Long-term Chemical Characterization of Aerosol at the **CVAO (Cape Verde)**

K. Müller, K.W. Fomba, E. Brüggemann, T. Gnauk, G. Spindler and H. Herrmann

Department of Chemistry, Leibniz Institute for Tropospheric Research, 04318 Leipzig, Germany Keywords: aerosol, chemical size distribution, seasonality, spatial distribution Presenting author email: konrad@tropos.de



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GOALS AND METHODS

Introduction

Cape Verde islands are the best place to investigate aerosols over the tropical NE Atlantic. At the CVAO the PM collection started in 2007, with one important goal being the input of nutrients into the Atlantic Ocean by Saharan dust.

Experimental approach



A PM₁₀ HV sampler (DHA-80, Digitel) and a five-stage low pressure impactor (LPI80/0.05, Hauke) were applied to collect PM for chemical analyses. The impactors were used only during intensive campaigns. In this study the PM samples and cloud water samples were investigated for water soluble ions, H_2O_2 , formaldehyde, OC and EC by ion chromatography

(ICS3000, Dionex) and a thermographic method (Cmat5500, Ströhlein).

Fig. 1. Typical back trajectories during a winterly (red) Saharan dust event and for a day with clean marine air. Filter samples during conditioning and sampler on the tower.







Seasonality of PM₁₀ particle mass concentration collected with DHA-80 HV-filter

121

10

22

Dec - Feb

176

15

22

30

68

Total

539

18

43

76

273

129

sampler between January 2007 and November 2011 on top of the tower*.

124

0

15

83

22

*Samples collected between 23 October 2009 and 09 July 2010 were not included here

Mar - Mav

118

53

because of the lower sampling height on the container roof.

Table 1:

Mass concentration

[µg/m³]

>200

90-200

50-90

20-50

<20

Number of samples

Fig. 2 Annual variability and monthly mean of PM10 mass concentration, non-sea-salt sulfate, EC, OM, nitrate and ammonium from all samples collected on top of the tower

Chloride depletion [in $\% \pm$ standard deviation] in the five size fraction Table 2: collected by Berner impactor characterized by the air mass origin

Size class Dp [µm]	Sahara	Marine - summer	Marine - winter	Europe	Mean of all
0.05 – 0.14	56±22	n.d.	n.d.	100±	81±13
0.14 – 0.42	50±32	89±11	62±14	100±	78±14
0.42 – 1.2	21±4	84±11	36.2±7	58±12	41±11
1.2 – 3.5	3.6±2.6	27.8±5.3	25.6±5.2	29.9±7.2	26±8
3.5 - 10	6.4±3.7	27.4±7.2	10.2±4.1	28.2±5.1	24±8

Major observations during HV sampling over five years

- Saharan dust season is from end of November to beginning of March (Tab. 1).
- > In the yearly mean Saharan dust was found to 25 μ g/m³ before sea salt with 20 μ g/m³ and secondary ions with 2.7 μ g/m³.
- Iron (Fig. 3) was found during dust events in concentrations up to 5µg/m³.
- Elevated EC is coupled to higher OC values (mainly in winters with air mass origin in Westafrica) but for OC during summer higher concentrations were observed from marine production.
- Ammonium, nss-sulfate, EC and OM were found in ultrafine and fine particles preferentially (Fig. 5).
- Ammonium was found with a spring maximum coupled to chlorophyll A.
- \succ In clean marine air masses nss-sulfate is correlated to oxalate (R² >0.9) in all other samples a lower correlation (Fig. 4) was found.
- Nitrate is found during summers to 97% in coarse mode and to 88% in winters.
- Chloride depletion (Tab. 2) was found in nearly all samples (depending on air mass):
 - Europe > marine summer > marine winter > Sahara

Fig. 5 Mean constitution of PM (without dust) during a summer and a winter campaign.



Fig. 6 Mean PM₁₀ constitution of 183 marine samples (left) and 49 mineral dust dominated samples (right).

SUMMARY AND OUTLOOK

The CVAO is an excellent place to investigate Saharan dust and marine aerosols. Saharan dust and sea salt are the dominating aerosol constituents. Non sea salt components were found from long range transports from Africa and Europe or were formed from marine precursors. Interannual variability and seasonality were observed for all components. In close cooperation with oceanic research the effects of PM deposition and the feeding by mineral dust and accompanied secondary species will be investigated.

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