

1 **Impact of the Asian Monsoon Anticyclone on the Variability of mid-**
2 **to-upper tropospheric methane above the Mediterranean Basin**

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24

24 Abstract

25 The space and time variabilities of methane (CH_4) total column and upper tropospheric mixing
26 ratios are analyzed above the Mediterranean Basin (MB) as part of the Chemical and Aerosol
27 Mediterranean Experiment (ChArME_x) programme. Since the analysis of the mid-to-upper
28 tropospheric CH_4 distribution from spaceborne sensors and model outputs is challenging, we have
29 adopted a climatological approach and have used a wide variety of datasets. We have combined
30 spaceborne measurements from the Thermal And Near infrared Sensor for carbon Observations-
31 Fourier Transform Spectrometer (TANSO-FTS) instrument on the Greenhouse gases Observing
32 SATellite (GOSAT) satellite, the Atmospheric InfraRed Spectrometer (AIRS) on the AURA
33 platform and the Infrared Atmospheric Sounder Interferometer (IASI) instrument aboard the
34 MetOp-A platform with model results from the Chemical Transport Model (CTM) MOCAGE, and
35 the Chemical Climate Models (CCMs) CNRM-AOCCM and LMDz-OR-INCA (according to
36 different emission scenarios). In order to minimize systematic errors in the spaceborne
37 measurements, we have only considered maritime pixels over the MB. The period of interest spans
38 from 2008 to 2011 considering satellite and MOCAGE data and, regarding the CCMs, from 2001 to
39 2010. Although CH_4 is a long-lived tracer with lifetime of ~ 12 years supposed to be well mixed in
40 the troposphere, an East-West gradient in CH_4 is observed and modelled in the mid-to-upper
41 troposphere with a maximum in the Western MB in all seasons except in summer when CH_4
42 accumulates above the Eastern MB. The peak-to-peak amplitude of the East–West seasonal
43 variation in CH_4 above the MB in the upper troposphere (300 hPa) is weak but almost twice greater
44 in the satellite measurements (~ 25 ppbv) than in the model data (~ 15 ppbv). The maximum of CH_4
45 in summer above the Eastern MB can be explained by a series of dynamical processes only
46 occurring in summer. The Asian monsoon traps and uplifts high amounts of CH_4 to the upper
47 troposphere where they build up. The Asian Monsoon Anticyclone redistributes these elevated CH_4
48 amounts towards North Africa and Middle East to finally reach and descent in the Eastern MB. In

49 the lower troposphere, the CH₄ variability is mainly driven by the local sources of emission in the
50 vicinity of the MB.

51

51 **1. Introduction**

52 During the last decades, major concerns have been caused by the impact and the role that
53 atmospheric trace gases play in climate and air pollution changes and the crucial issue was to
54 evaluate. In IPCC (2007), the ongoing changes of our atmosphere (composition, climate, air
55 pollution, radiation) are reported. Among trace gases, methane (CH₄), carbon dioxide (CO₂), and
56 nitrous oxide (N₂O) are predominant constituents which play an important role in atmospheric
57 changes because they are strongly influenced by human activities. In the frame of predicting the
58 future of the Earth's climate (IPCC, 2007), knowledge of today's CO₂, CH₄ and N₂O sources and
59 sinks, spatial distribution and time variability is essential and this study will be dedicated to CH₄.

60 The net positive radiative impact of the human activity on climate, starting from 1750, has been
61 evaluated to 1.6 [+0.6 to +2.4] Wm⁻² (IPCC, 2007). In the atmosphere, the long-lived greenhouse
62 gases account for 2.63 ± 0.26 Wm⁻² and are the predominant radiative terms. CO₂, with
63 tropospheric lifetime of 30–95 years, has a radiative efficiency of 1.4×10^{-5} Wm⁻²ppb⁻¹, but CH₄ and
64 N₂O, with tropospheric lifetimes of 12 and 114 years, respectively, are intensely more efficient by
65 3.7×10^{-4} and 3.03×10^{-3} Wm⁻²ppb⁻¹, respectively. IPCC (2007) estimated CH₄ and N₂O to be
66 responsible of 0.48 [+0.43 to 0.53] and 0.16 [+0.14 to 0.18] Wm⁻², respectively in the radiative
67 forcing changes.

68 The Mediterranean Basin (MB) is located in a transitional zone between subtropical and mid-
69 latitudes regimes (Lionello, 2012), highly sensitive to climate change. To illustrate, simulations
70 tend to show a pronounced decrease in precipitation (2000-2100), especially in the warm season
71 (Giorgi and Lionello, 2008), and Lionello (2012) reported on an observed summer West-East
72 dissymmetry in precipitation (1979-2002). In terms of anthropogenic pollution sources, the MB is
73 at the confluence of three continents, Europe, Africa and Asia. The impact of these distinct
74 continental sources such as from manufactures and densely populated coastal areas (e.g. Marseille,
75 Barcelona, Athens, Tunis, Cairo, Genoa or Roma) or forest fires (e.g. South East of France, Corsica,
76 Portugal, Greece) is still not well understood, especially on the O₃ and CO budgets. Besides these

77 regional sources, polluted air masses may originate from Asia during the summer monsoon period,
78 Africa through the Hadley cell and upper level anticyclone and North America through the
79 westerlies. The “Expérience sur Site pour CONtraindre les Modèles de Pollution atmosphérique et
80 de Transport d'Emissions” (ESCOMPTE) campaign (June-July 2001) in southeastern France aimed
81 to characterize the summer time pollution events over there (Cros et al., 2004). The goal of the
82 Mediterranean Intensive Oxidant Study (MINOS) campaign (July-August 2001) in the eastern
83 Mediterranean was to measure long-range transport of air pollution and aerosols from South East
84 Asia and Europe towards the MB (Ladstätter-Weissenmayer et al., 2003; Scheeren et al., 2003).
85 They have demonstrated the importance of coastal and synoptic transport mechanisms on the
86 variability of constituents but were not adapted to assess the budgets of O₃, CO and long-lived
87 species.

88 The ChArMEx (Chemistry and Aerosol Mediterranean Experiment) Project
89 (<http://charmex.lsce.ipsl.fr/>) is the atmospheric chemistry component of a large multidisciplinary
90 Mediterranean regional program proposed and conducted by France. It intends, among other
91 objectives, to quantify processes explaining the temporal evolution of chemical compounds and
92 aerosols in the troposphere above the Mediterranean Basin (MB). To achieve these goals over the
93 first phase (2010-2015), the program uses data from satellites, ground-based, sondes, aircraft,
94 models and assimilation in order to evaluate 1) the variabilities and recent trends of several species
95 (e.g. O₃, CO, N₂O) and aerosols, 2) the synoptic-scale circulation that controls their transport, and
96 3) the future chemical climate over the MB by 2100.

97 We have to keep in mind that it is rather challenging 1) to measure long-lived species from space
98 focusing on tropospheric layers and 2) to model these molecules over the entire troposphere.
99 Concentrations of long-lived atmospheric chemical species (CO₂, CH₄, and N₂O) are expected to be
100 very well-mixed and evenly distributed throughout the lower atmosphere. The past/present nadir-
101 viewing instruments able to actually measure CH₄ in the troposphere have been/are:

102 1) the Interferometric Monitor for Greenhouse gases (IMG) instrument operating in the Thermal
103 Infrared (TIR) aboard the ADvanced Earth Observing Satellite (ADEOS-1) platform in 1996-1997
104 (Clerbaux et al., 1998);

105 2) the near-IR (NIR) Scanning Imaging Absorption Spectrometer for Atmospheric Chartography
106 (SCIAMACHY) aboard the ENVIronment SATellite (ENVISAT) platform (Buchwitz et al., 2000)
107 from 2002 to 2012;

108 3) the Tropospheric Emission Spectrometer (TES) operating in the TIR aboard the Aura platform
109 (Worden et al., 2012) from 2004 to date;

110 4) the Thermal And Near infrared Sensor for carbon Observations – Fourier Transform
111 Spectrometer (TANSO-FTS) on the Greenhouse gases Observing SATellite (GOSAT) platform
112 (Yokota et al., 2009) both in the Short-Wave InfraRed (SWIR) and in the TIR from 2008 to date;

113 5) the Atmospheric InfraRed Sounder (AIRS) aboard the Aqua platform (Xiong et al., 2008)
114 measuring in the TIR from 2004 to date;

115 6) the Infrared Atmospheric Sounding Interferometer (IASI) instrument aboard the MetOp-A
116 and -B platforms (Hilton et al., 2012) operating in the TIR from 2008 to date, and aboard the
117 MetOp-C platform expected to be launched in 2016.

118 Table 1 synthesizes the above mentioned information and shows the nadir-viewing instrument
119 capability to measure tropospheric CH₄. The sensitivity of the TIR to measure CH₄ is rather weak
120 except on areas showing a high thermal contrast at the surface (vertical gradient of temperature
121 between the surface and the lowermost planetary boundary layer) as the ones encountered over the
122 tropics (Crevoisier et al., 2013) contrarily to the measurements performed in the SWIR (Yoshida et
123 al., 2013). In the NIR, analyses are essentially restricted to areas over land because the retrievals
124 over sea are considered less reliable due to fairly low surface albedo of water, which results in low
125 signals and thus in low signal-to-noise ratios (Georgoulias et al., 2011).

126 In parallel to the satellite data, models have also been used in order to assess the variability,
127 sources and sinks, and future trends of the long-lived species. Examples are: CH₄ emission and flux

128 estimates at global scales (Bergamaschi et al., 2009; Bousquet et al., 2011), future evolution of
129 long-lived species included in the international Atmospheric Chemistry and Climate Model
130 Intercomparison Project (ACCMIP) involving more than 10 different models (Lamarque et al.,
131 2013).

132 Numerous studies have examined the variabilities of atmospheric compounds above the MB to
133 highlight the processes (sources and sinks) associated by coupling surface, balloon-borne, airborne,
134 spaceborne measurements with models results at different scales, from mesoscales to global scales.
135 Constituents are for instance aerosols (Nabat et al., 2012), radionuclides (Masson et al., 2010),
136 ozone (Liu et al., 2009), carbon monoxide (Drori et al., 2012). The impact of 1) the different
137 meteorological regimes and 2) the seasonal variabilities of the emissions of atmospheric
138 constituents, e.g. CO emitted from fires in summers, produces a seasonal variation in all the
139 constituents. It also produces a longitudinal gradient between the Eastern and the Western MB,
140 together with a seasonal variation in the gradient. For example, European anthropogenic emissions
141 were found to significantly influence the Eastern MB surface CO concentrations, while European
142 biomass burning emissions were found to have only a small impact on Eastern MB surface CO
143 concentrations (Drori et al., 2012). Total columns of CH₄ as measured by SCIAMACHY over land
144 and the Eastern Mediterranean from 2003 to 2004 show latitudinal and seasonal variations that
145 cannot be attributed to volcano eruptions (Georgoulias et al., 2011).

146 The aim of the present paper is to assess the variability of CH₄ in the mid-to-upper troposphere
147 between the East and the West of the Mediterranean Basin and to attribute the seasonal variability
148 of the East-West gradient to different processes at both, synoptic and global scales depending on the
149 season and the altitude layer considered. We will study in detail the impact of the summer-time
150 long-range transport of CH₄ from Asia to the Eastern MB through the Asian Monsoon Anticyclone.
151 Since we have already underlined that measurement and modeling of the tropospheric CH₄
152 distribution are challenging, we will adopt a climatological approach and will use a wide variety of
153 space-borne measurements and model outputs to verify that they give consistent results. Our

154 approach is thus based on three different models: the CTM MOCAGE (Josse et al., 2004) and the
155 two Chemical Climate Models (CCMs) CNRM-AOCCM from Météo-France (Huszar et al., 2013)
156 and LMDz-OR-INCA (Hauglustaine et al., 2004; Szopa et al., 2013) from the Laboratoire des
157 Sciences du Climat et de l'Environnement (LSCE). To complete, we have considered the CH₄
158 profiles from AIRS and GOSAT, and the CH₄ total columns from IASI.

159 The manuscript is structured as follow. In section 2, we briefly present the spaceborne
160 instruments and datasets involved in this study, namely MetOp-A/IASI, AQUA/AIRS and
161 GOSAT/TANSO together with the models, namely MOCAGE, CNRM-AOCCM and LMDz-OR-
162 INCA. The meteorology and climatology of CH₄ inferred from the different datasets above the MB
163 are discussed in section 3. The CH₄ variability both in the East and in the West of the MB is
164 presented in section 4. A detailed discussion of the different processes involved in the CH₄
165 variability above the MB is presented in section 5 underlining the impact of the Asian Monsoon
166 Anticyclone to the distribution of the mid-to-upper CH₄ in the Eastern MB. Finally, section 6
167 concludes the paper.

168

169 **2. The datasets**

170 2.1. The satellite data

171 Our CH₄ study from satellite data analyses measurements from three different spaceborne TIR
172 sensors (IASI, AIRS and GOSAT) by only considering the pixels over the Mediterranean Sea in
173 order to minimize the systematic biases among pixels (day vs. night, land vs. sea). The sensitivity of
174 TIR retrievals strongly depends on surface parameters: emissivity, temperature and thermal contrast
175 (Claeyman et al., 2011). Compared to land surface, sea surface emissivity is relatively smaller in
176 magnitude and more spatially uniform. Sea surface temperature exhibits a diurnal amplitude weaker
177 than land surface temperature. Therefore, the vertical sensitivity of the TIR measurements, defined
178 as the full-width at half-maximum of the averaging kernels from the optimal estimation method
179 (Rodgers, 2000), over the sea is consistent during day and night and concentrated in the mid-

180 troposphere. Over the land, the vertical sensitivity is, on average, lower in the middle troposphere
181 during the day than during the night, depending on the actual value of the thermal contrast at the
182 surface.

183 Infrared sounders measurement errors can be rather large, e.g. up to 10% or more for a single
184 CH₄ total column IASI pixel (Turquety et al., 2004). Thus, by the use of time (monthly/seasonally)
185 and geographical averages, including more than a thousand measurements, we can lower the
186 random error to less than 1%. Systematic errors, if any, will of course be unchanged. For that
187 reason, our analysis relies on a differential method to highlight the CH₄ variability by considering
188 the difference between the Eastern MB (EMB) and the Western MB (WMB), assuming that the
189 systematic errors are of the same order of magnitude (although partially unknown) within each
190 geographical box that will be defined in section 3.

191

192 *2.1.1. The IASI data*

193 IASI, on board of MetOp-A, was launched in 2006 by the European Organisation for the
194 Exploitation of Meteorological Satellites (EUMETSAT). More specifications on platform and
195 instrument can be found on <http://smc.cnes.fr/IASI> and [http://www.eumetsat.int/Home/
196 Main/Satellites/Metop/Instruments/SP_2010053151047495](http://www.eumetsat.int/Home/Main/Satellites/Metop/Instruments/SP_2010053151047495)). The retrieval algorithm for CH₄ is
197 based on the neural network theory adapted from Turquety et al. (2004). The retrieval method is
198 embedded in the operational IASI Level 2 product processing facility at EUMETSAT
199 (EUMETSAT, 2004; Schlüssel et al., 2005, August et al., 2012). From the spectral bandwidth
200 1230-1347 cm⁻¹, the estimated accuracy of the CH₄ total column is about 2% and the estimated
201 precision is of the order of 10% (Turquety et al., 2004). The true accuracy cannot be stated without
202 reference to independent means of comparison, which are not available so far. Consequently, we
203 consider a random Gaussian error of ~10% associated with each single pixel of retrieved total
204 column of CH₄. At mid-latitudes, the vertical sensitivity of the total column CH₄ is peaking in the
205 mid-troposphere at ~8 km from 4 to 14 km (Razavi et al., 2009) and, in the tropics, at ~10 km from

206 5 to 15 km. Geophysical level 2 pre-operational data are provided by EUMETSAT (from version 4
207 to version 5 from 2008 to 2011). The CH₄ products, not yet validated, are only experimental
208 products, routinely generated for demonstration and evaluation. Note, the number of daily total
209 columns of CH₄ averaged in a 1°x1° bin is highly variable because of cloud-free IASI
210 considerations. The monthly-averaged IASI data within each of the East and West areas defined in
211 section 3 represent an average of 30000-70000 pixels depending on the month considered.

