# 1 Changes in black carbon emissions over Europe due to COVID-19

Nikolaos Evangeliou<sup>1,\*</sup>, Stephen M. Platt<sup>1</sup>, Sabine Eckhardt<sup>1</sup>, Cathrine Lund Myhre<sup>1</sup>, Paolo

Laj<sup>2,3,4</sup>, Lucas Alados-Arboledas<sup>5,6</sup>, John Backman<sup>7</sup>, Benjamin T. Brem<sup>8</sup>, Markus Fiebig<sup>1</sup>,

# 2 lockdowns

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6	Harald Flentje <sup>9</sup> , Angela Marinoni <sup>10</sup> , Marco Pandolfi <sup>11</sup> , Jesus Yus-Diez <sup>11</sup> , Natalia Prats <sup>12</sup> , Jean
7	P. Putaud <sup>13</sup> , Karine Sellegri <sup>14</sup> , Mar Sorribas <sup>15</sup> , Konstantinos Eleftheriadis <sup>16</sup> , Stergios Vratolis <sup>16</sup> ,
8	Alfred Wiedensohler <sup>17</sup> , Andreas Stohl <sup>18</sup>
9	
10	<sup>1</sup> Norwegian Institute for Air Research (NILU), Department of Atmospheric and Climate
11	Research (ATMOS), Kjeller, Norway.
12	<sup>2</sup> University of Grenoble Alpes, CNRS, IRD, Grenoble-INP, IGE, 38000 Grenoble, France.
13	<sup>3</sup> CNR-ISAC, National Research Council of Italy – Institute of Atmospheric Sciences and
14	Climate, Bologna, Italy.
15	<sup>4</sup> University of Helsinki, Atmospheric Science division, Helsinki, Finland.
16	<sup>5</sup> Andalusian Institute for Earth System Research (IISTA-CEAMA), Granada, Spain.
17	<sup>6</sup> Department of Applied Physics, University of Granada, Granada, Spain.
18	<sup>7</sup> Atmospheric Composition Research, Finnish Meteorological Institute, Helsinki, Finland.
19	<sup>8</sup> Laboratory of Atmospheric Chemistry, Paul Scherrer Institute, Villigen PSI, Switzerland.
20	<sup>9</sup> Deutscher Wetterdienst, Meteorologisches Observatorium Hohenpeissenberg, Albin-
21	Schwaiger-Weg 10, 82383 Hohenpeissenberg, Germany.
22	<sup>10</sup> Institute of Atmospheric Sciences and Climate, National Research Council of Italy (ISAC-
23	CNR), 40121, Bologna, Italy.
24	<sup>11</sup> Institute of Environmental Assessment and Water Research IDAEA-CSIC, C/Jordi Girona
25	18-26, Barcelona 08034, Spain.
26	<sup>12</sup> Izaña Atmospheric Research Center, State Meteorological Agency (AEMET), C/La Marina
27	20, 38001, Tenerife, Spain.
28	<sup>13</sup> European Commission, Joint Research Centre (JRC), Via Enrico Fermi 2749, Ispra (VA)
29	21027, Italy.
30	<sup>14</sup> Laboratoire de Météorologie Physique, UMR6016, CNRS/UBP, 63178 Aubière, France.
31	<sup>15</sup> El Arenosillo Atmospheric Sounding Station, Atmospheric Research and Instrumentation
32	Branch, National Institute for Aerospace Technology, 21130 Huelva, Spain.

- 33 <sup>16</sup>Environmental Radioactivity Lab, Institute of Nuclear & Radiological Sciences &
- 34 Technology, Energy & Safety, NCSR "Demokritos", Ag. Paraskevi, Athens, Greece.
- 35 <sup>17</sup>Department Experimental Aerosol and Cloud Microphysics, Leibniz Institute for
- 36 Tropospheric Research, Leipzig, Germany.
- 37 <sup>18</sup>Department of Meteorology and Geophysics, University of Vienna, UZA II,
- 38 Althanstraße 14, 1090 Vienna, Austria.
- 39
- 40 \* Corresponding author: N. Evangeliou (<u>Nikolaos.Evangeliou@nilu.no</u>)

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### 43 Abstract

44 Following the emergence of the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) responsible for COVID-19 in December 2019 in Wuhan (China) and its spread to the 45 46 rest of the world, the World Health Organization declared a global pandemic in March 2020. 47 Without effective treatment in the initial pandemic phase, social distancing and mandatory 48 quarantines were introduced as the only available preventative measure. In contrast to the 49 detrimental societal impacts, air quality improved in all countries that strict lockdowns were 50 applied, due to lower pollutant emissions. Here we investigate the effects of the COVID-19 51 lockdowns in Europe on ambient black carbon (BC), which affects climate and damages health, 52 using in-situ observations from 17 European stations in a Bayesian inversion framework. BC 53 emissions declined by 23 kt in Europe (20% in Italy, 40% in Germany, 34% in Spain, 22% in 54 France) during lockdowns compared to the same period in the previous five years, which is 55 partially attributed to COVID-19 measures. BC temporal variation in the countries enduring 56 the most drastic restrictions showed the most distinct lockdown impacts. Increased particle light 57 absorption in the beginning of the lockdown, confirmed by assimilated satellite and remote 58 sensing data, suggests residential combustion was the dominant BC source. Accordingly, in 59 Central and Eastern Europe, which experienced lower than average temperatures, BC was elevated compared to the previous five years. Nevertheless, an average decrease of 11% was 60 61 seen for the whole of Europe compared to the start of the lockdown period, with the highest 62 peaks in France (42%), Germany (21%), UK (13%), Spain (11%) and Italy (8%). Such a 63 decrease was not seen in the previous years, which also confirms an impact of COVID-19 on 64 the European emissions of BC.

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### 67 **1** Introduction

68 The identification of the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2 69 or COVID-19) in December 2019 (WHO, 2020) in Wuhan (China) and its subsequent 70 transmission to South Korea, Japan, and Europe (initially mainly Italy, France and Spain) and 71 the rest of the world led the World Health Organization to declare a global pandemic by March 72 2020 (Sohrabi et al., 2020). Although the symptoms are normally mild or not even detected for 73 most of the population, people with underlying diseases or elderly are very vulnerable showing 74 complications that can lead to death (Huang et al., 2020). Considering the lack of available 75 treatment and vaccination to combat further spread of the virus, the only prevention measures 76 included strict social, travel and working restrictions in a so-called lockdown period that lasted 77 for several weeks (mid-March to end of April 2020 for most of Europe). The most drastic 78 measures were taken in China, where the outbreak started, in Italy that faced large human losses 79 and later in the United States. Despite all these restriction, still six months after the first 80 lockdown, several countries are reporting severe human losses due to the virus (John Hopkins 81 University of Medicine, 2020).

82 Despite the dramatic health and socioeconomic consequences of COVID-19 lockdowns, 83 their environmental impact might be beneficial. Bans on mass gatherings, mandatory school 84 closures, and home confinement (He et al., 2020; Le Quéré et al., 2020) during lockdowns have 85 all resulted in lower traffic-related pollutant emissions and improved air quality in Asia, Europe 86 and America (Adams, 2020; Bauwens et al., 2020; Berman and Ebisu, 2020; Conticini et al., 87 2020; Dantas et al., 2020; Dutheil et al., 2020; He et al., 2020; Kerimray et al., 2020; Le et al., 88 2020; Lian et al., 2020; Otmani et al., 2020; Sicard et al., 2020; Zheng et al., 2020). The 89 restrictions also present an opportunity to evaluate the cascading responses from the interaction 90 of humans, ecosystems, and climate with the global economy (Diffenbaugh et al., 2020).

