# Evaluation of PM<sub>10</sub> and trace gas measurements with the MARGA during the HCCT 2010 campaign

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INTRODUCTION

During the Hill Cap Cloud Thuringia (HCCT) field campaign in September and October 2010 at Mt. Schmücke (Germany) the online system MARGA (Monitor for Aerosols & Gases in ambient Air [1,2]) was deployed at the upwind valley station in Goldlauter.

The instrument was developed by Metrohm Applikon, Netherlands and uses a combination of a wet-rotating denuder and a steam-jet-aerosol-collector to capture gases and particles in ambient air. During the HCCT campaign the MARGA was connected to a  $PM_{10}$  inlet-system to measure the concentrations of the inorganic components Cl<sup>-</sup>,  $NO_3^-$ ,  $SO_4^{2-}$ ,  $Na^+$ ,  $NH_4^+$ ,  $K^+$ ,  $Mg^{2+}$ ,  $Ca^{2+}$  in the particle phase and the corresponding trace gases HCl,  $HNO_2$ ,  $SO_2$ ,  $HNO_3$  and  $NH_3$  in a time resolution of one hour

## **OVERVIEW**

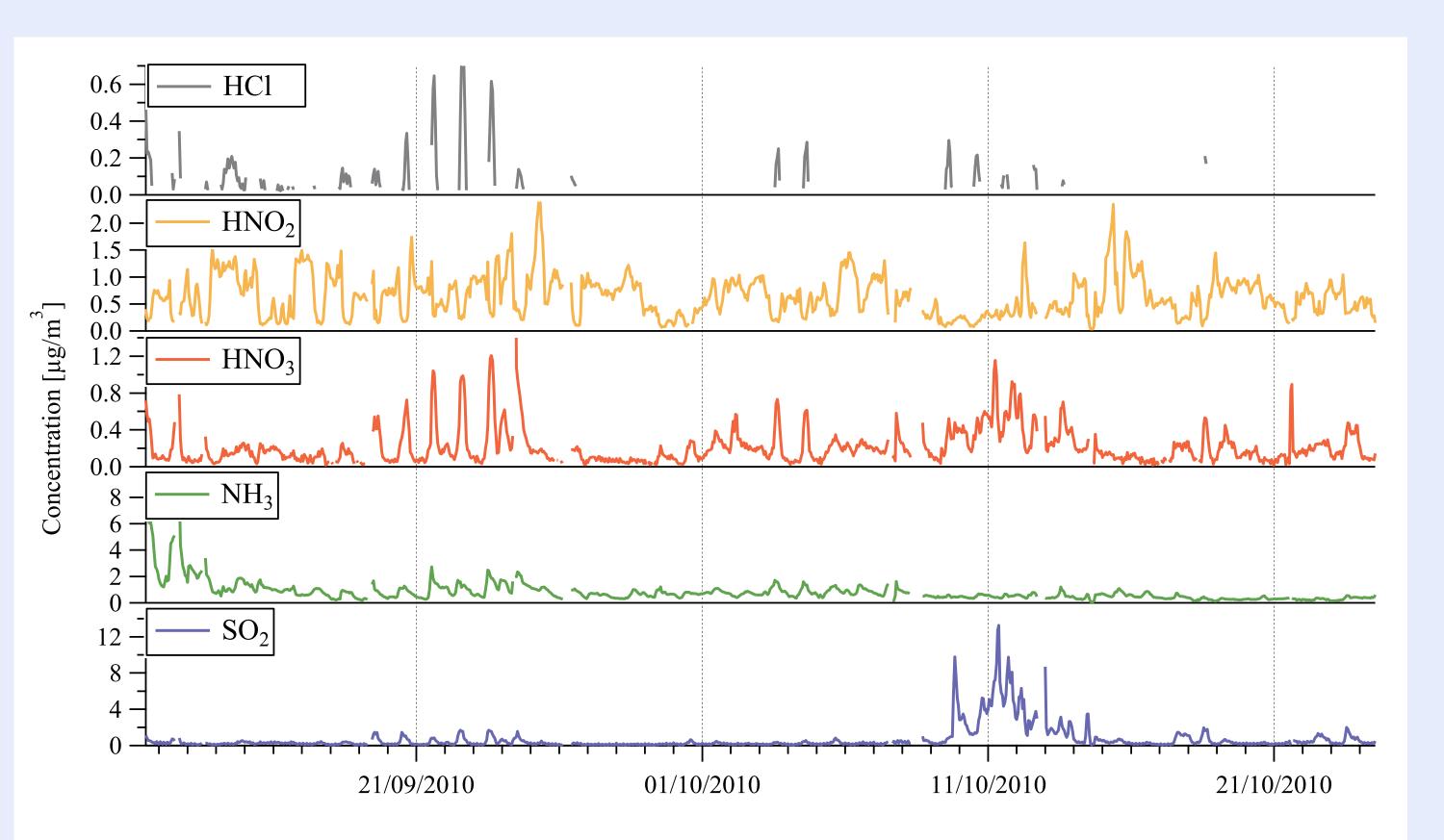
## **COMPARISON WITH AN AMS**

Table1 gives a summary of the measured particle and gas concentrations during HCCT in September and October 2010. Figures 1 and 2 show the time series of the main ions in the particle phase and the corresponding trace gases.

- Overall low gas phase concentrations, only in the mid of October high concentrations of SO<sub>2</sub> were observed due to the long range transport of continental air masses.
