



Western, L. M., Ramsden, A., Ganesan, A. L., Boesch, H., Parker, R. J., Scarpelli, T. R., Tunnicliffe, R. L., & Rigby, M. L. (2021). Estimates of North African Methane Emissions from 2010 to 2017 Using GOSAT Observations. *Environmental Science and Technology Letters*, 8(8), 626-632. <https://doi.org/10.1021/acs.estlett.1c00327>

Peer reviewed version

Link to published version (if available):
[10.1021/acs.estlett.1c00327](https://doi.org/10.1021/acs.estlett.1c00327)

[Link to publication record in Explore Bristol Research](#)
PDF-document

This is the author accepted manuscript (AAM). The final published version (version of record) is available online via American Chemical Society at <https://doi.org/10.1021/acs.estlett.1c00327>. Please refer to any applicable terms of use of the publisher.

University of Bristol - Explore Bristol Research

General rights

This document is made available in accordance with publisher policies. Please cite only the published version using the reference above. Full terms of use are available: <http://www.bristol.ac.uk/red/research-policy/pure/user-guides/ebr-terms/>

Estimates of North African methane emissions from 2010-2017 using GOSAT observations

Luke M. Western,^{*,†} Alice E. Ramsden,[‡] Anita L. Ganesan,[‡] Hartmut Boesch,^{¶,§}
Robert J. Parker,^{¶,§} Tia R. Scarpelli,^{||} Rachel L. Tunnicliffe,[†] and Matthew
Rigby[†]

[†]*School of Chemistry, University of Bristol, Bristol, BS8 1TS, UK*

[‡]*School of Geographical Sciences, University of Bristol, Bristol, BS8 1SS, UK*

[¶]*National Centre for Earth Observation, University of Leicester, Leicester, LE1 7RH, UK*

[§]*Earth Observation Science, School of Physics and Astronomy, University of Leicester,
Leicester, LE1 7RH, UK*

^{||}*Earth and Planetary Sciences, Harvard University, Cambridge, 02138, USA*

E-mail: luke.western@bristol.ac.uk

Abstract

Source characteristics of methane emissions in Africa are not well understood, despite methane's role as the second largest anthropogenic contributor to climate change. Here, we present monthly methane emission estimates from Algeria, Egypt, Libya, Morocco and Tunisia between 2010-2017, a region dominated by anthropogenic emissions. Emissions are estimated using observations from the GOSAT satellite and a Markov chain Monte Carlo inverse algorithm. Our top-down North African methane emissions are generally in line with inventory estimates and national reporting to the United Nations Framework Convention on Climate Change (UNFCCC). An exception is that summertime emissions from the Nile Delta region are considerably higher those that

11 predicted by inventory estimates, possibly due to agricultural practices and the influ-
12 ence of the Nile.

13 **Introduction**

14 Global atmospheric concentrations methane have been rising since a hiatus was observed
15 between the early 2000s and 2007¹. The reason for this pause and renewed growth is poorly
16 understood; previous studies have implicated the main drivers of change to be an increase in
17 anthropogenic emissions^{2,3}, natural sources⁴, reduced biomass burning and rising fossil fuel
18 emissions³ or a potential change in the main sink of methane, the global concentrations of
19 atmospheric OH radicals^{5,6}.

20 With very few ground-based measurements of methane on the African continent, there
21 is uncertainty about the potential role that African methane sources have played in recent
22 growth. There is evidence that methane emissions from tropical Africa could explain around
23 a third of a global emissions increase, primarily due to a rise in emissions from the Sudd in
24 South Sudan^{7,8}. This paper presents estimates of methane emissions between 2010-2017 from
25 North Africa, here defined as Algeria, Egypt, Libya, Morocco (including Western Sahara)
26 and Tunisia.

27 In this region, fugitive emissions from oil and natural gas production dominate and ac-
28 count for around half of the total bottom-up estimated emissions⁹. Between 2010-2017 North
29 Africa produced between 2.3 - 4.3 % of global petroleum and ~5 % of natural gas¹⁰. See the
30 supporting information for a further breakdown of North Africa's fossil fuel production.

31 As non-Annex I countries under the United Nations Framework Convention on Climate
32 Change (UNFCCC), these nations are not required to submit annual greenhouse gas reports.
33 However, all countries, except Libya, have submitted anthropogenic emissions estimates
34 through National Communications to the UNFCCC, although these emission estimates may
35 have high uncertainties. Algeria has estimated its methane emissions for 2000 (i.e. before

36 this period of study) as 1.58 Tg¹¹; Egypt has estimated its 2015 emissions as 1.98 Tg¹²;
37 Morocco as 0.51 Tg in 2010, 0.54 Tg in 2012, 0.55 Tg in 2014 and 0.59 Tg in 2016¹³; and
38 Tunisia as 0.28 Tg in 2010 and 0.29 Tg in 2012¹⁴.

39 Given the paucity of ground-based methane measurements in North Africa, we have
40 used observations of column-average methane concentrations from the GOSAT satellite,
41 combined with the atmospheric transport model NAME (Numerical Atmospheric dispersion
42 Modelling Environment)¹⁵ to infer methane emissions. Below, we outline the emissions
43 inference framework, present our result for each country, and discuss implications for the
44 methane budget of this region.

45 **Materials and Methods**

46 **GOSAT Observations**

47 We use observations of dry air column-average methane concentrations to infer methane
48 emissions in North Africa. These measurements are made from the Thermal And Near-
49 infrared Sensor for carbon Observation Fourier Transform Spectrometer (TANSO-FTS) on
50 board the GOSAT satellite¹⁶. The product comes from the University of Leicester GOSAT-
51 OPCR v7.2 proxy dry air column-average methane retrieval^{17,18}, XCH_4_{proxy} . The XCH_4_{proxy}
52 retrieval uses total column values of methane (XCH_4) and carbon dioxide (XCO_2) using
53 spectral windows at 1.65 and 1.61 μm for XCH_4 and XCO_2 respectively. The ratio of XCH_4
54 to XCO_2 is multiplied by an model ensemble-derived estimate of column-average carbon
55 dioxide mole fraction to estimate XCH_4_{proxy} . This method produces robust retrievals in the
56 presence of clouds and aerosols due to their common influence on XCH_4 and XCO_2 , however
57 may be subject to larger uncertainties in regions where CO_2 measurements are not prevalent.
58 We limit the XCH_4 observations to the spatial extent of -15° W to 38° E and 15° N to 34° N
59 and bin the observations to the resolution of the output of the atmospheric transport model
60 used in the inversion (NAME transport model section, 0.352° by 0.234°). Between April 2010

61 and the end of 2017, there is a mean of 1119 observations per month, with little seasonality
62 in observation coverage over this region (see Figure S4). Figure S5 shows the binned average
63 GOSAT XCH₄_{proxy} observations for November 2011 - April 2012 and May 2012 - October
64 2012.

65 **NAME transport model**

66 We use the approach outlined in Ganesan et al.¹⁹ and Tunnicliffe et al.²⁰ to relate XCH₄
67 observations to surface emissions. The sensitivity of the XCH₄_{proxy} measurements to surface
68 emissions are derived using the Lagrangian particle dispersion model NAME¹⁵ run in back-
69 ward mode. The sensitivities derived from the NAME model output have a resolution of
70 0.352° by 0.234° and we employ a simulation domain of -50 to 87 °E by -15 to 41 °N degrees
71 (Figure S6) to a height of 20 km.

72 Meteorological fields from the Met Office Unified Model model²¹ drive transport within
73 NAME, and have a temporal resolution of three hours and a spatial resolution that increases
74 throughout the study period from 0.352 by 0.234° in 2010 to 0.141 by 0.094° in 2017. We
75 run NAME for each of the 20 vertical levels defined within the XCH₄_{proxy} product. Parti-
76 cles are released from each level and the interaction with the surface (below 40 magl) and
77 boundaries are recorded²⁰. We combine the sensitivities from the vertical layers into a single
78 sensitivity of the XCH₄_{proxy} observation to emissions by weighting each level according to
79 the corresponding GOSAT averaging kernel and pressure weight from the retrieval^{19,22}. The
80 boundary sensitivities are combined into four scaling parameters each month in the inver-
81 sion, which uniformly scales the a priori boundary condition curtains on each horizontal
82 boundary.

83 We assume that the modelling error is stochastic with an uncertain standard deviation,
84 where the standard deviation is given a prior 95% uncertainty of 5.5-14.7 ppb.

85 **A priori methane emissions**

86 We base our a priori estimates of methane emissions on various sources, regridded to that of
87 the output of the transport model (NAME transport model section). Fugitive emissions from
88 oil, coal and gas, the dominant a priori emissions source in North Africa, are from Scarpelli
89 et al.⁹ and are a static annual climatology. The GFED v4.1 database²³ provides the emissions
90 for natural and anthropogenic biomass burning. Emissions from wetlands come from the
91 mean of the ensembles of the WetCHARTs database²⁴, where values for 2015 are repeated
92 for 2016 and 2017. Any emissions from rice paddies come from a monthly inventory for the
93 year 2000²⁵. Other anthropogenic emissions, such as enteric fermentation, landfills, road
94 transport, shipping and manufacturing, come from the EDGAR v.4.3.2 emissions inventory
95 by sector for 2012²⁶. We assume all other emissions (e.g. termites and geological sources) to
96 be negligible. Inventory estimates for Morocco are generally in line with emissions reported
97 to the UNFCCC. Egypt's 2015 reporting is around 50% higher than the a priori emissions,
98 and Tunisia's reporting is around 30-35% higher.

99 The a priori emissions for the region are 5.0 Tg year⁻¹ between 2010 - 2017 (5.1 Tg year⁻¹
100 in 2015). We treat these emissions as very uncertain, and assign a 95% uncertainty range for
101 the a priori emissions of 2.7-12.8 Tg year⁻¹ in 2010 to 2.7-13.0 Tg year⁻¹ in 2017, following
102 a log-normal distribution, where the mode of the distribution is the a priori emissions. This
103 distribution has been chosen as it approximately follows the geometric standard deviation
104 uncertainty (for a lognormal 68 % uncertainty) for Annex I countries in Scarpelli et al.⁹, the
105 dominant emissions source.

106 **Boundary conditions**

107 As we infer emissions in a limited domain, the sensitivity of the methane concentrations
108 to contributions from the domain edges must be quantified. The a priori boundary condi-
109 tion mole fraction curtains at the domain edges come from the ECMWF CAMS reanalysis
110 database (which has not been constrained using GOSAT data)²⁷. We take the a priori

111 estimate each month to be the mean state at the NAME domain edges. The derived emis-
 112 sions are mostly insensitive to the choice of a priori mole fraction at the boundary (see the
 113 supporting information).