212

213 *2.1.2. The AIRS data*

214 AIRS is onboard the space platform NASA EOS Aqua, launched in 2002
215 (<http://airs.jpl.nasa.gov/>). AIRS measures approximately 200 channels in the 7.66 μm absorption
216 band of CH₄, of which 71 channels are used to retrieve CH₄. A detailed description of the retrieval
217 algorithm can be found in Susskind et al. (2011). Note, the averaging kernels provided by NASA
218 will be considered further (section 4.2) in order to degrade the vertical resolution of the model
219 outputs. At mid-latitudes, the most sensitive layer of AIRS channels to CH₄ is at 300 hPa (~9 km)
220 with a vertical sensitivity from 700 to 100 hPa (Xiong et al., 2008), and, in the tropics, at 200 hPa
221 from 500 to 70 hPa consistently with the IASI TIR measurement sensitivity. Around 200-300 hPa,
222 considering the version V5 used in the present analysis (Xiong et al., 2008), the precision of AIRS
223 CH₄ is estimated to be 30 ppbv (1.7%) and validation using in situ aircraft measurements shows
224 that the accuracy of the retrieved CH₄ is 0.5–1.6%. Daily maritime profiles of CH₄ have been
225 averaged in 1°x1° bins over the MB. The monthly-averaged AIRS data within each of the East and
226 West areas defined in section 3 represent an average of 6000-10000 vertical profiles depending on
227 the month considered.

228

229 *2.1.3. The GOSAT data*

230 The Japanese Aerospace Exploration agency (JAXA) launched the GOSAT platform in 2009,
231 with the TANSO-FTS spectrometer, a nadir-viewing instrument designed for greenhouse gases

232 research, CO₂ and CH₄, operating in the TIR and SWIR domains [0.7-14.3 μm] (Kuze et al., 2009).
233 More specifications on platform and instrument can be found on <http://www.gosat.nies.go.jp/>. The
234 sensitivity of the SWIR CH₄ measurements at 1.67 μm (Yokota et al., 2009) at mid-latitudes over
235 the sea is very weak, thus few meaningful pixels could have been retrieved preventing the use of
236 such information in our analysis. The TIR measurements from Band 4 (5.5–4.3 μm) provide
237 vertical profiles of CH₄ along 7 vertical levels (Imasu et al., 2007) by using the optimal estimation
238 method with a vertical sensitivity in the tropics peaking at 10 km (higher than at mid-latitudes) from
239 5 to 15 km (Saitoh et al., 2012), consistently with the vertical sensitivity of IASI (Razavi et al.,
240 2009) and AIRS (Xiong et al., 2008) in the tropics. A selection by using Degree of Freedom of
241 Signal (DFS) is applied for the data having DFS values larger than 0.6 for CH₄. TIR data (L2
242 Version 0.10) were only available from 16 March to 24 November 2010 from the GOSAT User
243 Interface Gateway at the time the analysis has been performed. These retrievals provide vertical
244 profiles of mixing ratio of CH₄ from 1000 to 100 hPa. Comparisons with aircraft measurements
245 show that the average difference between the GOSAT (TIR) and aircraft CH₄ values (TIR –
246 aircraft) is –5 ppbv, and the 1σ standard deviation is 15 ppbv (Saitoh et al., 2012). Daily maritime
247 profiles of CH₄ have been averaged in 1°x1° bins over the MB. The monthly-averaged GOSAT data
248 within each of the East and West areas defined in section 3 represent an average of 100-300 vertical
249 profiles depending on the month considered, namely 20-30 times less than for AIRS.

250

251 2.2. The model data

252 2.2.1. The MOCAGE data

253 MOCAGE (MOdèle de Chimie Atmosphérique à Grande Echelle) (Peuch et al., 1999) is a 3D
254 CTM which covers the planetary boundary layer, the free troposphere, and the stratosphere for
255 different applications such as: operational chemical weather forecasting (Dufour et al., 2005);
256 tropospheric and stratospheric research studies (Claeyman et al., 2010; Ricaud et al., 2009); and
257 data assimilation research (El Amraoui et al., 2010; Claeyman et al., 2011). In our study, MOCAGE

258 is forced dynamically by wind and temperature fields from the analyses of the ARPEGE model
259 (Courtier et al., 1991). The MOCAGE horizontal resolution is $2^\circ \times 2^\circ$ and the model uses a semi-
260 Lagrangian transport scheme. It includes 47 levels from the surface up to 5 hPa with a vertical
261 resolution of about 800 m around the tropopause, 400-800 m in the troposphere and 40-400 m in the
262 7 levels of the boundary layer. Chemistry used within MOCAGE is a combination of tropospheric
263 (RACM described in Stockwell et al., 1997) and stratospheric (REPROBUS described in Lefèvre et
264 al., 1994) chemical schemes. Initial chemical conditions are taken from climatological fields over a
265 spin-up period of 3 months allowing the model to quickly bring chemical fields to realistic spatial
266 distributions. Surface emissions prescribed in MOCAGE are based upon yearly- or monthly-
267 averaged climatologies. More precisely, the CH_4 surface emissions are monthly averages and split
268 into anthropogenic sources taken from the Intergovernmental Panel on Climate Change (IPCC)
269 (Dentener et al., 2005), biomass burning (van de Werf et al., 2003) and biogenic sources (Michou
270 and Peuch, 2002). The CH_4 climatologies are representative of year 2000 for a total emission rate of
271 $534 \text{ Tg}(\text{CH}_4) \text{ yr}^{-1}$.

272

273 2.2.2. *The CNRM-AOCCM data*

274 The atmospheric model embedded in CNRM-AOCCM is presented in Huszar et al. (2013)
275 based on the Atmosphere-Ocean General Circulation Model (AOGCM) CNRM-CM5 described in
276 Voldoire et al. (2012). The main difference between CNRM-CM5 and CNRM-AOCCM resides in
277 the “online” coupling with a stratospheric chemistry which is based on the REPROBUS scheme.
278 This scheme is applied on the whole vertical column, except between the surface and the 560 hPa
279 level where long-lived chemical species are relaxed towards global average surface value following
280 the A1B scenario from IPCC (2007). The A1B scenario mainly describes a future world of very
281 rapid economic growth, global population that peaks in mid-century and declines thereafter, and the
282 rapid introduction of new and more efficient technologies. Convection of species is not considered.
283 In this chemistry version, the 3-D distribution of the seven absorbing gases (H_2O , CO_2 , O_3 , CH_4 ,

284 N₂O, CFC11, and CFC12) is then provided by the chemistry module of CNRM-AOCCM and
285 interacts with the radiative calculations. More details can be found in Michou et al. (2011). In the
286 present version, there are about 50 chemical species, and the horizontal resolution is 2.8°x2.8°.
287 Distribution of atmospheric constituents at the surface are zonally symmetric below 500 hPa (Fig.
288 10) and greenhouse gases follow the A1B scenario on atmospheric chemistry and climate for the
289 period of 1940-2100. In the present analysis, for this model, we only consider the climatological
290 period 2001-2010.

291

292 2.2.3. *The LMDz-OR-INCA data*

293 The Interaction between Chemistry and Aerosol (INCA) model is used to simulate the
294 distribution of aerosols and gaseous reactive species in the troposphere. In the present
295 configuration, the model includes 19 hybrid vertical levels extending up to 4 hPa, and a horizontal
296 resolution of 1.9° in latitude and 3.75° in longitude. INCA is coupled online to the LMDz General
297 Circulation Model (GCM) to account, with different degrees of complexity, for climate chemistry
298 interactions. In the simulations described here, LMDz is coupled with the ORCHIDEE (Organizing
299 Carbon and Hydrology in Dynamic Ecosystems) dynamic global vegetation model (Krinner et al.,
300 2005) for soil/atmosphere exchanges of water and energy (Hourdin et al., 2006), but not for
301 biogenic CO₂ or Volatile Organic Compounds (VOCs) fluxes. Together, these three models form
302 the LMDz-OR-INCA model. Fundamentals for the gas phase chemistry are presented in
303 Hauglustaine et al. (2004) and first results with the full tropospheric gaseous chemical scheme are
304 presented by Folberth et al. (2006). The model includes 223 homogeneous chemical reactions, 43
305 photolytic reactions and 6 heterogeneous reactions including non-methane hydrocarbon oxidation
306 pathways and aerosol formation. The LMDz-OR-INCA simulation covers four future projections of
307 emissions for the 2000–2100 period. The Representative Concentration Pathways (RCP) emissions
308 are used (Lamarque et al., 2011). They correspond to emission trajectories compatible with the
309 evolution of radiative forcing equivalent in 2100 to 2.6, 4.5, 6.0 and 8.5 Wm⁻² relative to pre-

310 industrial values (labelled therein after RCP 2.6, 4.5, 6.0 and 8.5). In the present analysis, for this
311 model, we only consider the climatological period 2001-2010.

312

313 **3. Atmospheric conditions controlling the spatial distribution of methane**

314 Figure 1 shows the CH₄ fields calculated by MOCAGE for summer (June-July-August, JJA)
315 2009 over the MB at 850, 500 and 200 hPa, superimposed with the wind fields from the ARPEGE
316 analyses averaged over the same period. Figure 2 presents the CH₄ vertical distribution as
317 calculated by MOCAGE in summer 2009 along an East-West axis above the MB. Similarly to
318 Figures 1-2, the Figures 3-4 present, in winter (December-January-February, DJF) 2009, the CH₄
319 fields as calculated by MOCAGE over the MB at 850, 500 and 200 hPa, and along an East-West
320 axis, respectively. On Figures 2 and 4, the MOCAGE CH₄ fields are superimposed with 1) the wind
321 fields from ARPEGE analyses and 2) the cold point tropopause pressure fields provided by the
322 National Oceanic and Atmospheric Administration (NOAA) National Centers for Environmental
323 Prediction (NCEP)/National Center for Atmospheric Research (NCAR) reanalyses, all these data
324 being averaged over the same period.

325 Considering the meteorology of the MB, we observe two different regimes. 1) In winter, and
326 more generally from autumn to spring (not shown), from the boundary layer to the upper
327 troposphere, air masses are essentially coming from either Europe or Eastern Atlantic Ocean. 2) In
328 summer, the meteorology of EMB and WMB is more complex and depends on the altitude
329 considered.

330 In the planetary boundary layer, cells develop in the WMB, and air masses come from Europe,
331 Northern Africa and Eastern Atlantic Ocean, whilst in the EMB, air masses are originated from four
332 major source regions: i) long fetch of maritime European air masses from NW throughout the whole
333 year, ii) North east continental flow originating in south Eastern Europe (Etesian winds) in summer,
334 iii) South-east flow from the Arabian Peninsula occurring in the fall, and iv) South-west flow along
335 the North-African coast most frequent during late winter and spring (Dayan, 1986). In the middle

336 troposphere, whatever the season, air masses are essentially coming from the west for both parts of
337 the basin. In summer, upper tropospheric air masses in the WMB are essentially coming from the
338 West, but in the EMB, they are also originated from Northern Africa and the Arabic Peninsula (Ziv
339 et al., 2004; Liu et al., 2009), and even farther away, from Asia (we will discuss this point in
340 sections 4 and 5). Note that, in summer, the EMB and WMB are also affected by the location of the
341 descending branch of the Hadley cell. These summer climatologies are all consistent with Millán et
342 al. (1997), Lelieveld et al. (2002), Ziv et al. (2004) and Schicker et al. (2010).

343 Seasonally-averaged wind fields from ARPEGE analyses show two different regimes in the
344 surface pressure values during the summer and the winter periods. During the summer in the WMB,
345 there is a higher pressure regime than in the EMB. In the lowermost troposphere (850 hPa), an
346 anticyclonic cell develops in the WMB that has an impact on the distribution of CH₄ by producing a
347 local minimum (Fig. 1). At 850 hPa, air masses are coming from Europe, North Africa and the
348 Atlantic Ocean. The CH₄ distribution shows a maximum over Europe, consistently with the
349 strongest emission zones (Fig. 10), and a strong minimum over North Africa. In the mid-
350 troposphere (500 hPa), air masses are coming from Europe, and the Atlantic Ocean. An East-West
351 gradient is detected with more CH₄ on the EMB. In the upper troposphere (200 hPa), air masses are
352 originated from the Atlantic Ocean (even North America) and from North Africa and Asia
353 producing over the MB an obvious North-South gradient with more CH₄ in the South (upper
354 troposphere) than in the North (lower stratosphere) attributed to the impact of long-range transport
355 of pollutants (as discussed in section 5). A systematic subsidence is present over the MB (Fig. 2)
356 whatever the longitudinal bin considered due to the presence of semi-permanent subtropical high
357 pressure systems which are centred over the tropical deserts. More precisely, in the WMB, the
358 descent is caused by the presence of a high pressure cell (Fig. 1) whilst, in the EMB, it is coming
359 from the Hadley cell that is further displaced over the Northern Africa producing a downward
360 branch in the area 30°N-35°N. The tropopause moves up from ~200 hPa in the WMB to ~175 hPa
361 in the EMB. The CH₄ distribution shows 1) an obvious transition at the tropopause and 2) a

362 minimum in the West and a maximum in the East in the low troposphere accentuated by the
363 systematic descent in the Eastern MB that brings CH₄-enriched air masses from the upper
364 troposphere to the mid-to-low troposphere.

365 In winter, the meteorological condition of the MB is much more homogeneous with westerlies
366 blowing whatever the pressure considered from 850 to 200 hPa (Figs. 3-4). North-South (and to a
367 lesser extent East-West) gradients in CH₄ can also be detected (Fig. 3) associated to the local
368 sources of emission over Europe at 850 hPa and to the stratosphere/troposphere transition at 200
369 hPa. The 500-hPa layer is a transition region between the low and the upper troposphere with
370 minima of CH₄ over North Africa and a cell of high CH₄ in the WMB. Contrarily to summer, since
371 the temperature of the Mediterranean Sea is greater than that of the surrounding continents, a
372 systematic upward motion is present (Fig. 4) whatever the longitudinal bin considered. The Hadley
373 cell is further displaced to the South (latitude < 30°N) and its downward branch does not affect
374 significantly the EMB. The tropopause pressure is rather stable from the WMB to the EMB, around
375 260 hPa. The CH₄ distribution shows minimum in the lowermost troposphere and a maximum in
376 the middle troposphere.

377

378 **4. CH₄ variability**

379 *4.1. CH₄ spatial distribution of the MB*

380 Figure 5 shows the distributions of 1) the CH₄ total columns from IASI over the MB averaged in
381 summer 2009 to compare with the MOCAGE results in time coincidence, and 2) the CH₄ mixing
382 ratios from AIRS at 260 hPa over the MB averaged in summer 2009 to compare with the
383 MOCAGE results at 200 hPa in time coincidence. The measured and modelled data are selected
384 only for the maritime pixels within the boxes [36°N-45°N, 1°E-12°E] and [30°N-37°N, 26°E-37°E]
385 to represent the WMB and the EMB (blue squares in each figure), respectively.

386 Due to its long lifetime (~12 years), CH₄ is considered as a well-mixed species in the
387 troposphere. Nevertheless the CH₄ spatial distribution over the MB in summer (JJA) 2009 shows

388 some gradients both in the East-West and the North-South directions. Indeed, in the middle
389 troposphere (inferred from the sensitivity of the IASI total columns) and in the upper troposphere
390 (200-260 hPa), an East-West gradient is observed in the model and satellite data of ~60 ppbv (~4%)
391 in total column and ~30-150 ppbv (~2-9%) in mixing ratio. A North-South gradient is also detected
392 in the MOCAGE and AIRS data but not in the IASI data set. Therefore, there is systematically a
393 maximum of CH₄ from the middle to the upper troposphere in the East of the MB compared to the
394 West. In the mid-to-upper troposphere, these East-West gradients are not originated from the CH₄
395 sources more intense in Europe than in Northern Africa or in Middle Asia (Fig. 1) but rather from
396 the long-range transport of Asian-origin air masses and the subsidence of air masses in the EMB
397 (Figs. 1-2 and detailed discussion in section 5).