- Slightly higher particle concentrations in October
- Periods with high Cl<sup>-</sup> concentrations indicate the influence of maritime air masses, but most of the time the chloride concentration was below the detection limit of the MARGA

#### Table 1: Summary of the aerosol concentrations during HCCT.

	Mean [µg/m³]	Stdev [µg/m³]	Ν		Mean [µg/m³]	Stdev [µg/m³]	Ν
HCI	0.13	0.14	164	SO4 <sup>2-</sup>	1.38	0.8	1003
HNO <sub>2</sub>	0.64	0.39	999	Na <sup>+</sup>	0.12	0.16	922
SO <sub>2</sub>	0.67	1.34	1002	$NH_4^+$	1.12	3.27	1001
HNO <sub>3</sub>	0.23	0.62	989	K⁺	0.1	0.04	888
NH <sub>3</sub>	0.81	0.84	1000	Mg <sup>2+</sup>	0.14	0.15	981
Cl-	0.19	0.21	402	Ca <sup>2+</sup>	0.59	0.36	992
$NO_3^-$	1.94	2.08	1003				



The MARGA measurements were compared with a HR-ToF-AMS (High Resolution – Time of Flight - Aerosol Mass Spectrometer, Aerodyne, USA [3]) during the HCCT campaign:

- Despite the different size cutting of the two instruments (MARGA: PM<sub>10</sub>, AMS: PM<sub>1</sub>) a good agreement was found for NO<sub>3<sup>-</sup></sub>, NH<sub>4</sub><sup>+</sup> and SO<sub>4<sup>2-</sup></sub> (figure 3) -> NO<sub>3<sup>-</sup></sub>, NH<sub>4</sub><sup>+</sup> and SO<sub>4<sup>2-</sup></sub> mainly present in the submicron fraction of particles.
- Exception: In the first week of the campaign MARGA detects more NO<sub>3<sup>-</sup></sub> than AMS -> high NO<sub>3<sup>-</sup></sub> concentrations in the particle coarse mode due to the influence of maritime air masses