91 Strongly light absorbing black carbon (BC, or 'soot'), is produced from incomplete 92 combustion of carbonaceous fuels e.g. fossil fuels, wood burning, biofuels (Bond et al., 2013). 93 By absorbing solar radiation, it warms the air, reduces tropical cloudiness (Ackerman, 2000) 94 and atmospheric visibility (Jinhuan and Liquan, 2000). BC causes pulmonary diseases (Wang 95 et al., 2014a), may act as cloud condensation nuclei affecting cloud formation and precipitation 96 (Wang et al., 2016) and contributes to global warming (Bond et al., 2013; Myhre et al., 2013; 97 Wang et al., 2014a). When deposited on snow, it reduces snow albedo (Clarke and Noone, 98 1985; Hegg et al., 2009) accelerating melting. Since BC is both climate relevant and strongly

99 linked to anthropogenic activity, it is important to determine the effects of the COVID-19100 lockdowns thereon.

Here, we present a rigorous assessment of temporal and spatial changes BC emissions over Europe (including Middle East and parts of North Africa), combining in situ observations from the Aerosol, Clouds and Trace Gases Research Infrastructure (ACTRIS) network and state-of-the-art emission inventories within a Bayesian inversion. We validate our results with independent satellite data and compare them to inventories and baseline and optimized emissions calculated for previous years.

### 107 2 Methods

108 This section gives a detailed description of all datasets and methods used for the 109 calculation of COVID-19 impact. Section 2.1 describes the instrumentation of the particle light 110 absorption measurements from Aerosol, Clouds and Trace Gases Research Infrastructure 111 (ACTRIS), and the networks European Monitoring and Evaluation Program (EMEP) and 112 Global Atmosphere Watch (GAW). These measurements were used in the inverse modelling 113 algorithm (dependent measurements) and to validate the optimised (posterior) emissions of BC 114 (independent measurements). For each of the observations and stations, the source - receptor 115 matrices (SRMs), also known as "footprint emission sensitivities" or "footprints", were 116 calculated as described in section 2.2. The latter together with the observations were fed in the 117 inversion algorithm described in section 2.3. To overcome classic inverse problems (Tarantola, 118 2005), prior (a priori) emissions of BC were used in the inverse modelling algorithm calculated 119 using bottom-up approaches (section 2.4). The optimised (a posteriori) emissions of BC were 120 compared with reanalysis data from MERRA-2 (Modern-Era Retrospective Analysis for 121 Research and Applications Version 2), which are described in section 2.5, while MERRA-2 122 Ångström exponent data together with absorption Ångström exponent from the aerosol robotic 123 network (AERONET) (section 2.6) were used to examine the presence of biomass burning 124 aerosols in Europe. A description of the statistical tests and the country definitions used in the 125 paper is given in sections 2.7 and 2.8, respectively.

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#### 2.1 Particle light absorption measurements

127 The measurement sites contributing data to this paper are regional background sites 128 (except for one site in Germany) and all contribute to the research infrastructure ACTRIS, and 129 the networks EMEP and GAW. The measurement data used for the period 2015 - May 2020 130 consist of hourly-averaged, quality-checked, particle light absorption measurements. The 131 quality assurance and quality control correspond to the Level 2 requirements for ACTRIS, 132 EMEP and GAW data, as described in detail in Laj et al. (2020).

133 All absorption measurements within ACTRIS and EMEP are taken using a variety of filter-based photometers: Multi-Angle Absorption Photometers (MAAP), Particle Soot 134 135 Absorption Photometers (PSAP) Continuous Light Absorption Photometers (CLAP), and the 136 Aethalometer (AE31). Information on instrument type at the various sites are included in Table 137 1 and procedures for harmonization of measurement protocols to produce comparable data seta 138 are described in Laj et al. (2020) in detail. Zanatta et al. (2016) suggested that a mass absorption cross-section (MAC) value of 10 m<sup>2</sup> g<sup>-1</sup> (geometric standard deviation of 1.33) at a wavelength 139 140 of 637 nm can be considered to be representative of the mixed boundary layer at European 141 ACTRIS background sites, where BC is expected to be internally mixed to a large extent. 142 Assuming an absorption Ångström exponent (AAE) is equal to unity, i.e. assuming no change 143 in MAC for different sources (Zotter et al., 2017), we extrapolated the MACs at 637 nm  $(MAC_{@\lambda 1})$  to the measurement wavelengths of our study  $(MAC_{@\lambda 2})$  using the following 144 145 equation:

## 146

 $MAC_{@\lambda 2} = MAC_{@\lambda 1}(\frac{\lambda 1}{\lambda 2})^{AAE} \xrightarrow{yields} MAC_{@\lambda 2} = 10(\frac{637}{\lambda 2})^1$ (1)following Lack and Langridge (2013). The resulting MAC values for each measurement 147

148 station are shown in Table 1.

#### 149 2.2 Source – receptor matrix (SRM) calculations

150 SRMs for each of the 17 receptor sites (Table 1) were calculated using the Lagrangian 151 particle dispersion model FLEXPART version 10.4 (Pisso et al., 2019). The model releases 152 computational particles that are tracked backward in time based on 3-hourly operational 153 meteorological analyses from the European Centre for Medium-Range Weather Forecasts 154 (ECMWF) with 137 vertical layers and a horizontal resolution of 1°×1°. The tracking of BC 155 particles includes gravitational settling for spherical particles with an aerosol mean diameter of  $0.25 \,\mu\text{m}$  and a logarithmic standard deviation of 0.3 and a particle density of 1500 kg m<sup>-3</sup> (Long 156 157 et al., 2013). FLEXPART also simulates dry and wet deposition (Grythe et al., 2017), 158 turbulence (Cassiani et al., 2014), unresolved mesoscale motions (Stohl et al., 2005) and 159 includes a deep convection scheme (Forster et al., 2007). SRMs were calculated for 30 days 160 backward in time, at temporal intervals that matched measurements at each receptor site. This 161 backward tracking is sufficiently long to include almost all BC sources that contribute to surface

162 concentrations at the receptors given a typical atmospheric lifetime of 3–11 days (Bond et al.,163 2013).

164 2.3 Bayesian inverse modelling

165 The Bayesian inversion framework FLEXINVERT+ described in detail in Thompson 166 and Stohl (2014) was used to optimize emissions of BC before (January to mid-March 2020) 167 and during the COVID-19 lockdown period in Europe (mid-March to end of April 2020). To 168 show potential differences on the signal from the 2020 restrictions, emissions were optimised 169 with the same set up during the same period (January to April) in the previous five years (2015– 2019). Note that the number of stations in the inversions of 2015–2019 was slightly higher (20 170 171 stations against 15 that were used in 2020), due to different data availability. The algorithm 172 finds the optimal emissions, which lead to FLEXPART modelled concentrations that better 173 match the observations considering the uncertainties for observations, prior emissions and SRMs. Specifically, the state vector of BC concentrations,  $y_{(M \times 1)}^{mod}$ , at M points in space and 174 175 time can be modelled given an estimate of the emissions,  $x_{(N \times 1)}$ , of the N state variables 176 discretised in space and time, while atmospheric transport and deposition are linear operations described by the Jacobian matrix of SRMs,  $\mathbf{H}_{(M \times N)}$ : 177

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$$v^{mod} = \mathbf{H}x + \epsilon \tag{2}$$

179 where  $\epsilon$  is an error associated with model representation, such as the modelled transport and 180 deposition or the measurements. Since H is not invertible or may not have unique inverse, 181 according to Bayesian statistics, the inverse problem can be described as the maximization of 182 the probability density function of the emissions given the prior information and observations. 183 This is equivalent to the minimum of the cost function:

184 
$$\mathbf{J}(x) = \frac{1}{2}(x - x_b)^T \mathbf{B}^{-1}(x - x_b) + \frac{1}{2}(y - \mathbf{H}x)^T \mathbf{R}^{-1}(y - \mathbf{H}x)$$
(3)

