

# An alternative way to evaluate chemistry-transport models variability

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**Abstract.** A simple and complementary model evaluation technique for regional chemistry-transport is discussed. The methodology is based on the concept that we can learn on models performances by comparing the simulation results with observational data available for other time periods than the period originally targeted. First, the statistical indicators selected in this study (spatial and temporal correlations) are computed for a given period, using co-localised observation and simulation data in time and space. Second, the same indicators are used to calculate scores for several other years by conserving only the spatial locations and Julian days of the year. The difference between the results provides complementary insights on the model capability to reproduce well the observed variability. In order to synthesise the large amount of results, a new indicator is proposed, designed to compare the several error statistics between all the years of validation and to quantify if the studied period was fairly modelled for the good reasons.

## 10 1 Introduction

Chemistry transport models (CTM) aim at simulating the atmospheric composition where humans and the environment can be affected by air pollution. Air pollution results from the presence of chemical components emitted into the atmosphere due to anthropogenic activities and natural sources (biogenic emissions from vegetation, soil erosion, sea salts, volcanic activity, and wild-land fires). CTMs are used to represent the dynamical and chemical processes that drive spatial and temporal features of the atmospheric composition.

To estimate the quality of CTMs, model output results are usually compared with available observations. These comparisons are performed since the models exist: this is crucial to quantify the ability of models to reproduce particular observed events or a general behaviour. The quantification of the model quality is performed in every research study: depending on the studied case, the modelled variables, the spatial and temporal resolution. The comparison between observations and model outputs is a complex task and has to take into account numerous factors, such as, for example the spatial representativeness of the monitoring stations (Valari and Menut, 2008; Solazzo and Galmarini, 2015). From many years, the way to evaluate a model

results is discussed and in the field of atmospheric composition, numerous methods were proposed. This is not possible to give an exhaustive list of all validation studies and we present here some examples.

Baldrige and Cox (1986) and Cox and Tikvart (1990) proposed the use of error statistics like correlation, bias, Root Mean Squared Error (RMSE) in the specific framework of air quality, *i.e.* the atmospheric composition when criteria pollutant concentrations exceed pre-defined limit values. Chang and Hanna (2004) also proposed an evaluation framework dedicated to air quality model performance and explained there is not "*a single best evaluation methodology*" and how important it is to use as much as possible evaluation criteria to really well understand model results. Later, and in order to ensure the use of systematic procedures in the evaluation process, dedicated tools were developed for the model evaluation. For example, Appel et al. (2011) and Galmarini et al. (2012) proposed complex statistical modules to extract all possible information related to the capability of a model to reproduce an observed event. In parallel, some studies were dedicated to revisit the way to evaluate models such as Thunis et al. (2012), dedicated to air quality in a policy framework. In this study, they proposed the "Target diagram" to have on the same plot the bias and the RMSE. Complementary to the definition of performance indicators to be used, Simon et al. (2012) use these indicators to compile photochemical models performances over a large set of data over several years of simulation. This kind of evaluation may also be done in dedicated projects such as the recent AQMEII (Air Quality Model Evaluation International Initiative), comparing chemistry-transport models running both in Europe and Northern America, (Vautard et al., 2012; Campbell et al., 2015) or the EURODELTA project, (Bessagnet et al., 2016) and in the EMEP (European Monitoring and Evaluation Programme) context in the frame of the United Nation Convention on Long-range Transboundary Air Pollution, (Prank et al., 2016). Using comparisons between observations and models outputs, some studies proposed methodologies to decompose the statistical scores in order to estimate the main source of errors, (Solazzo and Galmarini, 2016). Finally, other studies also use observations to adjust the result by implementing methods to unbiased simulation without changing the model, as in Porter et al. (2015) for ozone over the United States. The common point of all these studies is that they are always using, as best as possible, the observations corresponding in time and location to the model cell.

In the present study, a simple method is proposed to add information about the model performances more focussed on its variability. To reach this objective, we propose to use observations corresponding to the modelled period and geographical domain but also, to use observations for the same domain but other periods. By this way, we want to extract the information about the model variability and to answer the question: *Are the performances of the model satisfactory because the model is accurate or just because the model is able to reproduce a situation which is recurrent from year to year?* The issue to be solved and the tools developed are presented in section 2. The new methodology with the presentation of the indicator developed for this study are presented in section 3. The results and discussions to point out the drivers of model errors are presented in section 4 and section 5 for the new indicator.

## 2 Methodology

In the present study, a simple method is developed to improve the evaluation of models variability and to identify the processes responsible for discrepancies of models outputs *versus* observations. The methodology is general and could be applied to

55 all types of model. In this study, the methodology is presented for the specific case of the regional atmospheric composition modelling: a topic mixing meteorology and chemistry, with a high spatial and temporal variability, thus having a good potential to test the interest of our methodology.

## 2.1 Regional chemistry-transport modelling

In chemistry-transport modelling, several processes are involved, some of them directly influencing the others. When studying both meteorological and chemical variables, the dependencies between all variables are helpful to know how to better interpret the model results.