114 Inverse Method

115 Here we represent the problem of emissions and boundary condition inference as a statistical
 116 model. We denote the $\text{XCH}_4^{\text{proxy}}$ observations (see section GOSAT Observations) as the
 117 vector, \mathbf{y} , which can be modelled by the linear forward model

$$\mathbf{y} = \mathbf{H}\mathbf{x} + \mathbf{K}\mathbf{u} + \epsilon, \quad (1)$$

118 where \mathbf{H} is the sensitivity matrix to emissions at the surface and \mathbf{K} is the sensitivity to
 119 the boundaries, produced using NAME (section NAME transport model) with each row
 120 multiplied by the a priori estimates from sections and for \mathbf{H} and \mathbf{K} respectively, \mathbf{x} contains
 121 a vector which scales the a priori emissions by some factor, \mathbf{u} is as \mathbf{x} for the boundary
 122 conditions, and ϵ is the stochastic model-measurement error. We assume that all observations
 123 in \mathbf{y} are independent and identically distributed with a known stochastic measurement error
 124 σ_{obs} and unknown model error σ_{mod} , which combine as $\sigma_y = \sqrt{\sigma_{\text{obs}}^2 + \sigma_{\text{mod}}^2}$ (in ppb), where
 125 the resulting covariance matrix is $\mathbf{R} = \mathbf{I}\sigma_y^2$, and \mathbf{I} is the identity matrix. We follow a typical
 126 Bayesian framework,

$$p(\mathbf{x}, \mathbf{u}, \sigma_{\text{mod}} \mid \mathbf{y}) \propto p(\mathbf{y} \mid \mathbf{x}, \mathbf{u}, \sigma_{\text{mod}}) p(\mathbf{x}, \mathbf{u}, \sigma_{\text{mod}}). \quad (2)$$

Our hierarchical model is then

$$\mathbf{y} \mid \mathbf{x}, \mathbf{u}, \sigma_{\text{mod}} \stackrel{\text{iid}}{\sim} \mathcal{N}(\mathbf{H}\mathbf{x} + \mathbf{K}\mathbf{u}, \mathbf{R}), \quad (3)$$

$$\mathbf{x} \stackrel{\text{iid}}{\sim} \mathcal{LN}(0.16, 0.4^2), \quad (4)$$

$$\mathbf{u} \stackrel{\text{iid}}{\sim} \mathcal{LN}(0.004, 0.02^2), \quad (5)$$

$$\sigma_{\text{mod}} \sim \mathcal{LN}(2.2, 0.25^2), \quad (6)$$

127 where $\mathcal{N}(\cdot)$ and $\mathcal{LN}(\cdot)$ refer to the Normal and Lognormal distributions respectively.

128 We infer the emissions and influence from the boundary conditions using hierarchical
129 Bayesian inference²⁸. Sampling uses a two-stage sampler as in Say et al.²⁹. Firstly, a
130 No-U-Turn (NUTS) sampler³⁰ samples the latent field \mathbf{x} . A NUTS sampler is an exten-
131 sion to Hamiltonian Monte Carlo, which has previously been used for inference of trace-gas
132 emissions^{31,32}. A slice sampler³³, which computationally faster per iteration, samples the
133 hyperparameter σ_{mod} as a second step in the sampling process.

134 We infer methane emissions for each calendar month between April 2010 through 2017,
135 and assume that methane emissions are constant over each period of inference.

136 The elements of the latent field containing emissions are a basis function representation
137 of the NAME domain. We follow the approach of Say et al.²⁹, and optimise 100 basis
138 functions based on the a priori above-background mole fraction contribution in space using
139 a quadtree algorithm³⁴. The algorithm recursively divides the basis function into four new
140 basis functions until the desired number of basis functions is achieved, giving a higher spatial
141 resolution where there is a greater above-background a priori mole fraction contribution and
142 lower elsewhere (see Fig. S6).

143 We run the NUTS-slice sampler over a total of 250,000 iterations (burning the first
144 50,000), with multiple chains running in parallel. To check for convergence we use a Gelman-
145 Rubin diagnostic³⁵, ensuring all chains reach a criteria less than 1.05. The uncertainty in
146 the inferred emissions are quantified using the Highest Posterior Density (HPD) region (see

147 Box and Tiao³⁶ and supporting information).

148 Under this statistical model it is possible to infer the latent variables, namely the emis-
149 sions of methane and the boundary conditions, and the hyperparameter controlling the
150 uncertainty in the model error.

151 **Results and Discussion**

152 For Morocco, Algeria, Tunisia and Libya, the a priori estimates and UNFCCC National
153 Communications fall within the posterior 95% uncertainty, or are slightly underestimated,
154 for periods where reports are available. This would suggest that the inventories detailed in
155 the A priori methane emissions section, and the emissions estimates in UNFCCC National
156 Communications are largely consistent for these countries. Figure 1 shows the posterior
157 estimated emissions over the study period, where all emissions in the region were constrained
158 by the inversion (Figure S7). Figure S8 and Table S1 present these emissions as annual
159 means.

160 Estimated emissions from Egypt are consistently larger than the a priori estimates, al-
161 though they are smaller in 2015 than those reported in their National Communication. The
162 most noticeable discrepancy to the a priori estimate (Figure 1) is that Egypt has large, unex-
163 pected methane emission during the summer months, lasting from around June to October.
164 These summer emissions are uncorrelated with aerosol optical depth measurements in the
165 region³⁷. An increase in emissions for these months is observed in the a priori estimates,
166 attributed to rice cultivation²⁵, although to a much lesser extent than in the posterior es-
167 timates. These unexpected emissions come from north of Egypt, in the region containing
168 the Nile Delta, starting approximately north of the Aswan High Dam. Figure 3a shows
169 the the posterior mean flux estimate, and Figure 3b shows the a priori flux, as an average
170 for August over all years. These emissions in Egypt coincide with the summer agricultural
171 growing season, with high temperatures and high levels of irrigation. This is in contrast

172 to the rest of North Africa, where the main growing season is during the winter months,
173 and primarily driven by rainfall (see the supporting information for more discussion on agri-
174 cultural methane emissions outside of Egypt). Irrigation of crops in Egypt is generally fed
175 by branching canals, and drainage ditches³⁸, necessary due to the low levels of rainfall in
176 the region. As a result, high levels of irrigation are needed during the summer, with rice
177 and summer maize requiring the highest gross irrigation in the region³⁹ and sugarcane re-
178 quiring the highest levels of irrigation per square metre, followed by rice⁴⁰. Our a priori
179 annual mean estimate for agriculture (including rice) in Egypt using bottom up inventories
180 is 0.40 Tg, which is much smaller than 0.77 Tg in Egypt's 2015 UNFCCC reporting. Figure
181 2 shows the mean posterior estimated emissions for each month, alongside the daily mean
182 temperature at Cairo for each month, and the monthly satellite-estimated discharge from
183 Lake Nasser (feeding the Nile after the Aswan High Dam) from 2005-2008⁴¹. Although the
184 satellite-derived estimated discharge is not a direct indicator of water used for irrigation, it is
185 an indicator of the inflow to the Nile Delta. Figure 2 shows that the maximum in emissions
186 coincides within ± 1 month, with the maximum temperature or discharge. An exception is
187 that there is little to no summertime emissions increase in 2015, when North Africa expe-
188 rienced a drought⁴². Water levels of Lake Nasser⁴³ generally sharply drop in summertime,
189 coincident with larger discharge (Figure S9). Figure S9 shows that 2015 water levels had no
190 such drop and rise, perhaps indicating changes to discharge into the Nile Delta in 2015. This
191 is likely due to the droughts caused by the strong El Niño event in 2015, which impacted
192 the Nile river flow⁴⁴⁻⁴⁶ and global methane emissions⁴⁷. It is therefore likely that current
193 agricultural flooding practices, combined with high temperatures, are a major driver in the
194 Nile Delta's methane emissions. These findings corroborate evidence of the influence of the
195 (White) Nile on Africa's methane emissions⁷.

196 The biogeochemistry of methane release from fresh water is difficult to generalise, al-
197 though an increase in the area of water bodies within Egypt, coupled with warmer temper-
198 atures, would likely lead to increased methane emissions⁴⁸⁻⁵¹. Previous studies show that

199 river outflows from dammed reservoirs can release considerable levels of methane⁵², although
200 these processes have generally been observed at distances far closer to the dam itself than
201 from the Aswan High Dam to the bulk of emissions in the Nile Delta^{53,54}.

202 For validation of our results, we use weekly flask-air measurements from the Italian island
203 of Lampedusa^{55,56} (12.62°E, 35.52°N), which provides continuous measurements throughout
204 our period of study. Measured and forward modelled observations agree qualitatively (Fig.
205 S3), although the low-frequency of the measurements, which are primarily representative of
206 background air, make a more rigorous inter-comparison challenging.

207 Methane emissions from North Africa do not appear to have changed significantly (overall
208 trend of -0.2 ± 0.6 Tg year⁻¹, 95% uncertainty). However, our finding of a substantial and
209 unexpected seasonal source in the Nile Delta suggests that that agricultural emissions from
210 the region have been under-estimated in our a priori estimate. This may indicate a wider issue
211 of under-quantified emissions from agronomically managed temporary wetland ecosystems.

212 Ever increasing volumes of earth observation data at higher spatial resolution will allow
213 further regional scale studies to monitor changes in methane emission trends.

214 Code and data availability

215 ECMWF CAMS reanalysis data were downloaded from the Copernicus Atmosphere Monitor-
216 ing Service (CAMS) Atmosphere Data Store (ADS) [https://ads.atmosphere.copernicus.](https://ads.atmosphere.copernicus.eu/cdsapp#!/dataset/cams-global-reanalysis-eac4?tab=overview)
217 [eu/cdsapp#!/dataset/cams-global-reanalysis-eac4?tab=overview](https://ads.atmosphere.copernicus.eu/cdsapp#!/dataset/cams-global-reanalysis-eac4?tab=overview). Lake products cour-
218 tesy of the USDA/NASA G-REALM program at [https://ipad.fas.usda.gov/cropexplorer/](https://ipad.fas.usda.gov/cropexplorer/global_reservoir/)
219 [global_reservoir/](https://ipad.fas.usda.gov/cropexplorer/global_reservoir/). The latest version of the University of Leicester GOSAT Proxy v9.0
220 XCH₄ data are available from the Centre for Environmental Data Analysis data repository
221 at <https://doi.org/10.5285/18ef8247f52a4cb6a14013f8235cc1eb> (Parker and Boesch,
222 2020). The version used in this study (v7.2) is available from the Copernicus C3S Cli-
223 mate Data Store at <https://cds.climate.copernicus.eu>. Flask-air measurements from

224 Lampedusa are available at ftp://aftp.cmdl.noaa.gov/data/trace_gases/ch4/flask/
225 [surface/](#). The inversion results from this work, including all inputs to the inverse model,
226 is available at <https://osf.io/cdae3/> (Western 2021; DOI 10.17605/OSF.IO/CDAE3).
227 Access to the inversion code is available on request from the corresponding author.