185 where y is the vector of observed BC concentrations, x and  $x_b$  the vectors of optimized and 186 prior emissions, respectively, while **B** and **R** are the error covariance matrices that weight the 187 posterior-prior flux and observation-model mismatches, respectively. Based on the Bayes' 188 theorem, the most probable posterior emissions, x are given by the following equation 189 (Tarantola, 2005):

190

$$\mathbf{x} = \mathbf{x}_b + \mathbf{B}\mathbf{H}^{\mathrm{T}}(\mathbf{H}\mathbf{B}\mathbf{H}^{\mathrm{T}} + \mathbf{R})^{-1}(\mathbf{y} - \mathbf{H}\mathbf{x}_b)$$
(4)

Here, posterior emissions were calculated weekly between 1 January and 30 April 2020. The aggregated inversion grid (25°N–75°N and 10°W–50°E) and the average SRM for inversions are shown in Figure 1, while the measurement stations are listed in Table 1. The 194 variable grid uses high resolution at regions, where there are many stations and hence strong 195 contribution from emissions, while it lowers resolution at regions that lack measurement 196 stations following a method proposed by Stohl et al. (2010).

197 Prior emission errors **B** are correlated in space and time, but very little is known about 198 the true temporal and spatial error correlation patterns. The spatial error correlation for the 199 emissions is defined as an exponential decay over distance (we assume that emissions on land 200 and ocean are not correlated). The temporal error correlation matrix is described similarly using 201 the time difference between grid cells in different time steps. The full temporal and spatial 202 correlation matrix is given by the Kronecker product (see Thompson and Stohl, 2014). The 203 error covariance matrix for the emissions is the matrix product of correlation pattern and the 204 error covariance of the prior fluxes. We calculate the error on the emissions in each grid-cell 205 (on the fine grid) as a fraction of the maximum value out of that grid cell and the eight 206 surrounding ones.

207 The observation error covariance matrix **R** combines measurement, transport model and 208 representation errors. For the measurement errors, we use values given by the data providers. 209 Transport model errors are difficult to quantify and depend not only on the model but also on 210 the meteorological inputs. Therefore, we do not quantify the full transport error, but only the 211 part of it that can be estimated from FLEXPART, i.e. the stochastic uncertainty (see Stohl et 212 al., 2005). As regards to representation errors, we consider observation representation error and 213 model aggregation error. The observation representation error is calculated from the standard 214 deviation of all measurements available in a user-specified measurement averaging time 215 interval, based on the idea that if the measurements are fluctuating strongly within that interval 216 then their mean value is associated with higher uncertainty than if the measurements are steady 217 (Bergamaschi et al., 2010). The aggregation error is attributed to reduction of the spatial 218 resolution of the model and is calculated by projecting the loss of information in the state space 219 into the observation space (Kaminski et al., 2001). Hence, the observation error covariance 220 matrix is defined as the diagonal matrix with elements equal to the quadratic sum of the 221 measurement, transport model and measurement representation errors (Thompson and Stohl, 222 2014).

Theoretically, the algorithm can calculate negative posterior emissions, which are physically unlikely. To tackle this problem, an inequality constraint was applied on the emissions following the method of Thacker (2007) that applies the constraint as "error-free" observations:

227 
$$\hat{\mathbf{x}} = \mathbf{x} + \mathbf{A}\mathbf{P}^{\mathrm{T}}(\mathbf{P}\mathbf{A}\mathbf{P}^{\mathrm{T}})^{-1}(\mathbf{c} - \mathbf{P}\mathbf{x})$$
(5)

where **A** is the posterior error covariance matrix, **P** is a matrix operator to select the variables that violate the inequality constraint, and c is a vector of the inequality constraint, which in this case is zero.

231 We evaluated the assumptions made on the error covariance matrices for the prior emissions and the observations using the reduced  $\chi^2$  statistics (**B** and **R**). When  $\chi^2$  is equal to 232 unity, the posterior solution is within the limits of the prescribed uncertainties. The latter is the 233 234 value of the cost function at the optimum (Thompson et al., 2015). In the inversions performed here, the calculated  $\chi^2$  values were between 0.8 and 1.5 indicating that the chosen uncertainty 235 parameters are close to the ideal ones. The number of measurements used in each inversion was 236 237 equal to 12538 from 17 stations. To select the inversion that provides the most statistically 238 significant result, an evaluation of the improvement in the posterior modelled concentrations, 239 with respect to the prior ones, against the observations was performed (Figure 2). The resulting 240 values of each of the statistical measures that were performed are given in detail in Table 2. 241 Note that this is not a validation of the posterior emissions, because the comparison is only done for the observations that were included in the inversion (dependent observations), and the 242 243 inversion algorithm has been designed to reduce the model-observation mismatches. This 244 means that the reduction of the posterior concentration mismatches to the observations is 245 determined by the weighting that is given to the observations with respect to the prior emissions. 246 A proper validation of the posterior emissions is performed against observations that were not 247 included in the inversion (independent observations) in section 3.3.

#### 248 **2.4 Prior emissions**

As a priori emissions in the inversions, the ECLIPSE version 5 and 6 (Evaluating the 249 CLimate and Air Quality ImPacts of ShortlivEd Pollutants) (Klimont et al., 2017), EDGAR 250 251 (Emissions Database for Global Atmospheric Research) version HTAP v2.2 (Janssens-252 Maenhout et al., 2015), ACCMIP (Emissions for Atmospheric Chemistry and Climate Model 253 Intercomparison Project) version 5 (Lamarque et al., 2013) and PKU (Peking University) 254 (Wang et al., 2014b) were used (Figure 3). All inventories include the basic emission sectors 255 (e.g., waste burning, industrial combustion and processing, all means of transportation (aerial, 256 surface, ocean), energy conversion, residential and commercial combustion (see references 257 therein). Biomass burning emissions were adopted from the Global Fire Emissions Database, 258 Version 4.1s (GFEDv4.1s)(Giglio et al., 2013).

#### 259 **2.5** MERRA-2 (Modern-Era Retrospective Analysis for Research and Applications

#### 260 **Version 2)**

261 The MERRA-2 reanalysis dataset for BC (Randles et al., 2017) assimilates bias-corrected 262 AOD from Moderate Resolution Imaging Spectroradiometer (MODIS), Advanced Very High Resolution Radiometer (AVHRR) instruments, Multiangle Imaging SpectroRadiometer 263 264 (MISR) and Aerosol Robotic Network (AERONET) with the Goddard Earth Observing System 265 Model Version 5 (GEOS-5). BC and other aerosols in MERRA-2 are simulated with the 266 Goddard Chemistry, Aerosol, Radiation and Transport (GOCART) model and delivered in 267 hourly to monthly temporal resolution and 0.5°×0.625° spatial. The product has been validated 268 for AOD, PM and BC extensively (Buchard et al., 2017; Qin et al., 2019; Randles et al., 2017; 269 Sun et al., 2019). Ångström exponent (AE), a measure of how the AOD changes relative to the 270 various wavelength of light, is derived here from AOD469, AOD550, AOD670, and AOD865, 271 by fitting the data to the linear transform of Ångström's empirical expression:

$$\tau_{\lambda} = \tau_{\lambda_0} (\frac{\lambda}{\lambda_0})^{-a} \tag{6}$$

273 where  $\tau_{\lambda}$  is the known AOD at wavelength  $\lambda$  (in nm),  $\tau_{\lambda_0}$  is the AOD at 1000 nm, and  $\alpha$  stands 274 for AE (Gueymard and Yang, 2020).

