Continuous Methane Emissions from the Oil and Gas Industry in the Permian Basin

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Abstract

Emissions of methane (CH₄) in the Permian basin (U.S.A.) have been derived for 2019 and 2020 from satellite observations of the Tropospheric Monitoring Instrument (TROPOMI) using the divergence method, in combination with a data driven method to estimate the background concentrations. The resulting CH₄ emission data, which have been verified using model with known emissions, have a spatial resolution of approximately 10 km. The spatial patterns of the emissions are in a good agreement with the locations of oil and gas production and drilling activities in the Permian basin, as well as with emissions of nitrogen oxides (NO_x). Analysis of time-series of locations with large CH₄ emissions indicated that there are significant continuous emissions in this region. The CH₄ emissions can be characterized as a continuous area source, rather than as dominated by a few large unplanned releases. This is important considering possible CH₄ emission mitigation strategies. In addition to providing spatially resolved emissions, the divergence method also provides the total emissions of the Permian basin and its main sub-basins. The total CH₄ emission of the Permian is estimated as 3.0 ± 0.7 Tg yr⁻¹ for 2019, which agrees with other independent estimates based on TROPOMI data. For the Delaware sub-basin, it is estimated as 1.4 ± 0.3 Tg yr⁻¹ for 2019, and for the Midland sub-basin 1.2 ± 0.3 Tg yr⁻¹. In 2020 the emissions are 8% lower compared to 2019, which could be a result of strong decreases in drilling activities due to the COVID-19 crisis.

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15 Key Points:

- Methane emissions from the Permian basin in the U.S.A. can be derived from satellite
 data with a spatial resolution of approximately 10 km.
- The derived emissions are spatially consistent with satellite derived NO_x emissions and
 oil and gas industry activities.
- The dominant fraction of the methane emissions is from continuous emissions by the oil
 and gas industry.

22

23 Abstract

Emissions of methane (CH₄) in the Permian basin (U.S.A.) have been derived for 2019 and 2020 24 from satellite observations of the Tropospheric Monitoring Instrument (TROPOMI) using the 25 divergence method, in combination with a data driven method to estimate the background 26 concentrations. The resulting CH₄ emission data, which have been verified using model with 27 known emissions, have a spatial resolution of approximately 10 km. The spatial patterns of the 28 emissions are in a good agreement with the locations of oil and gas production and drilling 29 activities in the Permian basin, as well as with emissions of nitrogen oxides (NO_x). Analysis of 30 31 time-series of locations with large CH₄ emissions indicated that there are significant continuous emissions in this region. The CH₄ emissions can be characterized as a continuous area source, 32 rather than as dominated by a few large unplanned releases. This is important considering 33 possible CH₄ emission mitigation strategies. In addition to providing spatially resolved 34 emissions, the divergence method also provides the total emissions of the Permian basin and its 35 main sub-basins. The total CH₄ emission of the Permian is estimated as 3.0 ± 0.7 Tg yr⁻¹ for 36 2019, which agrees with other independent estimates based on TROPOMI data. For the 37 Delaware sub-basin, it is estimated as 1.4 ± 0.3 Tg yr⁻¹ for 2019, and for the Midland sub-basin 38 1.2 ± 0.3 Tg yr⁻¹. In 2020 the emissions are 8% lower compared to 2019, which could be a result 39 of strong decreases in drilling activities due to the COVID-19 crisis. 40

41 Plain Language Summary

42 Methane is a strong greenhouse gas that contributes to climate change. One of the main emissions sources of methane is from the oil and gas industry. To be able to reduce these 43 44 emissions we have to know the main sources and monitor if reduction measures work. In this study we estimated the emissions for the Permian basin in the U.S.A. using satellite observations. 45 This provides us with maps of the emissions with a spatial resolution of 10 km. The results 46 indicate that continuous emissions are important in the Permian basin. This may be caused by 47 48 small emissions from the thousands of wells in the Permian basin. Also, we were able to estimate the annual emissions from the basin, which correspond well with other studies. 49

50 1. Introduction

51 Methane (CH₄) is the second most important greenhouse gas after CO₂. As the 52 atmospheric lifetime of CH₄ is relatively short at 9.1 years and the global warming potential 53 large, a reduction in CH₄ emissions would lower the combined radiative forcing from greenhouse 54 gases on a timescale of years and is therefore a relatively efficient option to mitigate climate 55 change. For this reason, the Global Methane Pledge was initiated at the UN Climate Change 56 Conference (COP26) in November 2021 [*European Commission, United States of America*, 57 2021], which aims at reducing CH₄ emissions by 30% in 2030.

A significant fraction of global methane emissions comes from the oil and gas (O&G) 58 59 industry [IEA, 2021]. CH₄ is emitted during the construction of new wells, when operating wells, during storage and transportation of oil and gas, and when wells are abandoned. Some of the 60 emissions are intended releases, for example from venting, others are unintentional and caused 61 62 by malfunctioning equipment or by accidents. To be able to effectively reduce CH₄ emissions from the O&G industry, the largest contributions must be known, and which ones can be 63 mitigated with the least effort. In the recent literature there has been a strong focus on the 64 detection of large emissions (aka super-emitters) in O&G production regions using satellite data 65 [Cusworth et al., 2021; Irakulis-Loitxate et al., 2021; Lauvaux et al., 2022], for example in 66 Turkmenistan [Irakulis-Loitxate et al., 2022] and Algeria [Varon et al., 2021]. In this paper we 67 focus on the Permian basin, which is the largest oil and gas producing region in the U.S.A. The 68 Permian basin is located in Texas and New Mexico and covers an area of approximately 160,000 69 km². The exploitation of the Permian basin is mostly done using non-conventional technologies, 70 including hydraulic fracturing and horizontal drilling. The area is characterized by thousands of 71 production facilities and new ones are continuously developed, while others are abandoned when 72 no longer productive. Observations from the ground [Robertson et al., 2020], from aircraft and 73 satellites have shown significant emission of CH4, but also other gases like nitrogen dioxide 74 (NO₂) and formaldehyde [e.g. de Gouw et al., 2020]. For reducing the CH₄ emissions, a key 75 question is if these emissions are dominated by a few large point sources, or caused by many 76 small emissions, which together form a continuous area source. This is important because large 77 emissions from a few facilities will be easier to reduce than small emissions from many facilities 78 79 [Mayfield et al., 2017]. We address this question using satellite data from the Tropospheric Monitoring Instrument (TROPOMI) on board of the European Sentinel 5 Precursor (S5P) 80