228 **Acknowledgement**

229 We thank Eric Muala for supplying satellite estimated discharge data for Lake Nasser. We
230 thank the Japanese Aerospace Exploration Agency, National Institute for Environmental
231 Studies and the Ministry of Environment for the GOSAT data and their continuous support
232 as part of the Joint Research Agreement. This research used the ALICE High Perfor-
233 mance Computing Facility at the University of Leicester for the GOSAT retrievals. LMW,
234 RT and MR were supported by Natural Environment Research Council (NERC) grants
235 NE/N016548/1, NE/S016155/1, ALG was supported by a NERC Independent Research Fel-
236 lowship, NE/L010992/1, and AR was supported by a studentship from the NERC GW4+
237 Doctoral Training Partnership. Robert J. Parker and Hartmut Boesch are funded via the
238 UK National Centre for Earth Observation (NE/R016518/1 and NE/N018079/1).

239 **Supporting information**

240 The supporting information consists of sections describing North African petroleum and nat-
241 ural gas production; a description and presentation of the derived emissions using different a
242 priori boundary conditions; a description of the highest posterior density region; a discussion
243 on agricultural emissions outside of Egypt; validation of the results against measurements
244 made at Lampedusa in Italy and the supplementary tables and figures referred to in the
245 main text.

246 References

- 247 (1) Rigby, M.; Prinn, R. G.; Fraser, P. J.; Simmonds, P. G.; Langenfelds, R. L.;
248 Huang, J.; Cunnold, D. M.; Steele, L. P.; Krummel, P. B.; Weiss, R. F.; O’Doherty, S.;
249 Salameh, P. K.; Wang, H. J.; Harth, C. M.; Mühle, J.; Porter, L. W. Renewed growth
250 of atmospheric methane. *Geophysical Research Letters* **2008**, *35*.
- 251 (2) Schaefer, H.; Fletcher, S. E. M.; Veidt, C.; Lassey, K. R.; Brailsford, G. W.; Brom-
252 ley, T. M.; Dlugokencky, E. J.; Michel, S. E.; Miller, J. B.; Levin, I.; Lowe, D. C.;
253 Martin, R. J.; Vaughn, B. H.; White, J. W. C. A 21st-century shift from fossil-fuel to
254 biogenic methane emissions indicated by $^{13}\text{CH}_4$. *Science* **2016**, *352*, 80–84.
- 255 (3) Worden, J. R.; Bloom, A. A.; Pandey, S.; Jiang, Z.; Worden, H. M.; Walker, T. W.;
256 Houweling, S.; Röckmann, T. Reduced biomass burning emissions reconcile conflicting
257 estimates of the post-2006 atmospheric methane budget. *Nature Communications* **2017**,
258 *8*.
- 259 (4) Nisbet, E. G.; Dlugokencky, E. J.; Manning, M. R.; Lowry, D.; Fisher, R. E.;
260 France, J. L.; Michel, S. E.; Miller, J. B.; White, J. W. C.; Vaughn, B.; Bousquet, P.;
261 Pyle, J. A.; Warwick, N. J.; Cain, M.; Brownlow, R.; Zazzeri, G.; Lanoisellé, M.; Man-
262 ning, A. C.; Gloor, E.; Worthy, D. E. J. et al. Rising atmospheric methane: 2007-2014
263 growth and isotopic shift. *Global Biogeochemical Cycles* **2016**, *30*, 1356–1370.
- 264 (5) Rigby, M.; Montzka, S. A.; Prinn, R. G.; White, J. W. C.; Young, D.; O’Doherty, S.;
265 Lunt, M. F.; Ganesan, A. L.; Manning, A. J.; Simmonds, P. G.; Salameh, P. K.;
266 Harth, C. M.; Mühle, J.; Weiss, R. F.; Fraser, P. J.; Steele, L. P.; Krummel, P. B.;
267 McCulloch, A.; Park, S. Role of atmospheric oxidation in recent methane growth. *Pro-
268 ceedings of the National Academy of Sciences* **2017**, *114*, 5373–5377.
- 269 (6) Turner, A. J.; Frankenberg, C.; Wennberg, P. O.; Jacob, D. J. Ambiguity in the causes

- 270 for decadal trends in atmospheric methane and hydroxyl. *Proceedings of the National*
271 *Academy of Sciences* **2017**, *114*, 5367–5372.
- 272 (7) Lunt, M. F.; Palmer, P. I.; Feng, L.; Taylor, C. M.; Boesch, H.; Parker, R. J. An increase
273 in methane emissions from tropical Africa between 2010 and 2016 inferred from satellite
274 data. *Atmospheric Chemistry and Physics* **2019**, *19*, 14721–14740.
- 275 (8) Pandey, S.; Houweling, S.; Lorente, A.; Borsdorff, T.; Tzivlidou, M.; Bloom, A. A.;
276 Poulter, B.; Zhang, Z.; Aben, I. Using satellite data to identify the methane emission
277 controls of South Sudan’s wetlands. *Biogeosciences* **2021**, *18*, 557–572.
- 278 (9) Scarpelli, T. R.; Jacob, D. J.; Maasakkers, J. D.; Sulprizio, M. P.; Sheng, J.-X.; Rose, K.;
279 Romeo, L.; Worden, J. R.; Janssens-Maenhout, G. A global gridded ($0.1^\circ \times 0.1^\circ$)
280 inventory of methane emissions from oil, gas, and coal exploitation based on national
281 reports to the United Nations Framework Convention on Climate Change. *Earth System*
282 *Science Data* **2020**, *12*, 563–575.
- 283 (10) EIA, EIA: International Energy Statistics. <http://eia.gov/beta/international/>
284 (accessed 22 January 2021).
- 285 (11) Ministère de l’Amenagement du Territoire et de l’Environnement, Seconde Communi-
286 cation Nationale de l’Algerie sur les Changements Climatiques a la CCNUCC. 2010.
- 287 (12) Ministry of Environment: Egyptian Environmental Affairs Agency, Egypt’s First Bien-
288 nial Update Report to the United Nations Framework Convention on Climate Change.
289 2018.
- 290 (13) Département de l’Environnement, Deuxième Rapport Biennal actualisé du Maroc dans le
291 cadre de la CCNUCC. 2019.
- 292 (14) Ministère des Affaires Locales et de l’Environnement, Deuxième Rapport Biennal de la
293 Tunisie. 2016.

- 294 (15) Jones, A.; Thomson, D.; Hort, M.; Devenish, B. The U.K. Met Office’s Next-Generation
295 Atmospheric Dispersion Model, NAME III. *Air Pollution Modeling and Its Application*
296 *XVII* **2006**, 580–589.
- 297 (16) Kuze, A.; Suto, H.; Nakajima, M.; Hamazaki, T. Thermal and near infrared sensor for
298 carbon observation Fourier-transform spectrometer on the Greenhouse Gases Observing
299 Satellite for greenhouse gases monitoring. *Applied Optics* **2009**, *48*, 6716.
- 300 (17) Parker, R.; Boesch, H.; Cogan, A.; Fraser, A.; Feng, L.; Palmer, P. I.; Messerschmidt, J.;
301 Deutscher, N.; Griffith, D. W. T.; Notholt, J.; Wennberg, P. O.; Wunch, D. Methane
302 observations from the Greenhouse Gases Observing SATellite: Comparison to ground-
303 based TCCON data and model calculations. *Geophysical Research Letters* **2011**, *38*.
- 304 (18) Parker, R. J.; Webb, A.; Boesch, H.; Somkuti, P.; Barrio Guillo, R.; Di Noia, A.;
305 Kalaitzi, N.; Anand, J. S.; Bergamaschi, P.; Chevallier, F.; Palmer, P. I.; Feng, L.;
306 Deutscher, N. M.; Feist, D. G.; Griffith, D. W. T.; Hase, F.; Kivi, R.; Morino, I.;
307 Notholt, J.; Oh, Y.-S. et al. A decade of GOSAT Proxy satellite CH₄ observations.
308 *Earth System Science Data* **2020**, *12*, 3383–3412.
- 309 (19) Ganesan, A. L.; Rigby, M.; Lunt, M. F.; Parker, R. J.; Boesch, H.; Goulding, N.;
310 Umezawa, T.; Zahn, A.; Chatterjee, A.; Prinn, R. G.; Tiwari, Y. K.; van der Schoot, M.;
311 Krummel, P. B. Atmospheric observations show accurate reporting and little growth in
312 India’s methane emissions. *Nature Communications* **2017**, *8*.
- 313 (20) Tunnicliffe, R. L.; Ganesan, A. L.; Parker, R. J.; Boesch, H.; Gedney, N.; Poulter, B.;
314 Zhang, Z.; Lavrič, J. V.; Walter, D.; Rigby, M.; Henne, S.; Young, D.; O’Doherty, S.
315 Quantifying sources of Brazil’s CH₄ emissions between 2010 and 2018 from satellite
316 data. *Atmospheric Chemistry and Physics* **2020**, *20*, 13041–13067.
- 317 (21) Walters, D. N.; Williams, K. D.; Boutle, I. A.; Bushell, A. C.; Edwards, J. M.;
318 Field, P. R.; Lock, A. P.; Morcrette, C. J.; Stratton, R. A.; Wilkinson, J. M.; Wil-

- 319 lett, M. R.; Bellouin, N.; Bodas-Salcedo, A.; Brooks, M. E.; Copsey, D.; Earn-
320 shaw, P. D.; Hardiman, S. C.; Harris, C. M.; Levine, R. C.; MacLachlan, C. et al.
321 The Met Office Unified Model Global Atmosphere 4.0 and JULES Global Land 4.0
322 configurations. *Geoscientific Model Development* **2014**, *7*, 361–386.
- 323 (22) O’Dell, C. W.; Connor, B.; Bösch, H.; O’Brien, D.; Frankenberg, C.; Castano, R.;
324 Christi, M.; Eldering, D.; Fisher, B.; Gunson, M.; McDuffie, J.; Miller, C. E.; Na-
325 traj, V.; Oyafuso, F.; Polonsky, I.; Smyth, M.; Taylor, T.; Toon, G. C.; Wennberg, P. O.;
326 Wunch, D. The ACOS CO₂ retrieval algorithm: Part 1: Description and validation
327 against synthetic observations. *Atmospheric Measurement Techniques* **2012**, *5*, 99–121.
- 328 (23) van der Werf, G. R.; Randerson, J. T.; Giglio, L.; van Leeuwen, T. T.; Chen, Y.;
329 Rogers, B. M.; Mu, M.; van Marle, M. J. E.; Morton, D. C.; Collatz, G. J.; Yokel-
330 son, R. J.; Kasibhatla, P. S. Global fire emissions estimates during 1997-2016. *Earth*
331 *System Science Data* **2017**, *9*, 697–720.
- 332 (24) Bloom, A. A.; Bowman, K. W.; Lee, M.; Turner, A. J.; Schroeder, R.; Worden, J. R.;
333 Weidner, R.; McDonald, K. C.; Jacob, D. J. A global wetland methane emissions and
334 uncertainty dataset for atmospheric chemical transport models (WetCHARTs version
335 1.0). *Geoscientific Model Development* **2017**, *10*, 2141–2156.
- 336 (25) Yan, X.; Akiyama, H.; Yagi, K.; Akimoto, H. Global estimations of the inventory and
337 mitigation potential of methane emissions from rice cultivation conducted using the
338 2006 Intergovernmental Panel on Climate Change Guidelines. *Global Biogeochemical*
339 *Cycles* **2009**, *23*, n/a–n/a.
- 340 (26) Janssens-Maenhout, G.; Crippa, M.; Guizzardi, D.; Muntean, M.; Schaaf, E.; Den-
341 tener, F.; Bergamaschi, P.; Pagliari, V. EDGAR v4.3.2 Global Atlas of the three major
342 Greenhouse Gas Emissions for the period 1970-2012. *Open Access* **2017**, 55.
- 343 (27) Inness, A.; Ades, M.; AgustÃ-Panareda, A.; BarrÃ©, J.; Benedictow, A.; Blech-

