.

410 To account for both the fixed pattern and the random noise, we define the instrument noise as the standard deviation of a transmission spectrum filtered from all pixel correlation wider than 2 pixels and computed over most of the diffraction order. This calculation overestimates noise in case of spectrally narrow gaseous absorptions (since the broader ones are filtered) whose signature leaks in the pixel-to-pixel variation. We inferred directly the line-of-sight (LOS) density, which is the
415 number of molecules integrated along the line of sight N [cm^{-2}] of putative methane with that of CO_2 , simultaneously measured in diffraction order #178. CH_4 LOS density is estimated separately using either the CH_4 Q-branch in order #180 ($3015\text{-}3022\text{ cm}^{-1}$) or the deepest R-branch feature that can be found in order 182 ($3057\text{-}3060\text{ cm}^{-1}$), see Figure 2.

By considering that CH_4 can be loosely detected if the tentatively retrieved value exceeds its
420 error bar, an upper limit on CH_4 is deduced separately for the Q and R branches by assuming it to be equivalent to the error bar found for the CH_4 LOS density. A multivariate regression algorithm based on a Levenberg-Marquardt approach (MPFIT.pro IDL routine of C. Markwardt based on MINPACK-1) was used for all retrieval attempts. This algorithm provides as an output the covariance matrix of the fitted parameters whose diagonal values correspond to the square of the error bar of every
425 parameter. The upper limit on CH_4 vmr is then defined as the retrieved error of the CH_4 LOS density divided by the CO_2 LOS density. The uncertainty on CO_2 is ignored, as it accounts for only a minor fraction of the upper limit. A correspondence between the instrument SNR and the CH_4 LOS density detection limit can be theoretically established. The detection limit, and the step-by-step outputs of the retrieval are shown in Extended Data Figure 3 , with the resulting upper limits vs. altitude in
430 Extended Data Figure 4.

A similar simplified retrieval method was used to estimate the detection limit from NOMAD spectra. The forward model computes the optical column density for each spectrum separately, assuming a constant mixing ratio along the line-of-sight and using the most recent HITRAN³³ CH_4 line

list with CO₂ pressure broadening coefficients. The optical depth is then convoluted to the
435 instrument's PSF with FWHM of 0.15 cm⁻¹ and then multiplied by the grating blaze and AOTF functions.
The transmittance spectra are fit by minimizing the chi-squared using the Levenberg-Marquardt
algorithm (via the Python Scipy wrapper of MINPACKS *lmdr*³⁵) to determine an optimal polynomial
background and CH₄ mixing ratio. The standard error of the mixing ratio is derived from the covariance
matrix of the optimal fit parameters. This value can be thought of as the symmetric error bound on
440 the mixing ratio that can affect the transmittance within the measurement noise and should be a close
approximation to the detection limit.

Comparing the sensitivity of TGO solar occultations to that of SAM TLS.

The SAM laser spectrometer onboard Curiosity measures gaseous absorption using an
atmospheric sample in a 16.2 meters path length cell at ambient Mars pressure (~8 mbar)³⁶. The
445 number of air (CO₂) molecules interacting with the laser beam (the column density) is $N \approx 2 \cdot 10^{20}$ cm⁻².
Very high spectral resolution of TLS allows for resolving the natural, pressure-broadened linewidth of
two methane absorption lines of $\sim 10^{-2}$ cm⁻¹. For a subset of samplings, TLS was operated in a more
sensitive mode where CO₂, which constitutes 96% of the atmosphere, was progressively removed
from the sample enriching the remaining gases by a factor of 20-25. The achieved accuracy is 1-2 ppbv
450 of methane for the direct intake, and 50-100 pptv for the enrichment mode^{3,4,36}.

For solar occultation geometry, the number of CO₂ molecules along the line of sight at a slant
altitude of 20 km is $N \approx 10^{24}$ cm⁻², increasing as sounding closer to the surface (see Figure 6). The
spectral resolution, as confirmed in flight, is ~ 0.1 cm⁻¹ for the ACS MIR channel, and ~ 0.15 cm⁻¹ for
NOMAD SO. ACS and NOMAD spectra give access to multiple strong features of methane around the
455 3.3 μm range. Neglecting other factors, such as noise, systematic errors, contamination, etc.
potentially affecting both methods, one can estimate that the TGO occultation measurements are
theoretically a factor of ≥ 1000 x more sensitive than what can be achieved with the TLS direct intakes,
and of ≥ 30 x more sensitive comparing to measurements performed in the enrichment mode.