satellite [Veefkind et al., 2012], which was launched in 2017. The CH₄ observations of 81 TROPOMI have a spatial resolution of approximately 7x5.5 km² in nadir and larger towards the 82 edges of the 2600 km wide swath. The main contribution of TROPOMI for CH₄ emission 83 monitoring is the continuous mapping capability, providing a large number of overpasses over 84 any given region on Earth. The spatial resolution is not sufficient to detect the frequent but 85 relatively small individual plumes in the Permian. Instead of detecting individual plumes, the 86 aim of this work is to derive CH₄ emissions on a spatial resolution of approximately 10 km. 87 TROPOMI data have been used for quantifying emissions for the Permian basin using the wind 88 rotation method [Schneising et al., 2020], Bayesian inversion involving chemistry-transport 89 modelling [Zhang et al., 2020] and the divergence method [Liu et al., 2021]. The emission 90 estimates of these studies are in the range of 2-4 Tg yr⁻¹ CH₄ for the period 2018-2019. 91

In this work, we use the divergence method with a new data-driven approach to derive the large CH_4 background column, which removes the need for model estimates of the background. Before applying it to satellite data, we verify the method using model data. The spatial distribution of the TROPOMI derived emissions is compared to oil and gas production and drilling information and with TROPOMI derived NO_x emissions. A time series analysis for locations in the Permian basin with large emissions is presented, as well as the estimate of the emissions of the entire Permian basin.

99 2. Materials and Methods

In this section we describe the methods that are used to derive emissions from the column-averaged dry air mole fraction of methane (XCH4). This involves two steps: first the CH4 background and the enhanced column densities are computed, next the divergence is applied to estimate the emissions.

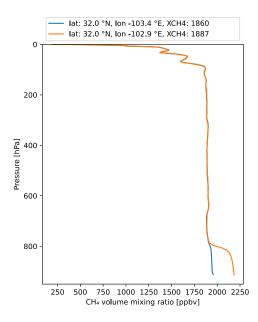
104 2.1 Background correction

105 The background correction first converts the XCH4 volume mixing ratio into the CH_4 106 concentration, *n*, in units of mole m⁻², by multiplying with the dry air number density:

$$n = 10^{-9} \text{ XCH4} \frac{p_{sfc}}{g m_{air}}$$
 Equation 1

107 where *n* is the CH₄ concentration in units of mole m⁻², p_{sfc} is the surface pressure in Pa, *g* 108 is gravitational constant, estimated as 9.81 m² s⁻¹, and m_{air} is the molar mass of dry air (28.9647 g 109 mole⁻¹).

110 Since this study focuses on a small domain, the daily transport in the upper atmosphere, which was estimated by the daily model re-analysis in Liu et al. (2021) can be simplified. To 111 estimate the background CH₄ concentration, a simple model is applied that describes the CH₄ 112 column concentration as the sum of a stratospheric contribution, a background tropospheric 113 concentration and a lower tropospheric enhancement. This model is illustrated in Figure 1, which 114 shows model data for a profile with and without lower tropospheric enhancement. The bulk of 115 116 the CH₄ column is determined by the tropospheric background, for which the concentration is almost constant from the surface to the tropopause (~100 hPa for this profile). The concentration 117 in the stratosphere decreases strongly with altitude. For the example shown in Figure 1, the 118 contribution of the lower tropospheric enhancement contributes 1.4% to the column integrated 119 120 XCH4.



121

- 122 Figure 1. CAMS model (see section 3) CH₄ volume mixing ratio profiles for two nearby locations
- 123 for 1 October 2020. The blue line represents background conditions and the orange line
- 124 *enhanced* CH₄ *concentrations in the lower troposphere.*
- 125

126 Using the model described above, the CH₄ column concentration can be written as:

$$n = k \operatorname{XCH4}_{s} p_{tp} + k \operatorname{XCH4}_{t} (p_{sfc} - p_{tp}) + \Delta n \qquad \text{Equation } 2$$

127 where $k = \frac{10^{-9}}{g m_{air}}$, XCH4_s is the mean CH₄ concentration in the stratosphere, XCH4_t is the mean 128 CH₄ concentration in the troposphere, p_{tp} is the tropopause pressure and Δn is the lower 129 tropospheric CH₄ enhancement.