- 344 schmidt, A.-M.; Dominguez, J. J.; Engelen, R.; Eskes, H.; Flemming, J.; Huijnen, V.;
345 Jones, L.; Kipling, Z.; Massart, S.; Parrington, M.; Peuch, V.-H.; Razinger, M.;
346 Remy, S.; Schulz, M.; Suttie, M. The CAMS reanalysis of atmospheric composition.
347 *Atmospheric Chemistry and Physics* **2019**, *19*, 3515–3556.
- 348 (28) Ganesan, A. L.; Rigby, M.; Zammit-Mangion, A.; Manning, A. J.; Prinn, R. G.;
349 Fraser, P. J.; Harth, C. M.; Kim, K.-R.; Krummel, P. B.; Li, S.; Mählhle, J.;
350 O’Doherty, S. J.; Park, S.; Salameh, P. K.; Steele, L. P.; Weiss, R. F. Characteri-
351 zation of uncertainties in atmospheric trace gas inversions using hierarchical Bayesian
352 methods. *Atmospheric Chemistry and Physics* **2014**, *14*, 3855–3864.
- 353 (29) Say, D.; Kuyper, B.; Western, L.; Khan, M. A. H.; Lesch, T.; Labuschagne, C.; Mar-
354 tin, D.; Young, D.; Manning, A. J.; O’Doherty, S.; Rigby, M.; Krummel, P. B.; Davies-
355 Coleman, M. T.; Ganesan, A. L.; Shallcross, D. E. Emissions and Marine Boundary
356 Layer Concentrations of Unregulated Chlorocarbons Measured at Cape Point, South
357 Africa. *Environmental Science & Technology* **2020**, *54*, 10514–10523.
- 358 (30) Hoffman, M. D.; Gelman, A. The No-U-Turn Sampler: Adaptively Setting Path Lengths
359 in Hamiltonian Monte Carlo. *Journal of Machine Learning Research* **2014**, *15*, 1593–
360 1623.
- 361 (31) Zammit-Mangion, A.; Cressie, N.; Ganesan, A. L.; O’Doherty, S.; Manning, A. J.
362 Spatio-temporal bivariate statistical models for atmospheric trace-gas inversion.
363 *Chemometrics and Intelligent Laboratory Systems* **2015**, *149*, 227–241.
- 364 (32) Zammit-Mangion, A.; Cressie, N.; Ganesan, A. L. Non-Gaussian bivariate modelling
365 with application to atmospheric trace-gas inversion. *Spatial Statistics* **2016**, *18*, 194–
366 220.
- 367 (33) Neal, R. M. Slice sampling. *The Annals of Statistics* **2003**, *31*, 705–767.

- 368 (34) Finkel, R. A.; Bentley, J. L. Quad trees a data structure for retrieval on composite keys.
369 *Acta Informatica* **1974**, *4*, 1–9.
- 370 (35) Gelman, A.; Rubin, D. B. Inference from Iterative Simulation Using Multiple Sequences.
371 *Statistical Science* **1992**, *7*, 457–472, Publisher: Institute of Mathematical Statistics.
- 372 (36) Box, G. E. P.; Tiao, G. C. *Bayesian inference in statistical analysis*, Wiley Classics
373 Library ed. ed.; A Wiley-Interscience publication; Wiley: New York, 1992; OCLC:
374 25247039.
- 375 (37) El-Metwally, M.; Alfaro, S. C.; Abdel Wahab, M. M.; Zakey, A. S.; Chatenet, B. Sea-
376 sonal and inter-annual variability of the aerosol content in Cairo (Egypt) as deduced
377 from the comparison of MODIS aerosol retrievals with direct AERONET measurements.
378 *Atmospheric Research* **2010**, *97*, 14–25.
- 379 (38) El-Agha, D. E.; Molden, D. J.; Ghanem, A. M. Performance assessment of irrigation
380 water management in old lands of the Nile delta of Egypt. *Irrigation and Drainage*
381 *Systems* **2011**, *25*, 215–236.
- 382 (39) Multsch, S.; Elshamy, M.; Batarseh, S.; Seid, A.; Frede, H.-G.; Breuer, L. Improving
383 irrigation efficiency will be insufficient to meet future water demand in the Nile Basin.
384 *Journal of Hydrology: Regional Studies* **2017**, *12*, 315–330.
- 385 (40) Osama, S.; Elkholy, M.; Kansoh, R. M. Optimization of the cropping pattern in Egypt.
386 *Alexandria Engineering Journal* **2017**, *56*, 557–566.
- 387 (41) Muala, E.; Mohamed, Y. A.; Duan, Z.; Van der Zaag, P. Estimation of Reservoir
388 Discharges from Lake Nasser and Roseires Reservoir in the Nile Basin Using Satellite
389 Altimetry and Imagery Data. *Remote Sensing* **2014**, *6*, 7522–7545.
- 390 (42) Bazza, M.; Kay, M.; Knutson, C. Drought characteristics and management in North
391 Africa and the Near East. *FAO Water Reports* **2018**,

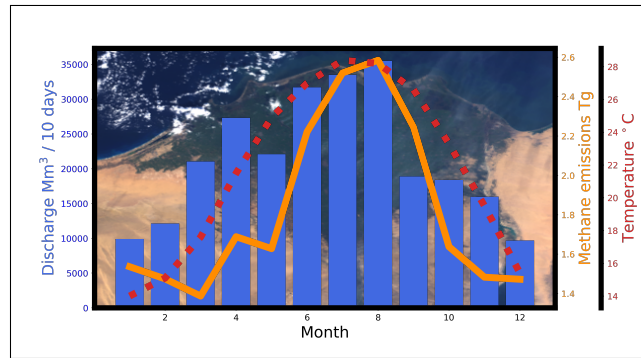
- 392 (43) Birkett, C.; Reynolds, C.; Beckley, B.; Doorn, B. In *Coastal Altimetry*; Vignudelli, S.,
393 Kostianoy, A. G., Cipollini, P., Benveniste, J., Eds.; Springer Berlin Heidelberg: Berlin,
394 Heidelberg, 2011; pp 19–50.
- 395 (44) Zaroug, M. A. H.; Eltahir, E. A. B.; Giorgi, F. Droughts and floods over the upper
396 catchment of the Blue Nile and their connections to the timing of El Niño and La Niña
397 events. *Hydrology and Earth System Sciences* **2014**, *18*, 1239–1249.
- 398 (45) Siam, M. S.; Eltahir, E. A. B. Climate change enhances interannual variability of the
399 Nile river flow. *Nature Climate Change* **2017**, *7*, 6.
- 400 (46) Conway, D. Future Nile river flows. *Nature Climate Change* **2017**, *7*, 319–320.
- 401 (47) Zhang, Z.; Zimmermann, N. E.; Calle, L.; Hurtt, G.; Chatterjee, A.; Poulter, B.
402 Enhanced response of global wetland methane emissions to the 2015-2016 El Niño-
403 Southern Oscillation event. *Environmental Research Letters* **2018**, *13*, 074009, Pub-
404 lisher: IOP Publishing.
- 405 (48) Bastviken, D.; Cole, J.; Pace, M.; Tranvik, L. Methane emissions from
406 lakes: Dependence of lake characteristics, two regional assessments,
407 and a global estimate. *Global Biogeochemical Cycles* **2004**, *18*, eprint:
408 <https://agupubs.onlinelibrary.wiley.com/doi/pdf/10.1029/2004GB002238>.
- 409 (49) Wilkinson, J.; Maeck, A.; Alshboul, Z.; Lorke, A. Continuous Seasonal River Ebulli-
410 tion Measurements Linked to Sediment Methane Formation. *Environmental Science &*
411 *Technology* **2015**, *49*, 13121–13129.
- 412 (50) Holgerson, M. A.; Raymond, P. A. Large contribution to inland water CO₂ and CH₄
413 emissions from very small ponds. *Nature Geoscience* **2016**, *9*, 222–226, Number: 3
414 Publisher: Nature Publishing Group.

- 415 (51) Aguilera, E.; Vila-Traver, J.; Deemer, B. R.; Infante-Amate, J.; Guzmán, G. I.;
416 González de Molina, M. Methane Emissions from Artificial Waterbodies Dominate the
417 Carbon Footprint of Irrigation: A Study of Transitions in the Food-Energy-Water-
418 Climate Nexus (Spain, 1900-2014). *Environmental Science & Technology* **2019**, *53*,
419 5091–5101, Publisher: American Chemical Society.
- 420 (52) Prairie, Y. T.; Alm, J.; Beaulieu, J.; Barros, N.; Battin, T.; Cole, J.; del Giorgio, P.; Del-
421 Sontro, T.; Guérin, F.; Harby, A.; Harrison, J.; Mercier-Blais, S.; Serça, D.; Sobek, S.;
422 Vachon, D. Greenhouse Gas Emissions from Freshwater Reservoirs: What Does the
423 Atmosphere See? *Ecosystems* **2018**, *21*, 1058–1071.
- 424 (53) Abril, G.; Guerin, F.; Richard, S.; Delmas, R.; GalyLacaux, C.; Gosse, P.;
425 Tremblay, A.; Varfalvy, L.; Santos, M. A. D.; Matvienko, B. Carbon dioxide
426 and methane emissions and the carbon budget of a 10-year old tropical reser-
427 voir (Petit Saut, French Guiana). *Global Biogeochemical Cycles* **2005**, *19*, eprint:
428 <https://agupubs.onlinelibrary.wiley.com/doi/pdf/10.1029/2005GB002457>.
- 429 (54) Kemenes, A.; Forsberg, B. R.; Melack, J. M. Methane release below a
430 tropical hydroelectric dam. *Geophysical Research Letters* **2007**, *34*, eprint:
431 <https://agupubs.onlinelibrary.wiley.com/doi/pdf/10.1029/2007GL029479>.
- 432 (55) Artuso, F.; Chamard, P.; Piacentino, S.; di Sarra, A.; Meloni, D.; Monteleone, F.;
433 Sferlazzo, D. M.; Thiery, F. Atmospheric methane in the Mediterranean: Analysis of
434 measurements at the island of Lampedusa during 1995-2005. *Atmospheric Environment*
435 **2007**, *41*, 3877–3888.
- 436 (56) Dlugokencky, E.; Lang, P.; Crotwell, A.; Mund, J.; Crotwell, M.; Thoning, K. At-
437 mospheric Methane Dry Air Mole Fractions from the NOAA ESRL Carbon Cycle
438 Cooperative Global Air Sampling Network, 1983-2016, Version: 2017-07-28. 2017;

439 ftp://aftp.cmdl.noaa.gov/data/trace_gases/ch4/flask/surface/, (accessed 19
440 January 2021).