398 Quantitatively, there is a positive bias in MOCAGE vs. IASI of less than 30 ppbv (2%) in CH₄
399 total column mixing ratio. The East-West gradient is rather consistent between IASI and MOCAGE
400 but the North-South modelled gradient is not detected in the IASI data set. In the upper troposphere
401 (200-260 hPa), MOCAGE and AIRS CH₄ mixing ratios are very consistent with gradients more
402 accentuated in the model (~150 ppbv) than in the AIRS data sets (~30 ppbv). A systematic negative
403 bias of MOCAGE compared to AIRS of ~100 ppbv up to 150 ppbv (10%) in the Northern MB is
404 detected. We discuss in the next section the consistency of the vertical profiles of CH₄ as measured
405 by the different spaceborne sensors and as calculated by MOCAGE together with the associated
406 biases.

407

408 *4.2. Vertical profiles*

409 Figure 6 shows the vertical profiles of CH₄ as measured by AIRS (750-100 hPa) and GOSAT
410 (1000-100 hPa) and as calculated by MOCAGE (1000-100 hPa) averaged over the EMB and the
411 WMB depending on the four seasons: winter (DJF), spring (March-April-May, MAM), summer
412 (JJA) and autumn (September-October-November, SON) 2010. There is a good agreement to within
413 20-30 ppbv between AIRS and GOSAT data in the vertical domain 750-200 hPa. GOSAT is

414 systematically greater than AIRS by about 20-30 ppbv for pressure greater than 300 hPa, whilst, for
415 pressure less than 300 hPa, AIRS is systematically greater than GOSAT by 20 ppbv degrading to
416 50-200 ppbv at 100 hPa. But the shape of the vertical spaceborne profiles is consistent between
417 AIRS and GOSAT. Separately, whatever the season considered, the MOCAGE low-to-mid
418 tropospheric CH₄ is low biased compared to the measured profiles by ~150-200 ppbv. Furthermore,
419 the MOCAGE vertical profiles systematically show a maximum at 300 hPa, that is not present in
420 any of the spaceborne measurements, and a strong decrease above.

421 In order to assess the impact of the vertical sensitivity of the spaceborne measurements to the
422 CH₄ profiles, we have applied the AIRS averaging kernels, derived from the AIRS retrieval method
423 (Susskind et al., 2011) and provided by NASA for each AIRS pixel, to the profiles calculated by
424 MOCAGE. Note that the AIRS a priori vertical profiles are not used in our study since we are only
425 interested in the vertical shape of the CH₄ profile and not the absolute amount of CH₄. Degrading
426 the vertical resolution of the MOCAGE profiles by the convolution of averaging kernels (Fig. 6)
427 does show a strong impact on the vertical shape of the CH₄ profiles since the strong maximum at
428 300 hPa is no longer present. Convolved MOCAGE CH₄ profiles are now consistent with AIRS
429 CH₄ profiles whatever the season considered but a systematic low bias of ~150-200 ppbv (8-10%)
430 between AIRS and MOCAGE convolved profiles is observed. This might be due to the fact that no
431 a priori information contributes to the convolved profile. This is also due to the overall
432 underestimation of CH₄ by global models. Indeed, due to coarse horizontal resolution and large
433 uncertainties in the estimated surface emissions, tropospheric CH₄ lifetimes, e.g. evaluated by the
434 multi-model intercomparison project ACCMIP, are about 5-13% lower than observation estimates
435 (Naik et al., 2013; Voulgarakis et al., 2013).

436 Along the vertical, it is almost impossible to validate the spaceborne profiles with an external
437 data set since, even within the Total Carbon Column Observing Network
438 (<http://www.tccon.caltech.edu/>) giving accurate and precise column-averaged abundances of CH₄
439 (Wunch et al., 2010) because no measurement sites are unfortunately available in the vicinity of the

440 MB. Near the surface, the amount of CH₄ is about 1700-1750 ppbv for MOCAGE, and is on
441 average less than the CH₄ GOSAT data by about 150-200 ppbv. At this stage, it is worthwhile
442 considering surface data within the MB. The NOAA Earth System Research Laboratory (ESRL) In
443 Situ Methane Measurements provide some surface CH₄ measurements within and/or in the vicinity
444 of the MB: Lampedusa, Italy (35.52°N, 12.62°E, 45 amsl), Centro de Investigacion de la Baja
445 Atmosfera (CIBA), Spain (41.81°N, 4.93°W, 845 amsl) and Negev Desert, Israel (30.86°N,
446 34.78°E, 477 amsl). On average, these three sites indicate (not shown) a surface CH₄ annual mean
447 of about 1 875 ppbv in 2010, with an annual oscillation of ~20 ppbv amplitude. Consequently, the
448 amount of surface CH₄ in the MOCAGE run for 2010 is actually low biased by about 150-200 ppbv
449 (8-10%) but is very consistent with the LMDz-OR-INCA surface data of ~1725-1750 ppbv over the
450 Mediterranean (Fig. 10). The slight differences between the EMB and the WMB according to the
451 season and height are studied in detail in the next sub-section.

452

453 *4.3. The East-West seasonal variations: measured and calculated differences*

454 The seasonal variations of the differences in CH₄ fields between the EMB and the WMB (i.e.,
455 EMB minus WMB, labelled as “E–W”) as measured by AIRS, GOSAT and IASI and as calculated
456 by LMDz-OR-INCA, CNRM-AOCCM and MOCAGE are presented in Figure 7 when considering
457 the upper troposphere (AIRS, GOSAT, LMDz-OR-INCA and CNRM-AOCCM at 260 and 300
458 hPa) and the middle troposphere (IASI and MOCAGE total column mixing ratios). In the middle
459 and upper troposphere (Fig. 7), despite the fact that spaceborne measurements and modelling of
460 CH₄ are challenging, the modelled and measured seasonal variations of E–W are consistent to each
461 other showing a maximum (peak) in summer and a wide minimum in winter.

462 If we consider the time evolution of the total column mixing ratios (namely focussing on the
463 middle troposphere), we note that both MOCAGE and IASI show a maximum in summer, although
464 3 times greater in MOCAGE (~60 ppbv) than in IASI (~20 ppbv) in July and August. The
465 minimum in January-February is close to zero but slightly positive in October (5-10 ppbv). The

466 much stronger maximum in August calculated by MOCAGE compared to IASI CH₄ total columns
467 might be attributable to the sensitivity of spaceborne measurements in the middle troposphere
468 whilst the MOCAGE tropospheric columns cover the entire troposphere from the surface to the top
469 of the model atmosphere, namely 5 hPa.

470 In the upper troposphere (300 hPa), the spaceborne instrument datasets show a E–W maximum
471 in summer of ~12 ppbv in August for AIRS and a E–W wide maximum of ~5 ppbv in July-
472 September for GOSAT. A E–W peak of ~10 ppbv in July-August is also calculated by CNRM-
473 AOCCM although, in the LMDz-OR-INCA dataset, the E–W maximum is slightly positive in
474 August (~2 ppbv). The minimum in the satellite datasets is observed in March-April and is negative
475 (from –15 to –20 ppbv) consistently with the LMDz-OR-INCA dataset whilst the CNRM-AOCCM
476 E–W minimum is less intense (–6 ppbv in February and April). The peak-to-peak amplitude of the
477 E–W seasonal variation is almost twice greater in the satellite measurements (~25 ppbv) than in the
478 model data (~15 ppbv). This represents a ~1.5-2.0% variation of CH₄ in the E–W over the entire
479 year. The difference in amplitude between satellite and model in the seasonal evolution of E-W may
480 be due to: a) the comparison technique. There is a broader vertical domain in the measurement data
481 than in the model data; b) regarding the processes in summer, we may have less CH₄ trapped in the
482 Asian Monsoon Anticyclone redistributed towards the EMB (see section 5) in the models compared
483 to the measurements; c) regarding the processes in winter, we may have too much CH₄ transported
484 over the Mediterranean Sea to the East compared to the West, leading to a too smooth E-W gradient
485 in the models compared to the measurements.

486 We have also to remind that statistically the number of spaceborne measurements used in our
487 analysis (see section 2) is ~5 times greater in IASI compared to AIRS, ~30 times greater in AIRS
488 compared to GOSAT. Consequently, GOSAT monthly-averaged data appear noisier than AIRS
489 monthly-averaged data. Note that IASI total columns are not and cannot be directly compared with
490 AIRS or GOSAT profiles in our analysis. Nevertheless, although IASI data are not operationally
491 produced, the IASI E-W seasonal variation is very consistent with the E-W seasonal variation as

492 deduced from all other datasets. The monthly random error attributed to the E–W IASI CH₄ is about
493 0.1%, much less than the observed peak-to-peak yearly variation. We estimate that the AIRS
494 monthly random error attributed to the E–W CH₄ is twice greater than the one calculated for IASI,
495 and that the GOSAT monthly random error is about 5 times greater than the ones calculated for
496 IASI. We discuss in the next section the origin of the summer peak in the E–W seasonal variation.
497

498 **5. Contribution of the Asian Monsoon Anticyclone**

499 As stated in sections 3 and 4, interpreting the temporal evolution of the E–W CH₄ seasonal
500 evolution along the vertical requires to consider the evolution of CH₄ over the Asian continent
501 because of the importance of long-range transport. From Rodwell and Hoskins (1996), it is known
502 that there is a meteorological link between monsoons and the dynamics of the deserts and more
503 precisely between the Asian monsoon and the EMB summer regime. The subsidence centre over the
504 EMB owes its location, timing of onset and intensity to the Asian monsoon, and not to the Hadley
505 circulation. Although it takes less than one day to reach the upper troposphere within the Asian
506 monsoon, back-trajectory calculation (Ziv et al., 2004) shows that it takes about 3–4 days for an air
507 parcel to reach and descent the upper tropospheric EMB from the vicinity of the anticyclone that
508 develops over the Asian monsoon. Inside the Asian Monsoon Anticyclone (AMA), pollutants like
509 CO originating from the surface constitute about 50% of the CO concentration at 100 hPa (Park et
510 al., 2009), with the reminder resulting from chemical production in the troposphere. Most of the CO
511 within the AMA comes from India and South East Asia, with an insignificant contribution from the
512 Tibetan Plateau. Randel and Park (2006), and Park et al. (2009) have analyzed in detail this
513 phenomenon over Asia by considering dynamical parameters (potential vorticity) and chemical
514 species (H₂O, CO and O₃).

515 Numerous studies have already evaluated the impact of transport vs. emission of pollutants and
516 aerosols over the MB and its temporal variability considering different pollutants, chemical

517 compounds and aerosols (Wanger et al., 2000; Lelieveld et al., 2002; Pfister et al., 2004; Kallos et
518 al., 2007). As stated in section 3, two main dynamic factors affect the EMB: 1) the upper to mid-
519 tropospheric subsidence, and 2) the lower-level cool Etesian winds (Ziv et al., 2004). Although the
520 EMB is characterized by strong descent in the middle and upper troposphere in summer, transport
521 from the boundary layer accounts for about 25% of the local Middle Eastern contribution to the
522 ozone enhancement in the middle troposphere (Liu et al., 2009). Elevated CO episodes in EMB
523 during summer can also be attributed to synoptic conditions prone to favorable transport from
524 Turkey and Eastern Europe towards the EMB rather than increased emissions (Drori et al., 2012).
525 Upper tropospheric longitudinal gradients in the EMB of CH₄, CO, hydrocarbons, including
526 acetone, methanol, and acetonitrile, halocarbons, O₃ and total reactive nitrogen (NO_y) were also
527 attributed in August 2001 to the chemical impact of the Asian plume (Scheeren et al., 2003).
528 Finally, Georgoulias et al. (2011) present some interesting results of CH₄ from space in the vicinity
529 of the Mediterranean Sea, but only over land and essentially over the Eastern Mediterranean. The
530 authors found, from the total columns of CH₄ as measured by SCIAMACHY in 2003 and 2004, an
531 obvious maximum in August that could not be attributed to any volcano eruptions although this
532 area hosts a significant number of geological formations that could potentially contribute to the total
533 CH₄ burden. Being given that the sensitivity of the SCIAMACHY CH₄ total columns covers the
534 vertical domain 1000-200 hPa from the vertical structure of the averaging kernels presented in
535 Buchwitz et al. (2005), we note that 1) this maximum localized in August is consistent with our
536 study, and 2) the impact of the AMA on the CH₄ fields in the mid-to-upper troposphere cannot be
537 ruled out.

538 In order to analyze the climalogical impact of the AMA onto the EMB, we have calculated (Fig.
539 8) the climatological six-day back-trajectories from the point at 33° N, 35° E located in the EMB
540 (red filled circle on Fig. 8) based on the British Atmospheric Data Centre trajectory service
541 (<http://badc.nerc.ac.uk/community/trajectory/>) from 1st July to 31st August (summer convective
542 period) from 2001 to 2010 every 12 hours at 5 different pressure levels: 850 and 700 hPa (lower

543 troposphere), 500 hPa (middle troposphere), and 300 and 200 hPa (upper troposphere). The position
544 of the gravity centre of each distribution at each level is represented every 24 hours by a star on
545 Figure 8. Data from European Centre for Medium-Range Weather Forecasts (ECMWF) archive (2.5
546 degree/pressure levels) are used in the present calculation. The methodology has been first used
547 over the Dome C (Concordia) station in Antarctica and presented in Ricaud (2014). We have also
548 performed the same analysis but for the winter period from 1st January to 31st March 2001-2010
549 (Fig. 8). Figure 8 undoubtedly shows that air parcels above the EMB during the Asian monsoon
550 period of July-August from 2001 to 2010 are originated: a) from Asia in the upper troposphere, b)
551 from Northern America and Northern Africa in the mid-troposphere and c) from Europe in the low
552 troposphere. The same Figure also shows that in winter (and all other seasons but summer, not
553 shown) air parcels above the EMB are originated from the West (Europe, Atlantic Ocean, North
554 America, Pacific Ocean) whatever the pressure level considered from 850 to 100 hPa.

555 We apply the same climatological approach based on the CNRM-AOCCM and LMDz-OR-
556 INCA CH₄ model results over the period 2001-2010. We consider (Fig. 9) the E–W CH₄ seasonal
557 evolution at pressure levels from the lowermost troposphere to the lowermost stratosphere (850,
558 700, 500, 300, 200 and 100 hPa) and different scenarios for LMDz-OR-INCA (RCPs 2.6, 4.5, 6.0
559 and 8.5) in order to check out whether the summer peak still persists. We also represent the fields of
560 CH₄ as specified and/or calculated in the lowermost level (surface level) by CNRM-AOCCM and
561 by LMDz-OR-INCA (4 scenarios) in summer averaged over the climatological period 2001-2010
562 over a wide area covering the MB and the Asian continent in Figure 10, whilst the CH₄ fields
563 calculated at 200 hPa are shown in Figure 11.

564 The E-W CH₄ seasonal variations from the two models (Fig. 9) behave distinctively in the entire
565 troposphere, and agree very well in the lowermost stratosphere. In the lower troposphere (850 and
566 700 hPa), the E–W CH₄ seasonal evolution from LMDz-OR-INCA exhibits a strong semi-annual
567 oscillation of ± 10 -15 ppbv peaking in winter and summer for the 4 RCPs whilst the evolution from
568 the CNRM-AOCCM shows a weak annual oscillation of 4-5 ppbv amplitude, with a strong

569 minimum in summer, namely out-of-phase relative to the LMDz-OR-INCA variation. In the middle
570 troposphere at 500 hPa, the 4 LMDz-OR-INCA outputs exhibit a net maximum in August of ~8
571 ppbv with minima ranging from -2 to -12 ppbv from October to June, whilst the output from
572 CNRM-AOCCM again shows a strong minimum in summer of about -4 ppbv. At this stage, it is
573 important to remind that the two models are Global Circulation Models (GCMs) with an on-line
574 chemistry. The emissions of CH₄ are time-, longitude- and latitude-dependent in LMDz-OR-INCA
575 with surface maxima over the Northern continent (Fig. 10). In CNRM-AOCCM, there is no
576 emission of CH₄ (Fig. 10) but mixing ratios of CH₄ between the surface and the 560 hPa level are
577 relaxed towards evolving global mean surface abundances. This explains why the two models
578 behave separately for pressures greater or equal to 500 hPa. Note that, regarding the shape of the
579 E-W CH₄ seasonal evolution, there is no significant difference within the different scenarios of the
580 LMDz-OR-INCA outputs since surface CH₄ show the same structures independently of the RCPs
581 considered (Fig. 10).

