

METEOROLOGISK INSTITUTT  
Norwegian Meteorological Institute

# Transboundary air pollution by main pollutants (S, N, O<sub>3</sub>) and PM in 2017

## Armenia

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# 1 User guide

This report is one in a series of country-specific notes, complementary to the EMEP Status Report 1/2019. It presents an overview of transboundary pollution of main pollutants (S, N, O<sub>3</sub>) and particulate matter (PM) for Armenia in 2017.

All model runs have been performed with the EMEP MSC-W model version rv4.33, using ECMWF-IFS meteorology. The transboundary contributions presented here are based on source-receptor calculations with the EMEP MSC-W model using meteorological and emission data for the year 2017.

As a basis for their correct interpretation, this section briefly explains what types of results are shown in this report and how they have been calculated.

## 1.1 The chapters of this report

**Emissions** (*Chapter 2*): The emissions for 2017 have been derived from the 2019 official data submissions to UNECE CLRTAP as of May 2019. The gridded distributions of the 2017 emissions have been provided by the EMEP Centre on Emission Inventories and Projections (CEIP). The emissions for the period of 2000–2016, too, have been derived from the data submissions to UNECE CLRTAP as of May 2019.

The gridded emission data used in the model calculations this year are available on WebDab at:

[http://www.ceip.at/webdab\\_emepdatabase/emissions\\_emepmodels](http://www.ceip.at/webdab_emepdatabase/emissions_emepmodels).

**Time series** (*Chapter 3*): Time series in depositions and air concentrations are presented for the period of 2000–2017. The calculations are based on a consistent series of model runs, all using the EMEP MSC-W model version rv4.33. For the years 2000–2017, the meteorology of the respective year is used. Thus, interannual variability in the model results is due to changes in both emissions and meteorology. It should also be noted that the emission data and model version are updated regularly (see respective chapters on emissions and model updates in EMEP status report 1/2019), which may lead to differences between results reported here and in earlier reports.

**Transboundary fluxes** (*Chapter 4*): Data are presented in the form of maps and pie charts. The data are generated by source-receptor calculations, where emissions for each emitter of one or more precursors are reduced by 15%. The results have been scaled up to represent the entire emission from an emitter.

**Transboundary concentrations** (*Chapters 5 and 6*): Data are presented in the form of maps and bar charts. Ozone and particulate matter are subject to significant non-linearities in chemistry. Therefore we calculate the effect of 15% reductions in emissions only.

The horizontal maps show the reduction in concentrations when emissions are reduced by 15% in Armenia. By convention, reductions in concentrations are represented by positive values in the maps. Thus, any negative values mean that concentrations increase as a result of an emission reduction (due to non-linearities in chemistry).

The bar charts identify the six most important emitter countries in terms of their effects on concentrations in Armenia that would result from a 15% reduction in emissions. In the bar charts, the sum of the *absolute values* of these effects corresponds to 100%. The percentage values (vertical scale in the bar charts) thus give an indication of the relative importance of the various emitter countries that influence concentrations in Armenia (positive or negative, large or small contributions). Again, reductions are represented by positive values. Hence, a negative bar in the chart means that a *reduction* in emissions from an emitter

country would lead to an *increase* in concentration in Armenia. In some countries this can occur because of strong non-linearities in chemistry.

In addition, for PM<sub>2.5</sub> and PM<sub>10</sub> we show total concentration along with the contribution from natural sources (sea salt and natural dust) to the total concentration.

**Comparison with observations** (*Chapter 7*) : The map of monitoring stations shows stations of Armenia in the EMEP measurement network with measurements in 2017 submitted to EMEP. The frequency analysis plots compare daily observation results with the model results. The measurement data are available from CCC: <http://www.nilu.no/projects/ccc/emepdata.html>. The table provides annual statistics of the comparison of model results with observations for each measured component. Comparison is done only for stations with a sufficiently consistent set of data available in weekly or higher time resolution.

Also shown this year is the evaluation against measurements from the European Environment Agency's Air Quality e-Reporting data base (in the scientific community often referred to as 'AirBase'). In countries with AirBase sites, scatter plots show model performance in regard to chemical species, for which measurements are available.

**Risks from ozone and PM** (*Chapter 8*) : The maps of ozone and PM values correspond to regional background levels and they are not representative of local point measurements, where these values can be much higher (i.e. in cities).

## 1.2 Country codes

Many tables and graphs in this report make use of codes to denote countries and regions in the EMEP area. Table 1 provides an overview of these codes and lists the countries and regions included in the source-receptor calculations for 2017.

Code	Country/Region/Source	Code	Country/Region/Source
AL	Albania	IS	Iceland
AM	Armenia	IT	Italy
AST	Asian areas	KG	Kyrgyzstan
AT	Austria	KZ	Kazakhstan
ATL	N.-E. Atlantic Ocean	LI	Liechtenstein
AZ	Azerbaijan	LT	Lithuania
BA	Bosnia and Herzegovina	LU	Luxembourg
BAS	Baltic Sea	LV	Latvia
BE	Belgium	MC	Monaco
BG	Bulgaria	MD	Moldova
BIC	Boundary/Initial Conditions	ME	Montenegro
BLS	Black Sea	MED	Mediterranean Sea
BY	Belarus	MK	North Macedonia
CH	Switzerland	MT	Malta
CY	Cyprus	NL	Netherlands
CZ	Czechia	NO	Norway
DE	Germany	NOA	North Africa
DK	Denmark	NOS	North Sea
DMS	Dimethyl sulfate (marine)	PL	Poland
EE	Estonia	PT	Portugal
ES	Spain	RO	Romania
EU	European Union (EU28)	RS	Serbia
EXC	EMEP land areas	RU	Russian Federation
FI	Finland	SE	Sweden
FR	France	SI	Slovenia
GB	United Kingdom	SK	Slovakia
GE	Georgia	TJ	Tajikistan
GL	Greenland	TM	Turkmenistan
GR	Greece	TR	Turkey
HR	Croatia	UA	Ukraine
HU	Hungary	UZ	Uzbekistan
IE	Ireland	VOL	Volcanic emissions

Table 1: Country/region codes used throughout this report.

### 1.3 Definitions, statistics used

The following definitions and acronyms are used throughout this note:

SOA - secondary organic aerosol, defined as the aerosol mass arising from the oxidation products of gas-phase organic species.

SIA - secondary inorganic aerosols, defined as the sum of sulphate ( $\text{SO}_4^{2-}$ ), nitrate ( $\text{NO}_3^-$ ) and ammonium ( $\text{NH}_4^+$ ). In the EMEP MSC-W model SIA is calculated as the sum:  
$$\text{SIA} = \text{SO}_4^{2-} + \text{NO}_3^- (\text{fine}) + \text{NO}_3^- (\text{coarse}) + \text{NH}_4^+$$

SS - sea salt.

MinDust - mineral dust.

PPM - primary particulate matter, originating directly from anthropogenic emissions. One usually distinguishes between fine primary particulate matter,  $\text{PPM}_{2.5}$ , with aerosol diameters below  $2.5 \mu\text{m}$  and coarse primary particulate matter,  $\text{PPM}_{\text{coarse}}$  with aerosol diameters between  $2.5 \mu\text{m}$  and  $10 \mu\text{m}$ .

$\text{PM}_{2.5}$  - particulate matter with aerodynamic diameter up to  $2.5 \mu\text{m}$ . In the EMEP MSC-W model  $\text{PM}_{2.5}$  is calculated as  $\text{PM}_{2.5} = \text{SO}_4^{2-} + \text{NO}_3^- (\text{fine}) + \text{NH}_4^+ + \text{SS}(\text{fine}) + \text{MinDust}(\text{fine}) + \text{SOA}(\text{fine}) + \text{PPM}_{2.5} + 0.27 \text{NO}_3^- (\text{coarse}) + \text{PM}_{25\text{water}}$ . ( $\text{PM}_{25\text{water}} = \text{PM}$  associated water).

$\text{PM}_{\text{coarse}}$  - coarse particulate matter with aerodynamic diameter between  $2.5 \mu\text{m}$  and  $10 \mu\text{m}$ . In the EMEP MSC-W model  $\text{PM}_{\text{coarse}}$  is calculated as  $\text{PM}_{\text{coarse}} = 0.73 \text{NO}_3^- (\text{coarse}) + \text{SS}(\text{coarse}) + \text{MinDust}(\text{coarse}) + \text{PPM}_{\text{coarse}}$ .