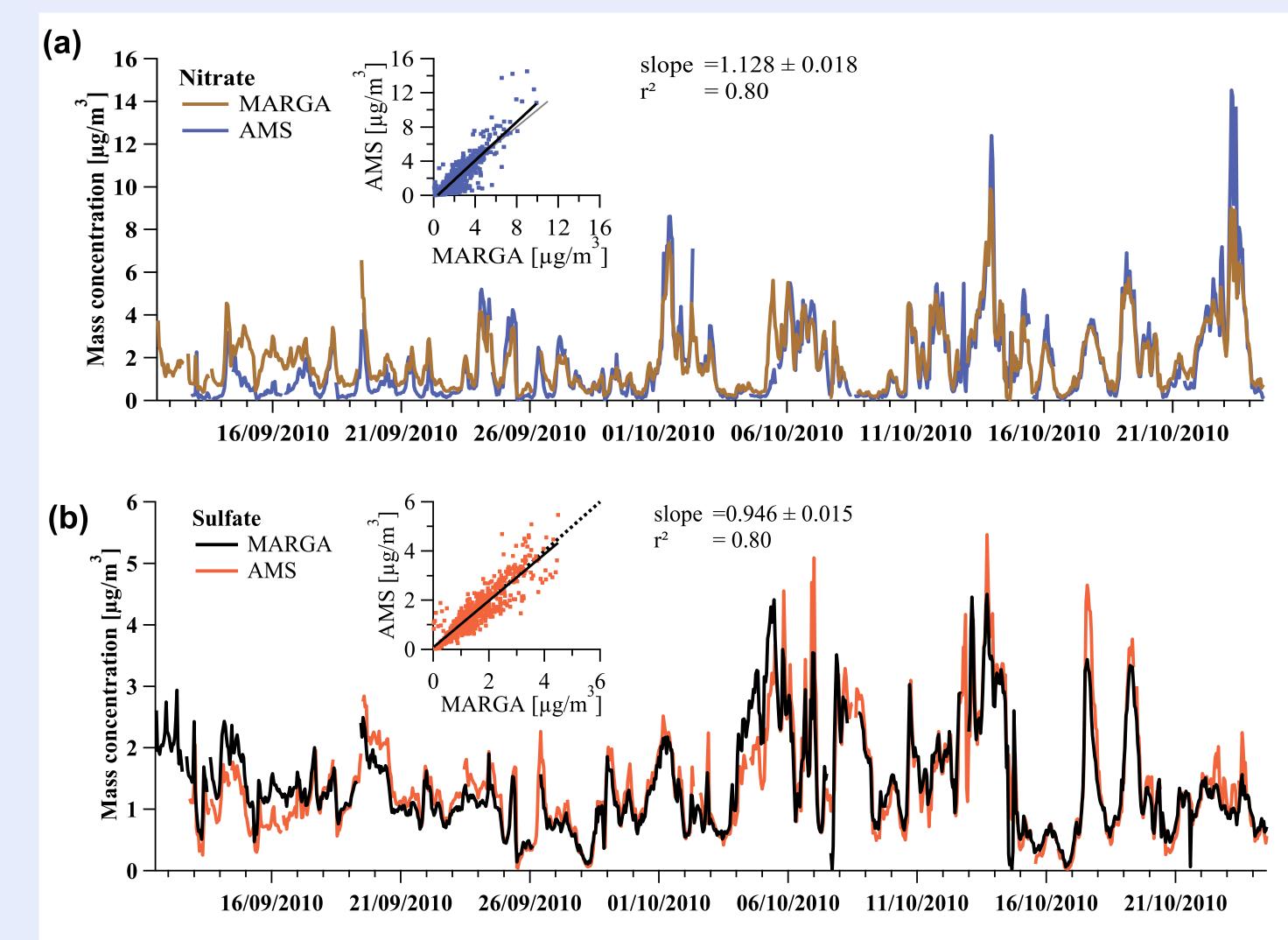


Fig. 1: Gas concentrations during HCCT 2010 measured with the MARGA.