#### **275 2.6 Absorption Ångström exponent from Aerosol Robotic Network (AERONET)**

276 data

277 Aerosol composition over Europe during the COVID-19 lockdown was confirmed using 278 the AERONET data (Holben et al., 1998). AERONET provides globally distributed 279 observations of spectral aerosol optical depth (AOD), inversion products, and precipitable 280 water in diverse aerosol regimes. The AE for a spectral dependence of 440-870 nm is related to 281 the aerosol particle size. Values less than 1 suggest an optical dominance of coarse particles 282 corresponding to dust, ash and sea spray aerosols, while values greater than one imply 283 dominance of fine particles such as smoke and industrial pollution (Eck et al., 1999). We chose 284 data from five stations covering Western, Central and Eastern Europe, for which cloud-free 285 measurements exist for the lockdown period, namely Ben Salem (9.91°E, 35.55°N), Minsk 286 (27.60°E, 53.92°N), Montsec (0.73°E, 42.05°N), MetObs Lindenberg (14.12°E, 52.21°N) and 287 Munich University (11.57°E, 48.15°N). We used Level 1.5 absorption AE (AAE) 288 measurements for the COVID-19 lockdown period (14 March to 30 April 2020).

289 2.7 Statistical measures

For the performance evaluation of the inversion results against dependent (observations that were included in the inversion) and independent observations (observations that were not included in the inversion), four different statistical quantities were used:

293 (1) Pearson's correlation coefficient:

294 
$$R_{mo} = \frac{n\sum_{i=1}^{n} m_{i}o_{i} - \sum_{i=1}^{n} m_{i}\sum_{i=1}^{n} o_{i}}{\sqrt{n\sum_{i=1}^{n} m_{i}^{2} - (\sum_{i=1}^{n} m_{i})^{2}} \sqrt{n\sum_{i=1}^{n} o_{i}^{2} - (\sum_{i=1}^{n} o_{i})^{2}}}$$
(7)

where n is sample size, m and o the individual sample points for model concentrations and observations indexed with i.

297 (2) The normalized root mean square error (nRMSE):

298 
$$nRMSE = \frac{\sqrt{\sum_{i=1}^{n} \frac{1}{n} (m_i - o_i)^2}}{o_i^{max} - o_i^{min}}$$
(8)

(3) The mean fractional bias *MFB* was selected as a symmetric performance indicator that gives
equal weights to under- or over-estimated concentrations (minimum to maximum values range
from -200% to 200%) and is defined as:

302 
$$MFB = \frac{1}{n} \frac{\sum_{i=1}^{n} (m_i - o_i)}{\sum_{i=1}^{n} (\frac{m_i + o_i}{2})}$$
(9)

303 (4) The mean absolute error was computed normalized (nMAE) over the average of all the 304 actual values (observations here), which is a widely used simple measure of error:

305 
$$nMAE = \frac{\sum_{i=1}^{n} |m_i - o_i|}{\sum_{i=1}^{n} o_i}$$
(10)

#### **306 2.8 Region definitions**

307 All country and regional masks are publicly available. Regions used for statistical 308 processing purposes were adopted from the United Nations Statistics Division 309 (https://unstats.un.org/home/). Accordingly, Northern Europe includes UK, Norway, Denmark, 310 Sweden, Finland, Iceland, Estonia, Latvia and Lithuania. Southern Europe includes Spain, Italy, 311 Greece, Slovenia, Croatia, Bosnia, Serbia, Albania and North Macedonia. Western Europe is defined by France, Belgium, Holland, Germany, Austria and Switzerland. Eastern Europe 312 313 includes Poland, Czechia, Slovakia, Hungary, Romania, Bulgaria, Moldova, Ukraine, Belarus 314 and Russia.

### 315 **3 Results**

316

#### 3.1 Optimized (posterior) emissions from Bayesian inversion

317 We performed five inversions for BC over Europe for 1st January- 30th April 2020, each 318 with different prior emissions from ECLIPSE version 5 and 6, EDGAR version HTAP v2.2, 319 ACCMIP version 5 and PKU (Figure 3). Total prior emissions of BC in Europe from the five 320 emission inventories for the period of the inversion ranged between 192-377 kt. We evaluated 321 the assumptions made on the error covariance matrices for the prior emissions and the observations using the reduced  $\chi^2$  statistic (**B** and **R**, see section 2.3). When  $\chi^2$  is equal to 322 unity, the posterior solution is within the limits of the prescribed uncertainties. The performance 323 324 of the inversions with the five different prior inventories was evaluated using four statistical 325 parameters (see section 2.7). The best performance of the inversions was achieved using 326 ECLIPSEv6 (Table 2 and Figure 2) with the smallest nRMSE (0.073) value, the largest Pearson's  $R^2$  (0.60), the closest to zero MFB value (0.03) and the smallest nMAE (714). 327 328 Therefore, all the results presented below correspond to this inversion.

329 Posterior emissions of BC were calculated to be 191 kt in the inversion domain (10°W-330 50°E, 25°N-75°N) or approximately 20% smaller than those in ECLIPSEv6 (239 kt) (Figure 331 4). Note that these numbers refer to the whole inversion domain (not only Europe) and the 332 whole study period (January - April 2020). The largest posterior differences were found in the 333 eastern part of the domain (20°E-50°E, 45°N-55°N), where emissions dropped from 35 to 29 334 kt. Emissions of BC in the western part of the inversion domain (10°W–20°E, 45°N–55°N) declined by almost 11% (from 45 to 40 kt), as those in the north part (5°W–35°E, 55°N–70°N) 335 336 that covers Scandinavian countries (from 8.7 to 6.4 kt). Finally, in the southern part  $(10^{\circ}W-$ 337 50°E, 35°N–45°N) of the domain (Spain, Italy, Greece) the posterior emissions also decreased by 21% relative to the priors (from 61 to 48 kt). The largest country decreases were seen in 338 339 France (from 14 to 8.2 kt), Italy (from 8.0 to 5.9 kt), UK (from 4.4 to 3.1 kt) and Germany 340 (from 4.5 to 4.1 kt). Surprisingly, BC emissions were slightly enhanced in Poland (from 21 to 341 23 kt), and in Spain (from 6.3 to 7.5 kt). In general, inversion algorithms reduce the mismatches 342 between modelled concentrations and observation by correcting emissions (section 2.3). If 343 decreased posterior emissions are calculated during the whole inversion period (before and 344 during the lockdowns), impact from the COVID-19 restrictions cannot be concluded and, most likely, the reduced emissions are due to errors in the prior emissions. In the next section (3.2), 345 346 we demonstrate that this decrease was due to the COVID-19 lockdowns, by comparing

posterior emissions with emissions from previous years, as well as with the respectiveemissions before and during the lockdown measures.

#### **349 3.2 Comparison with previous years**

350 We also performed inversions for 2015–2019 for the same period as the 2020 lockdowns 351 (January- April) using almost the same measurement stations and keeping the same settings. 352 The difference in BC emissions during the lockdown in 2020 (14 March to 30 April) from the 353 respective emissions during the same period in 2015–2019 (14 March to 30 April) are shown 354 in Figure 5 (a, emission anomaly) together with the gross domestic product (GDP) (Kummu et 355 al., 2020) (b), and (c) temperature anomaly from ERA-5 (Copernicus Climate Change Service 356 (C3S), 2020) for the same period as the emission anomaly. The difference in the 2020 emissions 357 of BC during the lockdown from the respective emissions in the same period in each of the 358 previous years (2015–2019) is illustrated in Supplementary Figure 1. As an independent source 359 of information, active fires from MODIS satellite product MCD14DL (Giglio et al., 2003) are 360 also shown in Figure 5a and Supplementary Figure 1.

