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APPLICATION OF WRF-CHEM TO FORECAST PM₁₀ CONCENTRATIONS OVER POLAND

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INTRODUCTION

In this study we apply the on-line model WRF-Chem version 3.5 (Grell et al. 2005; Skamarock & Klemp 2008) to forecast PM10 concentration over Poland, with a focus on the south-west part of the country called the Lower Silesia region. The PM10 forecasts are tested during the winter 2014 from 1st January to 28th February, as winters in general are favourable in Poland to high particulate matter concentrations due to both high coal consumption, which is commonly used fuel e.g. for heating houses, and meteorological conditions. The forecasts are evaluated by comparison with observations gathered by the Voivodeship Inspectorate of Environmental Protection in Poland.

DATA AND METHODS

The WRF-Chem model setup

WRF-Chem is used in nested mode with a summary of the model configuration in Table 1 and Fig. 1. The simulations are driven by the GFS meteorological data, available every 3h, with 0.5° x 0.5° spatial resolution. Emissions are the TNO MACC II data set with 1/8° x 1/16° spatial resolution (Pouliot et al. 2012). The first 48-h forecasting cycle on the 01 January 2014 uses a 2-week spin-up, with the model simulations started the 15th of December using the GFS-FNL meteorology for initial and boundary conditions. From the 2nd of January, the model uses chemistry cycling, and the WRF-Chem run for the previous day is used to initialize the next day's forecasting simulation. Temporal variations in emissions are restricted to emissions from nature, while the TNO MACC II emissions are assumed constant during the entire simulation.

Model evaluation

The PM₁₀ concentrations forecasts were compared with daily mean observation

Table 1. Model configuration

Category	Model setup	
Forecasts period	01st January – 28th February 2014	60°N —
Domains	Europe (36 km) – Poland (12 km) – SW Poland (4 km)	
Vertical resolution	35 layers	
PBL process	YSU	55°N —
Land-surface process	NOAH	
Cumulus	Kain-Fritch for d1 and d2	
Shortwave & Longwave radiation	RRTMG	50°N —
Microphysics	Lin	
Gas-phase mechanism	RADM2	45°N —
Aerosol model	MADE/SORGAM	
Photolysis scheme	Fast-J	
Wet deposition	Simplified parameterisation for wet scavenging	40°N —

Table 2. Model performance for the 24h and 48h lead time

Forecast range	Ν	FAC2	MB	NMB	RMSE
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gathered by the Voivodeship Inspectorate of Environmental Protection, from sites in Lower Silesia region in SW Poland (Table 2), separately for 24h and 48h lead time (Table 2). Time series and scatter plots for the two stations with the lowest MB and one station with the highest MB are given as examples (Fig. 3).

24h	977	0.67	-16.17	-0.38	29.17
48h	977	0.67	-16.38	-0.38	29.51



35°N



48h

CONCLUSION

Figure 3. Time series of PM10 concentration for the 48h lead time of the forecasts for Kłodzko, Działoszyn and Nowa Ruda. Scatter plots for these stations, presenting both 24h and 48h lead time; bosplots for 48h lead time.

We have found that WRF-Chem in general captures the variability in observed PM10 concentrations for most of the stations. However the highest observed peaks are underestimated by the model. In fact, the lowest performance was obtained for the Nowa Ruda station, which is located in a deep valley. This area has a high contribution of the emissions from coal fired residential heating. Such circumstances could cause high PM₁₀ observed concentrations peak during certain weather types such as winter time inversions. We argue that a higher resolution sector based emission data and temporal emission profile will be helpful for this analysis in connection with a focus on PBL processes in WRF-Chem and their impact on the initial distribution of emissions.

REFERENCES

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Grell, G. a., Peckham, S. E., Schmitz, R., McKeen, S. a., Frost, G., Skamarock, W. C., & Eder, B., 2005: Fully coupled "online" chemistry within the WRF model. Atmospheric Environment, 39(37), 6957-6975. doi:10.1016/j.atmosenv.2005.04.027. Pouliot, G., Pierce, T., Denier van der Gon, H., Schaap, M., Moran, M., & Nopmongcol, U., 2012: Comparing emission inventories and model-ready emission datasets between Europe and North America for the AQMEII project. Atmospheric Environment, 53, 4-14. doi:10.1016/j.atmosenv.2011.12.041

Skamarock, W. C., & Klemp, J. B., 2008: A time-split nonhydrostatic atmospheric model for weather research and forecasting applications. Journal of Computational Physics, 227, 3465–3485. doi:10.1016/j.jcp.2007.01.037.

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