The boundary conditions prescribe the concentrations of chemical species which may enter the simulation domain. Usually for large domains, they are issued from global models as monthly climatologies. They correspond to averaged values suitable to characterize the background concentrations of long-lived species such as ozone, carbon monoxide, mineral dust. Anthropogenic emissions are prescribed from databases and the influence of meteorology is limited in the model. Vegetation, fires and mineral dust emissions depend both on landuse data and meteorology. These emissions are not measurable, it is almost impossible to directly quantify their realism.

The meteorological variables influence transport and mixing processes, with a direct effect on gas and aerosol plumes locations and their vertical distribution. Cloudiness and temperature impact the photolysis efficiency, the boundary layer height impact the surface mixing of pollutants, rainfall impact the wet deposition. Moreover, meteorology impact emissions: wind variability is the prevalent driver for dust emissions, and it has also a major impact on wildfires emissions. Both temperature and solar irradiance influence the magnitude of biogenic emissions from vegetation. The spatial variability of landuse data has also a strong impact on all these natural emissions.

The chemistry-transport model is a numerical integration tool of all forcings and processes. The chemical mechanism handles the life cycle of chemical species (production and loss) when the deposition processes are the only net sink of species. With the model, the spatial (horizontal and vertical) and temporal resolutions are also defined, directly impacting the simulation representativeness and thus the realism of the modelled air pollutant concentrations when they are compared to available observations.

## 2.2 The studied case

25 The study focuses on the summer 2013 period (1st May to 31 August) over the Euro-Mediterranean region. This period is called "reference period" in this paper. This case has already been modelled (using the same models, WRF and CHIMERE) and the results were discussed in [Menut et al. \(2015\)](#). The same simulation is used in this study, all parameters are identical.

The observational data come from different sources depending on the variables, [Table 1](#). In this region, where the monitoring network are dense enough, comparisons are performed with observations from surface stations that provide hourly  $O_3$ ,  $NO_2$  surface concentrations for gases and  $PM_{2.5}$  and  $PM_{10}$  (particulate matter with mean mass median diameter lower than 2.5 and  $10\mu m$ , respectively) for particles. Complementary to surface concentrations data, evaluated using the EBAS database, [\(Tørseth et al., 2012\)](#), the meteorology is also evaluated for 2m temperature,  $T_{2m}$ , 10m wind speed,  $U_{10m}$ , and precipitation

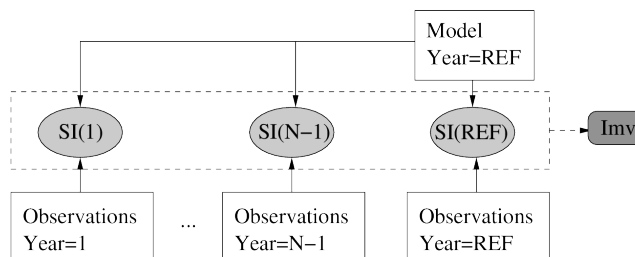
Variable	Network	Spatial coverage	Vertical coverage	Temporal frequency	Unit
O <sub>3</sub> , NO <sub>2</sub>	EBAS/EMEP	Europe	Surface	Hourly	ppb
PM <sub>2.5</sub> , PM <sub>10</sub>	EBAS/EMEP	Europe	Surface	Hourly	μg m <sup>-3</sup>
AOD, Angström	AERONET	Global	Column	Hourly	ad.
T <sub>2m</sub>	BADC	Global	Surface	Tri-hourly	°C
U <sub>10m</sub>	BADC	Global	Surface	Tri-hourly	m s <sup>-1</sup>
Precipitation	BADC	Global	Surface	Daily	mm day <sup>-1</sup>

**Table 1.** List of measurements data used for the statistical comparison with the model results. All data used are issued from surface stations, representative of their own environment. Originally provided hourly or three-hourly, they are used as daily averaged in this work.

rates (in mm day<sup>-1</sup>) from the BADC (British Atmospheric Data Centre). In order to quantify the transport of aerosols in dense plumes aloft, observations from AERONET (AErosol RObotic NETwork) program are used for the optical depth, AOD, and the Angström exponent. In this study, all variables are used as daily mean (except for precipitation corresponding to daily cumulated values) in order to (i) have homogeneous scores between the variables, (ii) be able to separate the systematic and the day-to-day variabilities. The use of an hourly time frequency was ruled out to avoid a too strong weight of the diurnal cycle in the temporal variability.

### 3 The proposed methodology

As discussed in the introduction, many Statistical Indicators (SI) exist to quantify the model ability to realistically simulate observed pollution events. The correlations (temporal and spatial), the Root Mean Squared Error (RMSE), its normalized expression nRMSE, and the bias (the difference between observations and modelled values) are widely used in regional air pollution modelling. The correlations are able to split the relative contributions of systematic meteorology or sources related variability and day-to-day variability. The RMSE and the bias are a direct quantification of the model error.



**Figure 1.** Principle of the multi-year variability indicator ( $I_{mv}$ ) calculation, using one modelled year and several year of observations. SI stands for "Statistical Indicator" and is related to spatial and temporal correlation.

