

GROUND-BASED CHARACTERISATION OF AEROSOL CHEMICAL COMPOSITION DURING THE SALTRACE CAMPAIGN

K. Wadinga Fomba, Konrad Müller, Thomas Müller and Hartmut Herrmann

Leibniz-Institute for Tropospheric Research, Leipzig, Germany

Email: fomba@tropos.de



TROPOS
Leibniz Institute for
Tropospheric Research

Member of the
Leibniz
Leibniz Association

Introduction

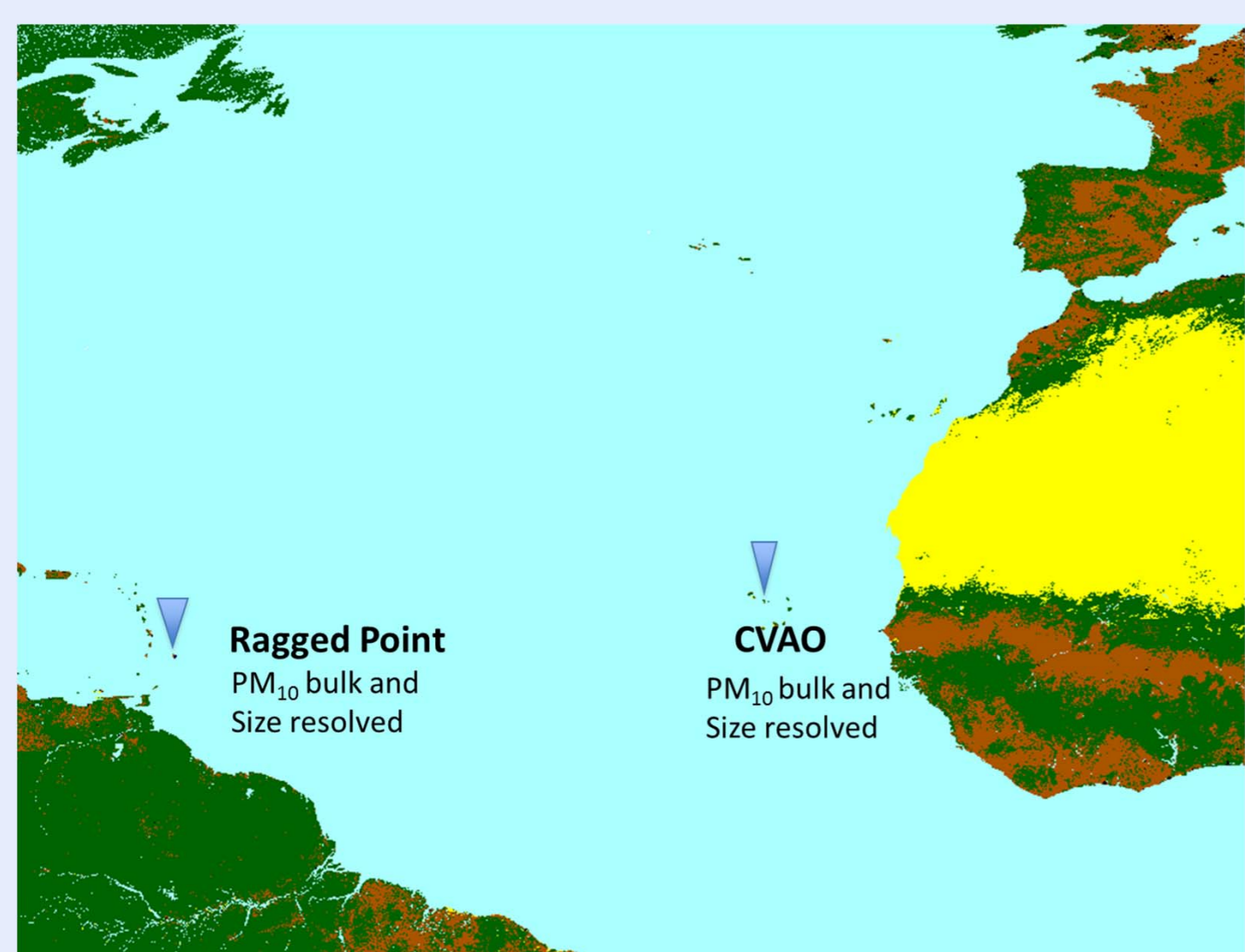


Fig. 1: Geographical depiction of ground-based stations during SALTRACE experiment

The transport of mineral dust over long distances can lead to changes in its chemical and physical properties. During the SALTRACE experiment, Saharan dust transport over the Atlantic was characterized using both ground-based and airborne measurements. Amongst others, one of the goals were to characterize the chemical properties close to the source regions (Cape Verde) and far away from the source region (Barbados, Ragged Point).

