The impact of shipping emissions on air pollution in the Greater North Sea region. Part II: Scenarios for 2030

Volker Matthias[1], Armin Aulinger[1], Anna Backes[1], Johannes Bieser[1], Beate Geyer[1], Markus Quante[1], and Malte Zeretzke[2]

[1]Helmholtz-Zentrum Geesthacht, Institute of Coastal Research, Max-Planck-Straße 1, 21502 Geesthacht, Germany
[2]DNV-GL, Brooktorkai 18, 20457 Hamburg, Germany

Correspondence to: V. Matthias
volker.matthias@hzg.de

Abstract. Scenarios for future shipping emissions in the North Sea have been developed in the framework of the Clean North Sea Shipping project. The effects of changing NO_x and SO_2 emissions were invesigated with the chemistry transport model CMAQ for the year 2030 in the North Sea area. It has been found that, compared to today, the contribution of shipping to the NO_2 and O_3 concentrations will increase due to the expected enhanced traffic by more than 20% and 5%, respectively, by 2030 if no regulation for further emission reductions will be implemented in the North Sea area. $PM_{2.5}$ will decrease slightly because the sulphur contents in ship fuels will be reduced as international regulations foresee. The effects differ largely between regions, seasons and date of the implementation of stricter regulations for NO_x emissions from new built ships.

1 Introduction

Shipping is an important contributor to air pollution in coastal areas. More than 90% of the global trade is done with ships. The total global transport work by ships (in tonne miles) has been tripled since the mid 1980s (Smith et al., 2014), corresponding to an average growth rate of 4% p.a., and the forecasts for the future are in the same order of magnitude (Smith et al., 2014). The North Sea is one of the areas with the highest ship densities in the world. Europe's three biggest harbours in Rotterdam, Hamburg and Antwerp are located in the North Sea region. At any time about 3000 ships are sailing in the North Sea (Aulinger et al., 2015). The steady increase in number and size of ships leads to an increasing contribution of ships to air pollution in North Sea coastal areas (Matthias et al., 2010; Hammingh et al., 2012; Jalkanen et al., 2012; Aulinger et al., 2015). Compared to other

20   modes of transport like trucks or trains, big container ships or tankers are very efficient in terms of fuel use per ton mile. However, NO_x, SO_2 and PM emissions are comparably high because of less strict regulations for the emissions of these pollutants from ships. This problem has already been recognized years ago leading to stricter regulations in some areas, the so called Emission Control Areas (ECAs). These regulations are the results of an agreement within the International Maritime

25   Organization (IMO) and they are laid down in MARPOL Annex VI (International Maritime Organisation, 2008). EU has implemented these rules in their directives 1999/32/EC (European Union, 1999) and 2012/33/EU (European Union, 2012) for the North and the Baltic Seas which are Sulphur Emission Control Areas (SECAs). This means that the fuel burned in these areas must not contain more than 0.1% sulphur (S) (until 31 Dec 2014 1.0% S). If fuels with higher sulphur content are

30   used, the exhaust gas has to be cleaned until it does not contain more sulphur than exhaust gas from a low-sulphur fuel. As a consequence, ships use low sulphur fuels or installed scrubbers on board that clean the exhaust gas from sulphur and other contaminants. Outside SECAs, the allowed sulphur content in ship fuels is currently at 3.5%, but it will be reduced to 0.5% in 2020 or 2025 at the latest. The exact date will be decided in 2018 when the availability of ship fuel containing less than

35   0.5% S will be reviewed.

   Nitrogen oxide emissions from ships are also regulated in MARPOL Annex VI. Since the year 2000 the NO_x limits for ships built after 1 January 2000 are 17 g/kWh NO_x (Tier I regulation) for slow speed engines (< 130 revolutions per minute (rpm)) and 9.8 g/kWh NO_x for high speed engines (> 2000 rpm). A power law following $45 \cdot rpm^{-0.2}$ g/kWh NO_x sets the maxiumum emissions at

40   intermediate engine speeds. In 2010 the second step (Tier II) of the NO_x regulations came into force with an emission factor of 14.4 g/kWh NO_x for new ships with slow speed engines. For high speed engines 7.7 g/kWh NO_x, for engines in between $44 \cdot rpm^{-0.23}$ g/kWh NO_x is valid. The third step (Tier III), where NO_x limits will be further reduced to 3.4 g/kWh NO_x (2.0 g/kWh NO_x for high speed engines, $9 \cdot rpm^{-0.2}$ g/kWh NO_x for engines in between), is planned for 2016 for the ECAs

45   around North America. However, this will only be applied to new ships sailing in designated Emission Control Areas while Tier I and Tier II represent global limits. The North Sea and the Baltic Sea are in discussion to become such an ECA for NO_x. However, Tier III rules will only be valid for ships built after the designation date (International Maritime Organisation, Marine Environment Protection Committee, 2014). Until now, it is unclear when this date will be.

50   In the European project Clean North Sea Shipping (CNSS) different technologies capable of reducing air emissions from ships in the North Sea were investigated. Among them are scrubbers that reduce sulphur emissions, catalysts that reduce NO_x emissions and the use of alternative fuels like liquefied natural gas (LNG). In order to estimate the effect of these technologies and of legislation on NO_x and SO_2 emissions from ships, emission scenarios were developed for the year 2030. These

55   scenarios consider the same development of the world fleet but different developments in legislation and the use of alternative fuels. The basis is a detailed emission inventory for the year 2011 which is

built upon AIS (Automatic Identification System) ship positions and a detailed ship characteristics data base (Aulinger et al., 2015). The scenarios are implemented as modified emission inventories for the year 2030. The inventories serve as input for the chemistry transport model CMAQ that is
60   setup for the North Sea region. CMAQ calculates transport and transformation of the emitted pollutants and finally yields concentration maps that illustrate the impact of shipping emissions on the air quality in the North Sea region.

2   Ship emission inventories

2.1   Reference emissions

65   The basis for the ship fleet and the ship movements on the North Sea is a data set with AIS positions of ships for the entire year 2011 combined with a ship characteristics data base that includes all ships given in the AIS data set. The data is used to calculate the energy demand of individual ships depending on the installed engine and their actual velocity. From this, fuel use as well as NO_x, SO_2, CO_2, CO, Hydrocarbon (HC), and Particulate Matter (PM) emissions are calculated
70   with load-dependent emission factors for the different species. For the first time, load dependent emission factors resulting from test bed measurements of about 250 different ship engines were used to calculate a ship emission inventory. For the details, the reader is directed to the accompanying paper by Aulinger et al. (2015).

2.2   Scenario description

75   The purpose of scenarios is to describe plausible and possible future developments. Scenarios are often used to describe the boundaries of possible future situations, e.g. a worst case and a best case. In our study we decided to create scenarios that describe the future development of policy and technology regarding exhaust gas emissions from ships in the North Sea area. We adopted the methodology described in (Eyring et al., 2005) for our scenarios and distinguish between traffic demand and future
80   technological and legislative developments. However, because we focus on the implementation of a NO_x emission control area in the North Sea, we take only one scenario for the fleet development into account as a basis as it is described in publications from IMO (Buhaug et al., 2009; Smith et al., 2014) and Det Norske Veritas (Det Norske Veritas, 2012). Taking multiple possible developments of the world trade into account would add too much complexity to the scenarios.
85   In brief, our fleet development scenario assumes an increase in the number of bigger ships while the number of smaller ships decreases in the North Sea area. This leads to an increase in ship number by 1.0% p.a. and an increase in transported cargo of 2.5% p.a. Additional to this increase in ship number it is assumed that per year 2.5% of all ships are replaced by new ones, no matter of what size they are. Older ships are replaced first. The main techniques under investigation are Liquefied Nat-
90   ural Gas (LNG) as an alternative fuel for shipping and end-of-the-pipe technologies like scrubbers

3

and Selective Catalytic Reduction (SCR) to reduce sulphur dioxide and nitrogen oxide emissions. The main drivers for changes in the use of ship fuels and in the amount of emissions to air are on the one hand regulations, and here mainly what is written in MARPOL Annex VI (International Maritime Organisation, 2008), and on the other hand the price of different fuels. Therefore, the main

95 scenarios include strict and less strict legislations as one axis and the price of LNG compared to Marine Gas Oil (MGO) or Heavy Fuel Oil (HFO) as the second axis. Some regulations in MARPOL Annex VI (those related to NO_x emissions) are only valid for newly built ships after a certain date, depending on the region where the ECA is located. The earliest date when Tier III NO_x regulations will come into force is 1 January 2016. For the North Sea and the Baltic Sea, it is likely that Tier III

100 NO_x regulations will be implemented significantly later than 2016.

Tier III NO_x regulations apply only to new ships. Therefore, it needs some time until a considerable number of ships in the fleet will have reduced NO_x emissions. Those regulations related to the sulphur content in ship fuels apply to all ships, and should have immediate effects on the total emissions of sulphur oxides. To particularly take into account the long term effects of new ships following Tier

105 III regulations, the year 2030 is used in the scenarios as target year. The development of the world fleet until 2030 compared to the reference year 2011 is considered.

These drivers are combined into six scenarios that can be arranged in a coordinate system with legislation on the x-axis and LNG price on the y-axis (see Fig. 1). Different implementation dates for NO_x Tier III rules (2016 and 2021) are chosen.

110 The stories behind these scenarios can be described as follows:

Scenario No ECA

The global economy suffers from low GDP growth rates and in order to avoid additional costs for the shipping industry some regulations will not be implemented (global SO_x limit, NO_x limits in ECAs). This can be considered as the worst case scenario, however 0.1% S in fuels in ECAs will

115 still be implemented.

Scenarios ECA SCR 16 and ECA SCR 21

All regulations currently given in MARPOL Annex VI will be in force. The global sulphur limit of 0.5% S in fuel will be in force by 2020, in ECAs a sulphur limit of 0.1% S will be implemented since 2015. A NO_x emission control area will be implemented in the North and Baltic Seas. Two

120 different years, 2016 and 2021, are considered as implementation dates. LNG is expensive and the LNG infrastructure is not built up to provide LNG to many ships. Therefore ship owners will prefer low sulphur fuels and catalysts (SCR) or exhaust gas recirculation systems (EGR) to comply with the rules. Some will use scrubbers only, if they do not have to follow the Tier III regulations (older ships).

125 *Scenario ECA LNG 16 and ECA LNG 21*

The legislation is the same as in scenarios ECA SCR 16 and ECA SCR 21 but LNG will be the

4

cheaper solution to comply with the rules. In 2030, about 6000 ships in the North Sea will run on LNG. Ships that sail more than 50% of the time in the North and Baltic Seas will preferably use LNG. Some newer ships will also be retrofitted with LNG engines.

130 *Scenario ECA opt*

This is built on scenario ECA SCR but it assumes that the strict rules for NO_x emissions for new-buildings will also apply to older ships in 2030. They will then be retrofitted with exhaust gas cleaning systems in order to follow these rules. This is regarded as the best case scenario and illustrates the reduction potential. However, exhaust gas cleaning systems increase the fuel consumption, 135 so the fuel use in this scenario will be higher than in the previous scenarios in which only parts of the ships are equipped with exhaust gas cleaning.

2.3 Future shipping emissions

The emission inventories that were constructed as input for the CMAQ model were developed from the ship emission inventory for 2011 which is based on AIS data and ship characteristics data. First, 140 the fleet development was applied. Then, the new emissions were calculated by using modified emission factors for the specific emissions of the ships. All emission factors are given in g/kWh for the different substances under investigation, they have been reduced depending on the scenario and taking into account the age of the different ships in the scenarios ECA SCR and ECA LNG. In particular, in all scenarios it has been taken into account that a fraction of the older ships, that do 145 not have to follow any of the Tier rules for NO_x emissions, are taken out of service until 2030 and will be replaced by ships following Tier II. In areas where no AIS data was available (e.g. west of France), shipping emissions as given in the EMEP inventory were used.

The LNG scenarios differ from the SCR scenarios in the following way: SO_2 emissions are zero while they are about 0.38 g/kWh for the SCR case. This a reduction by a factor of 10 compared to 150 the base case and corresponds to a sulphur content of 0.1% S, an average fuel consumption of 200 g/kWh and a ratio of 95% of the total sulphur emitted as SO_2. Also PM emissions are set to zero in the LNG case. For SCR use, sulphate emissions are again reduced by a factor of 10 according to the sulphur reduction from 1.0% S to 0.1% S in the fuel. Other PM emissions are kept constant. NO_x emissions were the same for both, LNG and SCR scenarios. They differ in the implementation date 155 of Tier III rules, only.

The change of the annual average emissions of NO_x and SO_x in comparison to the reference emissions of 2011 are given for the scenarios in Fig. 2 and Fig. 3, respectively. Regional differences in the emission changes are a result of ship types and ship sizes which undergo different temporal developments in fleet renewal. This depends on the actual age of the ships given in the ship 160 characteristics data base.

5

3 Chemistry transport modeling

3.1 CMAQ

The CMAQ model (Byun and Ching, 1999; Byun and Schere, 2006) was used in its version 4.7.1 with the CB05 chemistry mechanism. Compared to its previous version, the model update includes
165   several new features (Foley et al., 2010), among them are gas phase chlorine chemistry, improved secondary organic aerosol (SOA) formation (Edney et al., 2007) and an updated representation of sea salt that considers reactions with nitric acid and the formation of coarse mode nitrate (Kelly et al., 2010) in the so-called AE5 aerosol mechanism. The model was run for an entire year with a spinup time of 2 weeks. Standard profiles for the most important atmospheric pollutants were used as initial
170   conditions. However, their effect on the simulated atmospheric concentrations of the substances in focus in this paper is negligible after the spinup.

