



Evidence of successful methane mitigation in one of Europe's most important oil production region

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Abstract. Reducing methane emissions from the oil and gas production infrastructure is a cost-effective way for limiting global warming. In 2019, a measurement campaign in southern Romania found emission rates from the oil and gas sector substantially higher than the nationally reported emissions, with a few high-emitting sources (“super-emitters”) contributing disproportionately to total emissions. In 2021, our follow-up airborne remote sensing campaign, covering over 80 % of production sites, revealed a marked decrease in super-emitters. The observed change in the number of emitters is consistent with an emission reduction by 20 %–60 % from 2019 to 2021. This reduction is likely due to improvements in production infrastructure following the first campaign in 2019. This is further supported by additional site visits, which showed that many of the leaks identified in 2019 had indeed been mitigated. However, our top-down quantification remains higher than the bottom-up emission reports. Our study highlights the importance of measurement-based emission monitoring of climate change mitigation measures and illustrates the value of a multi-scale assessment integrating ground-based observations with large-scale airborne mapping to capture both the primary mode of emission sources and the rare, but significant, super-emitters.

1 Introduction

In 2015, the United Nations Climate Change Conference (UNFCCC) in Paris agreed to limit global warming well below 2 °C, which requires massive reductions in greenhouse gas (GHG) emissions (UNFCCC, 2015). Methane (CH₄) is a major contributor to global warming and an attractive near-term target for climate change mitigation. In 2021, this was acknowledged in the Global Methane Pledge (GMP) (European Commission and United States of America, 2021). In particular, CH₄ emission reductions related to oil and gas

(O&G) infrastructure are considered “no-regret” solutions, as they have mainly positive effects and can often be realized in a cost-effective way (Hopkins et al., 2016; McKain et al., 2015). In the 27 EU states, CH₄ emissions from O&G production constitute about one-third of the total reported emissions caused by the energy sector (UNFCCC, 2023a). The International Energy Agency (IEA) estimates that Romania’s onshore O&G production sector has the highest share of all EU countries, contributing 16 % to the total onshore O&G emissions (IEA, 2021), even after reported emissions have allegedly decreased by more than 85 % since 1990 (UNFCCC,

2023b), However, reported CH₄ emissions are highly uncertain as direct CH₄ emission measurements to independently verify these numbers are lacking.

To close this gap, the Romanian Methane Emissions from Oil and Gas (ROMEIO) measurement campaign provided the first independent estimates of CH₄ emission rates from O&G production in Romania. The campaign took place in October 2019 and covered the southern part of the country with ground, drone and aircraft in situ measurements (Delre et al., 2022; Korbeń et al., 2022; Stavropoulou et al., 2023; Maazallahi et al., 2025). The national-scale emissions derived from the campaign for oil production sites were already higher than the total reported emissions for the entire O&G sector in Romania. Furthermore, 10 % of the sites emitted more than 10 kg h⁻¹, accounting for more than 70 % of the total emissions (Stavropoulou et al., 2023). To accurately determine the total CH₄ emissions, it is therefore critical to detect and quantify a statistically robust number of “super-emitters”, i.e., those sources in the high-emitting tail of the emission distribution that contribute significantly to the total emissions. For the O&G sector in Romania, this threshold is about 10 kg h⁻¹. Identifying a robust number of super-emitters is challenging with ground-based surveys due to the large number of production sites and associated infrastructure distributed over a vast and sometimes difficult-to-access area. In 2021, we therefore deployed the airborne AVIRIS-NG (Airborne Visible InfraRed Imaging Spectrometer – Next Generation) imaging spectrometer in southern Romania. AVIRIS-NG provides excellent spatial coverage with a detection limit (> 10 kg h⁻¹) well-suited for detecting the emissions of super-emitters (e.g., Thorpe et al., 2013, 2017; Frankenberg et al., 2016; Foote et al., 2020; Borchart et al., 2021).

In this work, we present the results from the AVIRIS-NG measurement campaign in 2021. The AVIRIS-NG measurements are used to detect and quantify the largest CH₄ sources in the study area. The results were integrated with the ground-based data from 2019 to more accurately constrain the annual CH₄ emissions of the O&G sector in the country.

2 Data and methods

2.1 AVIRIS-NG campaign in 2021

The AVIRIS-NG flights were conducted on Thursday 29 and Friday 30 July 2021 over the same region as the ROMEIO campaign in 2019 (Fig. 1). AVIRIS-NG is a push-broom imaging spectrometer sampling 600 pixels in the across-flight direction over a 34° field of view (Hamlin et al., 2011). This results in a ground pixel size of about 5 m for an altitude of 6000 m above ground. The spectrometer samples a spectral range from 380 to 2510 nm at 5 nm with a resolution of 5–7.5 nm full width at half maximum (FWHM). The spectral range includes two spectral windows with CH₄ absorption

lines at 1.6 and 2.4 μm. The signal-to-noise ratio (SNR) in these windows is about 400 for a solar zenith angle of 30° and surface reflectance of 30 % (Cusworth et al., 2019).

The flights covered an area of approximately 3000 km² that includes 66 % of known processing facilities and 82 % of O&G production sites in the region. This translates to 582 processing facilities and 2805 oil and 299 gas production sites (Table S3 in the Supplement). Weather conditions were mostly cloud-free with low wind speeds of about 0.5 m s⁻¹ in the morning (08:00–10:00 UTC) of the first day that increased to 3.0 m s⁻¹ by noon and remained high on the second day. Some lines were flown twice due to the presences of scattered clouds during first data collection.

2.2 Methane retrieval algorithm

2.2.1 Matched filter

CH₄ enhancements were retrieved from the AVIRIS-NG radiance cube using the well-established matched filter method following the implementation by Foote et al. (2020). The matched filter \mathbf{q} is a linear filter that is applied to each radiance spectrum \mathbf{y}_i to obtain a scalar α_i , which is the enhancement in the CH₄ column density above the background (in units of ppm m). For a linear radiance model, we can write the product

$$\mathbf{q}^T \cdot \mathbf{y}_i = \mathbf{q}^T \cdot (\boldsymbol{\mu} + \alpha_i \mathbf{t} + \boldsymbol{\varepsilon}), \quad (1)$$

where $\boldsymbol{\mu}$ is the mean radiance spectrum, \mathbf{t} is the CH₄ target signature, and $\boldsymbol{\varepsilon}$ is the remaining clutter in the spectrum including instrument errors and variability in the surface reflectance. The solution for the optimal matched filter maximizes the signal (second term in brackets) and minimizes the clutter (third term). The CH₄ column enhancement is then

$$\alpha(\mathbf{y}_i) = \frac{(\mathbf{y}_i - \hat{\boldsymbol{\mu}})^T \cdot \hat{\mathbf{S}}^{-1} \cdot \mathbf{t}}{\mathbf{t}^T \cdot \hat{\mathbf{S}}^{-1} \cdot \mathbf{t}}, \quad (2)$$

where $\hat{\boldsymbol{\mu}}$ and $\hat{\mathbf{S}}$ are the mean vector and the covariance matrix estimated from the radiance cube.