For a limited-sized area we assume that the tropopause pressure, XCH4_s and XCH4_t are constant over the area, whereas Δn is expected to vary. Although stratospheric intrusions are common in the western U.S. [e.g. Lin et al., 2012], especially in spring, the Permian basin is at the eastern end of the affected area and therefore the impact of the assumption of a constant tropopause pressure will hold on most days. Under these assumptions, Equation 2 can be rewritten as:

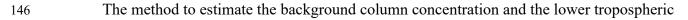
$$\Delta n = n - (c_0 + c_1 p_{sfc}) \qquad \text{Equation 3}$$

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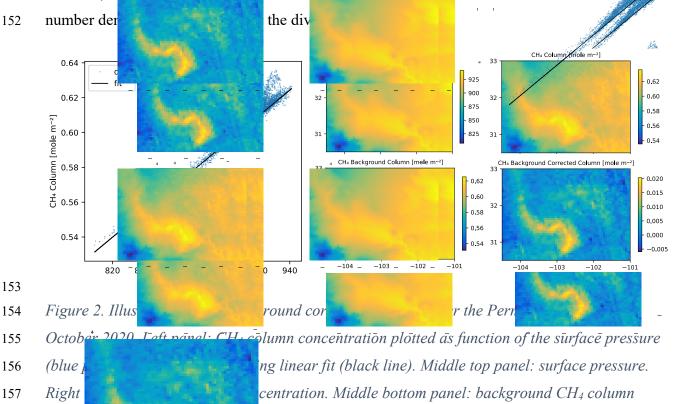
137 where $c_0 = k (XCH4_s - XCH4_t) p_{tp}$ and $c_1 = k XCH4_t$.

Thus, the background can be estimated by linearly fitting the CH_4 concentration *n* as 138 139 function of the surface pressure p_{sfc} , yielding the constants c_0 and c_1 . This is expected to provide an accurate estimate of the background CH₄ concentration when the lower tropospheric 140 enhancement Δn is close to zero for the larger part of the area. To make the fit less sensitive to 141 the enhanced CH₄ concentration values, we first bin the data based on the surface pressure. For 142 each bin we compute the 25th percentile of the CH₄ concentration and the median surface 143 pressure. These binned points are fitted using a linear least-squares fit, yielding the parameters c₀ 144 and c_1 of Equation 3, which describe the background CH₄ column concentration. 145

lages



- 147 enhancement is illustrated in Figure 2 over the Permian region for 6 October 2020. In the figure
- the CH₄ column concentration is shown along with the linear fit. A num
- significantly exceed the background concentration, which is represented
- show the constructed background column concentration and the backgr
- 151 column (Δr). It is noted that Δn will show similar anoticl features as XCH4, however it is not



kground corrected CH₄ column concentration.

Based on the conservation of mass, the emission can be computed as the horizontal flux divergence [Beirle et al., 2019], assuming that the sink term can be neglected due to the long atmospheric lifetime of CH₄ [Liu et al., 2021]. From the continuity equation for steady state conditions, the emission can be computed as:

$$E = \frac{\partial \Delta n_x u}{\partial x} + \frac{\partial \Delta n_y v}{\partial y}$$
 Equation 4,

where x and y are the two perpendicular horizontal directions and v=(u,v)

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2.2 Divergence Method

The divergence is computed using numerical derivatives calculated as the fourth-order or 165 second-order central-finite difference. The divergence method is extended here by introducing a 166 second estimate of the divergence. As illustrated in Figure S1, the standard divergence method 167 computes the divergence based on pixels in the S-N and E-W direction. While this method 168 already has proven very powerful, it doesn't make full use of the observations. A second 169 estimate of the divergence can be computed by using the ground pixels in the SW-NE and NW-170 SE direction (the orange pixels in Figure S1). We compute the wind vector (u', v') on these axes, 171 by applying a rotation of 45° with respect to the original wind vector (u,v). This provides two 172 estimates of the divergence, which are combined by computing a weighted average. The weight 173 depends on the wind direction and is computed for each grid box: when the wind is along the SN 174 or EW direction, more weight is given to the original divergence, and when the wind is in the 175 176 SW-NE or NW-SE direction more weight is given to the rotated divergence. The weight of the SN-EW direction is given by: 177

$$w_0 = \left|\frac{\varphi}{45} - 1\right|$$
 Equation 5,

178 where φ is the angle in degrees between the vectors (u, v) and $(\bar{u}, 0)$. The weight for the 179 NW-SE direction is $1 - w_{0.}$

Including the second estimate of the divergence has two advantages: firstly, it increases the data coverage because one of the divergence values maybe missing due to missing or invalid input data, secondly it potentially reduces the noise, because we use twice as much data to compute the divergence.

Figure S2 illustrates the full workflow that has been developed. The daily XCH4 mixing 184 ratios are background corrected yielding the tropospheric enhancements number densities. 185 Filtering is applied to remove grid boxes for which the terrain height is varying strongly, as the 186 background correction is expected to be inaccurate for these conditions. Additionally, this step 187 avoids the posterior correction on the background correction, which is important in Liu et al., 188 (2021). The tropospheric enhancements are used by the divergence method to compute daily CH₄ 189 190 emissions. As a default we use a boundary layer height of 500m for averaging of the wind and compute the divergence using a fourth-order finite difference. The daily emissions are filtered to 191 192 only include grid boxes which have valid neighbors to remove effects of cloud edges. Mean and

median emissions are computed from the daily data for the years 2019 and 2020 and for theentire period.

195 **3. Data**

196 S5P TROPOMI CH₄ mixing ratios are from the Weighting Function Modified

197 Differential Optical Absorption Spectroscopy (WFM-DOAS) algorithm version 1.5 [Schneising

198 et al., 2019]. The main data fields used are the methane column-averaged dry air mole fraction

199 (XCH4), the surface pressure, the geolocation and the data quality information. The WFMD-

200 DOAS data were downloaded from https://www.iup.uni-

201 <u>bremen.de/carbon_ghg/products/tropomi_wfmd</u> for the years 2019 and 2020.