The model was setup on a 72 x 72km^2 grid for entire Europe and subsequently on a nested 24 x 24km^2 grid for central Europe, see Fig. 4. The vertical model extent contains 30 layers up to 100 hPa in a sigma hybrid pressure coordinate system. 20 of these layers are below approx. 2km, the
175   lowest layer extends to ca. 36m above ground. The evaluation area was restricted to the Greater North Sea region and some neighbouring section of the NE Atlantic, covering approximately half of the central European domain (see the red box in Fig. 4).

3.2 COSMO-CLM

The meteorological fields that drive the chemistry transport model were simulated with the COSMO-
180   CLM mesoscale meteorological model (version 4.8) for the year 2008 (Geyer, 2014) using NCEP forcing data (Kalnay et al., 1996). This year was chosen because it does not contain very unusual meteorological conditions in Europe and can therefore be used to represent typical weather conditions in Europe. The same meteorological fields were used for the scenario runs, i. e. projected changes due to climate change were not considered in order to avoid a mixture of effects, from emissions and
185   meteorological data, in the resulting concentrations of air pollutants.

COSMO-CLM is the climate version of the regional scale meteorological community model COSMO (Rockel et al., 2008), originally developed by Deutscher Wetterdienst (DWD) (Steppeler et al., 2003; Schaettler et al., 2008). It has been run on a 0.22° x 0.22° grid using 40 vertical layers up to 20 hPa for entire Europe. COSMO-CLM uses the TERRA-ML land surface model (Schrodin and Heise,
190   2001), a TKE closure scheme for the planetary boundary layer (Doms and Schättler, 2002; Doms et al., 2011), cloud microphysics after Seifert and Beheng (2001, 2006), the Tiedtke scheme (Tiedtke, 1989) for cumulus clouds and a long wave radiation scheme following Ritter and Geleyn (1992). The meteorological fields were afterwards processed to match the CMAQ grid. As far as possible, CMAQ uses the information that is provided by the meteorological input fields to calculate transport,
195   transformation and loss of all gas phase and particulate species. The impact of the meteorological

fields on the output of the chemistry transport model was investigated in detail in the articles by Matthias et al. (2009) and Bieser et al. (2011a).

3.3   Boundary conditions

Chemical boundary conditions for the outer model domain were taken from monthly means of the TM5 global chemistry transport model system (Huijnen et al., 2010) and were provided by the Dutch Royal Meteorological Institute (KNMI). The model results have been interpolated in time and space to provide daily boundary conditions for the 72 x 72km^2 CMAQ grid for Europe. Boundary conditions for the nested 24 x 24km^2 grid were calculated on hourly basis from the outer coarse grid. They were kept the same for all scenarios in order to restrict the analysis of the effects of emission changes on shipping in North Western Europe.

3.4   Land based emissions

The model runs were performed with full emissions from all relevant sources in the model domain. Land based emissions in hourly temporal resolution were produced with SMOKE EU (Bieser et al., 2011a) for the year 2011. They are based on officially reported EMEP emissions which are distributed in time and space using appropriate surrogates like population density, street maps or land use. Point sources were considered as far as information from the European point source emission register was available. The vertical distribution of the emissions was calculated online with the SMOKE model, the results are given by Bieser et al. (2011b). The land based emissions were kept constant for all scenario model runs. Therefore the impact of reduced land based emissions, which can be expected for Europe in the year 2030, was not considered here. This was done to keep the the analysis clear and discuss only the effects of shipping instead of mixing it up with reductions of land based sources.

4   Impact of shipping on concentrations of pollutants

4.1   Situation today

The results for today's air pollution due to shipping serve as a reference case for this study. They are discussed in detail in the accompanying paper by Aulinger et al. (2015). In brief it can be said that ships contribute significant amounts to the concentrations of NO_2, particle bound nitrate (NO_3^-(p)) and particle bound sulphate (SO_4^{2-}(p)). In summer, ozone is enhanced, too. High contributions from shipping to the NO_2 and SO_2 concentrations are restricted to the open sea and the coastal areas in the southern North Sea and in Denmark (see the reference case in Fig. 5 and A.1). Nitrate and sulphate aerosol particles as well as ozone are secondary pollutants. They are transported far more inland but their relative contribution to concentrations at the coast is lower compared to NO_2 and SO_2.

There are large differences between summer and winter. Partly, they can be ascribed to seasonal

differences in the emissions with higher shipping emissions in summer. Most of the differences in
the concentrations are caused by atmospheric chemistry. As a photochemical pollutant, ozone is only
increased during the summer months. The situation is similar for sulphate and nitrate aerosol. Both
are formed via oxidation path ways that include the photochemically formed OH radical. Therefore,
the conversion rate of SO_2 into SO_4^{2-}(p) and of NO_2 into NO_3^-(p) in summer is higher than in
winter. This leads to higher contributions of shipping emissions to the concentrations of these aerosol
components in summer. On the other hand, total nitrate aerosol concentrations are much lower in
summer compared to winter, because the gas-to-particle partitioning between HNO_3 and NO_3^-(p) is
temperature dependent with higher particulate nitrate concentrations at low temperatures.

4.2 Scenarios for the North Sea in 2030

To derive the contribution of ships to the selected pollutant concentrations two model runs, one
including and one excluding shipping emissions, were performed. The difference is regarded as the
contribution of ships to the individual pollutant. For the scenarios the difference between two model
runs with different shipping emissions is regarded as the change in the contribution of ships in the
respective scenario. The evaluation is restricted to concentrations in the lowest model level, because
they are most relevant for the population.

We mainly discuss the consequences of changes in the NO_x emissions from ships because here we
see the main differences between the scenarios (Fig. 2). Additionally, the strict rules for SO_2 came
into force in the North Sea ECA on 1 January 2015, and there are only small differences between
the scenarios with respect to SO_2 emissions in the North Sea (Fig. 3). This will be further discussed
in section 4.2.4.

NO_x emissions from ships have an impact on the NO_2 concentrations, on nitrate aerosol and on
ozone. It can be expected that NO_2 concentrations increase due to ship emissions. The impact of
NO_x emissions from ships on ozone will be different between winter and summer. While in summer
increased NO_x emissions will lead to increased ozone under most weather conditions and in most
regions, this will have almost no effect in winter.

In the following, maps illustrating changes in the contribution of shipping on NO_2, nitrate aerosol
and ozone concentrations in 2030 will be shown for the scenarios No ECA, ECA SCR 16 (Tier III
in 2016), ECA SCR 21 (Tier III in 2021) and ECA opt. The color coded relative changes refer
to the impact of the shipping emissions on the concentrations of selected pollutants. The latter is
given in reference maps which are results of the reference model run considering shipping emissions
from 2011. The changes in the other scenarios will be illustrated in time series for different North
Sea coastal regions (see Figure 4), each of them comprising about 9200 km^2. These time series
nicely show the highly variable impact of shipping emissions, which depends to a large extent on
the weather conditions and the concentrations of pollutants from other sources.

4.2.1 Nitrogen dioxide

265 Scenario No ECA is reflecting a steady increase in shipping activity disregarding the implementation of stricter rules for NO_x emissions. This will lead to an increase in the contribution of shipping to the average NO_2 concentrations by more than 30 % in large areas of the North Sea region (Fig. 5b). The largest increase can be seen in the English Channel and the south western North Sea while Norway and Sweden exhibit lower increase rates of around 20%.

270 Figure 5c displays scenario ECA SCR 21 in which the Tier III rules for new ships come into force in 2021. This means that in 2030 a large part of the fleet will still follow the less strict Tier I and Tier II regulations. Very few ships older than 30 years will not have to comply with any of the NO_x regulations. As a consequence, the contribution of ships to average NO_2 concentratios will be higher than in the case with Tier III regulations from 2016 onwards (Fig. 5d). In large parts of the North

275 Sea the contribution of shipping to NO_2 concentrations will be higher than today.

Scenario ECA SCR 16 on average still shows a moderate increase in the NO_2 concentrations caused by ships compared to the situation today (Fig. 5d). While in the English Channel and the southern North Sea the concentrations decrease by a few percent only, they decrease by more than 10% in the north western parts of the North Sea and in particular at the British, Norwegian and Swedish coast.

280 This is caused by the fact that the traffic to the main North Sea ports in Rotterdam, Hamburg and Antwerp will still increase and ships will become bigger, resulting in a rather small concentration decrease in the English Channel. Today, smaller and older ships travel to the smaller harbours in the North Sea area. However, many of them will be replaced after 2016, which means that a large fraction of those ships will comply with Tier III. This will lead to a reduction of the contribution of

285 shipping emissions to NO_2 concentrations in the central and northern part of the North Sea.

The contribution of shipping to NO_2 concentrations will be drastically reduced in the case of scenario ECA opt when all ships comply with Tier III rules for NO_x emissions (Fig. 5e). The simulations show a reduction of approximately 80% all over the North Sea region compared to today.

All reductions in the contribution of shipping to NO_2 concentrations have a similar magnitude and

290 regional distribution in winter and summer. On average the impact of shipping is slightly higher in winter compared to summer with a larger increase in scenario No ECA and ECA SCR 21 and a smaller decrease in scenario ECA SCR 16 and ECA opt. All maps for the winter case can be seen in the appendix (Fig. A.3).

Figure 6 shows a time series of the contribution of shipping to the daily average NO_2 concentrations

295 at the coasts of Belgium and the Netherlands (see Fig. 4 for the region). Of the scenarios for 2030, scenario ECA SCR 16 shows a slight decrease in the contribution of shipping to the NO_2 concentrations compared to today. If Tier III will be implemented in 2021 (scenario ECA SCR 21) or not at all until 2030 (scenario No ECA), the contribution of shipping to the NO_2 concentration will be higher than today. Large reductions of NO_2 from shipping, on some days more than 4 μg/m^3,

300 are only achieved when all ships and not only new-buildings follow the Tier III regulations. Time

series for the other regions are included in the appendix (Fig. A.6).

4.2.2 Nitrate Aerosol

Nitrate aerosol (NO_3^- (p)) is formed in the atmosphere as a consequence of the oxidation of NO_2. The amount of aerosol particles formed highly depends on the presence of other pollutants, in particular on the availability of ammonia (NH_3). Ammonia mainly stems from agricultural activities. The regions with the highest ammonia emissions are western France, the Benelux countries, western Germany and Denmark. Particulate ammonium nitrate preferentially exists in winter, at low temperatures. At higher temperatures ammonium nitrate particles decompose into gaseous ammonia and nitric acid. Therefore, nitrate aerosol concentrations all over Europe are much lower in summer compared to winter. On the other hand, oxidation of NO_2 is much more effective in summer leading to a higher contribution of shipping to nitrate aerosol compared to winter.

In summer, the emission scenarios show very similar results for nitrate aerosol and for NO_2 (see Fig. 7). In scenario No ECA the contribution of shipping to nitrate aerosol concentrations increases by more than 30 % over sea and by 25 % or more in large aeras of central Europe and in southern Scandinavia (Fig. 7b). In scenario ECA SCR 21 (Figure 7c) large areas of the North Sea, and in particular northern France, show an increase in nitrate aerosol from shipping while in other areas the situation will remain unchanged. Scenario ECA SCR 16 (Fig. 7d) shows a decrease in the contribution of shipping to nitrate aerosol concentrations by 7-10% in the north eastern part of the North Sea while in the south western part a small increase by 5 - 10% can be observed. Again, in scenario ECA opt the contribution of shipping to nitrate aerosol will be reduced by 60 - 80%.

In winter, nitrate aerosol concentrations are only marginally affected by shipping emissions. For this reason the results of the scenario runs do not show reliable patterns of changes in NO_3^- (p) concentrations caused by changing shipping emissions when given as relative changes. Therefore, they are not shown.

4.2.3 Ozone

NO_x emissions from ships have a strong influence on the atmospheric ozone concentrations. Ozone is formed out of NO_2 and atmospheric oxygen in the presence of sunlight. Volatile organic compounds (VOC) help to transform emitted NO into NO_2, thereby enhancing the ozone formation significantly. On the other hand NO destroys ozone, leading to low ozone concentrations during night-time when no photolysis of NO_2 takes place. This leads to a strong diurnal cycle of the ozone concentration and a large difference between winter and summer levels with much higher ozone concentrations in the summer. Furthermore, increased NO_x emissions may cause additional ozone formation in presence of sufficiently high VOC concentrations. If the VOC levels are comparably low, more NO_x causes ozone destruction.

Here, we look at the impact of shipping emissions on the daily mean ozone values. Figure 8 shows

10

maps of the distribution of changes in the contribution of ships to mean ozone concentration for the different scenarios. Fig. 8a shows that shipping causes about 7 μg/m^3 additional ozone (summer average value) in large parts of the North Sea and in Denmark. On the other hand, there is only a small increase in ozone in the English Channel, where NO_x concentrations are high. The effect of

340   ozone destruction by additional NO_x emissions under low VOC conditions can be clearly seen in scenario No ECA (Fig. 8b). Reductions in ozone concentrations caused by shipping emissions, by partly more than 80%, are clearly noticeable in the English channel, the south western North Sea and the surrounding coast line. On the other hand, the shipping contribution to increased ozone concentrations will be enhanced by more than 20% far from the main shipping areas in Central Europe,

345   Ireland and the Northern UK.

Scenario ECA SCR 21 (Fig. 8c) contains higher NO_x emissions, leading to decreased ozone in the English channel and higher values in Central Europe, France, Ireland and the UK. Scenario ECA SCR 16 (Fig. 8d) shows a lower shipping contribution to ozone concentrations in the North East of the North Sea region and an almost unchanged situation in the South West. In the case of ECA

350   opt, ozone caused by shipping emissions is significantly reduced by 40-60% all over the modeling domain, except for the English channel where a significant increase is found.

Fig. 9 displays a time series of the daily average ozone concentrations in northern Germany. It can be seen that the ozone values would be lower in summer, on some days by more than 10 μg/m^3, if ships emitted as little NO_x as in the ECA opt scenario. On the other hand they would be slightly

355   higher in winter.