A suitable target signature \mathbf{t} can be obtained by computing the change in the at-sensor radiance spectrum L_ε due to an enhancement in CH₄ absorption using Lambert–Beer’s law:

$$L_\varepsilon(\lambda) = L_0(\lambda) \exp(-\alpha_\varepsilon s(\lambda)), \quad (3)$$

where $s(\lambda)$ is the unit absorption spectrum (in (ppm m)⁻¹) and $L_\varepsilon(\lambda)$ and $L_0(\lambda)$ are the radiance spectra with and without a CH₄ enhancement. For a small absorption, Eq. (3) can be approximated by

$$L(\lambda) \approx L_0(\lambda) + L_0(\lambda) \alpha s(\lambda), \quad (4)$$

which, if $L_0(\lambda)$ is approximated by the mean spectrum $\boldsymbol{\mu}(\lambda)$, gives the target signature from Eq. (1) as

$$\mathbf{t}(\lambda) = s(\lambda) \cdot \boldsymbol{\mu}(\lambda). \quad (5)$$

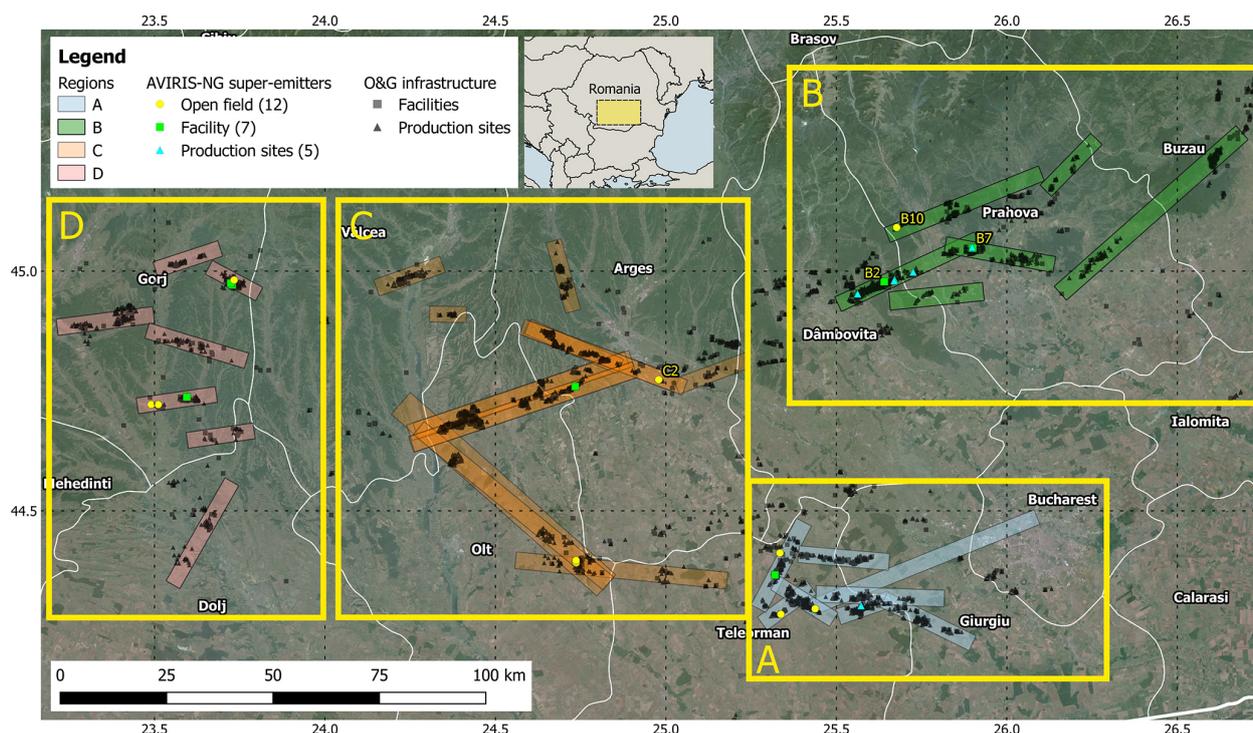


Figure 1. Map with AVIRIS-NG flight lines in 2021 for four regions (A–D) with locations of super-emitters and O&G infrastructure. Background map from the Copernicus Land Monitoring Service (GloLand/VHR_2018_WM).

The unit absorption spectrum s can be computed from Eq. (3) as the change in the natural logarithm of at-sensor radiance spectrum L_ε due to an enhancement of 1 ppm in the surface layer:

$$s(\lambda) = \frac{\ln L_0(\lambda) - \ln L_\varepsilon(\lambda)}{\alpha_\varepsilon}. \quad (6)$$

In this study, we used a simplified but fast forward model that ignores atmospheric scattering to compute the at-sensor radiances (L_0 and L_ε) for a Lambertian surface:

$$L(\lambda) = \mu_0 E_0(\lambda) \frac{\rho}{\pi} \cdot e^{-m \left(\left(1 + \frac{x_\varepsilon}{x_t}\right) \tau_t(\lambda) + \tau_{BG}(\lambda) \right)}, \quad (7)$$

with the cosine of the solar zenith angle μ_0 , solar irradiance spectrum E_0 (Coddington et al., 2021), surface reflectance ($\rho = 1.0$) and the geometric air mass factor for a nadir-viewing instrument m . The CH_4 optical depth in the surface layer (i.e., below 1000 m) is given by $\tau_t(\lambda)$. $\tau_{BG}(\lambda)$ is the optical depth of all other gases including the part of the CH_4 column above the surface layer. x_ε is the CH_4 enhancement (in ppmv) above the CH_4 background concentration x_t , i.e., $x_\varepsilon = 0$ ppm for L_0 and $x_\varepsilon = 1$ ppm for L_ε .

The optical depth profiles are computed for a mid-latitude summer reference atmosphere using the atmospheric radiative transfer simulator (arts, Version 2.2; Buehler et al., 2018) with absorption lines from the HITRAN 2012 database (Rothman et al., 2013). The profiles are scaled to obtain

column-dry-averaged CO_2 and CH_4 mixing ratios of 416 and 1.90 ppm for 2021, respectively (NOAA, 2022). Surface elevation and aircraft altitude for computing the geometric air mass factor are available from the AVIRIS-NG dataset.

Figure 2 shows an example of the unit absorption spectrum computed with the above model equation (dashed line) for the two spectral windows (1480–1800 and 2080–2450 nm) used for the CH_4 retrieval. In addition, the range of the unit absorption spectra computed with the detailed libRadtran radiative transfer model (Emde et al., 2016) is shown for a scattering atmosphere with varying aerosol optical depths (AODs) and surface reflectance (RHO).

Application to AVIRIS-NG radiance cubes

We apply the matched filter simultaneously to both spectral windows (i.e., 1480–1800 and 2080–2450 nm) shown in Fig. 2 to minimize the noise in the retrieved CH_4 field. We do not use a wider spectral window to avoid potential underestimation of CH_4 enhancements as reported by Roger et al. (2024).

The matched filter needs to be applied to each across-track line of the AVIRIS-NG independently to avoid stripes in the retrieved methane fields caused by small radiometric and spectral calibration differences between these lines, which is problem common to push-broom imagers. For short lines, we combined across-track positions to have at least 7500 spectra for an unbiased estimate of mean and covariance matrix.

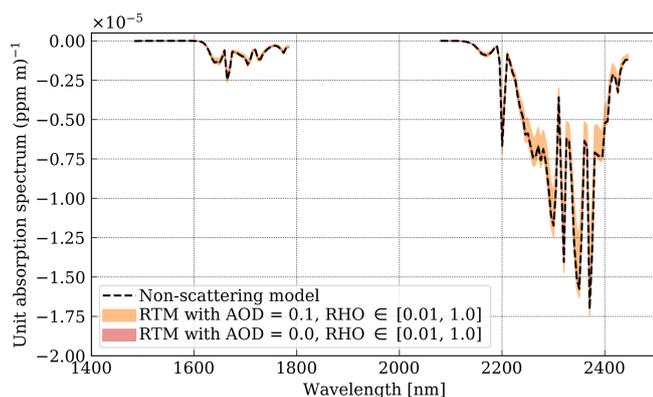


Figure 2. Unit absorption spectrum computed with the non-scattering model used in this study compared to libRadtran radiative transfer simulations using different surface reflectance (RHO) and aerosol optical depths (AODs) in two spectral windows.