$\text{PM}_{10}$  - particulate matter with aerodynamic diameter up to  $10 \mu\text{m}$ . In the EMEP MSC-W model  $\text{PM}_{10}$  is calculated as  $\text{PM}_{10} = \text{PM}_{2.5} + \text{PM}_{\text{coarse}}$ .

SOx - group of oxidized sulphur components ( $\text{SO}_2$ ,  $\text{SO}_4^{2-}$ ).

NOx - group of oxidized nitrogen components ( $\text{NO}$ ,  $\text{NO}_2$ ,  $\text{NO}_3^-$ ,  $\text{N}_2\text{O}_5$ ,  $\text{HNO}_3$ , etc.).

redN - group of reduced nitrogen components ( $\text{NH}_3$  and  $\text{NH}_4^+$ ).

SOMO35 is the Sum of Ozone Means Over 35 ppb is an indicator for health impact assessment recommended by WHO. It is defined as the yearly sum of the daily maximum of 8-hour running average over 35 ppb. For each day the maximum of the running 8-hours average for  $\text{O}_3$  is selected and the values over 35 ppb are summed over the whole year.

If we let  $A_8^d$  denote the maximum 8-hourly average ozone on day  $d$ , during a year with  $N_y$  days ( $N_y = 365$  or  $366$ ), then SOMO35 can be defined as:

$$\text{SOMO35} = \sum_{d=1}^{d=N_y} \max(A_8^d - 35 \text{ ppb}, 0.0)$$

where the max function ensures that only  $A_8^d$  values exceeding 35 ppb are included. The corresponding unit is ppb·days (abbreviated also as ppb·d).

AOT40 is the accumulated amount of ozone over the threshold value of 40 ppb, i.e.:

$$\text{AOT40} = \int \max(\text{O}_3 - 40 \text{ ppb}, 0.0) dt$$

where the max function ensures that only ozone values exceeding 40 ppb are included. The integral is taken over time, namely the relevant growing season for the vegetation concerned, and for daytime only. The corresponding unit is ppb-hours (abbreviated to ppb·h).

Although the EMEP model generates a number of AOT-related outputs, in accordance with the recommendations of the UNECE Mapping Manual we will concentrate in this report on two definitions:

**AOT40<sub>f</sub><sup>uc</sup>** - AOT40 calculated for forests using estimates of O<sub>3</sub> at forest-top (*uc*: upper-canopy). This AOT40 is that defined for forests by the UNECE Mapping Manual, but using a default growing season of April-September.

**AOT40<sub>c</sub><sup>uc</sup>** - AOT40 calculated for agricultural crops using estimates of O<sub>3</sub> at the top of the crop. This AOT40 is close to that defined for agricultural crops by the UNECE Mapping Manual, but using a default growing season of May-July, and a default crop-height of 1 m.

POD<sub>Y</sub> - Phyto-toxic ozone dose, is the accumulated stomatal ozone flux over a threshold *Y*, i.e.:

$$\text{POD}_Y = \int \max(F_{st} - Y, 0) dt \quad (1)$$

where stomatal flux  $F_{st}$ , and threshold,  $Y$ , are in  $\text{nmol m}^{-2} \text{s}^{-1}$ , and the max function evaluates  $\max(A - B, 0)$  to  $A - B$  for  $A > B$ , or zero if  $A \leq B$ . This integral is evaluated over time, from the start of the growing season (SGS), to the end (EGS).

For the generic crop and forest species, the suffix “gen” can be applied, in this report e.g. POD<sub>Y,gen</sub> (or  $AF_{st}^{1.6_{gen}}$ ) is used for forests and POD<sub>3.0,gen-CR</sub> (or  $AF_{st}^{3_{gen}}$ ) is used for crops.

## 2 Emissions

### 2.1 Emissions used in the EMEP MSC-W model calculations

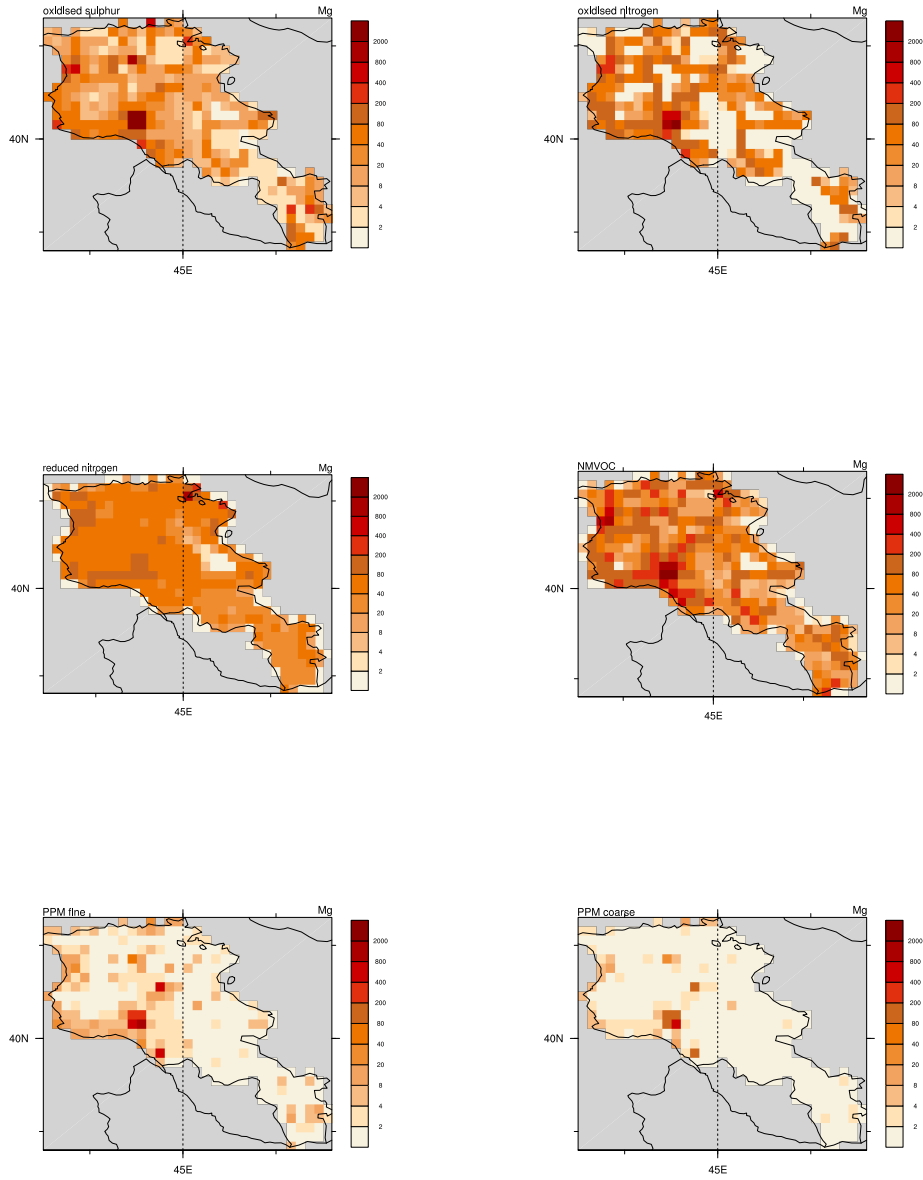


Figure 1: Spatial distribution of emissions from Armenia in 2017.



### 3 Time series

Important: For correct interpretation of the results shown in this chapter please read the paragraph on *Time series* in Section 1.1.

	2000	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
SO <sub>x</sub>	8	18	22	26	27	27	28	29	30	31	32	35	39	39
NO <sub>x</sub>	10	19	21	24	23	23	23	23	22	22	22	20	18	20
NH <sub>3</sub>	14	16	16	17	17	17	18	18	18	18	19	20	20	19
NMVOc	16	32	33	35	35	35	34	34	34	34	34	35	36	36
CO	110	116	114	112	111	110	109	108	107	106	105	106	108	112
PM <sub>2.5</sub>	4	4	4	4	4	4	4	4	4	4	4	4	4	4
PM <sub>10</sub>	5	5	5	5	5	5	5	5	5	5	6	6	6	6

Table 2: Emissions from Armenia. Unit: Gg. (SO<sub>x</sub> given as SO<sub>2</sub>, and NO<sub>x</sub> as NO<sub>2</sub>).