An analysis of the different regions reveals that the days with concentrations higher than 120 μg/m^3 (a value recommended by the World Health Organization, WHO) would decrease significantly by 50% or more without shipping emissions in all regions except the Netherlands (see Table 1). The scenarios for 2030 do not show big differences in the number of days with concentrations above 120

360   μg/m^3. While small increases in the number of days can be expected if Tier III rules were not implemented (scenario No ECA), the only case with a strong decrease in exceedance days is scenario ECA opt.

4.2.4 Sulphur dioxide and sulphate aerosol

Sulphur dioxide (SO_2) emissions from ships are directly related to the sulphur content of ship fuels.

365   In the scenarios for 2030 all ships will follow the same rules for sulphur, which allow 0.1% S in the fuel in the North and Baltic Sea ECA and 0.5% outside of it. Therefore, the scenarios do not differ much in terms of sulphur emissions. The main difference is between the ECA SCR and ECA LNG scenarios, because LNG does not contain any sulphur at all which makes the sulphur emissions from these ships even lower than for all other ships that comply with the 0.1% S rule inside the North Sea

370   area.

In Fig. 10 changes in the contribution from shipping to the SO_2 and SO_4 concentrations for the

11

scenario ECA SCR 16 and ECA LNG 16 are shown for summer. In the ECA LNG 16 scenario sulphur dioxide and sulphate aerosol concentrations are even further reduced than in the ECA SCR 16 scenario. The reductions are between 40% and 60% for the SCR case and between 60% and 80%

375   for the LNG case. The reductions are slightly higher for SO_2 compared to sulphate, however the reductions for sulphate are more widespread than those for SO_2. The results for the other scenarios are very close to those in Fig. 10 which is why they are not shown here. More maps are included in the appendix (Fig. A.1, A.2, A.4, A.5).

4.2.5   PM2.5

380   Particulate matter with a diameter less than 2.5 μm ($PM_{2.5}$) originating from shipping emissions is mainly formed through a conversion of gaseous SO_2 and NO_2 into particulate nitrate (NO_3^-(p)) and sulphate (SO_4^{2-}(p)). The amount of these secondary aerosol components depends critically on the level of NH_3 emissions, which are a prerequisite for the formation of ammonium sulphate and annomium nitrate in the atmosphere. The area where the highest contribution of shipping to the

385   $PM_{2.5}$ concentrations is noticeable is south east of the main shipping lanes, in North West France, Belgium, The Netherlands and North Germany. These are areas with high ammonia emissions from agricultural activities.

Reductions in the contribution of shipping to $PM_{2.5}$ are visible in all scenarios (see Fig. 11). This is caused by the significant effect the sulphur reductions in the ship fuel has on sulphate aerosol

390   concentrations. In scenario ECA opt the $PM_{2.5}$ reduction is the largest. Here, also nitrate aerosol is significantly reduced.

The time series for PM2.5 concentrations and the respective reductions in the different scenarios can be seen for Northern Germany in Fig. 12. All scenarios except for No ECA show reductions in $PM_{2.5}$ on almost all days. Again, the largest reductions can be seen for ECA opt. There is no clear

395   seasonal trend for the reductions although total $PM_{2.5}$ is higher in winter compared to summer.

5   Conclusions

This paper investigates the effects of different future developments of shipping emissions in the North Sea area on air quality in the North Sea region. The main differences between the scenarios for 2030 concern nitrogen oxide emissions. They could be significantly lowered by using ex-

400   haust cleaning techniques or alternative fuels like LNG. Additionally, international regulations for a mandatory reduction of nitrogen oxide emissions in the North and Baltic Sea areas are under debate in the International Maritime Organization. To avoid misinterpretations of the results, land based emissions and meteorological conditions were the same in the scenario runs and in the base case.

It was found that the expected increase in ship traffic in the North Sea will lead to enhanced levels

405   of NO_2, nitrate aerosol and ozone in large areas of North West France, Belgium, The Netherlands,

North Germany and Denmark if no emission reduction measures will be taken. For secondary pollutants like ozone and aerosols, this effect is more pronounced in summer compared to winter. In winter shipping does not contribute much to elevated levels of aerosol concentrations.

The effect of emission reduction measures depends on the year of implementation. If already in 2016 new ships needed to follow the new Tier III rules for new-buildings, the concentrations of NO_2, nitrate aerosol and ozone in 2030 would be on the same level as today. This means that the emission reductions of the new ships are compensated by an increased ship traffic. If it took another five years until Tier III regulations were in place, the concentrations of NO_2, nitrate aerosol and ozone which are caused by ship traffic would be higher in 2030 compared to today. The difference to the implementation of the Tier III rules in 2016 would be about 10-15%. Significant reductions can only be achieved if all ships, not only new-buildings, followed strict NO_x emission limitations. The situation is different for sulphur dioxide, sulphate aerosol particles and also for $PM_{2.5}$. Regulations on lower sulphur levels in ship fuels have been implemented on 1 January 2015 for the North Sea and will be in place for all seas in 2020 (or the latest in 2025). This will significantly reduce the impact of shipping on SO_2 and sulphate aerosol concentrations. As a consequence of lower sulphate aerosol concentrations $PM_{2.5}$ concentrations will also be reduced. The use of LNG as alternative fuel would further reduce sulphur emissions and therefore also SO_2 and $PM_{2.5}$ concentrations.

Our model study shows that all effects of shipping emissions on air quality differ largely by region and season, depending on the pollutant in focus. Gaseous primary pollutants like NO_2 and SO_2 have a short life time. Consequently, their effects can mainly be seen close to the shipping lanes. Aerosols, which are formed through oxidation in the atmosphere can be transported over large distances. Contributions of shipping to nitrate, sulphate and PM2.5 concentrations can be seen far inland. For ozone, future emission reductions of NO_x could even lead to enhanced concentrations in regions that already today have high NO_x and low VOC concentrations like in the English Channel. However, this will depend on the future development of other NO_x emission sources, too. These were not taken into account here, as has been done for climate change neither, in order to focus on shipping effects and facilitate the interpretation of emission changes in this sector. Because it can be expected that in particular land based anthropogenic NO_x emissions in Europe will further decrease until 2030, the relative contribution of shipping emissions to NO_2 and nitrate pollution levels in the North Sea area will be higher than demonstrated here. Ozone might increase in regions where it was found to decrease in the scenarios No ECA and ECA SCR 16, namely in the English Channel. When this region, where ozone formation is currently VOC limited, turns into a NO_x limited region, NO_x emissions from shipping will enhance ozone concentrations in the entire study area.

13

ment of the emission scenarios.

445 References

Aulinger, A., Matthias, V., Zeretzke, M., Geyer, B., Bieser, J., and Quante, M.: The impact of shipping emissions on air pollution in the Greater North Sea region. Part I: Current emissions and concentrations, Atmos. Chem. Phys., submitted, 2015.

Bieser, J., Aulinger, A., Matthias, V., Quante, M., and Builtjes, P.: SMOKE for Europe - adaptation, modifi-
450   cation and evaluation of a comprehensive emission model for Europe, Geoscientific Model Development, 4, 47–68, doi:http://dx.doi.org/10.5194/gmd-4-47-201110.5194/gmd-4-47-2011, 2011a.

Bieser, J., Aulinger, A., Matthias, V., Quante, M., and Denier van der Gon, H. A. C.: Vertical emission profiles for Europe based on plume rise calculations, Environmental Pollution, 159, 2935–2946, doi:http://dx.doi.org/10.1016/j.envpol.2011.04.03010.1016/j.envpol.2011.04.030, 2011b.

455 Buhaug, Ø., Corbett, J., Endresen, Ø., Eyring, V., Faber, J., Hanayama, S., Lee, D., Lee, D., Lindstad, H., Markowska, A., Mjelde, A., Nelissen, D., Nilsen, J., Pålsson, C., Winebrake, J., Wu, W., and Yoshida, K.: Second IMO GHG Study, Tech. rep., International Maritime Organization (IMO), London, UK, 2009.

Byun, D. and Ching, J.: Science Algorithms of the EPA Models-3 Community Multiscale Air Quality Modeling System, Epa/600/r-99/030, US Environmental Protection Agency, Office of Research and Development,
460   Washington DC, 1999.

Byun, D. and Schere, K.: Review of the Governing Equations, Computational Algorithms, and Other Components of the Models-3 Community Multiscale Air Quality (CMAQ) Modeling System, Applied Mechanics Reviews, 59, 51–77, 2006.

Det Norske Veritas: DNV Report Shipping 2020, Tech. rep., DNV, Høvik, Norway, 2012.

465 Doms, G. and Schättler, U.: A Description of the Nonhydrostatic Regional Model LM. Part I: Dynamics and Numerics., Tech. rep., Deutscher Wetterdienst, www.cosmomodel.org/content/model/documentation/core/ default.htm, 2002.

Doms, G., J., F., Heise, E., Herzog, H.-J., Mrionow, D., Raschendorfer, M., Reinhart, T., Ritter, B., Schrodin, R., Schulz, J.-P., and Vogel, G.: A Description of the Nonhydrostatic Regional COSMO Model. Part II:
470   Physical Parameterization, Tech. rep., Deutscher Wetterdienst, http://www.cosmo-model.org/content/model/ documentation/core/cosmoPhysParamtr.pdf, accessed: 2014-04-09, 2011.

Edney, E. O., Kleindienst, T. E., Lewandowski, M., and Offenberg, J. H.: Updated SOA chemical mechanism for the Community Multi-Scale Air Quality model, Tech. Rep. EPA 600/X-07/025, U.S. EPA, Research Triangle Park, NC., 2007.

475 European Union: Council Directive 1999/32/EC of 26 April 1999 relating to a reduction in the sulphur content of certain liquid fuels and amending Directive 93/12/EEC, http://eur-lex.europa.eu/legal-content/EN/TXT/ ?uri=celex:31999L0032, last accessed 31 August 2015, 1999.

European Union: Directice 2012/33/EU of the European Parliament and of the Council of 21 November 2012 amending Council Directive 1999/32/EC as regards the sulphur content of marine fuels, http://eur-lex.europa.
480   eu/legal-content/EN/TXT/?uri=celex:32012L0033, last accessed 31 August 2015, 2012.

Eyring, V., Kohler, H. W., Lauer, A., and Lemper, B.: Emissions from international shipping: 2. Impact of future technologies on scenarios until 2050, Journal Of Geophysical Research-Atmospheres, 110, D17 306, 2005.

Foley, K. M., Roselle, S. J., Appel, K. W., Bhave, P. V., Pleim, J. E., Otte, T. L., Mathur, R., Sarwar, G.,

485   Young, J. O., Gilliam, R. C., Nolte, C. G., Kelly, J. T., Gilliland, a. B., and Bash, J. O.: Incremental testing of the Community Multiscale Air Quality (CMAQ) modeling system version 4.7, Geoscientific Model Development, 3, 205–226, doi:http://dx.doi.org/10.5194/gmd-3-205-201010.5194/gmd-3-205-2010, http://www.geosci-model-dev.net/3/205/2010/, 2010.

Geyer, B.: High-resolution atmospheric reconstruction for Europe 1948-2012: coastDat2, Earth System
490   Science Data, 6, 147–164, doi:http://dx.doi.org/10.5194/essd-6-147-201410.5194/essd-6-147-2014, http://www.earth-syst-sci-data.net/6/147/2014/, 2014.

Hammingh, P., Holland, M., Geilenkirchen, G., Jonson, J., and Maas, R.: Assessment of the environmental impacts and health benefits of a nitrogen emission control area in the North Sea, Tech. rep., PBL Netherlands Environmental Assessment Agency, The Hague, 2012.

495   Huijnen, V., Williams, J., van Weele, M., van Noije, T., Krol, M., Dentener, F., Segers, A., Houweling, S., Peters, W., de Laat, J., Boersma, F., Bergamaschi, P., van Velthoven, P., Le Sager, P., Eskes, H., Alkemade, F., Scheele, R., Nedelec, P., and Paetz, H. . W.: The global chemistry transport model TM5: description and evaluation of the tropospheric chemistry version 3.0, Geoscientific Model Development, 3, 445–473, doi:http://dx.doi.org/10.5194/gmd-3-445-201010.5194/gmd-3-445-2010, 2010.

500   International Maritime Organisation: Amendments to the Annex of the Protocol of 1997 to Amend the International Convention for the Prevention of Pollution from Ships, 1973, as Modified by the Protocol of 1978 Relating thereto (MARPOL Annex VI), http://www.imo.org/blast/blastDataHelper.asp?data_id=23760&filename=176%2858%29.pdf, last access 28 January 2015, 2008.

International Maritime Organisation, Marine Environment Protection Committee: http://www.imo.org/
505   MediaCentre/MeetingSummaries/MEPC/Pages/MEPC66.aspx, last access on 25 January 2015, 2014.

Jalkanen, J. . P., Johansson, L., Kukkonen, J., Brink, A., Kalli, J., and Stipa, T.: Extension of an assessment model of ship traffic exhaust emissions for particulate matter and carbon monoxide, Atmospheric Chemistry and Physics, 12, 2641–2659, doi:http://dx.doi.org/10.5194/acp-12-2641-201210.5194/acp-12-2641-2012, 2012.

510   Kalnay, E., Kanamitsu, M., Kistler, R., Collins, W., Deaven, D., Gandin, L., Iredell, M., Saha, S., White, G., Woollen, J., Zhu, Y., Chelliah, M., Ebisuzaki, W., Higgins, W., Janowiak, J., Mo, K. C., Ropelewski, C., Wang, J., Leetmaa, A., Reynolds, R., Jenne, R., and Joseph, D.: The NCEP/NCAR 40-year reanalysis project, Bulletin of the American Meteorological Society, 77, 437–471, 1996.