361 Overall, BC emissions decreased by ~46 kt during the COVID-19 lockdown in the 362 inversion domain (10°W–50°E, 25°N–70°N) as compared with the same period in the previous 363 five years. We record a significant decrease in BC emissions in Central Europe (Northern Italy, 364 Austria, Germany, Spain and some Balkan countries) (Figure 5). On average, emissions were 365 23 kt lower (63 to 40 kt) over Europe during the lockdown in 2020 than in the same period of 366 2015–2019 (Figure 5). The decrease has the same characteristics when compared to each of 367 previous years since 2015 (Supplementary Figure 1) based on measurements of BC at similar regions as those used for the 2020 inversion. The countries that showed drastic reductions in 368 369 BC emissions during the lockdowns were those that suffered from the pandemic dramatically, 370 with many human losses, strict social distancing rules and consequently less transport. 371 Specifically, comparing with the previous five years, the 2020 emissions of BC during the 372 lockdowns dropped by 20% in Italy (3.4 to 2.7 kt), 40% in Germany (3.3 to 2.0 kt), 34% in 373 Spain (4.7 to 3.1 kt), 22% in France (3.5 to 2.7 kt) and remained the same or were slightly 374 enhanced in Poland (~9.2 kt), and Scandinavia (~1.2 kt). Overall, BC emissions during the 2020 lockdowns in Western Europe declined by 32% (8.8 to 6.0 kt), in Southern Europe by 42% (17 375 376 to 9.9 kt) and in Northern Europe by 29% (5.4 to 3.8 kt) as compared with the 2015–2019 377 period. BC emissions in Eastern Europe were slightly increased during the 2020 lockdown as 378 compared to the same period in the last five years (28 to 31 kt). The hot-spot emissions in 379 Eastern Europe coincide with the presence of active fires as revealed from MODIS (Figure 5a).

380 Note that these numbers correspond to BC emissions during the COVID-19 lockdown period
381 only (mid-March – April 2020).

382 Some localised areas of increased BC emissions exist in Southern France, Belgium, 383 Northern Germany and Eastern Europe (Figure 5), which are observed relative to almost every 384 year since 2015 (Supplementary Figure 1). While some hotspots in France cannot be easily 385 explained, increased emissions in Eastern European countries are likely due to increased 386 residential combustion, as people had to stay home during the lockdown. The combination of 387 the financial consequences of the COVID-19 lockdown with the relatively low GDP per capita 388 in these countries and the fact that from mid-March to end of April 2020 surface temperatures 389 in these countries were significantly lower than in previous years is suggestive of increased 390 emissions due to residential combustion. This source is most important in Eastern Europe 391 (Klimont et al., 2017). Although residential combustion can be performed for heating or 392 cooking needs in poorer countries, it is also believed to provide a more natural type of warmth 393 and a comfortable and relaxing environment. Hence, it should not be assumed as an emission 394 source in countries with lower GDPs only, especially as people spent more time at home. 395 Moreover, the prevailing average temperatures over Europe during the lockdown were below 396 15°C (Supplementary Figure 2), a temperature used as a basis temperature below which 397 residential combustion increases (Quayle and Diaz, 1980; Stohl et al., 2013).

#### **3**98 **3.3**

399 One of the basic problems when dealing with inverse modelling is that changing model, 400 observational, or prior uncertainties can have drastic impacts on posterior emissions. We addressed this issue by finding the optimal parameters, in order to have a reduced  $\chi^2$  statistic 401 402 around unity (see section 2.3). However, there are two other sources of uncertainty that, 403 although not linked with the inversion algorithm, could affect posterior emissions drastically. 404 The first is the use of different prior emissions; to estimate this type of uncertainty, we 405 performed five inversions for January to April 2020 using each of the prior emission datasets 406 (ECLIPSEv6 and v5, EDGAR HTAPv2.2, ACCMIPv5 and PKU). The uncertainty was 407 calculated as the gridded standard deviation of the posterior emissions resulting from the five 408 inversions. The second type of uncertainty concerns measurement of BC, which is defined as a 409 function of five properties (Petzold et al., 2013). However, as of today, no single instrument 410 exists that could measure all of these properties at the same time. Hence, BC is not a single 411 particle conistuent, rather an operational definition depending on the measurement technique 412 (Petzold et al., 2013). Here we use light absorption coefficients (Petzold et al., 2013) converted

Uncertainty and validation of the posterior emissions

413 to equivalent BC (eBC) using the mass absorption cross section (MAC). The MAC is 414 instrument specific and wavelength dependent. The site-specific MAC values used to convert the filter-based light absorption to eBC can be seen in Table 1. It has been reported that MAC 415 values vary from  $2 - 3 \text{ m}^2 \text{ g}^{-1}$  up to 20 m<sup>2</sup> g<sup>-1</sup> (Bond and Bergstrom, 2006). To estimate the 416 417 uncertainty of the posterior fluxes associated with the variable MAC, we performed a sensitivity study for January to April 2020 using MAC values of 5, 10 and 20 m<sup>2</sup> g<sup>-1</sup> in all stations, as well 418 419 as variable MAC values for each station (Table 1). Since these values are lognormally 420 distributed, the uncertainty is calculated as the geometric standard deviation. The impact of 421 other sources of uncertainty, such as those referring to scavenging coefficients, particle size and 422 density that are used in the model have been studied before and significantly smaller than the 423 sources of uncertainty that are considered here (Evangeliou et al., 2018; Grythe et al., 2017).

The posterior emissions are less sensitive to the use of different MACs than the use of different prior inventories (Figure 6). The relative uncertainty due to different use of MAC values was up to 20–30% in most of Europe and increases dramatically (~100%) far from the observations. The emission uncertainty of BC from the use of different priors was estimated to be up to 40% in Europe and shows very similar characteristics (same hot-spot regions and larger values where measurements lack). Overall, the combined uncertainty of BC emissions was ~60% in Europe.

Validation of top-down emissions obtained by inversion algorithms can be proper only if measurements that were not included in the inversion are to be used (independent observations). For this reason, we left out of the inversion observations from two stations (DE0054K and DE0066R, Table 1). Due to the higher measurement station density in Central Europe, we randomly selected two German stations, rather than from a country that is adjacent to regions that lack observations.

437 The prior, optimized and measured concentrations are shown in Figure 7 together with 438 MERRA-2 surface BC concentrations at the same stations. The average footprint emission 439 sensitivities are also given for the period of the lockdown. At station DE0054K, prior emissions 440 represent observations very well until the beginning of the lockdown and then fail (Figure 7). 441 On the other hand, the posterior emissions represent the variant concentrations during the 442 lockdown effectively and also manage to capture some concentration peaks, which is reflected 443 by lower *nRMSE*. Backward modelling showed that the enhanced concentrations originate 444 from Northern Germany and the Netherlands, where posterior emissions were increased 445 compared with the prior ones (Figure 4). A similar pattern was seen at station DE0066K,

446 although this station showed concentrations up to 4 mg m<sup>-3</sup> (Figure 7). Again, the optimized 447 emissions managed to represent the peaks at the end of January 2020 and at the beginning of 448 the lockdown, which is again reflected by a half *nRMSE* values and *MFB* close to zero as 449 compared to the priors. The larger concentrations during the lockdown result from increased 450 emissions over Eastern Germany, Poland and the Netherlands, as well as in oil industries in the 451 North Sea (Figure 4b). In all these regions the footprint emissions sensitivities corresponding 452 to the two independent stations were the highest.