441 **Graphical TOC Entry**

442



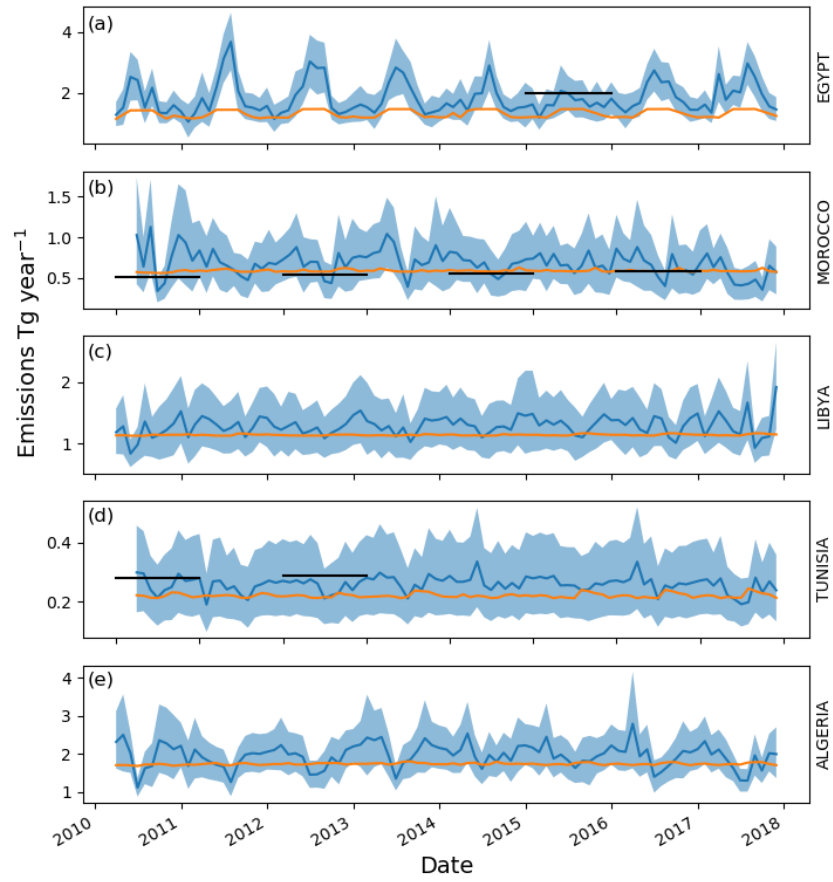


Figure 1: Emissions from countries in North Africa between 2010-2017: (a) Egypt, (b) Morocco, (c)Libya, (d) Tunisia and (e) Algeria. The blue line shows the posterior mean emissions for each month, and the blue shading shows the 95 % HPD region. The orange line shows the a priori emissions for each country, and the black lines show annual emissions reported to the UNFCCC.

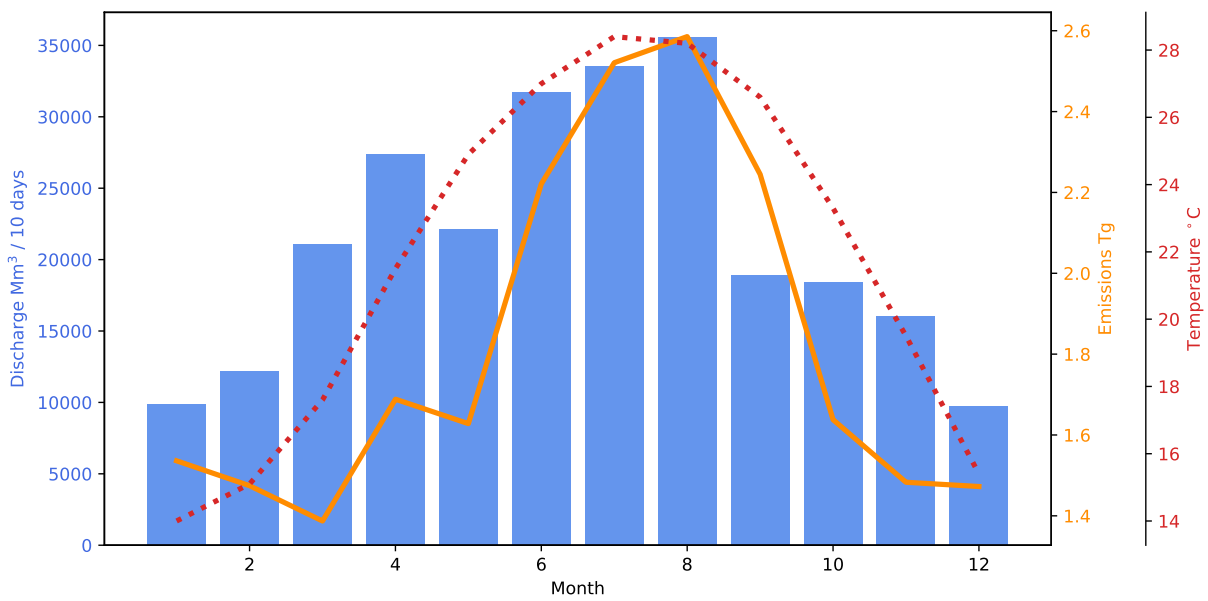


Figure 2: The mean posterior estimated emissions for Egypt for each month between 2010-2017 (orange), the daily mean temperature at Cairo for each month (red), and the monthly satellite-estimated discharge from Lake Nasser (feeding the Nile after the Aswan High Dam) from 2005-2008⁴¹ (blue).

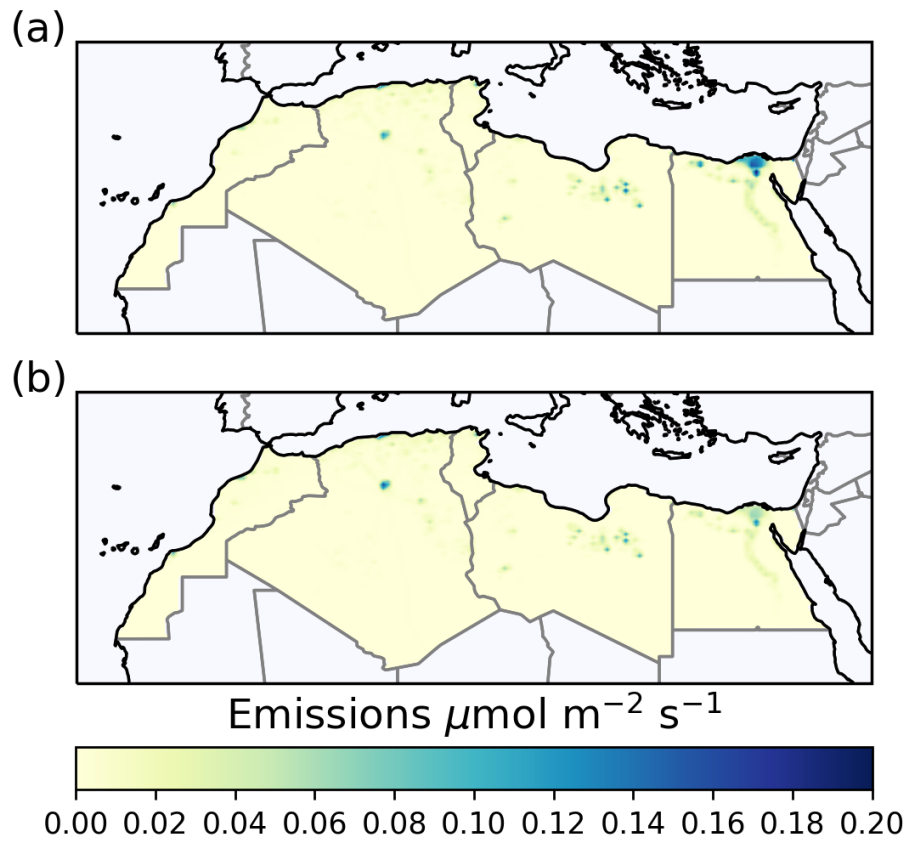


Figure 3: A map showing (a) the posterior mean estimated flux and (b) the a priori flux from the North African region as an average for August over all years.

Supplement to: Estimates of North African methane emissions from 2010-2017 using GOSAT observations

Luke M. Western,^{*,†} Alice E. Ramsden,[‡] Anita L. Ganesan,[‡] Hartmut Boesch,^{¶,§}
Robert J. Parker,^{¶,§} Tia R. Scarpelli,^{||} Rachel L. Tunnicliffe,[†] and Matthew
Rigby[†]

[†]*School of Chemistry, University of Bristol, Bristol, BS8 1TS, UK*

[‡]*School of Geographical Sciences, University of Bristol, Bristol, BS8 1SS, UK*

[¶]*National Centre for Earth Observation, University of Leicester, Leicester, LE1 7RH, UK*

[§]*Earth Observation Science, School of Physics and Astronomy, University of Leicester,
Leicester, LE1 7RH, UK*

^{||}*Earth and Planetary Sciences, Harvard University, Cambridge, 02138, USA*

E-mail: luke.western@bristol.ac.uk

North African petroleum and natural gas production

Production of petroleum products in Algeria, the main producer in the region, have remained steady over the period of study, at around 1881 thousand barrels per day (TBPD) in 2010, reducing to around 1637 TBPD by the end of 2017¹ (Figure S1). Libya, the second largest producer, has had large fluctuations in production since the Arab Spring², starting in February 2011. Production has not recovered since 2010, where annual production was 1844 TBPD, and hit an annual low of 478 TBPD in 2016. Production in Egypt has remained

fairly constant over the period of study, with annual production varying between 655 - 714 TBPD. Natural gas production is again largest in Algeria, followed by Egypt. Production in Algeria remained largely unchanged between 2010-2015 (~ 185 billion cubic metres, BCM), and declined slightly in Egypt from 67 to 49 BCM between 2010-2015. Libya's post-2010 gas production is much smaller in comparison (10-18 BCM), but was 30 BCM in 2010. Tunisia and Morocco are comparatively minor producers of oil and gas.