Experiment and Sampling

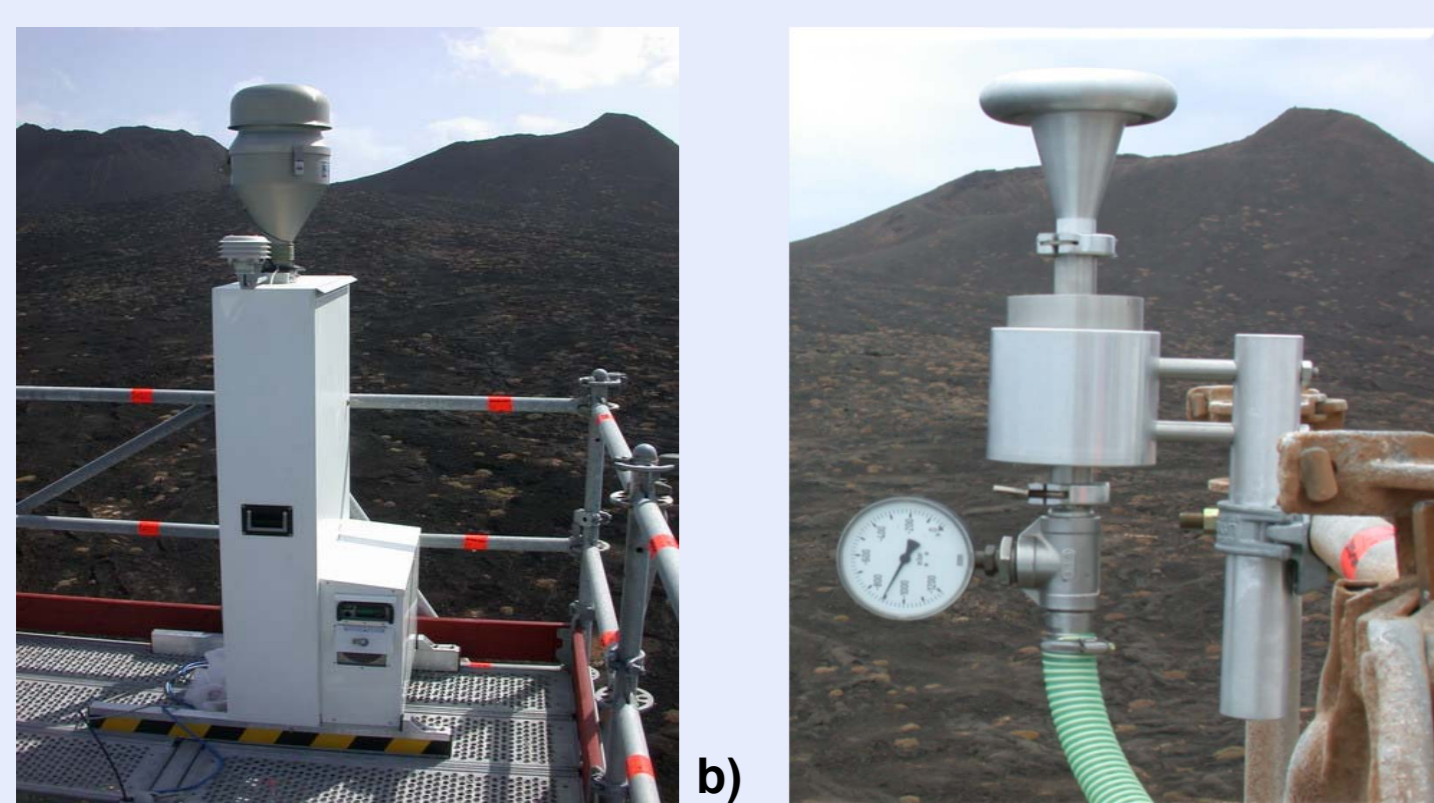


Fig. 2: a) Digital DHA-80 high volume sampler with PM_{10} -inlet, 500 l/min sampling rate b) 5-stage BERNER impactor with 75 l/min, sampling rate, stage cut-offs: 0.05 - 0.14 - 0.42 - 1.2 - 3.5 - 10 μm .