The assumption that L_0 can be approximated by the mean spectrum $\hat{\mu}$ results in a bias in enhancement α that needs to be corrected using a scaling factor. Foote et al. (2020) computed the correction factor as the deviation of the radiance spectrum from the mean spectrum:

$$\alpha(y_i) = \frac{1}{R_i} \frac{(y_i - \hat{\mu})^T \cdot \hat{S}^{-1} \cdot t}{t^T \cdot \hat{S}^{-1} \cdot t} \quad \text{with } R_i = \frac{y_i^T \cdot \hat{\mu}}{\hat{\mu}^T \cdot \hat{\mu}}. \quad (8)$$

Since we apply the matched filter to two spectral windows that can have different surface reflectance, we extend their approach by computing a correction factor for each spectral window separately, which removes biases if surface reflectance strongly differs in the two bands based on tests with synthetic spectra. While the correction factor reduces biases in the CH₄ enhancements, it increases retrieval noise for dark surfaces with a low signal-to-noise ratio (e.g., forests and water). Note that the second effect can cause the CH₄ retrieval to appear to depend on surface reflectance.

2.3 Emission quantification

CH₄ emissions plumes were identified by visual inspection of the retrieved CH₄ maps. Local CH₄ enhancements were classified as plume if they (a) were a cluster of pixels with plume-like shape, (b) had significant enhancement above the (local) background variability, and (c) did not correlate spatially with surface reflectance or surface features. Once a source had been identified, a plume detection algorithm was used for plume segmentation (Kuhlmann et al., 2019, 2021). The algorithm uses a threshold to identify pixels where the local mean of the signals is significantly enhanced above the background. The threshold was manually adjusted for each plume to fully cover the visible plume and to avoid false positives. A center line was fitted through the ridge of the detected plume to provide natural coordinates of along- and across-plume directions. The algorithms are part of the open-

source Python library for data-driven emission quantification (ddeg; Kuhlmann et al., 2024a).

The CH₄ emissions were determined from the detected plume using the integrated mass enhancement (IME; Frankenberg et al., 2016; Varon et al., 2018) method implemented in the *ddeg* library (Kuhlmann et al., 2024a). The IME method is a mass-balance approach derived from a Gaussian plume model that computes the emission rate Q from wind speed U , length L , and the integrated mass enhancement M in the emission plume \mathcal{P} under the assumption of steady-state conditions:

$$Q = \frac{U}{L} \cdot M = \frac{U}{L} \cdot \sum_{i \in \mathcal{P}} A_i \cdot (V_i - V_{i,BG}), \quad (9)$$

where A_i is the pixels size and V_i and $V_{i,BG}$ are enhancement and background CH₄ columns of the i th pixel (in kg m⁻²). L is the length of the plume, which is obtained from the arc length of the center line from the source to the last detectable pixel. U is the effective wind speed that would be computed by weighting the wind profile with the CH₄ concentration profile (Kuhlmann et al., 2024a). Since concentration profiles are not known in our study but many sources identified are located at small vent stacks, the wind speeds were taken from the 10 m wind of the ECMWF operational analysis product at ~ 10 km resolution. The uncertainty due to these assumptions is discussed below.

While the matched filter retrieves CH₄ enhancements above the background in the AVIRIS-NG lines, locally the CH₄ background can deviate from zero, for example, due to diffuse CH₄ emissions in the area or small systematic errors in the retrieval. Therefore, the local background field $V_{i,BG}$ was computed from the pixels surrounding the detected plume using a normalized convolution with a Gaussian kernel ($\sigma = 3$ pixels), where we masked the convex hull of the detected plume dilated by a disk kernel with a radius of 3 pixels. The dilation was used to avoid overestimating the background by including pixels from the emission plume that were below the detection limit.

The integration area \mathcal{P} contains all pixels of the plume from the source location up to plume length L . We computed the integration area as the convex hull of the detected pixel mask dilated by a disk kernel (radius: 3 pixels), selecting an integration area larger than the detected plume. This avoids overestimating the integrated mass M when excluding pixels below the detection limit due to random noise inside the detectable plume. Furthermore, it includes CH₄ mass below the detection limit at the plume edges to avoid underestimating the mass M . In the along-plume direction, the integration area was limited to pixels with an along-plume coordinate between 0 and L to not include pixels upstream of the source and further downstream than L .

2.4 Estimation of uncertainty

The errors in the estimated emissions are due to errors in the retrieved CH₄ enhancements as well as errors in the emission quantification method resulting from errors in the background field, wind speed and plume length as well as methodological limitations of mass-balance approaches such as the assumption of steady-state conditions. These errors can be spatially and temporally correlated, in which case they would not reduce when, for example, integrating CH₄ pixels or averaging multiple estimates of the same source. We assume that errors in emission estimates of the same source are mostly correlated because the error is dominated by the representation error in the wind speed, i.e., the limited resolution of the analysis product, which should be mainly driven by local topography. As a consequence, we do not reduce the uncertainty of estimated emissions for multiple estimates of the same source but use the mean uncertainty instead.

2.4.1 Uncertainties in matched filter

The single sounding precision of CH₄ enhancements was computed from the AVIRIS-NG SNR by propagation of uncertainty in the matched filter. A SNR of about 400 results in a radiance uncertainty of about $12.5 \mu\text{W m}^{-2} \text{nm}^{-1} \text{sr}^{-1}$. The CH₄ uncertainty depends on radiance levels due to the scaling of the albedo correction resulting in larger values over dark surfaces and lower values over bright surfaces. The median uncertainty is about 450 ppm m, in good agreement with the standard deviation of the retrieved CH₄ field.

Errors in the CH₄ enhancements are caused by simplifications used in the matched filter. We conducted detailed sensitivity tests using synthetic spectra to quantify these errors. The unit absorption spectrum s was computed by Eq. (7) using mean values for each line, while surface elevation, solar and viewing zenith angles, aircraft altitude, and XCH₄ background can vary during data acquisition. Figure 3a–e show that the variability in these parameters during data acquisition does not have a strong impact (< 5 %) on the unit absorption spectrum and consequently the CH₄ enhancements. Note that solar zenith angle (SZA) does not vary significantly for a line and the range shown here is considering all lines in the campaign.

For a non-scattering atmosphere, surface reflectance does not impact the unit absorption spectrum. To analyze the impact of scattering, synthetic spectra were simulated with the libRadtran model (Emde et al., 2016) for varying surface reflectance and two scattering scenarios. The first scenario only includes molecular scattering setting aerosol optical depth (AOD) to zero. The second scenario uses libRadtran's default aerosol scenario with rural-type aerosols below 2 km and background aerosols above 2 km under spring–summer conditions and a visibility of 50 km. AODs were set to 0.10 in the two spectral windows (1.6 and 2.4 μm) to simulate the high aerosol load (up to 0.15 at 1.6 μm at the AERONET

station in Bucharest) due to the presence of Saharan dust in the atmosphere (well-mixed below 4–5 km based on CAMS forecast). Figure 3f shows no significant impact of molecular scattering alone, while an AOD of 0.10 results in a ratio varying between 1.10 and 0.95 for a surface reflectance ranging from 0.02 to 1.0. Since over darker surfaces (< 0.02), CH₄ single sounding precision is too high for identifying plumes, errors due to aerosols are expected to be smaller than $\pm 5 \%$ for the detected plumes during the campaign.