Kelly, J. T., Bhave, P. V., Nolte, C. G., Shankar, U., and Foley, K. M.: Simulating emission and chemical
515   evolution of coarse sea-salt particles in the Community Multiscale Air Quality (CMAQ) model, Geoscientific Model Development, 3, 257–273, doi:http://dx.doi.org/10.5194/gmd-3-257-201010.5194/gmd-3-257-2010, http://www.geosci-model-dev.net/3/257/2010/, 2010.

Matthias, V., Quante, M., and Aulinger, A.: Determination of the optimum MM5 configuration for long term CMAQ simulations of aerosol bound pollutants in Europe, Environmental Fluid Mechanics, 9, 91–108,
520   doi:http://dx.doi.org/10.1007/s10652-008-9103-610.1007/s10652-008-9103-6, 2009.

Matthias, V., Bewersdorff, I., Aulinger, A., and Quante, M.: The contribution of ship emissions to air pollution in the North Sea regions, Environmental Pollution, 158, 2241–2250, doi:http://dx.doi.org/10.1016/j.envpol.2010.02.01310.1016/j.envpol.2010.02.013, 2010.

Ritter, B. and Geleyn, J. F.: A Comprehensive Radiation Scheme For Numerical Weather Prediction

525   Models With Potential Applications In Climate Simulations, Monthly Weather Review, 120, 303–
       325, doi:http://dx.doi.org/10.1175/1520-0493(1992)120%3C0303:ACRSFN%3E2.0.CO;210.1175/1520-
       0493(1992)120<0303:ACRSFN>2.0.CO;2, 1992.

Rockel, B., Will, A., and Hense, A.: The Regional Climate Model COSMO-CLM(CCLM), Meteorologische
       Zeitschrift, 17, 347–348, 2008.

530   Schaettler, U., Doms, G., and Schraff, C.: A Description of the Nonhydrostatic Regional COSMO-Model Part
       VII: User's Guide, Tech. rep., Deutscher Wetterdienst, 2008.

Schrodin, R. and Heise, E.: The multi-layer-version of the DWD soil model TERRA/LM, Consortium for
       Small-Scale Modelling (COSMO) Tech. Rep, 2, 16, 2001.

Seifert, A. and Beheng, K. D.: A double-moment parameterization for simulating autoconversion, ac-
535   cretion and selfcollection, Atmospheric Research, 59, Int Union Geodesy & Geophys, ICCP, IAMAS,
       doi:http://dx.doi.org/10.1016/S0169-8095(01)00126-010.1016/S0169-8095(01)00126-0, 2001.

Seifert, A. and Beheng, K. D.: A two-moment cloud microphysics parameterization for mixed-phase clouds.
       Part 1: Model description, Meteorology And Atmospheric Physics, 92, 45–66, 2006.

Smith, T. W. P., Jalkanen, J. P., Anderson, B. A., Corbett, J. J., Faber, J., Hanayama, S., O'Keeffe, E., Parker,
540   S., Johansson, L., Aldous, L., Raucci, C., Traut, M., Ettinger, S., Nelissen, D., Lee, D. S., Ng, S., Agrawal,
       A., Winebrake, J. J., Hoen, M., Chesworth, S., and Pandey, A.: Third IMO GHG Study 2014, Tech. rep.,
       International Maritime Organization (IMO), London, UK, 2014.

Steppeler, J., Doms, G., Schattler, U., Bitzer, H. W., Gassmann, A., Damrath, U., and Gregoric, G.: Meso-
       gamma scale forecasts using the nonhydrostatic model LM, Meteorology and Atmospheric Physics, 82,
545   75–96, doi:http://dx.doi.org/10.1007/s00703-001-0592-910.1007/s00703-001-0592-9, 2003.

Tiedtke, M.: A Comprehensive Mass Flux Scheme For Cumulus Parameter-
       ization In Large-scale Models, Monthly Weather Review, 117, 1779–1800,
       doi:http://dx.doi.org/10.1175/1520-0493(1989)117%3C1779:ACMFSF%3E2.0.CO;210.1175/1520-
       0493(1989)117<1779:ACMFSF>2.0.CO;2, 1989.

17
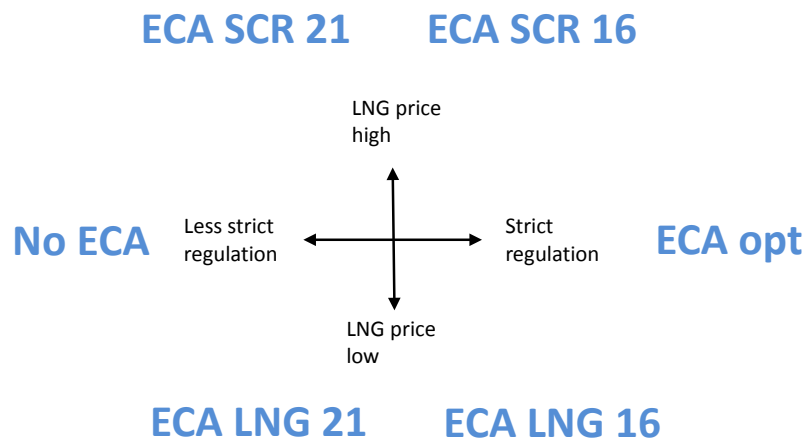
Fig. 1. Scenario identifiers for technical developments and legislation with respect to ship emissions for 2030.
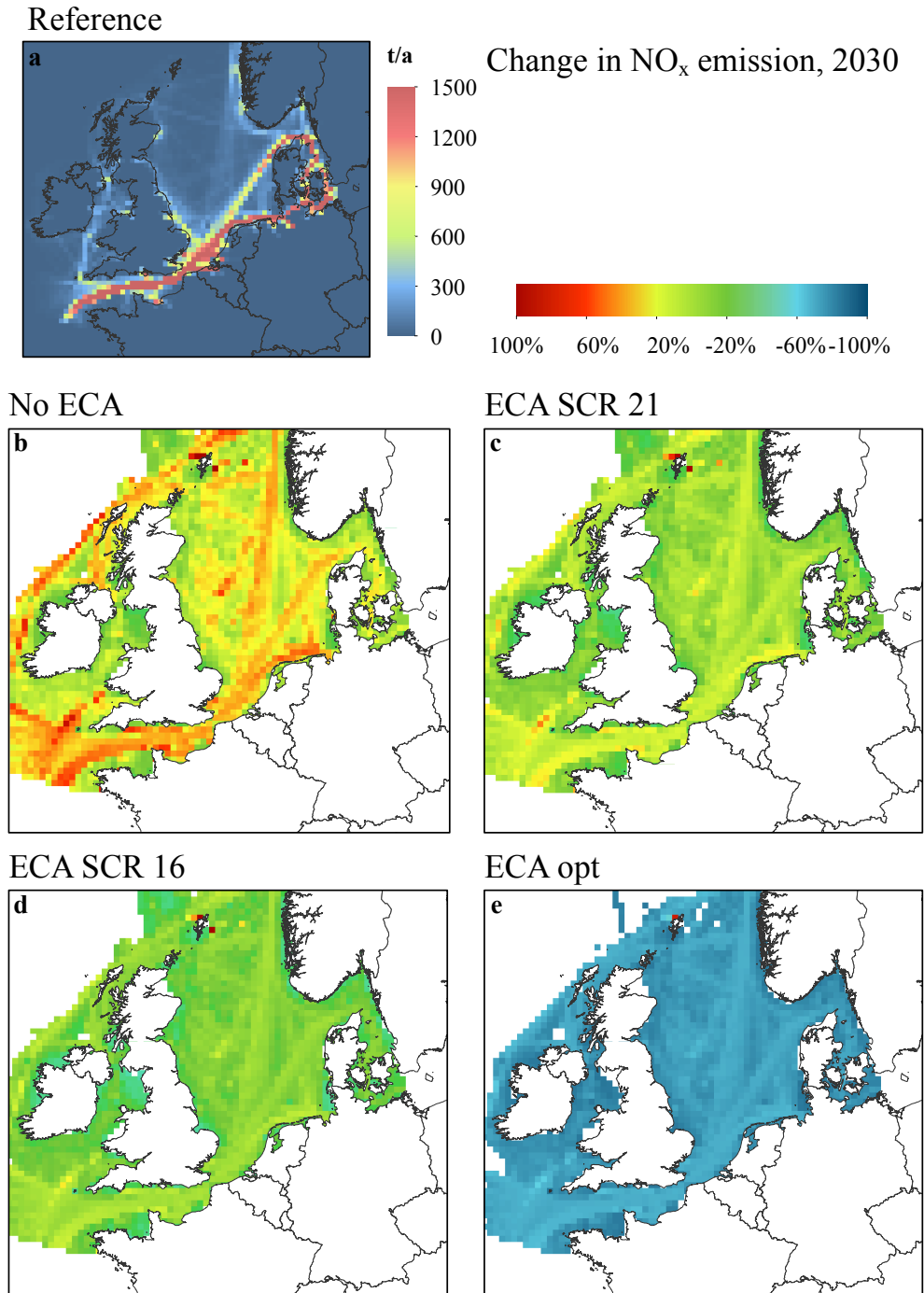
**Fig. 2.** NO$_x$ emissions from ships, a) annual totals in t per grid cell of 24 x 24 km^2. Emission changes for the scenarios b) No ECA, c) ECA LNG, d) ECA SCR, and e) ECA opt for 2030. No values are shown in grid boxes where the NO$_x$ emissions from ships were below 0.5 t/a per grid cell of 24 x 24 km^2.
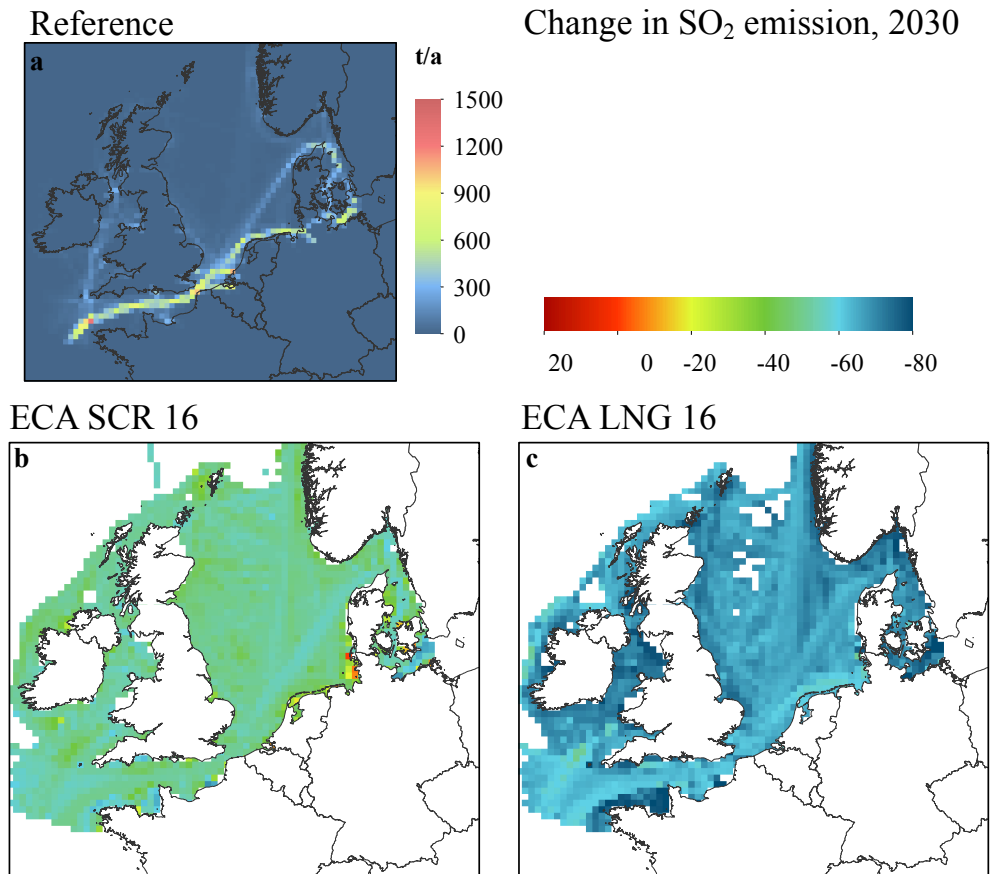
Fig. 3. SO$_2$ emissions from ships, a) annual totals in t per grid cell of 24 x 24 km^2. Emission changes for the scenarios b) ECA SCR 16, and c) ECA LNG 16 for 2030. No values are shown in grid boxes where the SO$_2$ emissions from ships were below 0.5 t/a per grid cell of 24 x 24 km^2.
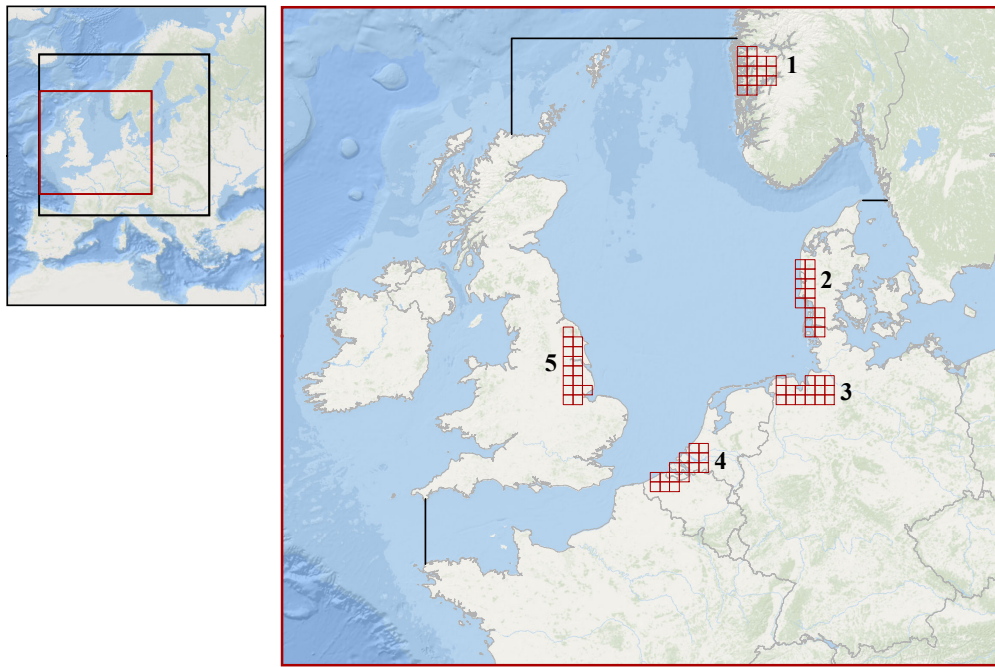
Fig. 4. Left: Modelling domains, outer domain with 72 x 72 km^2 resolution (outer black line), inner domain with 24 x 24 km^2 resolution (inner black line) and the evaluation area (red line). Right: Evaluation area including the greater North Sea region illustrating also the five regions (namely 1 to 5) for which time series of pollutant concentrations have been derived from CMAQ modelling results.
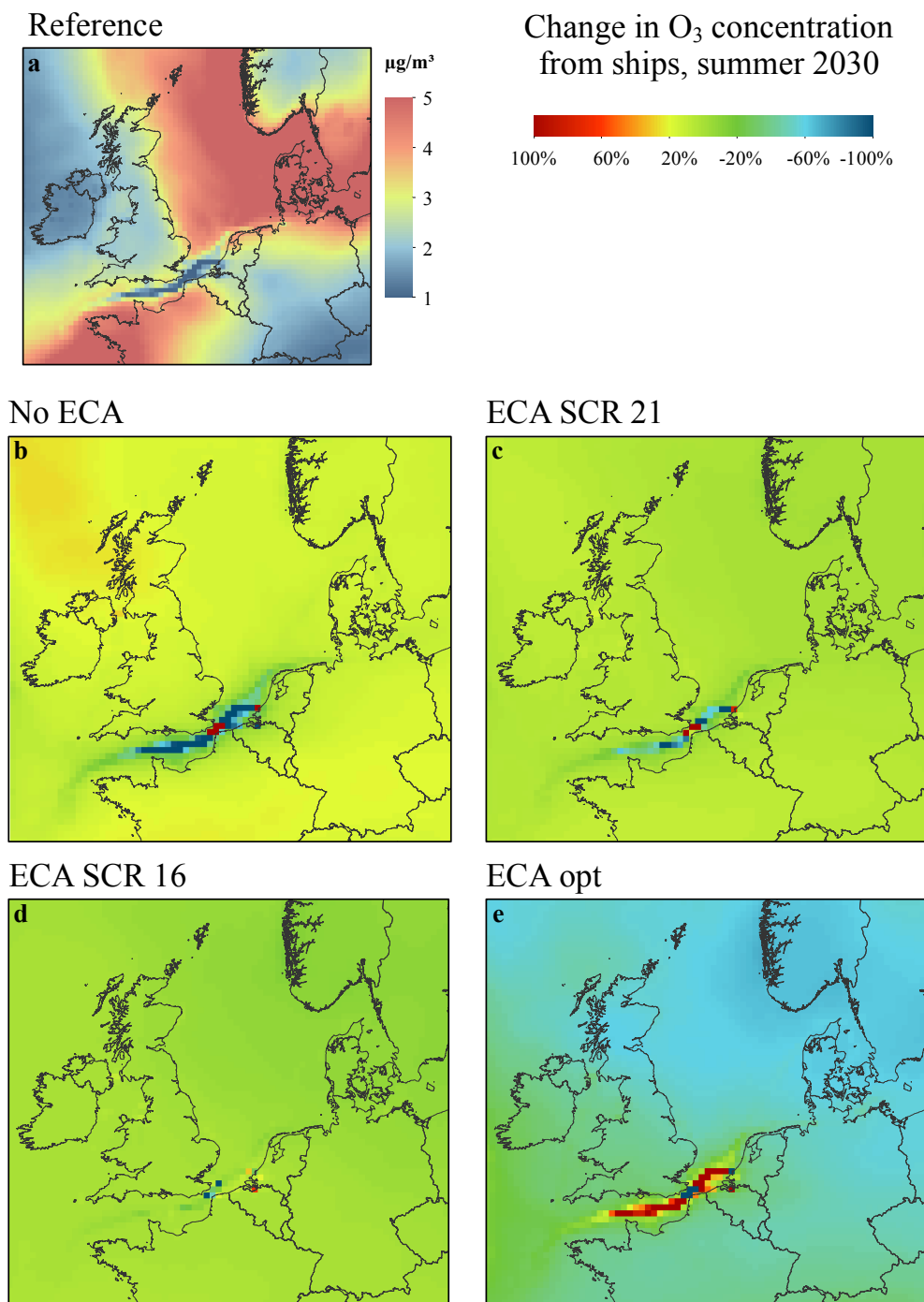
Reference