Can we reconcile TGO and MSL measurements?

460 The mass of CH₄ in Gale crater if filled to the lowest rim can be calculated as follows: The Gale crater diameter is 154 km and its lowest rim is 2 km high, resulting in 6.7·10¹¹ kg of air or 9.3·10³⁶ molecules. The amount of CH₄ corresponding to 0.41 ppbv content is 0.41·9.3·10²⁷ molecules or 100kg of CH₄. More accurately, using MOLA topography and GCM atmospheric profiles we find 30 kg of CH₄, as Gale is not a perfect cylinder and the air density decreases with height.

465 There is no reason to consider the 10 random measurements by SAM's TLS of 0.41 ppbv on average in a two year period⁴ to coincide with transient methane releases, so we assume that amount is present at all days during the martian year. Ref.⁴ estimated the mixing rate to be 1 sol⁻¹, meaning that the methane in the crater is replenished every sol. So the total mass of CH₄ emitted in Gale crater over one martian year is 20 tons or 7.5·10²⁹ molecules. The full mass of the atmospheres is 2.6·10¹⁶
470 kg, giving 2·10⁻¹² or 2 pptv of CH₄ well mixed. In 24 martian years (or 44 Earth years) this accumulates to 50 pptv, and this just with a single source in Gale. Transient enrichments of methane, observed remotely², and spikes detected by SAM's TLS³ further accelerate the accumulation rate.

 To make the TGO and TLS-SAM results consistent (i.e. an accumulation of 50 pptv over 300 years), the mixing inside/outside Gale would have to be lower than 1 sol⁻¹ by a factor of ~7. This large
475 factor is in full contradiction with mesoscale simulations, and with the strong slope winds^{24, 25}. A newer simulation³⁷ also shows that one should not expect Gale Crater to trap any trace gas for longer than 1 sol. They were looking at water vapour, which was largely released by sublimation of overnight surface ice, and was then mixed up above 3 km (compared to MOLA, so already several km above the crater's highest rim) by noon and very well mixed by 4pm on the same day. A trace gas mixed over this height
480 range is then rapidly transported out of Gale Crater by the afternoon upslope winds, combined with the lower branch of the Hadley Circulation (northward or southward, depending on time of year). The majority of any release at 6am had left the crater by 6am on the next day.

 Finally, the dust event 2018A affecting a subset of our measurements could not substantially bias the conclusions of present study. First, we establish stringent upper limits on methane before the
485 event, and during the event in the polar areas. Second, the fact that the H₂O₂ measured in dusty

conditions is well reproduced by conventional models (despite the strong sensitivity of that species to the oxidizing capacity of the atmosphere) does not suggest the existence of any major chemical mechanism on mineral dust^{26, 38}.

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Data availability

515 The datasets generated by the NOMAD and ACS instruments and analysed during the current study
will be available in the ESA PSA repository, <https://archives.esac.esa.int/psa>, after the six-months prior
access period following ESA Rules on Information, Data and Intellectual Property. The data used for
the figures are available on request from the corresponding author O.K.

Code availability

520 The computer codes used to decipher the upper limits of CH₄ are available on request from the
corresponding author O.K.

