

1 Dear Dr. Nicholas Henry Savage,

2 We appreciate valuable comments, which have helped improve the paper. We revised the text
3 according to the suggested corrections and would like to thank you for the thorough reading of the
4 paper. Below we provide our point-by-point replies, where for clarity the comments are displayed in
5 bold italics.

6 ***Title. As per the instruction of GMD, please include the version of WRF-Chem in the title.***

7 Thank you for this reminder, model version is now included in the title: »Evaluation of the high
8 resolution WRF-Chem (v3.4.1) air quality forecast and its comparison with statistical ozone
9 predictions«.

10 ***Abstract. Please specify the resolution of the model configuration applied (high resolution
11 is rather a relative term)***

12 We included the information about model resolution in the first sentence of the abstract, which is
13 now: »An integrated modelling system based on the regional on-line coupled meteorology-
14 atmospheric chemistry WRF-Chem model configured with two nested domains with horizontal
15 resolution 11.1 km and 3.7 km has been applied for numerical weather prediction and for air quality
16 forecast in Slovenia.«

17 ***2.1 WRF-Chem forecast system. Please state the height of the model top.***

18 The height of the model top is 50 hPa, this information is now included in the paper in the following
19 sentence: » The vertical structure of the atmosphere is resolved with 42 vertical levels extending up
20 to 50 hPa, with the highest resolution of ~25 m near the ground.«

21 ***Please provide a reference (even if it is only a report) for the emissions inventory.***

22 We added the reference to the project presentation at Slovenian Environment Agency (report is not
23 yet available).

24 ***2.2 Statistical ozone daily maximum forecast. Please provide references for the statistical
25 model.***

26 We added the reference to the final report about statistical model (also available online).

27 ***2.3 Evaluation methodology. What is the height of the lowest model level, and how does
28 that compare to a typical inlet height?***

29 We added this information to the paper the following way: »In the case of air pollutants, the
30 instantaneous lowest model level mixing ratios (with grid point center about 12 m above model
31 orography - an exception is KRV station as explained below) are compared to the hourly averaged
32 concentrations measured at monitoring stations (which have a typical inlet height of 3 m) from the
33 national network and some other environmental information systems in Slovenia. Figure 3 shows
34 locations of these AQ monitoring stations, and Tab. 1 lists the basic characteristics, including
35 comparison of the station altitude, the height of model orography, model analysis height, and
36 pollutants with higher than 75% availability of valid data during the analyzed time period for each of
37 the AQ monitoring site«

1 ***Have you considered using data from above level 1 - in very mountainous terrain, an***
2 ***observation site can be well above the model orography at the relevant grid point and it is***
3 ***more appropriate to use data from level 2 or above.***

4 Thank you for this question. In the case of AQ variables we usually use results from a higher model
5 level for the KRV station. The altitude of this station is well above model topography (model height:
6 1272 m, model grid point at the lowest level: 1284 m, station altitude: 1740 m). In the present paper
7 we originally included results for all stations (also KRV) at the lowest model level, because the
8 correlation coefficient at the lowest model level is highest (CORR decreases with increasing the
9 model level), showing that in spite of the negative bias due to too low model topography, the near
10 surface processes still play an important role in ozone dynamics. In the review process we
11 reconsidered this and decided to use model data from the 5th model level for KRV (model grid point
12 center: 1414 m), but stay with the lowest model level for all other stations. For KRV the 5th model
13 level is still well below the station altitude, but this reduces the bias for KRV from $-12 \mu\text{g m}^{-3}$ to -2
14 $\mu\text{g m}^{-3}$ for ozone hourly values, and from $-16 \mu\text{g m}^{-3}$ to $-7 \mu\text{g m}^{-3}$ for ozone daily maxima (which lowers
15 the impact of KRV bias on overall model performance). Unfortunately also CORR then decreases from
16 0.76 to 0.74 for ozone daily maxima (which has a negligible impact on overall model performance).
17 For other stations the differences between model height and station altitude are smaller. Also for
18 some of the stations model height is too low (e.g. VNA, model height: 468 m, station altitude: 630
19 m), but for other stations the model height is too high (e.g. HRA, model height: 540 m, station
20 altitude: 290 m), related to very complex topography in sub-alpine region of Slovenia. Consequently,
21 by increasing the model levels we could reduce the negative bias for stations of the first group (with
22 too low model orography), but cannot decrease the positive bias for the stations of the second group
23 with too high model orography. This makes an approach of using higher model levels for stations
24 with too low model orography questionable, also in the light that also CORR decreases with
25 increasing model levels. We thus support the approach of using the data on the lowest model level
26 and make a posterior bias correction, which does not impact the ozone dynamics and can be applied
27 for all stations. We only made an exception for KRV station, for which the height in the model was
28 significantly underestimated, as well as the station is known to be influenced by the conditions of the
29 free troposphere (except during hot summer daytime conditions), which is not the case for other
30 stations.

31 For meteorological variables we did not explore the impact of using results from higher model levels.
32 This would be far beyond the scope of this study, focused on ozone prediction, also because the
33 impact of using the higher layer data depends on meteorological variable, as well as the set of
34 meteorological stations is not the same as in the case of AQ stations.

35 In the paper due to using results for KRV on the 5th level we corrected all of the AQ statistics and
36 also the text throughout the paper accordingly. We included the following text:

37 »In the case of the elevated alpine KRV station, AQ variables are evaluated for the 5th model layer
38 instead of the first model layer. We made this exception for KRV, since the height of the model
39 topography was significantly underestimated there (Tab. 1), as well as the station is known to be
40 strongly influenced by the conditions of the free troposphere. The selection of the 5th model layer for
41 KRV station is based on analyses performed for different model layers (results not shown) and was
42 found to reduce the negative bias for O₃ due to too low WRF-Chem topography at this location.

1 Although even for this model layer the location of the grid point representing KRV station (1414 m) is
2 still well below the true station altitude (1740 m), the O₃ bias for KRV station is significantly smaller
3 than for the first layer, while the correlation coefficient between the measured and simulated O₃
4 levels remains similar in both cases (the 5th or the lowest model layer). Taking results from higher
5 model layers would further decrease the negative model bias, but would also worsen the correlation
6 coefficient for O₃ at this station due to decreased impact of surface processes.«

7 Later in text also:

8 Instead of: »The elevated alpine KRV station is the only one with negative bias (-12 µgm⁻³) in
9 forecasted 1-hour O₃ concentrations, which can be explained by the too low altitude of the KRV
10 station in model topography, since the mean O₃ concentration increases with height.«

11 We added: » In Fig. 4a the elevated alpine KRV station is the only one with high negative bias (-12
12 µgm⁻³) in forecasted 1-hour O₃ concentrations at the lowest model layer, which can be explained by
13 the too low altitude of the KRV station in model topography. The high negative bias for hourly O₃
14 concentrations at KRV station is reduced to a value of only -2 µgm⁻³ by using the 5th model layer
15 concentrations as explained in chapter 2.3. The 5th model level predictions will be used for KRV in all
16 analyses that follow.

17 We added also: » For sites with highest positive bias in 1-hour O₃ concentrations (TRB, ZAG, HRA and
18 ISK, with bias of 36 µgm⁻³, 31 µgm⁻³, 26 µgm⁻³ and 32 µgm⁻³, respectively), this can also be partly
19 explained by too high altitude of the stations in model orography (Tab. 1), since the mean O₃
20 concentration increases with height.«

21 Later in text we deleted: »or Alpine stations (KRV)« and added: » Here we recall that high negative
22 bias in WRF-Chem forecast for alpine KRV site due to too low altitude of the station in model
23 topography was compensated by taking prediction from the 5th model level.«

24 Also the values of statistics in text and figures are changed throughout the paper.

25 ***3.1 Evaluation of meteorological variables. There is a large decrease in the precipitation***
26 ***bias from day 1 to day 2 - is this a model spin up issue? If so would a different initialisation***
27 ***improve this error?***

28 We agree. Additional circumstance here is also that in the 3.4.1 model version it was not possible to
29 include the information about hydrometeors at the boundaries of the nested domain (in the applied
30 1-way nesting procedure). Since the intensity of (relatively rare) summertime precipitation events
31 was expected to have a less significant impact on ozone concentrations, we considered this issue less
32 problematic (in our study focused on ozone). We added the following text: "It must also be taken
33 into account that the 3.4.1 model version does not allow to include the information about
34 hydrometeors at the boundaries of the nested domain (in the applied 1-way nesting procedure),
35 which contributes to the negative simulated bias of precipitation. A large decrease in the
36 precipitation bias from day 1 to day 2 suggests that different initialization methodology (e.g. using 1
37 day spin-up for meteorology) could improve the prediction of precipitation events."

38 ***Please provide some evidence for the statement "the main precipitation events were well***
39 ***predicted and simulated" or remove this statement.***

1 Although we performed analyses and produced some plots we think that including additional
2 material here is beyond the scope of the paper. We thus decided to remove this statement.

3 **3.3 Evaluation and comparison of different methods for O₃ daily maximum predictions.**
4 **Please correct the statement "ideal forecast would lie in the right-bottom corner". It fact**
5 **the ideal model would have correlation coefficient of 1 and a standard deviation equal to**
6 **the observations, i.e. it would be co-located with the black dot which indicates the model.**
7 **The black dot is not always in the bottom right corner on these plots.**

8 Thank you, we corrected this statement. The statement that is now included is: » The ideal model
9 would have a correlation coefficient of 1 and a standard deviation equal to the observations, which
10 means that it would be co-located with the black dot on the diagram. «

11 **In the section on the evaluation of the model's ability to predict episodes, too much weight**
12 **is given to accuracy. For example, the statement "Accuracy ... increases with threshold**
13 **level" is misleading. A model which always forecasts "no event" will have an increasing**
14 **accuracy as the number of events decreases. To compare skill at different thresholds you**
15 **need to use a differnt metric e.g. Critical Success Index or Equitable Threat Score. These**
16 **would be better choices in general than accuracy in this section. There is no harm in**
17 **including accuracy in the tables, but it should not be the primary criterion for judging**
18 **forecast skill.**

19 In the revised paper we replaced Accuracy (A) measure by Equitable Thread score (ETS), we also
20 changed the order of categorical statistics in Tab. 5, so that ETS is shown in the first column, followed
21 by CSI, B, FAR and POD. We corrected the text, to give most weight to the ETS and briefly mention
22 the rest of them. The text that we now have in the paper regarding the categorical evaluations is the
23 following: »Equitable Threat Score (ETS) measures the fraction of observed and/or correctly
24 predicted events, adjusted for the frequency of hits that would be expected to occur by random
25 chance. Although this score takes into account the climatology it is not truly equitable. It ranges from
26 -1/3 to 1, where the minimum value depends on climatology (it is near 0 for rare events). Looking at
27 Tab. 5 ETS shows equal skill for WRF-Chem and statistical forecast, higher than persistence for the
28 120 $\mu\text{g m}^{-3}$ threshold (1-day and 2-day forecast). ETS decreases with increasing the threshold for both
29 WRF-Chem and statistical forecast, indicating the challenge that both models have to accurately
30 predict the extremes. In the case of 140 $\mu\text{g m}^{-3}$ threshold, WRF-Chem has the same ETS as
31 persistence, higher than the statistical model for 1-day forecast, while for 2-day forecast WRF-Chem
32 outperforms the statistical model, followed by persistence. In the case of 160 $\mu\text{g m}^{-3}$ threshold
33 persistence has the highest ETS for a 1-day forecast, followed by statistical model and WRF-Chem,
34 while in the case of 2-day predictions, statistical model shows the highest skill and WRF-Chem the
35 lowest. Another measure, the critical success index (CSI), is similar to ETS, except that it does not
36 take into account the climatology of the events and thus gives poorer scores for rarer events. It
37 measures the percentage of cases that are correctly forecasted out of those either forecasted or
38 observed, and ranges from 0 to 1 (1 indicating the perfect forecast). Similar as ETS, CSI gives higher
39 scores for persistence in the case of 1-day forecast for the higher two thresholds, while on the
40 second day WRF-Chem or the statistical model already performs better. Bias (B) determines whether
41 the same fraction of events are both forecasted and observed. A tendency of the statistical model
42 and of WRF-Chem to under-predict O₃ threshold exceedances shows as a B below 1 for these two
43 models. The false alarm ratio (FAR) that measures the percentage of forecast high O₃ events that

1 turn out to be false alarms, gives highest skill for WRF-Chem, followed by statistical model and
2 persistence. The probability of detection (POD) is a measure of how often a high threshold
3 occurrence is actually predicted to occur, and is relatively low for WRF-Chem with respect to other
4 models. «

5 ***Also why were these specific three thresholds chosen?***

6 There was no specific reason for these certain three thresholds. We also performed the calculations
7 for different thresholds, e.g. 130 $\mu\text{g m}^{-3}$ or 150 $\mu\text{g m}^{-3}$, distinguishing between higher and lower ozone
8 maxima, and the conclusions were similar. We included some thresholds which present an elevated
9 ozone levels and pose a greater risk to human health, and decided to exclude the statistics for a
10 higher threshold (180 $\mu\text{g m}^{-3}$, a legislation limit value) due to a very low number of exceedances for
11 this threshold. In the paper we extended the following sentence: »Table 5 summarizes the
12 categorical evaluation results for three different thresholds (120, 140, 160 $\mu\text{g m}^{-3}$) of elevated ozone
13 levels, which pose a greater risk to human health.«

14 ***Grammatical and other minor corrections.***

15 ***p1030 line 22, "The first RT-AQF systems.."***

16 ***p1030 line 25, delete "existing"***

17 ***p1032 line 13, "during summertime conditions"***

18 ***p1032 line 21, "a one-way"***

19 ***p1032 line 22, "evaluated a forecast"***

20 ***p1033 line 2, "based on WRF-Chem are implemented worldwide"***

21 ***p1033 line 4, "over the topographically complex"***

22 ***p1033 line 6, "with a statistical model"***

23 ***p1033 line 6, "at the Slovenian"***

24 ***p1036 line 19, "a southwestern"***

25 ***p1036 line 24, "shows a mean O3 daily mean"***

26 ***p1037 line 27, "is a mountainous station"***

27 ***p1037 line 27, "As well as the elevated station KRV, the ISK, OTL and VNA stations
28 area are also influenced by regional transport of pollutants."***

29 ***p1038 line 7, "information about the AQ forecast can also be gained by the evaluation
30 of meteorological forecasts for these stations."***

31 ***p1038 line 16, "index of agreement"***

32 ***p1041 line 3, "with a range of 0.64 to 0.90 for 1 day forecasts"***

33 ***p1041 line 7, "On average"***

34 ***p1042 line 8, "3 month accumulations by"***

35 ***p1042 line 3, "has problems simulating the"***

36 ***p1043 line 1, "the model over-predicts"***

37 ***p1043 line 5, "explained by model error in"***

38 ***p1043, line 16, "poorly reproduced meteorological"***

39 ***p1043, line 26, "Also interesting to discuss are the results"***

40 ***p1045, line 3, "In this section we want to answer the question: 'how accurate is the
41 1 h O3 daily maximum WRF-Chem forecast in comparison to the statistical model
42 prediction or to persistence?'"***

1 *p1045, line 8 "which is, along with their computational efficiency, "*
2 *p1045, line 9 "Among the strengths of the deterministic models are that they give"*
3 *p1045 line 12, "Furthermore, they also allow forecasts for"*
4 *p1045 line 14, "descriptions of"*
5 *p1045 line 27, "because a statistical"*
6 *p1046 line 1, "with an available"*
7 *p1046 line 5, "already beats persistence"*
8 *p1046 line 12, "than the statistical forecast"*
9 *p1046 line 25, "MNBE in Fig. 8 has very similar results to ME."*
10 *p1047 line 13, "also contingency-table-based statistics are an important metric of"*
11 *p1047 line 15, "It is important to take into account"*
12 *p1048 line 9, "were to be applied to"*
13 *p1049 line 7, "local emissions result in model underestimations of NO2"*
14 *p1049 line 12, "show good WRF-Chem model performance"*
15

16 We revised the text according to the suggested corrections and would like to thank again for the
17 thorough reading of the paper.

18

19

1 Dear Dr. Georg A. Grell,

2 We appreciate and would like to thank you for all the comments and raised questions, which have
3 helped to improve the quality of the paper. Below we provide our point-by-point replies, where for
4 clarity the comments are displayed in bold italics.

5 ***This paper describes the use of the community version of WRF-Chem for real-time ozone
6 and aerosol predictions. The authors perform statistical evaluations over a 3 month period,
7 comparing the model forecasts with observations as well as statistical forecast methods. In
8 general his paper is well written and should be published in GMD. This can be done with
9 only minor modifications.***

10 We thank for this comment.

11 ***Although the authors provide much information on model set-up there are a few details
12 that I was looking for and couldn't find. Is this 2-way nesting or 1-way nesting? If it is 1-
13 way nesting, how was it applied?***

14 It is a 1-way nesting applied by two consecutive simulations (using ndown). We added this
15 information the following way (section 2.1): »A 1-way nesting is applied by two separate consecutive
16 simulations, where outputs from the coarse grid integration are processed to provide boundary
17 conditions for the nested run every 15 minutes.«

18 ***Is the choice of physics parameterization the same on both domains?***

19 Yes, schemes are the same on both domains. To include this information in the paper we changed in
20 Section 2.1: »We decided to apply the same schemes as were used...« to »In both domains we
21 decided to apply the same schemes as were used...«.

22 ***Which photolysis model have you been using?***

23 Fast-J photolysis scheme (Fast et al., 2006), this information is now added in section 2.1.

24 ***All evaluations I am assuming are done on the high resolution domain.***

25 Yes. We included this information in the first sentence of Section 2.3: »We evaluate the 1-day and 2-
26 day WRF-Chem meteorological and AQ forecasts on the high resolution domain during a 3-month
27 period (June - August 2013).«

28 ***Also, the color choice for figures 5, 6, and 7 is unfortunate. The two blue colors are almost
29 impossible to separate – at least with my aging eyes. Why not a different color? Figure 5 is
30 even more difficult to read, a bit too small for me.***

31 We replotted these figures with two different colors. Still it is hard to distinguish between 1-day and
32 2-day forecast (Fig. 5-7), but the purpose of these figures is more to separate model forecast from
33 observations. 1-day and 2-day forecast are more easily distinguished by the use of statistics. Figure 5
34 is now divided into two parts.