Change in NO$_2$ concentration from ships, summer 2030

No ECA

ECA SCR 21
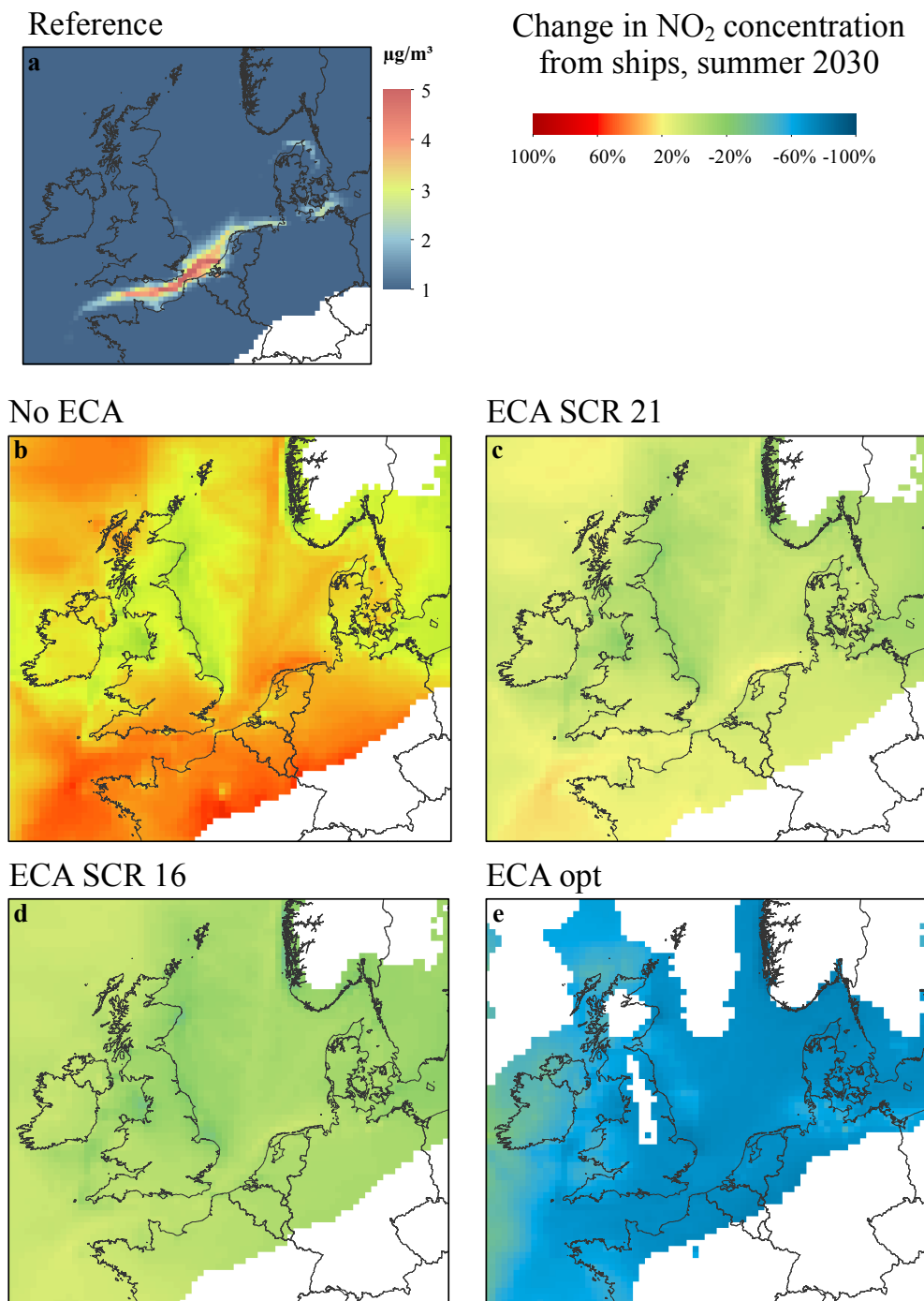
ECA SCR 16

ECA opt

Fig. 5. Contribution of shipping to the total NO$_2$ concentrations in summer (JJA) (a) today (Reference) and change in the scenarios (b) No ECA, (c) ECA SCR 21, (d) ECA SCR 16, (e) ECA opt. No values are shown in grid boxes where the contribution from shipping to the NO$_2$ concentrations in either the reference or the scenario case was below 0.05 μg/m^3.

22
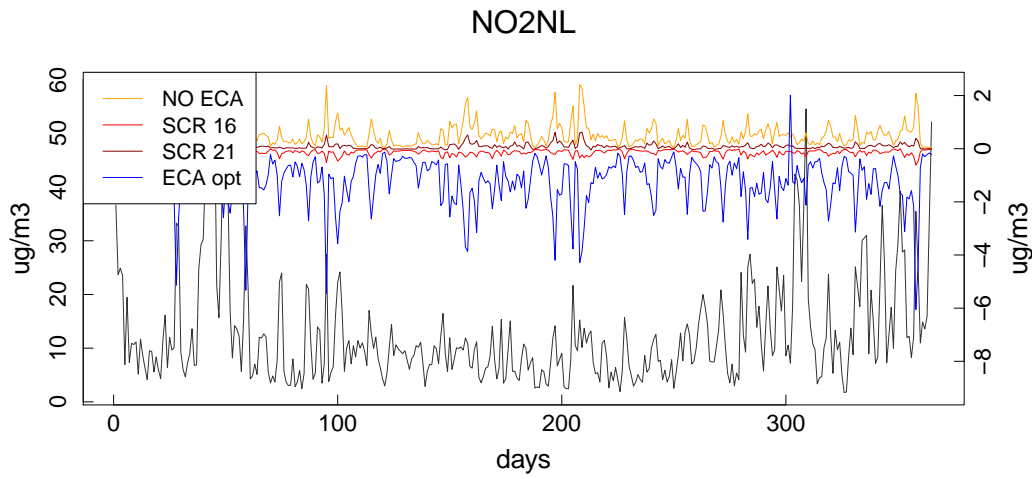
Fig. 6. Time series of daily average NO$_2$ concentrations in μg /m^3 (black, left y-axes) and change in the contribution of shipping to the NO$_2$ concentrations in the coastal areas of Belgium and the Netherlands (region 4) for all scenarios (colored, right y-axes).
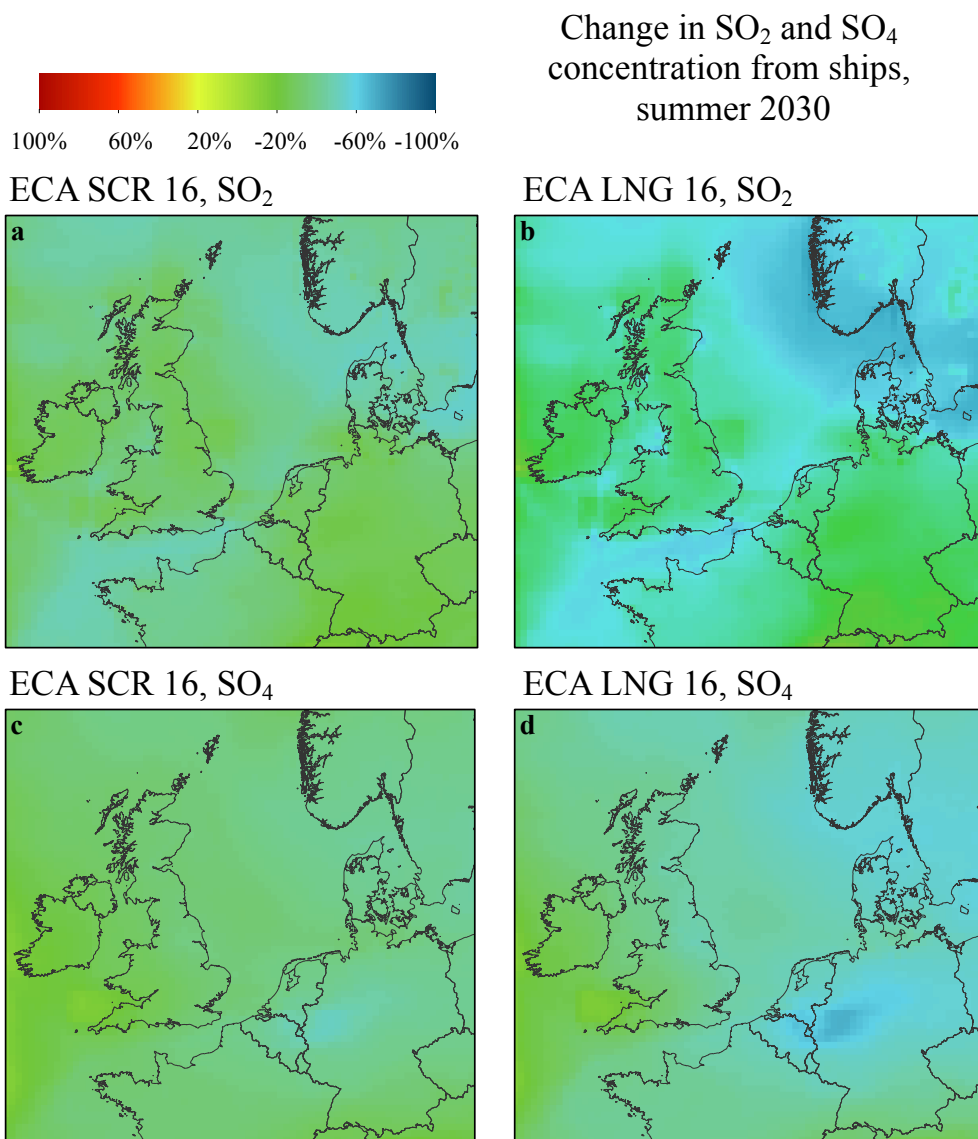
Reference