582 In the upper troposphere (200 and 300 hPa), the outputs from the two models show a peak in
583 summer in the E-W CH₄ seasonal evolution (Fig. 9), but this differs from the RCPs considered for
584 LMDz-OR-INCA. The maximum is much more intense in CNRM-AOCCM (~8 ppbv in July-
585 August and ~30 ppbv in June-July at 300 and 200 hPa, respectively) than in LMDz-OR-INCA (~1
586 and ~10 ppbv in August for RCP 4.5 but only -4 and +4 ppbv in August for RCP 8.5 at 300 and
587 200 hPa, respectively; one peak at -4 ppbv in August for RCP 6.0 at 300 hPa but no peak at 200
588 hPa; no peak for RCP 2.6 neither at 300 nor at 200 hPa). On average, from 500 to 200 hPa, only the
589 RCP 4.5 scenario from LMDz-OR-INCA shows a positive maximum in summer. At 300 and 200
590 hPa, the LMDz-OR-INCA summer peak is much less intense than the CNRM-AOCCM summer
591 peak.

592 It is not obvious to understand why the E-W seasonal variation at 200 hPa is positive in summer
593 for RCP 4.5 and not for the other RCPs (except RCP 8.5 in August). The horizontal distribution of
594 CH₄ calculated by the two models at 200 hPa (Fig. 11) drastically differs but local maxima are

595 centred within the AMA. A zonally-symmetric structure showing a strong South-North gradient in
596 CH₄ is modelled by CNRM-AOCCM with maxima in the tropics (1800 ppbv) and minima at high
597 latitudes (1700 ppbv) and a local maximum centred within the core of the AMA with values greater
598 than 1807 ppbv elongated towards two axis: 1) South-East Asia and 2) Middle East and EMB. The
599 CH₄ field calculated by LMDz-OR-INCA considering the 4 scenarios also shows two maxima over
600 Northern India and over North-East Asia but the horizontal distribution is not zonally-symmetric
601 due to a zonally-asymmetric CH₄ surface field. In all the scenarios considered, the CH₄ maxima
602 within the AMA range from 1710 to 1750 ppbv with increasing RCPs from 2.6 to 8.5. An elongated
603 tongue of enriched CH₄ enters the EMB. More precisely, we can argue that in RCPs 2.6, 6.0 and
604 8.5, the primary maximum of CH₄ is located northward at 50°N, 135°E (CH₄ values greater than
605 1720, 1730 and 1750 ppbv, respectively) although it is a secondary maximum in RCP 4.5 (CH₄
606 values less than 1720 ppbv). Through long-range transport, this mid-latitude maximum is
607 transported Eastward within a band 40°N-50°N enriching CH₄ in the WMB and producing a E-W
608 minimum in summer for RCPs 2.6, 6.0 and 8.5. Since there is a North-South gradient with a
609 maximum in the South for CNRM-AOCCM, CH₄-depleted air masses reach the WMB although
610 CH₄-enriched air masses from the AMA reach the EMB producing a systematic peak in summer,
611 consistently with RCP 4.5.

612 In the lower stratosphere (100 hPa, Fig. 9), all the model outputs are consistent to each other
613 showing an annual oscillation, with a wide maximum in summer (60-80 ppbv) and a wide minimum
614 in winter (20-35 ppbv). This is apparently surprising keeping in mind that both models significantly
615 differ from the surface (see Fig. 10) to ~500 hPa. But, in the WMB, the 100-hPa pressure
616 corresponds to 420-K potential temperature both in summer (Fig. 2) and in winter (Fig. 4) whilst, in
617 the EMB, it corresponds to 390 K in summer and 400 K in winter, namely closer to the tropopause
618 in summer than in winter. Consequently, whatever the model considered, the E-W CH₄ seasonal
619 variation at 100 hPa a) is always positive and b) shows a peak in the summer period. We note that
620 the summer peak in E-W seasonal evolution from the middle to the upper troposphere has also been

621 observed and calculated by considering other constituents like CO and O₃ (not shown). This is the
622 main topic of a forthcoming paper.

623 In conclusion, a schematic representation of the summertime processes impacting mid-to-upper
624 CH₄ in the EMB is presented in Figure 12. In our study, whatever the amount of CH₄ at the surface
625 and its horizontal distribution, 1850-2000 ppbv for LMDz-OR-INCA consistently with the emission
626 sources (Asia, Northern and Eastern Europe, Central Africa) or 1820 ppbv uniformly spread (Fig.
627 10), the Asian monsoon traps elevated amounts of CH₄ that converge through the depression, and
628 are uplifted up to the upper troposphere at 200 hPa (Fig. 11) where they build up. At this level, the
629 AMA re-distributes elevated amounts of CH₄ towards Middle East, North Africa and the EMB
630 through long-range transport. Finally, elevated amounts of CH₄ build up in the EMB where they
631 descend to the middle troposphere.

632

633 **6. Conclusions**

634 The present study is part of the Chemical and Aerosol Mediterranean Experiment (ChArMEX)
635 programme. The aim is to investigate the tropospheric CH₄ time and space variations above the
636 Mediterranean Basin (MB) and to attribute the variability to differing synoptic and global scales
637 depending on the season and the altitude layer considered. Since the analysis of the mid-to-upper
638 tropospheric CH₄ distribution from spaceborne sensors and model outputs is challenging, we have
639 used a wide variety of datasets. 1) The spaceborne measurements from Thermal Infrared (TIR)
640 instruments: Thermal And Near infrared Sensor for carbon Observations - Fourier Transform
641 Spectrometer (TANSO-FTS) instrument on the Greenhouse gases Observing SATellite (GOSAT)
642 satellite, the Atmospheric InfraRed Spectrometer (AIRS) on the Aura platform and the Infrared
643 Atmospheric Sounder Interferometer (IASI) instrument aboard the MetOp-A platform. 2) The
644 model results from the Chemical Transport Model (CTM) MOCAGE, and the two Chemical
645 Climate Models (CCMs) CNRM-AOCCM and LMDz-OR-INCA (the later considering different
646 emission scenarios, RCPs 2.6, 4.5, 6.0 and 8.5).

647 Since CH₄ is a long-lived tracer with lifetime of ~12 years and is supposed to be well mixed in
648 the troposphere, we had to adopt a climatological approach to highlight the weak expected
649 variability. Spaceborne measurements and the model results were selected and monthly-averaged
650 only over the Mediterranean Sea. The period under interest spans from 2008 to 2011 for the satellite
651 measurements and the MOCAGE model results whilst, regarding the CCMs, we have averaged the
652 model outputs over the climatological period from 2001 to 2010.

653 From both satellite and model results, our study obviously demonstrates the persistence of an
654 East-West gradient in CH₄ from the middle to the upper troposphere with a maximum in the
655 Western MB whatever the season considered except in summer when larger amounts of CH₄
656 accumulate above the Eastern MB. In winter, air masses mainly originating from Atlantic Ocean
657 and Europe tend to favour an elevated amount of mid-to-upper tropospheric CH₄ in the West
658 compared to the East of the MB, with a general upward transport above the MB. In summer, the
659 meteorological condition of the MB is changed, favouring air from Northern Africa and Middle
660 East together with Atlantic Ocean and Europe, with a general descent above the Eastern MB.

661 Our analysis shows that, in the upper troposphere (300 hPa), the peak-to-peak amplitude of the
662 East–West seasonal variation in CH₄ above the MB is weak but almost twice greater in the satellite
663 measurements (~25 ppbv) than in the model data (~15 ppbv).. The maximum of CH₄ in summer
664 above the Eastern MB can be explained by a series of dynamical processes only occurring in
665 summer. The Asian monsoon traps and uplifts high amounts of CH₄ to the upper troposphere where
666 they build up. The Asia monsoon Anticyclone redistributes these elevated CH₄ amounts towards
667 North Africa and Middle East to finally reach and descent in the Eastern MB. Consequently, the
668 seasonal variation of the difference in CH₄ between the East and the West MB shows a maximum in
669 summer for pressures from 500 to 100 hPa considering both spaceborne measurements and model
670 results whatever the emission scenarios used for the CCMs. But only the RCP 4.5 scenario gives
671 systematically a positive summer peak whatever the pressure level considered, consistently with the
672 measurements.

673 From this study, we conclude that CH₄ in the mid-to-upper troposphere over the MB is mainly
674 affected by long-range transport, particularly intense in summer from Asia. Conversely, in the
675 lower troposphere, the CH₄ variability is driven by the local sources of emission in the vicinity of
676 the MB. Other constituents can also be affected by this summer mechanism e.g. O₃ and CO (not
677 shown). In a forthcoming paper, the time evolution of the CH₄, O₃ and CO fields above the MB and
678 at the Asian scale is being studied by considering the outputs from different CCMs in the
679 contemporary period (2000-2010) in order to study the future evolution of the chemical climate
680 over the MB by 2100. Finally, despite the fact that IASI CH₄ data as delivered by EUMETSAT are
681 not operational, the seasonal variation of the East-West difference in CH₄ total columns is
682 nevertheless consistent with theoretical results and measurements from AIRS and IASI.

683

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698 **References**

- 699 August, T., Klaes, D., Schlüssel, P., Hultberg, T., Crapeau, M., Arriaga, A., O'Carroll, A., Coppens,
700 D., Munro, R., and Calbet, X.: IASI on Metop-A: Operational Level 2 retrievals after five years
701 in orbit, *J. Quant. Spectrosc. Rad. Transfer*, 113:11, 1340-1371, 2012.
- 702 Bergamaschi, P., Frankenberg, C., Fokke Meirink, J., Krol, M., Villani, M. G., Houweling, S.,
703 Dentener, F., Dlugokencky, E. J., Miller, J. B., Gatti, L. V., Engel, A., and Levin, I.: Inverse
704 modeling of global and regional CH₄ emissions using SCIAMACHY satellite retrievals, *J.*
705 *Geophys. Res.*, 114, D22301, doi:10.1029/2009JD012287, 2009.
- 706 Bousquet, P., Ringeval, B., Pison, I., Dlugokencky, E. J., Brunke, E.-G., Carouge, C., Chevallier,
707 F., Fortems-Cheiney, A., Frankenberg, C., Hauglustaine, D. A., Krummel, P. B., Langenfelds, R.
708 L., Ramonet, M., Schmidt, M., Steele, L. P., Szopa, S., Yver, C., Viovy, N., and Ciais, P.: Source
709 attribution of the changes in atmospheric methane for 2006–2008, *Atmos. Chem. Phys.*, 11,
710 3689-3700, doi:10.5194/acp-11-3689-2011, 2011.
- 711 Buchwitz, M., Rozanov, V. V., and Burrows, J. P.: A near-infrared optimized DOAS method for
712 the fast global retrieval of atmospheric CH₄, CO, CO₂, H₂O, and N₂O total column amounts
713 from SCIAMACHY Envisat-1 nadir radiances, *J. Geophys. Res.*, 105(D12), 15231–15245,
714 doi:10.1029/2000JD900191, 2000.
- 715 Buchwitz, M., de Beek, R., Burrows, J. P., Bovensmann, H., Warneke, T., Notholt, J., Meirink, J.
716 F., Goede, A. P. H., Bergamaschi, P., Körner, S., Heimann, M., and Schulz, A.: Atmospheric
717 methane and carbon dioxide from SCIAMACHY satellite data: initial comparison with
718 chemistry and transport models, *Atmos. Chem. Phys.*, 5, 941-962, doi:10.5194/acp-5-941-2005,
719 2005.
- 720 Claeysman, M., Attié, J.-L., El Amraoui, L., Cariolle, D., Peuch, V.-H., Teyssèdre, H., Josse, B.,
721 Ricaud, P., Massart, S., Piacentini, A., Cammas, J.-P., Livesey, N. J., Pumphrey, H. C., and
722 Edwards, D. P.: A linear CO chemistry parameterization in chemistry-transport models:
723 evaluation and application to data assimilation, *Atmos. Chem. Phys.*, 10, 6097-6115, 2010.

- 724 Claeysman, M., Attié, J.-L., Peuch, V.-H., El Amraoui, L., Lahoz, W. A., Josse, B., Ricaud, P., von
725 Clarmann, T., Höpfner, M., Orphal, J., Flaud, J.-M., Edwards, D. P., Chance, K., Liu, X.,
726 Pasternak, F., and Cantié, R.: A geostationary thermal infrared sensor to monitor the lowermost
727 troposphere: O₃ and CO retrieval studies, *Atmos. Meas. Tech.*, 4, 297-317, 2011.
- 728 Clerbaux, C., Chazette, P., Hadji-Lazaro, J., Mégie, G., Müller, J.-F., and Clough, S. A.: Remote
729 sensing of CO, CH₄, and O₃ using a spaceborne nadir-viewing interferometer, *J. Geophys. Res.*,
730 103, 18999–19013, doi:10.1029/98JD01422, 1998.
- 731 Courtier, P., Freydier, C., Geleyn, J. F., Rabier, F., and Rochas, M.: The ARPEGE project at
732 METEO-FRANCE. In: Proc ECMWF Workshop. Numerical methods in atmospheric modelling,
733 9–13 Sept 1991, 2, 193–231. ECMWF, Shinfield Park, Reading, UK, 1991.
- 734 Crevoisier, C., Nobileau, D., Armante, R., Crépeau, L., Machida, T., Sawa, Y., Matsueda, H.,
735 Schuck, T., Thonat, T., Pernin, J., Scott, N. A., and Chédin, A.: The 2007–2011 evolution of
736 tropical methane in the mid-troposphere as seen from space by MetOp-A/IASI, *Atmos. Chem.*
737 *Phys.*, 13, 4279-4289, doi:10.5194/acp-13-4279-2013, 2013.
- 738 Cros, B., Durand, P., and Cachier, H.: An overview of the ESCOMPTE campaign, *Atmos. Res.*,
739 69(3-4), 241-279, 2004.
- 740 Dayan, U.: Climatology of Back Trajectories from Israel Based on Synoptic Analysis. *J. Climate*
741 *Appl. Meteor.*, 25, 591–595. doi: [http://dx.doi.org/10.1175/1520-0450\(1986\)](http://dx.doi.org/10.1175/1520-0450(1986)25<591:CLBTIAES.2.TX;2-0), 1986.
- 742 Dentener, F., Stevenson, D., Cofala, J., Mechler, R., Amann, M., Bergamaschi, P., Raes, F., and
743 Derwent, R.: The impact of air pollutant and methane emission controls on tropospheric ozone
744 and radiative forcing: CTM calculations for the period 1990–2030, *Atmos. Chem. Phys.*, 5,
745 1731–1755, <http://www.atmos-chem-phys.net/5/1731/2005/>, 2005.
- 746 Drori, R., Dayan, U., Edwards, D. P., Emmons, L. K., and Erlick, C.: Attributing and quantifying
747 carbon monoxide sources affecting the Eastern Mediterranean: a combined satellite, modelling,
748 and synoptic analysis study, *Atmos. Chem. Phys.*, 12, 1067-1082, doi:10.5194/acp-12-1067-
749 2012, 2012.