Further potential sources of errors are the computation of mean vector $\hat{\mu}$ and covariance matrix \hat{S} . Since $\hat{\mu}$ and \hat{S} are computed from a single radiance cube acquired along a flight line, it is necessary to have a sufficiently large number of spectra for unbiased estimates. Since the matched filter is applied for across-track position, the number can be small for short lines. Sensitivity tests with synthetic spectra showed that 7500 spectra are sufficient (i.e., < 2 % bias) for an unbiased estimate of 100 pixels with CH₄ enhancements of 5000 ppm m.

The matched filter assumes that the unit absorption depends linearly on the CH₄ enhancement, which is not the case for optical thick absorbers. The linearization results in an error in the estimated CH₄ enhancements that depends on surface reflectance and the true enhancement. The bias is generally small for enhancements measured during the campaign (< 5 %) but can get large for high surface reflectance (> 0.50) and strong enhancements (> 7000 ppm m), leading to an underestimation of CH₄ enhancements by 10 %–20 %. Other trace gases (CO₂ and H₂O) were found to have no significant impact on the unit absorption spectrum.

Other sources of errors in CH₄ retrievals are correlations between the unit absorption spectrum and the wavelength-dependent surface reflectance. These errors are non-negligible due to the relative low spectral resolution of the AVIRIS-NG instrument but are difficult to quantify (e.g., Ayasse et al., 2018). To avoid impact of surface features on the result, visual inspection of the plumes included ensuring that plumes are not correlated with surface features.

Overall, we estimate that the uncertainty in the matched filter is less than 5 % for CH₄ fields near identified plumes, taking into account, for example, the surface reflectance in the area. We expect the uncertainty for a plume to be spatially correlated. However, the correlation between sources should be small, and even for multiple observations of the same source, we expect only a weak correlation because the plumes were detected in the overlapping region of different AVIRIS-NG lines with different spatial coverage. For comparison, the single sounding precision is about 500 ppm m, i.e., 10 % for an average plume size of 20 pixels and an enhancement of 1000 ppm m.

2.4.2 Uncertainties in the IME approach

The errors in the emission quantification method result from errors in the CH₄ retrieval, the background field, wind speed

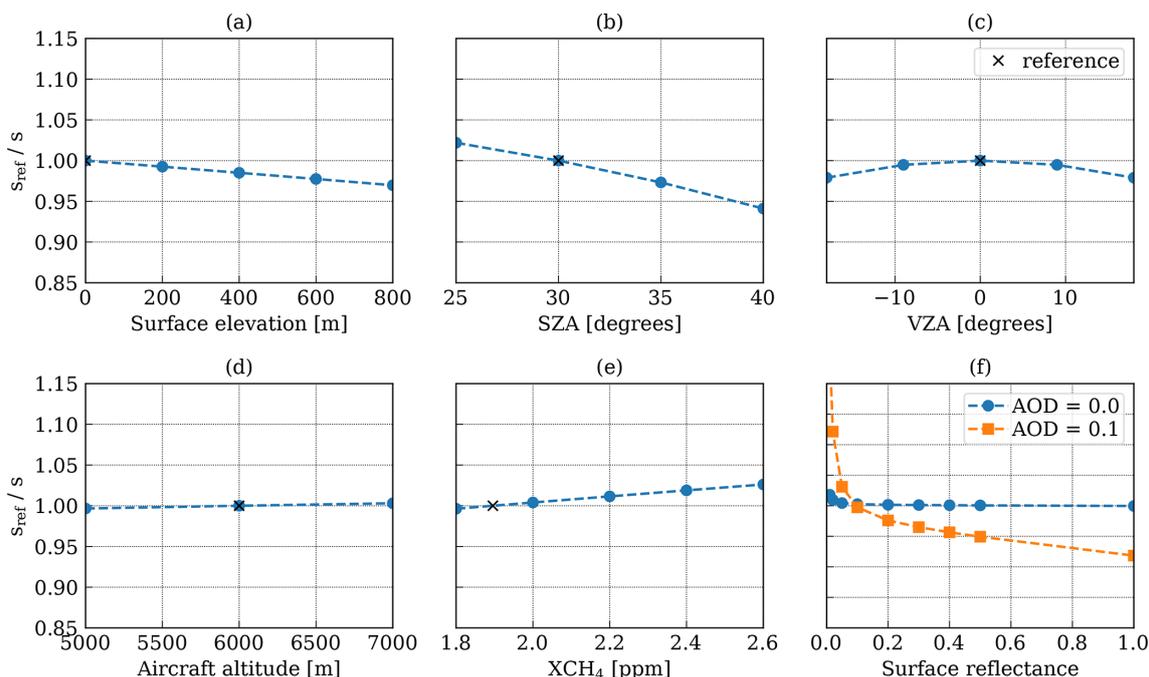


Figure 3. Sensitivity of the unit absorption spectrum s to variation in the input parameters computed by varying the parameters of the forward model (Eq. 7). The sensitivity is compared to a reference using surface elevation of 0.0 m, SZA of 30°, viewing zenith angle (VZA) of 0°, no atmospheric scattering, XCH_4 of 1.90 ppm and aircraft altitude of 6000 m. To test the impact of surface reflectance, s was simulated with libRadtran radiative transfer model using a Rayleigh atmosphere (AOD = 0.0) and aerosol scenario (AOD = 0.1). A value larger (smaller) than 1 would result in overestimated (underestimated) CH_4 enhancements.

and plume length as well as methodological limitations of mass-balance approaches such as the assumption of steady-state conditions and the assumption that the plume is within the convex hull of the detected pixels. The total uncertainty is computed from the uncertainty of M , L and U by propagation of uncertainty assuming independent variables, i.e.,

$$\sigma_Q = \frac{1}{L} \sqrt{U^2 \sigma_M^2 + M^2 \sigma_U^2 + \left(\frac{UM}{L}\right)^2 \sigma_L^2}. \quad (10)$$

The uncertainty of the plume length σ_L was set to 10 % and to at least half a pixel size (i.e., about 5 m), which is a rough estimate considering how the plume length can vary when modifying the threshold for the plume detection algorithm.

The uncertainty of the integrated mass σ_M was computed from the uncertainty of the CH_4 retrieval. For the plumes identified in the campaign, the mean uncertainty computed from the single sounding precision is 11 %, ranging from 3 % to 32 % depending on plume size. The mass M is sensitive to the size of dilation kernel used for computing the CH_4 background and for defining the integration area \mathcal{P} . To quantify the impact, we varied the size of the kernel from 1 to 5 for all detected plumes. As a result, IMEs varied between 2 % for large plumes and 49 % for small plumes. In addition, the mean and scatter of the local background is 100 ± 111 ppm m.

If the emissions were computed without subtracting the background, integrated masses would be 10 ± 15 % larger.

To account for the uncertainty in the estimated background, for the impact of surrounding pixels on the IME in the error budget, we double the variance σ_M^2 computed from the single sounding precision. To account for spatially correlated errors from the matched filter, we add an uncertainty of 5 %.

The error in the effective wind speed σ_U arises from errors in the ECMWF analysis and the effective plume height. The error in the ECMWF analysis consists of errors related to the modeling system and errors due to the interpolation from model resolution (10 km hourly) to source location and measurement time. To estimate these errors, we compare the 10 m model winds with observations from 29 meteorological stations in the study area (Meteomanz.com, 2025). The root mean square deviation (RMSD) is 1.0 m s^{-1} , which is consistent with other evaluation studies of 10 m winds reporting RMSDs of 1.0 – 2.0 m s^{-1} (Vanella et al., 2022; Potisomporn et al., 2023). We therefore use an uncertainty of 1.0 m s^{-1} . To estimate the uncertainty of the wind direction for visualization, we use the ensemble spread in the ERA-5 reanalysis (Hersbach et al., 2018).

