

## Introduction

Background air measurements of persistent organic pollutants (POPs) at many sites occurs only at fixed intervals (e.g. one day per week) without any a priori consideration of air mass transport (i.e., whether the air is likely to be polluted or not).

The intermittent sampling approach may miss key long-range atmospheric transport (LRAT) episodes, which are often associated with the highest POP concentrations<sup>1</sup>.

## Objectives

- I. To develop a forecast system using the FLEXPART model to predict long-range atmospheric transport episodes of POPs using PCB-28 as a model compound,
- II. To evaluate the capability of the forecast system to capture specific LRAT episodes at a background site in southern Norway (Birkenes) through targeted sampling (i.e. when LRAT episodes are predicted),
- III. To assess whether predicted LRAT episodes for PCB-28 coincide with elevated concentrations of additional PCBs, and
- IV. To identify source regions of PCBs during individual episodes.

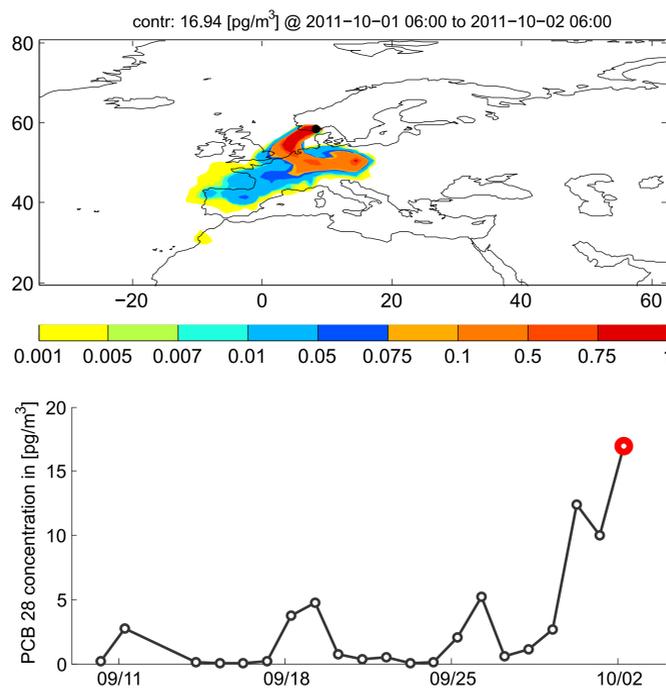


Figure 1: Forecasts of predicted concentrations of PCB-28 at Birkenes were made one day ahead using FLEXPART. Example shows predictions for October 1<sup>st</sup>, 2011.

## Results and Discussion

- Forecasts were made on day ahead to decide whether targeted samples should be collected during suspected LRAT events (Figure 1)
- Three predicted LRAT episodes (E) in 2011, which occurred in January (E1), February (E2) and late September/early October (E3a,b,c,d,e), were analyzed (Table 1).
- Measured concentrations of PCBs in all targeted samples (N=7) were above the 75th percentile of the concentrations obtained from the regular monitoring program (N=52) and included the highest measured values of all samples (Figure 2).
- A retrospective evaluation of the episodes with highest measured concentrations of PCB-28 in 2011 provides information on source regions (Figure 3).

## Conclusions

- This study most likely represent the first attempt to both (i) use model predictions driven by a priori information on emissions of POPs to trigger air sampling as well as (ii) retrospectively evaluate the source regions for measurements collected during predicted episodes.
- Observations targeted at strong pollution episodes (as in this study) or on transport from specific source regions with highly uncertain emissions (as could be done in a very similar forecasting framework) could significantly enhance our understanding of POP sources.
- For details, see Halse et al.<sup>5</sup>