35 ***Some other questions I have:***

36 ***(1) There is a negative temperature bias, but a positive short wave bias? Since you are
37 using the interaction flag for convection/radiation the SW bias could be interpreted as not
38 enough cloud cover, which could give you a low bias at night, but at day? Are you cycling***

1 ***soil temperature and soil moisture or is that always a new initialization with coarse***
2 ***resolution GFS data?***

3 All meteorological variables, including soil temperature and soil moisture are always initialized with
4 GFS data, which is now mentioned in the paper. This explains higher negative bias for T2m during the
5 first day of simulation (not valid for daily maxima, where bias is the same on the first and the second
6 day of simulation). For all hourly values T2m bias decreases from -2.1 C to 0.8 C due to reduced bias
7 for nighttime temperatures on the second day of simulation. Looking at results station by station the
8 link between T2m and SW bias is not straightforward (they appear not to be directly correlated). On
9 the first day of simulation higher SW is due to less cloudy conditions (more cloud cover on the
10 second day).

11 ***(2) The statistics I assume are always over domain 2. The fact that the precipitation under-***
12 ***forecast is a lot less on day 2 may indicate some spin-up issues, especially also when taking***
13 ***into consideration the coarse initial conditions (did you use .5 degree data from GFS?)***

14 Yes, we used the 0.5 degree data from GFS, this information is now added in section 2.1 as »...with
15 meteorological initial (ICs) and lateral boundary conditions (BCs) taken from the 0.5° data from the
16 Global Forecast System (GFS)...«. We also agree that under-prediction of precipitation indicates some
17 spin-up problem, where it must also be taken into account that in 3.4.1 model version ndown
18 procedure does not allow to include the information about hydrometeors at boundaries of the
19 nested domain. Since the intensity of (rare) summertime precipitation events was expected to have a
20 less significant impact on ozone concentrations, we considered this issue less problematic in our
21 study focused on ozone. But we agree that applying a different initialization methodology should
22 reduce the precipitation error. The following text was added: »It must be mentioned that the 3.4.1
23 model version does not allow to include the information about hydrometeors at the boundaries of
24 the nested domain (in the applied 1-way nesting procedure), which contributes to the negative
25 simulated bias of precipitation. A large decrease in the precipitation bias from day 1 to day 2
26 suggests that different initialization methodology (e.g. using 1 day spin-up for meteorology) could
27 improve the prediction of precipitation events.«

28 ***(3) On page 1047, line 22 you talk about WRF-Chem under-predicting Ozone maxima, while***
29 ***before you had a positive bias. Do you mean under-predict exceedances?***

30 We replaced »ozone maxima« to »threshold exceedances«.

31 ***(4) In the summary and conclusions you should mention again (you have that hidden***
32 ***somewhere in section 2.1, pg 1034) that different choice of physical or chemical***
33 ***parameterization will influence and possibly change outcomes. However I think your***
34 ***choices are good choices, since they are well documented in other real-time applications.***

35 We added the following sentence to the conclusions: »Since the selection of physical or chemical
36 parameterization schemes influences and possibly changes the outcomes, we decided to apply the
37 schemes that are well documented and have previously been used in other applications (e.g.
38 AQMEII).«

39 ***(5) Pg. 1031, line 7: The MM5 reference should be 1994, not 1995 – if I remember correctly***

40 This error is now corrected.

1 **(6) Pg 1032, line 11: 2011 should not be a reference for WRF-Chem. Just 2005 is good**
2 **enough.**

3 We deleted the 2011 reference.

4 **(7) Pg. 1049, last line: If you want you could add the recent Pagowski et al publication in**
5 **GMD (also WRF-Chem special issue) as an example of chemical data assimilation**

6 The following sentence was added: »For WRF-Chem model a technical note on the implementation
7 of the aerosol assimilation and a guidance for prospective users has been recently published by
8 Pagowski et al. (2014).«

9

10

Evaluation of the high resolution WRF-Chem (v3.4.1) air quality forecast and its comparison with statistical ozone predictions

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Abstract

An integrated ~~high-resolution~~ modelling system based on the regional on-line coupled meteorology-atmospheric chemistry WRF-Chem model configured with two nested domains with horizontal resolution 11.1 km and 3.7 km has been applied for numerical weather prediction and for air quality forecast in Slovenia. In the study an evaluation of the air quality forecasting system has been performed for summer 2013. In the case of ozone (O₃) daily maxima, the first ~~day~~ and second day model predictions have been also compared to the operational statistical O₃ forecast and to the persistence. Results of discrete and categorical evaluations show that the WRF-Chem based forecasting system is able to produce reliable forecasts, which depending on monitoring site and the evaluation measure applied can outperform the statistical model. For example, the correlation coefficient shows the highest skill for WRF-Chem model O₃ predictions, confirming the significance of the non-linear processes taken into account in an on-line coupled Eulerian model. For some stations and areas biases were relatively high due to highly complex terrain and unresolved local meteorological and emission dynamics, which contributed to somewhat lower WRF-Chem

1 skill obtained in categorical model evaluations. Applying a bias-correction could further
2 improve WRF-Chem model forecasting skill in these cases.

3 **Key words:** Air quality, forecast, ozone, WRF-Chem, online-coupled model, statistical model

4

5 **1 Introduction**

6 Real-time air quality forecasting (RT-AQF) is a relatively new discipline in atmospheric
7 sciences, which has evolved as a response to societal and economic needs, reflecting the
8 progress in scientific understanding of physical processes and numerical and computational
9 technologies (Zhang et al., 2012a). ~~The F~~first RT-AQF systems, developed for forecasting air
10 pollution in exposed urban regions, were either empirical methods based on persistence,
11 climatology, human expertise and meteorological forecast (e.g. Wolff and Liroy, 1978), or
12 statistical models taking advantage of ~~existing~~ links between pollutant concentrations,
13 meteorological variables (wind speed and direction, temperature, cloudiness, moisture etc.)
14 and physical (emissions) parameters (e.g. McCollister and Wilson, 1975; Cobourn, 2007;
15 Vlachogianni et al., 2011). The next step in evolution of RT-AQF systems was the use of
16 sophisticated chemical transport models that represent all major processes (meteorological
17 and chemical) that lead to the formation and accumulation of air pollutants. Many of these
18 RT-AQF systems consist of an offline coupled meteorological model and a chemical-
19 transport model, where the meteorological model (e.g., ALADIN, ALADIN International
20 Team, 1997; MM5, Grell et al., 1994~~5~~; WRF, Skamarock et al., 2008) provides
21 meteorological input for the chemical-transport model (e.g., EMEP, van Loon et al., 2004;
22 CMAQ, Byun and Schere, 2006; CAMx, ENVIRON, 2011; CHIMERE, Menut et al., 2013)
23 with an output time interval typically around 1 hour. Examples are the EURAD
24 (http://db.eurad.uni-koeln.de/index_e.html), SILAM (<http://silam.fmi.fi/>), ForeChem
25 (<http://atmoforum.aquila.infn.it/forechem/>), CALIOPE (<http://www.bsc.es/caliope/>) forecast
26 systems and others. The new generation of an online coupled models (e.g., MCCM, Grell et
27 al., 2000; GATOR-GCMM, Jacobson 2001; Meso-NH-C, Tulet et al. 2003; WRF-Chem,
28 Grell et al., 2005; Enviro-HIRLAM, Baklanov et al., 2008; GEM-AQ, Kaminski et al. 2008;
29 COSMO-ART, Vogel et al., 2009; WRF-Chem-MADRID, Zhang et al., 2010a) presents an
30 alternative approach with one unified modelling system, in which meteorological and air
31 quality variables are simulated together within the same model. The online approach permits
32 the simulation of two-way interactions between different atmospheric processes including

1 emissions, chemistry, clouds and radiation, and a better response of the simulated pollutant
2 transport to changes of the wind field (Grell et al., 2004), and can thus provide a more
3 realistic representation of the atmosphere. The use of online coupled models can be
4 particularly important in regions with high aerosol loadings and cloud coverage (Otte et al.,
5 2005; Eder et al., 2006), where physical processes in the atmosphere may be modified by the
6 aerosol direct effect on radiation or by aerosol cloud interactions. Several reviews
7 summarized the strengths and limitations of offline and online coupled models (e.g. Zhang
8 2008; Klein, 2012; Baklanov et al., 2014). There is an increasing awareness that an integrated
9 online approach is needed not only for assessment, forecasting and communication of air
10 quality, but also for weather forecasting (e.g. Baklanov, 2010; Grell and Baklanov, 2011;
11 Klein et al., 2012; Zhang et al., 2012b; Baklanov et al., 2014). Nevertheless, there are several
12 issues regarding the inclusion of chemistry into numerical weather prediction models. More
13 evidence is required whether an integrated model can produce a good climatology of the most
14 important chemical species, and if such a model is, considering many uncertainties, able to
15 beat persistence forecasts of these species (Grell and Baklanov, 2011). These questions are
16 calling for further research and studies exploring the performance of the models with an
17 online coupled chemistry.

18 In recent years extensive efforts have been devoted to develop air quality (AQ) forecasting
19 systems for Slovenia. In this study we explore the use of the state-of-the-science WRF-Chem
20 model (Grell et al., 2005, ~~2011~~) with coupled meteorological, microphysical, chemical, and
21 radiative processes for forecasting AQ in Slovenia during ~~the~~ summertime conditions. In last
22 decade WRF-Chem has been increasingly applied to many areas worldwide (e.g., Misenis and
23 Zhang, 2010; Fast et al., 2009; Zhang et al., 2010a, 2010b; Li et al., 2011; Tie et al., 2009; Hu
24 et al., 2012; Forkel et al., 2012, Žabkar et al., 2011, [a](#), 2013). In most of these studies WRF-
25 Chem model has been successfully used to simulate historical poor AQ conditions in hindcast
26 approach. To our knowledge, only a few studies focused on using WRF-Chem for forecasting
27 AQ, most of these have applied WRF-Chem forecast before and during field campaigns
28 (McKeen et al., 2005, 2007, 2009; Yang et al., 2011). Takigawa et al. (2007) evaluated O₃
29 forecast for a 1 month time period from ~~a~~ one-way nested global-regional RT-AQF system
30 with full chemistry based on the global CHASER (Sudo et al. 2002) and regional WRF-Chem
31 models, while Saide et al. (2011) evaluated [a](#) forecast system based on ~~the~~ WRF-Chem model
32 for simulating carbon monoxide (CO) as a PM₁₀/PM_{2.5} surrogate over Santiago de Chile for
33 wintertime conditions. WRF-Chem-MADRID (Zhang et al., 2010a) with two additional gas-

1 phase mechanisms, sectional representation for particle size distribution and more advanced
2 model treatments compared to WRF-Chem, was applied by Chuang et al. (2011) and by
3 Yahya et al. (2014) for forecasting AQ over the Southeastern U.S.. In spite of a limited
4 number of evaluation studies published in the literature, an increasing number of real-time
5 weather and air quality forecasting systems based on WRF-Chem ~~is performed~~
6 are implemented worldwide (http://ruc.noaa.gov/wrf/WG11/Real_time_forecasts.htm).

7 In our study we explore the forecasting skill of WRF-Chem model over the topographically
8 complex and geographically diverse area of Slovenia for three summer months (June - August
9 2013). Furthermore, in the case of O₃ we compare WRF-Chem predictions with a statistical
10 model for predicting O₃ daily maxima, currently used at the Slovenian Environment Agency
11 (SEA). Both first day (1-day) and second day (2-day) forecasts are considered, while a
12 persistence model, which assumes that pollutant level today and tomorrow will be the same as
13 yesterday, is used as a threshold for useful model prediction. Since the availability of accurate
14 and reliable forecasting system could be useful to the local authorities and could help to
15 advise the public the proper preventive actions, we want to answer the question whether
16 WRF-Chem model outperforms the statistical model or persistence. Namely, considering
17 many uncertainties related to one unified model, it may not be easy for models with online
18 chemistry to be able to perform well enough to meet the required standards, and more
19 research and studies are needed to investigate that (Grell and Baklanov, 2011). Due to the
20 limited number of previous studies focused on online coupled forecasting systems, the aim of
21 our study is also to provide a greater insight into potential that lies in the approach based on
22 an unified model for forecasting weather and air pollution. Finally, identified strengths,
23 limitations and deficiencies of analyzed RT-AQFs, are expected to present the basis for
24 further research.

25 **2 Methodology**

26 **2.1 WRF-Chem forecast system**

27 The RT-AQF system for Slovenia based on the WRF-Chem model version 3.4.1 is configured
28 with two nested domains (Fig.1) with horizontal resolution 11.1 km and 3.7 km, and 151×100
29 and 181×145 grid points, respectively. A 1-way nesting is applied by two separate
30 consecutive simulations, where outputs from the coarse grid integration are processed to
31 provide boundary conditions for the nested run every 15 minutes. The vertical structure of the

1 | atmosphere is resolved with 42 vertical levels extending up to 50 hPa, with the highest-near
2 | ground resolution of ~25 m near the ground. About 15 levels are located within the lowest 2
3 | km to assure high vertical resolution of the daytime planetary boundary layer (PBL). To
4 | produce the 48-hour forecast, the model is run every day, starting at 00 UTC, with
5 | meteorological initial (ICs) and lateral boundary conditions (BCs) taken from the 0.5° data
6 | from the Global Forecast System (GFS), ~~a global numerical weather prediction system~~
7 | operated by the US National Weather Service (NWS). For chemical BCs forecasts from
8 | global MOZART-4/ GEOS-5 (Emmons et al., 2010) RT-AQF system with temporal
9 | availability of 6 h are used. The instantaneous outputs at the 24th hour of the previous day
10 | forecast are used to initialize next day's forecasting simulation. An exception is the very first
11 | day of the first 48-hour forecasting cycle, when global MOZART-4/ GEOS-5 fields were used
12 | also to initialize chemistry. A three day spin-up ahead of the first analyzed forecast day is
13 | then taken into account to allow pollutants to accumulate in the air masses.

14 | In the WRF-Chem model, several choices for parameterizations of physical and chemical
15 | processes are available (Grell et al., 2005; Skamarock et al., 2008; Peckham et al., 2011), and
16 | their choice can have a strong impact on the model predictions. In both domains wWe decided
17 | to apply the same schemes as were used in simulation SII for Phase-2 of the Air Quality
18 | Model Evaluation International Initiative (AQMEII) (e.g., Balzarini et al., 2014, Baró et al.,
19 | Curci et al., 2014, Forkel et al., 2014, Im et al., 2014a and 2014b, Kong et al., 2014, 2014,
20 | San José et al., 2014). These include Yonsei University (YSU) PBL scheme (Hong et al.,
21 | 2006), NOAH land-surface model (Chen and Dudhia, 2001), Rapid Radiative Transfer
22 | Method for Global (RRTMG) long-wave and short-wave radiation scheme (Iacono et al.
23 | 2008), Grell 3D ensemble cumulus parameterization scheme (Grell and Devenyi, 2002) with
24 | radiative feedback, Morrison double-moment cloud microphysics (Morrison et al., 2008),
25 | Fast-J photolysis scheme (Fast et al., 2006), RADM2 gas phase chemistry (Stockwell et al.,
26 | 1990) and the MADE/SORGAM aerosol module (Ackermann et al., 1998, Schell et al.,
27 | 2001). Current model implementation includes a modified RADM2 gas phase chemistry
28 | solver as described in Forkel et al. (2014), which avoids under-representation of nocturnal O₃
29 | titration in areas with high NO emissions. According to Forkel et al. (2014) the modified
30 | solver tends to over-estimate the low NO₂ concentration for pristine regions and in the free
31 | troposphere, which results in an overestimation of O₃. Due to the focus on polluted regions
32 | this deficiency was considered as less important than the advantage of better description of
33 | the titration. In addition, the comparatively small modelling domain (D1) ensures that the

1 boundary conditions constrain the high bias of the modified solver for O₃ and NO₂ in the free
2 troposphere. Also according to our sensitivity tests (results not shown) the modified solver
3 showed better performance for O₃ daily maxima and O₃ nighttime minima than the QSSA
4 RADM2 solver supplied originally with WRF-Chem model.

5 Among feedbacks only the aerosol direct effects on radiation according to Fast et al. (2006)
6 and Chapman et al. (2009) are taken into account. As shown by Kong et al. (2014) for two air
7 pollution episodes, this degree of aerosol-meteorology interactions in 3.4.1 version of the
8 WRF-Chem improved model performance for high aerosol loads, while the representation of
9 the indirect effects needs to be further improved to be able to outperform simulations with
10 direct effects only.

11 Biogenic emissions are estimated using MEGAN (Model of Emissions of Gases and Aerosols
12 from Nature; Guenther et al., 2006) online model calculations, while dust emissions are
13 modelled according to Shaw et al. (2008) with an adjustment to avoid high dust fluxes from
14 some Dalmatian islands in Croatia. A detailed anthropogenic inventory for pollutants CO,
15 NH₃, NO_x, SO₂, and NMVOC, which has been for the purpose of AQ forecasting constructed
16 for year 2009 by SEA (SEA, 2014), is used to estimate anthropogenic emissions in Slovenia.
17 For areas outside Slovenia the recently updated anthropogenic emissions for the year 2009
18 based on the TNO-MACC-II (Netherlands Organization for Applied Scientific Research,
19 Monitoring Atmospheric Composition and Climate – Interim Implementation), the same as
20 prepared for phase-2 of the AQMEII exercise (Pouliot et al., 2014), are being used. Daily
21 updates of the WRF-Chem based experimental AQ forecast are provided at
22 <http://meteo.fmf.uni-lj.si/onesnazenje>.

23 **2.2 Statistical ozone daily maximum forecast**

24 The statistical O₃ model (Žabkar, 2011b), currently used at SEA for forecasting O₃ daily
25 maxima at 8 measuring sites in Slovenia (Fig.3), is a multivariate regression tool combined
26 with clustering algorithms to take into account measured data, weather forecast data, as well
27 as the predicted backward trajectories of each monitoring site. As regards measurements,
28 yesterday (at 12, 15, 18 and 21 local time, daily maximum, daily minimum, daily average)
29 and today early morning (7 local time) meteorological (pressure, relative humidity, direct and
30 diffusive solar radiation, wind speed) and AQ data (O₃, NO_x, NO₂, CO, PM₁₀, SO₂) are used.
31 For meteorological predictions the 24-h ECMWF forecast variables at 12 UTC of the forecast

1 day at different vertical levels (1000 hPa, 925 hPa, 850 hPa, 500 hPa, 300 hPa) above the
2 measuring sites are taken into account. Among all these variables by the use of stepwise
3 technique, based on the F-statistic only significant variables were selected to be included in
4 multivariate regression equations for different monitoring sites (from 15 to 26 variables,
5 depending on monitoring site).