	2000	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
SO <sub>x</sub> dep.	15	16	16	17	18	18	18	17	17	14	19	20	24	21
NO <sub>x</sub> dep.	6	9	8	9	9	8	9	8	9	8	10	10	10	10
redN dep.	13	15	14	15	16	16	16	16	16	16	18	18	20	18

Table 3: Estimated deposition of Sulphur (S) and Nitrogen (N) in Armenia. Unit: Gg(S) or Gg(N).

	2000	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
mean ozone	40	41	42	41	41	41	41	41	40	42	41	42	42	42
max ozone	47	48	49	48	49	48	48	48	48	49	49	49	48	50
AOT40 <sub>f</sub> <sup>uc</sup>	24835	23755	27771	24138	27481	23559	27945	23511	26487	27217	27041	28512	24456	30597
SOMO35	3723	4014	4384	3977	4295	3952	4105	3955	3928	4338	4248	4432	4171	4689
POD <sub>1.0,gen-DF</sub>	20	25	24	25	25	25	24	25	27	24	23	23	27	24
PM <sub>2.5</sub> anthrop.	7	7	10	8	9	8	10	7	8	7	11	9	8	10
PM <sub>10</sub> anthrop.	11	11	17	13	14	13	17	11	11	11	17	14	12	16

Table 4: Estimated yearly mean values of air quality indicators averaged over Armenia. Unit: daily mean ozone (ppb), daily max ozone (ppb), AOT40<sub>f</sub><sup>uc</sup> (ppb-h), SOMO35 (ppb-d), POD<sub>1.0,gen-DF</sub> (mmol/m<sup>2</sup>), PM<sub>2.5</sub> (μg/m<sup>3</sup>) and PM<sub>10</sub> (μg/m<sup>3</sup>).

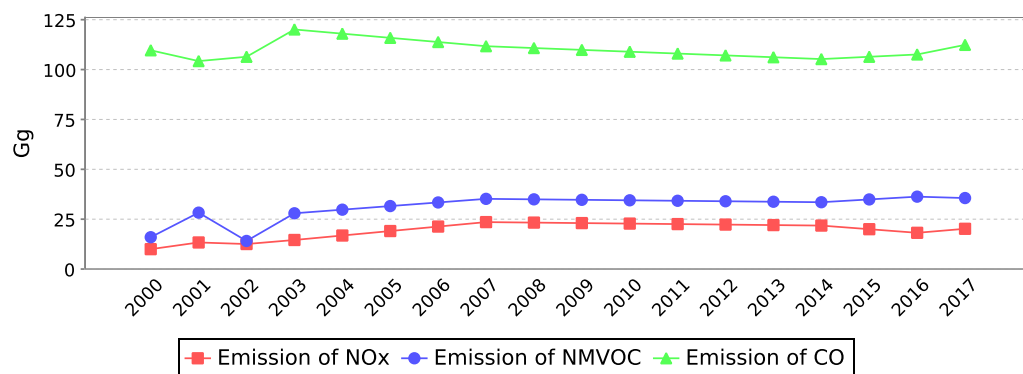


Figure 2: Trends in emissions of photo-oxidant pollution precursors. Unit: Gg (note that NO<sub>x</sub> is here given as NO<sub>2</sub>).

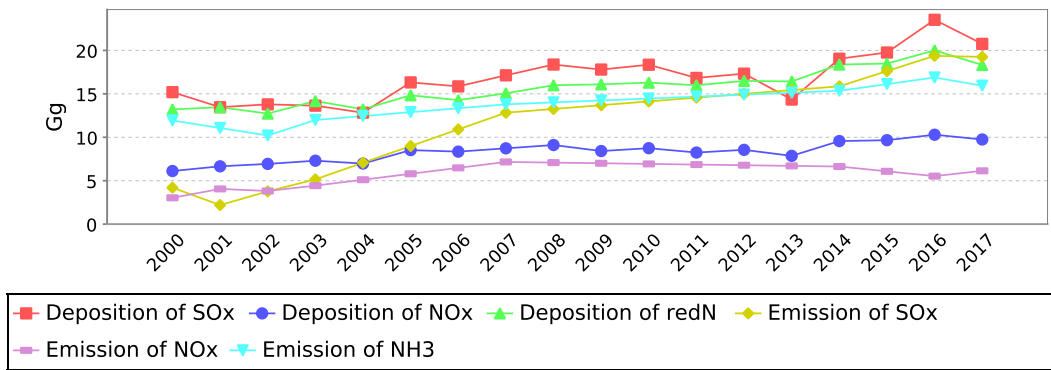


Figure 3: Trends in emissions and depositions of oxidised sulphur, oxidised nitrogen and reduced nitrogen. Unit: Gg(S) or Gg(N).

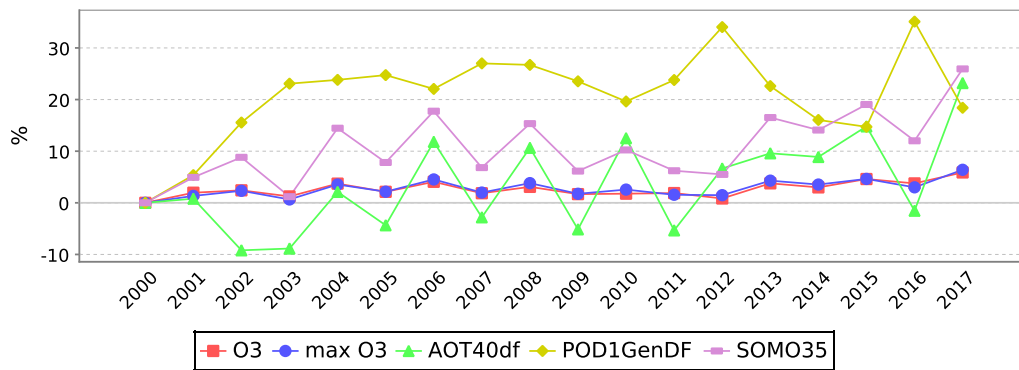


Figure 4: Changes in ozone related pollution relative to 2000. Unit: %. The large changes from year to year in some countries are mainly related to meteorological variability.

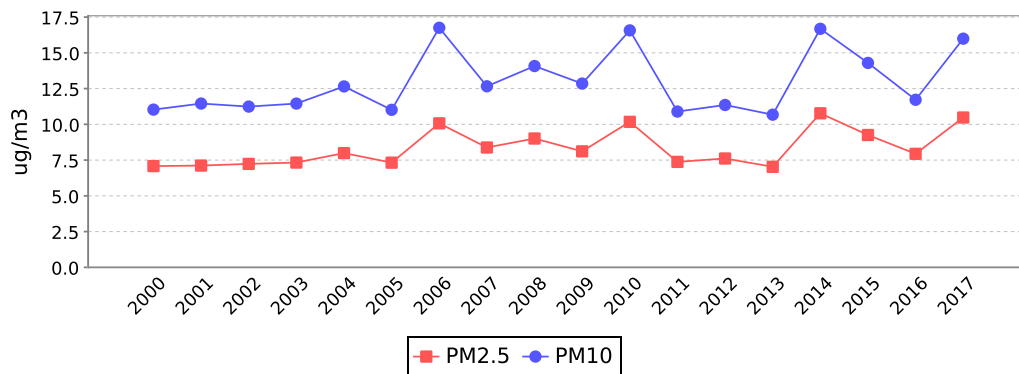


Figure 5: Trends in mean concentrations of particulate matter. Unit:  $\mu\text{g}/\text{m}^3$ .