In addition, we add 15 % uncertainty to account for the height dependency of the plume. To calculate this number, we assumed a logarithmic wind profile for a neutrally strati-

fied surface layer ($\bar{u}(z) = \frac{u_*}{\kappa} \ln(z/z_0)$) with von Karman constant ($\kappa = 0.40$), surface roughness ($z_0 = 0.10$ m) and friction wind speed u_* chosen to match the 10 m wind speed (Jacobson, 2005). For this profile, the effective wind speed for a plume at 5 and 20 m above the ground would be 15 % smaller or higher, respectively, than the 10 m wind speed.

2.5 Additional site visits

To further investigate the sources identified by AVIRIS-NG, a team with an optical gas imaging (OGI) camera carried out follow-up visits to each identified location in November 2022. The team searched the area where the plume was detected to identify the origin of the emissions. A protocol with photos and videos was provided for each site.

In 2023, additional site visits were conducted visiting the largest leaks identified in 2019. The primary goal of these site visits was to check if the leaks, mostly open ends, were fixed as stated by the operator.

2.6 Top-down estimates of total emissions from oil and gas in southern Romania

To improve the top-down estimates of the total emissions from oil and gas in southern Romania, we merge the datasets from the ground- and drone-based campaign in 2019 with the airborne AVIRIS-NG campaign in 2021. We calculate emissions for three source categories: “oil production sites”, “gas production sites” and “processing facilities”. As we found that emissions changed between 2019 and 2021, we compute emissions for 2019 and for four scenarios in 2021 that account for potential changes in emissions distribution between the 2 years.

2.6.1 Merging of datasets for calculating annual emissions

Since AVIRIS-NG only observed super-emitters above its detection limits, we used an approach similar to Sherwin et al. (2024) to calculate the total emissions per category, day and region (A–D in Fig. 1) within the AVIRIS-NG flight lines:

$$E_l = (1 - z) \cdot N_l \cdot f_{\leq DL}^{\text{emis}} \cdot \text{EF} + E_{\text{ANG}}, \quad (11)$$

where z is the zero mode, i.e., the fraction of sites with zero emissions; N_l is the number of sites in the flight lines; $f_{\leq DL}^{\text{emis}}$ is the fraction of emissions below the AVIRIS-NG detection limit; EF is the emission factor, i.e., the arithmetic mean of the emissions (in kg h^{-1}) of all sites; and E_{ANG} is the emissions retrieved from the AVIRIS-NG measurements. Note that for 2019, E_{ANG} is zero and $f_{\leq DL}^{\text{emis}}$ is one because no AVIRIS-NG campaign was flown. To compute annual emissions (in kt), we assume that the observed AVIRIS-NG emissions are constant in time or at least a representative sample of emissions throughout the year.

Since $f_{\leq DL}^{\text{emis}}$ depends on the AVIRIS-NG detection limit, which depends on wind speed, and wind speed varied between regions and days, it is necessary to calculate emissions separately by region and day, and we thus obtained emission estimates for each region with two estimates for region C as measurements were conducted on both days. Total emissions within the AVIRIS-NG flight lines are computed by averaging the two estimates for region C and adding up the estimates per region.

To compute the total emissions in the study area E_b , we scale the emissions inside the flight lines with the total number of sites in region N_b :

$$E_b = \frac{N_b}{N_l} \cdot E_l. \quad (12)$$

The number of sites in the study area was provided by the operators (Table S3). The zero mode for oil production sites was determined as 0.25 ± 0.10 from ground-based measurements in 2019. The uncertainty of the zero mode was estimated from the sensitivity tests conducted by Stavropoulou et al. (2023). The emission factor EF was computed from the emission distribution using different scenarios, and the emission fraction $f_{\leq DL}^{\text{emis}}$ was computed from the AVIRIS-NG detection limits and the emission distributions.

2.6.2 AVIRIS-NG detection limits

The AVIRIS-NG detection limit for CH_4 plumes depends on pixel size, wind speed and uncertainty in the retrieved CH_4 maps. In our study, CH_4 emission plumes were detected by visual inspection. The smallest plumes identified were about 10 m wide and 30–50 m long for a pixel size of about 5 m. These small plumes were generally compact and did not show visible dispersion in the across-plume direction (e.g., Fig. S4 in the Supplement). To model such plumes, we simulated plumes using a Gaussian plume model with a fixed standard width of 5 m and a random uncertainty of 500 ppm m.

Figure S1 shows these plumes for varying source strength and wind speed. We compute the noise-free peak signal-to-noise ratio (PSNR) for each plume. Comparing the synthetic plumes with the plumes identified in the AVIRIS-NG lines, we estimate that a PSNR of about 2.0 is a good threshold for the detection limit. For the non-dispersive Gaussian plume model given above, the detection limit can be computed as a function of wind speed from

$$c_{\text{peak}} = \frac{Q_{\text{DL}}}{\sqrt{2\pi}\sigma_y U} = 2\sigma_{\text{CH}_4}, \quad (13)$$

which gives

$$Q_{\text{DL}} = 2\sqrt{2\pi}\sigma_y\sigma_{\text{CH}_4}U \approx 30.6 \frac{\text{kg h}^{-1}}{\text{m s}^{-1}} \cdot U. \quad (14)$$

The uncertainty of the detection limit was computed from the variability in the wind speed in the study area during data acquisition. The wind speed and its variability were computed

as the mean and standard deviation of the 10 m wind speeds from the ECMWF analysis in the region during measurement time (Table S2).

2.6.3 Scenarios for changes in the emission distribution for oil production sites

For the campaign in 2019, Stavropoulou et al. (2023) determined emission distributions for oil production sites using probability density functions that follow log-normal distributions: a mean distribution and two extremes corresponding to a lower and upper limit of the estimated emissions (95 % CI – confidence interval). The distributions were fitted as a normal distribution with mean μ and standard width σ by taking the natural logarithm of the CH_4 emissions. The emission factor is calculated as $\text{EF} = \exp(\mu + 0.5\sigma^2)$ (Stavropoulou et al., 2023).

Since the number of super-emitters identified with AVIRIS-NG was lower in 2021 than expected from these distributions (see results), we considered four scenarios of how the distribution may have changed between 2019 and 2021: Scenario 1 uses the mean distribution from 2019 and Scenario 2 uses the distribution from 2019 that corresponds to the lower limit of estimated emissions. Scenarios 1 and 2 assume that the emission distribution is still valid below the AVIRIS-NG detection limit, meaning any emission reduction results from fewer than expected AVIRIS-NG detections. Scenarios 3 and 4 assume that emissions also changed below the AVIRIS-NG detection limit. To account for this we reduce the standard width σ for Scenario 3 and mean μ for Scenario 4 of the 2019 distribution such that the number of expected emitters above the detection limit matches the six emitters found with AVIRIS-NG. Scenario 3 reduces the width of the emission distribution, meaning there are fewer low- and high-emitting sources. Scenario 4 shifts the whole emission distribution assuming all emissions were reduced equally instead of addressing only the largest sources. The emission distributions for the four scenarios are shown Fig. 4.

2.6.4 Emissions from gas production sites and processing facilities

Stavropoulou et al. (2023) only provided the emission distribution and the emission factors for oil production sites but did not provide an estimate for gas production sites and processing facilities, as the sample size was smaller. To estimate the emissions from gas production sites and processing facilities, we assume that the emission distributions for the four scenarios are also applicable for other source categories. In practice, this means that the fraction of emissions below the detection limit $f_{\leq \text{DL}}^{\text{emis}}$ is the same for all source categories. The emission factors for gas production sites and processing facilities were calculated as the arithmetic mean of emissions measured during the 2019 campaign. These measurements are taken from Table S16 in Stavropoulou et al. (2023), ex-

cluding the estimates using the BDL (below detection limit) method.