### 453 **4 Discussion**

454 The improved air quality that Europe during the lockdown was also evident from the 455 assimilated MERRA-2 satellite-based BC data. The latter are plotted in Supplementary Figure 456 3 (left axis) for 2015–2020, together with the posterior emissions calculated in the present study 457 (right axis). For instance, weekly average concentrations of BC over Europe in MERRA-2 458 (Supplementary Figure 3, bottom). Many of the ACTRIS stations reported increased light 459 absorption in the beginning of the lockdown (e.g., Figure 7); MERRA-2 data show the same 460 patterns in France, Italy, UK and in Spain, and in all of Europe, in general. This can be explained 461 by residential combustion considering that the surface temperature during the lockdown was 462 lower than in previous years (Figure 5). The latter was confirmed by MERRA-2 reanalysis 463 Ångström Exponent (AE) parameter at 470–870 nm, which shows higher values over Central 464 and Eastern Europe during the lockdown in 2020 than in the same period of the previous years 465 (Figure 8a,b). Larger AE values confirm the presence of wood burning aerosols (Eck et al., 466 1999). The fact that during the COVID-19 lockdown, residential combustion was a significant 467 aerosol source in Europe, as compared to the previous years, was also confirmed by real-time 468 observations of absorption AE from the AERONET data in five selected stations over Europe 469 (Figure 8c). Measured absorption AE was higher during mid-March to April 2020 than in the 470 same period of the last five years.

Emissions of BC calculated with Bayesian inversion for the lockdown period dropped substantially in most of the countries that suffered from further spread of the virus and, accordingly, from strict lockdown measures, as compared to the respective emissions before the lockdowns (Supplementary Figure 3). Specifically, the decrease in France was as high as 42%, 8% in Italy, 21% in Germany, 11% in Spain and 13% in the UK. Emissions also declined in Scandinavia by 5%, although Sweden did not enforce a lockdown. Overall, a reduction in BC emissions of about 11% can be concluded for Europe as a whole due to the lockdown. 478 Stronger decreases in Eastern Europe were likely partly compensated by increased residential479 combustion in resulting from the prevailing low temperatures.

480 We report a 23 kt decrease in BC emissions in Europe during the lockdown that partially 481 resulted from the COVID-19 outbreak, as compared to the same period in all previous years 482 since 2015, based on particle light absorption measurements. We highlight these changes in BC emissions partially as a result of COVID-19 restrictions by plotting the temporal variability of 483 484 the BC emissions in the 5 previous years (2015 – 2019) for France, Italy, Germany, Spain, 485 Scandinavia and Europe (Figure 9). We record decreases in BC emissions in France, Italy, 486 Germany and Scandinavia in mid-March to April 2020, opposite to what was estimated for all 487 years between 2015 and 2019, which is obviously due to COVID-19. The UK and Spain showed 488 a similar decrease in mid-March to April 2020 emissions as in all previous years (2015–2019). 489 However, the estimated posterior BC emissions during the 2020 lockdowns were significantly 490 lower than those of the same period in any of the previous years. Overall, emissions declined 491 by 20% in Italy, 40% in Germany, 34% in Spain, 22% in France and remained the same and 492 slightly enhanced in Scandinavia or Poland as compared to those of the last five years.

### 493 **5** Conclusions

494 The impact of the COVID-19 lockdowns over Europe on the BC emissions, in response 495 to the pandemic was assessed in the present manuscript. Particle light absorption measurements 496 from 17 ACTRIS stations all around Europe were rapidly gathered and cleaned to produce a 497 high-quality product. The latter was used in a well-established Bayesian inversion framework 498 and BC emissions were optimised over Europe to better capture the observations. However, 499 one should be careful not to overinterpret the emission changes at regional scales, due to the 500 poor station data density used and the high resolution timesteps of the inversions (weekly 501 posterior emissions). We calculate that the optimised (posterior) BC emissions declined from 502 63 to 40 kt (23%) during the lockdowns over Europe, as compared to the same period in the 503 previous five years (2015-2019). The largest reductions were calculated for countries that 504 suffered from the pandemic dramatically, such as Italy (3.4 to 2.7 kt), Germany (3.3 to 2.0 kt), 505 Spain (4.7 to 3.1 kt), France (3.5 to 2.7 kt). BC emissions in Western Europe during the 2020 506 lockdowns were decreased from 8.8 to 6.0 kt (32%), in Southern Europe from 17 to 9.9 kt 507 (42%) and in Northern Europe from 5.4 to 3.8 kt (29%) as compared to the same period in the 508 last five years. BC emissions were slightly enhanced in Eastern Europe (from 28 to 31 kt) and 509 remained unchanged in Scandinavia during the lockdown, due to increased residential

510 combustion, as people had to stay home and temperatures at that time were the lowest of the 511 last five years. The presence of wood burning aerosols during the lockdowns was confirmed by 512 large MERRA-2 AE, as well as from absorption AE measurements from AERONET that were 513 higher in the lockdowns than in the same period of the last five years. The impact of the 514 European lockdowns on BC emissions was also confirmed by a 11% decrease of the posterior 515 emissions over Europe during the lockdowns, as compared to the period before, opposite to 516 what was calculated in the previous years, which is obviously due to COVID-19. This decrease 517 was more pronounced in France (42%), Italy (8%), Germany (21%), Spain (11%), UK (13%) 518 and in Scandinavian countries (5%). The full impact of the disastrous pandemic will likely take 519 years to assess. Nevertheless, with COVID-19 cases once again increasing in many countries, 520 the information presented here are essential to understand the full health and climate impacts

521 of lockdown measures.

522 *Data availability*. All measurement data and model outputs used for the present publication are 523 open and can be downloaded from <u>https://doi.org/10.21336/gen.b5vj-sn33</u> or upon request to 524 the corresponding author. All prior emission datasets are also available for download. ECLIPSE 525 emissions can be obtained from 526 <u>http://www.iiasa.ac.at/web/home/research/researchPrograms/air/Global\_emissions.html</u>,

527 EDGAR version HTAP\_V2.2 from <u>http://edgar.jrc.ec.europa.eu/methodology.php#</u>, ACCMIP 528 version 5 from <u>http://accent.aero.jussieu.fr/ACCMIP\_metadata.php</u>) and PKU from 529 <u>http://inventory.pku.edu.cn</u>. FLEXPART is open in public and can be downloaded from 530 <u>https://www.flexpart.eu</u>, so as FLEXINVERT+ from <u>https://flexinvert.nilu.no</u>. MERRA-2 re-531 analysis data can be obtained from <u>https://disc.gsfc.nasa.gov</u>, so as AERONET measurements 532 from https://aeronet.gsfc.nasa.gov.

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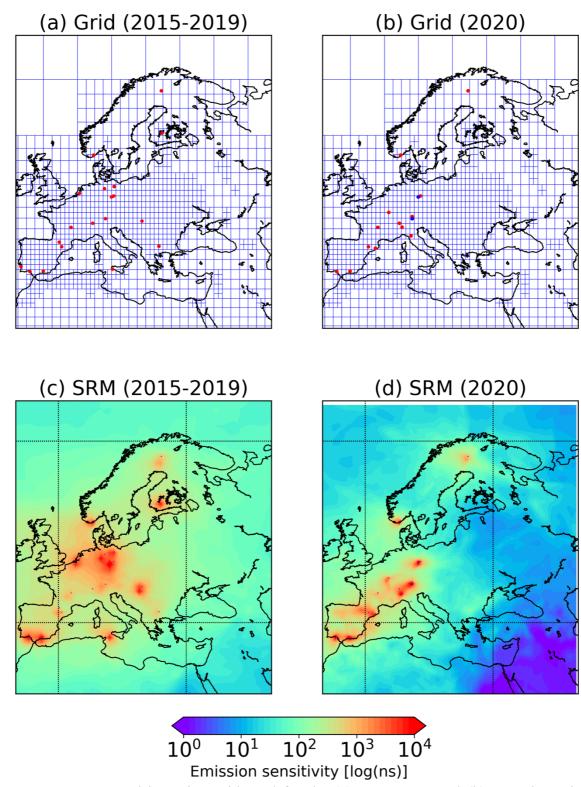
#### 810 TABLES & FIGURES

**Table 1.** Observation sites from the ACTRIS platform used to perform the inversions (dependent observations) and to validate the posterior emissions (independent observations) (the altitude indicates the sampling height in meters above sea level). Multi-Angle Absorption Photometers (MAAP) were used at all sites, except El Arenosillo (ES0100R) where a Continuous Light Absorption Photometer (CLAP) was used, Birkenes (NO0002R), where a Particle Soot Absorption Photometer (PSAP) and Observatoire Perenne de l' Environnement (FR0022R) and Zeppelin (NO0042G) where Aethalometers (AW31) were used.