## 4 Transboundary fluxes

### 4.1 Deposition of oxidised sulphur

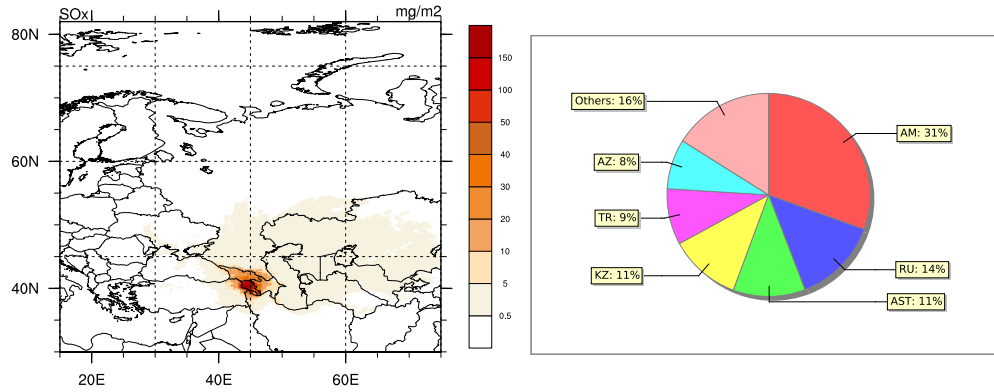


Figure 6: Contribution of emissions from Armenia to deposition of oxidised sulphur in the EMEP domain. Unit:  $\text{mg}(\text{S})/\text{m}^2$ . The pie chart shows the six main receptor areas where oxidised sulphur from Armenia is deposited. Unit: %.

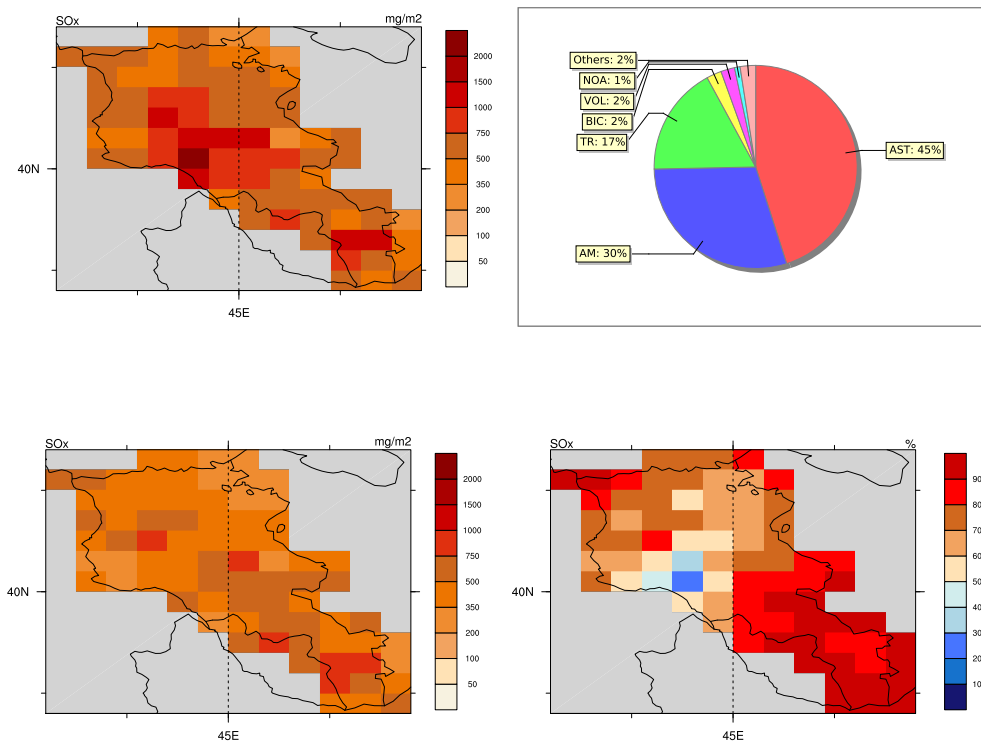


Figure 7: Top left: Deposition of oxidised sulphur in Armenia. Unit:  $\text{mg}(\text{S})/\text{m}^2$ . Top right: The six main contributors to oxidised sulphur deposition in Armenia. Unit: (%). Bottom left: Oxidised sulphur deposition from transboundary sources. Unit:  $\text{mg}(\text{S})/\text{m}^2$ . Bottom right: Fraction of transboundary contribution to total deposition. Unit: %.

## 4.2 Deposition of oxidised nitrogen

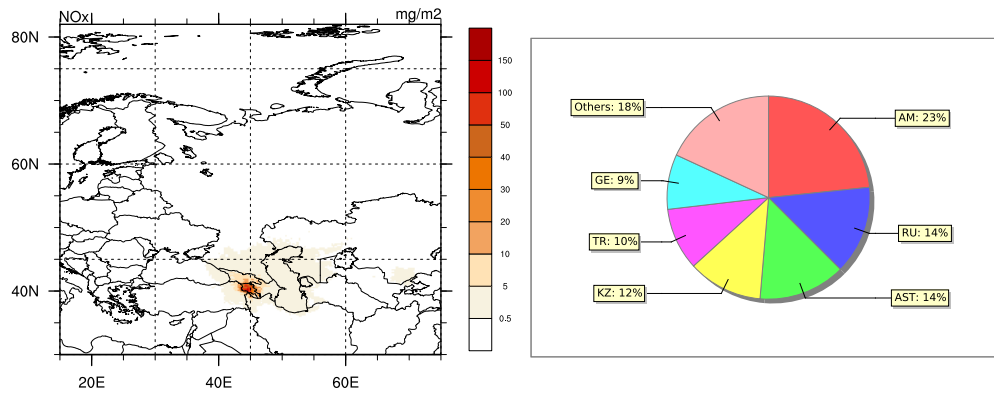


Figure 8: Contribution of emissions from Armenia to deposition of oxidised nitrogen in the EMEP domain. Unit:  $\text{mg(N)/m}^2$ . The pie chart shows the six main receptor areas where oxidised nitrogen from Armenia is deposited. Unit: %.

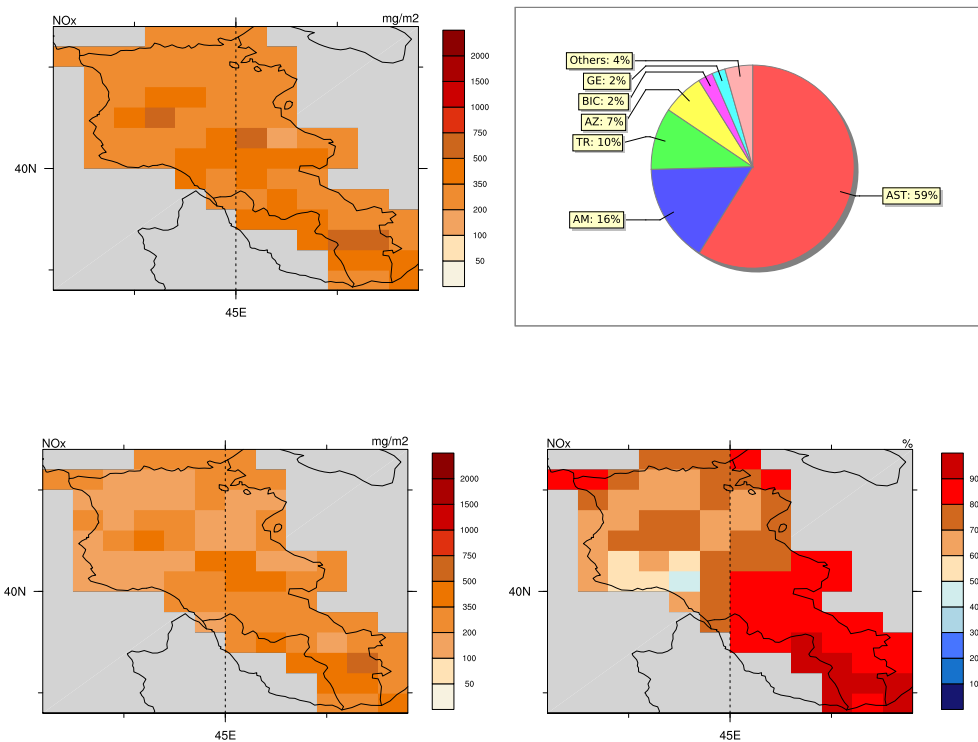


Figure 9: Top left: Deposition of oxidised nitrogen in Armenia. Unit:  $\text{mg(N)/m}^2$ . Top right: The six main contributors to oxidised nitrogen deposition in Armenia. Unit: %. Bottom left: Oxidised nitrogen deposition from transboundary sources. Unit:  $\text{mg(N)/m}^2$ . Bottom right: Fraction of transboundary contribution to total deposition. Unit: %.