The emission factors are $11.2 \pm 4.0 \text{ kg h}^{-1}$ for 31 gas production sites and 13.0 ± 3.0 for 60 kg h^{-1} processing facilities. The 1σ uncertainty was estimated using bootstrapping. These emission factors were used for Scenario 1, while for the other scenarios, the emission factors were scaled considering the change in the emission factor for oil production sites for the four scenarios (i.e., 5.4, 3.6, 2.3 and 1.2 kg h^{-1}).

2.6.5 Estimation of uncertainty

Monte Carlo simulations were used to compute the median and the 95 % confidence intervals (CIs) for the annual emissions estimated by Eq. (11). For each region and day, we created an ensemble with 5000 members using normally distributed errors for zero mode, the emission factors and the AVIRIS-NG measurements. The fraction of emissions below the AVIRIS-NG detection limit was computed from an ensemble of detection limits considering the spatial and temporal variability in wind speed in the region. To calculate the AVIRIS-NG emissions and their uncertainties, we sum the mean estimates per source assuming uncorrelated errors, as the uncertainty is dominated by the uncertainty in the wind speed, for which no bias was found. The uncertainty in the change in the emission distribution is accounted for by the four scenarios.

2.7 Bottom-up estimates of total emissions from oil and gas in the entirety of Romania

Bottom-up reports of oil and gas from energy production in the entirety of Romania were taken from the UNFCCC submission (Table 1.B.2 in UNFCCC, 2023b). We assumed emission categories in Table 1.B.2.a with a description “oil produced” to represent emissions from oil production sites and categories in Table 1.B.2.b with a description “gas produced” to represent emissions from gas production sites. Emissions from venting and flaring correspond to Table 1.B.2.c.

The IEA’s estimates for Romanian emissions in 2021 are the sum of categories including “Onshore Oil”, “Onshore Gas” and “Gas pipelines and LNG [liquefied natural gas] facilities” (IEA, 2021). We group fugitive emissions from onshore oil and gas to oil and gas production sites, respectively. All other emissions are included in “venting and flaring”.

3 Results

3.1 Methane super-emitters identified with AVIRIS-NG

In total, we identified 35 emission plumes at 25 locations, with some sources being observed up to three times due to overlapping flight lines. Maps and photos for all sources are provided in the Supplement. The sources were located at oil

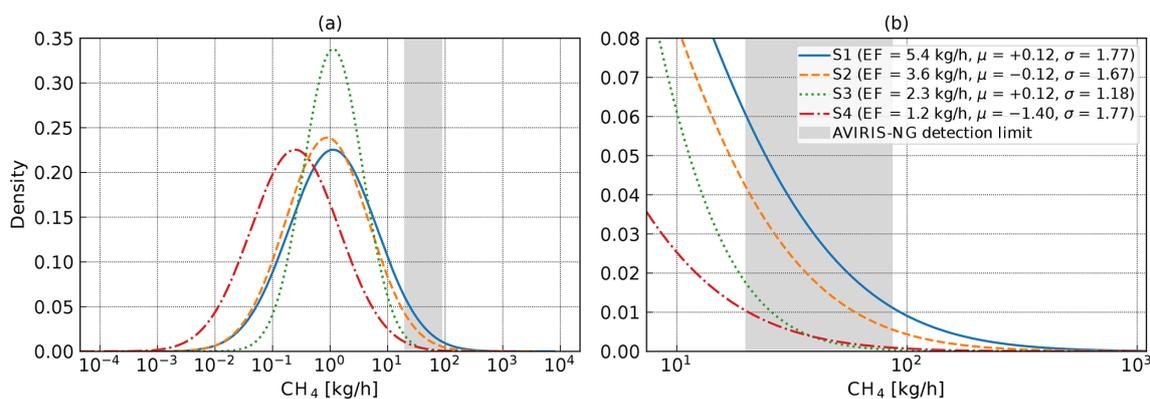


Figure 4. The emission distribution of oil production sites for the four scenarios. The AVIRIS-NG detection limit is shown in gray based on varying wind speed during the campaign. Panel (b) presents a zoomed-in view for the long tail of the distribution.

production sites (6 sources); at processing facilities (7); and, unexpectedly, in the open field (12), i.e., not directly linked to O&G infrastructure visible in aerial images. Closer inspection of the sites during the site visits and from images using Google Street View and Google Earth suggests that at least six of the open-field sources are linked to O&G processing facilities in the proximity.

Figure 5 and Table S1 show the estimated emission rates for all 25 sources. The source strengths ranged from 16 to 501 kg h^{-1} with a median of 80 kg h^{-1} (Fig. 1b). The total measured emissions were 2975 kg h^{-1} with 16 % from oil production sites, 30 % from the processing facilities and 54 % from sources located in the open field. The OGI team successfully identified emissions from 14 sources (5 at oil production sites, 6 at processing facilities and 3 in the open field linked to facilities), indicating that these sites continued to emit over a period of at least 1 more year. The mean uncertainty of the estimated emission rate is 84 %, ranging from 40 % to 241 %. Our total uncertainty in the estimated emissions is dominated by the uncertainty in the wind speed, which has a mean uncertainty of 76 % (32 %–239 %). The mean uncertainty in the integrated mass, i.e., the uncertainty in the CH_4 retrieval, is 25 % (9 %–64 %).

Figure 6 shows four examples for sources identified in the campaign. For each source, we show an aerial image, the AVIRIS-NG CH_4 map with inferred emissions and a site photograph. Figure 6a shows a CH_4 plume at an oil production site in the northeastern part of the study area. The plume shape (gray dots) is consistent with the westerly wind from the ECMWF analysis product indicated by the arrow. The emission rate was $81 \pm 35 \text{ kg h}^{-1}$. The plume was also observed 1 h earlier in an overlapping flight line with a consistent emission rate ($69 \pm 45 \text{ kg h}^{-1}$). The site was visited by the OGI team on 22 November 2022. They confirmed the presence of a large CH_4 leak originating from an open-ended line as indicated in the photo. An emission plume at an oil processing facility is shown in Fig. 6b. The estimated source strength was $90 \pm 48 \text{ kg h}^{-1}$. The OGI team identified emis-

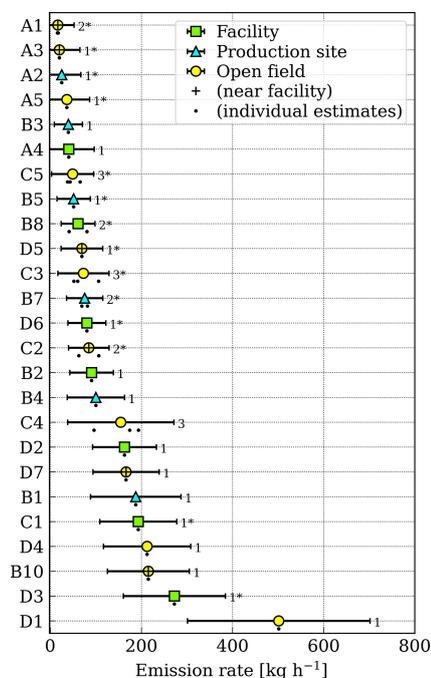


Figure 5. Methane super-emitter identified during the airborne campaign with emission rates of the sources with uncertainty (1σ) and the number of detections. The asterisk marks sources that were also detected by follow-up site visits in 2022.

sions from a vent on the roof of the building on 22 November 2022. Source B10 is located about 100 m north of an oil processing facility in the open field (Fig. 6c). The emissions were quantified at $215 \pm 90 \text{ kg h}^{-1}$. Investigation by the OGI team on 22 November 2021 showed strong emissions originating from a vent stack installed in the field.