6 The important part of the statistical forecast is calculation of 24-h backward trajectories on
7 meteorological fields of ALADIN/SI forecast. The inclusion of 24-h predicted trajectories
8 into statistical model is based on the study (Žabkar et al., 2008) which showed, that the
9 highest O₃ daily maxima at monitoring sites in Slovenia are in general associated with short
10 (slow-moving) backward trajectories with ~~at~~the southwestern origin, while the lowest
11 measured daily maximum O₃ values for all the stations are associated with the clusters of long
12 northwestern trajectories. Clusters of similar trajectories were for the purpose of statistical
13 forecast calculated by *k*-means clustering algorithms (Moody and Galloway, 1988; Žabkar et
14 al., 2008) on 6 years (2004-2010) of data (ALADIN/SI trajectories). As an example, Fig. 2
15 ~~shows a mean is showing mean~~ O₃ daily maxima for clusters of similar trajectories for one of
16 the monitoring sites. The same 6-year time period of training data was used in the stepwise
17 multiple regression procedure to determine the multiple regression prognostic equations
18 associated with monitoring sites and trajectory clusters, from measurements, ECMWF
19 forecast data, average cluster O₃ daily maximum, and day-of-the-year variable.

20 The first step of the statistical O₃ prediction is the calculation of trajectories approaching the
21 monitoring stations at 12 UTC of the forecast day. In the next step these backward trajectories
22 of each monitoring site are associated to the nearest pre-calculated cluster of similar
23 trajectories. Finally, the multiple regression equation of the associated group of trajectories is
24 used to calculate the O₃ daily maximum prediction. It must also be noted, that the decision on
25 declaring O₃ episodes is only partially based on the results from this statistical model; it also
26 involves a decision made by AQ forecasters.

27 **2.3 Evaluation methodology**

28 We evaluate the 1-day and 2-day WRF-Chem meteorological and AQ forecasts on the high
29 resolution domain during a 3-month period (June - August 2013). The main focus is on O₃
30 predictions. In the case of air pollutants, the instantaneous lowest model level mixing ratios
31 (with grid point center about 12 m above model orography - an exception is KRV station as

1 explained below) are compared to the hourly averaged concentrations measured at monitoring
2 stations (which have a typical inlet height of 3 m) from the national network and some other
3 environmental information systems in Slovenia. Figure 3 shows locations of these AQ
4 monitoring stations, and Tab. 1 lists the basic characteristics, including comparison of the
5 station altitude, the height of model orography, model analysis height, and pollutants with
6 higher than 75% availability of valid data during the analyzed time period for each of the AQ
7 monitoring site. In the case of the elevated alpine KRV station, AQ variables are evaluated for
8 the 5th model layer instead of the first model layer. We made this exception for KRV, since
9 the height of the model topography was significantly underestimated there (Tab. 1), as well as
10 the station is known to be strongly influenced by the conditions of the free troposphere. The
11 selection of the 5th model layer for KRV station is based on analyses performed for different
12 model layers (results not shown) and was found to reduce the negative bias for O₃ due to too
13 low WRF-Chem topography at this location. Although even for this model layer the location
14 of the grid point representing KRV station (1414 m) is still well below the true station altitude
15 (1740 m), the O₃ bias for KRV station is significantly smaller than for the first layer, while
16 the correlation coefficient between the measured and simulated O₃ levels remains similar in
17 both cases (the 5th or the lowest model layer). Taking results from higher model layers would
18 further decrease the negative model bias, but would also worsen the correlation coefficient for
19 O₃ at this station due to decreased impact of surface processes.

20 All AQ stations are background, 7 of them are measuring urban background, 1 suburban and
21 9 rural conditions. Valid O₃ measurements are for the analyzed time period available for 13
22 AQ stations. When studying the general model performance, data from additional 4 stations
23 for two other pollutants (NO₂, PM10) are also analyzed to get a better picture of model
24 behavior over the domain, known for its large topographical and climate diversity. The
25 coverage of three climate zones in Slovenia (Mediterranean, sub-alpine and mountainous)
26 with monitoring stations is the following: NG, KOP and OTL are Mediterranean sites, KRV is
27 a mountainous station, and the remaining stations are sub-alpine. As well as the
28 elevated station KRV, the also-ISK, OTL and VNA stations are also influenced by measuring
29 regional transport of pollutants.

30 For evaluation of predicted meteorological variables, data from SEA meteorological stations
31 (MET, Fig. 3) for 2m temperature (T2m), 10 m wind speed (W10m), relative humidity (RH),
32 incoming shortwave radiation (SR) and precipitation (RR) are used. It must be noted, that

1 MET stations with lower spatial representativeness (e.g. alpine stations) were not a priori
2 excluded from the analyses, which needs to be taken into account when looking at evaluation
3 results. The reason for not excluding these stations was that some ~~interesting~~ information
4 ~~about the~~ AQ forecast can also be gained ~~also~~ by the evaluation of meteorological forecast
5 for these stations.

6 Basic statistical measures (correlation coefficient (CORR), mean error (ME), mean absolute
7 error (MAE) and root mean square error (RMSE)) are used for evaluating model's forecasting
8 skills of meteorological and AQ variables. In the case of O₃, correlation coefficients are
9 presented also by Taylor diagrams (Taylor, 2001), which graphically summarize the similarity
10 between model forecasts and observations not only in terms of their correlation, but also with
11 their centered root-mean-square difference and the amplitude of their variations, represented
12 by their standard deviations. Furthermore, some additional discrete statistical measures,
13 including index ~~of~~ agreement (IOA), the mean normalized bias error (MNBE), and the mean
14 normalized gross error (MNGE) are calculated for O₃ daily maximum concentrations
15 predicted by the different models. Finally, to evaluate the model's ability to predict
16 exceedances and non-exceedances also several categorical indices including Equitable Threat
17 Score accuracy (ETSA), Critical Success Index (CSI), ~~b~~Bias (B), ~~F~~false aAlarm rRatio (FAR)
18 ~~and, P~~probability Oof Ddetection (POD) ~~and critical success index (CSI)~~ are calculated for
19 different thresholds. Definitions of statistical measures are shown in Appendix A.

20 **2.4 Meteorology and air quality of June-August 2013**

21 The analyzed period was marked by three heat wave events, which contributed to the summer
22 characterized by high temperatures, sunny weather and lack of precipitation in Slovenia. The
23 first heat wave event with measured temperature daily maxima up to 35 °C occurred after a
24 rather cold beginning of the month and lasted from June 15 – 21. The event was terminated by
25 a cold front passage and followed by the pronounced cold episode during the end of June and
26 the beginning of July. Another heat wave event with temperatures above 35 °C observed in
27 the lowland, started on July 26 and was briefly interrupted on July 29, when thunderstorms
28 related to frontal passage were accompanied by exceptionally strong wind gusts. The most
29 remarkable of three extraordinary hot episodes was recorded from August 1 – 8. On the last
30 day of this episode, August 8, temperatures reached 40 °C at some measuring sites in
31 Slovenia, and many of them observed their highest temperature ever recorded.

1 As expected for summertime conditions, measured concentrations of most air pollutants,
2 including PM₁₀, were in general low during the analyzed time period. The only exception
3 was O₃ with exceedances of 8-hour target value (120 µg m⁻³) measured at all AQ monitoring
4 stations during the three heat wave events, which is the reason why the main focus of the
5 present study is on this pollutant. During the second two events (in July and August) also
6 threshold exceedances of 1-hour daily maxima were recorded for O₃. In spite of the hot and
7 sunny conditions during the first heat wave event in June 2013, measured daily O₃ maxima at
8 the Slovenian stations did not exceed the 1-hour information threshold value (1h ITV; 180
9 µg m⁻³), but reached 171 µg m⁻³ at the Mediterranean OTL and the elevated alpine KRV
10 stations. During the second heat wave event 1-hour daily maxima exceeded 180 µg m⁻³ at
11 KRV, OTL, NG and KP (July 23 – 28), while the highest number of 1-hour exceedances (20)
12 has been in July measured at OTL station. Similarly, during the August heat wave event O₃
13 concentrations exceeded the 1h ITV at LJ, MB, OTL, NG and KP from August 2 – 7. To
14 summarize, the Mediterranean stations (NG, OTL, KP) due to very high O₃ concentrations
15 measured during the heat wave events (especially the second two events) exhibited the
16 poorest AQ in Slovenia during the analyzed time period, while the legislation limit values
17 have been exceeded only occasionally for the sub-alpine stations.

18

19 **3 Results and discussion**

20 **3.1 Evaluation of meteorological variables**

21 Table 2 shows conventional statistical scores evaluating the 1-day WRF-Chem forecast for
22 the basic meteorological variables, 2m temperature (T2m; for hourly values and daily
23 maxima), 10 m wind speed (W10m), relative humidity (RH) and incoming solar radiation
24 (SR). Results for three selected measuring sites (LJ, NG, MS) and overall result for all 24
25 MET monitoring sites (shown in Fig. 3) are presented separately.

26 Incoming solar radiation is the main energy source that drives all atmospheric processes,
27 including PBL processes, and has a critical role also in atmospheric chemistry. For almost all
28 sites the mean SR was overestimated by the model, with an overall ME of 16 W/m² and 11
29 W/m² for 1-day and 2-day forecast, respectively. CORR was higher for 1-day (0.77) than for
30 2-day (0.71) forecast, with a range of 0.64 to 0.90 for 1-day forecasts at different stations. The

1 larger positive bias during the first day than for the second day can be attributed to less cloudy
2 conditions during the first day of simulation.

3
4 In the case of T2m 1-day (2-day) WRF-Chem meteorological forecast showed an overall
5 correlation with measurements of 0.93 (0.94) for all 1-hour values and 0.97 (0.96) for 1-hour
6 daily maxima. With an exception of three alpine stations with higher simulated positive bias,
7 daily T2m maxima were simulated with ME between -3.9 °C and -0.6 °C, depending on
8 station spatial representativeness. All meteorological variables, including soil temperature and
9 soil moisture, are always initialized with GFS data. This explains higher negative bias for
10 T2m during the first day of simulation in spite of the overestimated of solar radiation. A while
11 aAn average systematic underestimation of T2m daily maxima was -2.1 °C both for 1-day and
12 2-day forecast. Nighttime T2m minima showed lower systematic bias for 2-day forecast,
13 which resulted in overall bias for all hourly T2m values of -1.3 °C for 1-day and -0.8 °C for 2-
14 day forecast. Predominant weak wind conditions with variable direction at stations located in
15 complex topography were challenging to simulate. The general model tendency was to
16 overestimate W10m with overall ME of 0.8 m/s for 1-day and 2-day forecast, where for some
17 stations bias can be very low (e.g. LJ; Tab. 2) and much higher for some other stations due to
18 their local positioning in complex topography (e.g. HRA located in valley with ME of 1.9
19 m/s). For hourly values the correlation is lower (Tab. 2), but for mean daily W10m values
20 Pearson correlation coefficient between 0.4 and 0.9 has been simulated, depending on
21 monitoring site. Relative humidity shows slightly better results for 1-day than for 2-day
22 forecast with CORR of 0.77 and low overall ME of 2 % for 1-day forecast, which for
23 particular stations can be positive (e.g. KRV) or negative (e.g. LJ, NG; Tab. 2).

24 ~~Incoming solar radiation is the main energy source that drives all atmospheric processes,~~
25 ~~including PBL processes, and has a critical role also in atmospheric chemistry. For almost all~~
26 ~~sites the mean SR was overestimated by the model, with an overall ME of 16 W/m² and 11~~
27 ~~W/m² for 1 day and 2 day forecast, respectively. CORR was higher for 1 day (0.77) than for~~
28 ~~2 day (0.71) forecast, with span from 0.64 to 0.90 for 1 day forecast at different stations.~~

29 Precipitation (RR) has an important role in cleansing of the atmosphere by wet deposition and
30 scavenging. OIn average, the predicted precipitation underestimated the measured 3-month
31 accumulations byfor -55 mm (1-day) or -8 mm (2-day forecast), where the station averaged
32 predicted 3-month precipitation was 145 mm for 1-day, and 194 mm for 2-day forecast

(results not shown). ~~It must also be taken into account that the 3.4.1 model version does not allow to include the information about hydrometeors at the boundaries of the nested domain (in the applied 1-way nesting procedure), which contributes to the negative simulated bias of precipitation. A large decrease in the precipitation bias from day 1 to day 2 suggests that different initialization methodology (e.g. using 1 day spin-up for meteorology) could improve the prediction of precipitation events. Although the WRF-Chem simulations sometimes failed to correctly predict the true amount and location of the more randomly spread summertime convective precipitation, the main precipitation events (e.g. those terminating three heat wave events) were well predicted and simulated.~~

3.2 Evaluation of air quality variables

In this section we evaluate WRF-Chem predictions for O₃, NO₂ and PM₁₀, as three of the most problematic pollutants in terms of harm to human health and compliance with EU limit values (EEA, 2012). Table 3 shows the domain wide performance statistics for 1-day and 2-day forecasts of these pollutants, where in the case of O₃ 1-hour and 8-hour averages and daily maxima are analyzed separately. The comparison of 1-day and 2-day forecasts shows that concentrations of air pollutants were somewhat better forecasted 1-day than 2-days ahead by means of almost all of statistics shown in Tab. 3, with higher impact on O₃ predictions. Although the 2-day prediction was generally not worse for the majority of meteorological variables, the reason for better 1-day prediction in the case of O₃ could be somewhat stronger simulated winds on the second day of simulation. Stronger winds impact the transport and dispersion of pollutants, and have the greatest consequence for secondary pollutants (like O₃) which need time to be formed.

As shown in Tab. 3 the WRF-Chem simulations tend to overestimate the 1-hour and 8-hour O₃ values with ME of ~~14.53.7~~ μgm^{-3} and ~~14.63.8~~ μgm^{-3} , respectively. Looking at MAE, RMSE and CORR statistics, agreement with measurements is better for 8-hour (~~22.69~~ μgm^{-3} , ~~28.15~~ μgm^{-3} and 0.69) than for 1-hour O₃ values (~~25.15~~ μgm^{-3} , ~~32.15~~ μgm^{-3} and 0.65), which is in line with results of previous studies (e.g. Tong and Mauzerall, 2013) and suggests that the current modeling system has problems ~~simulatingto simulate~~ the small-scale fluctuations of O₃. On the other hand evaluations of predicted 8-hour and daily O₃ maxima, which are of most concern, show a nice model performance (ME, MAE RMSE and CORR of ~~-2.73.4~~ μgm^{-3} , ~~13.37~~ μgm^{-3} , ~~16.77.1~~ μgm^{-3} and 0.81 for daily maxima, respectively), in line or even better

1 than obtained in some previous studies (e.g. Tong and Mauzerall, 2006; Chuang et al., 2011;
2 Yahya et al., 2014), which could be to some extent related to higher model resolution.

3 To understand results of the domain wide statistics (in Tab. 3) we further analyze spatial and
4 temporal characteristics of model O₃ predictions. Figure 4 shows a spatial pattern of average
5 simulated 1-day predictions for O₃, NO₂ and PM₁₀ overlaid with measured averages, where
6 in the case of O₃ results for all hourly values and for daily maxima are shown separately.
7 Examples of forecasted and measured time series for O₃ at different stations are shown in Fig.
8 5. In Fig. 4a the elevated alpine KRV station is the only one with high negative bias (-12
9 µgm⁻³) in forecasted 1-hour O₃ concentrations at the lowest model layer, which can be
10 explained by the too low altitude of the KRV station in model topography. The high negative
11 bias for hourly O₃ concentrations at KRV station is reduced to a value of only -2 µgm⁻³ by
12 using the 5th model layer concentrations as explained in chapter 2.3. The 5th model level
13 predictions will be used for KRV in all analyses that follow, since the mean O₃ concentration
14 increases with height. Besides KRV also the Mediterranean KOP and OTL stations, as well as
15 the rural ZAV site, are stations with comparatively high measured nighttime O₃ levels, which
16 results in low overall bias for all hourly O₃ values for these stations (from -23 to -7 µgm⁻³).
17 Namely, WRF-Chem model cannot capture well the profound nighttime O₃ reductions (shown
18 also by Žabkar et al, 2013; Im et al., 2014a), which contributes to the overall over-prediction
19 of hourly O₃ concentrations (from 10 to 36 µgm⁻³) for stations with very low measured
20 nighttime O₃ concentrations. For sites with highest positive bias in 1-hour O₃ concentrations
21 (TRB, ZAG, HRA and ISK, with bias of 36 µgm⁻³, 31 µgm⁻³, 26 µgm⁻³ and 32 µgm⁻³,
22 respectively), this can also be partly explained by too high altitude of the stations in model
23 orography (Tab. 1), since the mean O₃ concentration increases with height.

24 Looking at O₃ daily maxima (Fig. 4b), the under-predictions occur at alpine KRV (-16 µgm⁻³
25 for the lowest model level shown in Fig.4) and at three Mediterranean stations (OTL, NG,
26 KOP; from -14 to -11 µgm⁻³). For Mediterranean stations the underestimations of daily
27 maxima are most probably due to inaccurate representation of coastal processes in model,
28 which are crucial for PBL height evolution and accumulation of pollution in the near ground
29 air layers. For TRB station located in narrow valley of the very complex terrain that cannot be
30 appropriately resolved in the current model topography, the model over-predicts O₃ daily
31 maxima for 14 µgm⁻³. For other sub-alpine stations the bias of O₃ daily maxima predictions is
32 lower.

1 To some extent the previously mentioned model over-predictions of nighttime O₃ minima
2 could be explained by model ~~incongruity-error~~ in predicted NO₂ levels. When evaluating the
3 primary pollutants one must be aware that in the model the instantaneous emissions are spread
4 over an entire grid box, which results in underestimated emissions and concentrations close to
5 the source regions and overestimated emissions and concentrations at rural locations adjacent
6 to the source regions, and can thus cause a combined effect of negative and positive biases at
7 urban and rural sites. Comparisons of WRF-Chem predicted NO₂ levels with measurements
8 show that in spite of the high spatial resolution the concentrations of the small urban areas are
9 insufficiently represented by the model (Fig. 4c). In Slovenia many towns are located in
10 basins or very narrow valleys, usually poorly or even not resolved in model topography.
11 Smoothed local emissions for these towns show significant underestimations of NO₂
12 concentrations (e.g. ZAG in Fig. 6). In combination with ~~poorlydeficiently~~ reproduced
13 meteorological processes (calm and stable nighttime conditions in valleys and basins) this
14 results in an underestimation of the O₃ loss by titration. This can explain the positive
15 nighttime bias of O₃ found at these sites. The situation is better for bigger cities, located in
16 wider basins, like LJ or CE (LJ; Fig. 6), while at rural sites NO₂ is either well simulated (e.g.
17 MOH; Fig. 6), or slightly over-predicted due to increased emissions from adjacent urban area
18 (e.g. ZAD; Fig. 6). The overall agreement of hourly NO₂ predictions with measurements was
19 good for rural sites, while urban sites experienced under-predictions, which were highest for
20 small cities, especially for NG (ME of -13 µgm⁻³) and ZAG (ME of -14 µgm⁻³).