Aerosol particles were collected at the top of a 30 m tower at the CVAO and 25 m tower at Ragged Point using a high volume DIGITEL DHA-80 (Fig. 2a) for bulk PM_{10} particles and a five-stage BERNER impactor (Fig. 2b) with PM_{10} cutoff 0.05-10 μm for size resolved analysis.

The size-resolved particles were collected on aluminum foils while the bulk particles were collected on pre-heated quartz fiber filters. Inorganic ions, soluble and total trace metals, OC/EC were analyzed from these filters.

Trace metals

	CVAO (ng/m^3)	Ragged Point (ng/m^3)
Si	392 \pm 196	1495 \pm 850
K	157 \pm 55	265 \pm 99
Ca	295 \pm 158	555 \pm 231
Fe	119 \pm 52	578 \pm 284
Ti	11 \pm 5.4	62 \pm 31
Mn	2 \pm 1.2	12 \pm 6
Cr	4.71 \pm 3.09	2.83 \pm 0.85
Cu	4.21 \pm 2.29	0.55 \pm 0.2
V	1.25 \pm 0.51	1.68 \pm 0.84
Zn	3.24 \pm 1.3	1.88 \pm 0.72
Ni	0.86 \pm 0.24	0.76 \pm 0.3
Pb	0.96 \pm 0.31	0.84 \pm 0.34

Table 1: Trace metal concentrations observed at CVAO and Ragged Point.