- 750 Dufour, A., Amodei, M., Ancellet, G., and Peuch, V.-H.: Observed and modelled “chemical
751 weather” during ESCOMPTE, *Atmos. Res.*, 74, 161–189, 2005.
- 752 El Amraoui, L., Attié, J.-L., Semane, N., Claeysman, M., Peuch, V.-H., Warner, J., Ricaud, P.,
753 Cammas, J.-P., Piacentini, A., Cariolle, D., Massart, S., and Bencherif, H.: Midlatitude
754 stratosphere – troposphere exchange as diagnosed by MLS O₃ and MOPITT CO assimilated
755 fields, *Atmos. Chem. Phys.*, 10, 2175–2194, 2010.
- 756 EUMETSAT, IASI Level 2 Products Guide, EUM/OPS-EPS/MAN/04/0033. Available on
757 <http://oiswww.eumetsat.org/WEBOPS/eps-pg/IASI-L2/IASIL2-PG-0TOC.htm>, 2004.
- 758 Folberth, G. A., Hauglustaine, D. A., Lathière, J., and Brocheton, F.: Interactive chemistry in the
759 Laboratoire de Météorologie Dynamique general circulation model: model description and
760 impact analysis of biogenic hydrocarbons on tropospheric chemistry, *Atmos. Chem. Phys.*,
761 6:2273–2319. www.atmos-chem-phys.net/6/2273/2006/, 2006.
- 762 Georgoulias, A. K., Kourtidis, K. A., Buchwitz, M., Schneising, O., and Burrows, J. P.: A case
763 study on the application of SCIAMACHY satellite methane measurements for regional studies:
764 the Greater Area of Eastern Mediterranean, *Int. J. Remote Sens.*, 32(3), 787–813,
765 doi:10.1080/01431161.2010.517791, 2011.
- 766 Giorgi, F., and Lionello, P.: Climate change projections for the Mediterranean region, *Global and
767 Planetary Change*, 63(2), 90–104, doi:10.1016/j.gloplacha.2007.09.005, 2008.
- 768 Hauglustaine, D. A., Hourdin, F., Jourdain, L., Filiberti, M.-A., Walters, S., Lamarque, J.-F., and
769 Holland, E. A.: Interactive chemistry in the Laboratoire de Meteorologie Dynamique general
770 circulation model: description and background tropospheric chemistry evaluation, *J. Geophys.
771 Res.*, 109:D04314. doi:10.1029/2003JD003957, 2004.
- 772 Hilton, F., Armante, R., August, T., Barnet, C., Bouchard, A., Camy-Peyret, C., Capelle, V.,
773 Clarisse, L., Clerbaux, C., Coheur, P.-F., Collard, A., Crevoisier, C., Dufour, G., Edwards, D.,
774 Faijan, F., Fourrié, N., Gambacorta, A., Goldberg, M., Guidard, V., Hurtmans, D., Illingworth,
775 S., Jacquinet-Husson, N., Kerzenmacher, T., Klaes, D., Lavanant, L., Masiello, G., Matricardi,

- 776 M., McNally, A., Newman, S., Pavelin, E., Payan, S., Péquignot, E., Peyridieu, S., Phulpin, T.,
777 Remedios, J., Schlüssel, P., Serio, C., Strow, L., Stubenrauch, C., Taylor, J., Tobin, D., Wolf,
778 W., and Zhou, D.: Hyperspectral Earth Observation from IASI: Five Years of Accomplishments,
779 B. Am. Meteor. Soc., 93, 347–370, doi:10.1175/BAMS-D-11-00027.1, 2012.
- 780 Hourdin, F., Musat, I., Bony, S., Braconnot, P., Codron, F., Dufresne, J.-L., Fairhead, L., Filiberti,
781 M.-A., Friedlingstein, P., Grandpeix, J.-Y., Krinner, G., LeVan, P., Li, Z.-X., and Lott, F.: The
782 LMDZ4 general circulation model: climate performance and sensitivity to parametrized physics
783 with emphasis on tropical convection, *Clim. Dyn.*, 27, 787–813, doi:10.1007/s00382-006-0158-
784 0, 2006.
- 785 Huszar, P., Teyssèdre, H., Cariolle, D., Olivié, D. J. L., Michou, M., Saint-Martin, D., Senesi, S.,
786 Voldoire, A., Salas y Melia, D., Alias, A., Karcher, F., Ricaud, P., and Halenka, T.: Modeling
787 the present and future impact of aviation on climate: an AOGCM approach with online coupled
788 chemistry, *Atmos. Chem. Phys.*, 13, 10027-10048, doi:10.5194/acp-13-10027-2013, 2013.
- 789 Imasu, R., Saitoh, N., and Niwa, Y.: Retrieval performance of GOSAT thermal infrared FTS sensor
790 for measuring CO₂ concentrations, *Proc. SPIE 6744, Sensors, Systems, and Next-Generation*
791 *Satellites XI*, 67440F, <http://dx.doi.org/10.1117/12.737796>, 2007.
- 792 IPCC: Climate change 2007: The physical science basis. Contribution of working group I to the
793 fourth assessment report of the intergovernmental panel on climate change, Cambridge, UK and
794 New York, USA, Cambridge University Press, 996 pp., 2007.
- 795 Josse, B., Simon, P., and Peuch, V.-H.: Rn-222 global simulations with the multiscale CTM
796 MOCAGE, *Tellus*, 56B, 339–356, 2004.
- 797 Kallos, G., Astitha, M., Katsafados, P., and Spyrou, C.: Long-Range Transport of
798 Anthropogenically and Naturally Produced Particulate Matter in the Mediterranean and North
799 Atlantic: Current State of Knowledge, *J. Appl. Meteorol. Clim.*, 46, 1230-1251, 2007.
- 800 Krinner, G., Viovy, N., de Noblet-Ducoudré, N., Ogée, J., Polcher, J., Friedlingstein, P., Ciais, P.,
801 Sitch, S., and Prentice, I. C.: A dynamic global vegetation model for studies of the coupled

- 802 atmosphere-biosphere system, *Global Biogeochem. Cycles*, 19, GB1015,
803 doi:10.1029/2003GB002199, 2005.
- 804 Kuze, A., Suto, H., Nakajima, M., and Hamazaki, T.: Thermal and near infrared sensor for carbon
805 observation Fourier-transform spectrometer on the Greenhouse Gases Observing Satellite for
806 greenhouse gases monitoring, *Appl. Opt.*, 48, 6716-6733, 2009.
- 807 Ladstätter-Weißmayer, A., Heland, J., Kormann, R., von Kuhlmann, R., Lawrence, M. G.,
808 Meyer-Arnek, J., Richter, A., Wittrock, F., Ziereis, H., and Burrows, J.-P.: Transport and build-
809 up of tropospheric trace gases during the MINOS campaign: comparison of GOME, in situ
810 aircraft measurements and MATCH-MPIC-data, *Atmos. Chem. Phys.*, 3, 1887–1902, 2003.
- 811 Lamarque, J.-F., Kyle, G. P., Meinshausen, M., Riahi, K., Smith, S. J., van Vuuren, D. P., Conley,
812 A. J., and Vitt, F.: Global and regional evolution of short-lived radiatively-active gases and
813 aerosols in the Representative Concentration Pathways, *Climatic Change*, 109:191–212 DOI
814 10.1007/s10584-011-0155-0, 2011.
- 815 Lamarque, J.-F., Shindell, D. T., Josse, B., Young, P. J., Cionni, I., Eyring, V., Bergmann, D.,
816 Cameron-Smith, P., Collins, W. J., Doherty, R., Dalsoren, S., Faluvegi, G., Folberth, G., Ghan,
817 S. J., Horowitz, L. W., Lee, Y. H., MacKenzie, I. A., Nagashima, T., Naik, V., Plummer, D.,
818 Righi, M., Rumbold, S. T., Schulz, M., Skeie, R. B., Stevenson, D. S., Strode, S., Sudo, K.,
819 Szopa, S., Voulgarakis, A., and Zeng, G.: The Atmospheric Chemistry and Climate Model
820 Intercomparison Project (ACCMIP): overview and description of models, simulations and
821 climate diagnostics, *Geosci. Model Dev.*, 6, 179-206, 2013.
- 822 Lefèvre, F., Brasseur, G. P., Folkins, I., Smith, A. K., and Simon, P.: Chemistry of the 1991–1992
823 stratospheric winter: Three-dimensional model simulations, *J. Geophys. Res.*, 99, 9183–8195,
824 1994.
- 825 Lelieveld, J., Berresheim, H., Borrmann, S., Crutzen, P. J., Dentener, F. J., Fischer, H., Feichter, J.,
826 Flatau, P. J., Heland, J., Holzinger, R., Kormann, R., Lawrence, M. G., Levin, Z., Markowicz,

- 827 K. M., Mihalopoulos, N., Minikin, A., Ramanathan, V., de Reus, M., Roelofs, G. J., Scheeren,
828 H. A., Sciare, J., Schlager, H., Schultz, M., Siegmund, P., Steil, B., Stephanou, E. G., Stier, P.,
829 Traub, M., Warneke, C., Williams, J., and Ziereis, H.: Global Air Pollution Crossroads over the
830 Mediterranean, *Science*, 298, 5594, 794-799, DOI: 10.1126/science.1075457, 2002.
- 831 Lionello, P. (Ed.): *The Climate of the Mediterranean Region: From the past to the future*. 592p,
832 Elsevier, 2012.
- 833 Liu, J. J., Jones, D. B. A., Worden, J. R., Noone, D., Parrington, M., and Kar, J.: Analysis of the
834 summertime buildup of tropospheric ozone abundances over the Middle East and North Africa
835 as observed by the Tropospheric Emission Spectrometer instrument, *J. Geophys. Res.*, 114,
836 D05304, doi:10.1029/2008JD010993, 2009.
- 837 Masson, O., Piga, D., Gurriaran, R., and D'Amico, D.: Impact of an exceptional Saharan dust
838 outbreak in France: PM10 and artificial radionuclides concentrations in air and in dust deposit,
839 *Atmos. Env.*, 44, 20, 2478–2486, <http://dx.doi.org/10.1016/j.atmosenv.2010.03.004>, 2010.
- 840 Michou, M., and Peuch, V.-H.: Surface exchanges in the MOCAGE multiscale Chemistry and
841 Transport Model, *J. Water Sci.*, 15, 173–203, 2002.
- 842 Michou, M., Saint-Martin, D., Teyssèdre, H., Alias, A., Karcher, F., Olivié, D., Voldoire, A., Josse,
843 B., Peuch, V.-H., Clark, H., Lee, J. N., and Cheroux, F.: A new version of the CNRM
844 Chemistry-Climate Model, CNRM-CCM: description and improvements from the CCMVal-2
845 simulations, *Geosci. Model Dev.*, 4, 873–900, doi:10.5194/gmd-4-873-2011, 2011.
- 846 Millán, M. M., Salvador, R., Mantilla, E., and Kallos, G.: Photooxidant dynamics in the
847 Mediterranean basin in summer: Results from European research projects, *J. Geophys. Res.*, 102,
848 8811–8823, doi:10.1029/96JD03610, 1997.
- 849 Nabat, P., Solmon, F., Mallet, M., Kok, J. F., and Somot, S.: Dust emission size distribution impact
850 on aerosol budget and radiative forcing over the Mediterranean region: a regional climate model
851 approach, *Atmos. Chem. Phys.*, 12, 10545-10567, 2012.

- 852 Naik, V., Voulgarakis, A., Fiore, A. M., Horowitz, L. W., Lamarque, J.-F., Lin, M., Prather, M. J.,
853 Young, P. J., Bergmann, D., Cameron-Smith, P. J., Cionni, I., Collins, W. J., Dalsøren, S. B.,
854 Doherty, R., Eyring, V., Faluvegi, G., Folberth, G. A., Josse, B., Lee, Y. H., MacKenzie, I. A.,
855 Nagashima, T., van Noije, T. P. C., Plummer, D. A., Righi, M., Rumbold, S. T., Skeie, R.,
856 Shindell, D. T., Stevenson, D. S., Strode, S., Sudo, K., Szopa, S., and Zeng, G.: Preindustrial to
857 present-day changes in tropospheric hydroxyl radical and methane lifetime from the
858 Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP), *Atmos. Chem.*
859 *Phys.*, 13, 5277-5298, doi:10.5194/acp-13-5277-2013, 2013.
- 860 Park, M., Randel, W. J., Emmons, L. K., and Livesey, N. J.: Transport pathways of carbon
861 monoxide in the Asian summer monsoon diagnosed from Model of Ozone and Related Tracers
862 (MOZART), *J. Geophys. Res.*, 114, D08303, doi:10.1029/2008JD010621, 2009.
- 863 Peuch, V.-H., Amodei, M., Barthet, T., Cathala, M.-L., Josse, B., Michou, M., and Simon, P. :
864 MOCAGE: Modèle de Chimie, Atmosphérique A Grande Echelle, Actes des Atelier de
865 Modélisation de l'Atmosphère, Météo-France, 33–36, 1999.
- 866 Pfister, G., Petron, G., Emmons, L. K., Gille, J. C., Edwards, D. P., Lamarque, J.-F., J.-L. Attié, C.
867 Granier, and Novelli, P. C: Evaluation of CO simulations and the analysis of the CO budget for
868 Europe, *J. Geophys. Res.-Atmos.* (1984–2012), 109(D19), 2004.
- 869 Randel, W. J., and Park, M.: Deep convective influence on the Asian summer monsoon anticyclone
870 and associated tracer variability observed with Atmospheric Infrared Sounder (AIRS), *J.*
871 *Geophys. Res.*, 111, D12314, doi:10.1029/2005JD006490, 2006.
- 872 Razavi, A., Clerbaux, C., Wespes, C., Clarisse, L., Hurtmans, D., Payan, S., Camy-Peyret, C., and
873 Coheur, P. F.: Characterization of methane retrievals from the IASI space-borne sounder, *Atmos.*
874 *Chem. Phys.*, 9, 7889–7899, 2009.
- 875 Ricaud, P., Attié, J.-L., Teyssède, H., El Amraoui, L., Peuch, V.-H., Matricardi, M., and
876 Schluessel, P.: Equatorial total column of nitrous oxide as measured by IASI on MetOp-A:

- 877 implications for transport processes, *Atmos. Chem. Phys.*, 9, 3947-3956, doi:10.5194/acp-9-
878 3947-2009, 2009.
- 879 Ricaud, P.: Variabilités de la vapeur d'eau et de la température troposphérique mesurées par le
880 radiomètre micro-onde HAMSTRAD au Dôme C, Antarctique. Partie II : Résultats scientifiques,
881 *La Météorologie*, 85, 35-46, doi: 10.4267/2042/53749, 2014.
- 882 Rodgers, C. D., *Inverse Methods for Atmospheric Sounding: Theory and Practice*, 1st ed.
883 Singapore: World Scientific, 2000.
- 884 Rodwell, M. J., and Hoskins, B. J.: Monsoons and the dynamics of deserts, *Q. J. R. Meteorol. Soc.*,
885 122, 1385-1404, 1996.
- 886 Saitoh, N., Touno, M., Hayashida, S., Imasu, R., Shiomi, K., Yokota, T., Yoshida, Y., Machida, T.,
887 Matsueda, H., and Sawa, Y.: Comparisons between XCH₄ from GOSAT Shortwave and Thermal
888 Infrared Spectra and Aircraft CH₄ Measurements over Guam, *SOLA*, 8, 145–149,
889 doi:10.2151/sola.2012-036, 2012.
- 890 Scheeren, H. A., Lelieveld, J., Roelofs, G. J., Williams, J., Fischer, H., de Reus, M., de Gouw, J. A.,
891 Bolder, M., van der Veen, C., and Lawrence, M.: The impact of monsoon outflow from India
892 and Southeast Asia in the upper troposphere over the eastern Mediterranean, *Atmos. Chem.*
893 *Phys.*, 3, 1589–1608, 2003.
- 894 Schicker, I., Radanovics, S., and Seibert, P.: Origin and transport of Mediterranean moisture and
895 air, *Atmos. Chem. Phys.*, 10, 5089-5105, doi:10.5194/acp-10-5089-2010, 2010.
- 896 Scheeren, H. A., Lelieveld, J., Roelofs, G. J., Williams, J., Fischer, H., de Reus, M., de Gouw, J. A.,
897 Warneke, C., Holzinger, R., Schlager, H., Klüpfel, T., Bolder, M., van der Veen, C., and
898 Lawrence, M.: The impact of monsoon outflow from India and Southeast Asia in the upper
899 troposphere over the eastern Mediterranean, *Atmos. Chem. Phys.*, 3, 1589-1608,
900 doi:10.5194/acp-3-1589-2003, 2003.
- 901 Schlüssel, P., Hultberg, T. H., Phillips, P. L., August, T., and Calbet, X.: The operational IASI
902 Level 2 Processor, *Adv. Space Res.*, 36, 982-988, 2005.