21 ~~Also fi~~ interesting to discuss are ~~thealso~~ results for predicted PM₁₀ concentrations (Tab. 3 and
22 Fig. 4d), showing slight over-prediction of daily PM₁₀ levels at all stations which is
23 somewhat surprising due to the fact that nearly all current off-line and on-line coupled
24 chemical transport models show large systematic PM₁₀ underestimations. For example,
25 within AQMEII exercise, where seventeen modeling groups from Europe and North America
26 were brought together, running eight operational online-coupled air quality models over
27 Europe and North America, the rural PM₁₀ concentrations over Europe were underestimated
28 by all models (model configurations) by up to 66% while for the urban PM₁₀ concentrations
29 the underestimations were even much larger (up to 75%) (Im et al., 2014b). The reason for
30 slight over-prediction of PM₁₀ levels could be to some extent attributed to the high model
31 spatial resolution used in our study. Further, CORR for daily PM₁₀ concentrations is rather
32 low (0.34 and 0.37 for 1-day and 2-day forecasts, respectively; Tab. 3), which is partly due to
33 the low temporal dynamics of measured daily PM₁₀ concentrations during the analyzed time

1 | period (no recorded PM10 ~~exceeding exceedances~~), and partly due to the simulated PM10
2 | overestimations during the heat wave events. These over-predictions contributed also to the
3 | overall positive bias of predicted PM10 levels. As shown in Fig. 7 for two monitoring sites,
4 | there was a significant PM10 over-prediction simulated on June 10 (day 8 in Fig. 7), related to
5 | the pre-frontal advection of polluted air-masses coming from the north-western part of the
6 | domain D2 (coming from domain D1). The next significant PM10 over-prediction occurred
7 | during the first heat wave episode (June 17-22), when during the hot and low wind conditions
8 | (after June 17) the PM10 levels started to build up in the PBL over entire domain D2 (and
9 | over southwestern parts of domain D1), and reached the maximum concentrations in Slovenia
10 | again with prefrontal advection of polluted air masses. Both over-predictions contributed to
11 | an overall positive bias in forecasted PM10 concentrations. Detailed analyses showed that
12 | high concentrations in domain D1 originated from boundary conditions, and appear to be a
13 | consequence of overestimated advection of Saharan dust in MOZART model predictions. The
14 | increase in PM10 concentrations over Slovenia was also simulated during the prefrontal
15 | advection related to the cold front which terminated the next two heat wave events in July and
16 | August (days 56-57 and days 67-68 in Fig. 7), but during these days predicted PM10 levels
17 | were close to the measured PM10 concentrations.

18 | **3.3 Evaluation and comparison of different methods for O₃ daily maximum** 19 | **predictions**

20 | In this section we want to answer the question: “–how accurate is the 1-hour O₃ daily
21 | maximum WRF-Chem forecast in comparison ~~with to~~ the statistical model prediction or ~~with~~
22 | to persistence?”. According to Zhang et al. (2012a) statistical models are known to be
23 | generally more suitable for complex site-specific relations between concentrations of air
24 | pollutants and predictors. With appropriate and accurate predictors they have a higher
25 | accuracy as compared to deterministic models, which is, along with their ~~–beside the~~
26 | computational efficiency their main advantage (Zhang et al., 2012a). Among the strengths of
27 | the deterministic models are that they give prognostic time- and spatially-resolved
28 | concentrations under typical and atypical scenarios, and can give scientific insights into
29 | pollutant formation processes (Zhang et al., 2012a). Furthermore, they also allow forecasts
30 | also for locations which are not monitored due to their complete spatial coverage. In spite of
31 | simplified descriptions of physical and chemical processes in the deterministic models and
32 | inaccuracies and uncertainties in model inputs (in particular the emissions), some previous

1 studies already suggested that deterministic models can also have skills close to statistical
2 forecasting tools (e.g. Manders et al., 2009). In addition to evaluation and comparison of O₃
3 daily maxima predictions with WRF-Chem and the statistical model, we decided to add a
4 persistence model as a threshold for useful model prediction. Persistence works well under
5 stationary conditions, but because it cannot handle changes in weather and emissions, fails at
6 the beginning and at the end of the episodes (Zhang et al., 2010a). Regarding the extremes,
7 models of all types are known to have problem to accurately predict them, while persistence
8 predicts extremes with a 1-day (2-day) time lag.

9 Figure 8 compares discrete statistics site by site for 1-day and 2-day model predictions of 1-
10 hour O₃ daily maxima. Similarly, Tab. 4 shows these statistics for all data with different
11 thresholds applied (only for WRF-Chem and persistence, because a statistical forecast is not
12 available for all stations), and separately for different types of stations (sub-alpine urban,
13 rural, Mediterranean urban) with an available statistical forecast. Looking at ME persistence
14 gives results close to zero as long as no threshold is applied, while with threshold of 140 μgm⁻³
15 ³ (Tab. 4) ME of 1-day persistence (-10.2 μgm⁻³) is very close to the WRF-Chem model for 1-
16 day predictions (-11.29 μgm⁻³), and for 2-day predictions WRF-Chem (-13.84.6 μgm⁻³)
17 already beats ~~the~~ persistence (-19.4 μgm⁻³). Site-by-site comparison (Fig. 8) shows that for
18 most stations the statistical forecast has a lower ME than WRF-Chem forecast, but there are
19 also stations (ISK, HRA, LJ, KRV) with lower or equal ME for WRF-Chem than for
20 statistical model, indicating the possible occurrence of atypical conditions not resolved by the
21 statistical model. Looking at MAE and RMSE, at all stations except those with highest ME
22 (KRV, TRB, KOP) WRF-Chem outperforms the persistence already in the 1-day forecast.
23 Among sites with available statistical forecast there are only two (OTLKRV, KOP) with
24 WRF-Chem performing worse than the statistical forecast. CORR is one of the parameters
25 that suggest how much the model is able to follow the true nature of processes regardless the
26 possible bias. For almost all stations WRF-Chem shows higher CORR than persistence for 1-
27 day and 2-day forecasts. Only at the KRV station the 1-day statistical forecast (CORR=0.80)
28 somewhat outperforms WRF-Chem (0.746), and at NG and KOP CORR for WRF-Chem and
29 statistical model is very similar.

30 The Taylor diagrams in Fig. 9 show CORR together with the centered root-mean-square
31 difference (RMSD) between model forecasts and observations, and the amplitude of their
32 variations (standard deviation), ~~The ideal where ideal model forecast would lie in the right~~

~~bottom corner~~ have a correlation coefficient of 1 and a standard deviation equal to the observations, which means that it would be co-located with the black dot on the diagram.

WRF-Chem gives higher CORR and lower RMSD for all types of stations, while standard deviation of WRF-Chem O₃ daily maxima predictions is underestimated and lower than for other model forecasts. The latter shows that the variability in WRF-Chem model predictions is not as large as that in observed values. MNBE in Fig. 8 has ~~a course~~ very similar results to ME. For all forecasts except WRF-Chem for the TRB site (with MNBE of 16%) which is located in a narrow valley that is not resolved in the current model resolution, MNBE is below the ±10-15%, which is the U.S. EPA (US EPA, 1991) recommended threshold for the models used for regulatory applications. For MNGE the U.S. EPA recommendation below 30-35% for O₃ applications is met by all forecasts, even in the case of 2-day persistence model. With exception of the MS and KOP sites MNGE is lower for WRF-Chem than for statistical forecast, while for KOP and KRV sites ~~with highest negative bias in WRF-Chem predictions~~, 1-day persistence gives best results, followed by the statistical forecast ~~and or~~ WRF-Chem. Very similar are results for IOA with the range of 0-1, and score 1 indicating perfect model agreement with the observations. We can conclude that for most stations the WRF-Chem predictions are in line or even outperform the statistical model. With the exception of the stations with high bias due to very complex local topography (TRB) ~~or~~ unresolved coastal processes (KOP) ~~or alpine stations (KRV)~~, the WRF-Chem forecasts are more accurate than persistence. Here we recall that high negative bias in WRF-Chem forecast for alpine KRV site due to too low altitude of the station in model topography was compensated by taking prediction from the 5th model level.

The key requirement for a forecast system is to be able to predict O₃ concentration levels greater than a given threshold. Thus, in addition to the discrete evaluation just presented, also the contingency-table-based statistics are an important metrics of forecast performance. Table 5 summarizes the categorical evaluation results for three different thresholds (120, 140, 160 μgm⁻³) of elevated O₃ levels, which pose a greater risk to human health. Namely, it is important to take ~~should be taken~~ into account that results of categorical statistics are very sensitive to the threshold chosen, as well as to the overall pollution levels during the analyzed months. Equitable Threat Score (ETS) measures the fraction of observed and/or correctly predicted events, adjusted for the frequency of hits that would be expected to occur by random chance. Although this score takes into account the climatology it is not truly equitable. It ranges from -1/3 to 1, where the minimum value depends on climatology (it is

1 near 0 for rare events). Looking at Tab. 5 ETS shows equal skill for WRF-Chem and
2 statistical forecast, higher than persistence for the 120 μgm^{-3} threshold (1-day and 2-day
3 forecast). ETS decreases with increasing the threshold for both WRF-Chem and statistical
4 forecast, indicating the challenge that both models have to accurately predict the extremes. In
5 the case of 140 μgm^{-3} threshold, WRF-Chem has the same ETS as persistence, higher than the
6 statistical model for 1-day forecast, while for 2-day forecast WRF-Chem outperforms the
7 statistical model, followed by persistence. In the case of 160 μgm^{-3} threshold persistence has
8 the highest ETS for a 1-day forecast, followed by statistical model and WRF-Chem, while in
9 the case of 2-day predictions, statistical model shows the highest skill and WRF-Chem the
10 lowest. Accuracy (A), which measures how often the forecasts are correct either above or
11 below the threshold, increases with threshold level. Looking at 1-day forecast A is highest for
12 statistical forecast at 120 μgm^{-3} threshold, for WRF-Chem forecast at 140 μgm^{-3} threshold,
13 and in the case of 160 μgm^{-3} threshold applied, for persistence. Another measure, the critical
14 success index (CSI), is similar to ETS, except that it does not take into account the
15 climatology of the events and thus gives poorer scores for rarer events. It measures the
16 percentage of cases that are correctly forecasted out of those either forecasted or observed,
17 and ranges from 0 to 1 (1 indicating the perfect forecast). Similar as ETS, CSI gives higher
18 scores for persistence in the case of 1-day forecast for the higher two thresholds, while on the
19 second day WRF-Chem or the statistical model already performs better. There is Bias (B)
20 determines whether the same fraction of events are both forecasted and observed. A tendency
21 of the statistical model and of WRF-Chem to under-predict O_3 threshold exceedances shows
22 as a B below 1 for these two models. a tendency of the statistical model and of WRF-Chem to
23 under predict O_3 daily maxima. This shows as a bias (B) below 1 for these two models, where
24 B determines whether the same fraction of events are both forecasted and observed. The false
25 alarm ratio (FAR) that measures the percentage of forecast high O_3 events that turn out to be
26 false alarms, gives highest skill for WRF-Chem, followed by statistical model and
27 persistence. The probability of detection (POD) is a measure of how often a high threshold
28 occurrence is actually predicted to occur, and is relatively low for WRF-Chem with respect to
29 other models. Another useful measure, the critical success index (CSI), measures the
30 percentage of cases that are correctly forecasted out of those either forecasted or observed,
31 and is for higher two thresholds best for persistence in the case of 1-day forecast, while on the
32 second day WRF-Chem or the statistical model already perform better.

1 It must be noted, that in categorical evaluations systematic biases like those obtained with
2 WRF-Chem for some stations (e.g. KOP, KRV), significantly impact the model performance.
3 For example, if KOP ~~and KRV~~ stations ~~was~~were excluded from categorical evaluations,
4 WRF-Chem performance improved by means of all statistical measures (results not shown). If
5 correction techniques, based on observations and the previous day's forecast (e.g., McKeen et
6 al., 2005, 2007; Kang et al., 2008) ~~were to~~would be applied to correct the systematic biases,
7 WRF-Chem forecasts might outperform the other two models even in categorical evaluations.

9 **4 Summary and conclusion**

10 A high resolution modelling system based on an on-line coupled WRF-Chem has been
11 applied for numerical weather prediction and for forecasting air quality in Slovenia. In the
12 study the evaluation of the forecasting system has been conducted for three summer months.
13 Since the selection of physical or chemical parameterization schemes influences and possibly
14 changes the outcomes, we decided to apply schemes which are well documented and have
15 previously been used in other applications (e.g. AQMEII). -Both 1-day and 2-day predictions
16 of meteorological and air quality variables have been analyzed. The focus has been on O₃ as
17 the only pollutant with recorded exceedances of legislation limit values during the three heat
18 wave events in June, July and August 2013. WRF-Chem daily O₃ maximum predictions have
19 also been compared to the operational statistical model and persistence forecasts to answer the
20 question how skillful are the WRF-Chem model predictions compared to these two models.

21 1-day and 2-day WRF-Chem PM₁₀ forecasts showed a very low bias. Exceptions were two
22 events with significantly over-predicted PM₁₀ levels due to prefrontal advection of polluted
23 air masses from neighboring regions. Knowing that majority of the current chemical transport
24 models show large negative biases in simulated PM₁₀ concentrations, these results present a
25 good starting point for studying the importance of aerosol feedbacks with realistic model
26 aerosol concentrations, left for future research.

27 The overall agreement of WRF-Chem NO₂ forecast with measurements was good for rural
28 sites, while urban sites experienced model under-predictions, which were highest for small
29 towns. One important reason is that many small towns are located in basins or very narrow
30 valleys, usually poorly presented in model topography. Smoothed local emissions result
31 showin ~~-as~~ model underestimations of NO₂ concentrations for these towns. This in
32 combination with insufficiently reproduced calm meteorological conditions in basins and

1 valleys during the nighttime hours explains also WRF-Chem over-predictions of nighttime O₃
2 concentrations.

3 Evaluations of predicted 1-hour and 8-hour daily O₃ maxima, which are in the case of this
4 pollutant of the highest interest, show ~~good~~^{nice} WRF-Chem model performance.
5 Nevertheless, there are also stations which experience high over- or under-predictions of O₃
6 daily maximum levels. For Mediterranean sites the under-predictions of the daily maxima are
7 most probably due to inaccurate representation of coastal processes in model, which are crucial
8 for the PBL height evolution and accumulation of pollution in the near ground air layers. For
9 some sub-alpine stations the reason for the higher bias in O₃ daily maximum predictions is
10 their location either at elevated mountainous or coastal regions, or in narrow valleys which
11 cannot be appropriately resolved in the current model resolution - that impacts how accurately
12 model simulates the local processes responsible for the level of local pollution. Comparisons
13 of WRF-Chem O₃ daily maximum forecasts with persistence and with statistical model
14 predictions show that with respect to some statistical parameters the deterministic WRF-
15 Chem forecast can outperform the other two for both 1-day and 2-day predictions. For
16 example, correlation coefficient shows highest skill for WRF-Chem model, confirming the
17 importance of complex processes as taken into account in an on-line coupled Eulerian model.
18 Further improvement of WRF-Chem forecasting skill could be obtained by applying one of
19 the bias-correction methods in order to account for unresolved topographical and coastal
20 effects, as well as emission patterns. Chemical data assimilation, although currently still in its
21 infancy for online coupled meteorology-chemistry models (Bocquet et al., 2014), could in
22 future also be used as an efficient method for improving prediction of chemical concentration
23 fields. [For WRF-Chem model a technical note on the implementation of the aerosol](#)
24 [assimilation and a guidance for prospective users has been recently published by Pagowski et](#)
25 [al. \(2014\).](#)

26

27 **Appendix A: Statistical measures**

28 For *i*-th observed (O_i) and the corresponding modelled (M_i) value of variable, discrete
29 statistical measures are calculated as follows:

30 Mean error:

$$31 \quad ME = \frac{1}{N} \sum_{i=1}^N (M_i - O_i)$$

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Mean absolute error:

$$MAE = \frac{1}{N} \sum_{i=1}^N |M_i - O_i|$$

Root mean square error:

$$RMSE = \sqrt{\frac{1}{N} \sum_{i=1}^N (M_i - O_i)^2}$$

Correlation coefficient:

$$r = \frac{\sum_{i=1}^N (M_i - \bar{M})(O_i - \bar{O})}{\sqrt{\sum_{i=1}^N (M_i - \bar{M})^2 (O_i - \bar{O})^2}}$$

Index of agreement:

$$IOA = 1 - \frac{\sum_{i=1}^N (M_i - O_i)^2}{\sum_{i=1}^N (|M_i - \bar{O}| + |O_i - \bar{O}|)^2}$$

Mean normalized bias error:

$$MNBE = \frac{1}{N} \sum_{i=1}^N \frac{M_i - O_i}{O_i} \times 100$$

Mean normalized gross error:

$$MNGE = \frac{1}{N} \sum_{i=1}^N \frac{|M_i - O_i|}{O_i} \times 100$$

For categorical evaluation all model predictions are first classified into four groups (a , b , c and d):

- a prediction is above, but observation is below the threshold
- b prediction and observation are above the threshold
- c prediction and observation are below the threshold
- d prediction is below, but observation is above the threshold

Categorical statistics are calculated as follows:

Equitable threat score: $ETS = \frac{b - a_r}{a + b + d - a_r}$, where $a_r = \frac{(a + b)(b + d)}{a + b + c + d}$