45 The main goal of this study is to separate the contributions due to systematic and sporadic events. The systematic events correspond to yearly phenomena when the sporadic events correspond to event observed during one year but not the others. In addition, complementary to the model variability quantification, the model error is also important to estimate. The key point of this study is to (i) study the model variability which is statistically represented by the correlations, and (ii) add complementary information on the model errors, here that could be represented by the RMSE (or the nRMSE).

50 First, as presented in **Figure 1**, the SI are calculated between observations data and model outputs for the simulation year (*i.e.* the reference year). Second, the SI are calculated between the observations data for other years and the model output for the reference year. Logically, the scores calculated for the reference year for observations and model outputs would give the better results. By difference with the scores calculated for other years (with the observations only), we expect to conclude if the model is able to catch the observed variability and for the good reasons. Using this approach, the goal is to give complementary information to those usually obtained when using only SI calculated for a single year, the studied year.

We apply this methodology for the simulation of the year 2013 and using observations data for years ranging from 2008 to 2013. In order to give some synthetic answers, the different SI scores are aggregated into a single indicator, called  $I_{mv}$  and presented in detail in the next section. Of course it seems apparently awkward to evaluate day by day a model with observational data from another year. For a given station at a given day of the reference year air concentrations will be affected by a different local meteorology, emissions and also long range transport of chemical species. But we can consider that to take the same date for another year is strictly the same that to choose randomly a date in the same season. This trivial method can emphasize how a model is affected by large scale patterns and long term temporal cycles.

### 3.1 Calculation of correlations and nRMSE

In this study, we focus on three Statistical Indicators: the spatial correlation, the temporal correlation and the normalized RMSE. For these three indicators, it is important that, for all years of validation, the same list of stations with valid measurements is used.

The correlation used in this study is the Pearson's' correlation. Each correlation provides specific information on the quality of the simulation. The temporal correlation, noted  $R_t$ , is estimated station by station and using daily averaged data in order to have homogeneous comparisons between all variables. This correlation is directly related to the variability from day to day, for each station.  $O_{t,i}$  and  $M_{t,i}$  represent the observed and modelled values, respectively, at time  $t$  and for the station  $i$ , for a total of  $T$  days and  $I$  stations. The mean time averaged value  $\overline{X}_i$  is:

$$\overline{X}_i = \frac{1}{T} \sum_{t=1}^T X_{t,i} \quad (1)$$

The temporal correlation  $R_{t,i}$  for each station  $i$  is calculated as:

$$R_{t,i} = \frac{\sum_{t=1}^T (M_{t,i} - \overline{M}_i)(O_{t,i} - \overline{O}_i)}{\sqrt{\sum_{t=1}^T (M_{t,i} - \overline{M}_i)^2 \sum_{t=1}^T (O_{t,i} - \overline{O}_i)^2}} \quad (2)$$

75 The mean temporal correlation,  $R_t$ , used in this study is thus:

$$R_t = \frac{1}{I} \sum_{i=1}^I R_{t,i} \quad (3)$$

with  $I$  the total number of stations. The spatial correlation, noted  $R_s$ , uses the same formula type except it is calculated from the temporal mean averaged values of observations and model for each location where observations are available. A good correlation shows that the model correctly locates the largest horizontal gradients as known sources and long range transport  
80 plumes.

The spatio-temporal mean averaged value is estimated as:

$$\bar{X} = \frac{1}{I} \sum_{i=1}^I \bar{X}_i \quad (4)$$

and the spatial correlation is thus expressed as:

$$R_s = \frac{\sum_{i=1}^I (\bar{M}_i - \bar{M})(\bar{O}_i - \bar{O})}{\sqrt{\sum_{i=1}^I (\bar{M}_i - \bar{M})^2 \sum_{i=1}^I (\bar{O}_i - \bar{O})^2}} \quad (5)$$

85 The normalized Root Mean Square Error is expressed as:

$$nRMSE = \sqrt{\frac{1}{T} \frac{1}{I} \sum_{t=1}^T \sum_{i=1}^I \left( \frac{O_{t,i} - M_{t,i}}{O_{t,i}} \right)^2} \quad (6)$$

for all stations  $i$  and all times  $t$ .

### 3.2 Definition of the $I_{mv}$ indicator

For the specific purpose of the model variability (and not the model error), we define an indicator,  $I_{mv}$ , dedicated to express  
90 in one value the results obtained with the temporal and spatial correlations. The goal of this indicator is to quantify how the correlation between measurements data (for different years) and model outputs (for the reference year) evolves from a year to another one. This indicator does not replace the usual statistical indicators but aims at providing complementary information about the variability between years.

We first define the differences,  $D$ , between all years as:

$$95 \quad D = \frac{1}{N-1} \left( \sum_{i=1}^{N-1} |s_i - s_N| \right) \quad (7)$$

with  $s_N$  the score of the indicator for the reference year being modelled and  $s_i$  the score of the indicator computed using observations corresponding to other meteorological years (from 1 to  $N - 1$  if there is  $N - 1$  other available years for the observations).