From the global data, a daily gridded dataset for the years 2019 and 2020 is derived on an 202 equirectangular grid with a resolution of 0.05° in both latitude and longitude, which corresponds 203 to 5.5 km in the N-S direction and for a latitude of 30° to 6.4 km in the E-W direction. Note that 204 the resolution is similar to the nadir resolution of the TROPOMI CH₄ observations (5.5 km in 205 flight direction and 7 km in the cross-flight direction). The gridding processing uses the ground 206 pixel corners provided to calculate the overlap with the grid boxes, which are used to compute 207 weighted averages. The gridding yields daily fields of XCH4 and surface pressure. Only ground 208 pixels with the recommended data quality are included in the gridded fields. 209

Meridional and zonal wind components are from the ERA-5 reanalysis from the European Centre for Medium-Range Weather Forecasts [*Hersbach et al.*, 2020]. The wind data is downloaded for all 14 pressure levels between 600 and 1000 hPa and has a spatial resolution of $0.25^{\circ} \ge 0.25^{\circ}$ latitude/longitude and an hourly temporal resolution. In the divergence method we take the wind history into account by averaging the wind data for the time steps of 17, 18 and 19 hrs UTC.

Model XCH4 and surface pressure data have been used from the Copernicus Atmosphere Monitoring Service (CAMS) global forecasting system (IFS cycle 47R1) [*Agustí-Panareda et al.*, 2019], experiment he9h [*Barré et al.*, 2021]. These data include an adjustment of the concentrations using satellite data assimilation, whereas the emissions and surface fluxes remain unchanged by the satellite data assimilation procedure. Due to their respective vertical sensitivities, the satellite data mainly provide a correction to the concentrations in the free

troposphere and above. At lower altitudes, the emissions are the dominant influence on CH4 222 concentration. The CAMS data have a spatial resolution of 0.1° x 0.1° latitude x longitude. Data 223 for each day of the year 2020 at 18:00 UTC were obtained and resampled to the same spatial grid 224 as the TROPOMI XCH4 data. Also, all grid boxes for which the TROPOMI data has fill values 225 were removed in the CAMS XCH4 data set, to generate representative pseudo-observations. In 226 addition to the CAMS XCH4 data, the CAMS emissions version 4.2 with a spatial resolution of 227 0.1° x 0.1° latitude x longitude have been used [Granier et al., 2019]. The anthropogenic 228 emissions in this data set, including fossil fuel, agricultural and landfill/waste emissions, are 229 from EDGARv4.2FT2010 [Olivier and Janssens-Maenhout, 2012]. From the original monthly 230 data, average emissions for the sum of all sectors for 2019 and 2020 have been constructed. 231 232 Oil and gas production data and drill rig counts are from the Enverus Drilling Info and

Rig Analytics data base tools (<u>https://www.enverus.com/drillinginfo-and-rigdata/</u>, last accessed 25-2-2022), respectively. Oil and gas production volumes are reported monthly for each well location and gridded to match the TROPOMI CH₄ maps. The locations of drill rigs are reported daily. Monthly gridded maps are created by counting the number of drill rigs within each grid cell weighted by the number of days on location per month.

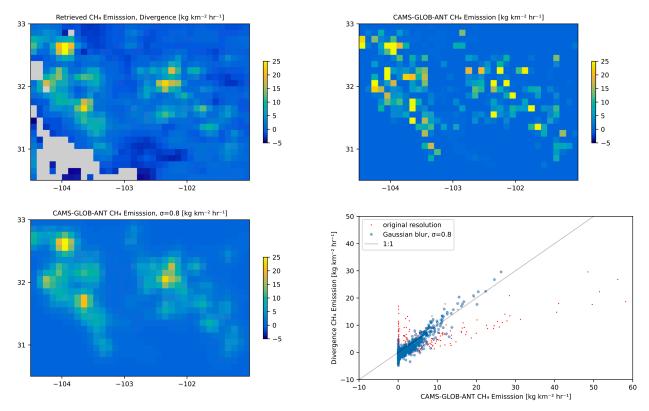
The NO_x emission data are from [*Dix et al.*, 2022]. These data represent a mean NO_x emission for the time period May 2018 until December 2020. The data have been regridded to the same latitude-longitude grid as the TROPOMI XCH4 data.

241 4. Verification using CAMS model data

The divergence method has been verified using CAMS model data, for which the input 242 emissions are known. As described above, the CAMS data for 2020 were gridded and sampled 243 for the grid boxes for which also TROPOMI XCH4 is available. Thus, it has the same coverage 244 as the TROPOMI, including missing data due to for example cloud contamination. On this 245 dataset we apply the same background correction and derive the emissions using the divergence 246 method. Figure 3 shows the median emission for 2020 derived from the CAMS data as well as 247 the input emissions. The retrieved emissions show generally the same spatial features as the 248 input emissions, however at a lower spatial resolution. To test this, we applied a Gaussian blur to 249 the input emissions. We manually varied the standard deviation σ of the Gaussian kernel and 250 found that for a value of approximately 9 km the resolution of the retrieved and input emissions 251