- 903 Stockwell, W. R., Kirchner, F., Kuhn, M., and Seefeld, S.: A new mechanism for regional
904 atmospheric chemistry modelling, *J. Geophys. Res.: Atmospheres* (1984–2012), 102(D22),
905 25847-25879, 1997.
- 906 Susskind, J., Blaisdell, J. M., Iredell, L. and Keita, F.: Improved Temperature Sounding and Quality
907 Control Methodology Using AIRS/AMSU Data: The AIRS Science Team Version 5 Retrieval
908 Algorithm, *IEEE Trans. Geosc. Remote Sens.*, 49, 883-907, 2011.
- 909 Szopa, S., Balkanski, Y., Schulz, M., Bekki, S., Cugnet, D., Fortems-Cheiney, A., Turquety, S.,
910 Cozic, A., Déandreis, C., Hauglustaine, D., Idelkadi, A., Lathière, J., Lefèvre, F., Marchand, M.,
911 Vuolo, R., Yan, N., and Dufresne, J.-L.: Aerosol and ozone changes as forcing for climate
912 evolution between 1850 and 2100, *Clim. Dynam.*, 40, 2223-2250, 2013. DOI:10.1007/s00382-
913 012-1408-y
- 914 Turquety, S., Hadji-Lazaro, J., Clerbaux, C., Hauglustaine, D. A., Clough, S. A., Cassé, V.,
915 Schlüssel, P., and Mégie, G.: Operational trace gas retrieval algorithm for the Infrared
916 Atmospheric Sounding Interferometer, *J. Geophys. Res.*, 109, D21301,
917 doi:10.1029/2004JD004821, 2004.
- 918 van der Werf, G. R., Randerson, J. T., Collatz, G. J., and Giglio, L.: Carbon emissions from fires in
919 tropical and subtropical ecosystems, *Global Change Biol.*, 9, 547–562, 2003.
- 920 Voldoire, A., Sanchez-Gomez, E., Salas y Melia, D., Decharme, B., Cassou, C., Sénési, S., Valcke,
921 S., Beau, I., Alias, A., Chevallier, M., Deque, M., Deshayes, J., Douville, H., Fernandez, E.,
922 Madec, G., Maisonnave, E., Moine, M.-P., Planton, S., Saint-Martin, D., Szopa, S., Tyteca, S.,
923 Alkama, R., Belamari, S., Braun, A., Coquart, L., and Chauvin, F.: The CNRM-CM5.1 global
924 climate model: description and basic evaluation, *Clim. Dynam.*, doi:10.1007/s00382-011-1259-
925 y, 2012.
- 926 Voulgarakis, A., Naik, V., Lamarque, J.-F., Shindell, D. T., Young, P. J., Prather, M. J., Wild, O.,
927 Field, R. D., Bergmann, D., Cameron-Smith, P., Cionni, I., Collins, W. J., Dalsøren, S. B.,
928 Doherty, R. M., Eyring, V., Faluvegi, G., Folberth, G. A., Horowitz, L. W., Josse, B.,

- 929 MacKenzie, I. A., Nagashima, T., Plummer, D. A., Righi, M., Rumbold, S. T., Stevenson, D. S.,
930 Strode, S. A., Sudo, K., Szopa, S., and Zeng, G.: Analysis of present day and future OH and
931 methane lifetime in the ACCMIP simulations, *Atmos. Chem. Phys.*, 13, 2563-2587,
932 doi:10.5194/acp-13-2563-2013, 2013.
- 933 Wanger, A., Peleg, M., Sharf, G., Mahrer, Y., Dayan, U., Kallos, G., Kotroni, V., Lagouvardos, K.,
934 Varinou, M., Papadopoulos, A., and Luria, M.: Some Observational and Modelling Evidence of
935 Long Range Transport of Air Pollutants from Europe Towards the Israeli Coast, *J. Geophys.*
936 *Res.*, 105, 7177-7186, 2000.
- 937 Worden, J., Kulawik, S., Frankenberg, C., Payne, V., Bowman, K., Cady-Peirara, K., Wecht, K.,
938 Lee, J. E., and Noone, D.: Profiles of CH₄, HDO, H₂O, and N₂O with improved lower
939 tropospheric vertical resolution from Aura TES radiances, *Atmos. Meas. Tech.*, 5, 397–411,
940 www.atmos-meas-tech.net/5/397/2012/doi:10.5194/amt-5-397-2012, 2012.
- 941 Wunch, D., Toon, G. C., Wennberg, P. O., Wofsy, S. C., Stephens, B. B., Fischer, M. L., Uchino,
942 O., Abshire, J. B., Bernath, P., Biraud, S. C., Blavier, J.-F. L., Boone, C., Bowman, K. P.,
943 Browell, E. V., Campos, T., Connor, B. J., Daube, B. C., Deutscher, N. M., Diao, M., Elkins, J.
944 W., Gerbig, C., Gottlieb, E., Griffith, D. W. T., Hurst, D. F., Jiménez, R., Keppel-Aleks, G.,
945 Kort, E. A., Macatangay, R., Machida, T., Matsueda, H., Moore, F., Morino, I., Park, S.,
946 Robinson, J., Roehl, C. M., Sawa, Y., Sherlock, V., Sweeney, C., Tanaka, T., and Zondlo, M. A.:
947 Calibration of the Total Carbon Column Observing Network using aircraft profile data, *Atmos.*
948 *Meas. Tech.*, 3, 1351-1362, doi:10.5194/amt-3-1351-2010, 2010.
- 949 Xiong, X., Barnet, C., Maddy, E., Sweeney, C., Liu, X., Zhou, L., and Goldberg, M.:
950 Characterization and validation of methane products from the Atmospheric Infrared Sounder
951 (AIRS), *J. Geophys. Res.*, 113, G00A01, doi:10.1029/2007JG000500., 2008.
- 952 Yokota, T., Yoshida, Y., Eguchi, N., Ota, Y., Tanaka, T., Watanabe, H., and Maksyutov, S.: Global
953 Concentrations of CO₂ and CH₄ Retrieved from GOSAT: First Preliminary Results, *SOLA*, 5,
954 160-163, doi:10.2151/sola.2009, 2009.

- 955 Yoshida, Y., Kikuchi, N., Morino, I., Uchino, O., Oshchepkov, S., Bril, A., Saeki, T., Schutgens,
956 N., Toon, G. C., Wunch, D., Roehl, C. M., Wennberg, P. O., Griffith, D. W. T., Deutscher, N.
957 M., Warneke, T., Notholt, J., Robinson, J., Sherlock, V., Connor, B., Rettinger, M., Sussmann,
958 R., Ahonen, P., Heikkinen, P., Kyrö, E., Mendonca, J., Strong, K., Hase, F., Dohe, S., and
959 Yokota, T.: Improvement of the retrieval algorithm for GOSAT SWIR XCO₂ and XCH₄ and
960 their validation using TCCON data, *Atmos. Meas. Tech.*, 6, 1533-1547, doi:10.5194/amt-6-
961 1533-2013, 2013.
- 962 Ziv, B., Saaroni, H., and Alpert, P.: The factors governing the summer regime of the eastern
963 Mediterranean, *Int. J. Climatol.*, 24, 1859–1871, doi: 10.1002/joc.1113, 2004.
- 964

964 **Table 1.** Nadir-viewing instruments having the capabilities to measure CH₄ in the troposphere.

965 Please, refer to the text for the acronyms.

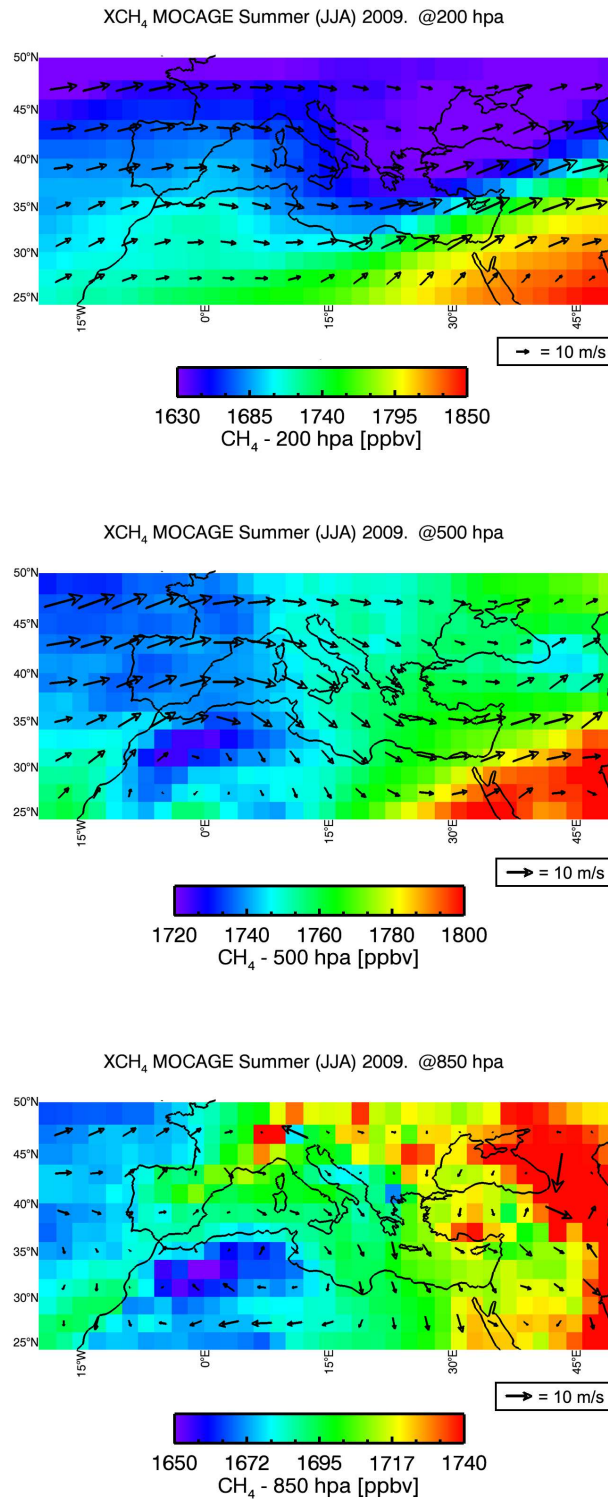
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Platform	Instrument	Operation time	Wavelength
ADEOS-1	IMG	1996-1997	TIR
ENVISAT	SCIAMACHY	2002-2012	NIR
Aura	TES	2004-date	TIR
GOSAT	TANSO-FTS	2008-date	SWIR & TIR
Aqua	AIRS	2004-date	TIR
MetOp-A	IASI	2008-date	TIR
MetOp-B	IASI	2012-date	TIR
MetOp-C	IASI	Expected in 2016	TIR

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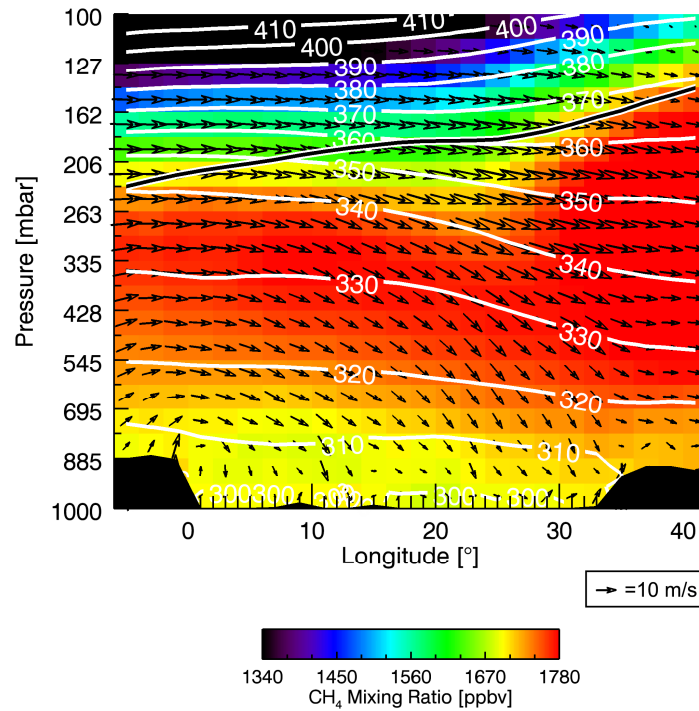
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970 **Figure 1.** (From bottom to top) Fields of CH₄ as calculated by MOCAGE and averaged for summer
 971 (JJA) 2009 at 850, 500 and 200 hPa. Superimposed are the horizontal winds from ARPEGE
 972 averaged over the same period. In order to highlight the CH₄ horizontal gradients, the range of the
 973 colour scale changes from top to bottom.

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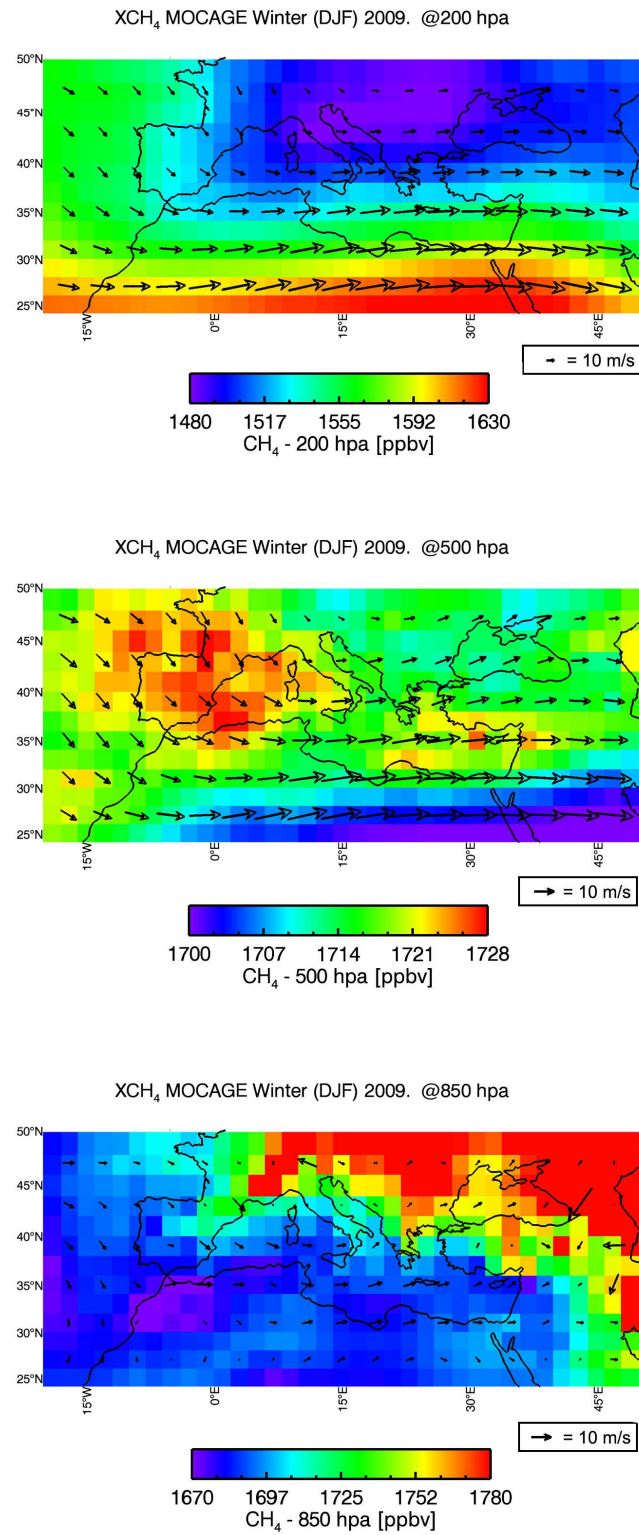
MOCAGE summer (JJA) 2009. CH₄ [ppbv]

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976 **Figure 2.** Vertical distribution of CH₄ as calculated by MOCAGE and averaged for JJA 2009 as a
 977 function of longitude along the red line represented above the Figure. Superimposed are the
 978 associated longitudinal and vertical winds from ARPEGE, together with the isentropes (white lines)
 979 and the cold point tropopause from NOAA NCEP/NCAR reanalyses (black line) averaged over the
 980 same period.