Change in NO$_3$ concentration from ships, summer 2030

µg/m³

No ECA

ECA SCR 21
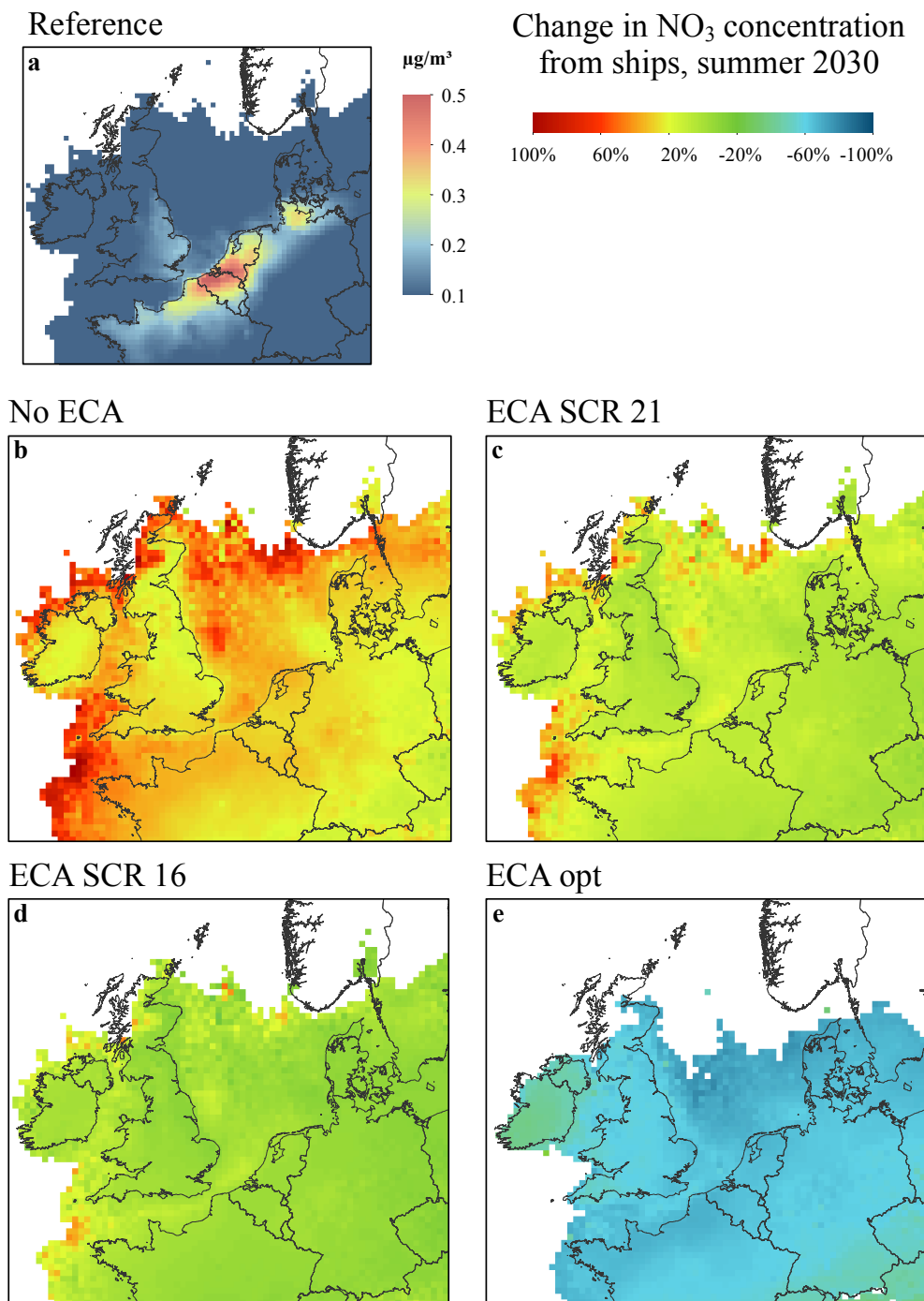
ECA SCR 16

ECA opt

Fig. 7. Contribution of shipping to the total NO$_3^-$ concentrations in summer (JJA) (a) today (Reference) and change in the scenarios (b) No ECA, (c) ECA SCR 21, (d) ECA SCR 16, (e) ECA opt. No values are shown in grid boxes where the contribution from shipping to the NO$_3$ concentrations in either the reference or the scenario case was below 0.005 µg/m³.

24

Reference

Change in O$_3$ concentration from ships, summer 2030

No ECA

ECA SCR 21

ECA SCR 16

ECA opt

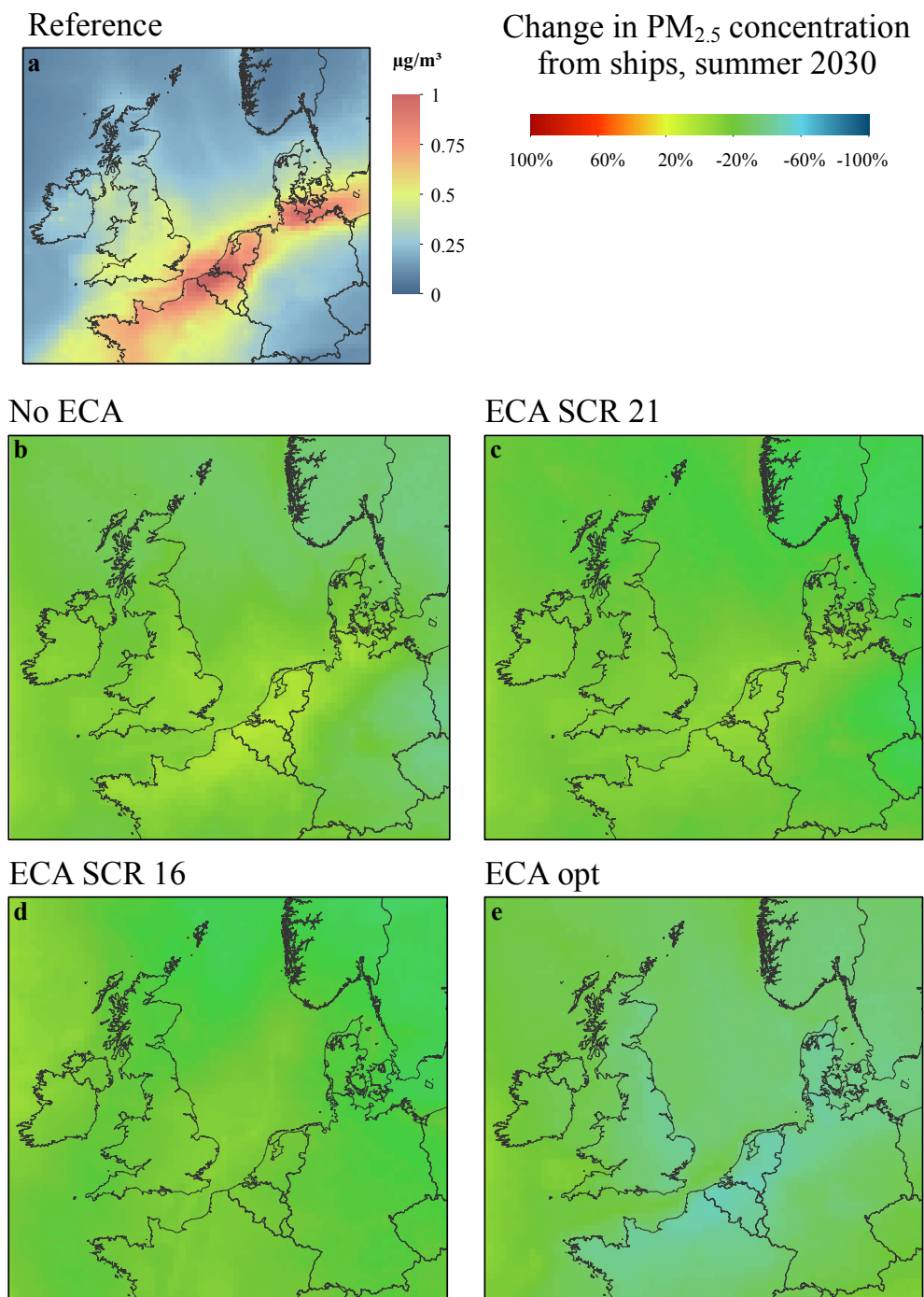**Fig. 8.** Contribution of shipping to the mean O$_3$ concentrations in summer (JJA) (a) today (Reference) and change in the scenarios (b) No ECA, (c) ECA SCR 21, (d) ECA SCR 16, (e) ECA opt.

O3DE
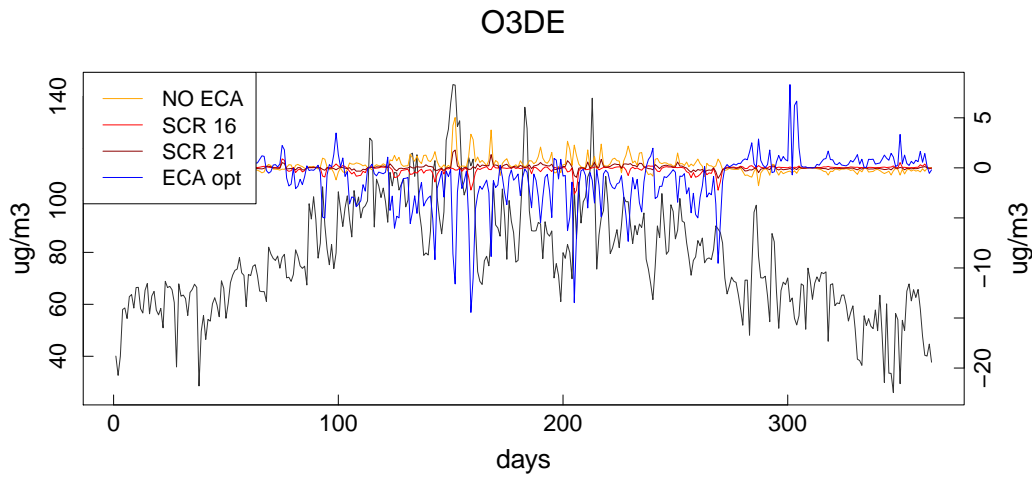
Fig. 9. Time series of daily average ozone concentrations in μg /m^3 (black, left y-axes) and change in the contribution of shipping to the ozone concentrations in the coastal areas of Germany (region 3) for all scenarios (right y-axes).
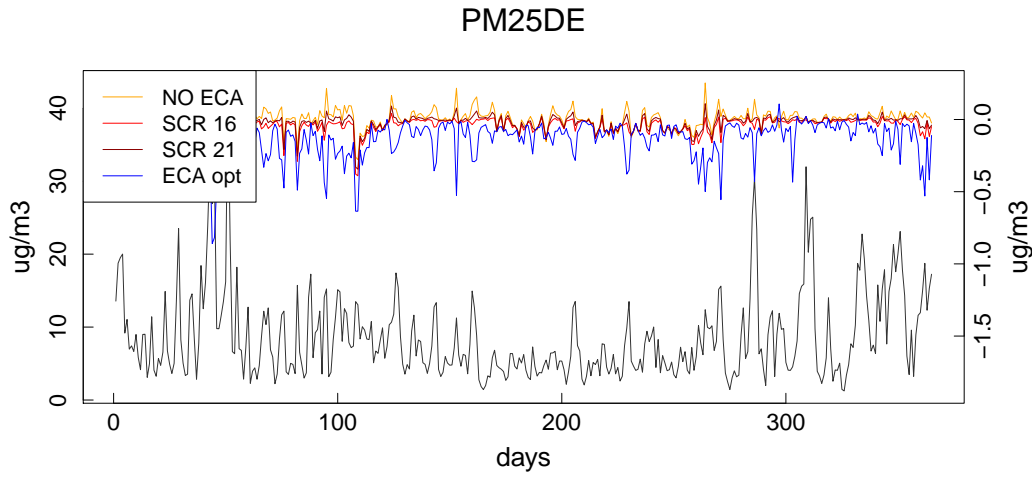
Fig. 10. Change in the contribution of shipping to the total (a and b) SO_2 and (c and d) SO_4 concentrations in summer (JJA) for the scenarios ECA SCR 16 (left), ECA LNG 16 (right), in relation to the reference case. See Figures A.1 and A.2 for the reference concentrations

Reference

Change in PM$_{2.5}$ concentration from ships, summer 2030

No ECA

ECA SCR 21

ECA SCR 16

ECA opt

Fig. 11. Contribution of shipping to the total PM$_{2.5}$ concentrations in summer (JJA) (a) today (Reference) and change in the scenarios (b) No ECA, (c) ECA SCR 21, (d) ECA SCR 16, (e) ECA opt.