match well. The spatial resolution has a strong effect on the slope between the retrieved and 252 input emissions, and the correlation coefficient increases from 0.75 to 0.93 when the Gaussian 253 blur is applied. Whereas the divergence method favorably retrieves the spatial variability of the 254 emissions, retrieving the total emission is more complicated. We computed the total emission for 255 the Delaware and Midland sub-basins, as well as for the entire Permian basin (the definition of 256 these regions is shown in *Figure S3*). The estimated emission for the Delaware sub-basin shows 257 a bias of -16% lower compared to the CAMS inputs, for the Midland sub-basin -40% and for the 258 entire Permian basin -42%. The total emission estimates are sensitive to a possible offset: 259 applying an offset of 1.0 kg km⁻² hr⁻¹ to the retrieved emissions is sufficient to close the gap 260 between the retrieval and the input. The bias with this offset applied is 7%, -8% and -1%, for the 261 Delaware sub-basin, the Midland sub-basin and the entire Permian basin. The mean emission of 262 263 the Delaware sub-basin is almost a factor of 2.0 larger compared to the mean for the entire Permian basin. For the Midland basin this is a factor of 2.6. For larger emissions, a bias will have 264 a smaller relative impact than for lower emissions. This agrees with the finding that the bias is 265 lower for the Delaware sub-basin. Thus, these results point to a possible low bias of the method, 266 which is a potential limitation of the method for estimating the total emission of a region, 267 especially when the average emissions are small. A possible complication of this analysis is that 268 in the CAMS dataset the concentrations and emissions are not consistent, because only the 269 concentrations are adjusted using satellite observations. However, it is not straightforward to 270 271 assess the sign or magnitude that this may have on the emissions derived with the divergence 272 method.

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Figure 3. Divergence method applied to CAMS model data. Top-left panel: CH₄ emission derived from CAMS model data with the divergence method. Top-right panel: CAMS input emissions on the original resolution. Bottom-left panel: CAMS input emission with a Gaussian blur with σ =0.8. Bottom-right panel: retrieved emissions plotted as function of the CAMS input emissions, for the original and blurred data.

The CAMS data have also been used to test the sensitivity for variations in the setup. The 280 default divergence setup uses a boundary layer height of 500 m and a fourth-order central-finite 281 difference method for calculating the derivatives. We have tested the impact of boundary layer 282 heights of 250 and 1000 m, the use of second-order central-finite difference, and the use of a 283 least-squares fit (instead of the fit based on the 25th percentile) to compute the background. The 284 boundary layer height is important because the wind is computed as the mean over this layer. As 285 can be seen in Table 1, the largest impact on the mean emission is due to the background method 286 (-19% compared to the default) and increasing the boundary layer to 1000 m (-11%). The impact 287 of changing the boundary layer to 250 m and the impact of the central-finite difference method 288 order is marginal (<2%). These results are similar to the results presented in [Liu et al., 2021]. 289

290 Based on this sensitivity analyses, we estimate the uncertainty of the mean emissions of the order

291 25%.

Emission [kg m ⁻² hr ⁻¹]	Default	250 m	1000 m	2 nd order	Lstsq
Mean	1.44	1.47	1.28	1.43	1.17
Median	0.16	0.23	0.01	0.21	-0.20
P1	-0.90	-0.84	-1.04	-0.85	-1.08
P3	2.20	2.19	2.10	2.24	1.90

292

293 Table 1. Sensitivity of the emissions for 2020 derived from the CAMS data for the entire domain,

for variations in the assumed boundary layer height, the order of the central-finite difference

295 method and the method to compute the background. The first column represents the default case,

which uses a boundary layer height of 500m, 4^{th} order central finite difference and a fit of the

297 background concentration based on the 25th percentiles. Variations with respect to the default

are a 250 m or 1000m boundary layer height, a 2^{nd} order the central-finite difference method,

299 and a least-squares fit (Lstsq) for the background concentration. The table lists the mean,

300 *median, 25th percentile (P1) and 75th percentile (P3) of the difference with the default case.*

301

5. CH₄ derived from TROPOMI Data

The median CH₄ emissions derived by applying the divergence method to TROPOMI 302 WFMD data for 2019-2020 are shown in Figure 4. Overall, the emission is highest in regions 303 304 where there are activities related to the O&G industry. The spatial outline of the Delaware and 305 Midland sub-basins can be distinguished clearly in this figure. Whereas the main outlines correspond to the emission data derived from the CAMS model data (Figure 3, top left panel), 306 there are significant differences between these maps. The TROPOMI results show significantly 307 higher spatial variability. This may be caused by the instrument noise, which is not accounted for 308 309 in the CAMS analysis. In addition, the CAMS model assumes a fixed pattern, whereas in reality, the emissions will vary significantly in space and time. Also, for the CAMS data the wind 310 information and the advection in the model are consistent, whereas for the application on 311 TROPOMI data there may be significant errors in the wind fields. Some regions in *Figure 4* 312 show negative emissions, which are considered artefacts of the method. In regions with 313 significant variability in orography, the assumed vertical model (Equation 2) may not hold, 314 315 leading to such artefacts.

The NO_x emissions in the Permian basin are to a large extent related to the O&G 316 industry. Although the NO_x emissions come from different sources, mainly generators and 317 engines, they are expected to come from the same sites as where the CH₄ is emitted [Warneke et 318 al., 2014]. The NO_x and CH₄ emissions derived from TROPOMI (Figure 4) show similar spatial 319 variations. The spatial correlation is higher in the western Delaware basin. NO_x also has 320 significant contributions from road transportation and power generation. In the NO_x emission, 321 spatial features related to the main cities, Midland and Odessa and the Interstate I-20 can be 322 distinguished [Dix et al., 2022]. As expected, these features are not found in the CH₄ emission 323 data. 324

To further link the satellite derived CH₄ emission data with the O&G industry activities, 325 we used data on the oil and gas production and on the drilling days. For each grid box we define 326 a score of 0-3 for oil production, gas production and drilling. A score of 0 is given when the 327 activity data are less than 1% of the median, a score of 1 when the data are higher than this value 328 but less than the 25th percentile, a score of 2 when the data are between the 25th and 75th 329 percentile, and a score 3 when the data are higher than the 75th percentile. Finally, the scores of 330 oil production, gas production and drilling are combined, which gives 10 categories ranging from 331 0 to 9. The map of the categories and the distribution of the CH_4 emissions over the categories 332 are shown in Figure 4. As can be seen in the figure, the overall spatial variation of the production 333 and drilling data shows good agreement with both the CH₄ and NO_x emissions. Especially for the 334 categories 7-9 significantly higher CH₄ and NO_x emissions are found. The lower categories show 335 an average value of near zero, which is also a sign that the satellite retrievals are in 336 correspondence with the oil and gas activity data. 337