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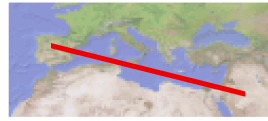
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983 **Figure 3.** Same as Fig. 1, but for winter (DJF) 2009.

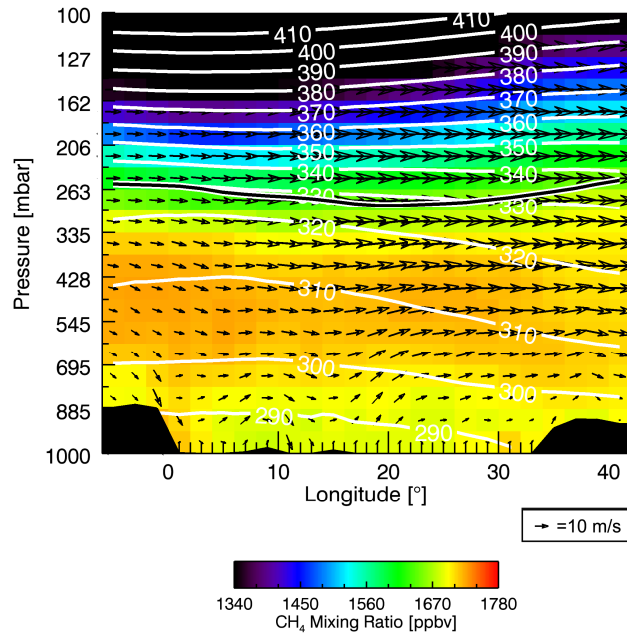
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MOCAGE winter (DJF) 2009. CH₄ [ppbv]



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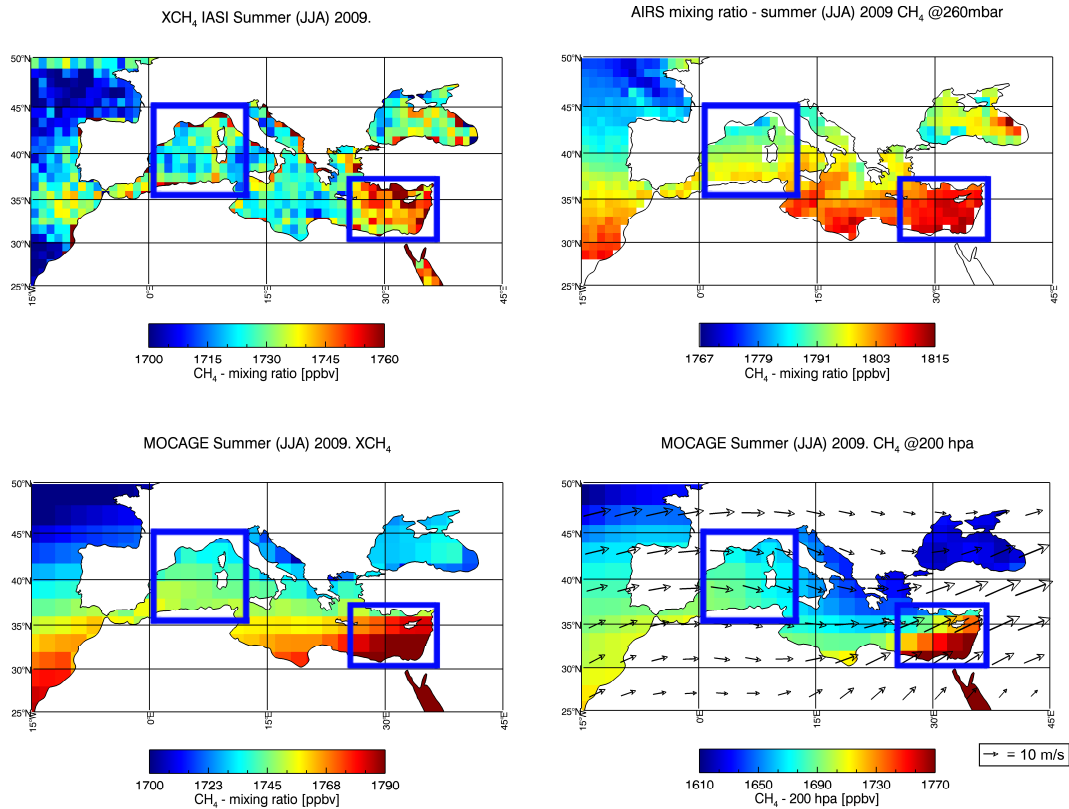
987 **Figure 4.** Same as Fig. 2, but for winter (DJF) 2009.

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992 **Figure 5.** Field of total columns of CH₄ as measured by IASI and averaged for summer (JJA) 2009

993 (top left), and field of CH₄ at 260 hPa as measured by AIRS and averaged for JJA 2009 (top right).

994 (Bottom) Same as above but as calculated by MOCAGE. Satellite data are represented in a 1°x1°

995 resolution whilst model data are shown in a 2°x2° resolution. The two blue squares in the lower left

996 Figure represent the Western and Eastern Mediterranean Basins where the measured and modelled

997 data are selected over the Mediterranean Sea. Superimposed are the horizontal winds from

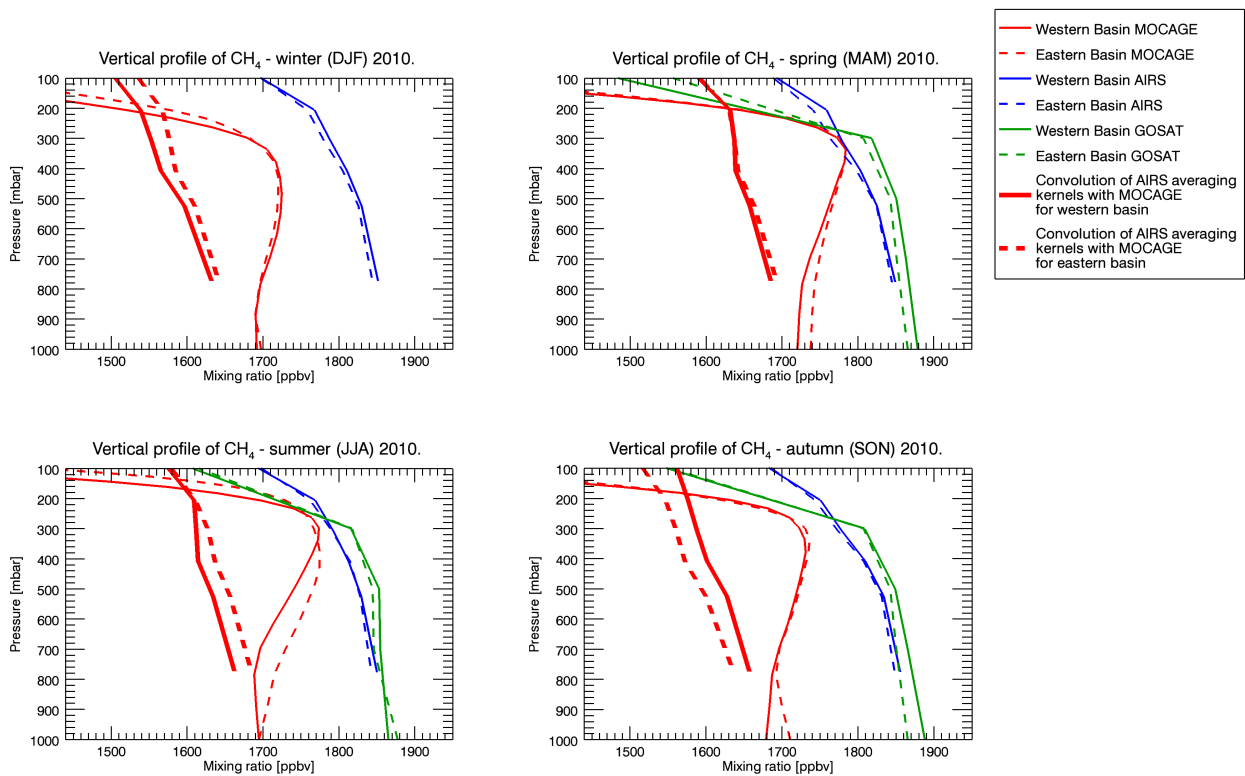
998 ARPEGE at 200 hPa averaged over the same period (bottom right). In order to highlight the CH₄

999 horizontal gradients, the range of the colour scale changes for each figure.

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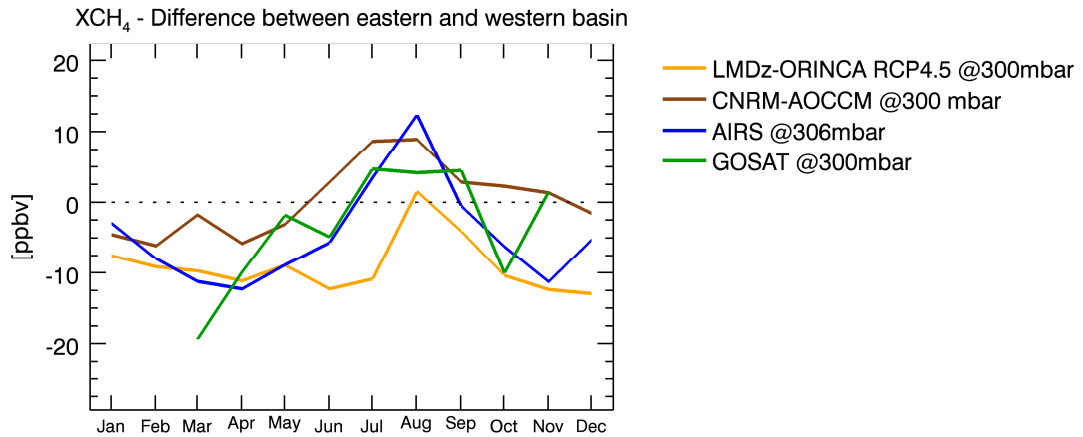
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1003 **Figure 6.** (From top to bottom and from left to right) Seasonally-averaged vertical profiles of CH₄
 1004 as measured by AIRS (blue lines) and GOSAT (green lines), and as calculated by MOCAGE (thin
 1005 red lines) over the Eastern (dashed lines) and Western (solid lines) MBs in winter, summer, spring
 1006 and autumn 2010. Also shown are the seasonally-averaged MOCAGE profiles convolved with the
 1007 AIRS averaging kernels (thick red lines) for the four seasons over the Eastern (dashed lines) and
 1008 Western (solid lines) MBs.

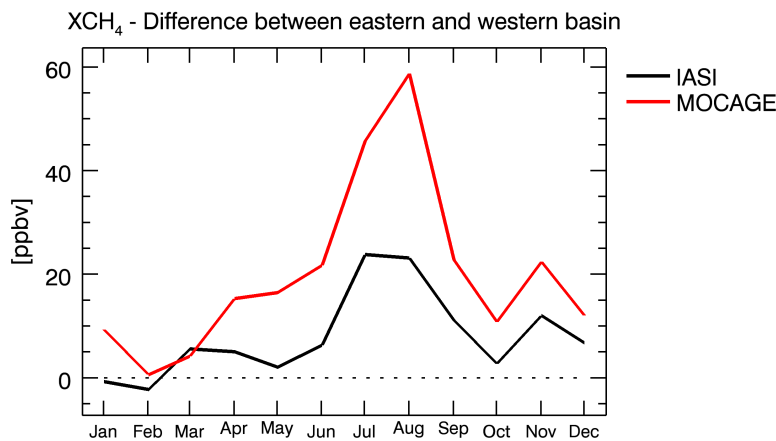
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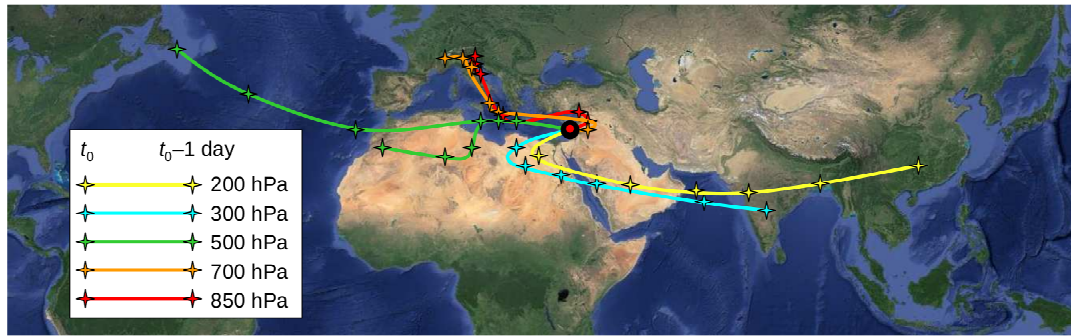
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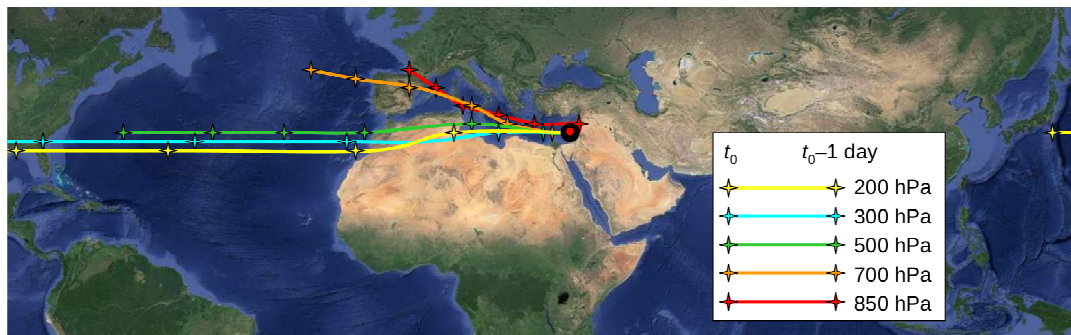
1013 **Figure 7.** (Top) Seasonal evolution of the difference in the CH₄ fields between the Eastern and
 1014 Western MB as measured by AIRS (blue line) and GOSAT (green line) at 306 and 300 hPa,
 1015 respectively and as calculated by LMDz-OR-INCA (yellow line) and CNRM-AOCCM (brown
 1016 line). (Bottom) Seasonal evolution of the difference in the CH₄ total columns between the Eastern
 1017 and Western MB as measured by IASI (black line) and as calculated by MOCAGE (red line). The
 1018 LMDz-OR-INCA and CNRM-AOCCM data sets cover the climatological period 2001-2010. The
 1019 MOCAGE and IASI data sets cover the period 2008-2011 whilst the satellite AIRS and GOSAT
 1020 data sets are representative of the year 2010.