### 4.3 Deposition of reduced nitrogen

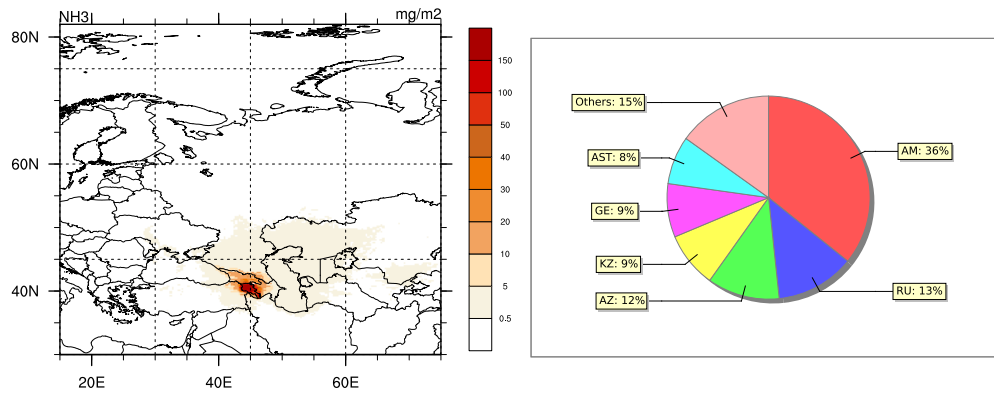


Figure 10: Contribution of emissions from Armenia to deposition of reduced nitrogen in the EMEP domain. Unit:  $\text{mg(N)/m}^2$ . The pie chart shows the six main receptor areas where reduced nitrogen from Armenia is deposited. Unit: %.

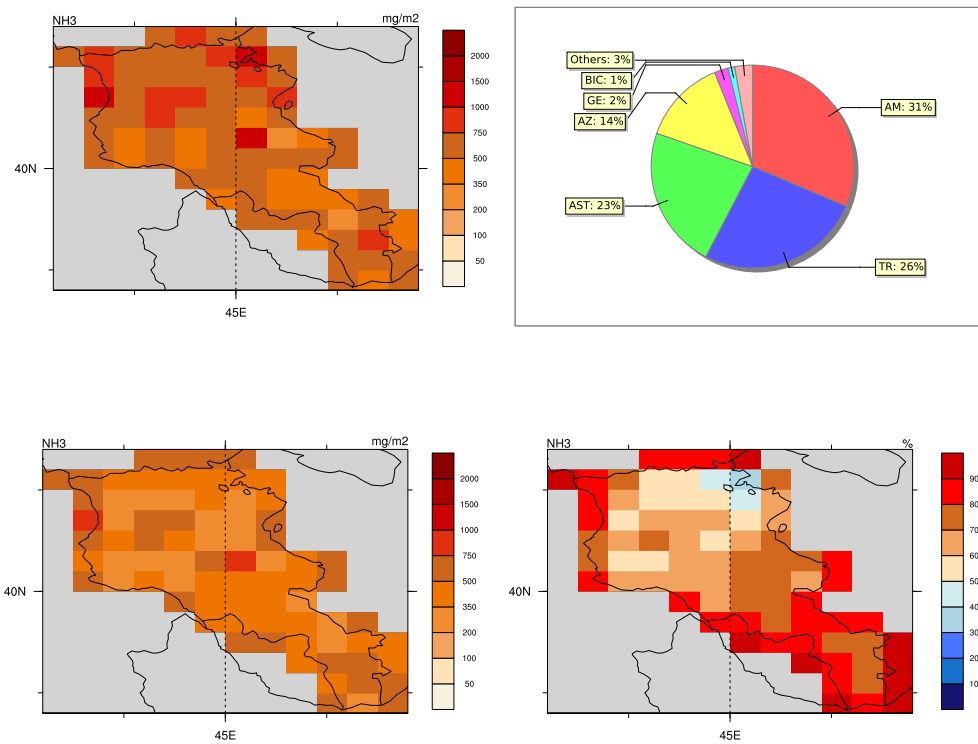


Figure 11: Top left: Deposition of reduced nitrogen in Armenia. Unit:  $\text{mg(N)/m}^2$ . Top right: The six main contributors to deposition of reduced nitrogen in Armenia. Unit: %. Bottom left: Deposition of reduced nitrogen from transboundary sources. Unit:  $\text{mg(N)/m}^2$ . Bottom right: Fraction of transboundary contribution to total deposition. Unit: %.

## 5 Transboundary concentrations of ozone

### 5.1 AOT40<sup>UC</sup><sub>f</sub>

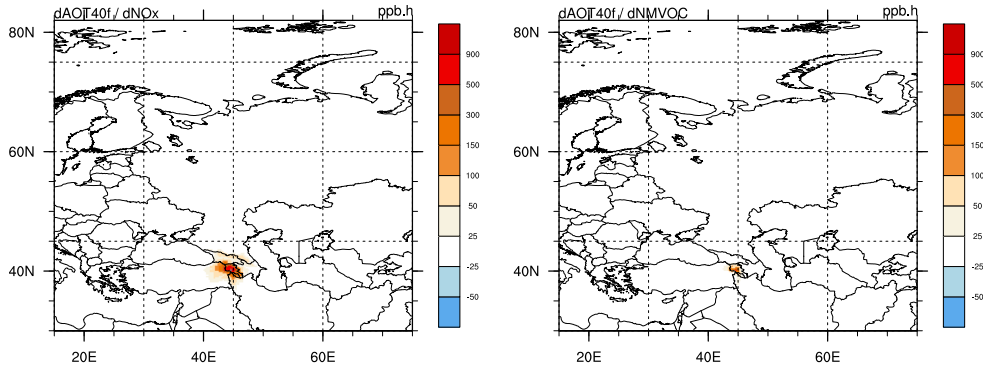


Figure 12: Reduction in AOT40<sup>UC</sup><sub>f</sub> that would result from a 15% reduction in emissions of NO<sub>x</sub> (left) and NMVOC (right) from Armenia. Unit: ppb·h.

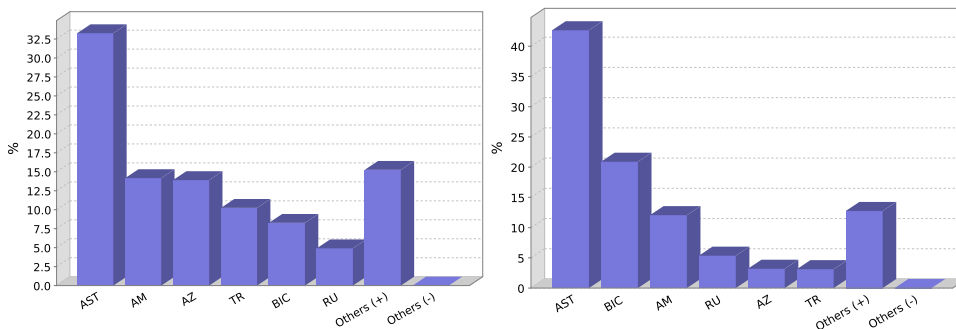


Figure 13: The six most important emitter countries or regions, with respect to their effects on AOT40<sup>UC</sup><sub>f</sub> in Armenia that would result from reductions in NO<sub>x</sub> emissions (left) or NMVOC emissions (right). The sum of the absolute values of the effects of all emitter countries corresponds to 100%. See Section 1.1 for more information.

## 5.2 $POD_{1.0,gen-DF}$ – Ozone fluxes to deciduous forests

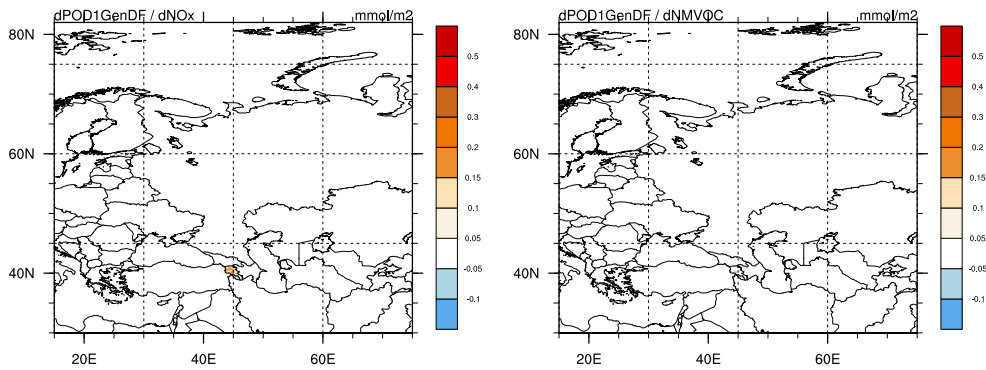


Figure 14: Reduction in  $POD_{1.0,gen-DF}$  that would result from a 15% reduction in emissions of  $NO_x$  (left) and NMVOC (right) from Armenia. Unit:  $mmol/m^2$ .