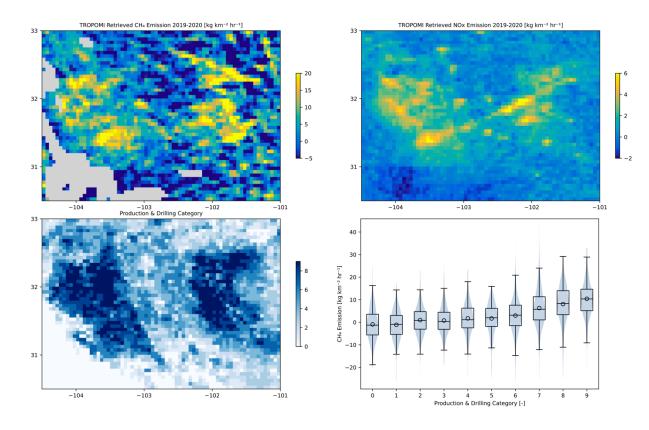


Figure 4. Panel top left: Median CH₄ emission derived using the divergence method
applied to TROPOMI data from 2019-2020. Top-right NO_x emission for 2018-2020 retrieved
from TROPOMI [REF Dix]. Bottom left: production drilling categories. Bottom right, combined
violin and boxplot showing the distribution of retrieved CH₄ emissions for each
production/drilling category.

338

For five locations with high median emissions, labeled A to E in *Figure S3*, a time series 344 345 of the CH₄ emission for 2019 and 2020 are shown in *Figure 5*. These five locations have been selected by hand from the median emission map (Figure S3). For all locations, except for 346 location C, the 30-day running mean and 30-day running median are above zero for almost the 347 entire time period. As can be seen in Figure S3, there is an area with negative median emissions 348 just north of the location C. These negative emissions are probably an artefact of the orography 349 and are also affect the time series for the location C. For all five locations in Figure 5 the mean 350 emission over the whole time period is larger than the median value, indicating that the 351 distribution is skewed towards the larger values. Although the mean value is much more 352 sensitive to outliers compared to the median, the mean falls well in the interquartile range. 353 Furthermore, the difference between the mean and median is less than 30% for the five locations. 354

355 The median value is more representative for the continuous emissions and less sensitive to

extreme values compared to the mean. For reference, *Figure S4* shows a similar plot as *Figure 5*,

but for five locations with background emission values. For these background locations the

difference between the mean and median over the entire time period is smaller as compared to

the locations with high emissions.

We also analyzed the distribution of the daily emissions for the Permian basin and the sub-basins. Overall, the distributions are heavy tailed and skewed towards the large values. For the 2019-2020 period the mean is 30% higher than then median for the entire Permian and 24% and 27% for the Delaware and the Midland sub-basins.

Super emitting events that are short in duration will have a much larger effect on the mean than on the median. The fact that the mean and median differ by less than 32% for both the time series for high emitting locations as well as for the entire Permian basin, indicates that a large fraction of the emissions is continuous rather than episodic. This is further supported by the inspection of monthly maps, which show similar spatial patters of the main CH₄ emission hotspots.

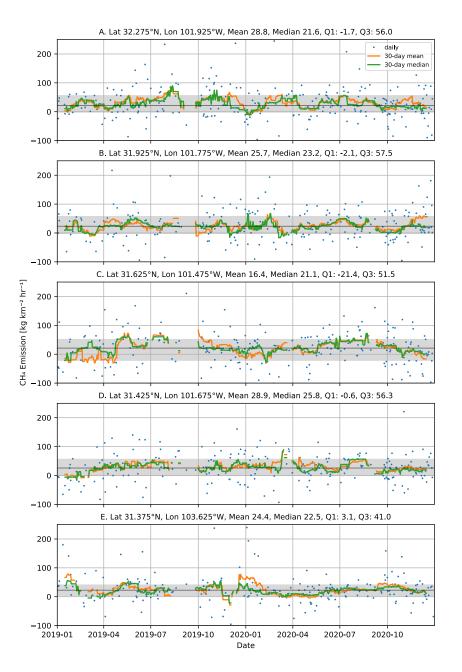


Figure 5. Time series for five locations with high CH₄ emissions. The blue dots are the daily data, the orange line the 30-day running mean and the green line represents the 30-day running median. Running mean and medians are only shown when at least 5 of the 30 days contain valid data. The grey area indicates the interquartile range and the black line the median over the whole time period.

370

From the daily emissions we estimated the annual emissions for three regions: the
Delaware sub-basin, the Midland sub-basin and the entire Permian. The boundaries used for the

378 Delaware and Midland basin are shown in *Figure S3*, the Permian basin covers the entire map.

The annual emissions were calculated by summing the yearly mean values of a basin and were

converted to Tg yr⁻¹. The derived annual emissions are listed in *Table 2* for 2019, 2020 and

2019-2020. Based on the sensitivity analysis presented in Section 2, we estimate the uncertainty in these numbers as 25%. This uncertainty is dominated by biases caused by the configuration of the method. Therefore, the uncertainties in the annual emissions are expected to be significantly

384 larger compared to differences in the annual emissions between the years.

The annual CH₄ emissions of the Delaware and the Midland basin are found to be comparable. It is estimated that these sub-basins contribute 70-90% to the entire CH₄ emissions of the Permian basin. The derived emissions are found to be 8 to 27 % lower in 2020 compared to 2019, which could be an indication of the impact of the COVID-19 crisis [*Lyon et al.*, 2021].