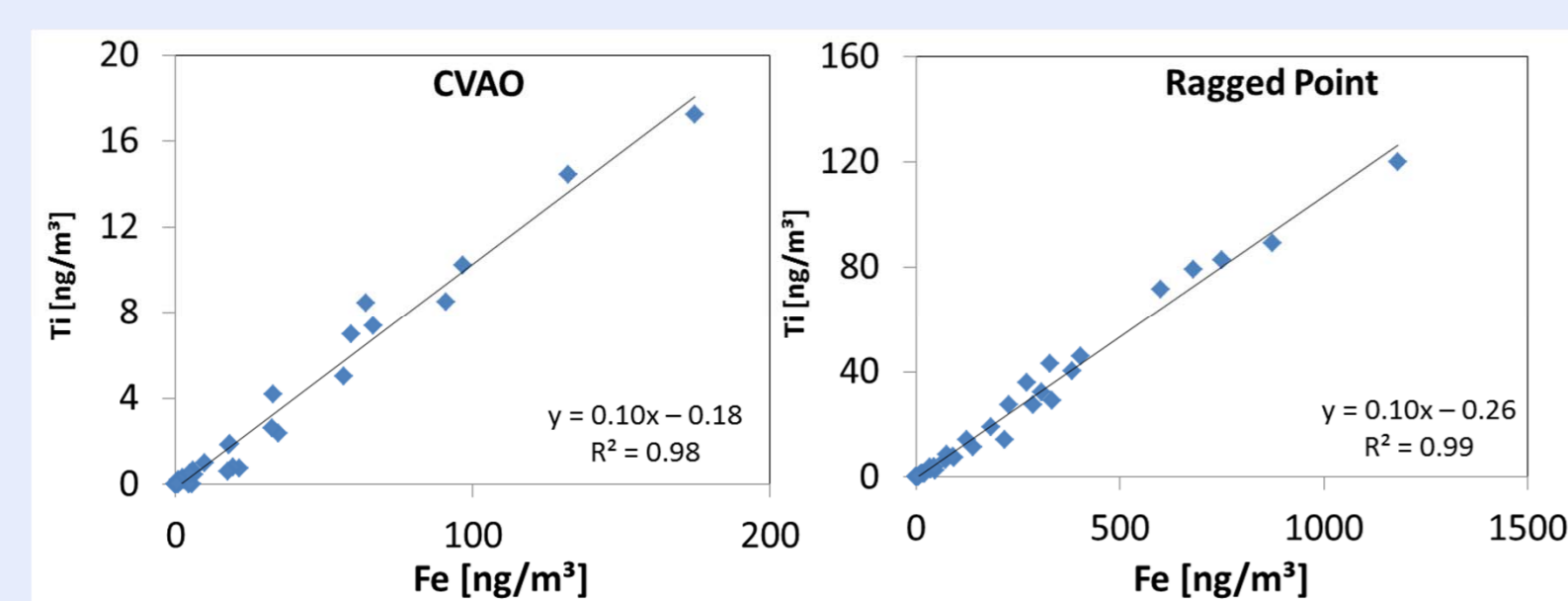


Fig. 7: Ti used as dust traces showed good correlation with iron and also similar ratios at both stations

Z/Ti	CVAO 03/07/13	Ragged Point 10/07/13
Si	33.34	29.94
Mn	0.25	0.21
Co	0.02	0.015

Table 2: Mineral dust elemental ratios observed during dust events at CVAO and Ragged Point.

Acknowledgement

The authors will like to thank the TROPOS for financing this experiment and also their technical staff for their kind assistance. Many thanks also to DLR for organizing the experiment.

Results

Strong variation in aerosol mass concentration with peak concentrations of about 50 $\mu g/m^3$ indicating days of strong Saharan dust influence on aerosol loadings. However, aerosol mass at both stations was concentrated in the coarse mode. Average mass concentrations were $23.58 \pm 9.1 \mu g/m^3$ and $28.04 \pm 11.1 \mu g/m^3$ at CVAO and Ragged Point, respectively.