We now aim to develop a simple indicator, called  $I_{mv}$ , which is a combination of the statistical indicator for the reference year and the differences between years. This  $I_{mv}$  corresponds in fact to the SI itself weighted by the differences between the SI scores of all years. We want that  $I_{mv}$  follows these rules:

- $I_{mv}$  has the same evolution than the studied SI. If the correlation increases,  $I_{mv}$  also increases.
- $I_{mv}$  is bounded between 0 and 1 such as the correlation. This enables to compare the results for different variables (with different metrics).
- In case of high correlation value found for the studied year i.e ideally  $s_N$  tends to 1:
  - If the differences between the other years are low ( $D$  tends to 0), it means that the model is correct for the studied year, but possibly because it reproduces a recurrent phenomena. In this case, we want that  $I_{mv}$  decreases and tends to 0.
  - If the differences between the other years are high ( $D$  tends to 1): in this case, the model gives good results for the studied year and this is not because it simulates a systematic event. In this case, we want that  $I_{mv}$  remains close to the indicator value. With  $s_N \approx 1$  and  $I_{mv} \approx 1$ , we can conclude that the model is very good for the studied year and this is not due to an easy phenomenon to model.
- In case of low correlation value, and whatever the magnitude of differences between years, the model is not correct.  $I_{mv}$  must be low, as the indicator value.

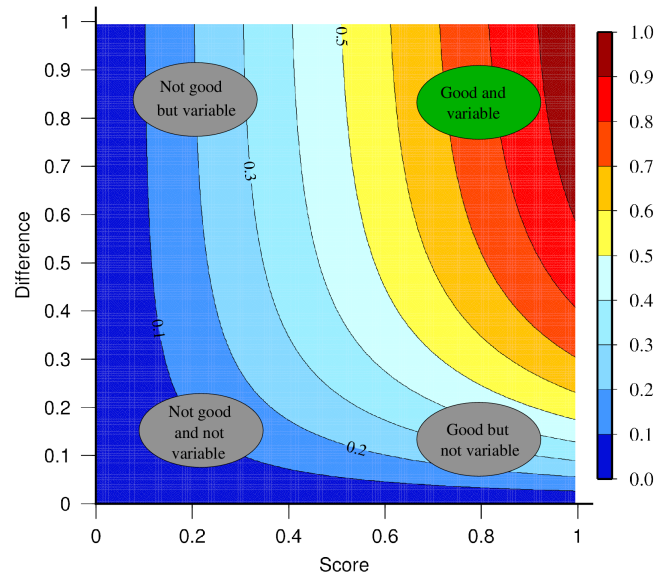
These constraints induce to define an indicator having this kind of formulation:

$$I_{mv} = s_N (1 - \exp(-D_s)^4) \quad (8)$$

s

This means that  $I_{mv}$  has always, as maximum value, the value of the indicator itself. The power 4 is here defined to have a specific shape for  $I_{mv}$  respecting the rules presented below. Finally, this expression gives an indicator variability presented in **Figure 2**. Considering the state-of-the art of chemistry-transport modelling, the model is considered as accurate and to have an acceptable variability for  $I_{mv} > 0.4$ : this means that the correlation is at least 0.5 and the differences are also at least greater than 0.5.

Finally, this indicator is not calculated for nRMSE and bias. Two reasons explain this choice: (i) contrarily to correlations, RMSE and bias are not bounded between 0 and 1. This leads to indicators values possibly varying a lot between several years and thus difficult to compare between years. (ii) The goal of the indicators is to extract a message from the model variability of



**Figure 2.** Scheme of the  $I_{mv}$  values as a function of the studied year correlations values and the multi-years differences  $D$ .

the studied year compared to the other years. In this case, the correlations represent statistical indicators more adapted to this evaluation.