389 Table 2. Annual Emissions derived with the divergence method for the Delaware,

390 Midland sub-basins and for the entire Permian basin for 2019, 2020 and 2019-2020. The

391	Delaware and Midland basins are sub-basins	of the Permian (see Figure S3).
571		

Annual Emission [Tg yr ⁻¹]	Delaware	Midland	Permian
2019	1.4	1.2	3.0
2020	1.1	0.9	2.7
2019-2020	1.3	1.0	2.9
Difference 2020 - 2019	-18%	-27%	-8%
Estimated 1- σ uncertainty of t	the emission is 25%		

392

393 6. Conclusions

We have investigated the CH₄ emissions from the oil and gas industry in the Permian basin for 2019 and 2020 using satellite data. We used the divergence method to derive daily CH₄ emissions. Compared to previous applications of the divergence method for CH₄ emissions [*Liu et al.*, 2021], we applied a model-independent correction to derive the CH₄ background concentration. This method assumes that the background CH₄ concertation in the troposphere and the tropopause height is spatially constant and is therefore limited to regional applications.

400 Using model data for 2020 we demonstrated that the divergence method can retrieve the spatial

401 variability of the emissions at a reduced spatial resolution of approximately $10x10 \text{ km}^2$. Based on 402 a sensitivity analyses, we estimate that the uncertainty in the yearly mean emissions is of the 403 order 25%.

The divergence method to TROPOMI CH₄ data was applied for the years 2019-2020: 404 The spatial patterns between these years agree well, indicating that the method gives consistent 405 results. The spatial distribution shows generally the same structures as oil and gas activity data 406 407 (production and drilling) and NO_x emissions, also derived using TROPOMI. However, there is a weak spatial correlation between the CH₄ emissions and the activity and NO_x emissions. 408 Therefore, constructing CH₄ emissions from these data sets, as suggested by [de Gouw et al., 409 2020] will require additional information, for example locally varying conversion factors. The 410 spatially variability of the retrieved emissions differs clearly from the CAMS emission data set 411 [*Granier et al.*, 2019]. 412

Time series for locations with high CH₄ emissions and background values were analyzed. 413 The background locations have near zero or negative median emissions over the time series. The 414 interquartile range in the data is typically 40 kg km⁻² hr⁻¹. The time series of high emission 415 locations shows for four out of five locations a skewed distribution towards higher values. 416 However, the difference of the mean and median over the time series differ less than 32%, 417 indicating that the mean value is not dominated by a few outliers. From this analysis we 418 speculate that the emissions in the Permian basin are driven by continuous emissions, rather than 419 by a few large unplanned releases. This is important, because it means that the emissions may be 420 caused by the daily operations and given the large number of facilities in the Permian, they may 421 be hard to reduce. The continuous emission was also found by [Schneising et al., 2020] for the 422 entire Permian basin, and here we confirm that this also applies to spatially resolved data, 423

The total emissions were estimated for the Delaware and Midland basins and for the entire Permian basin. For the entire basin we find values 3.0 ± 0.7 Tg yr⁻¹ for 2019, For the Delaware basin we find emissions of 1.4 ± 0.3 Tg yr⁻¹ and for the Midland basin 1.2 ± 0.2 Tg yr⁻¹ ¹. The estimated 25% uncertainty is based on a sensitivity analysis of the retrieval method. The estimated emission for the Permian basin agrees within the uncertainties with estimates from [*Liu et al.*, 2021] (2.82 to 3.78 Tg yr⁻¹), [*Schneising et al.*, 2020] (3.18±1.13 Tg yr⁻¹ for 20182019) and [*Zhang et al.*, 2020] (2.9±0.5 Tg yr⁻¹ for March 2018 – March 2019).

Annual emissions in the Permian basin for the year 2020 are 8% to 27% lower compared to 2019. Updated results using the method of [*Schneising et al.*, 2020] using the same data version as used in this work, show agreement for 2019 (2.92 ± -1.61 Tg yr⁻¹), 2020 (2.27 ± -1.75 Tg yr⁻¹) and confirm the reduction in emission in 2020 compared to 2019. A possible explanation is the drop in demand for oil and gas, leading to significantly reduced drilling activity and reduced production in the Permian basin in 2020.

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national contributions from The Netherlands, Germany, and Belgium.

449

450 **Open Research**

451 The main data sets that are used in this research are:

452	•	The TROPOMI WFM-DOAS data available at https://www.iup.uni-
453		bremen.de/carbon_ghg/products/tropomi_wfmd/
454	٠	The ERA-5 meteorological information available at
455		https://www.ecmwf.int/en/forecasts/datasets/reanalysis-datasets/era5
456	•	The CAMS model data are available from ECMWF.

457	Upon publication all Python scripts and data files used to generate and analyze the data
458	will be made available for public access.
459	
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531