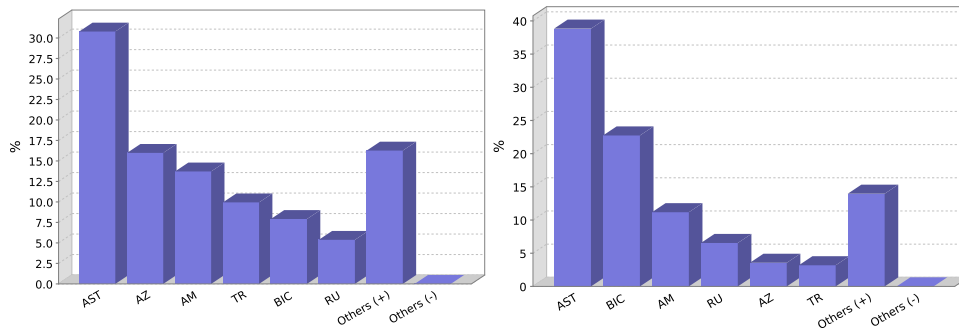


Figure 15: The six most important emitter countries or regions, with respect to their effects on  $POD_{1.0,gen-DF}$  in Armenia that would result from reductions in  $NO_x$  emissions (left) or NMVOC emissions (right). The sum of the absolute values of the effects of all emitter countries corresponds to 100%. See Section 1.1 for more information.

### 5.3 SOMO35 – Risk of ozone damages to human health

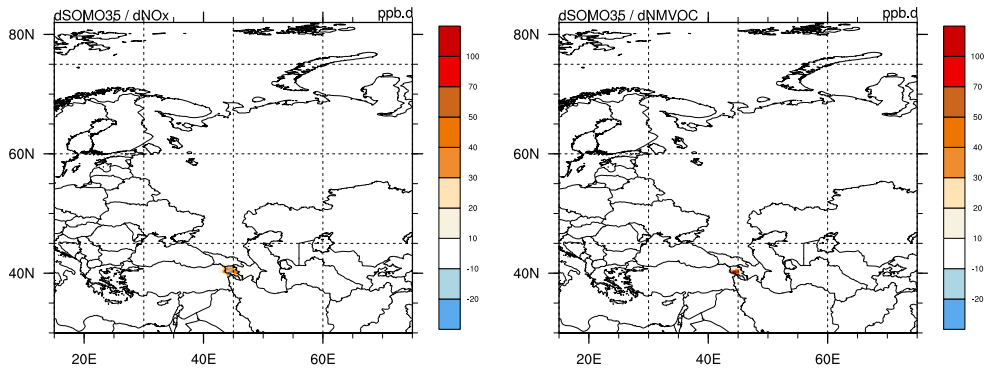


Figure 16: Reduction in SOMO35 that would result from a 15% reduction in emissions of  $\text{NO}_x$  (left) and NMVOC (right) from Armenia. Unit: ppb-day.

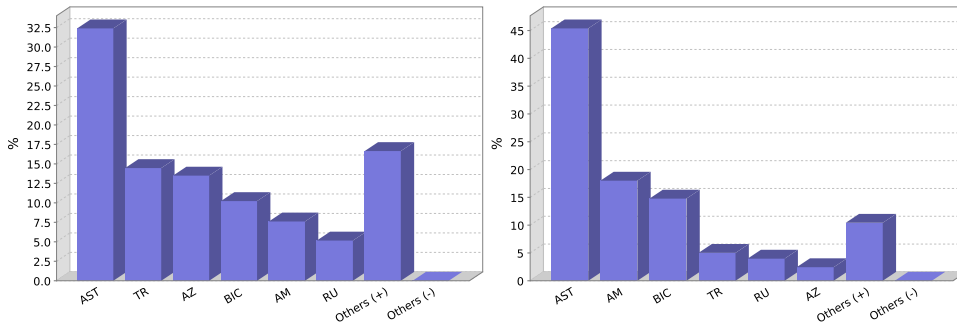


Figure 17: The six most important emitter countries or regions, with respect to their effects on SOMO35 in Armenia that would result from reductions in  $\text{NO}_x$  emissions (left) or NMVOC emissions (right). The sum of the absolute values of the effects of all emitter countries corresponds to 100%. See Section 1.1 for more information.



## 6 Transboundary concentrations of particulate matter

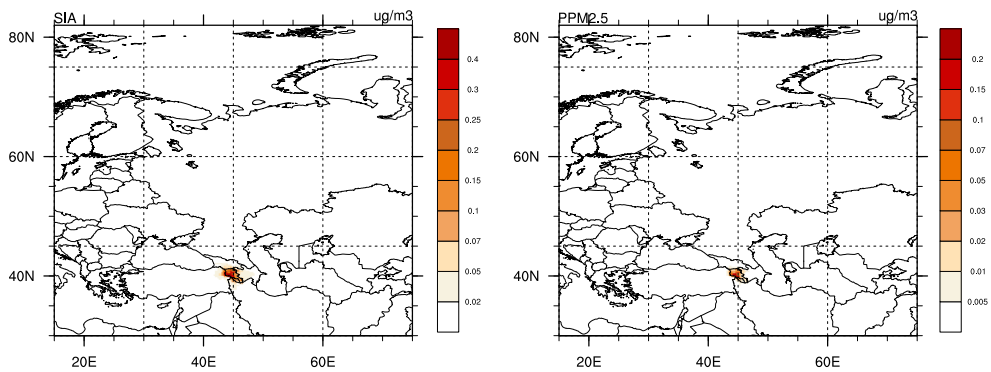


Figure 18: Reduction in concentrations of SIA (left) and  $PPM_{2.5}$  (right) that would result from a 15% reduction in emissions from Armenia. Unit:  $\mu\text{g}/\text{m}^3$ . Note the difference in scales.

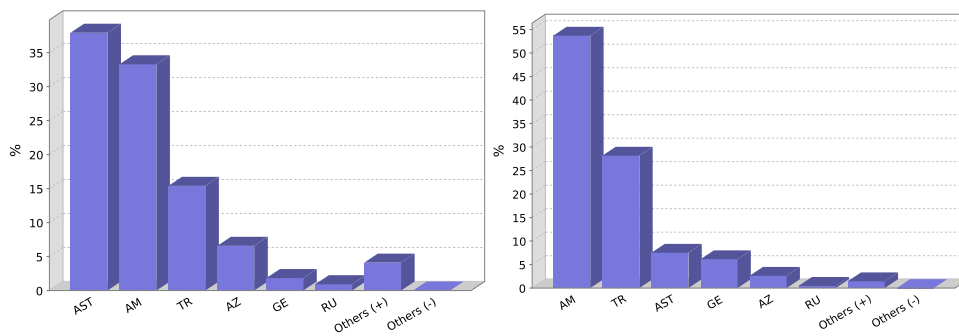


Figure 19: The six most important emitter countries or regions, with respect to their effects on SIA (left) or  $PPM_{2.5}$  (right) in Armenia that would result from reductions in emissions. The sum of the absolute values of the effects of all emitter countries corresponds to 100%. See Section 1.1 for more information.

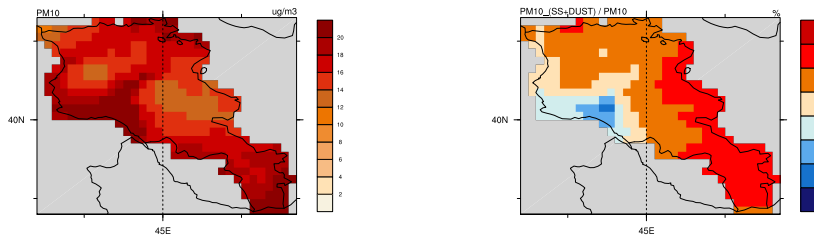


Figure 20:  $PM_{10}$  concentration (left) and fraction of natural contributions of  $PM_{10}$  (sea salt and natural dust) to total  $PM_{10}$  (right) in Armenia.