## 25 4 Time series of statistical indicators

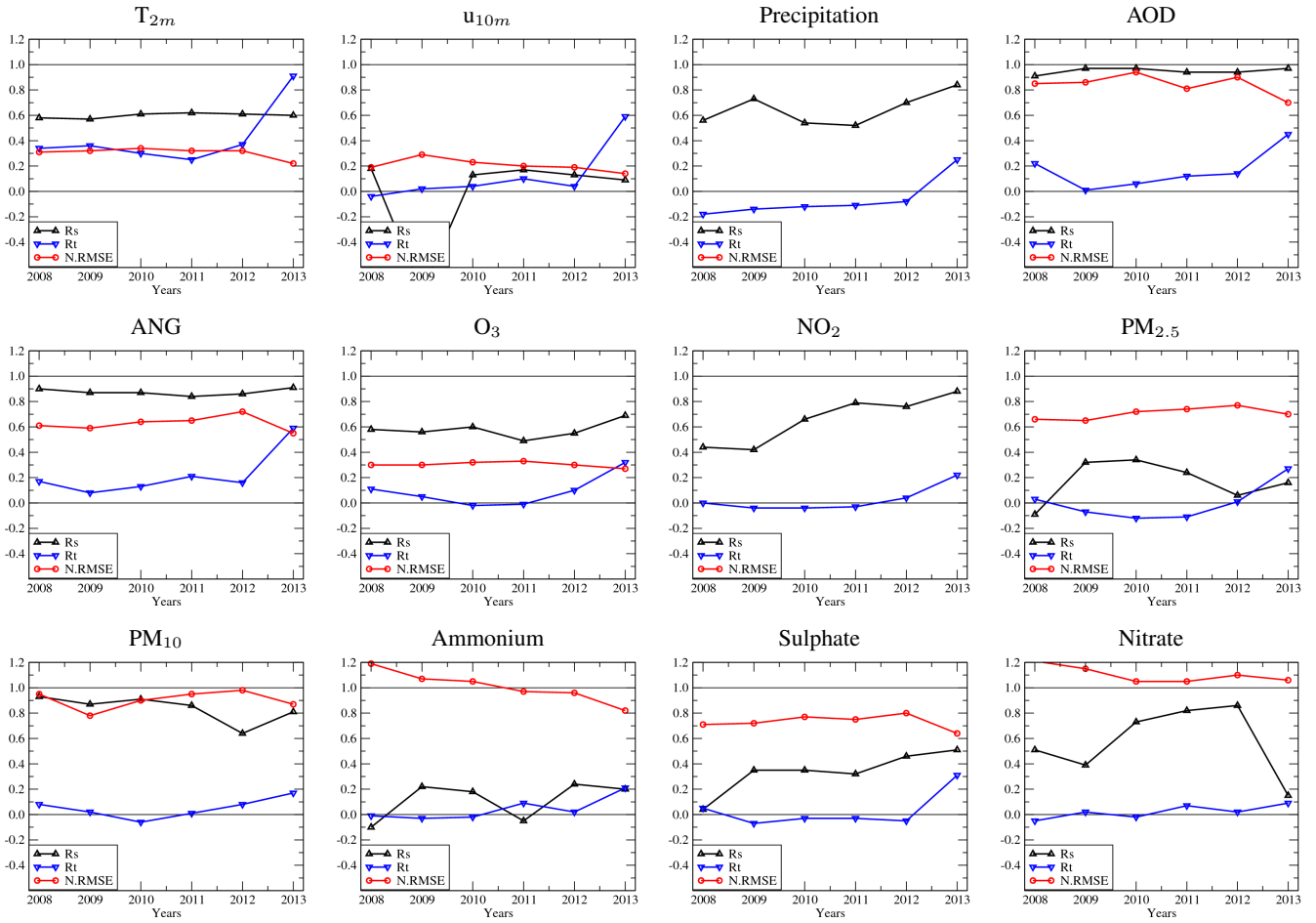
The calculations of differences are performed for the correlations and the nRMSE. These values are calculated for all variables described in Table 1 and for the years 2008 to 2013. For each year, it is reminded that only the May to August period is considered. Results are presented as time series in Figure 3 and discussed in the following sections. Note also that some values discussed in these sections are also reported in the synthetic Table 4.

### 30 4.1 Meteorological variables

The meteorological variables are  $T_{2m}$ ,  $u_{10m}$  and the precipitation rate. The values of the Statistical scores are provided, year by year, in Figure 3. As an example, the same values are reported for  $T_{2m}$  in Table 2.

$T_{2m}$  is a meteorological variable, constraining processes both for meteorology and chemistry. Its diurnal cycle is well marked as its latitudinal variability (for large model domains), often ensuring a good spatial correlation. In general, this variable is the less uncertain of all modelled meteorological parameters. The spatial correlation is good for all years, ranging from 0.57 (2009) to 0.62 (2011). For the studied year (2013), the score is 0.60, slightly lower than for 2011. Even if the correlation for the selected year is good, it is not significantly better than for the other year, with  $D=0.02$ . This means that the model reproduces fairly well a spatial pattern that is observed every year. Indeed, the simulation domain is large and the temperature has a latitudinal





**Figure 3.** Multi years scores for  $T_{2m}$ ,  $u_{10m}$ , the precipitation rate, Aerosol Optical Depth (AOD), Angström exponent (ANG), surface concentrations of  $O_3$ ,  $NO_2$ ,  $PM_{2.5}$ ,  $PM_{10}$ , Ammonium, Sulphate and Nitrate. The correlations and the nRMSE are calculated between the observations (2008-2013) and the model results (2013). The spatial correlation,  $R_s$ , is in black, the temporal correlation,  $R_t$  in blue, the nRMSE in red..

variability larger than between each measurements stations. The temporal correlation ranges from 0.25 to 0.91 (2013). The nRMSE is less evolving than the correlations, with values ranging from 0.22 (2013) to 0.34 (2010). The lowest value is found for 2013, highlighting the fact that the model error is the lowest for the reference year. The model is thus performing well in capturing the day to day variability for  $T_{2m}$  and for the good reasons.