28

Fig. 12. Time series of daily average PM2.5 concentrations in μg/m^3 (black, left y-axis) and change in the contribution of shipping to the PM2.5 concentrations in the coastal areas of Germany (region 3) for the main scenarios (colored, right y-axis).

Table 1. Number of days with 8-hour maximum ozone concentrations greater than 120 μg/m^3 in 2030 in selected regions around the North Sea for the scenarios No ECA, ECA SCR 16, ECA SCR 21, and ECA opt.

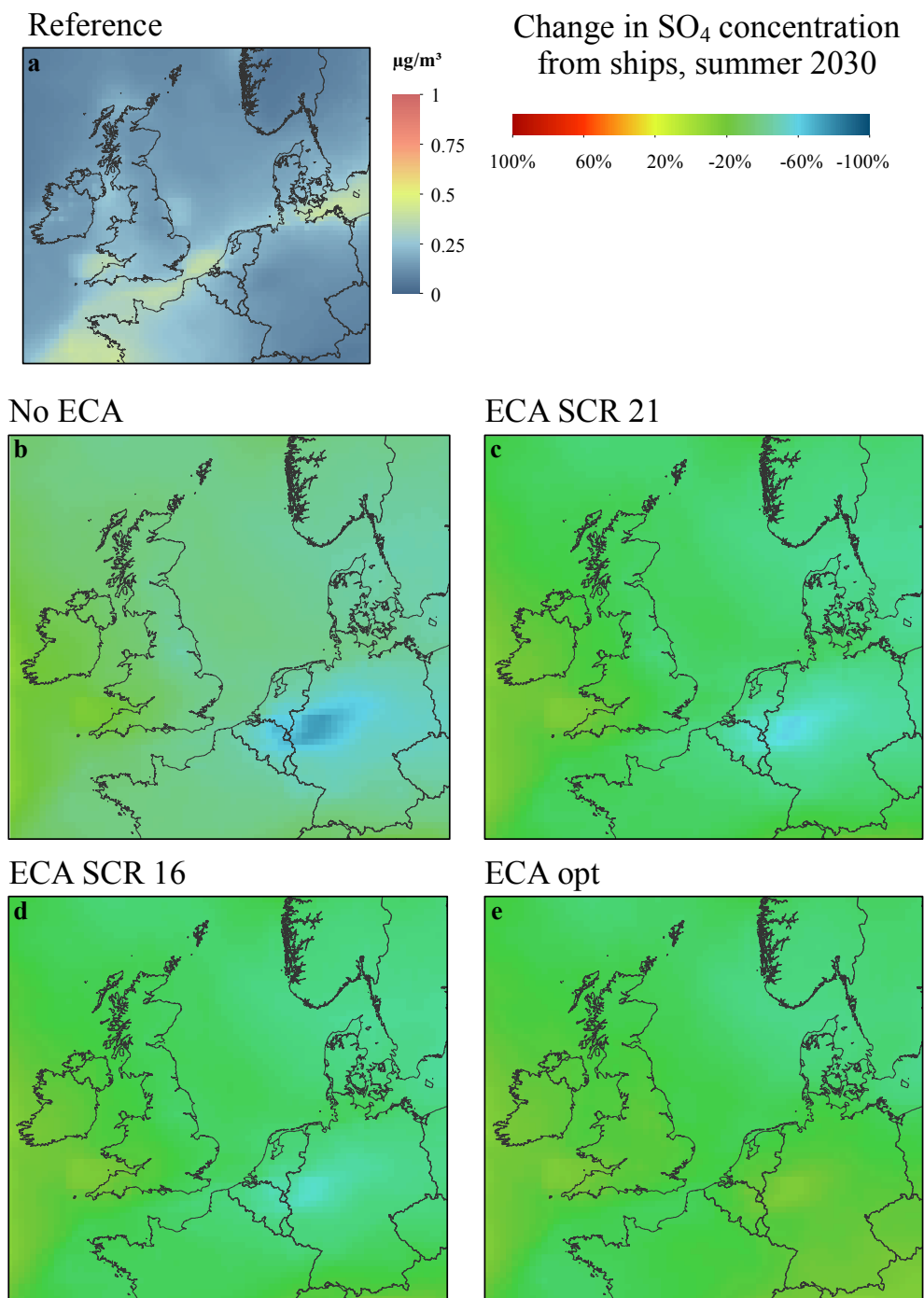|	1	2	3	4	5
2011	9	19	27	46	29
No ECA	9	20	29	46	29
ECA SCR 16	8	18	26	46	29
ECA SCR 21	8	18	27	46	29
ECA opt	6	12	16	45	22

550 **Appendix A**

Supplementary information on other seasons and other pollutants

A1   Sulphur containing species in summer
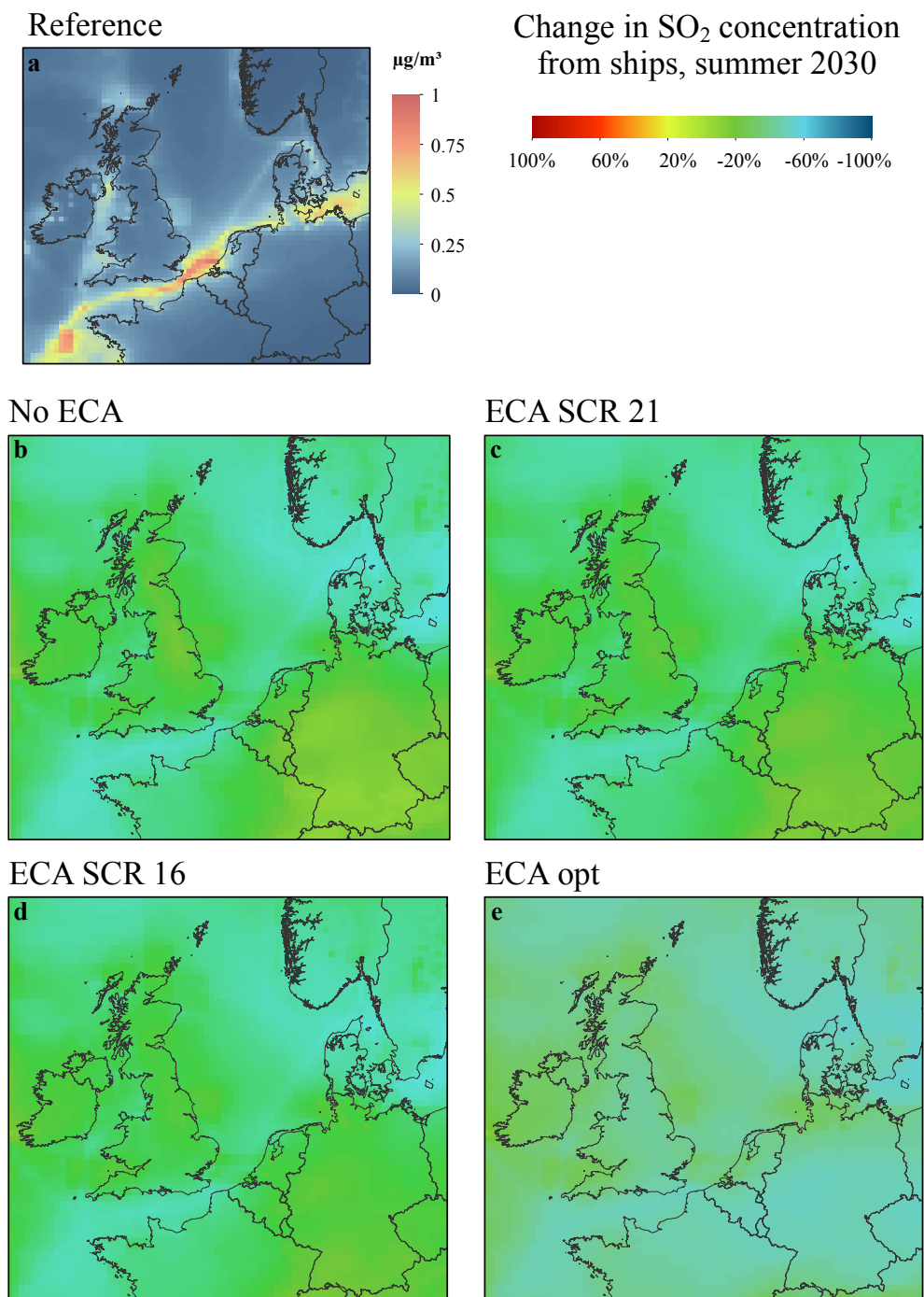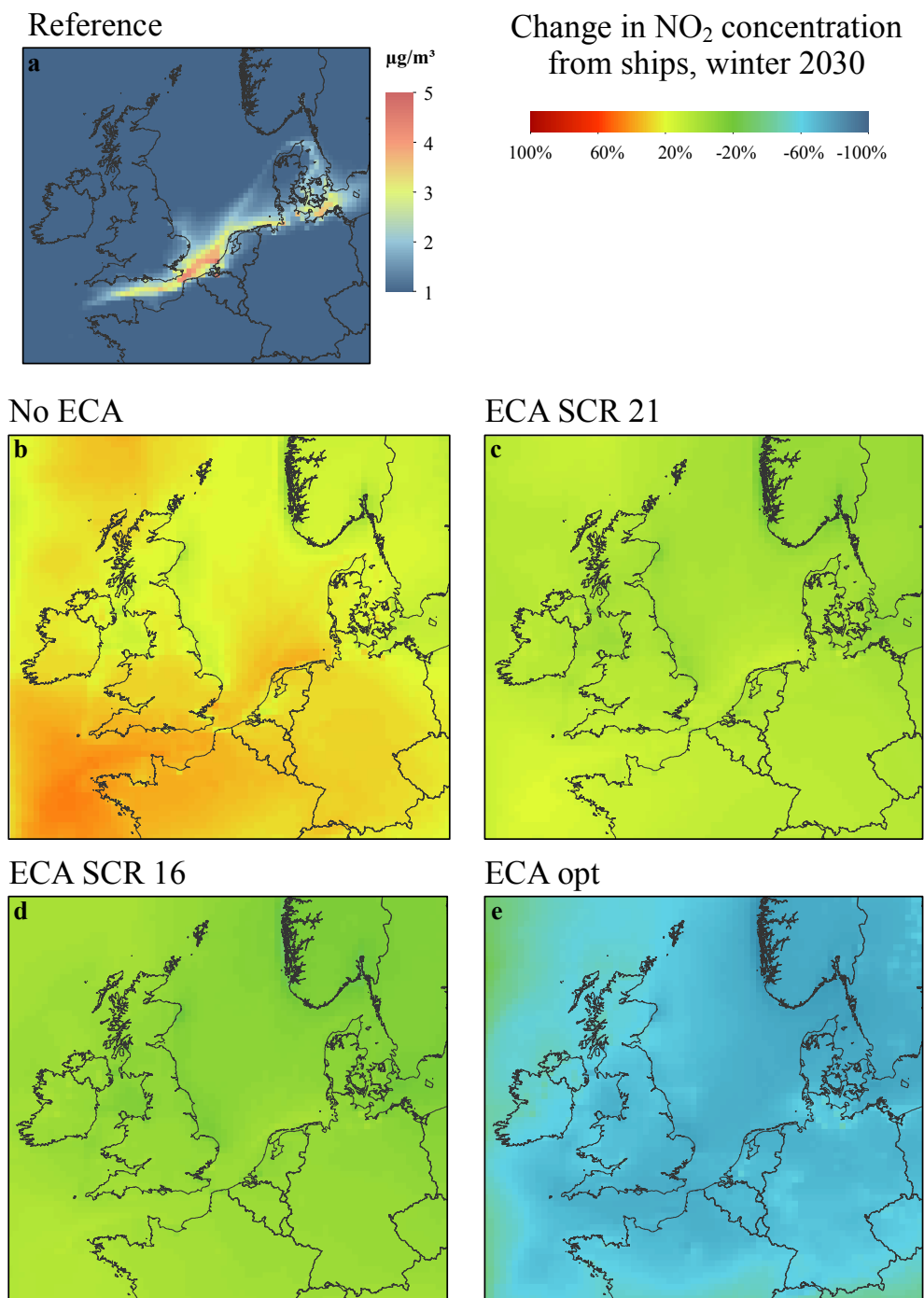
Fig. A.1. Change in the contribution of shipping to the total SO₂ concentrations in summer (JJA) compared to the reference case (a) for the scenarios (b) No ECA, (c) ECA SCR 16, (d) ECA SCR 21, and (e) ECA opt.
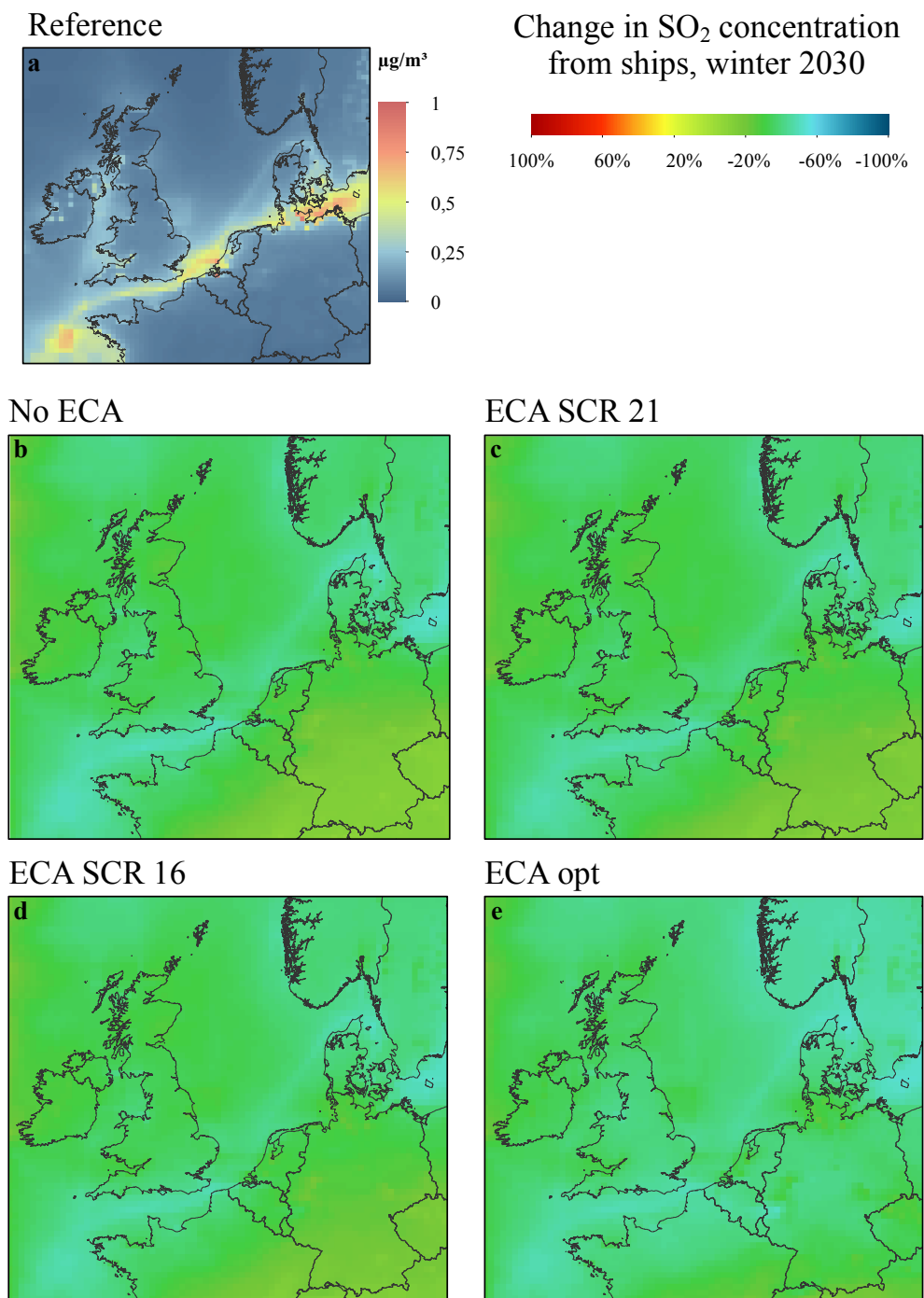
Reference