Extended data figure captions

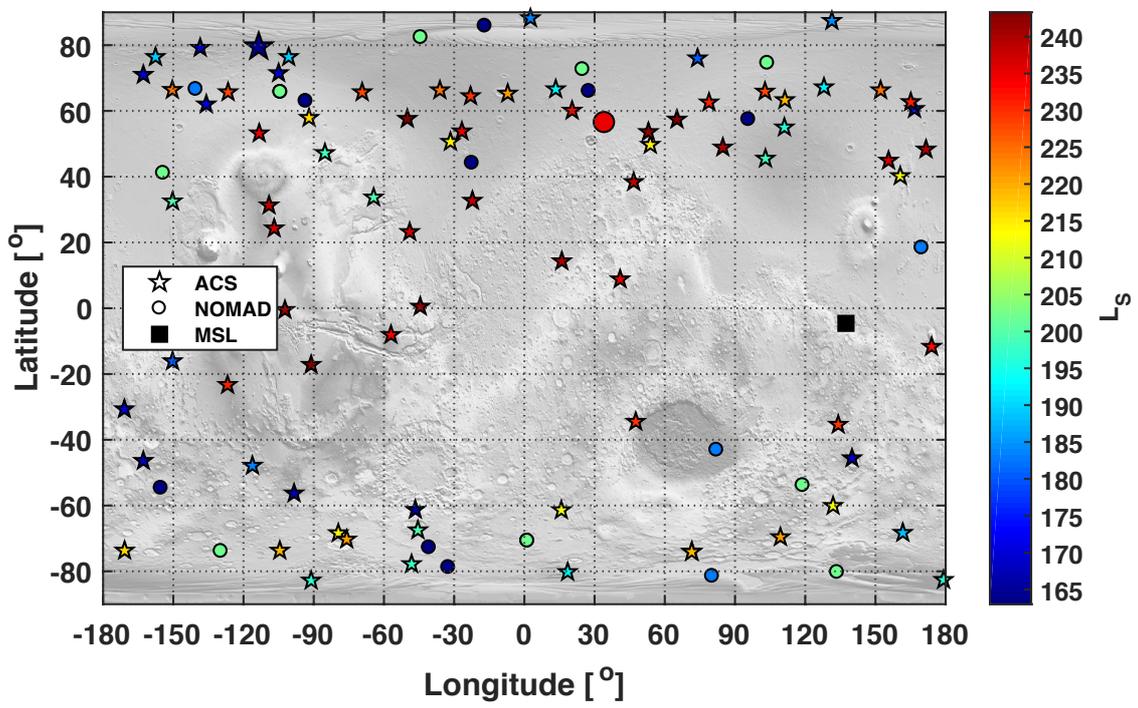
Extended Data Figure 1. A sequence of transmittance spectra measured with ACS MIR channel for
an example orbit ($L_s=180.9^\circ$) obtained using the secondary grating position #12¹⁷. Different
525 diffraction orders are denoted by changing colour of the transmission curves, and their numbers are
indicated at the upper scale. Diffraction orders used in this study are #178 (for CO₂), #180 (CH₄ QR-
branch), and #182 (CH₄ RQ-branch). Enhanced extinction on the short wavelength edge of the
spectra is due to H₂O ice absorption.

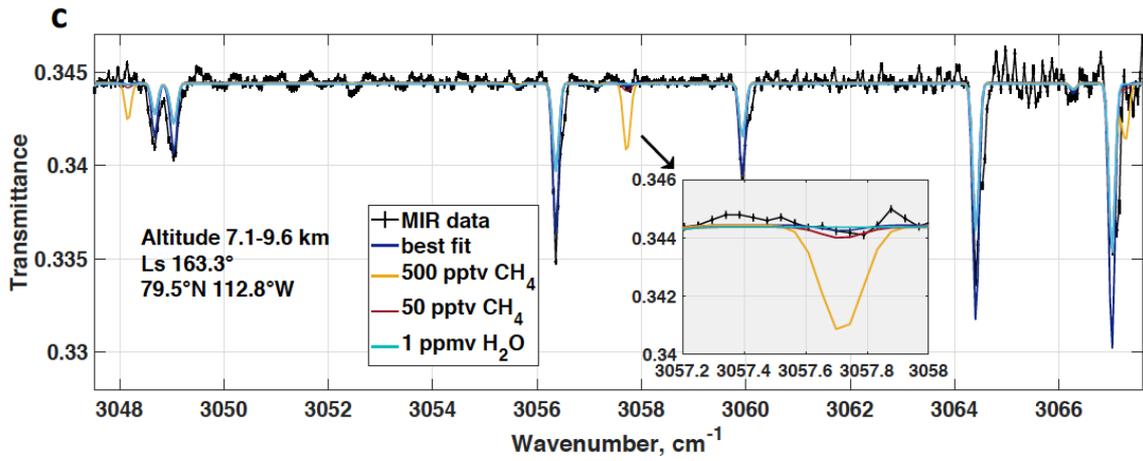
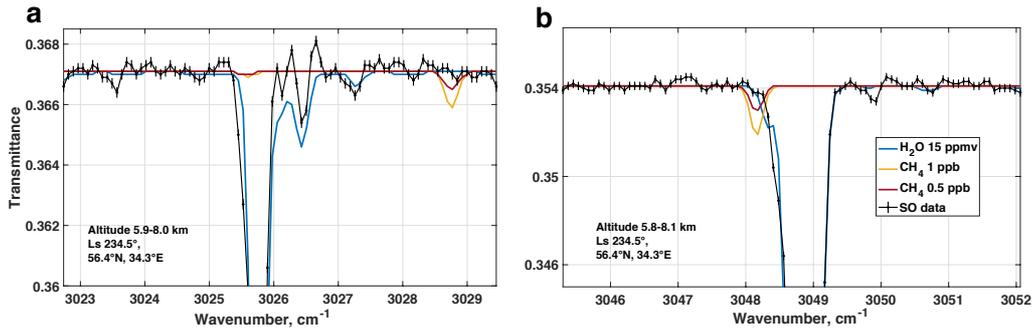
Extended Data Figure 2. Retrieval of trace gases from ACS MIR spectra using Rodger's regression
530 illustrated for one aerosol-free polar case (upper panels), and for a more cloudy low-latitude case
(lower panels). Both occultations were observed before the global dust event. Left panels: water
vapour profiles retrieved using faint H₂O absorption lines separately in diffraction orders #180 and
#182. Middle panels: attempts to retrieve CH₄ in the same diffraction orders. Curves with error bars
indicate the regularized profiles. The error bars give the 1- σ uncertainty on the retrieved