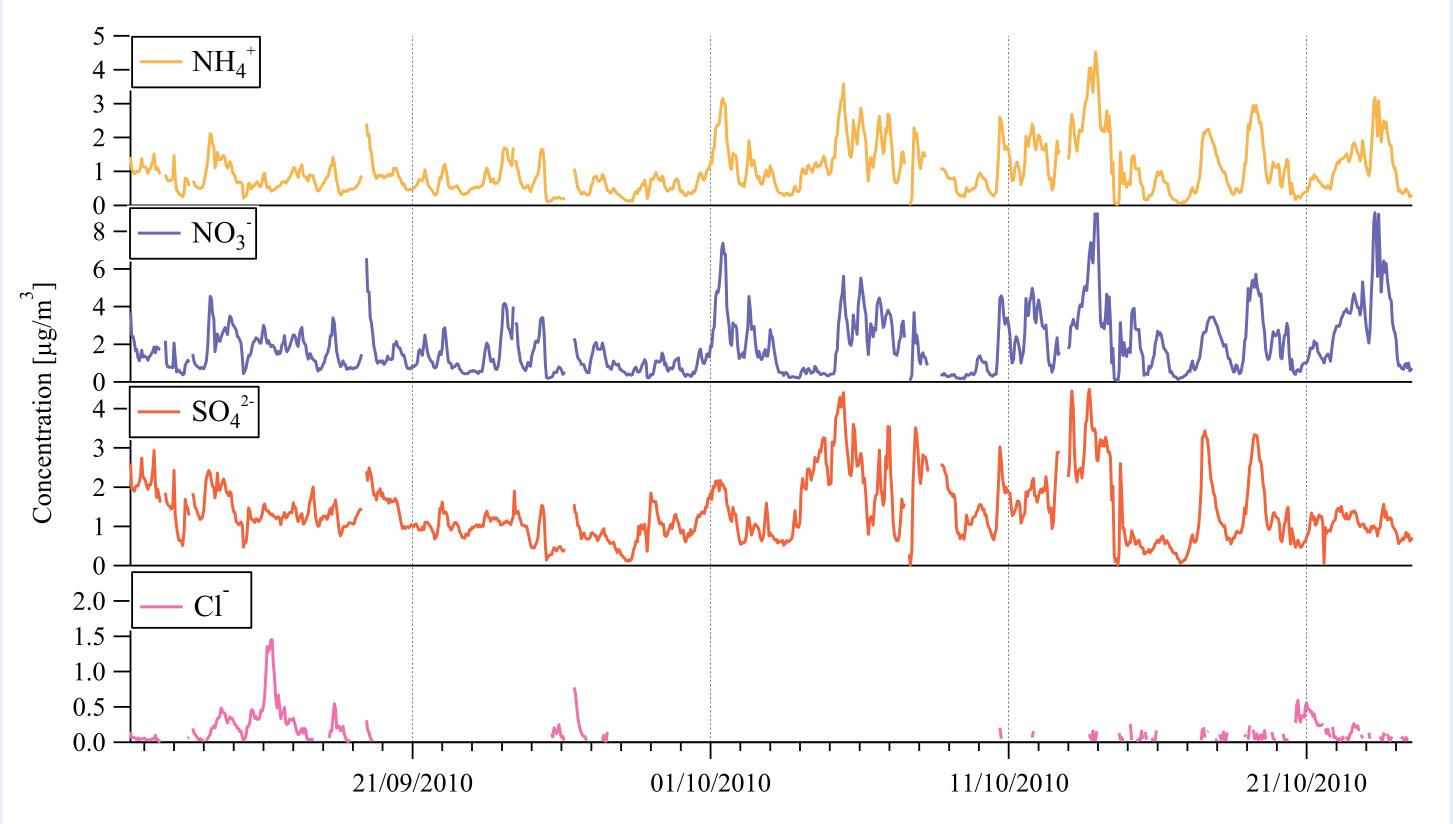


Fig. 3: Comparison between MARGA and HR-ToF-AMS measurements during HCCT 2010. (a) Comparison of the nitrate concentration. (b) Comparison of the sulfate concentration.

## **DIURNAL VARIATIONS**

Figure 4 shows the diurnal variation of selected gases and particles as box plots averaged over the whole HCCT campaign.

- Maximum concentration for NH<sub>3</sub> and SO<sub>2</sub> is observed in the early afternoon, minimum concentration during the night.
- SO<sub>4</sub><sup>2-</sup> shows no diurnal variation.
- NO<sub>3</sub><sup>-</sup> shows a weak diurnal variation with highest concentration in the morning and lowest concentration in the afternoon