Finally, Source C2 is also located in the open field about 50 m west of a processing facility. The plume was detected on both 29 and 30 July 2021. The site was visited on 24 November 2022, but no source was found. We used the history of



Figure 6. Four examples of CH_4 plumes at (a) an oil production site, (b) a processing facility and (c, d) from vent stacks in the open field observed in 2021. The panels from top to bottom show ©Google Earth imagery, AVIRIS-NG CH_4 maps with estimated emissions Q and site photos with a yellow arrow indicating the emission point. M , U and L are integrated mass, wind speed and plume length. All uncertainties are 1σ . The maps show the detected plumes as dots and the plume center curve as line. The arrow in the top-right corner shows the wind direction from IFS analysis including the spread (2σ) from the ERA-5 reanalysis as shaded area.

Google Street View and Google Earth aerial images to reconstruct the evolution of this facility. On images taken between July 2012 and May 2021, a vent stack in the open field can be seen, which was no longer present in July 2022. Aerial images from February 2021 show heavy construction work on the site, including the installation of a gas flare, which was completed by November 2021. It is likely that the vent stack was dismantled during the same months but after the AVIRIS-NG flight in July. Consequently, the source could not be located during the site visit in November 2022.

3.2 Number of super-emitters in 2019 and 2021

AVIRIS-NG can only measure emissions above its detection limit, which varied between 15 and 107 kg h^{-1} during data acquisition due to varying wind speeds on the campaign days (Eq. 14). Table S6 compares the number of super-emitters expected at oil production sites in each region with the ac-

tual number of AVIRIS-NG detections, considering the spatial coverage of the flight campaign. For the mean distribution obtained for 2019 (Scenario 1), we would expect to find between 30 and 190 (95 % CI) oil production sites with emissions above the AVIRIS-NG detection limits. However, AVIRIS-NG only detected emissions at six oil production sites, which is also a smaller number than the 15 to 142 emitters expected from the distribution corresponding to the lower limit of estimated emissions in 2019 (Scenario 2). The number of high-emitting oil production sites was thus significantly lower in 2021, which suggests that the emission distribution changed between 2019 and 2021. For Scenarios 3 and 4, we adjusted the emission distribution to match the expected number of super-emitters with the detection in 2021.

3.3 Total emission in 2019 and 2021

Figure 7a shows our estimates of total emission of CH₄ from oil and gas in 2019 and 2021 in southern Romania. Detailed numbers with intermediate steps are shown in the Supplement.

For 2019, Stavropoulou et al. (2023) estimated emissions of 120 kt (79–180 kt) from oil production sites. In addition, we estimated 28 kt (8–52 kt) from gas production sites and 75 kt (38–119 kt) from processing facilities in 2019. In 2019, the total emissions from O&G infrastructure were therefore 224 kt (157–295 kt for 95 % CI). This estimate from ground- and drone-based measurements agrees well with the independent estimate from airborne in situ measurements, which is about 227 ± 87 kt for 2019 (Maazallahi et al., 2025).

We estimate that emissions in the study area were 20 %–60 % lower in 2021 compared to 2019 with estimates of 175 kt (142–209 kt) in Scenario 1 and 76 kt (58–94 kt) in Scenario 4. Depending on the scenario, super-emitters from AVIRIS-NG account for only 5 %–14 % of total emissions at O&G production sites but are a major contributor at processing facilities (28 %–56 %). Consequently, emission reductions are greater at O&G production sites (35 %–75 %) than at processing facilities (10 %–50 %).

To quantify the 95 % CIs associated with the expected number of super-emitters and the measurement-based emission estimates, we conducted extensive Monte Carlo simulations, accounting for the uncertainty in the AVIRIS-NG measurements, the AVIRIS-NG detection limits, the emission factors and the fraction of non-emitting sites. The results of this analysis show that the observed differences between 2019 and 2021 are greater than what can be attributed to the measurement and methodology uncertainties (95 % CI), leading us to conclude that the emission reductions from 2019 to 2021 were significant.

Despite these reductions, the actual methane emissions from oil and gas in the study area in 2021 still exceed the reported emissions for the entirety of Romania (Fig. 7b). The national inventory report of Romania states that emissions from O&G production were 71 kt in 2019 (not shown) and 66 kt in 2021 split between oil production (14 %), gas production (32 %), and venting and flaring (54 % in 2021) (Table 1.B.2 in UNFCCC Submission 2023 v2). Figure 7b also shows emissions estimated by the IEA, which include onshore oil and gas, pipelines, and LNG facilities. They are 14 kt higher than the emissions reported to the UNFCCC but still at the lower end of the emissions determined from the measurement campaigns in the study area alone.

4 Discussions

The quantification of CH₄ emissions from oil and gas is challenging for both bottom-up and top-down approaches due to the limited amount of information available.

The quantification of emissions using the IME method can result in systematic uncertainties from the assumptions made in the implementation. The two largest uncertainties are the effective wind speed and the background field. The effective plume height will generally be higher than the source location due to plume rise and vertical mixing. In this study, we assume a plume height between 5 and 20 m because 11 out of 25 super-emitters were elevated, mainly, at vent stacks, which are about 5 m high. The remaining source locations are closer to the ground (five) or unknown (nine). To estimate the background, we decided to subtract the background field to obtain the local enhancement. The motivation here is that the matched filter retrieves the CH₄ enhancement above the AVIRIS-NG line. It is possible that near potential sources, local background concentrations are increased. However, these diffuse emissions cannot be accounted for using the IME method. If we do not subtract the local background, AVIRIS-NG emissions would be 10 ± 15 % higher, which is within the uncertainty budget assigned to AVIRIS-NG in this study.

The strongest evidence for a change in emissions from 2019 to 2021 is the lower number of super-emitters detected by AVIRIS-NG at oil production sites. The number depends on the AVIRIS-NG detection limit. Figure S2 shows the sources identified with AVIRIS-NG as a function of wind speed together with the detection limit and its uncertainty from the variability in wind speed. The emission rates are mostly at or above the detection limit. An exception are sources that were already identified in another flight line at lower wind speed (i.e., B7b, C2b and C5c). We therefore conclude that the detection limit is a robust estimate. Another reason for the lower number of super-emitters might be seasonal variability in emissions because the campaigns in 2019 and 2021 were conducted in July and October, respectively. Data to determine the influence of seasonal effects are unfortunately not available. However, Varon et al. (2023) estimate week-to-week variability for CH₄ emissions in the Permian Basin to be about 25 %, which, if we assume similar variability in Romania, would be too small to explain the difference between 2019 and 2021 in Romania.

The most likely explanation for the reductions from 2019 to 2021 is that operators implemented improvements in the production infrastructure after the 2019 campaign. The operators were informed about the locations of the highest-emitting sites and the likely origin of the emissions (mainly open-ended lines) already in 2020, well before the flights in 2021. During the 2-year gap between the campaigns, operators therefore had the opportunity to implement emission mitigation measures. In fact, the operators have communicated to us that the infrastructure has been upgraded and leaks have been significantly reduced. The additional site visits in 2023 also found that several open-ended lines detected in 2019 had indeed been sealed.