From **Figure 3**, the calculation of  $u_{10m}$  also gives satisfactory results with  $R_t=0.60$ . The spatial correlation,  $R_s=0.09$ , is not correct and very variable from one year to another. As for  $T_{2m}$ , we also have an effect of the model resolution and the representativeness of the variable.

Year	$R_s$	$R_t$	nRMSE
2008	0.58	0.34	0.31
2009	0.57	0.36	0.32
2010	0.61	0.30	0.34
2011	0.62	0.25	0.32
2012	0.61	0.37	0.32
2013	0.60	0.91	0.22
<i>D</i>	0.02	0.59	0.10

**Table 2.** Scores for  $T_{2m}$ . The correlations and nRMSE are calculated between the observations (2008-2013) and the model results (2013).

Scores for the precipitation are correct, with a very good spatial correlation, always exceeding 0.6. As for the temperature, the latitudinal effect plays a major role in the variability. Both the spatial and temporal correlations increase significantly for the reference year. The nRMSE is not on the plot, the values being larger than 1.2. The model is biased in absolute values and overestimates the amount of daily precipitation. But the day to day variability is correct and this is what is requested in  
50 atmospheric composition modelling (the lower atmosphere is scavenged when a precipitation occurs, whatever its value).

For the meteorological variables, these scores showed that the meteorological forcing is well retrieved, and always better for the year being considered compared to other years.

## 4.2 Optical properties

The optical properties are directly linked to the atmospheric composition of aerosol and may be quantified using the Aerosol  
55 Optical Depth (AOD) and the Angström exponent (ANG).

For the AOD, the spatial correlation is very good for 2013,  $R_s=0.97$  but it is as good or better for other years. This means that we model a rather recurring phenomenon: every year the same stations are on average exposed to aerosol plumes. The temporal correlation is lower with  $R_t=0.45$  but much better than for other years. This indicates that the model partly reproduces the observed temporal variability but the events are changing from one year to another and the model captures well these changes.  
60 In the studied region, the AOD are sensitive to desert dust outbreaks in summer. This means that large scale systems are driving the aerosol plumes; they are spatially recurrent and temporally better estimated for the year being considered than for other years.

For the ANG, the spatial correlation is very good,  $R_s=0.91$  but also persistent in time. The temporal correlation is much better for 2013 than other years. This is probably due to a size distribution that is not necessarily well simulated from one day  
65 to another (showed by AOD and explained in (Menut et al., 2016)) but the relative contributions of fine and coarse aerosol atmospheric load are fairly reproduced. This feature highlights the high sensitivity of the AOD calculation to the modelled aerosol size distribution, although the overall mass emitted and transported is realistic.

Globally, the AOD and ANG reflect the model's ability to retrieve the long range transport of long-lived aerosols which depends on several processes (emissions, transport, and deposition). These scores show the model is able to retrieve these yearly recurrent plumes but the model size distribution of particles clearly requires improvements.

### 4.3 Surface concentrations

For the surface concentrations of gaseous and aerosol species, the variability is much more related to local effects. As an example, the detailed values of the statistical indicators and the differences between years are fully presented for NO<sub>2</sub>.

Year	R <sub>s</sub>	R <sub>t</sub>	nRMSE
2008	0.44	0.00	1.56
2009	0.42	-0.04	1.76
2010	0.66	-0.04	1.82
2011	0.79	-0.03	2.07
2012	0.76	0.04	2.84
2013	0.88	0.22	1.76
<i>D</i>	0.27	0.23	0.33

**Table 3.** Scores for NO<sub>2</sub>. The correlations and nRMSE are calculated between the observations (2008-2013) and the model results (2013).

NO<sub>2</sub> is both primary and secondary in origin. Mostly emitted in urbanized areas, the diurnal cycle of this species is well constrained. Depending on meteorological conditions, its lifetime may vary significantly, from hours to days. Modelling this species with CTMs is challenging because several uncertainties are acting at the same time, including the spatial representativeness of the model cell. The scores show if the sources are properly placed and if the photochemistry and transport processes have been well simulated. In general, at coarse model resolution, the model results for this species are worse than for ozone. The spatial correlation gives a score of R<sub>s</sub>=0.88 for 2013. This corresponds to the best correlation compared to the other years. The anthropogenic emissions being strongly linked to industrial activities and traffic, and these activity sectors being spatially fixed, the good spatial correlation is more due to anthropogenic sources such as biogenic and vegetations fires. The temporal correlation is low for 2013, R<sub>t</sub>=0.22, but is close to 0 for other years, then significantly better for the reference year compared to the others. These two correlations values show that the model certainly captures the right location of emission sources (low variability of R<sub>s</sub>). The nRMSE is large and shows that the concentrations are overestimated by the model. But this overestimation appears for all years and can be due to the representativeness of the surface measurements compared to the surface of model cells.