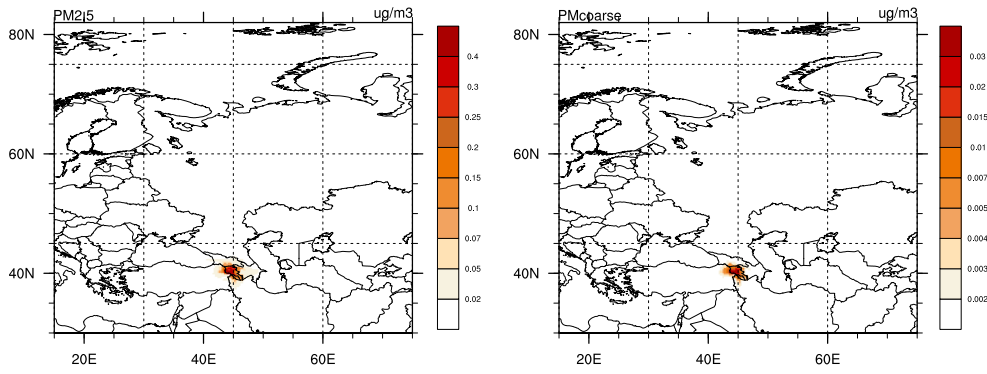


Figure 21: Reduction in  $PM_{2.5}$  and  $PM_{coarse}$  concentrations that would result from a 15% reduction of emissions from Armenia. Unit:  $\mu g/m^3$ . Note the different color scales.

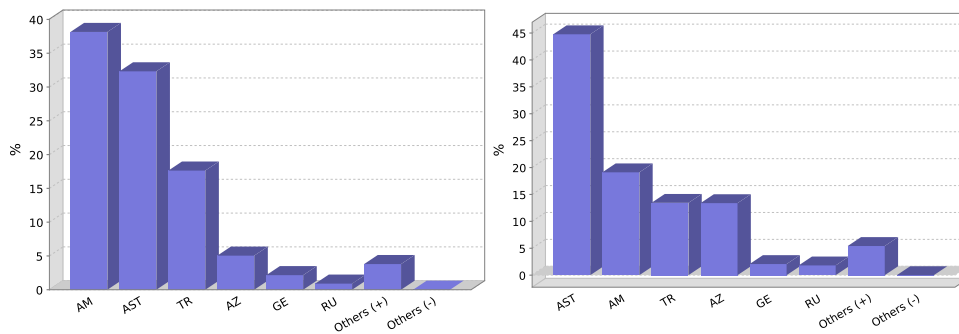


Figure 22: The six most important emitter countries or regions, with respect to their effects on  $PM_{2.5}$  (left) or  $PM_{coarse}$  (right) in Armenia that would result from reduction in emissions. The sum of the absolute values of the effects of all emitter countries corresponds to 100%. See Section 1.1 for more information.

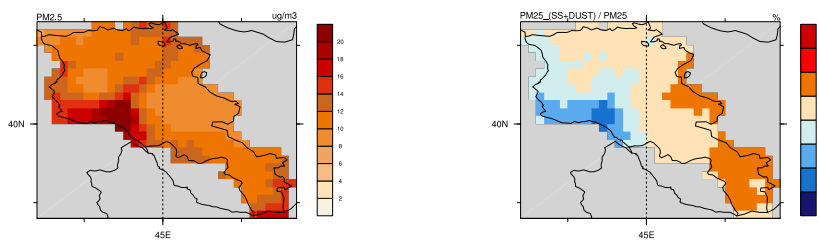


Figure 23:  $PM_{2.5}$  concentration (left) and fraction of natural contributions of  $PM_{2.5}$  (sea salt and natural dust) to total  $PM_{2.5}$  (right) in Armenia.

## 7 Comparison with observations



Figure 24: Location of stations in Armenia.

A sufficiently consistent set of daily ozone observations in AM for 2017 is not available for this analysis.

Figure 25: Frequency analysis of ozone in Armenia at the stations that reported O<sub>3</sub> for 2017 (Model, Observations).

Component	No.	Bias	Correlation	RMSE
SO2 in Air	1	206%	0.20	0.79
Sulfate in Air	1	45%	0.59	0.36
NO2 in Air	1	127%	-0.33	0.44
NO3- in Air	1	29%	0.55	0.27
NH3+NH4+ in Air	1	-34%	-0.40	2.35
PM10	0			
PM2.5	0			
Ozone daily max	0			
Ozone daily mean	0			
SO4 wet dep.	0			
Nitrate wet dep.	0			
Ammonium wet dep.	0			
Precipitation	0			

Table 5: Annual statistics of comparison of model results with observations in Armenia for stations with a sufficiently consistent set of data available in weekly or higher time-resolution. Standard deviations provide variability ranges between stations.

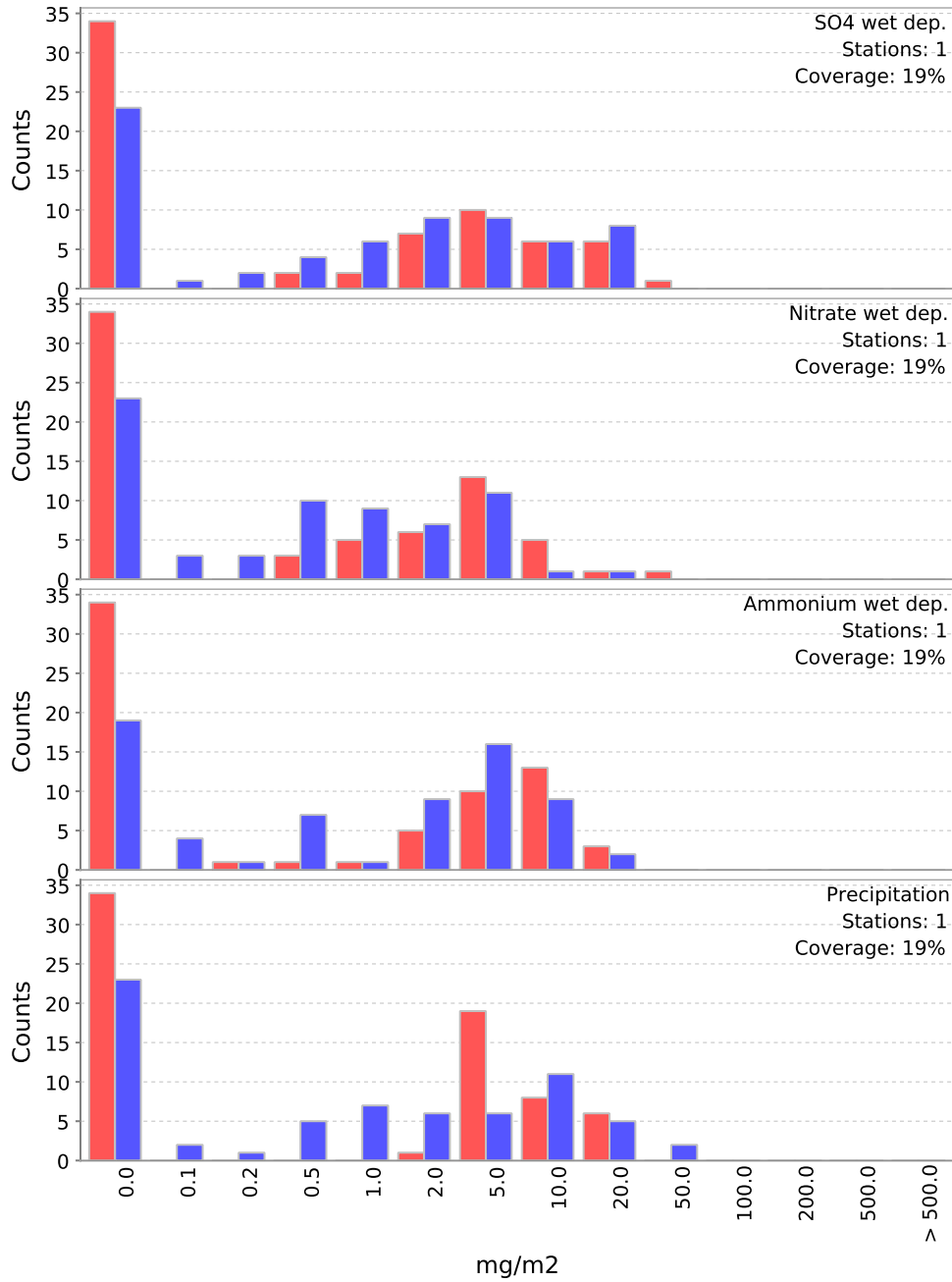


Figure 26: Frequency analysis of depositions in precipitation in Armenia (Model, Observations).