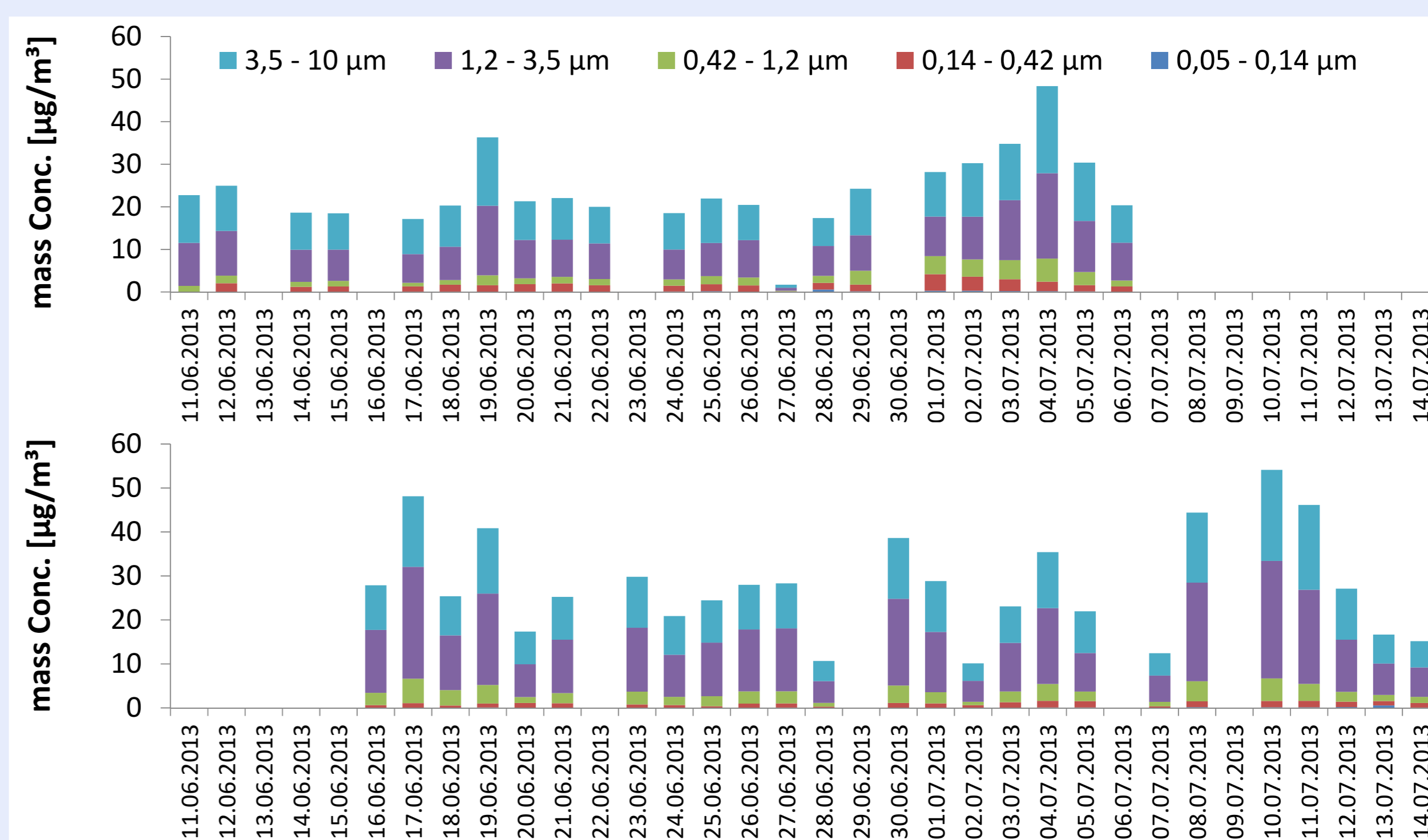


Fig. 3: Size resolved aerosol mass concentration at CVAO (top) and Ragged Point (bottom) during four weeks of parallel measurements at both stations.

Aerosol mass was dominated (80%) by sea salt and mineral dust with higher mineral dust than sea salt observed at Ragged Point in comparison to CVAO.