Critical success index: $CSI = \frac{b}{a + b + d}$

Bias: $B = \frac{a + b}{b + d}$

False alarm ratio: $FAR = \frac{a}{a + b}$

1 Probability of detection: $POD = \frac{b}{b+d}$

2

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1 **References**

- 2 Ackermann, I. J., Hass, H., Memmesheimer, M., Ziegenbein, C., Ebel, A.: The
3 parameterization of the sulfate-nitrate-ammonia aerosol system in the long-range transport
4 model EURAD. *Meteorological Atmospheric Physics*, 57, 101-114, 1995.
- 5 ALADIN International Team, The ALADIN project: Mesoscale modelling seen as a basic
6 tool for weather forecasting and atmospheric research. *WMO Bull.*, 46, 317–324, 1997.
- 7 Baklanov, A., Korsholm, U., Mahura, A., Petersen, C., Gross, A.: Enviro-HIRLAM: on-line
8 coupled modelling of urban meteorology and air pollution. *Adv. Sci. Res.*, 2, 41-46.
- 9 Baklanov, A., 2010. Chemical weather forecasting: a new concept of integrated modeling.
10 *Adv. Sci. Res.*, 4, 23-27, 2008.
- 11 Baklanov, A., Schlünzen, K., Suppan, P., Baldasano, J., Brunner, D., Aksoyoglu, S.,
12 Carmichael, G., Douros, J., Flemming, J., Forkel, R., Galmarini, S., Gauss, M., Grell, G.,
13 Hirtl, M., Joffre, S., Jorba, O., Kaas, E., Kaasik, M., Kallos, G., Kong, X., Korsholm, U.,
14 Kurganskiy, A., Kushta, J., Lohmann, U., Mahura, A., Manders-Groot, A., Maurizi, A.,
15 Moussiopoulos, N., Rao, S. T., Savage, N., Seigneur, C., Sokhi, R. S., Solazzo, E., Solomos,
16 S., Sørensen, B., Tsegas, G., Vignati, E., Vogel, B., Zhang, Y.: Online coupled regional
17 meteorology chemistry models in Europe: current status and prospects. *Atmospheric*
18 *Chemistry and Physics*, 14, 317-398, 2014.
- 19 Balzarini, A., Pirovano, G., Honzak L., Žabkar, R., Curci, G., Forkel R., Hirtl, M., San José ,
20 R., Tuccella, P., Grell, G.: WRF-Chem model sensitivity to chemical mechanisms choice in
21 reconstructing aerosol optical properties. *Atmospheric Environment*,
22 doi:10.1016/j.atmosenv.2014.12.033, 2014.
- 23 Baró, R., Jiménez-Guerrero, P., Balzarini, A., Curci, G., Forkel, R., Hirtl, M., Honzak, L., Im,
24 U., Lorenz, C., Pérez, J.L., Pirovano, G., San José, R., Tuccella, P., Werhahn, J., Žabkar, R.:
25 Sensitivity analysis of the microphysics scheme in WRF-Chem contributions to AQMEII
26 phase 2. *Atmospheric Environment*, submitted, 2014.
- 27 M. Bocquet, Elbern, H., Eskes, H., Hirtl, M., Žabkar, R., Carmichael, G.R., Flemming, J.,
28 Inness, A., Pagowski, M., Pérez Camaño, J.L., Saide, P.E., San Jose, R., Sofiev, M., Vira, J.,
29 Baklanov, A., Carnevale, C., Grell, G., Seigneur, C.: Data Assimilation in Coupled Chemistry
30 Meteorology Models, submitted to ACP/GMD Special Issue, 2014.

1 Byun, D. W., Schere, K. L.: Review of the governing equations, computational algorithms,
2 and other components of the Models- 3 Community Multiscale Air Quality (CMAQ)
3 Modeling System. *Appl. Mech. Rev.*, 59, 51–77, 2006.

4 Chapman, E. G., Gustafson Jr., W. I., Easter, R.C., Barnard, J. C., Ghan, S. J., Pekour, M. S.,
5 Fast J. D.: Coupling aerosol-cloud-radiative processes in the WRF-Chem model: Investigating
6 the radiative impact of elevated point sources. *Atmospheric Chemistry and Physics*, 9, 945-
7 964, 2009.

8 Chen, F., Dudhia, J.: Coupling an Advanced Land Surface–Hydrology Model with the Penn
9 State NCAR MM5 Modeling System. Part I: Model Implementation and Sensitivity. *Monthly*
10 *Weather Review*, 129, 569–585, 2001.

11 Chuang, M.T., Zhang, Y., Kang, D.W.: Application of WRF-Chem-MADRID for real-time
12 air quality forecasting over the southeastern United States. *Atmospheric Environment*, 45,
13 6241-6250, 2011.

14 Cobourn, W.G.: Accuracy and reliability of an automated air quality forecast system for
15 ozone in seven Kentucky metropolitan area. *Atmospheric Environment* 41, 5863-5875, 2007.

16 [Curci, G., Hogrefe, C., Bianconi, R., Im, U., Balzarini, A., Baro, R., Brunner, D., Forkel, R.,](#)
17 [Giordano, L., Hirtl, M., Honzak, L., Jimenez-Guerrero, P., Knote, C., Langer, M., Makar,](#)
18 [P.A., Pirovano, G., Perez, J.L., San Jose, R., Syrakov, D., Tuccella, P., Werhahn, J., Wolke,](#)
19 [R., Zabkar, R., Zhang, J., Galmarini, S. \(2014\), Uncertainties of simulated aerosol optical](#)
20 [properties induced by assumptions on aerosol physical and chemical properties: an AQMEII-](#)
21 [2 perspective, Atmospheric Environment, doi: 10.1016/j.atmosenv.2014.09.009.](#)

22 EC/2008/50. Directive 2008/50/EC of the European Parliament and of the Council of 21 May
23 2008 on ambient air quality and cleaner air for Europe. *Official Journal of the European*
24 *Union*, L152, 44 pp., 2008.

25 Eder, B.K., Kang, D., Mathur, R., Yu, S., Schere, K.: An operational evaluation of the Eta-
26 CMAQ air quality forecast model. *Atmos. Environ.* 40, 4894-4905, 2006.

27 EEA: The application of models under the European Union's Air Quality Directive: A
28 technical reference guide, 72 pp., 2011.

29 EEA: Air Quality in Europe – 2012 Report, ISBN 978-92-9213-328-3, Luxembourg: Office
30 for Official Publications of the European Union, 108pp., 2012.

1 Emmons, L. K., Walters, S., Hess, P. G., Lamarque, J.-F., Pfister, G. G., Fillmore, D.,
2 Granier, C., Guenther, A., Kinnison, D., Laepple, T., Orlando, J., Tie, X., Tyndall, G.,
3 Wiedinmyer, C., Baughcum, S. L., and Kloster, S.: Description and evaluation of the Model
4 for Ozone and Related chemical Tracers, version 4 (MOZART-4), *Geosci. Model Dev.*, 3, 43-
5 67, doi:10.5194/gmd-3-43-2010, 2010.

6 ENVIRON: CAMx User's Guide, Comprehensive Air Quality Model With Extensions
7 Version 5.40, ENVIRON International Corporation, Novato, California, 2011.

8 Fast, J., et al.: Evaluating simulated primary anthropogenic and biomass burning organic
9 aerosols during MILAGRO: implications for assessing treatments of secondary organic
10 aerosols. *Atmos. Chem. Phys.*, 9, 6191-6215, 2009.

11 Fast, J. D., Gustafson Jr., W. I., Easter, R. C., Zaveri, R. A., Barnard, J. C., Chapman, E. G.,
12 Grell, G. A., Peckham, S. E.: Evolution of ozone, particulates, and aerosol direct radiative
13 forcing in the vicinity of Houston using a fully coupled meteorology-chemistry-aerosol
14 model. *J. Geophys. Res.*, 111, D21305, 2006.

15 Forkel, R., Werhahn, J., Buus Hansen, A., McKeen, S., Peckham, S., Grell, G., Suppan, P.:
16 Effect of aerosol-radiation feedback on regional air quality – A case study with WFR/Chem.
17 *Atmospheric Environment* 53, 202-211, 2012.

18 Forkel, R., Balzarini, A., Baró, R., Curci, G., Jiménez-Guerrero, P., Hirtl, M., Honzak, L., Im,
19 U., Lorenz, C., Pérez, J.L., Pirovano, G., San José, R., Tuccella, P., Werhahn, J., Žabkar, R.:
20 Analysis of the WRF-Chem contributions to AQMEII phase2 with respect to aerosol radiative
21 feedbacks on meteorology and pollutant distribution. *Atmospheric Environment*,
22 doi:10.1016/j.atmosenv.2014.10.056, 2014.

23 Grell, G., Dudhia, J., Stauffer, D.: A description of the fifth-generation Penn State/NCAR
24 Mesoscale model (MM5). TN-398+STR,NCAR, Boulder, CO, 1995~~4~~.

25 Grell, G.A., Emeis, S., Stockwell, W.R., Schoenemeyer, Forkel, R., Michalakes, J., Knoche,
26 R., Seidl, W.: Application of a multiscale, coupled MM5/chemistry model to the complex
27 terrain of the VOTALP valley campaign, *Atmospheric Environment* 34, 1435– 1453, 2000.

28 Grell, G. A., Devenyi, D.: A generalized approach to parameterizing convection combining
29 ensemble and data assimilation techniques, *Geophys. Res. Lett.* 29, 14, 2002.

1 Grell, G. A., Knoche, R., Peckham, S. E., and McKeen, S. A.: Online versus offline air
2 quality modeling on cloud-resolving scales, *Geophysical Research Letters* 31, L16117,
3 doi:10.1029/2004GL020175, 2004.

4 Grell, G.A., Peckham, S.E., Schmitz, R., McKeen, S.A., Frost, G., Skamarock, W., Eder, B.:
5 Fully coupled “online” chemistry within the WRF model. *Atmospheric Environment*, 39,
6 6957-6975, 2005.

7 Grell, G., Baklanov, A.: Integrated modeling for forecasting weather and air quality: A call
8 for fully coupled approaches. *Atmospheric Environment*, 45, 6845-6851, 2011.

9 Guenther, A., Karl, T., Harley, P., Wiedinmyer, C., Palmer, P. I., Geron, C.: Estimates of
10 global terrestrial isoprene emissions using MEGAN (Model of Emissions of Gases and
11 Aerosols from Nature). *Atmospheric Chemistry and Physics*, 6, 3181-3210, 2006.

12 Hong, S., Noh, Y., Dudhia, J.: A new vertical diffusion package with an explicit treatment of
13 entrainment processes. *Monthly Weather Review*, 134, 2318-2341, 2006.

14 Hu, X.-M., Doughty, D., Sanchez, K.J., Joseph, E., and Fuentes, J. D.: Ozone variability in
15 the atmospheric boundary layer in Maryland and its implications for vertical transport model,
16 *Atmos. Environ.*, 46, 354-364, 2012.

17 Iacono, M. J., Delamere, J. S., Mlawer, E. J., Shephard, M. W., Clough, S. A., and Collins,
18 W. D.: Radiative forcing by long-lived greenhouse gases: Calculations with the AER radiative
19 transfer models, *J. Geophys. Res.*, 113, D13103, 2008.

20 Im, U., Bianconi, R., Solazzo, E., Kioutsioukis, I., Badia, A., Balzarini, A., Baro, R., Bellasio,
21 R., Brunner, D., Chemel, C., Curci, G., Flemming, J., Forkel, R., Giordano, L., Jimenez-
22 Guerrero, P., Hirtl, M., Hodzic, A., Honzak, L., Jorba, O., Knote, C., Kuenen, J.J.P., Makar,
23 P.A., Manders-Groot, A., Neal, L., Perez, J.L., Pirovano, G., Pouliot, G., San Jose, R.,
24 Savage, N., Schroder, W., Sokhi, R.S., Syrakov, D., Torian, A., Tuccella, P., Werhahn, K.,
25 Wolke, R., Yahya, K., Žabkar, R., Zhang, Y., Zhang, J., Hogrefe, C., Galmarini, S.:
26 Evaluation of operational online-coupled regional air quality models over Europe and North
27 America in the context of AQMEII phase 2. Part I: Ozone. *Atmospheric Environment*,
28 doi:10.1016/j.atmosenv.2014.09.042, 2014a.

29 Im, U., Bianconi, R., Solazzo, E., Kioutsioukis, I., Badia, A., Balzarini, A., Baro, R., Bellasio,
30 R., Brunner, D., Chemel, C., Curci, G., Denier van der Gon, H.A.C., Flemming, J., Forkel, R.,
31 Giordano, L., Jimenez-Guerrero, P., Hirtl, M., Hodzic, A., Honzak, L., Jorba, O., Knote, C.,

1 Makar, P.A., Manders-Groot, A., Neal, L., Perez, J.L., Pirovano, G., Pouliot, G., San Jose, R.,
2 Savage, N., Schroder, W., Sokhi, R.S., Syrakov, D., Torian, A., Tuccella, P., Werhahn, K.,
3 Wolke, R., Yahya, K., Žabkar, R., Zhang, Y., Zhang, J., Hogrefe, C., Galmarini, S.:
4 Evaluation of operational online-coupled regional air quality models over Europe and North
5 America in the context of AQMEII phase 2. Part II: Particulate Matter. *Atmospheric*
6 *Environment*, doi:10.1016/j.atmosenv.2014.08.072, 2014b.

7 Jacobson, M. Z.: GATOR-GCMM: A global through urban scale air pollution and weather
8 forecast model. 1. Model design and treatment of subgrid soil, vegetation, roads, rooftops,
9 water, sea ice, and snow. *J. Geophys. Res.* 106, 5385-5402. 2001

10 Kaminski J.W., Neary L., Struzewska J., McConnell J.C., Lupu A., Jarosz J., Toyota K.,
11 Gong S.L., Cote J., Liu X., Chance K., Richter A.: GEM-AQ, an on-line global multiscale
12 chemical weather modelling system: model description and evaluation of gas phase chemistry
13 processes. *Atmos Chem Phys* 8:3255–3281, 2008.

14 Kang, D., Mathur, R., Rao, S.T., Yu, S.: Bias adjustment techniques for improving ozone air
15 quality forecasts. *J. Geophys. Res.* 113, D23308, 2008.

16 Klein, T., Kukkonen, J., Dahl, A., Bossioli, E., Baklanov, A., Vik, A.F., Agnew, P., Karatzas,
17 K.D., Sofiev, M.: Interactions of Physical, Chemical, and Biological Weather Calling for an
18 Integrated Approach to Assessment, Forecasting, and Communication of Air Quality.
19 *AMBIO*, 41, 851–864, 2012.

20 Kong, X., Forkel, R., Sokhi, R., Suppan, P., Baklanov, A., Gauss, M., Brunner, D., Baro
21 Esteban, R., Balzarini, A., Chemel, C., Curci, G., Galmarini, S., Jiménez Guerrero, P., Hirtl,
22 M., Honzak, L., Im, U., Pérez, J. L., Piravano, G., San Jose, R., Schlünzen, H., Tsegas, G.,
23 Tuccella, P., Werhahn, J., Žabkar, R.: Investigation of meteorology and chemistry interactions
24 and their representations in online coupled models with the supported case Studies from
25 AQMEII phase2. *Atmospheric Environment*, doi:10.1016/j.atmosenv.2014.09.020, 2014.

26 Li, G., Zavala, M., Lei, W., Tsimpidi, A.P., Karydis, V.A., Pandis, S.N., Canagaratna, M.R.,
27 Molina, L.T.: Simulations of organic aerosol concentrations in Mexico City using the WRF-
28 CHEM model during the MCMA-2006/MILAGRO campaign. *Atmos. Chem. Phys.* 11, 3789-
29 3809, 2011.

1 Manders, A.M.M., Schaap, M., Hoogerbrugge, R.: Testing the capability of the chemistry
2 transport model LOTOS-EUROS to forecast PM10 levels in the Netherlands. *Atmos.*
3 *Environ.*, 43, 4050-4059, 2009.

4 McCollister, G., Wilson, K.: Linear stochastic models for forecasting daily maxima and
5 hourly concentrations of air pollutants. *Atmos. Environ.*, 9, 417-423, 1975.

6 McKeen, S., Wilczak, J., Grell, G., Djalova, I., Peckham, S., Hsie, E.-Y., Gong, W., Bouchet,
7 V., Ménard, S., Moffet, R., McHenry, J., McQueen, J., Tang, Y., Carmichael, G.R.,
8 Pagowski, M., Chan, A., Dye, t., Frost, G., Lee, P., Mathur, R.: Assessment of an ensemble of
9 seven real-time ozone forecasts over eastern North America during the summer of 2004. *J.*
10 *Geophys. Res.*, 110, D21307. <http://dx.doi.org/10.1029/2005JD005858>, 2005.

11 McKeen, S., Chung, S.H., Wilczak, J., Grell, G., Djalalova, I., Peckham, S., Gong, W.,
12 Bouchet, V., Moffet, R., Tang, Y., Carmichael, G.R., Mathur, R., Yu, S.: Evaluation of
13 several PM2.5 forecast models using data collected during the ICARTT/NEAQS 2004 field
14 study. *J. Geophys. Res.*, 112, D10S20. <http://dx.doi.org/10.1029/2006JD007608>, 2007.

15 McKeen, S., et al.: An evaluation of real-time air quality forecasts and their urban emissions
16 over eastern Texas during the summer of 2006 Second Texas Air Quality Study field study. *J.*
17 *Geophys. Res.*, 114, D00F11. <http://dx.doi.org/10.1029/2008JD011697>, 2009.

18 Menut, L., Bessagnet, B., Khvorostyanov, D., Beekmann, M., Blond, N., Colette, A., Coll, I.,
19 Curci, G., Foret, G., Hodzic, A., Mailler, S., Meleux, F., Monge, J.J., Pison, I., Siour, G.,
20 Turquety, S., Valari, M., Vautard R., Vivanco, M.G.: CHIMERE 2013: a model for regional
21 atmospheric composition modelling, *Geoscientific Model Development*, 6, 981-1028,
22 [doi:10.5194/gmd-6-981-2013](https://doi.org/10.5194/gmd-6-981-2013), 2013.

23 Misennis, C., Zhang, Z.: An examination of sensitivity of WRF-Chem predictions to physical
24 parameterizations, horizontal grid spacing, and nesting options. *Atmospheric Research*, 97,
25 315–334, 2010.

26 Moody, J., Galloway, J.: Quantifying the relationship between atmospheric transport and the
27 chemical composition of precipitation on Bermuda, *Tellus*, 40B, 436–479, 1988.

28 Morrison, H., Thompson, G., Tatarskii, V.: Impact of cloud microphysics on the development
29 of trailing stratiform precipitation in a simulated squall line: Comparison of one- and two-
30 moment schemes. *Monthly Weather Review*, 137, 991-1007, 2009.

1 Otte, T.L., Pouliot, G., Pleim, J.E., Young, J.O., Schere, K.L., Wong, D.C., Lee, P.C.S.,
2 Tsidulko, M., McQueen, J.T., Davidson, P., Mathur, R., Chuang, H.-Y., DiMego, G., Seaman,
3 N.L.: NCEP Notes: linking the Eta model with the community multiscale air quality (CMAQ)
4 modeling system to build a national air quality forecasting system. *Weather Forecast*, 20, 367-
5 384, 2005.

