The chemical composition of the marine aerosol and its relation to hygroscopic properties over Atlantic Ocean

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MEASUREMENT

Marine aerosol particles play an important role in global climate regulation and marine biogenic system. For a better understanding of its importance, much research was performed to investigate its physical and chemical properties by ship campaigns, but mostly focused on coastal regions or part of ocean areas. In our project the physicochemical properties of marine boundary layer aerosol were measured on board of the research vessel Polarstern during a series of cruises from North Atlantic to South Atlantic. This study shows results of 1) spatial distribution of main chemical compositions along the ship track, and 2) the closure study utilizing the HTDMA and AMS data from Cruise ANT-XXVII/4.

High Resolution Time of Flight Aerosol Mass Spectrometer (HR-ToF-AMS) Size - resolved chemical composition

Aerodynamic Particle Sizer (APS) Number size distribution (D_p> 500nm)





ANT-XXVII/4

Expedition: from Cape Town, South Africa to Bremerhaven, Germany Duration: April 20, 2011 - May 20, 2011 Total distance: 6633 nm



Germany Research Vessel Polarstern (Handbook Polarstern (draft), AWI, 2007)

Scanning Mobility Particle Sizer (SMPS) Number size distribution (D_p: 10-800nm)

Hygroscopicity Tandem Differential Mobility Analyzer (HTDMA) Hygroscopic growth factors at different RH

Differential Mobility Analyzer-Cloud Condensation Nuclei Counter (DMA-CCNc) Particle activation

Integrating Nephelometer + Multi Angle Absorption Photometer(MAAP) Light scattering and absorption

RESULTS

Back trajectories and spatial distribution of non-sea-salt sulfate, organics and ammonium



Aerosol acidity estimation

Main inlet

Digitel filter

sampler

24h filter

sampling for

chemical

composition

Assume that 1) particles are completely neutralized, 2) ammonium is only present in the form of NH_4NO_3 , $(NH_4)_2SO_4$, and NH_4CI ,

 $Predicted [Ammonium] = 18 \times (2 \times \frac{[Sulfate]}{96} + \frac{[Nitrate]}{62} + \frac{[Chloride]}{35.5})$

The relation between predicted and measured ammonium showed that aerosol measured during the cruise is neither pure acidic, nor pure neutralized, indicating influence from different air mass.



Three main compositions non-sea-salt (nss) sulfate, organics and ammonium show very low concentrations (< 1 µg m⁻³) in the most part of cruise, but higher concentration when the ship was close to Europe and North Africa, indicating the possible influence from land. Gaps correspond to period with/of ship exhaust contamination and have been deleted.

Closure Method

The hygroscopic growth factor (HGF) for 200 nm particles measured by the HTDMA was directly compared with the estimated one based on the Zdanovskii-Stokes-Robinson (ZSR) method and the AMS and MAAP mass concentration. Predicted HGF is calculated as below:

$$HGF = \left(\sum_{i} \varepsilon_{i} HGF_{i}^{3}\right)^{1/3}$$

 ε_i are volume fraction of composition *i* in the mixture under the assumption of no change in the volume after compounds are mixed.

	$(NH_4)_2SO_4$	NH_4NO_3	NH ₄ HSO ₄	H_2SO_4	Organics	soot
Density (g·cm ⁻³)	1.77	1.72	1.78	1.83	1.4	1.77
HGF (D _{p-dry} =200nm)	1.72	1.82	1.8	2.06	1.19	1

Closure Study

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The predicted hygroscopic growth factor (HGF) based on AMS chemical composition could generally fit well with measured HGF on 200 nm.

SUMMARY

During the cruise ANT XXVII/4, the mass concentration of main chemical compositions was very low. The higher concentration when the ship was close to European and African continents indicated potential continental influences. The closure study with ZSR method showed good fitting between predicted and measured HGFs in general. More detailed classification of chemical compositions need to be done to reduce the uncertainties of closure study.

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