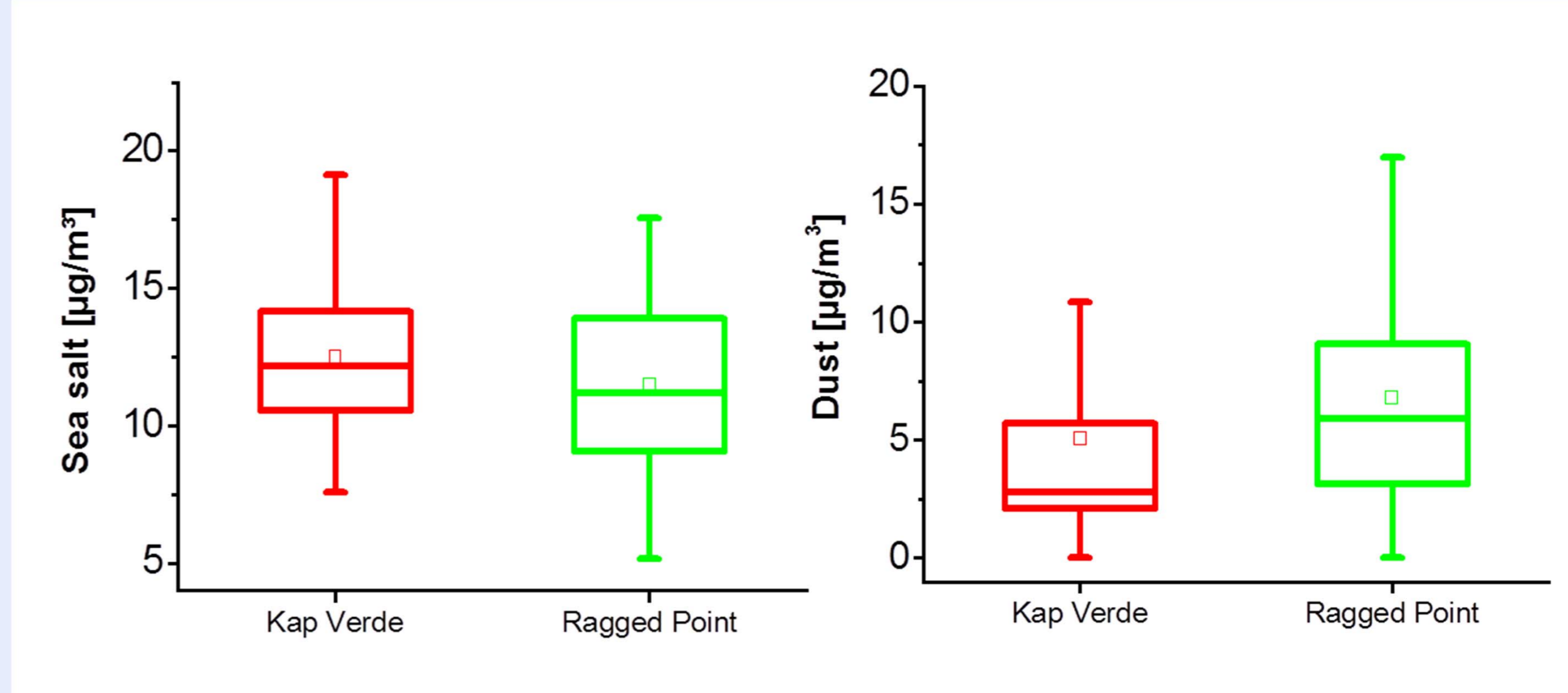


Fig. 4: Sea Salt and mineral dust aerosol compositions at Ragged Point and CVAO

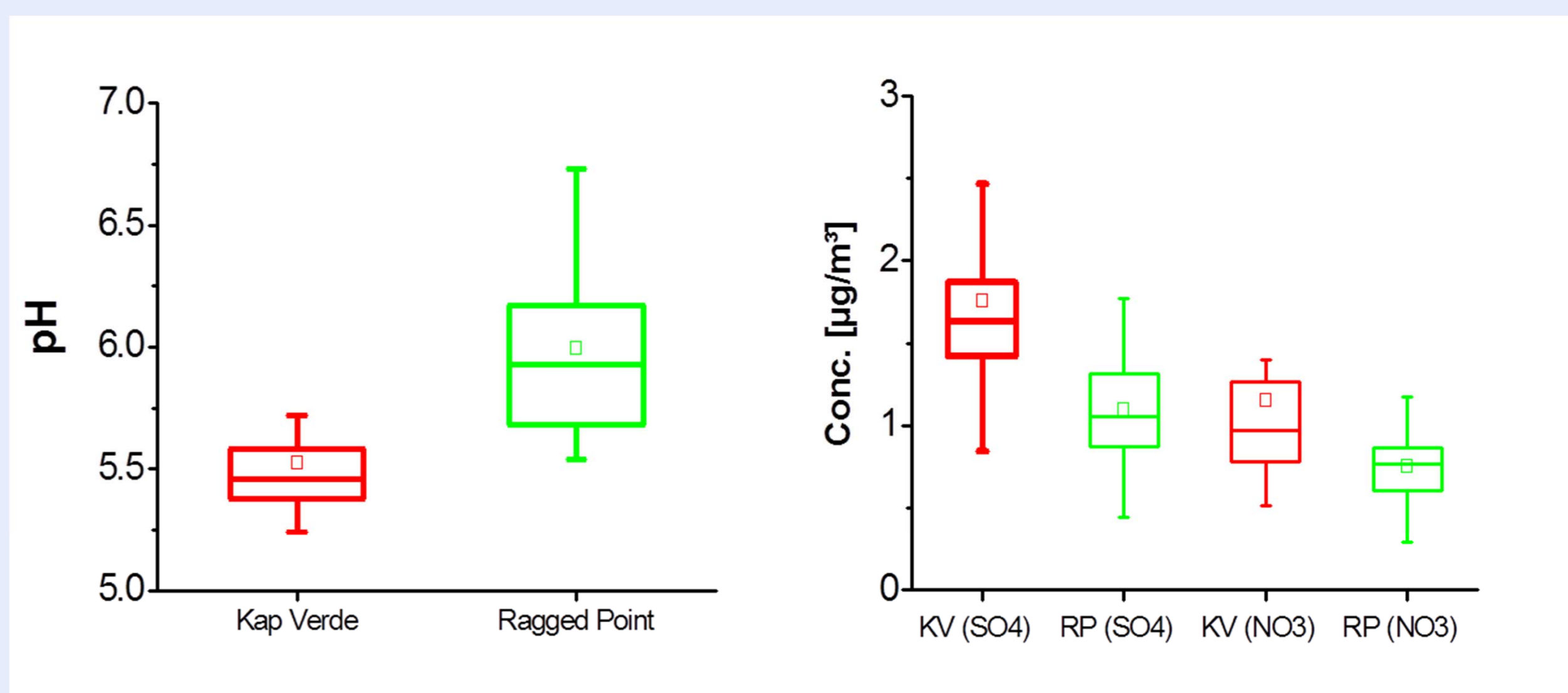


Fig. 5: Aerosol acidic components (sulfates and nitrates) and pH at Ragged Point and CVAO