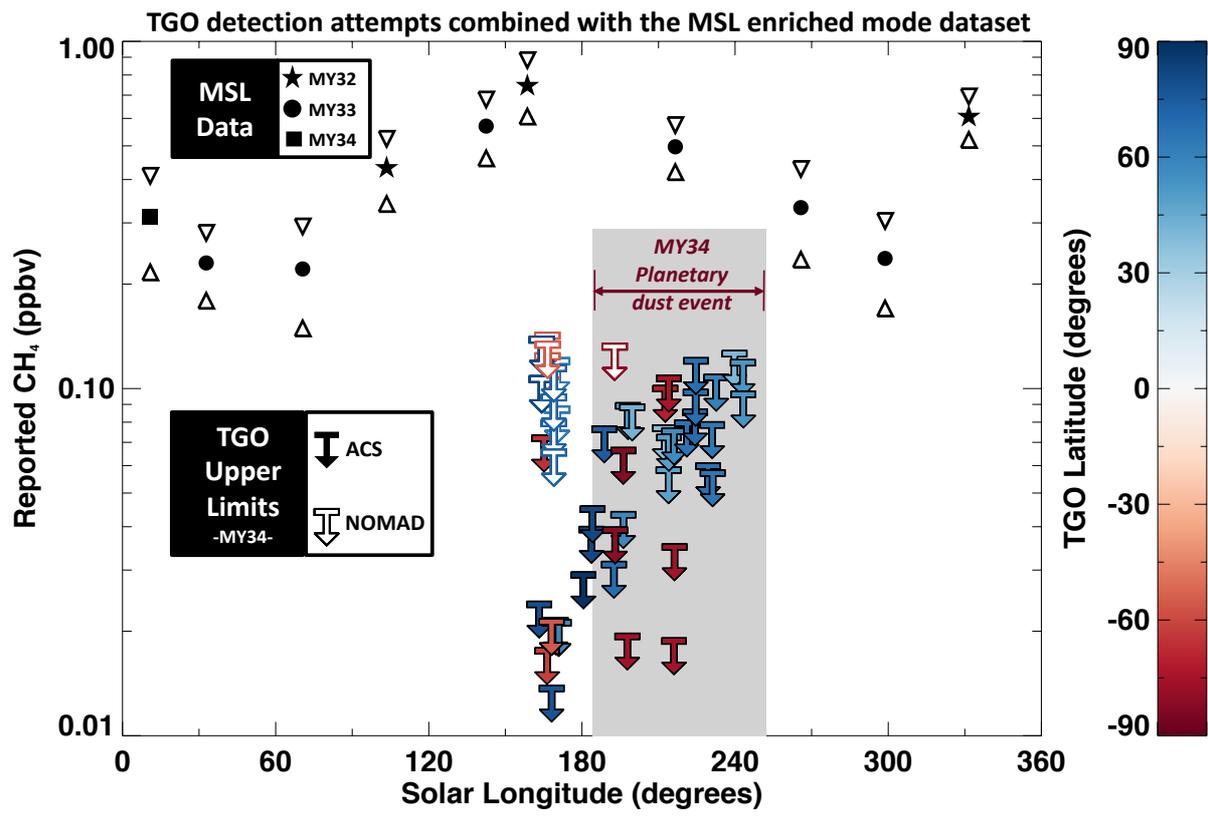
535 parameters. To illustrate the accuracy for the individual spectra, the regularization was also turned off (scatter points). Right panels: profiles of the optical depth on the line of sight, and of the signal-to-noise ratio (SNR) in the MIR spectra. The SNR for each spectrum was calculated over the whole diffraction order, excluding spectral intervals with gaseous features.