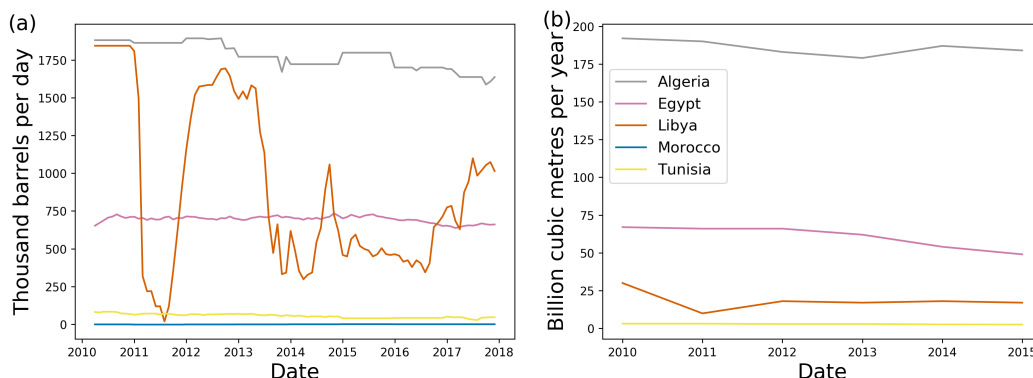


Figure S1: Oil (a) and natural gas (b) production in Algeria, Egypt, Libya, Morocco and Tunisia. Any natural gas production in Morocco was negligible. Production data for natural gas were only available until 2015. The data are from the U.S. Energy Information Administration¹.

Results using different a priori boundary conditions

We repeat our results to test the influence of the chosen a priori mole fraction at the model domain boundary. The inverse method and sensitivity to the boundaries was treated as in the main text, but with the a priori mole fraction at the domain boundary taken from the ECMWF CAMS CH₄ flux inversion product v17r1 (<https://ads.atmosphere.copernicus.eu/cdsapp#!/dataset/cams-global-greenhouse-gas-inversion?tab=overview>, accessed 22 April 2021). The data product is produced using surface observations only. See https://atmosphere.copernicus.eu/sites/default/files/2018-12/CAMS73_2015SC3_D73.2.4.4-2017_201811_validation_1990-2017_v1.pdf for more information (accessed 22 April

2021).

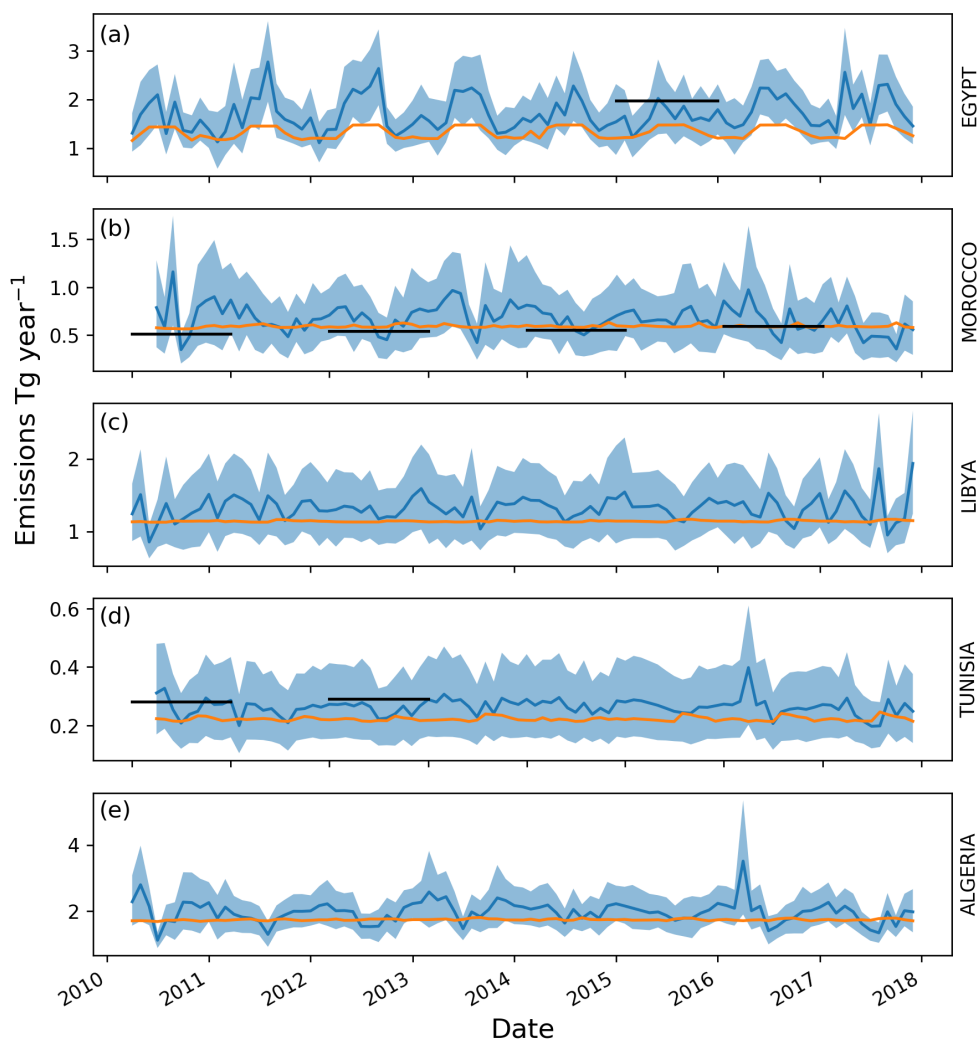


Figure S2: Emissions from countries in North Africa between 2010-2017 using different a priori mole fractions at the boundary to that in the main text. (a) Egypt, (b) Morocco, (c) Libya, (d) Tunisia and (e) Algeria. The blue line shows the posterior mean emissions for each month, and the blue shading shows the 95 % HPD region. The orange line shows the a priori emissions for each country, and the black lines show annual emissions reported to the UNFCCC.

Figure S2 shows the emissions time series from the five North African countries (as Figure 1, main text) using the alternate a priori mole fraction at the boundary, as described. Emissions do not change substantially from those presented in the main text and do not change any of the conclusions drawn, although some small differences exist. Most notably, summertime emissions peaks in Egypt are not as high as in the main text, but still remain

elevated above that observed in the a priori emissions estimates and exhibit a seasonal cycle.

Highest Posterior Density region

The Highest Posterior Density (HPD) region is the narrowest region, R , in the total posterior parameter space that holds probability content $(1 - \alpha)$. This is achieved if the following conditions are fulfilled,

1. $p\{\mathbf{x} \in R \mid \mathbf{y}\} = (1 - \alpha)$
2. for $\mathbf{x}_1 \in R$ and $\mathbf{x}_2 \notin R$, $p(\mathbf{x}_1 \mid \mathbf{y}) \geq p(\mathbf{x}_2 \mid \mathbf{y})$.

Agricultural emissions outside of Egypt

Unlike in Egypt, where irrigation for agriculture is dependent on the Nile, agricultural practices in the rest of North Africa are dependent on precipitation as their main water source³. The agricultural sector is the largest contributor to gross domestic product in both Algeria and Libya³. There is more rainfall in North Africa during the Northern Hemisphere winter, when temperatures are lower, with little rainfall and high temperatures during the Northern Hemisphere summer⁴⁻⁶. Therefore, it is expected, given the observations in Egypt, that emissions from agronomically managed temporary wetlands in North Africa, other than Egypt, would have higher emissions in winter months (during their crop growing season with maximum rainfall), than in the summer months. In addition, livestock in North Africa are slaughtered during times of fodder scarcity, leading to lower emissions from ruminants during times of low rainfall³.

A pattern of higher North African methane emissions in winter months during the agricultural season is somewhat apparent in our inferred emissions (see Figure 1, main text), particularly in Algeria, although to a much lesser extent than the seasonality in Egypt. This may be explained by the lower temperatures during the agricultural season, in contrast

to high summer temperatures during Egypt’s agricultural season, leading to lower rates of methanogenesis^{7–10}.

Validation against measurements made at Lampedusa

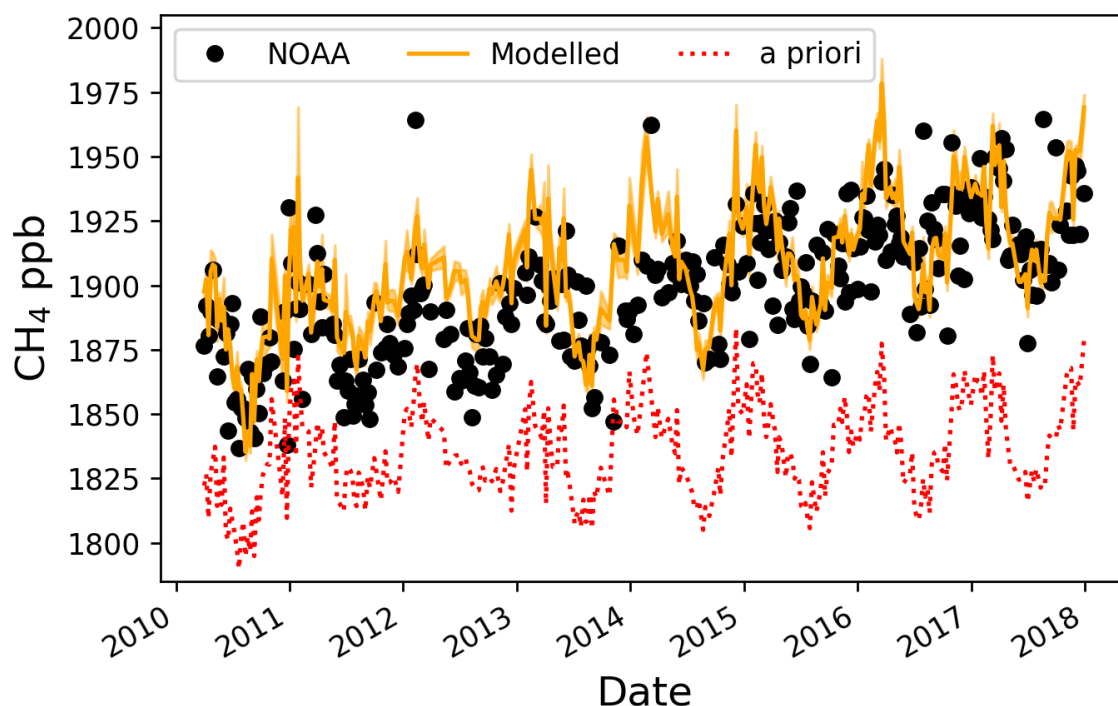


Figure S3: Methane concentrations (black circles) measured at Lampedusa measurement station (12.62°E, 35.52°N) and forward modelled concentrations (orange lines) using the estimated emissions and NAME-derived sensitivities. Also shown are the forward modelled a priori concentrations (dotted red line), modelled using the a priori emissions and NAME-derived sensitivities.