The four scenarios employed to modify the emission distribution reflect different possible effects of emission mitigation: the most pessimistic is Scenario 1, which assumes

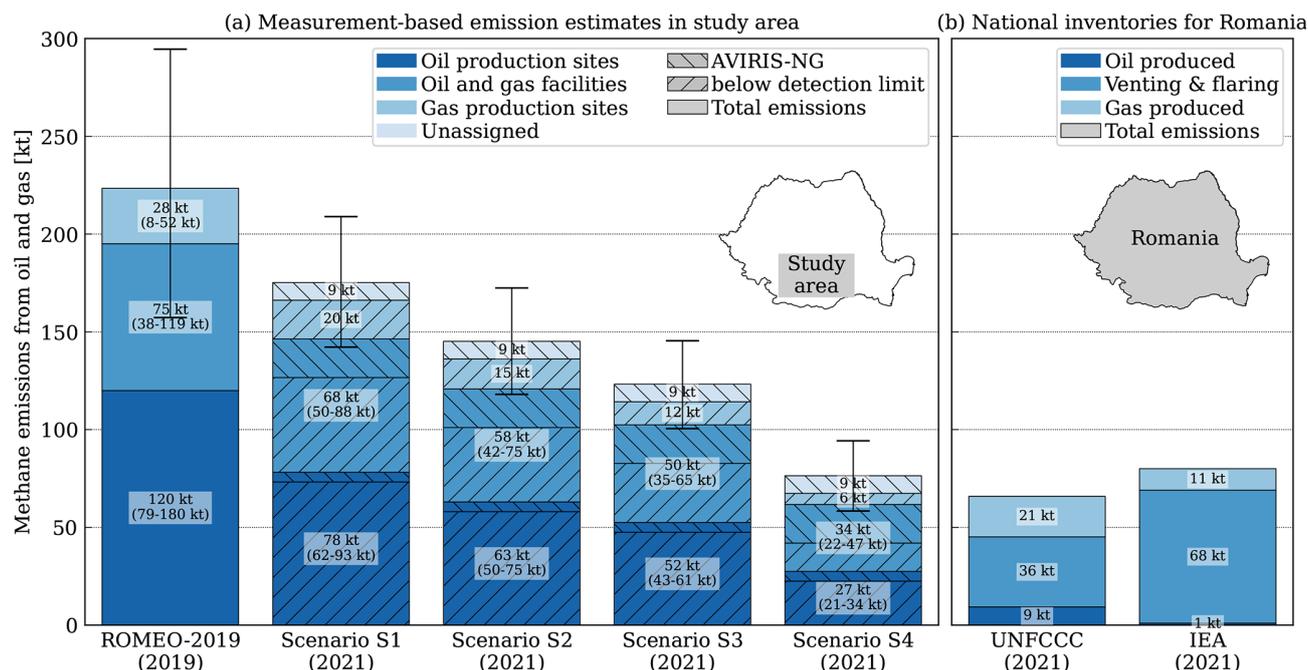


Figure 7. Comparison of top-down CH_4 emission estimates from O&G infrastructure in the study region with emissions reported to UNFCCC (Table 1.B.2 originating from oil and gas production, UNFCCC, 2023b; IEA estimates for oil and gas production including pipelines and LNG facilities in the entirety of Romania, IEA, 2021). Top-down estimates are provided for 2019 and, for the four scenarios described in the text, for 2021. Error bars show the 95 % confidence interval. “Unassigned” sources are open-field sources that have not been assigned to a processing facility.

that only the emissions from the largest sources were curbed, while those below the detection limit of AVIRIS-NG remained unaffected. The most optimistic is Scenario 4, which assumes that all emissions were reduced equally. This is quite unlikely given the considerable effort of controlling all production sites rather than addressing only the largest leaks. Unfortunately, direct emission measurements below the AVIRIS-NG detection limits are lacking in 2021 to constrain these scenarios, but they encompass the full range of potential mitigation efforts and were supported by site visits in subsequent years.

The situation at the processing facilities is complex. The number of super-emitters observed in 2021 is lower than expected from the measurements conducted in 2019. However, the uncertainty in this estimate is large due to the limited sample size. We have strong evidence that at least one super-emitter (C2) was closed after the AVIRIS-NG flights, and it is possible that others were addressed already before then. A complication is the possibility that mitigation measures at the production sites (e.g., sealing of open-ended lines) might increase the emissions from venting in the surroundings, which, according to our study, was the largest source of emissions at and near processing facilities. In 2021, we identified six “open-field” sources that we assigned to nearby processing facilities. Four of these were identified during the 2022 survey as vent stacks, which were already visible in

Google imagery 10 years earlier (A1, A3, C2 and B10). Due to the lack of information regarding the location and number of these stacks in the dataset provided by the operator, these sources were not visited during the ground-based campaign in 2019. Since venting is generally associated with high emissions, the emission factors estimated from the 2019 measurements might be biased low. Additional measurement campaigns would be necessary to better constrain emissions in Romania. While we see some evidence for mitigation at processing facilities, the estimated emissions remain highly uncertain.

Our top-down estimates in the study area still exceed emissions reported to the UNFCCC for the entirety of Romania. In particular, we find that reported emissions from oil production (Fig. 7b) are substantially lower than our measurement-based estimates. In contrast, reported emissions from gas production are higher than top-down estimates, likely because most gas production occurs outside our study area in northern Romania. In the reported emission, CH_4 emissions from venting and flaring are mainly due to oil production. These emissions can occur at oil production sites or at processing facilities. If all venting and flaring would occur at production sites, the reported emissions would be consistent with our measurement-based estimates (Scenarios 3 and 4). However, while the ground-based survey in 2019 found venting primarily at production sites, the small number

of super-emitters at production sites and the high emissions from vent stacks in 2021 indicate that venting shifted towards processing facilities. In this case, top-down estimates at oil production sites would be substantially greater than reported emissions, suggesting that the emission factors used to compile the reports are too low. However, this assumes that the diurnal variability in emissions is negligible, while studies in other regions show evidence for a diurnal cycle in emissions (e.g., Vaughn et al., 2018). Our measurements were conducted exclusively during daytime, so additional measurements would be necessary to constrain the diurnal cycle of emissions.

5 Conclusions

Our study highlights the importance of monitoring emissions using measurement-based techniques to improve emission estimates, identify their main causes and monitor the effectiveness of mitigation measures. It also demonstrates the need to combine large-scale mapping surveys using instruments like AVIRIS-NG with ground-based surveys to cover the full range of sources from small leaks to super-emitters. Despite the high detection limit, AVIRIS-NG directly measured in only 2 d CH₄ emissions of 2964 kg h⁻¹ (i.e., 26 kt), which is 15 %–34 % of our bottom-up estimates for the basin and 40 % of the reported emissions for the entirety of Romania. The emissions originated at only 25 locations, thus providing targets for effective and efficient future reductions in emissions in Romania. Additional ground-based measurements and airborne campaigns, e.g., with the new AVIRIS-4 instruments with better detection limits (Green et al., 2022), would be needed to ensure continuous monitoring of emission reduction measures in the future.

Code and data availability. The codes for CH₄ retrieval and up-scaling are available on request. The ddeq Python library for data-driven emission quantification (Kuhlmann et al., 2024a) is available on Gitlab.com (<https://gitlab.com/empa503/remote-sensing/ddeq>, last access: 13 May 2025). AVIRIS-NG Level 1 data are available on the AVIRIS-NG data portal (<https://avirisng.jpl.nasa.gov/dataportal/>, Jet Propulsion Laboratory, 2025). The data used for up-scaling are available in the Supplement. The AVIRIS-NG CH₄ maps and the results from the emission quantification are published on the Zenodo data repository (<https://doi.org/10.5281/zenodo.14974368>, Kuhlmann et al., 2024b).

Supplement. The supplement related to this article is available online at <https://doi.org/10.5194/acp-25-5371-2025-supplement>.

Author contributions. GK developed and applied the methane retrieval and the emission quantification methods. GK, FS and DZA developed the methods for merging and upscaling the 2019 and 2021 datasets. StS, AT and AH collected and prepared the raw data used in the study. The study was conceptualized by GK, AC, TR and DB and supervised by TR and DB. Funding was acquired by StS, LE, AC, TR and DB. The paper was prepared and revised by GK with input from all co-authors.

Competing interests. The contact author has declared that none of the authors has any competing interests.

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