The spatial correlation is good for O<sub>3</sub>, NO<sub>2</sub> and PM<sub>10</sub>, with R<sub>s</sub>=0.69, 0.88 and 0.81 respectively. For PM<sub>2.5</sub> this correlation is low with R<sub>s</sub>=0.16. The PM<sub>10</sub> shows that the largest particles are well modelled over the whole domain, and this was also the conclusion for the AOD and ANG. The low score for PM<sub>2.5</sub> indicates that for the aerosol distribution, the fine mode is not as well modelled as the coarse mode. This is confirmed by the scores of the aerosol inorganic species, Ammonium, Sulphate

and Nitrate that contributes to a large fraction of the fine fraction of particles. Except for Sulphate (with  $R_s=0.51$ ), the spatial correlations are 0.15 for Nitrate and 0.20 for Ammonium. Thus, the fine part of the aerosol is not well modelled mainly due to a deficiency in the modelling of nitrates.

The temporal correlations have a completely different behaviour than the spatial correlations. The values are generally low, from  $R_t=0.09$  for Nitrate to  $R_t=0.32$  for  $O_3$ . Surprisingly, the  $PM_{10}$  concentrations display a good spatial correlation but a poor temporal correlation. This is due to the long lifetime in the atmosphere of non-reactive species such as mineral dust: large plumes are correctly modelled over regions but the day to day variability needs improvements. Another point is the good spatial correlation for  $NO_2$  but its low temporal correlation with  $R_t=0.22$ . In this case, this means we have a correctly spatialized anthropogenic emissions inventory (mainly for  $NO_2$  sources) but difficulties to model the day to day chemistry.

For the surface concentrations, we can conclude that  $O_3$ ,  $NO_2$  and  $PM_{10}$  concentrations are spatially well modelled and this is not due to a recurrent behaviour. For particles, the problem is more related to the fine mode, where  $PM_{2.5}$  concentrations are not well located. This modelling problem is highlighted by the low correlations and  $I_{mv}$  values for the inorganic species. For the temporal correlations, the scores are always lower than for the spatial correlation but also always higher for the reference year than for the other years.

## 5 Estimation of the $I_{mv}$ indicator for all variables

To summarize the results obtained for each statistical indicator and the values of differences between all years, we apply the  $I_{mv}$  formulation. This enables to have one values for each SI ( $R_s$  and  $R_t$ ) and each variable. Results are presented in Table 4 and are also displayed on single plots in Figure 4.

In Table 4, the  $I_{mv}$  larger than 0.4 are highlighted. This threshold is clearly subjective but mentioned here to better highlight the variables being well modelled and with a correct variability from a year to another. As discussed in detail, the best scores are obtained for the meteorological variables, and more temporally than spatially.

In Figure 4, The x-axis represents the correlation (spatial or temporal), the y-axis represents the differences between all years,  $D$ . For each studied variables, their values are reported on the figure where the colours represent the value of  $I_{mv}$ . The interpretation of these results follows the quality criteria presented in the academic scheme in Figure 2. This presentation shows an important spread for the spatial correlation results. If the relative differences  $D$  range from 0 to 0.6, the correlations range from 0.09 (for the 10m wind speed) to 0.97 (for AOD). The common point is that there is no variable with differences above 0.5. This means that, spatially, the studied problem shows systematic patterns from year to year. The low values of correlations show that some variables are systematically badly estimated. This means that some meteorological structures (for  $u_{10m}$ ) or emission sources (contributing to the  $PM_{2.5}$  surface concentrations) are systematically mis-located.

The representation of temporal correlations shows a specific linear pattern. The largest correlation values are positively correlated with differences. This temporal correlation represents the day to day variability at each location. This means that the studied problem is based on high day to day variability without similar consecutive days (in this case, one would have high

Variable	$R_s$			$R_t$		
	Value	D	$I_{mv}$	Value	D	$I_{mv}$
$T_{2m}$	0.60	0.02	0.04	0.91	0.59	<b>0.82</b>
$u_{10m}$	0.09	0.23	0.05	0.59	0.56	<b>0.53</b>
precip	0.89	0.20	<b>0.49</b>	0.08	0.07	0.02
AOD	0.97	0.02	0.09	0.45	0.34	0.33
ANG	0.91	0.04	0.14	0.59	0.44	<b>0.49</b>
$O_3$	0.69	0.13	0.29	0.32	0.27	0.21
$NO_2$	0.88	0.27	<b>0.58</b>	0.22	0.23	0.13
$PM_{2.5}$	0.16	0.15	0.07	0.27	0.32	0.20
$PM_{10}$	0.81	0.10	0.27	0.17	0.14	0.07
Ammonium	0.20	0.13	0.08	0.21	0.20	0.12
Sulphate	0.51	0.21	0.29	0.31	0.34	0.23
Nitrate	0.15	0.51	0.13	0.09	0.08	0.03

**Table 4.** The  $I_{mv}$  values for all variables: the meteorology with  $T_{2m}$ ,  $u_{10m}$  and precipitation rate, the vertically integrated column of aerosols with the Aerosol Optical Depth (AOD) and the Angström exponent (ANG), the surface concentrations of all aerosols in term of size distribution with  $PM_{2.5}$  and  $PM_{10}$  and for the inorganic species with  $D_p < 10 \mu m$ . Values of  $I_{mv}$  above 0.3 are bolded. Units of the variables are detailed in [Table 1](#).