The paucity of in situ measurement data in the region of study, which motivates the use of satellite-observation derived emissions, makes detailed validation against an independent dataset difficult. Here we validate our emissions by comparing our posterior model to weekly flask-air samples from the Italian island of Lampedusa^{11,12} (12.62°E, 35.52°N).

The modelled concentrations are calculated from NAME footprints, simulated using

20,000 particles released for a 1-hour period when each measurement was made at a height of 10 ± 10 metres above ground level. The particles were tracked for 30 days and their interaction with the surface and boundaries were recorded as described for GOSAT in the main text.

Given the low frequency of the measurements, which are made to sample background conditions, a rigorous validation of the posterior emissions is difficult. In addition, errors in the transport and meteorology using NAME may compound errors in the forward modelled concentrations using GOSAT- and NAME-derived emissions estimates. It is clear, however, that the posterior mean predicted mole fractions greatly improve the fit to observations over the a priori predicted concentrations, as can be seen in Figure , where the a priori predicted concentration falls far below the observations. Some simple statistics using the posterior mean predicted and a priori predicted mole fraction shows an improvement to the root-mean-square error from 66 ppb to 21 ppb, and an improvement to the R^2 statistic from 0.37 to 0.54.

Supplementary Tables and Figures

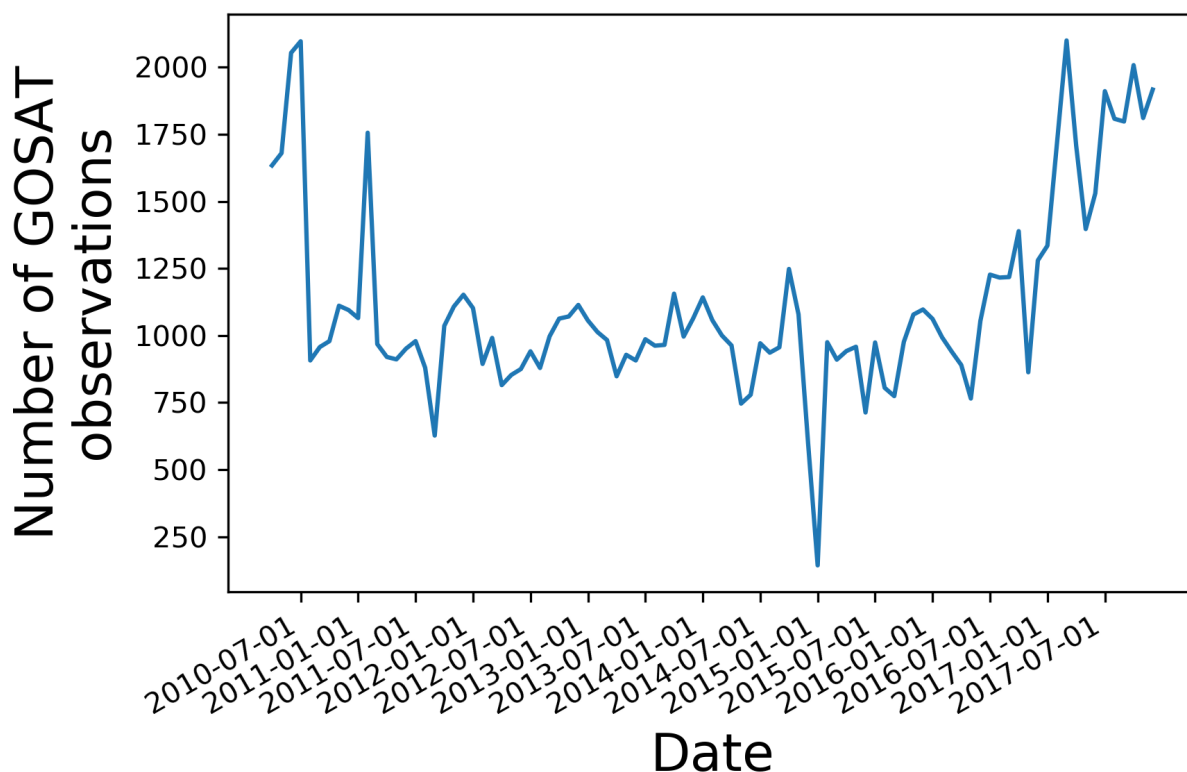


Figure S4: The number of GOSAT $\text{XCH}_4_{\text{proxy}}$ observations per month used in the inversion.

Table S1: Annual estimates of mean posterior methane emissions and the 95% uncertainty range. The annual emissions are estimated from the monthly estimates within each year, assuming that the posterior distributions are independent, which will most likely underestimate the true annual uncertainty.

Year	Annual emissions (Tg year^{-1})				
	Egypt	Morocco	Libya	Tunisia	Algeria
2010	1.50 (1.35, 1.66)	0.78 (0.62, 0.94)	1.16 (1.01, 1.30)	0.27 (0.22, 0.31)	2.00 (1.75, 2.25)
2011	1.76 (1.60, 1.93)	0.65 (0.56, 0.74)	1.31 (1.18, 1.45)	0.24 (0.21, 0.28)	1.80 (1.65, 1.96)
2012	1.64 (1.49, 1.80)	0.67 (0.58, 0.77)	1.26 (1.13, 1.39)	0.25 (0.22, 0.29)	1.85 (1.71, 2.00)
2013	1.64 (1.48, 1.79)	0.75 (0.64, 0.86)	1.31 (1.17, 1.45)	0.26 (0.22, 0.30)	2.07 (1.88, 2.26)
2014	1.61 (1.47, 1.75)	0.66 (0.57, 0.77)	1.32 (1.19, 1.45)	0.27 (0.23, 0.31)	2.01 (1.84, 2.18)
2015	1.49 (1.34, 1.64)	0.69 (0.59, 0.79)	1.33 (1.19, 1.47)	0.27 (0.23, 0.30)	2.00 (1.83, 2.17)
2016	1.77 (1.63, 1.92)	0.65 (0.55, 0.75)	1.30 (1.17, 1.43)	0.26 (0.22, 0.29)	1.95 (1.77, 2.14)
2017	1.82 (1.67, 1.98)	0.55 (0.48, 0.63)	1.33 (1.20, 1.46)	0.23 (0.20, 0.27)	1.82 (1.68, 1.96)

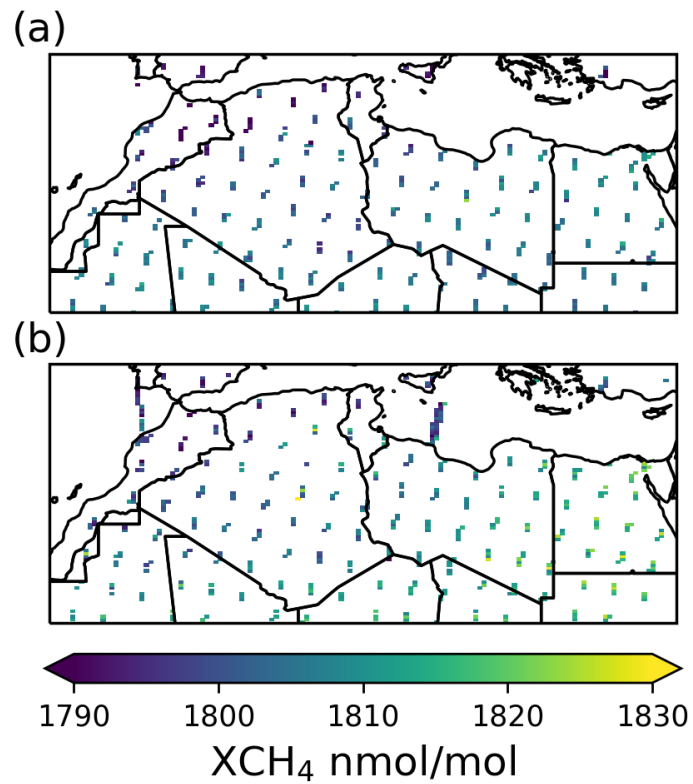


Figure S5: The average GOSAT XCH₄_{proxy} observations binned to the resolution of the NAME dispersion model output for (a) November 2011 - April 2012 and (b) May 2012 - October 2012.

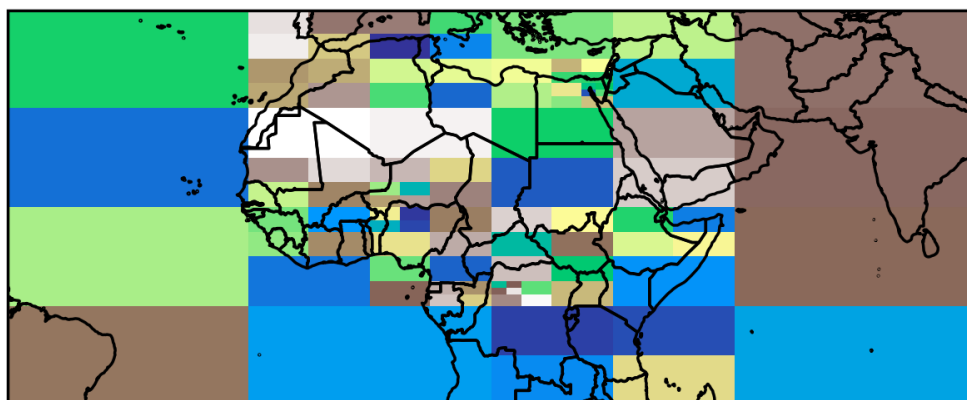


Figure S6: The basis functions representation of the emissions within the model domain for March 2013.

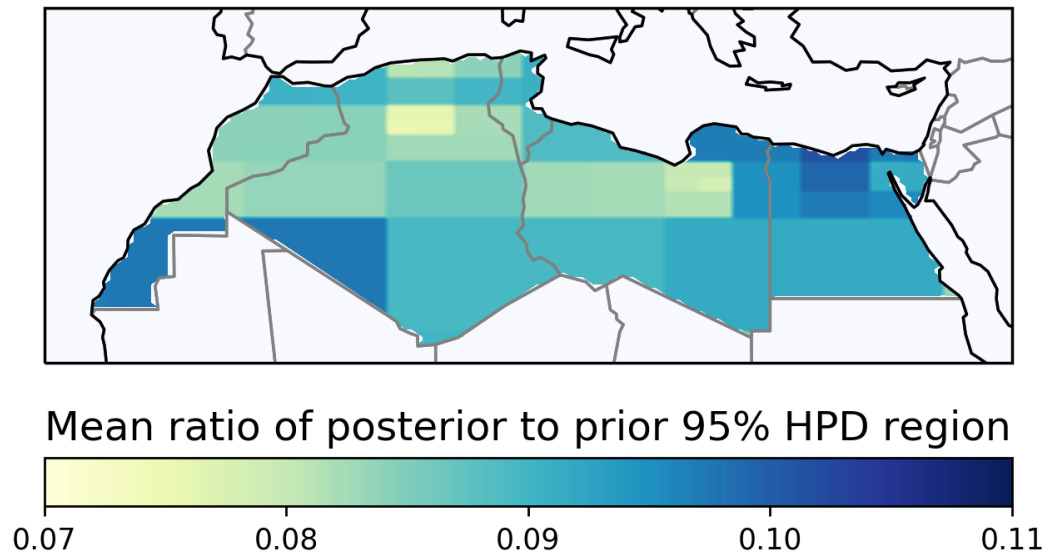


Figure S7: The mean ratio of the range of the upper and lower bounds of the posterior to prior 95% HPD region in space. This mean is over all months estimated within the study. The map shows that, on average in the region of study, the range between the 97.5% and 2.5% uncertainty bound in the posterior distribution is, at most, 10% of that in the prior, or, equivalently, shows at least a 90% uncertainty reduction.