6 Peckham, S.E., Grell, G.A., McKeen, S.A., Barth, M., Pfister, G., Wiedinmyer, C., Fast, J.D.,
7 Gustafson, W.I., Ghan, S.J., Zaveri, R., Easter, R.C., Barnard, J., Chapman, E., Hewson, M.,
8 Schmitz, R., Salzman, M., Freitas, S.R.: WRF-Chem Version 3.3 User's Guide, 2011.

9 [Pouliot, G., Pierce, T., Denier van der Gon, Kuenen, J., Zhang, J., M., Moran, M., Makar, P.:](#)
10 [Analysis of the emission inventories and model-ready emission datasets of Europe and North](#)
11 [America for phase 2 of the AQMEII project, Atmospheric Environment,](#)
12 <http://dx.doi.org/10.1016/j.atmosenv.2014.10.061>, 2014.

13 Saide, P.E., Carmichael, G.R., Spak, S.N., Gallardo, L., Osses, A.E., Mena-Carrasco, M.A.,
14 Pagowski, M.: Forecasting urban PM10 and PM2.5 pollution episodes in very stable
15 nocturnal conditions and complex terrain using WRF-Chem CO tracer model. *Atmospheric*
16 *Environment*, 45, 2769-2780, 2011.

17 [San José, R., Pérez, J.L., Balzarini, A., Barò, R., Curci, G., Forkel, R., Galmarini, S., Grell,](#)
18 [G., Hirtl, M., Honzak, L., Im, U., Jimenez-Guerrero, P., Langer, M., Pirovano, G., Tuccella,](#)
19 [P., Werhahn, J., Zabkar, R. \(2015\), Sensitivity of feedback effects in CBMZ/MOSAIC](#)
20 [chemical mechanism, Atmospheric Environment, doi: 10.1016/j.atmosenv.2015.04.030.](#)

21 [SEA: Upgrade of the system for monitoring air pollution, determining the causes of excessive](#)
22 [burdening and analysis of the effects of improvement measures. Project presentation.](#)
23 [Slovenian Environment Agency \(http://www.arso.gov.si/en/\), June, 2014.](#)

24 Schell, B., Ackermann, I.J., Hass, H., Binkowski, F.S., Ebel, A.: Modeling the formation of
25 secondary organic aerosol within a comprehensive air quality model system, *Journal of*
26 *Geophysical Research*, 106, 28275-28293, 2001.

27 Shaw, W.J., Allwine, K, Fritz, B.G., Rutz, F.C., Rishel, J.P., Chapman, E.G.: An evaluation
28 of the wind erosion module in DUSTAN. *Atmospheric Environment*, 42, 1907–1921, 2008.

29 Skamarock, W.C., Klemp, J.B., Dudhia, J., Gill, D.O., Barker, D.M., Duda, M.G., Huang,
30 X.Y., Wang, W., Powers, J.G.: A Description of the Advanced Research WRF Version 3.
31 NCAR Technical Note, NCAR/TN-475pSTR, 113 pp, 2008.

1 Stockwell, W. R., Middleton, P., Chang, J. S., and Tang, X.: The second generation regional
2 acid deposition model chemical mechanism for regional air quality modeling. *J. Geophys.*
3 *Res.*, 95, 16343-16367, 1990.

4 Sudo, K., Takahashi, M., Kurokawa, J., and Akimoto, H.: CHASER: A global chemical
5 model of the troposphere 1. Model description. *J. Geophys. Res.*, 107,
6 doi:10.1029/2001JD001113, 2002.

7 Takigawa, M., Niwano, M., Akimoto, H., Takahashi, M.: Development of a One-way Nested
8 Global-regional Air Quality Forecasting Model. *SOLA*, 2007, Vol. 3, 081-084, 2007.

9 Tulet P., Crassier V., Solmon F., Guedalia D., Rosset R.: Description of the mesoscale
10 nonhydrostatic chemistry model and application to a transboundary pollution episode between
11 northern France and southern England. *J Geophys Res* 108(D1):4021, 2003.

12 US Environmental Protection Agency (US EPA): Guidance for regulatory application of the
13 Urban Airshed Model. EPA-450/4-91-013, July 1991, United States Environmental Protection
14 Agency, Research Triangle Park, NC 27711, 1991.

15 Taylor, K.E.: Summarizing multiple aspects of model performance in a single diagram. *J.*
16 *Geophys. Res.*, 106, 7183-7192, 2001.

17 Tie, X., Geng, F.H., Peng, L., Gao, W., Zhao, C.S.: Measurement and modeling of O₃
18 variability in Shanghai, China; application of the WRF-Chem model. *Atmospheric*
19 *Environment*, 43, 4289-4302, 2009.

20 Tong, D.Q., Mauzerall, D.L.: Spatial variability of summertime tropospheric ozone over the
21 continental United States: Implications of an evaluation of the CMAQ model. *Atmospheric*
22 *Environment*, 40, 3041–3056, 2006.

23 van Loon M., Roemer M.G.M., Builtjes P.J.H., Bessagnet B., Rouil L., Christensen J.H.,
24 Brandt J., Fagerli H., Tarrason L., Rodgers I.: Model inter-comparison in the framework of
25 the review of the unified EMEP model. Technical report R2004/282, TNO, 2004.

26 Vlachogianni, A., Kassomenos, P., Karppinen, A., Karakitsios, S., Kukkonen, J.: Evaluation
27 of a multiple regression model for the forecasting of the concentrations of NO_x and PM₁₀ in
28 Athens and Helsinki. *Science of the Total Environment*, 409, 1559–1571, 2011.

29 Vogel, B., Vogel, H., Bäumer, D., Bangert, M., Lundgren, K., Rinke, R., Stanelle, T.: The
30 comprehensive model system COSMO-ART – Radiative impact of aerosol on the state of the

1 atmosphere on the regional scale, *Atmos. Chem. Phys.*, 9, 8661–8680, doi:10.5194/acp-9-
2 8661-2009, 2009.

3 Wolff, G.T., Lioy, P.J.: An empirical model for forecasting maximum daily ozone levels in
4 the northeastern United States. *J. Air Pollut. Control Assoc.*, 28, 1034-1038, 1978.

5 Yahya, K., Zhang, Y., Vukovich, J.M.: Real-time air quality forecasting over the southeastern
6 United States using WRF/Chem-MADRID: Multiple-year assessment and sensitivity studies,
7 92, 318–338, 2014.

8 Yang, Q., Gustafson Jr., W. I., Fast, J. D., H. Wang, H., Easter, R. C., Morrison, H.:
9 Assessing regional scale predictions of aerosols, marine stratocumulus, and their interactions
10 during VOCALS-REx using WRF-Chem. *Atmospheric Chemistry and Physics*, 11, 11951–
11 11975, doi:10.5194/acpd-11-22663-2011, 2011.

12 Zhang, Y.: Online-coupled meteorology and chemistry models: history, current status, and
13 outlook. *Atmos. Chem. Phys.*, 8, 2895–2932, 2008.

14 Zhang, K., Wan, H., Wang, B., Zhang, M., Feichter, J., Liu, X.: Tropospheric aerosol size
15 distributions simulated by three online global aerosol models using the M7 microphysics
16 module. *Atmos. Chem. Phys.*, 10, 6409-6434, 2010.

17 Zhang, Y., Pan., Y., Wang, K., Fast, J.D., Grell, G.A.: WRF-Chem-MADRID: incorporation
18 of an aerosol module into WRF-Chem and its initial application to the TexAQS2000 episode.
19 *J. Geophys. Res.*, 115, D18202, 2010a.

20 Zhang, Y., Wen, X.-Y., Jang, C.J.: Simulating climate-chemistry-aerosol-cloud radiation
21 feedbacks in continental U.S. using online-coupled WRF-Chem. *Atmospheric Environment*,
22 44, 3568-3582, 2010b.

23 Zhang, Y., Bocquet, M., Mallet, V., Seigneur, C., Baklanov, A.: Real-time air quality
24 forecasting, part I: History, techniques, and current status. *Atmospheric Environment*, 60,
25 632-665, 2012a.

26 Zhang, Y., Bocquet, M., Mallet, V., Seigneur, C., Baklanov, A.: Real-time air quality
27 forecasting, part II: State of the science, current research needs, and future prospects.
28 *Atmospheric Environment*, 60, 656-676, 2012b.

29 Žabkar, R., Rakovec, J., Gaberšek, S.: A trajectory analysis of summertime ozone pollution in
30 Slovenia. *Geofizika*, 25, 179-202, 2008.

1 Žabkar, R., Rakovec, J., Koračin, D.: The roles of regional accumulation and advection of
2 ozone during high ozone episodes in Slovenia: a WRF-Chem modelling study. Atmospheric
3 Environment, 45, 1192-1202, 2011a.

4 Žabkar, R.: Nadgradnja modela statističnega napovedovanja ozona s predhodnim
5 razvrščanjem trajektorij v skupine, final report. Available online:
6 <http://www.arso.gov.si/zrak/kakovost%20zraka/poro%C4%8Dila%20in%20publikacije/poro>
7 [%C4%8Dila%20o%20projektih/Porocilo_2011%20napoved_ozona.pdf](http://www.arso.gov.si/zrak/kakovost%20zraka/poro%C4%8Dila%20in%20publikacije/poro%C4%8Dila%20o%20projektih/Porocilo_2011%20napoved_ozona.pdf), 2011b.

8 Žabkar, R., Koračin, D., Rakovec, J.: A WRF-Chem sensitivity study using ensemble
9 modelling for a high ozone episode in Slovenia and the Northern Adriatic area. Atmospheric
10 Environment, 77, 990-1004, 2013.

11

1 Table 1: AQ monitoring sites.

Monitoring site	Abbreviation	Type of zone	Altitude (m)	<u>Model orography (m)</u>	<u>Model analysis height (m)</u>	Pollutants
Celje	CE	Urban	240	<u>300</u>	<u>313</u>	O ₃ , PM10, NO ₂
Hrastnik	HRA	Urban	290	<u>540</u>	<u>552</u>	O ₃ , SO ₂
Iskrba	ISK	Rural	540	<u>579</u>	<u>591</u>	O ₃ , NO ₂
Koper	KOP	Urban	56	<u>72</u>	<u>85</u>	O ₃ , PM10
Kovk	KOV	Rural	608	<u>516</u>	<u>528</u>	NO ₂
Krvavec	KRV	Rural	1740	<u>1272</u>	<u>1414</u>	O ₃
Ljubljana	LJ	Urban	299	<u>287</u>	<u>300</u>	O ₃ , PM10, NO ₂ ,
Murska Sobota	MS	Rural	188	<u>189</u>	<u>202</u>	O ₃ , PM10, NO ₂
Nova Gorica	NG	Urban	113	<u>150</u>	<u>163</u>	O ₃ , PM10, NO ₂
Otlica	OTL	Rural	918	<u>874</u>	<u>886</u>	O ₃
Sv. Mohor	MOH	Rural	394	<u>254</u>	<u>266</u>	NO ₂
Trbovlje	TRB	Suburban	250	<u>459</u>	<u>471</u>	O ₃ , PM10, NO ₂
Velenje	VEL	Urban	389	<u>461</u>	<u>474</u>	O ₃ , SO ₂
Vnajnarje	VNA	Rural	630	<u>468</u>	<u>480</u>	NO ₂
Zadobrova	ZAD	Rural	280	<u>275</u>	<u>287</u>	PM10, NO ₂
Zagorje	ZAG	Urban	241	<u>431</u>	<u>443</u>	O ₃ , PM10, NO ₂
Zavodnje	ZAV	Rural	765	<u>678</u>	<u>690</u>	O ₃ , NO ₂

2

1 Table 2: Statistical scores for 1-hour values of 2m temperature (T2m), 10 m wind speed
2 (W10m) and relative humidity (RH), and for daily average incoming solar radiation (SR).
3 Shown are results for 1-day forecast, calculated separately for three measuring sites (LJ, NG,
4 MS) and for 24 MET monitoring stations (ALL) during the 3-month period. In the case of
5 temperature results for daily maxima are also shown.

Variable	Station	NoCases	Mean	ME	MAE	RMSE	CORR
T2m 1h (°C)	LJ	2129	20.3	-1.6	2.3	2.9	0.91
	NG	2184	21.8	-1.1	2.1	2.5	0.94
	MS	2184	19.2	-2	2.3	2.8	0.95
	ALL	47836	18.7	-1.3	2.3	2.9	0.93
T2m max (°C)	LJ	89	26.5	-1.6	1.8	2.1	0.98
	NG	90	26.8	-3	3	3.3	0.96
	MS	90	26.2	-1.7	1.8	2	0.98
	ALL	1976	24.2	-2.1	2.7	3.2	0.97
W10m (m/s)	LJ	2129	1.5	0	0.7	1	0.58
	NG	2183	2.7	1	1.4	1.9	0.35
	MS	2184	2.3	0.4	1.1	1.4	0.53
	ALL	43378	2.4	0.8	1.4	1.9	0.36
RH (%)	LJ	2066	62	-2	8	10	0.85
	NG	2121	62	-1	12	15	0.75
	MS	2121	69	3	8	11	0.88
	ALL	48556	68	2	11	14	0.77
SR (W/m ²)	LJ	90	276	19	31	43	0.84
	NG	90	278	4	32	43	0.77
	MS	90	273	15	26	37	0.9
	ALL	1710	273	16	35	49	0.77

1 Table 3: Domain wide performance statistics for 1-day and 2-day forecast in μgm^{-3} . For
 2 different pollutants statistics for all hourly (hour), 8-hour averages (8h), 8-hour daily
 3 maximum (8h max), daily maximum (max) or daily average (day) concentrations are shown.

		NoCases	Mean	ME	MAE	RMSE	CORR
O ₃ (hour)	1 day	28391	94. <u>8</u>	14.53.7	25. <u>15</u>	32. <u>51</u>	0.65
	2 day	28391	9 <u>5.04.2</u>	14.53.8	25. <u>58</u>	32. <u>59</u>	0.64
O ₃ (8h)	1 day	28072	94. <u>81</u>	14.63.8	22. <u>69</u>	28. <u>15</u>	0.69
	2 day	28072	9 <u>5.04.2</u>	14.63.8	23. <u>03</u>	28. <u>59</u>	0.68
O ₃ (8h max)	1 day	1157	11 <u>1.50.7</u>	-0. <u>17</u>	13. <u>26</u>	1 <u>6.57</u>	0.77
	2 day	1157	11 <u>1.60.9</u>	-0. <u>21</u>	1 <u>3.74.1</u>	17. <u>04</u>	0.75
O ₃ (max)	1 day	1170	11 <u>6.55.8</u>	-2. <u>73.4</u>	13. <u>37</u>	1 <u>6.77.1</u>	0.81
	2 day	1170	11 <u>6.65.8</u>	-3. <u>19</u>	14. <u>04</u>	17. <u>59</u>	0.7 <u>89</u>
NO ₂ (hour)	1 day	26178	7.3	-5.1	7.5	10.8	0.3
	2 day	26178	7.5	-4.9	7.6	10.8	0.3
PM10 (day)	1 day	718	29. <u>0</u>	7.1	12. <u>0</u>	18.8	0.34
	2 day	718	29.1	7.2	12. <u>0</u>	19.1	0.37

4

1 Table 4: Discrete evaluation of 1-hour daily maximum ozone predictions.

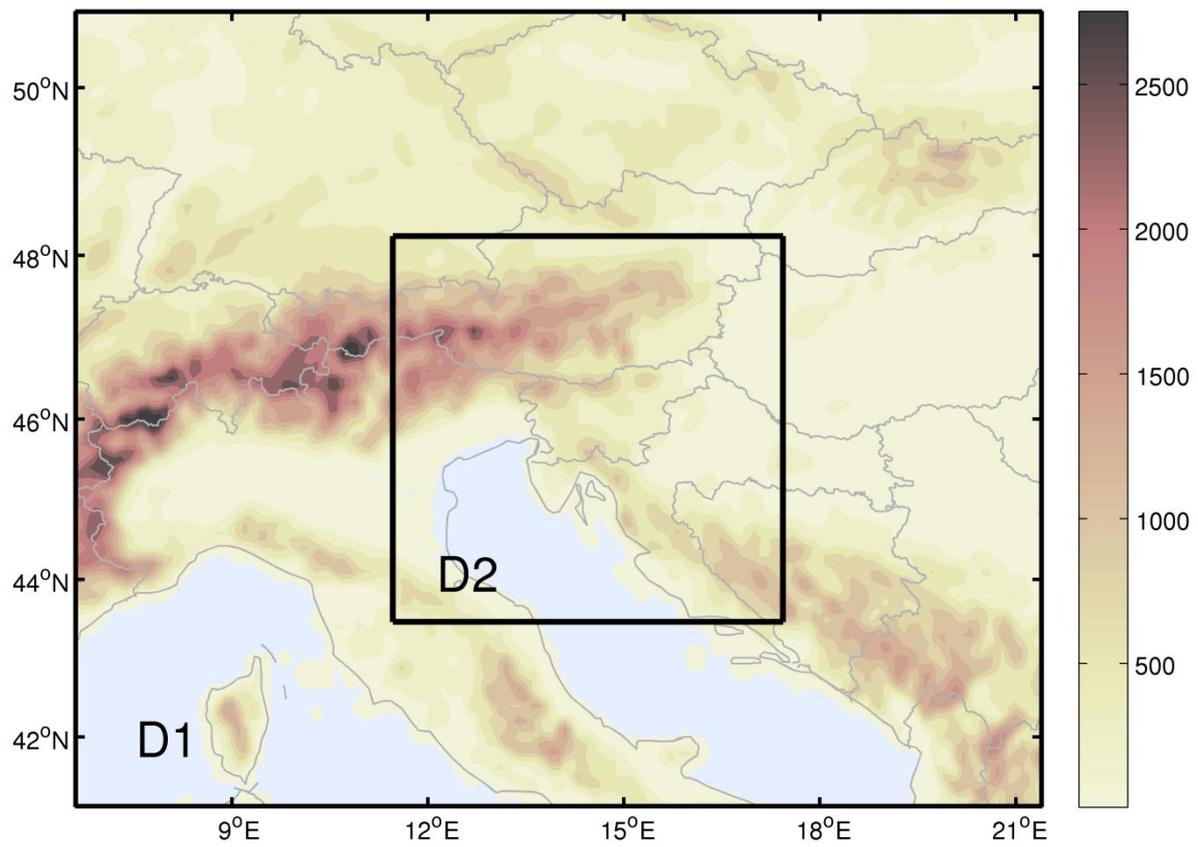
Stations	Threshold, NoCases	Forecast	Mean (μgm^{-3})	ME (μgm^{-3})	MAE (μgm^{-3})	RMSE (μgm^{-3})	CORR	MNBE (%)	MNGE (%)	IOA
All	> 0 1170	F 1day	11 56.85	-32.64	13.37	16.774	0.81	-0.05	11.720	0.86
		F 2day	11 65.68	-3.91	14.04	17.59	0.789	-0.71	12.36	0.84
		PER 1day	119.5	-0.4	15.8	21.1	0.65	1.6	14.5	0.81
		PER 2day	119.8	-0.4	21.7	27.7	0.39	2.8	19.6	0.65
	> 140 1102	F 1day	14 43.13	-11.29	15.27	178.94	0.52	-6.874	9.59	0.57
		F 2day	14 10.46	-134.86	167.51	1920.40	0.421	-8.691	10.58	0.487
		PER 1day	145.0	-10.2	15.6	19.6	0.41	-6.5	10.0	0.52
		PER 2day	135.8	-19.4	24.76	29.2	0.31	-12.4	15.9	0.38
Sub-alpine urban with SF (LJ, HRA)	> 0 180	F 1day	115.3	1.1	10.7	14.0	0.84	3.4	11.1	0.91
		F 2day	115.4	0.8	12.0	15.2	0.80	3.5	12.2	0.88
		PER 1day	114.3	-0.3	16.7	21.7	0.64	2.2	16.5	0.80
		PER 2day	114.6	-0.3	21.9	27.8	0.41	3.9	21.6	0.65
		SF 1day	114.0	-0.5	11.9	15.7	0.81	1.6	11.2	0.88
		SF 2day	116.2	0.6	13.4	17.1	0.75	3.2	12.7	0.84
Rural with SF (MS, ISK, KRV, OTL)	> 0 360	F 1day	117.652	-5.681	13.346	16.376	0.80	-35.0	10.817	0.865
		F 2day	117.448	-6.488	14.255	17.489	0.767	-35.4	11.424	0.841
		PER 1day	123.6	-0.3	15.0	20.7	0.65	1.4	13.1	0.81
		PER 2day	124.1	-0.4	21.6	27.8	0.37	2.4	18.5	0.64
		SF 1day	121.5	-2.9	15.0	19.4	0.74	-0.7	12.2	0.83
		SF 2day	122.9	-1.8	15.8	20.5	0.67	0.5	13.2	0.79
Mediterranean urban with SF (KOP, NG)	> 0 179	F 1day	123.5	-11.8	17.4	22.5	0.76	-6.9	12.5	0.80
		F 2day	124.5	-11.2	17.2	21.8	0.77	-6.5	12.4	0.82
		PER 1day	135.9	-0.5	17.4	23.0	0.68	1.2	13.8	0.83
		PER 2day	136.0	-0.2	25.2	31.5	0.41	2.8	19.7	0.66
		SF 1day	129.3	-7.0	15.9	20.7	0.75	-3.6	11.6	0.83
		SF 2day	131.6	-4.5	15.6	20.4	0.74	-1.6	11.6	0.84