35 correlations but low differences). This illustrates the fact that the studied problem is primarily an issue of sporadic events and the model is able to correctly find this variability from one day to another.

## 6 Conclusions

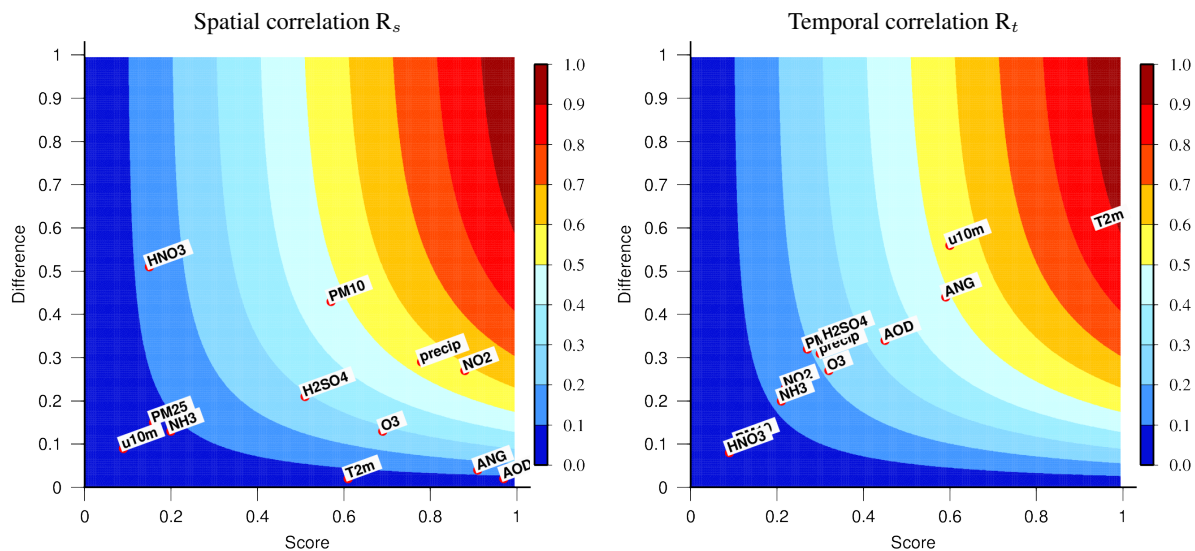
At first glance, using a different year than the simulated one for the day to day evaluation seems awkward. However, we can learn more about the performances of chemistry transport models than using a single year for the usual statistical indicators.

40 Of course, this approach will never replace a strict evaluation of a pollution case analysis using time series, vertical profiles and usual error statistics. However, it offers a very fast and integrated vision of the strengths and weaknesses of a model with very little calculation. This methodology can also be deployed in inter-comparison exercises.

To answer the questions presented in the introduction, and for this particular model and simulated period, the following conclusions can be drawn. The model always simulates better the studied year than any other meteorological year and it is able

45 to reproduce the day to day variability for high concentrations of pollutants.

The spatial correlation is good for 2m temperature and precipitation rate, but not for wind speed: this highlights the fact that the modelled domain is large and the resolution not optimized for small scale processes. The spatial correlation is also very good for the long-range transport of particles as demonstrated with  $R_s=0.97$  and  $0.90$  for AOD and ANG. But, since this



**Figure 4.** Results of the  $I_{mv}$  scores for the spatial and temporal correlations. For each model variable its value is represented using the correlation on the x-axis and the difference between the studied year and the others on the y-axis. The colours represent the  $I_{mv}$  values.

feature occurs every year, this leads to low  $I_{mv}$  values. This means that for a large domain, the main spatial patterns of particle concentrations are recurrent and well modelled. The chemical species that are best modelled are either species with a long atmospheric lifetime ( $PM_{10}$ ) or species well spatially constrained on the domain (such as  $NO_2$  mainly due to anthropogenic emissions). For particles, the results depend on the size distribution: the largest particles are better simulated than the finest ones.

The conclusions are different for the temporal correlation. The scores are calculated using daily observations and modelled outputs. Thus, these scores reflect the ability of the model to retrieve the day to day variability. As for the spatial correlation, scores are good for the meteorological variables. For the aerosol, and mainly for the long-lived species (such as mineral dust), the temporal correlation is also correct as the  $I_{mv}$  values:  $I_{mv}=0.33$  and  $0.49$  for AOD and ANG respectively. But for the short-live species the temporal correlation and the  $I_{mv}$  values are low. This means that improvements are required in priority for the day to day variability compared to the locations of emissions. This may probably be due to the atmospheric transport, the spatial variability of 10m wind speed being poorly simulated. But, on overall, the temporal correlation is better for the studied year than for the others, showing that the problem is highly variable from year to year, but the model is significantly able to catch the evolution of the atmospheric composition.

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## **7 Code and/or data availability**

- 10 This study presenting a methodology using existing data and models, all required information are already included in this article.

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