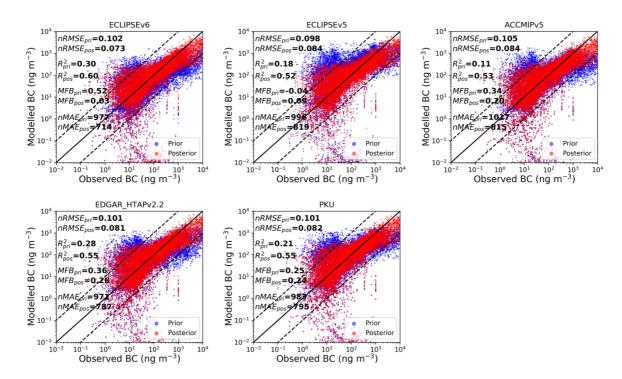
Name	Latitude	Longitude	Altitude	Туре	Wavelength (nm)	$\frac{MAC_{@637}}{(m^2 g^{-1})}$
Jungfraujoch (CH0001G)	46.55	7.99	3578	Dependent	637	10
Hohenpeissenberg (DE0043G)	47.80	11.01	985	Dependent	660	9.65
Melpitz (DE0044K)	51.53	12.93	86	Dependent	670	8.78
Zugspitze- Schneefernerhaus (DE0054R)	47.42	10.98	2671	Independent	670	9.51
Leipzig- Eisenbahnstrasse (DE0066K)	51.35	12.41	120	Independent	670	9.51
Izaña (ES0018G)	28.41	-16.50	2373	Dependent	670	9.51
Granada (ES0020U)	37.16	-3.61	680	Dependent	670	9.51
Montsec (ES0022R)	42.05	0.73	1571	Dependent	670	9.51
El Arenosillo (ES0100R)	37.10	-6.73	41	Dependent	652	13.64
Montseny (ES1778R)	41.77	2.35	700	Dependent	670	8.48
Pallas (FI0096G)	67.97	24.12	565	Dependent	637	10.00
Observatoire Perenne de l' Environnement (FR0022R)	48.56	5.51	392	Dependent	880	7.24
Puy de Dôme (FR0030R)	45.77	2.96	1465	Dependent	670	9.51
Ispra (IT0004R)	45.80	8.63	209	Dependent	880	6.96
Mt Cimone (IT0009R)	44.18	10.70	2165	Dependent	670	9.51
Birkenes II (NO0002R)	58.39	8.25	219	Dependent	660	7.59
Zeppelin mountain (NO0042G)	78.91	11.89	474	Dependent	880	7.24

- 820 Table 2. Statistical measures (*RMSE*, Pearson's  $R^2$ , *MFB* and *nMAE*) for each of the prior
- 821 and posterior concentrations against dependent observations (observations that were used in the
- 822 inversion algorithm) for BC (eBC). Note that the inversion using ECLIPSEv6 prior emission
- 823 dataset gave the best agreement with the observations and therefore the results of this inversion
- 824 are presented here.

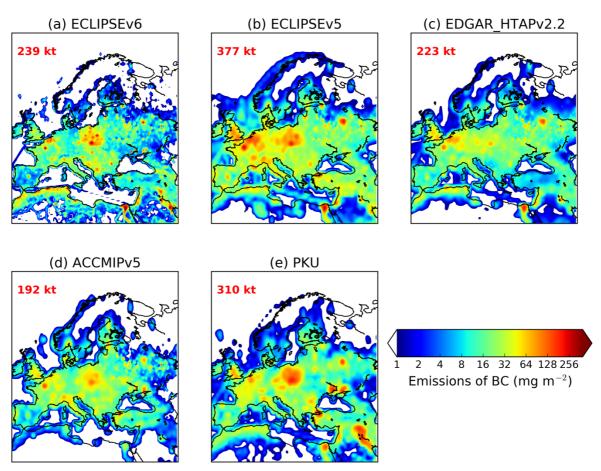
	nRMSE	Pearson's R <sup>2</sup>	MFB	nMAE
Prior ECLIPSEv6	0.102	0.30	0.52	997
Prior ECLIPSEv5	0.098	0.18	-0.04	996
Prior EDGAR_HTAPv2.2	0.105	0.11	0.34	1017
Prior ACCMIPv5	0.101	0.28	0.36	971
Prior PKU	0.101	0.21	0.25	983
Posterior ECLIPSEv6	0.073	0.60	0.03	714
Posterior ECLIPSEv5	0.084	0.52	0.09	819
Posterior EDGAR_HTAPv2.2	0.084	0.53	0.20	815
Posterior ACCMIPv5	0.091	0.55	0.26	787
Posterior PKU	0.082	0.55	0.24	795



827 828 Figure 1. Aggregated inversion grid used for the (a) 2015–2019 and (b) 2020 inversions, 829 respectively. The dependent measurements that were used in the inversion were taken from 830 stations highlighted in red. The two independent stations that were used for the validation are 831 shown in blue. (c, d) Footprint emission sensitivity (i.e. SRM) averaged over all observations 832 and time steps for each of the inversions. Red points denote the location of each measurement 833 site.



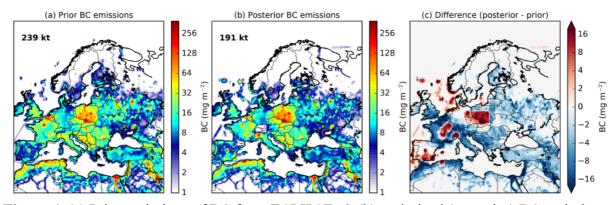
834 835 Figure 2. Scatter plots of prior and posterior concentrations against dependent observations (observations that were included in the inversion framework) from ACTRIS from January to 836 837 April 2020. Four statistical measures (*nRMSE*, Pearson's  $R^2$ , *MFB* and *nMAE*) were used to 838 assess the performance of each inversion using five different prior emission inventories for BC 839 (ECLIPSEv5, v6, ACCMIPv5, EDGAR\_HTAPv2.2 and PKU).



### **PRIOR EMISSIONS (JAN-APR 2020)**

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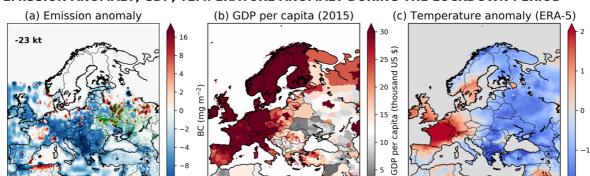
842 Figure 3. Prior emissions of black carbon (BC) used in the inversions. BC emissions from 843 anthropogenic sources were adopted from ECLIPSE version 5 and 6 (Evaluating the CLimate 844 and Air Quality ImPacts of ShortlivEd Pollutants) (Klimont et al., 2017), EDGAR (Emissions 845 Database for Global Atmospheric Research) version HTAP v2.2 (Janssens-Maenhout et al., 846 2015), ACCMIP (Emissions for Atmospheric Chemistry and Climate Model Intercomparison 847 Project) version 5 (Lamarque et al., 2013) and PKU (Peking University) (Wang et al., 2014b). 848 Biomass burning emissions of BC from Global Fire Emissions Database (GFED) version 4.1 849 (Giglio et al., 2013) were added in each of the aforementioned inventories.