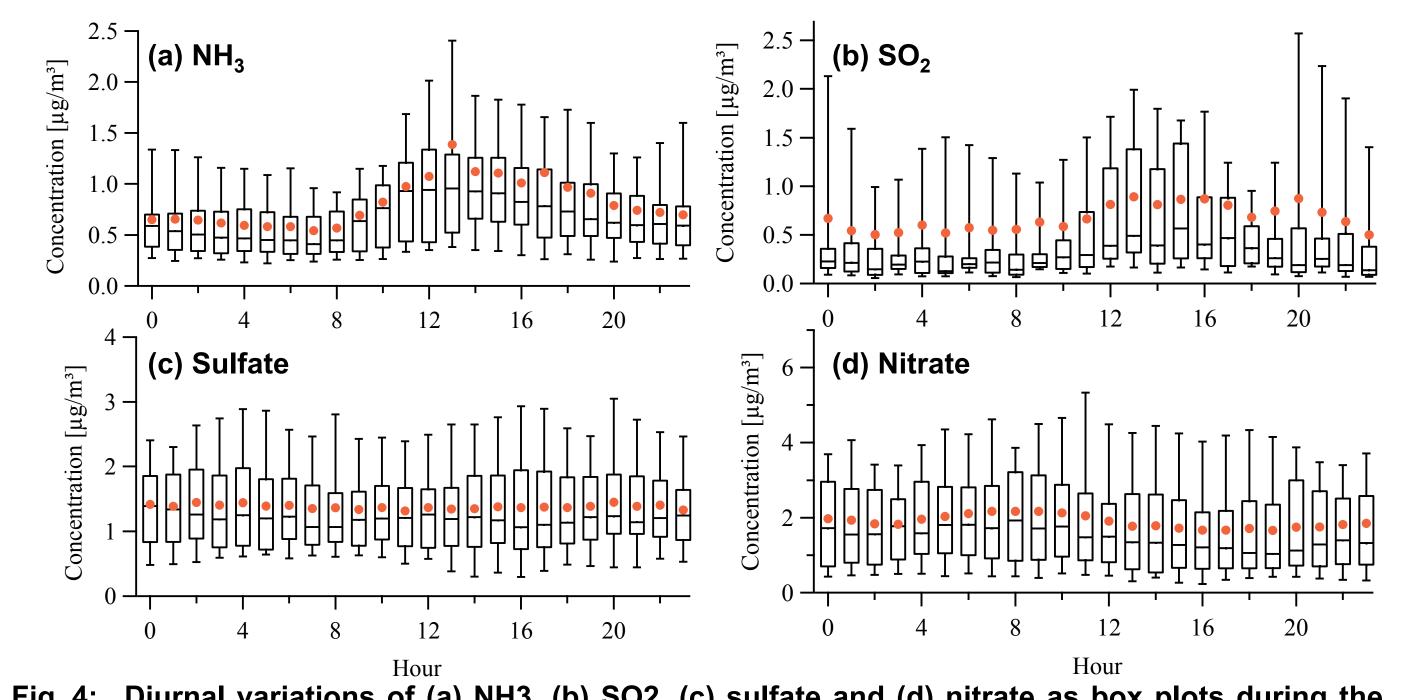


Fig. 2: Concentrations of inorganic ions during HCCT measured with the MARGA.

## SUMMARY

A MARGA was used to quantify the water-soluble inorganic ions in the particle phase and the corresponding trace gases during the HCCT campaign in autumn 2010. A data yield of over 90% was achieved for the main ions and gases.

The comparison of the MARGA with a HR-ToF-AMS shows a good agreement for nitrate, ammonium and sulfate. Only in the first week of the campaign a significant difference in the nitrate concentrations was visible.

Fig. 4: Diurnal variations of (a) NH3, (b) SO2, (c) sulfate and (d) nitrate as box plots during the HCCT campaign. The red dots represent the median concentration of the measured aerosols.

### References

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[2] Makkonen, U., A. Virkkula, Mäntykenttä, J., Hakola, H., Keronen, P., Vakkari, V., Aalto, P. P. (2012). Semi-continuous gas and inorganic aerosol measurements at a Finnish urban site: comparisons with filters, nitrogen in aerosol and gas phases, and aerosol acidity. *Atmos. Chem. Phys. Discuss.* 12(2): 4755-4796.
[3] Poulain, L., Spindler, G., Birmili, W., Plass-Dülmer, C., Wiedensohler, A., Herrmann, H. (2011). Seasonal and diurnal variations of particulate nitrate and organic matter at the IfT research station Melpitz. *Atmos. Chem. Phys.* 11(24): 12579-12599.