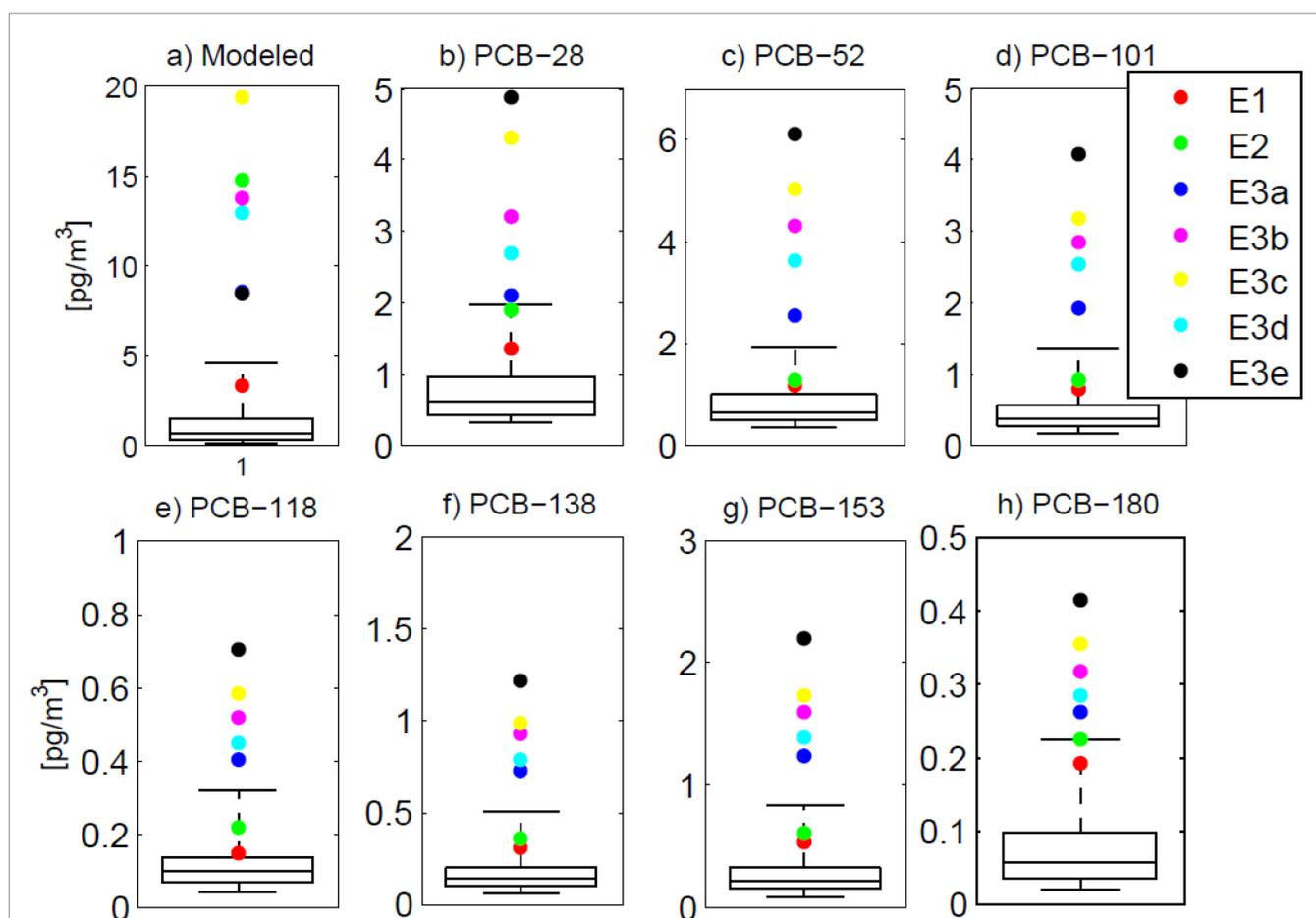


Figure 2: Modeled PCB-28 (a) and measured PCB concentrations (b-h) in units of  $\text{pg}/\text{m}^3$ , for both the annual sampling program and the targeted samples. The box and whisker plots show the annual results (2011) for Birkenes, based on weekly samples (N=52). The line shows the median, while the box and whiskers delineates the 25 and 75 percentiles and the 5 and 95 percentiles, respectively. Targeted samples (N=7) are represented by colored dots (see Table 1).