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1 Table 5: Categorical evaluation of 1-hour daily maximum ozone predictions for different
 2 thresholds, calculated for 8 monitoring sites with available statistical forecast.

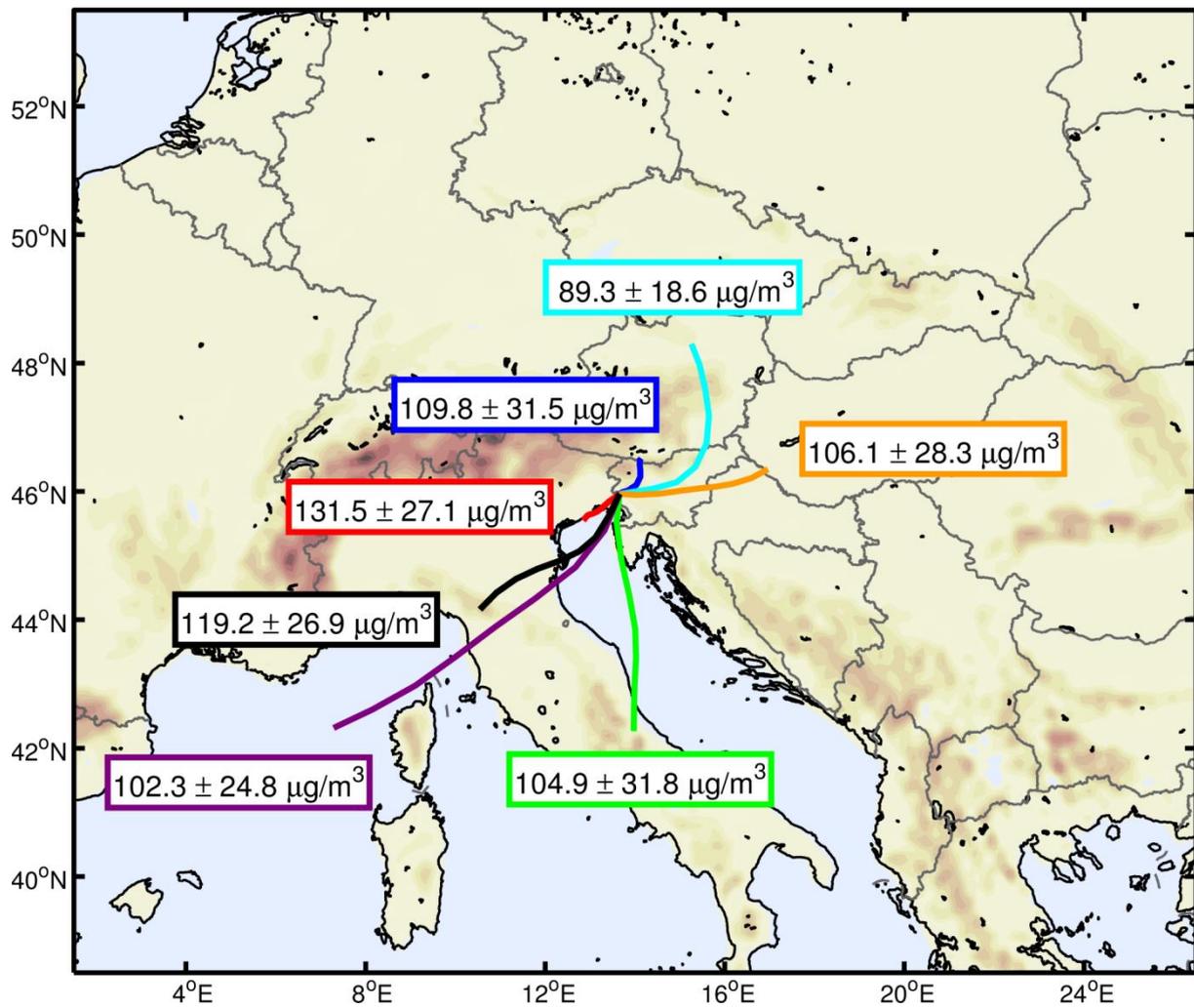
Threshold	Forecast	ETSA	CSI	B	FAR	POD	a	b	c	d
> 120	F 1day	0. 4278	0. 631	0. 8176	0.13	0. 7066	395	25339	3137	10721
	F 2day	0. 3977	0. 6159	0. 7975	0.14	0. 684	4137	24533	3037	11527
	PER 1day	0. 3174	0.59	0.99	0.25	0.74	91	267	249	93
	PER 2day	0. 1764	0.49	1.00	0.34	0.65	123	235	209	124
	SF 1day	0. 4280	0.67	1.02	0.21	0.81	67	257	243	61
	SF 2day	0. 3877	0.65	1.03	0.23	0.80	77	264	225	66
	> 140	F 1day	0. 4084	0. 5047	0. 6459	0.1 54	0.5 51	197	11103	4902
F 2day		0. 3782	0.4 74	0.6 60	0.1 98	0.5 349	252	10899	4769	95104
PER 1day		0. 4082	0.53	1.00	0.31	0.69	62	141	435	62
PER 2day		0. 1972	0.35	1.00	0.48	0.52	97	106	391	97
SF 1day		0. 3079	0.43	0.73	0.29	0.52	40	99	398	91
SF 2day		0. 3079	0.43	0.70	0.27	0.51	37	98	403	94
> 160		F 1day	0. 1991	0.22	0.3 87	0.3 42	0.25	910	19	6267
	F 2day	0. 1791	0.2 01	0.3 40	0.3 526	0.22	96	17	61922	59
	PER 1day	0. 4092	0.45	1.00	0.38	0.62	29	47	595	29
	PER 2day	0. 2288	0.28	1.00	0.56	0.43	43	33	572	43
	SF 1day	0. 2390	0.27	0.49	0.35	0.32	13	24	539	52
	SF 2day	0. 2590	0.29	0.63	0.41	0.37	19	27	540	46



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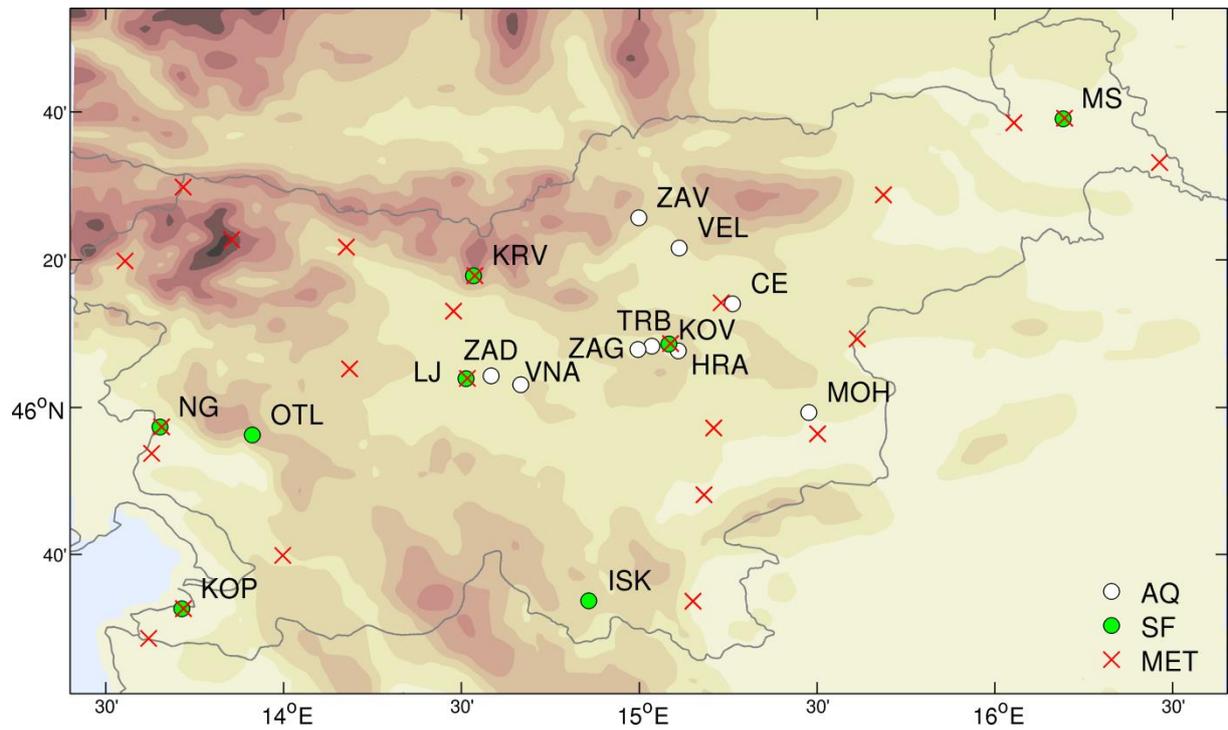
2 Figure 1: Modelling domains (D1, D2) used in WRF-Chem RT-AQF system. Orography (in
3 meters) is shown in resolution of D1 domain (11.1 km).

4



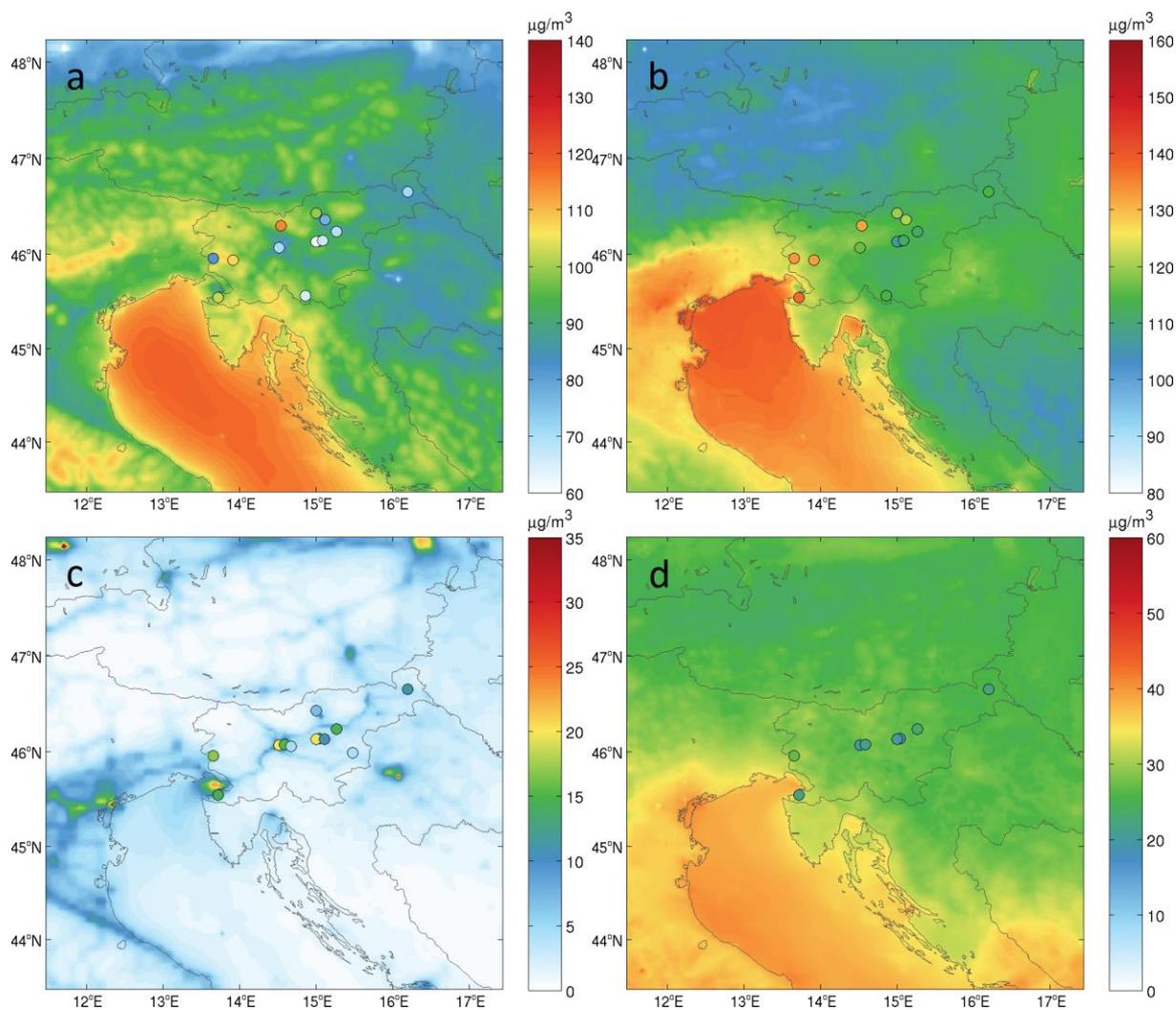
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Figure 2: Example of ozone analysis for the Nova Gorica (NG) monitoring site (average daily maximum \pm standard deviation) for 7 clusters of similar trajectories, as used in the statistical ozone daily maximum forecast for the NG station.

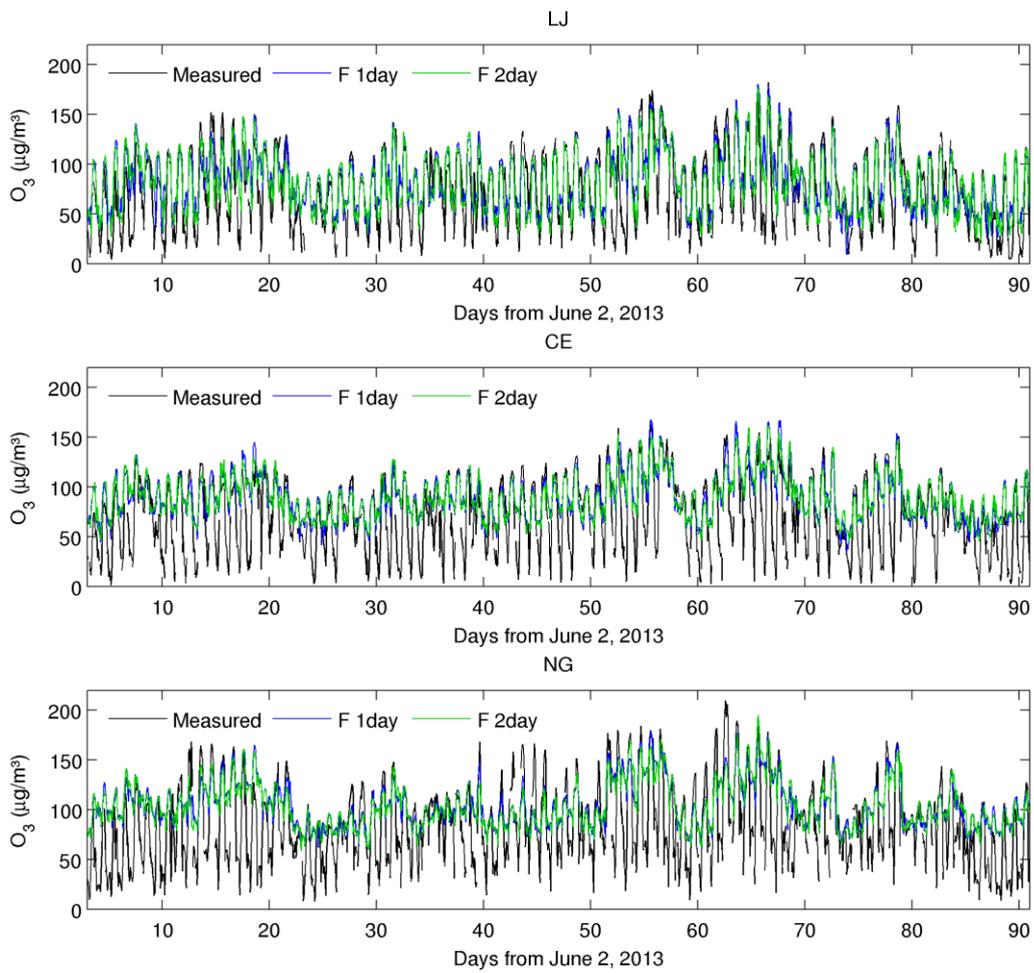


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Figure 3: Locations of monitoring stations used in evaluation of air quality variables (AQ stations; shown are also station abbreviations) and meteorological variables (MET stations). Green dots indicate measuring sites with available ozone daily maximum statistical forecast (SF). For the meaning of abbreviations of AQ sites see Tab. 1.