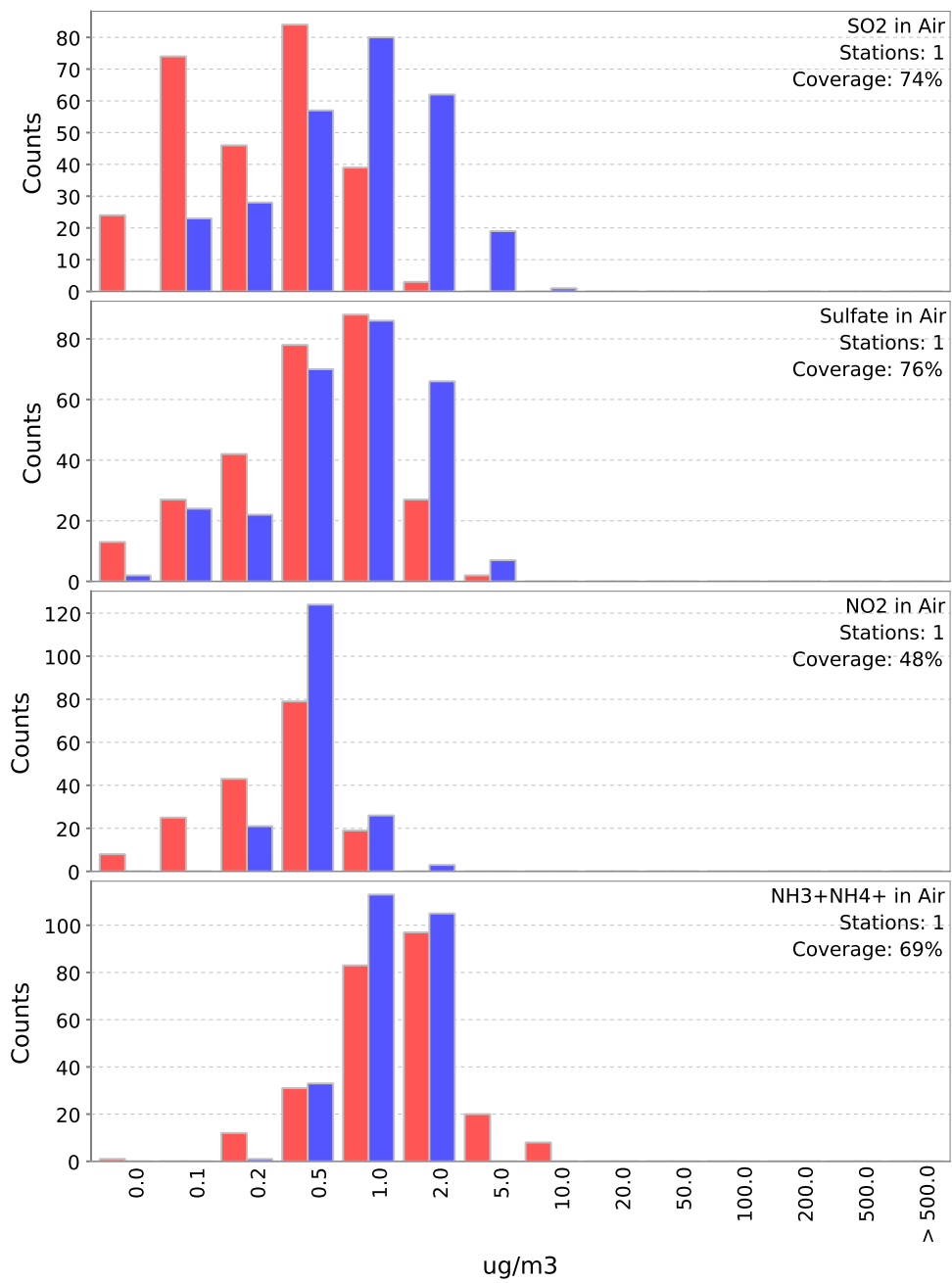


Figure 27: Frequency analysis of air concentrations in Armenia (Model, Observations).

No daily Airbase observations of  $\text{NO}_2$  in AM for 2017 available for this analysis.

No daily Airbase observations of  $\text{SO}_2$  in AM for 2017 available for this analysis.

No daily Airbase observations of  $\text{PM}_{10}$  in AM for 2017 available for this analysis.

No daily Airbase observations of  $\text{PM}_{2.5}$  in AM for 2017 available for this analysis.

No daily Airbase observations of  $\text{O}_3$  in AM for 2017 available for this analysis.

Figure 28: Daily model results versus AirBase observations in Armenia for  $\text{NO}_2$ ,  $\text{SO}_2$ ,  $\text{O}_3$ ,  $\text{PM}_{10}$  and  $\text{PM}_{2.5}$  if available.

## 8 Risk of damage from ozone and particulate matter in Armenia

### 8.1 Ecosystem-specific AOT40 values

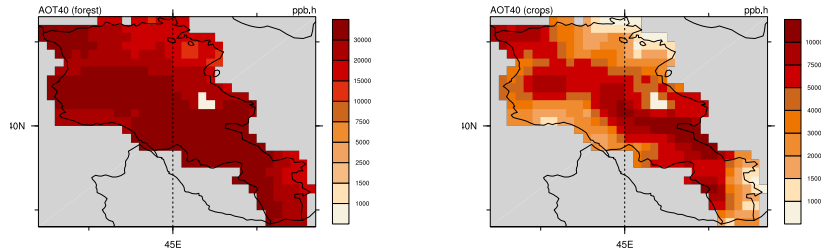


Figure 29:  $AOT40_f^{uc}$  and  $AOT40_c^{uc}$  in Armenia in 2017. ( $AOT40_f^{uc}$ : growing season April-September, critical level for forest damage = 5000 ppb-h;  $AOT40_c^{uc}$ : growing season May-July, critical level for agricultural crops = 3000 ppb-h.)

### 8.2 Ecosystem-specific ozone fluxes

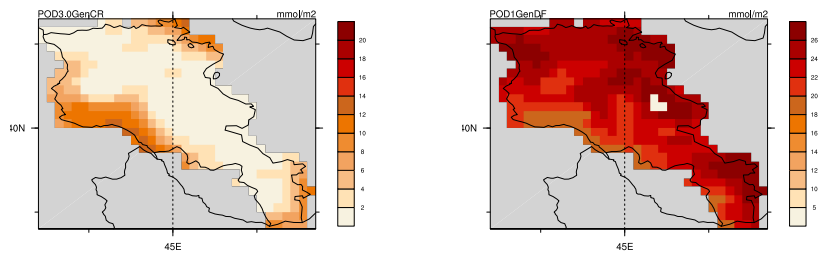


Figure 30:  $POD_{3.0,gen-CR}$  and  $POD_{1.0,gen-DF}$  in Armenia in 2017.

### 8.3 Health impacts from ozone and particulate matter

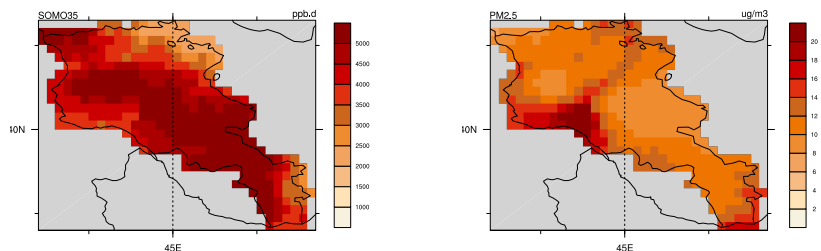


Figure 31: Regional scale  $SOMO_{35}$  and  $PM_{2.5}$  in Armenia in 2017.