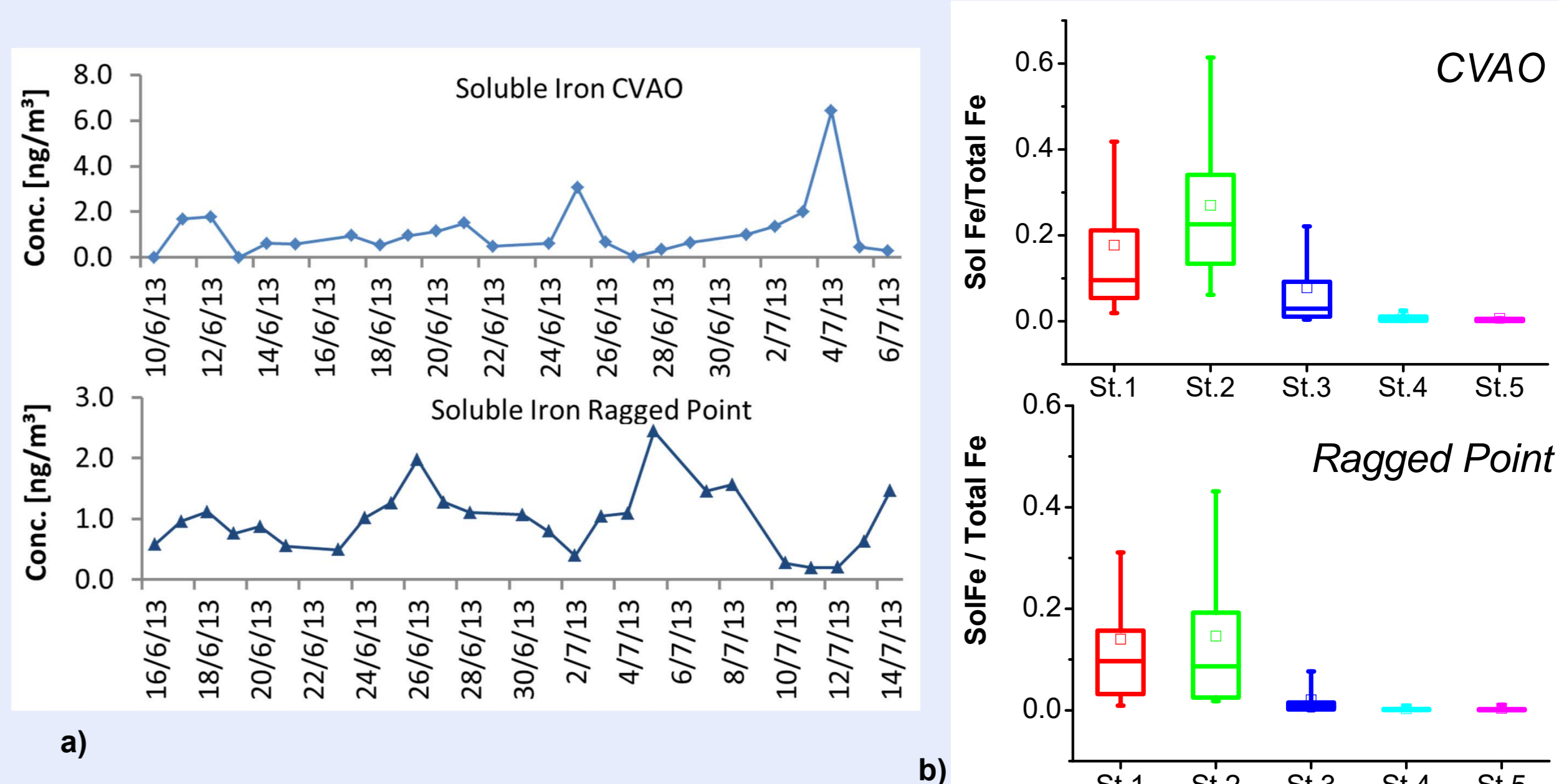


Fig. 6: a) Soluble iron Variation at CVAO and ragged point and b) size resolved fractional soluble iron at both stations.