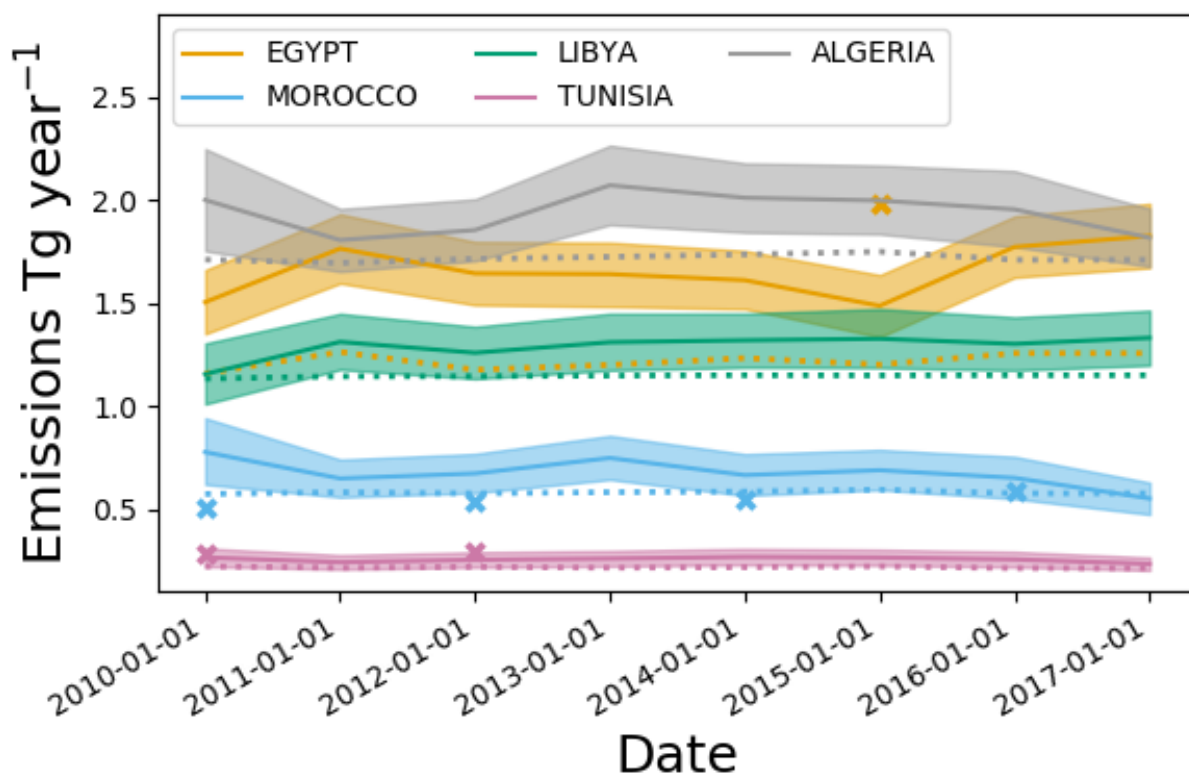


Figure S8: Methane emissions estimates in North Africa presented as annual means. The lines are posterior mean emissions estimates, and the shading shows the 95% posterior uncertainty. The dashed line shows the annual mean a priori emissions estimate and the crosses show estimates of methane emissions submitted as national reporting to the UNFCCC. The emissions estimates are tabulated in Table S1. The annual emissions are estimated from the monthly estimates within each year, assuming that the posterior distributions are independent, which will most likely underestimate the true annual uncertainty.

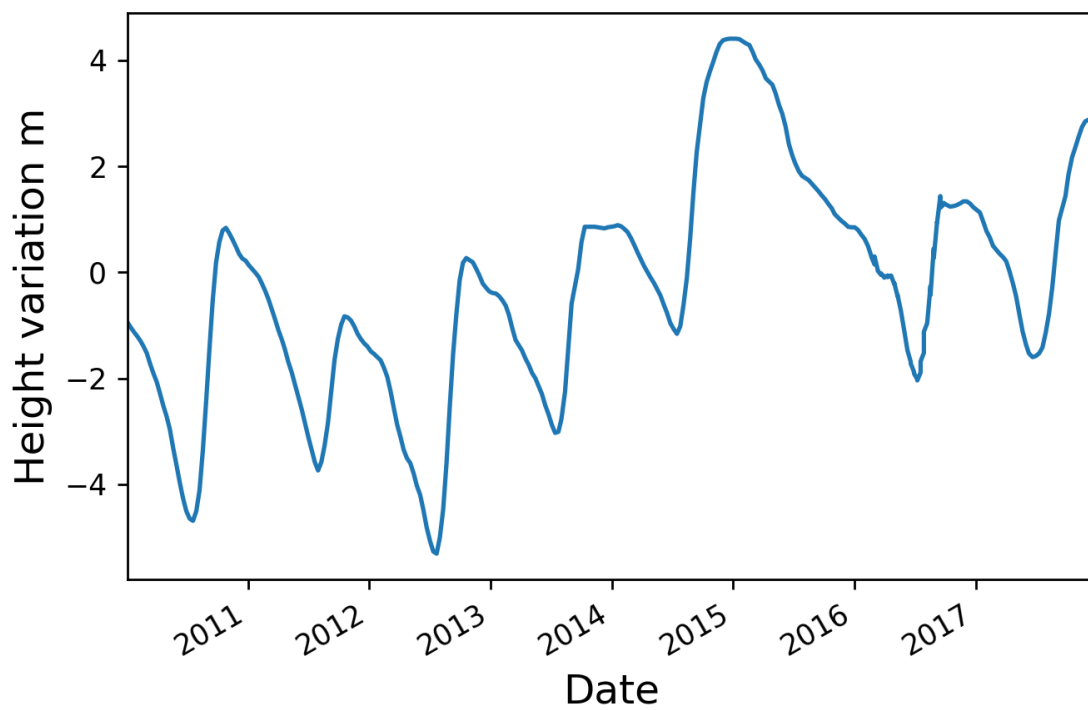


Figure S9: Lake Nasser water level from satellite radar and altimetry¹³. The water level generally drops and rises during midsummer, although 2015 seems anomalous in this trend.

References

- (1) EIA, EIA: International Energy Statistics. <http://eia.gov/beta/international/> (accessed 22 January 2021).
- (2) Bahgat, G. The impact of the Arab spring on the oil and gas industry in North Africa: a preliminary assessment. *The Journal of North African Studies* **2012**, *17*, 503–514.
- (3) Schilling, J.; Freier, K. P.; Hertig, E.; Scheffran, J. Climate change, vulnerability and adaptation in North Africa with focus on Morocco. *Agriculture, Ecosystems & Environment* **2012**, *156*, 12–26.
- (4) Chourghal, N.; Lhomme, J. P.; Huard, F.; Aidaoui, A. Climate change in Algeria and its impact on durum wheat. *Regional Environmental Change* **2016**, *16*, 1623–1634.
- (5) Salhi, A.; Martin-Vide, J.; Benhamrouche, A.; Benabdelouahab, S.; Himi, M.; Benabdelouahab, T.; Casas Ponsati, A. Rainfall distribution and trends of the daily precipitation concentration index in northern Morocco: a need for an adaptive environmental policy. *SN Applied Sciences* **2019**, *1*, 277.
- (6) Aieb, A.; Lefsih, K.; Scarpa, M.; Bonaccorso, B.; Cicero, N.; Mimeche, O.; Madani, K. Statistical modeling of monthly rainfall variability in Soummam watershed of Algeria, between 1967 and 2018. *Natural Resource Modeling* **2020**, *33*, e12288, eprint: <https://onlinelibrary.wiley.com/doi/pdf/10.1111/nrm.12288>.
- (7) Bastviken, D.; Cole, J.; Pace, M.; Tranvik, L. Methane emissions from lakes: Dependence of lake characteristics, two regional assessments, and a global estimate. *Global Biogeochemical Cycles* **2004**, *18*, eprint: <https://agupubs.onlinelibrary.wiley.com/doi/pdf/10.1029/2004GB002238>.
- (8) Wilkinson, J.; Maeck, A.; Alshboul, Z.; Lorke, A. Continuous Seasonal River Ebulli-

- tion Measurements Linked to Sediment Methane Formation. *Environmental Science & Technology* **2015**, *49*, 13121–13129.
- (9) Holgerson, M. A.; Raymond, P. A. Large contribution to inland water CO₂ and CH₄ emissions from very small ponds. *Nature Geoscience* **2016**, *9*, 222–226, Number: 3
Publisher: Nature Publishing Group.
- (10) Aguilera, E.; Vila-Traver, J.; Deemer, B. R.; Infante-Amate, J.; Guzmán, G. I.; González de Molina, M. Methane Emissions from Artificial Waterbodies Dominate the Carbon Footprint of Irrigation: A Study of Transitions in the Food-Energy-Water-Climate Nexus (Spain, 1900-2014). *Environmental Science & Technology* **2019**, *53*, 5091–5101, Publisher: American Chemical Society.
- (11) Artuso, F.; Chamard, P.; Piacentino, S.; di Sarra, A.; Meloni, D.; Monteleone, F.; Sferlazzo, D. M.; Thiery, F. Atmospheric methane in the Mediterranean: Analysis of measurements at the island of Lampedusa during 1995-2005. *Atmospheric Environment* **2007**, *41*, 3877–3888.
- (12) Dlugokencky, E.; Lang, P.; Crotwell, A.; Mund, J.; Crotwell, M.; Thoning, K. Atmospheric Methane Dry Air Mole Fractions from the NOAA ESRL Carbon Cycle Cooperative Global Air Sampling Network, 1983-2016, Version: 2017-07-28. 2017; ftp://aftp.cmdl.noaa.gov/data/trace_gases/ch4/flask/surface/, (accessed 19 January 2021).
- (13) Birkett, C.; Reynolds, C.; Beckley, B.; Doorn, B. In *Coastal Altimetry*; Vignudelli, S., Kostianoy, A. G., Cipollini, P., Benveniste, J., Eds.; Springer Berlin Heidelberg: Berlin, Heidelberg, 2011; pp 19–50.