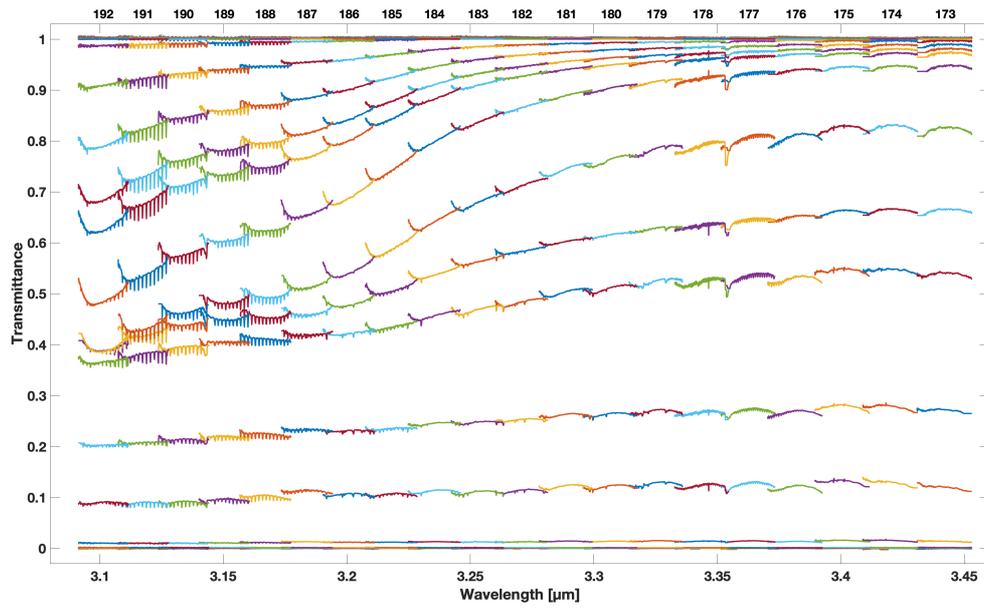
Extended Data Figure 3. Top: the theoretical relation between SNR/pixel and the retrieved 1- σ 540 uncertainty on the CH₄ line-of-sight density expressed in molecules·cm⁻². At SNR > 1,000/pixel, the associated uncertainty is 10¹⁴ molecules·cm⁻², which would yield an equivalent vmr uncertainty of 0.1 ppbv of CH₄ in the 10 km altitude range where the CO₂ density is usually around 10²⁴ molecules·cm⁻². Bottom, from left to the right: Altitude profiles of (left) SNR / pixel, (middle) CO₂ and error on CH₄ LOS density in cm⁻², and (right) resulting upper limits retrieved for CH₄ molecules. The displayed 545 observation corresponds to high northern latitudes after equinox (Ls 192°). The prevailing clear conditions allowed sounding very close to the surface with high SNR, yielding optimal conditions for the retrieval of CH₄. The black, red and blue colours refer respectively to CO₂, (in order #178), CH₄ (in order #180) and CH₄ (in order #182).