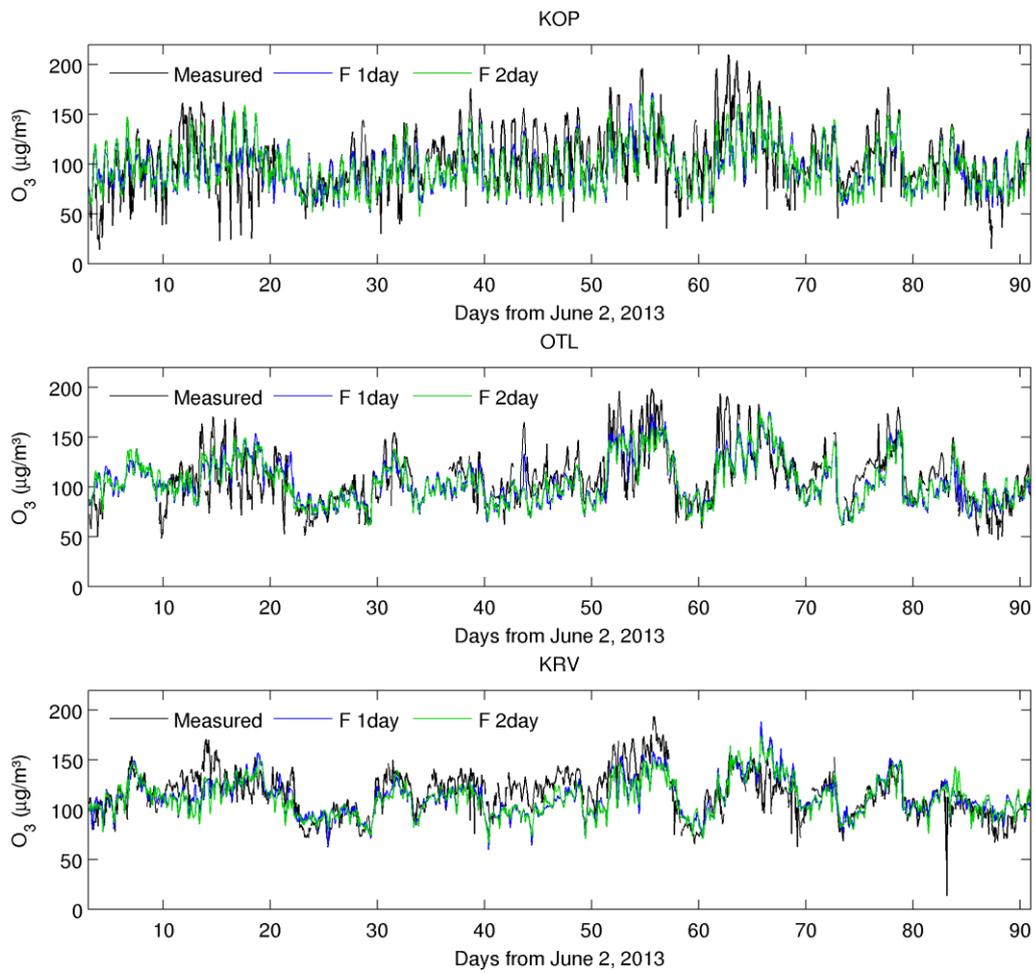
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 2 Figure 4: 3-month average 1-day predictions of a) hourly O_3 , b) O_3 daily maximum, c) hourly
 3 NO_2 , and d) daily PM_{10} concentrations for the first model layer, overlaid with measurements.
 4



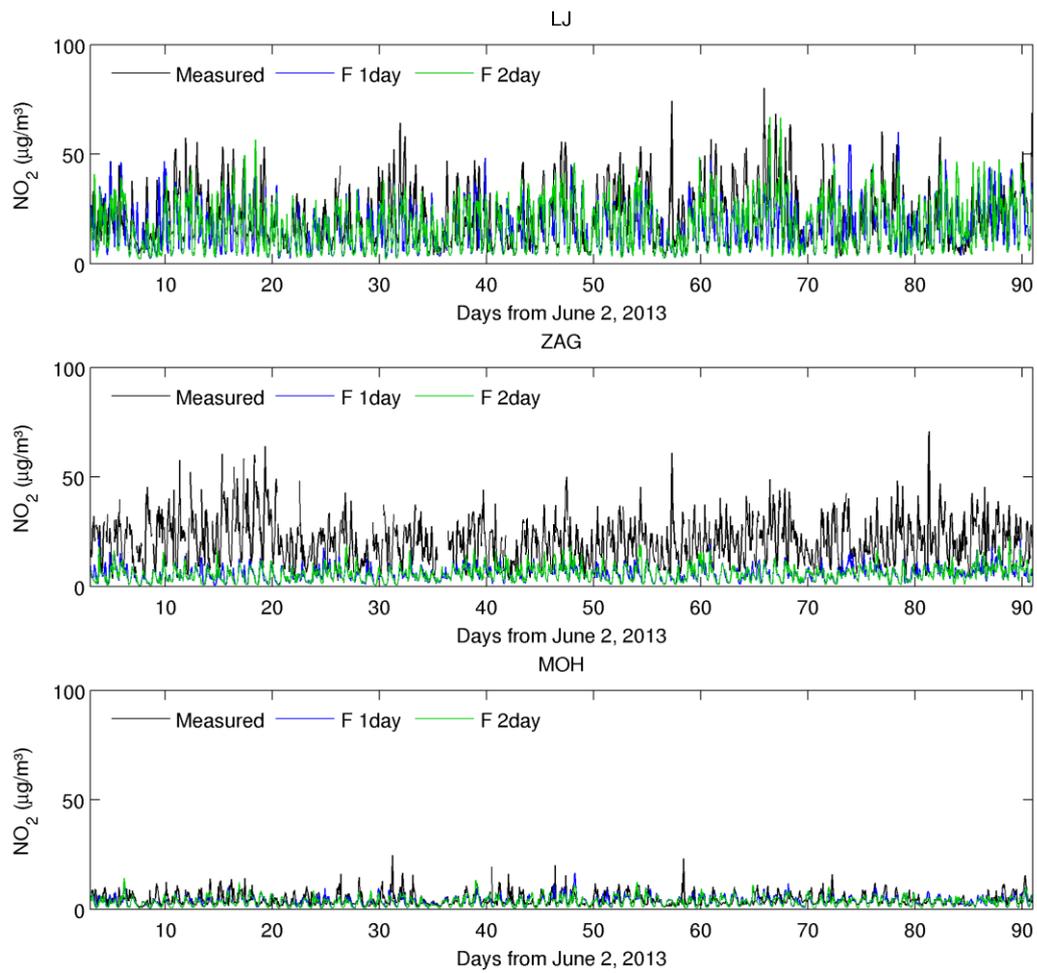
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2 Figure 5: Time evolution of hourly ozone concentrations for 1-day (F 1day) and 2-day (F
 3 2day) WRF-Chem predictions and measurements for some stations during the 3-month
 4 period. (continued)

5

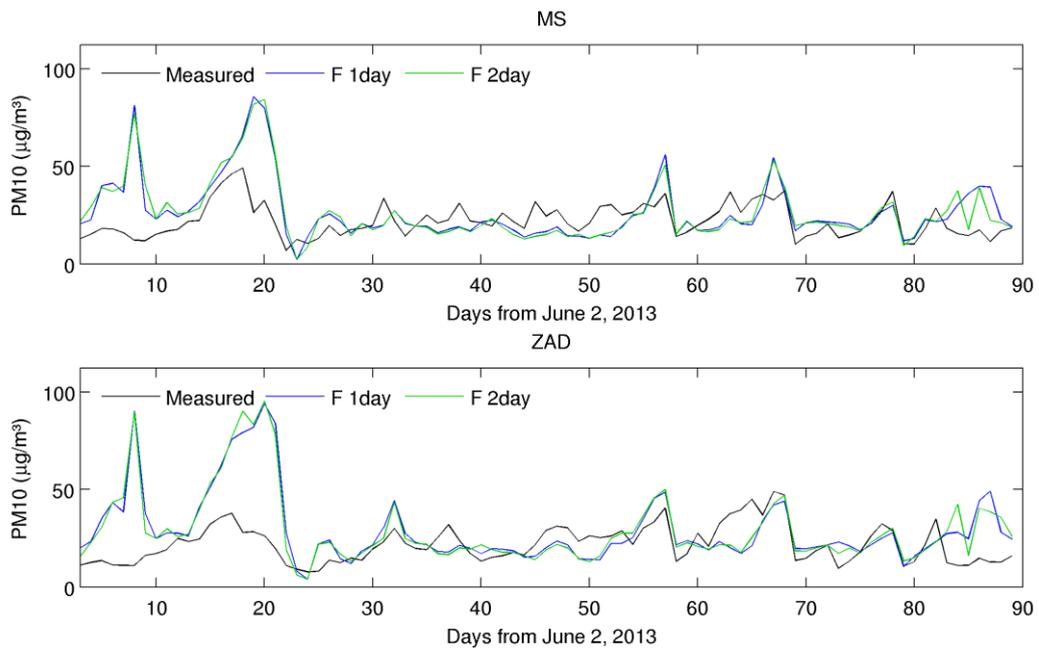


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2 Figure 5: (continued)



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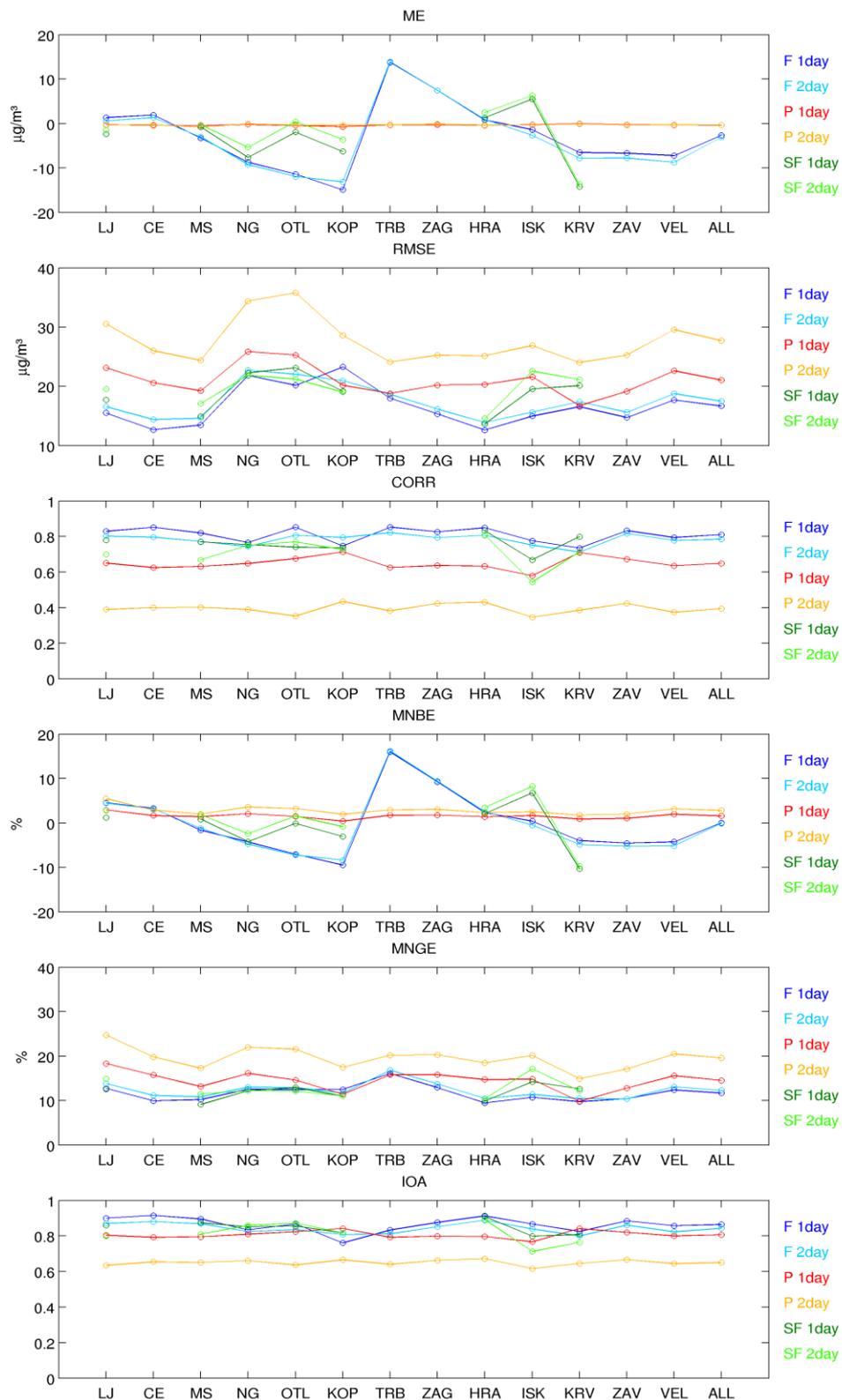
2 Figure 6: The same as Fig. 5 but for NO₂ at LJ, ZAG and MOH stations.



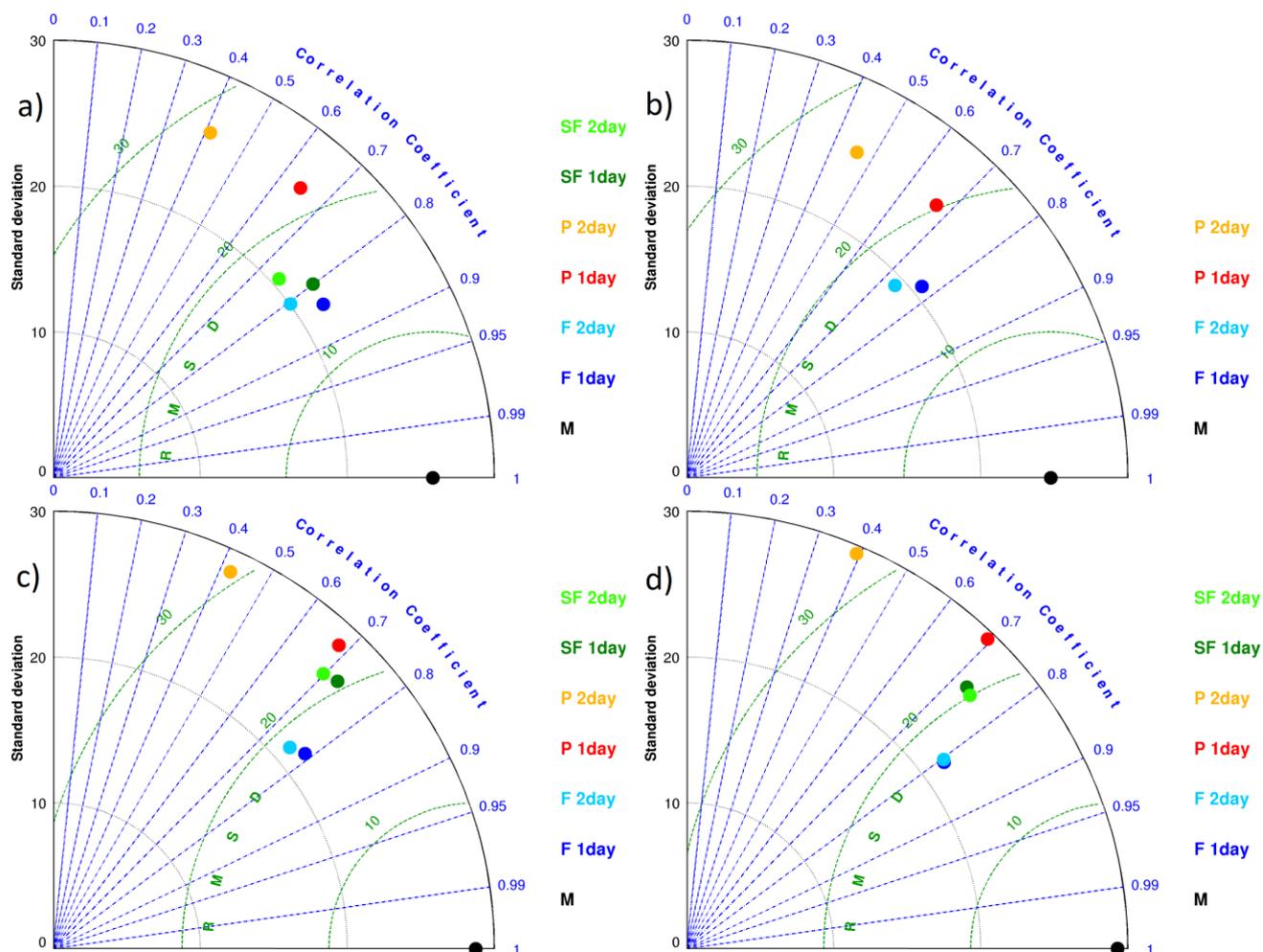
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2 Figure 7: The same as Fig. 5, but for daily PM10 concentrations at MS and ZAD stations.

3



1
 2 Figure 8: Site-by-site comparison of discrete statistics for 1-day and 2-day WRF-Chem (F
 3 1day, F 2day), statistical (SF 1day, SF 2 day) and persistence model (P 1day, P 2day)
 4 predictions of ozone daily maxima during the 3 analyzed summer months.
 5



1
 2 Figure 9: Taylor diagrams comparing 1-day and 2-day ozone daily maximum statistical
 3 forecast (SF), persistence (P) and WRF-Chem forecast (F) for a) sub-alpine urban stations
 4 with SF (LJ, HRA), b) sub-alpine urban stations without SF (CE, TRB, ZAG, VEL), c) rural
 5 stations with SF (MS, ISK, KRV, OTL) and d) Mediterranean urban stations (NG, KOP).