Change in SO$_4$ concentration from ships, summer 2030

μg/m³

1
0.75
0.5
0.25
0

100%   60%   20%   -20%   -60% -100%

No ECA

ECA SCR 21

ECA SCR 16

ECA opt

Fig. A.2. Change in the contribution of shipping to the total SO$_4$ concentrations in summer (JJA) compared to the reference case (a) for the scenarios (b) No ECA, (c) ECA SCR 16, (d) ECA SCR 21, and (e) ECA opt.
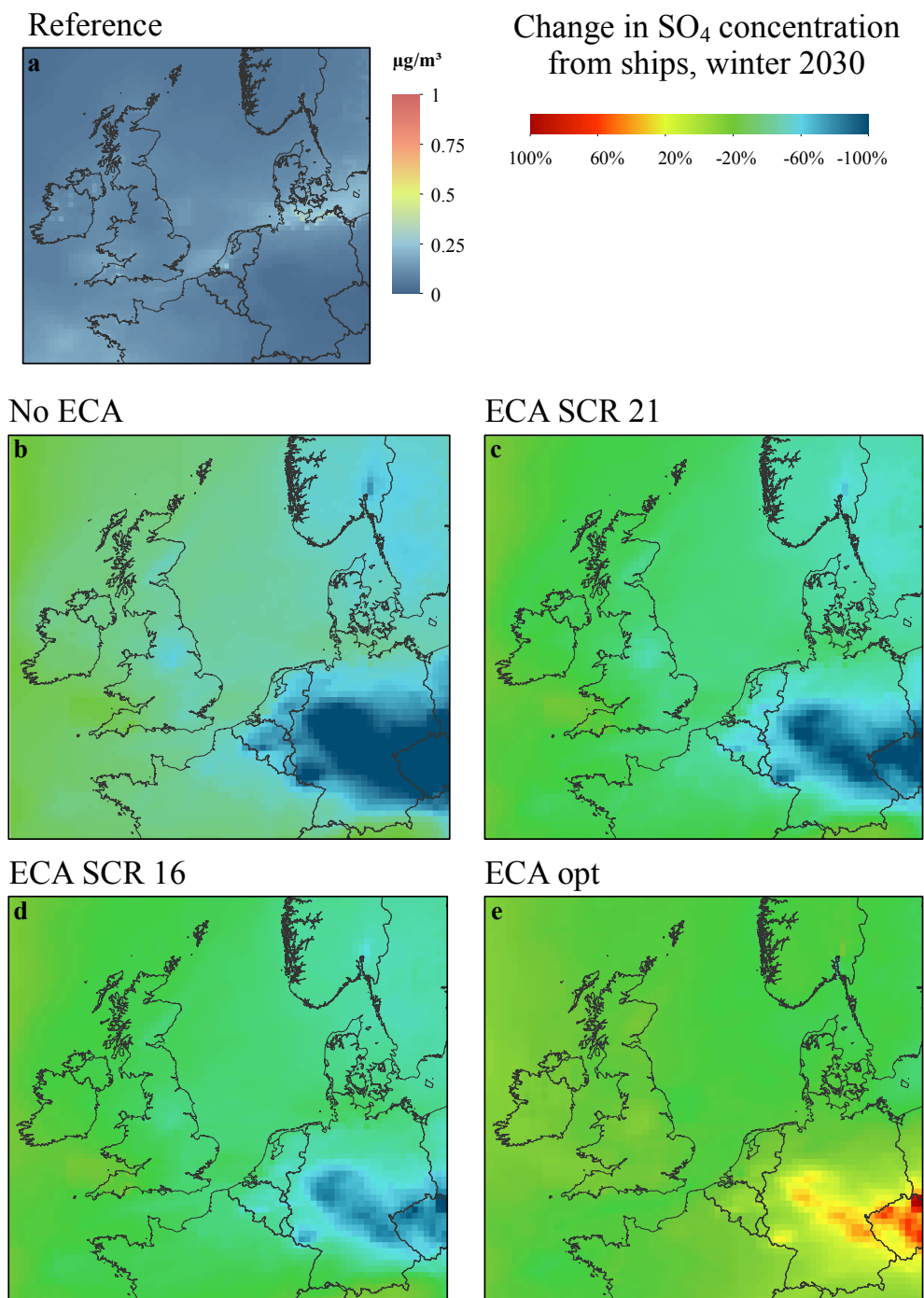
A2   Contribution of shipping to air pollution in winter

Reference

Change in NO$_2$ concentration from ships, winter 2030

100%   60%   20%   -20%   -60%   -100%

No ECA

ECA SCR 21

ECA SCR 16

ECA opt

Fig. A.3. Change in the contribution of shipping to the total NO$_2$ concentrations in winter (DJF) compared to the reference case (a) for the scenarios (b) No ECA, (c) ECA SCR 16, (d) ECA SCR 21, and (e) ECA opt.

Reference

Change in SO$_2$ concentration from ships, winter 2030

No ECA

ECA SCR 21

ECA SCR 16

ECA opt

Fig. A.4. Change in the contribution of shipping to the total SO$_2$ concentrations in winter (DJF) compared to the reference case (a) for the scenarios (b) No ECA, (c) ECA SCR 16, (d) ECA SCR 21, and (e) ECA opt.

Reference

Change in SO₄ concentration from ships, winter 2030

No ECA

ECA SCR 21

ECA SCR 16

ECA opt

Fig. A.5. Change in the contribution of shipping to the total SO₄ concentrations in winter (DJF) compared to the reference case (a) for the scenarios (b) No ECA, (c) ECA SCR 16, (d) ECA SCR 21, and (e) ECA opt.
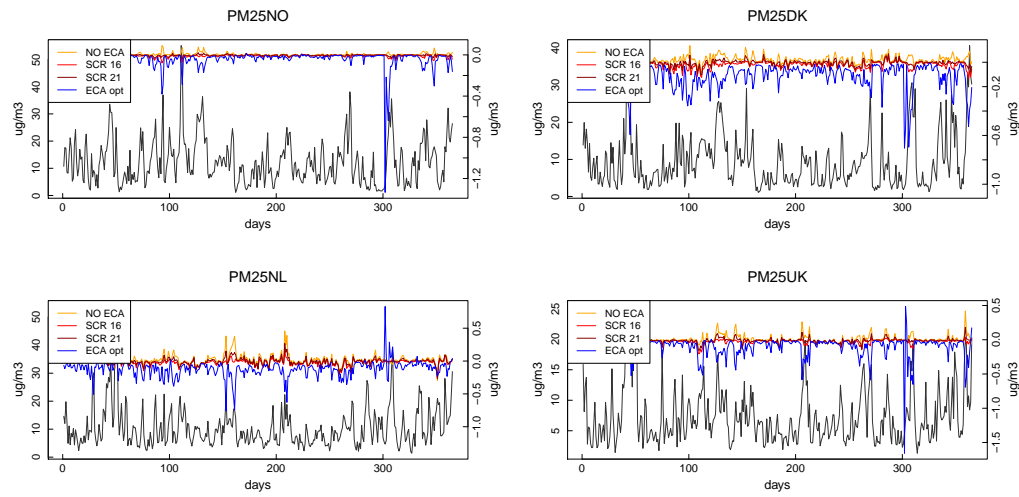
A3 Time series in selected areas
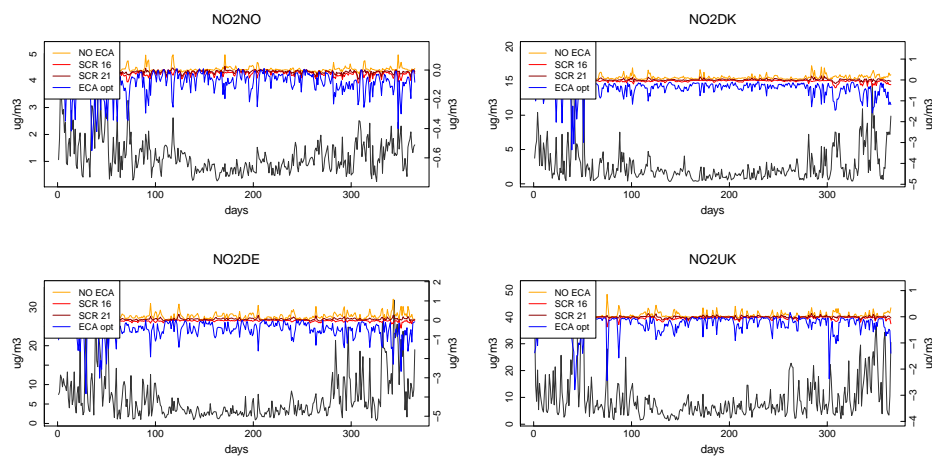
Fig. A.6. Time series of daily average NO_2 concentrations in $\mu g/m^3$ (black, left y-axes) and change in the contribution of shipping to the NO_2 concentrations in the coastal areas of Norway (region 1), Denmark (region 2), Germany (region 3), and Great Britain (region 5) for all scenarios (right y-axes).
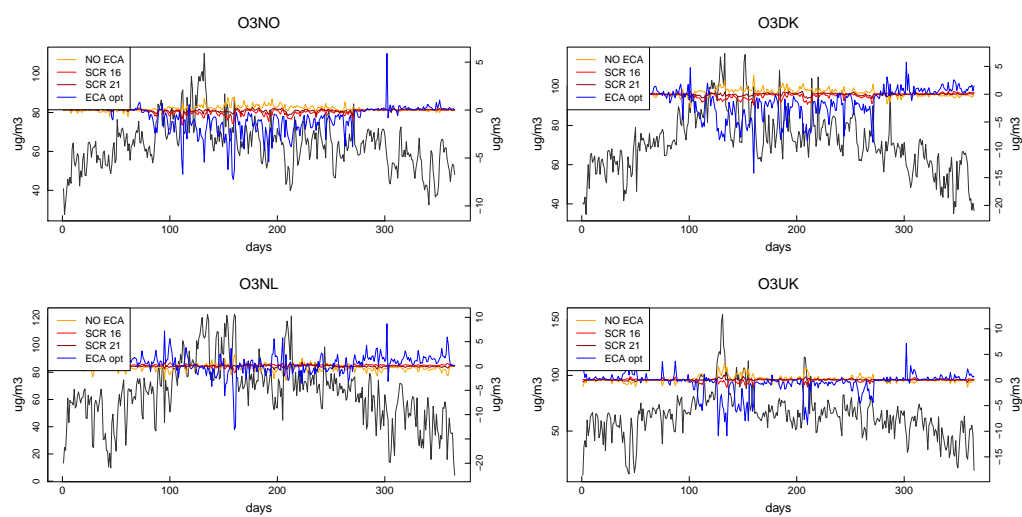
Fig. A.7. Time series of daily average O$_3$ concentrations in μg/m^3 (black, left y-axes) and change in the contribution of shipping to the O$_3$ concentrations in the coastal areas of Norway (region 1), Denmark (region 2), Belgium and The Netherlands (region 4), and Great Britain (region 5) for all scenarios (right y-axes).

Fig. A.8. Time series of daily average PM$_{2.5}$ concentrations in μg/m^3 (black, left y-axes) and change in the contribution of shipping to the PM$_{2.5}$ concentrations in the coastal areas of Norway (region 1), Denmark (region 2), Belgium and The Netherlands (region 4), and Great Britain (region 5) for all scenarios (right y-axes).

39