## Methods

Sampling was carried out at the Birkenes observatory in 2011, a background station located in southern Norway (N 58°23, E 08°15). Regular monitoring samples were collected over 24 h once per week (N=52). Targeted LRAT samples were collected over 12 to 25 hours.

The atmospheric transport of PCB-28 was simulated by use of the Lagrangian particle dispersion model FLEXPART<sup>2-3</sup> which can be operated in forward mode (for forecasting) and backward mode (for retrospective evaluation of LRAT episodes). PCB-28 was chosen as our model compound, following an earlier study for Birkenes<sup>4</sup>.

Table 1: Sampling times for targeted samples collected during predicted episodes in 2011.

| Episode | Start date | Start time | End date | End time |
|---------|------------|------------|----------|----------|
| E1      | 06.01      | 11:50      | 07.01    | 23:49    |
| E2      | 24.02      | 17:29      | 25.02    | 05:49    |
| E3a     | 29.09      | 08:05      | 29.09    | 21:10    |
| E3b     | 29.09      | 21:15      | 30.09    | 15:30    |
| E3c     | 30.09      | 15:30      | 01.10    | 05:35    |
| E3d     | 01.10      | 05:38      | 02.10    | 07:04    |
| E3e     | 02.10      | 07:08      | 02.10    | 18:49    |

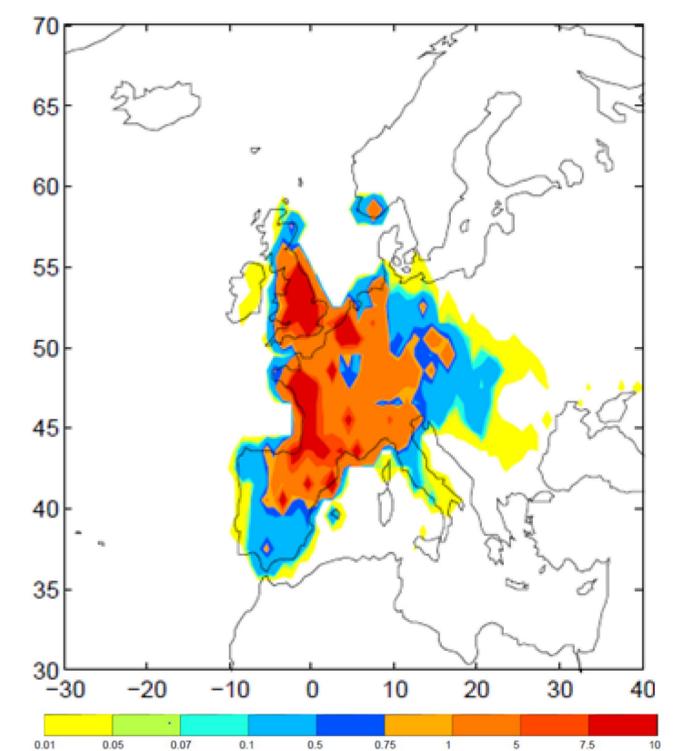


Figure 3: Map of EC (emission contributions,  $1\text{E}-12$   $[\text{pg}/\text{m}^3]$ ) for PCB-28 for the episode with highest measured concentrations at Birkenes during 2011 (E3e).

## Acknowledgements

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## References

1. Y. Yao, T. Harner, J. M. Ma, L. Tuduri and P. Blanchard, *Environ. Sci. Technol.*, 2007, **41**, 688-694.
2. A. Stohl, M. Hittenberger and G. Wotawa, *Atmospheric Environment*, 1998, **32**, 4245-4264.
3. A. Stohl, C. Forster, A. Frank, P. Seibert and G. Wotawa, *Atmospheric Chemistry and Physics*, 2005, **5**, 2461-2474.
4. S. Eckhardt, K. Breivik, Y. F. Li, S. Manø and A. Stohl, *Atmospheric Chemistry and Physics*, 2009, **9**, 6597-6610.
5. A. K. Halse, S. Eckhardt, M. Schlabach, A. Stohl and K. Breivik, *Atmospheric Environment*, 2013, **71**, 335-339.