532 Figure Captions

- Figure 1. CAMS model (see section 3) CH₄ volume mixing ratio profiles for two nearby locations for 1 October 2020.
 The blue line represents background conditions and the orange line enhanced CH₄ concentrations in the lower
 troposphere.
- Figure 2. Illustration of the background correction for data over the Permian region for 6 October 2020. Left panel:
 CH₄ column concentration plotted as function of the surface pressure (blue points) and the corresponding
 linear fit (black line). Middle top panel: surface pressure. Right top panel: CH₄ column concentration. Middle
 bottom panel: background CH₄ column concentration. Right bottom: background corrected CH₄ column
- 539bottom panel: background CH4 column concentration. Right bottom: background corrected CH4 column540concentration.
- Figure 3. Divergence method applied to CAMS model data. Top-left panel: CH₄ emission derived from CAMS model
 data with the divergence method. Top-right panel: CAMS input emissions on the original resolution. Bottom left panel: CAMS input emission with a Gaussian blur with σ=0.8. Bottom-right panel: retrieved emissions
 plotted as function of the CAMS input emissions, for the original and blurred data.
- Figure 4. Panel top left: Median CH₄ emission derived using the divergence method applied to TROPOMI data from
 2019-2020. Top-right NOx emission for 2018-2020 retrieved from TROPOMI [REF Dix]. Bottom left: production
 drilling categories. Bottom right, combined violin and boxplot showing the distribution of retrieved CH₄
 emissions for each production/drilling category.
- Figure 5. Time series for five locations with high CH₄ emissions. The blue dots are the daily data, the orange line the 30-day running mean and the green line represents the 30-day running median. Running mean and medians
- are only shown when at least 5 of the 30 days contain valid data. The grey area indicates the interquartile
- range and the black line the median over the whole time period.
- 553

554 Table Captions

- 555 Table 1. Sensitivity of the emissions for 2020 derived from the CAMS data for the entire domain, for variations in 556 the assumed boundary layer height, the order of the central-finite difference method and the method to 557 compute the background. The first column represents the default case, which uses a boundary layer height of 558 500m, 4th order central finite difference and a fit of the background concentration based on the 25th 559 percentiles. Variations with respect to the default are a 250 m or 1000m boundary layer height, a 2nd order the central-finite difference method, and a least-squares fit (Lstsq) for the background concentration. The 560 561 table lists the mean, median, 25th percentile (P1) and 75th percentile (P3) of the difference with the default 562 case. 563 Table 2. Annual Emissions derived with the divergence method for the Delaware, Midland sub-basins and for the
- and the divergence method for the Delaware, Midland sub-basins and for the
 entire Permian basin for 2019, 2020 and 2019-2020. The Delaware and Midland basins are sub-basins of the
 Permian (see Figure S3).
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Supporting Information for

Continuous Methane Emissions from the Oil and Gas Industry in the Permian Basin

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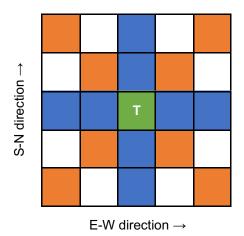


Figure S1. Schematic view of the ground pixels used to compute the divergence. For the green target pixel the standard divergence is computed based on the blue pixels. The rotated divergence is computed based on the orange pixels.

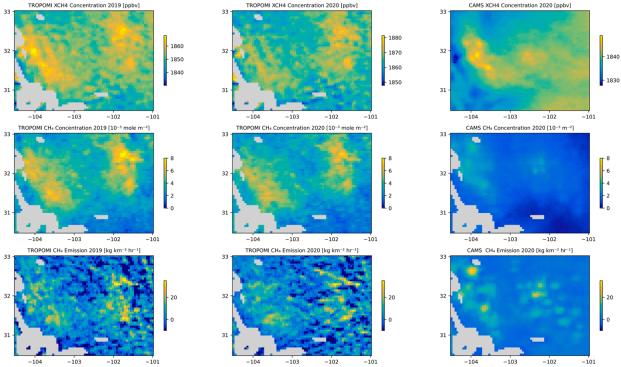


Figure S2. Top row: median XCH4 for Tropomi for 2019 (left), 2020 (middle) and for the CAMS model data for 2020. Middle row: same as top row, but for the background corrected CH₄ column concentrations. Bottom row: same as top row, but for the CH4 emissions derived with the divergence method.

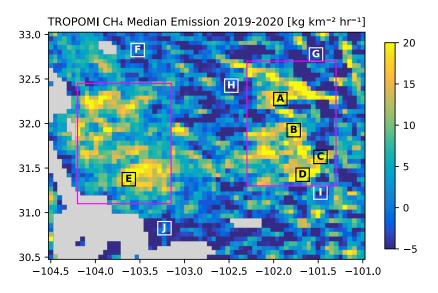


Figure S3. CH4 median emission with the five selected locations with high emissions (labeled A-E in black) and background emissions (labeled F-J in white). The pink boxes indicate the areas used for computing the statistics for the Delaware and Midland sub-basins

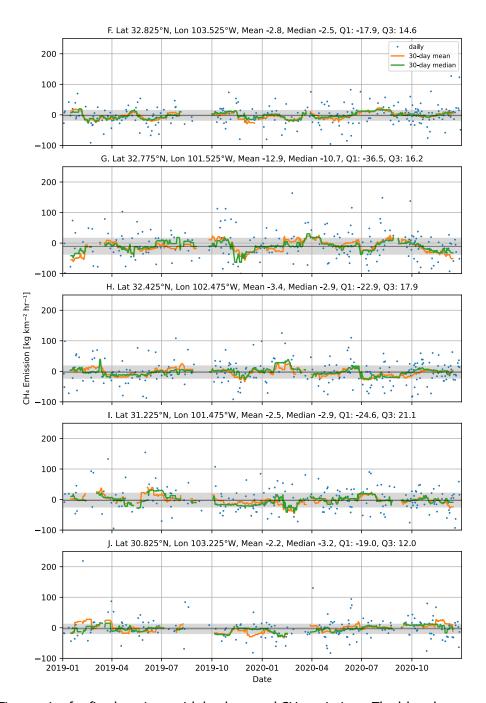


Figure S4. Time series for five locations with background CH₄ emissions. The blue dots are the daily data, the orange line the 30-day running mean and the green line represents the 30-day running median. Running mean and medians are only shown when at least 5 of the 30 days contain valid data. The grey area indicates the interquartile range and the black line the median over the whole time period.