VOCALS REx: Aerosol Physics at the Ocean Surface On the NOAA RV Ronald H. Brown October, November 2008 Catherine Hoyle, NOAA PMEL Derek Coffman, NOAA PMEL Tim Bates, NOAA PMEL Trish Quinn, NOAA PMEL Lelia Hawkins, Scripps Institution of Oceanography David Covert, UW Dept. of Atmospheric Sciences

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VOCALS REX GOAL

Better understand aerosol, clouds and precipitation as interactive processes that in turn influence cloud properties - cloud cover, thickness and particle size and optics over the South East Pacific.

Hypotheses we are addressing – two of many.

1a .

"Variability in the physico-chemical properties of aerosols has a measurable impact upon the formation of drizzle in stratocumulus clouds over the SEP."

1c.

"The small effective radii measured from space over the SEP are primarily controlled by anthropogenic, rather than natural, aerosol production; entrainment of polluted air from the lower free-troposphere is an important source of cloud condensation nuclei (CCN)."

Panama Canal, entry from the Atlantic





Instrument Summary

Optics: light scattering and absorption coefficients nephelometer and absorption photometers, 3 wavelength, humidified nephelometer

Chemistry: 12 to 24 hour impactor, filter samples for major ions.

Physics: number-size distribution 20nm to 10μ m diameter DMPS, Differential Mobility Particle Sizer, 20 to 800nm APS, Aerodynamic particle sizer, 800nm to 10μ m

Cloud Condensation Nuclei Concentration DMT, Thermal Gradient CCNC 0.10, 0.15, 0.20, 0.30 and 0.60% SS.

DMPS is a **differential** measurement 5 minute time resolution CCNC is a **integral** with 60 minute resolution.

In order to relate the two:

Integrate DMPS number-size distribution from 800nm to critical diameter, for a given CCNC supersaturation assuming a chemical composition - eg., ammonium sulfate.

Integral (over size) of particle number concentration from DMPS vs. droplet concentration from CCNC at 0.2% and 0.6% supersaturation



Chemical assumption: Ammonium sulfate



0.6% supersaturation R = 0.980 Intercept = -11 ± 6 Slope = 1.15 ± 0.018





Coastal, continental aerosol

Dominant accumulation mode Minor Aitken mode (shoulder)

Strongly influenced by urban industrial sources in region 500 km, several days upwind Combustion Biomass burning Copper smelter SO_2



Remote marine aerosol

Dominant accumulation mode Minor Aitken mode (shoulder)

Strongly influenced by urban industrial sources in region 500 to 1000 km or more upwind

Mixing ratio of species SH-SO2t at 200 m asl for age class 0 - 10.00 DAYS Latest analysis time 20081108. 0 Actual time 20081121. 0 Mean value 0.111E+00 Maximum value 0.494E+01 Minimum value 0.000E+00 Distance of grid lines 5.0 deg





VOCALS date





Summary:

Aerosol number-size distribution is consistent with past results in the SEP in terms of relation to POCs, i.e. depleted accumulation and Aitken mode concentrations.

A spatial relation between number-size distribution features and POCs is evident, but a causal relationship is not yet clear.

At times the effect of continental, pollution, aerosol extends far to the west of the continent into the SEP stratus field.

Evidence for new particle formation in the MBL is minimal, it occurs in a very small volume fraction.

The chemistry implies a relatively soluble aerosol. I.e. size, not chemistry, is the main controlling factor in terms of effective CCN.











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POC-like aerosol fields are extensive in time and space in the SEP MBL.

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The source of gas phase compounds for mass growth of Aitken mode to effective CCN size seems to be marine DMS and organic compounds, as well as SO_2 and organic compounds advected from the continent.

The chemistry implies a highly soluble aerosol. I.e. size, not chemistry, is the controlling factor in terms of effective CCN.

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