851 852

**Figure 4.** (a) Prior emissions of BC from ECLIPSEv6, (b) optimized (posterior) BC emissions

- 853 after processing the ACTRIS data into the inversion algorithm, and (c) difference between
- 854 posterior and prior emissions. All the results correspond to the inversion yielding the best results
- 855 (Table 2 and Figure 2).
- 856



## EMISSION ANOMALY, GDP, TEMPERATURE ANOMALY DURING THE LOCKDOWN PERIOD

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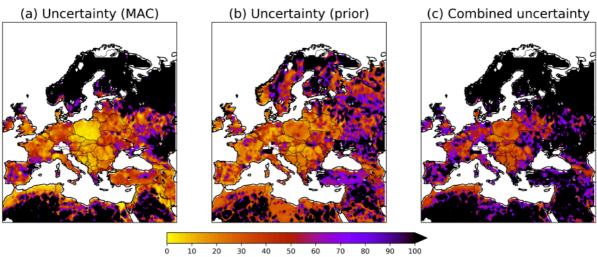
Figure 5. (a) Difference in posterior BC emissions during the lockdown (14 March to 30 April 859 2020) in Europe from the respective emissions during the same period in 2015 - 2019, (b) GDP 860 from Kummu et al. (2020), and (c) temperature anomaly from ERA-5 (Copernicus Climate Change Service (C3S), 2020) for the same period as the emission anomaly. The base GDP value 861 862 below which a low income can be assumed was set to 12 thousand US dollars. Active fires from

863 MODIS are plotted together with emission anomaly (green dots).

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Degrees Celsiu:

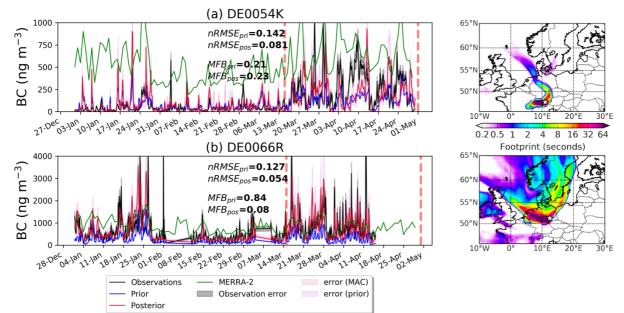


20 30 40 50 60 70 80 9 Relative emission uncertainty (%)

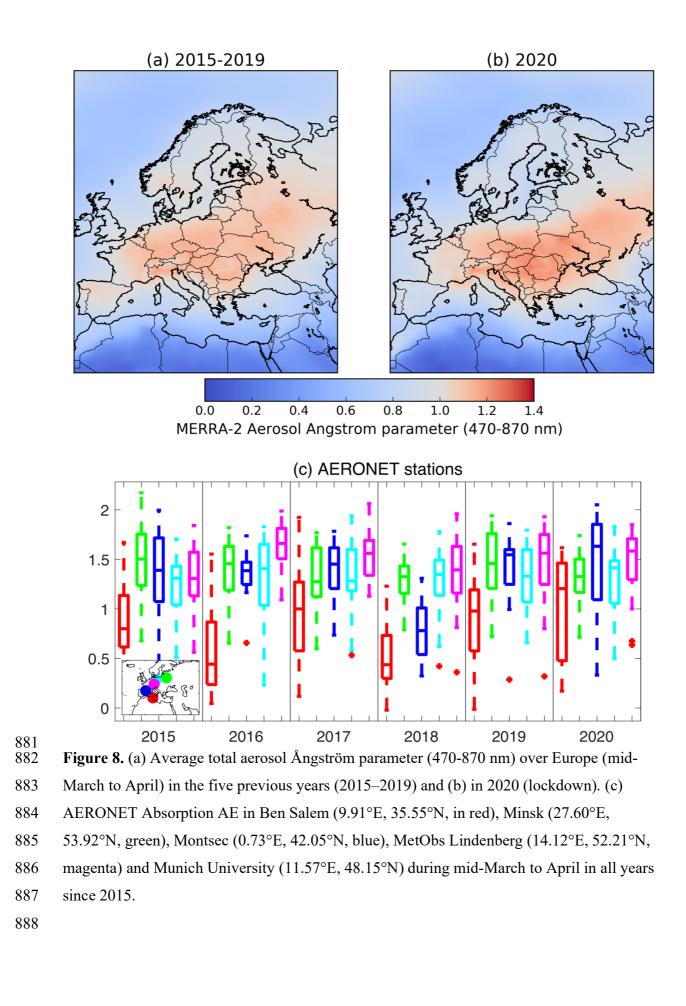
865 866 Figure 6. (a) Uncertainty of BC emissions due to the use of variable MAC values to convert 867 from aerosol absorption to eBC concentrations that are used by the inversion algorithm. (b)

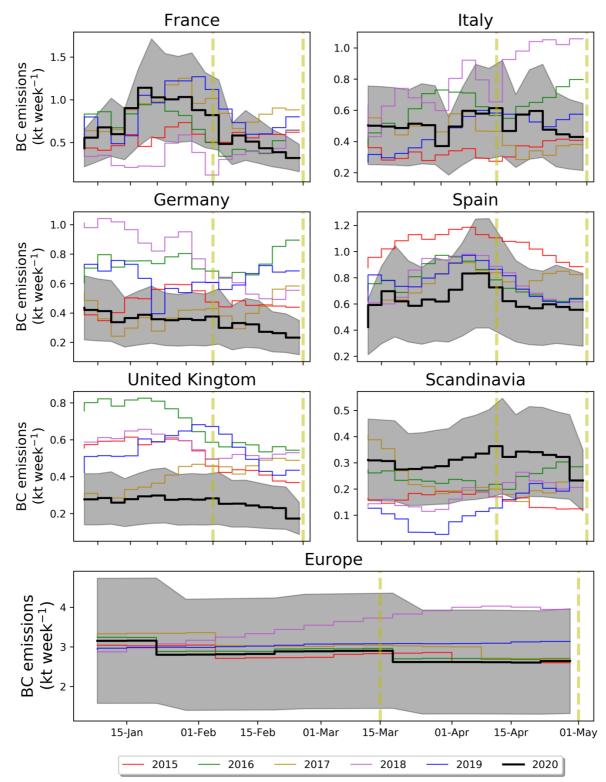
Uncertainty due to the use of five different prior emissions inventories for BC. (c) Combined 868

- uncertainty. 869
- 870



871 872 Figure 7. Prior and posterior BC concentrations at DE0054K and DE0066R stations that were 873 not included in the inversion are compared with observations. The validation is done by 874 calculating the *nRMSEs* and *MFBs* for the prior and posterior concentrations. The uncertainty of the observations is also given together with the posterior uncertainties in the 875 876 concentrations calculated from the use of different MAC and prior emissions. For comparison, 877 we plot the concentrations from MERRA-2 at the same two stations. The vertical dashed lines 878 denote the period of the lockdown in most of Europe. On the right, the average footprint 879 emission sensitivities are given at each independent station for the period of the lockdown.







889 890 Figure 9. Posterior BC emissions in the most highly affected European countries (France, Italy, 891 Germany, Spain and UK), Scandinavia and Europe from the COVID-19 pandemic (2020). 892 Posterior BC emissions for every year since 2015 are also plotted in the same temporal 893 resolution to show changes in BC emissions characteristics during the 2020 COVID-19 894 pandemic. The grey shaded area corresponds to the BC emission uncertainty, while the vertical 895 yellow dashed lines correspond to the beginning and end of the 2020 lockdown.