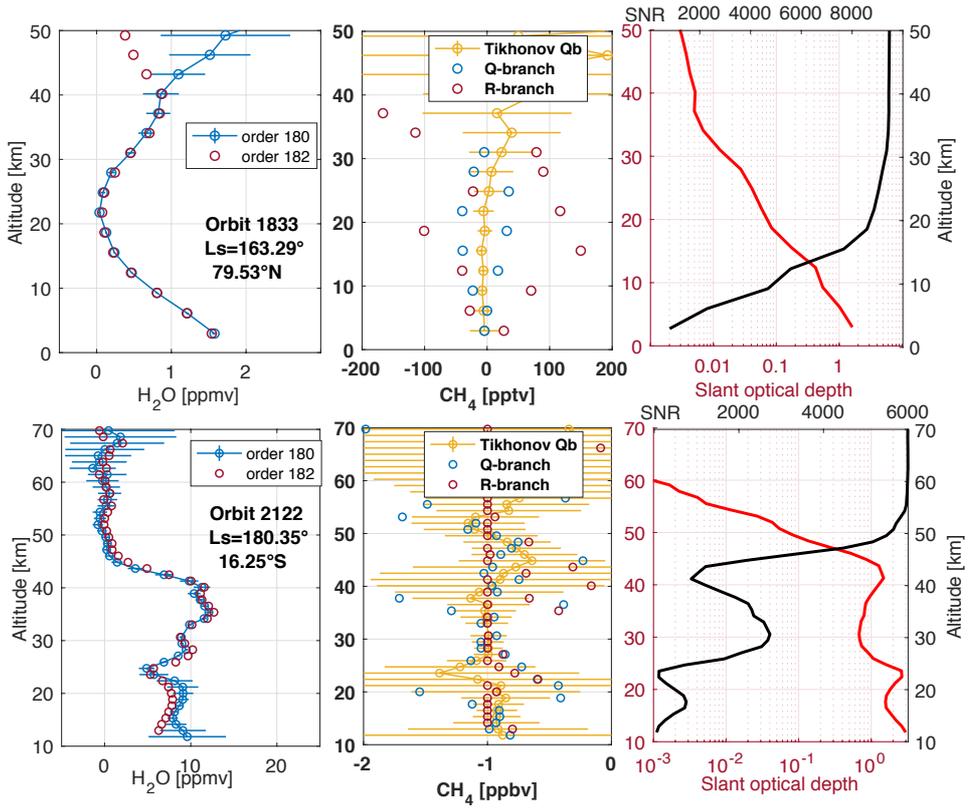
Extended Data Figure 4. A compilation of all the retrieved upper limits from the ACS-MIR dataset 550 covering the period from April 21st to September 4th 2018 using (Top) the Q-branch, and (bottom) the R-branch of CH₄ absorption. The colour scale denotes L_s. Symbols of alike colour denote individual profiles.



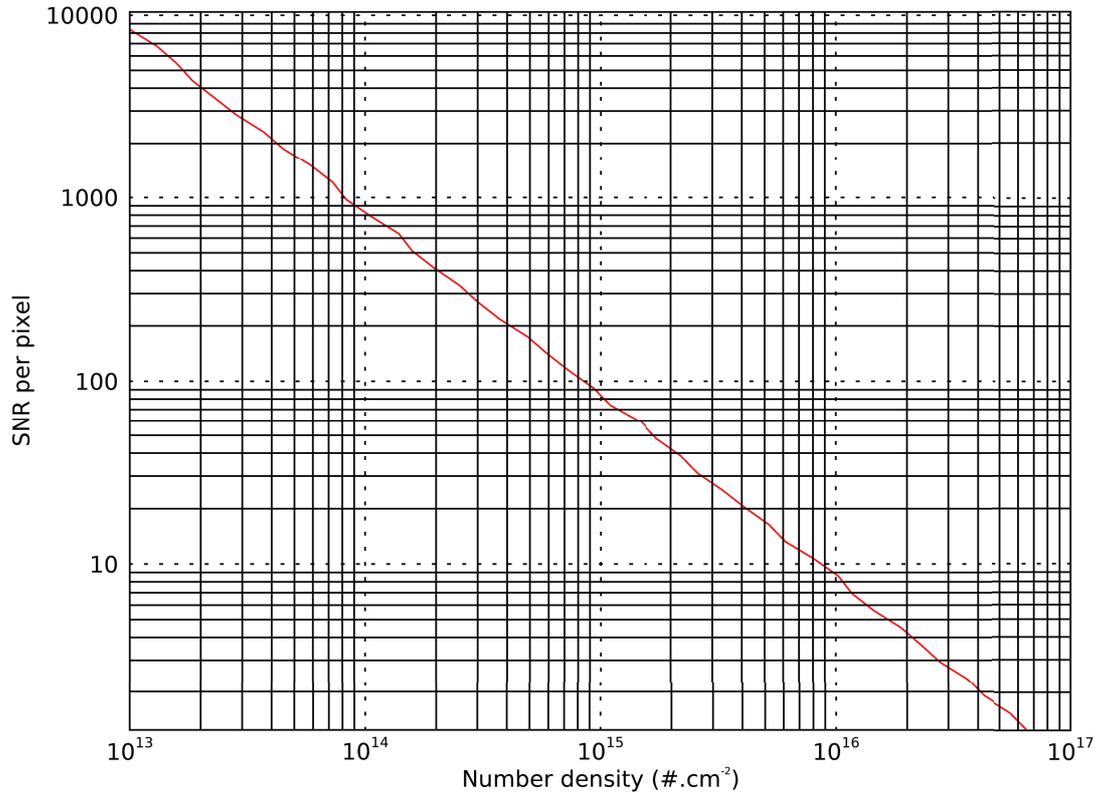








CH₄ 1-σ detection threshold



565