6-Day Backtrajectories from the Eastern Mediterranean Basin
July-August 2001-2010



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6-Day Backtrajectories from the Eastern Mediterranean Basin
January-February-March 2001-2010

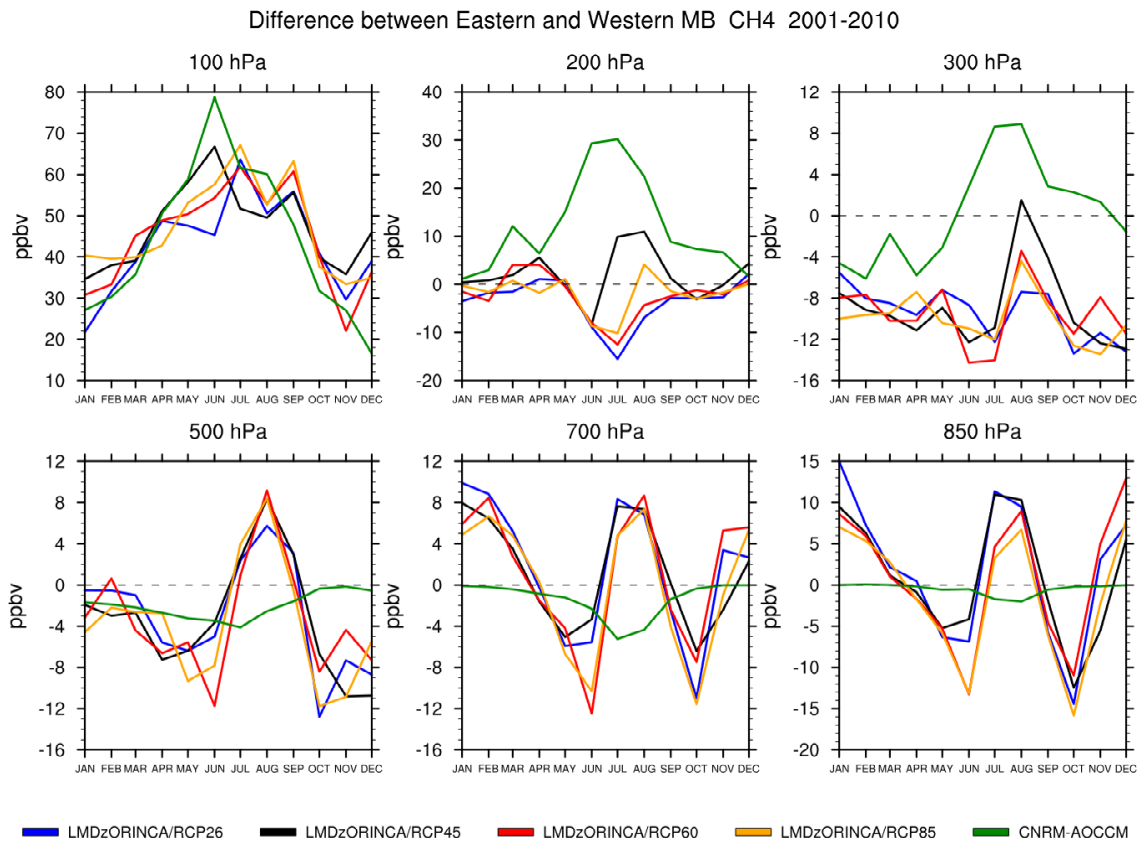


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1023 **Figure 8.** (Top) Climatological six-day back-trajectories from the point at 33° N, 35° E located in
 1024 the Eastern Mediterranean Basin (red filled circle) calculated from the British Atmospheric Data
 1025 Centre trajectory service (<http://badc.nerc.ac.uk/community/trajectory/>) from 1st July to 31st
 1026 August from 2001 to 2010 every 12 hours at 850 (red line), 700 (orange line), 500 (green line), 300
 1027 (blue line) and 200 hPa (yellow line). The position of the gravity center of each distribution at each
 1028 level is represented every 24 hours by a star. (Bottom) Same as top, but calculated from 1st January
 1029 to 31st March 2001-2010.

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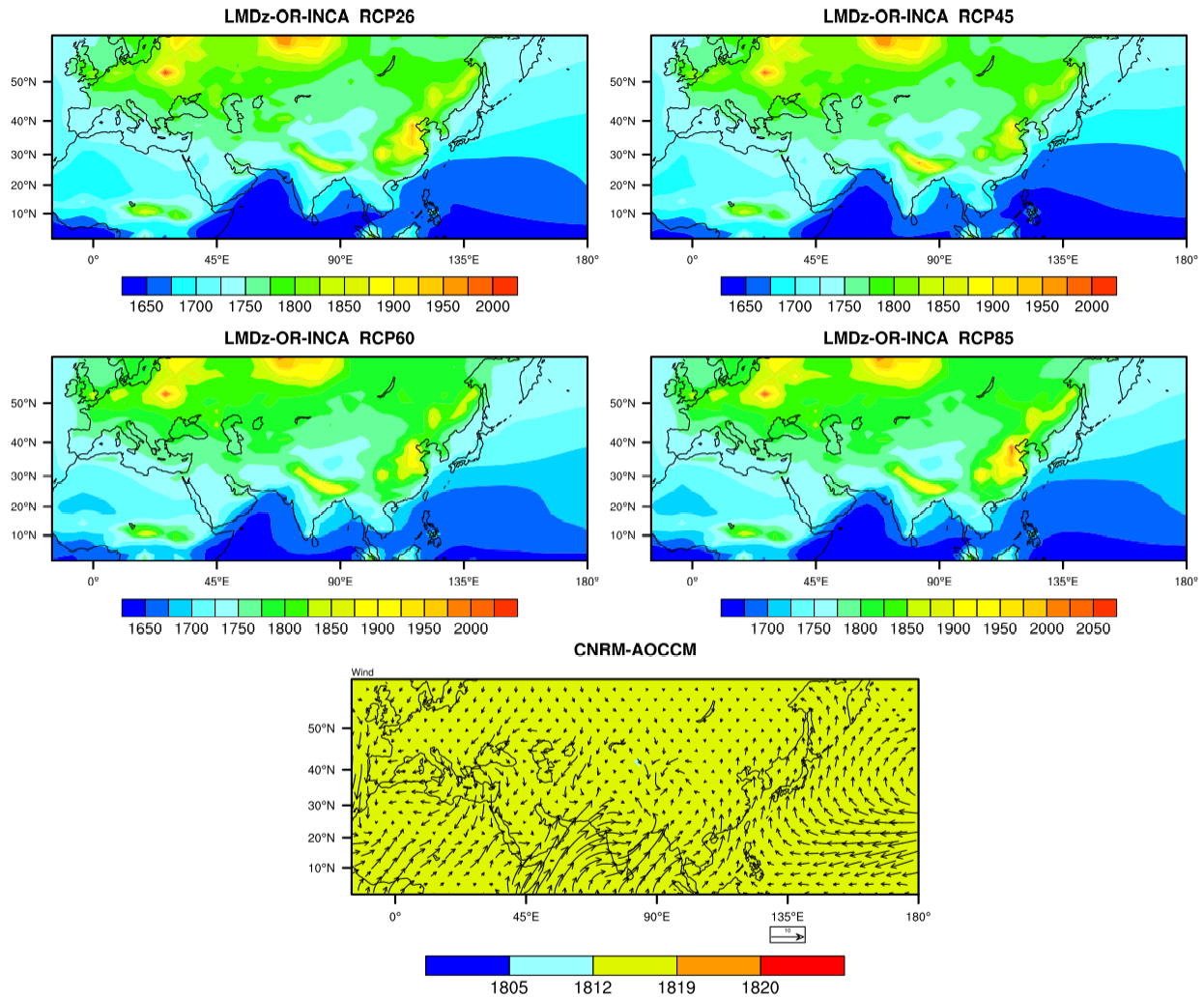
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1032 **Figure 9.** (From top to bottom and from left to right) Seasonal evolution of the difference in the
 1033 CH₄ fields between the Eastern and Western MB over the climatological period 2001-2010 at 100,
 1034 200, 300, 500, 700 and 850 hPa as calculated by CNRM-AOCCM (green) model and LMDz-OR-
 1035 INCA according to the 4 IPCC scenarios: RPCs 2.6 (blue), 4.5 (black), 6.0 (red) and 8.5 (yellow).
 1036 See section 2.2 for more details.

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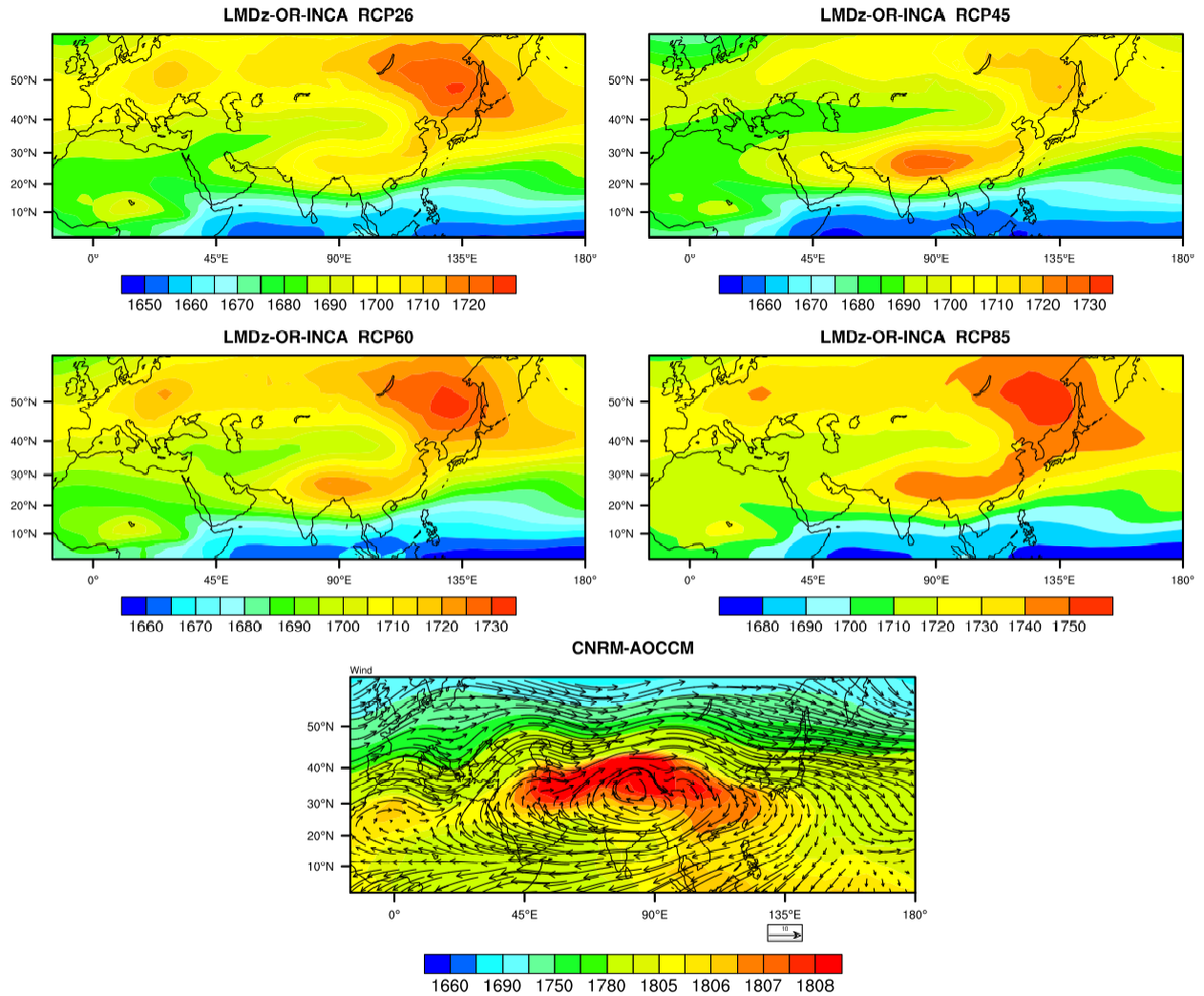
CH₄ (ppbv) surface level JJA 2001-2010

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1039 **Figure 10.** Fields of surface CH₄ as calculated by the CNRM-AOCCM model (bottom) and the
 1040 LMDz-OR-INCA model (top and centre) according to the 4 IPCC scenarios (RCPs 2.6 (top left),
 1041 4.5 (top right), 6.0 (centre left) and 8.5 (centre right)) averaged over the summer season (JJA) and
 1042 the climatological period 2001-2010. Superimposed to the CNRM-AOCCM CH₄ fields (bottom) is
 1043 the wind field at the surface averaged over the same period. Note that the range of the colour scale
 1044 changes for each figure and that the surface CH₄ for CNRM-AOCCM (bottom) is constant.

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CH₄ (ppbv) 200 hPa JJA 2001-2010

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1047 **Figure 11.** Fields of CH₄ as calculated by the CNRM-AOCCM model (bottom) and the LMDz-OR-
 1048 INCA model (top and centre) considering the 4 IPCC scenarios (RCPs 2.6 (top left), 4.5 (top right),
 1049 6.0 (centre left) and 8.5 (centre right)) at 200 hPa averaged over the summer season (JJA) and the
 1050 climatological period 2001-2010. Superimposed to the CNRM-AOCCM CH₄ fields (bottom) is the
 1051 wind field at 200 hPa averaged over the same period. Note that the range of the colour scale
 1052 changes for each figure and that the colour scale for the CNRM-AOCCM model (bottom) is non
 1053 linear.

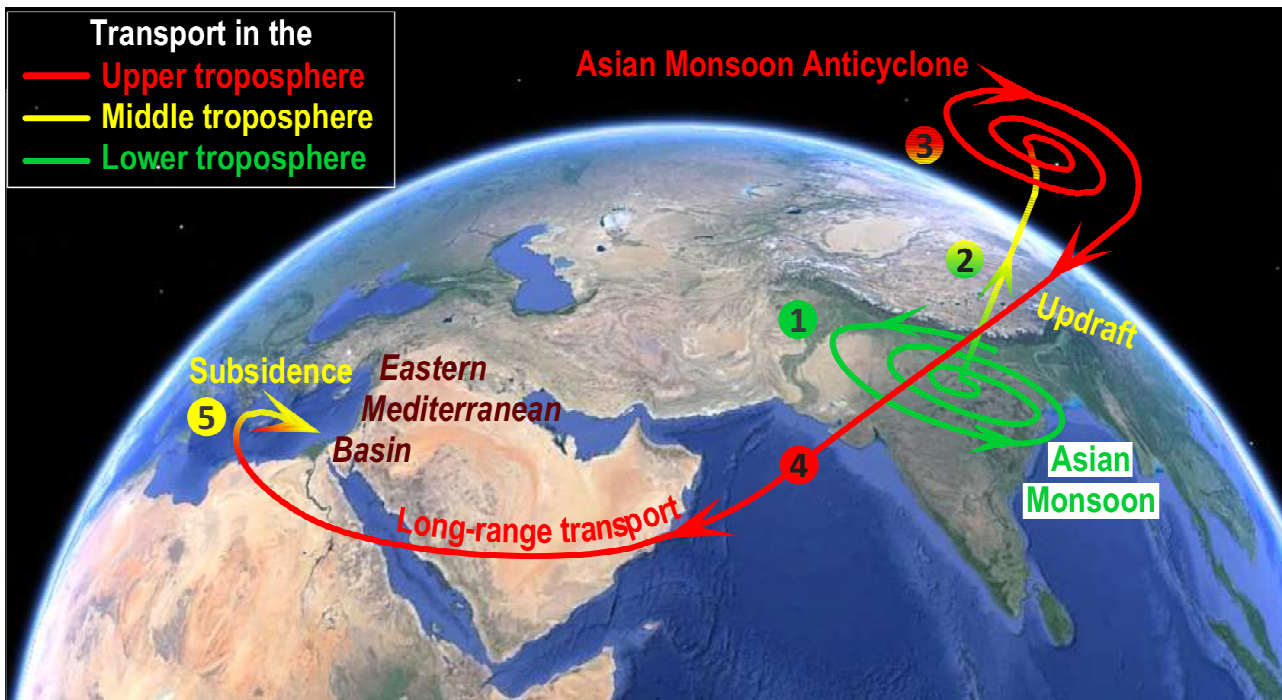
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1059 **Figure 12.** Schematic representation of the processes impacting the mid-to-upper tropospheric
 1060 pollutants, including CH_4 , above the Eastern Mediterranean Basin in summer (July-August). (1)
 1061 Trapping of lower tropospheric pollutants in the Asian monsoon. (2) Updraft of pollutants in the
 1062 Asian monsoon up to the upper troposphere. (3) Build-up of pollutants within the Asian monsoon in
 1063 the upper troposphere. (4) Large-scale re-distribution of pollutants by the Asian Monsoon
 1064 Anticyclone to the Middle East and North Africa in the upper troposphere. (5) Build-up of
 1065 pollutants through descent down to the middle troposphere above the Eastern Mediterranean Basin.

1066