

DRI Cloud condensation nuclei (CCN) spectrometers.

Produce a field of supersaturations ( $S$ ) by thermal diffusion of temperature and water vapor between two parallel plates, where cloud droplets grow on hygroscopic sample particles.

More hygroscopic (e.g., larger) particles produce larger cloud droplets.

Continuous flow through the cloud chamber ( $\sim 30$ s) then into an optical particle counter (OPC). CCN spectrum is deduced from the OPC droplet spectrum. A calibration curve relates OPC droplet size to particle hygroscopicity (critical supersaturation— $S_c$ ).

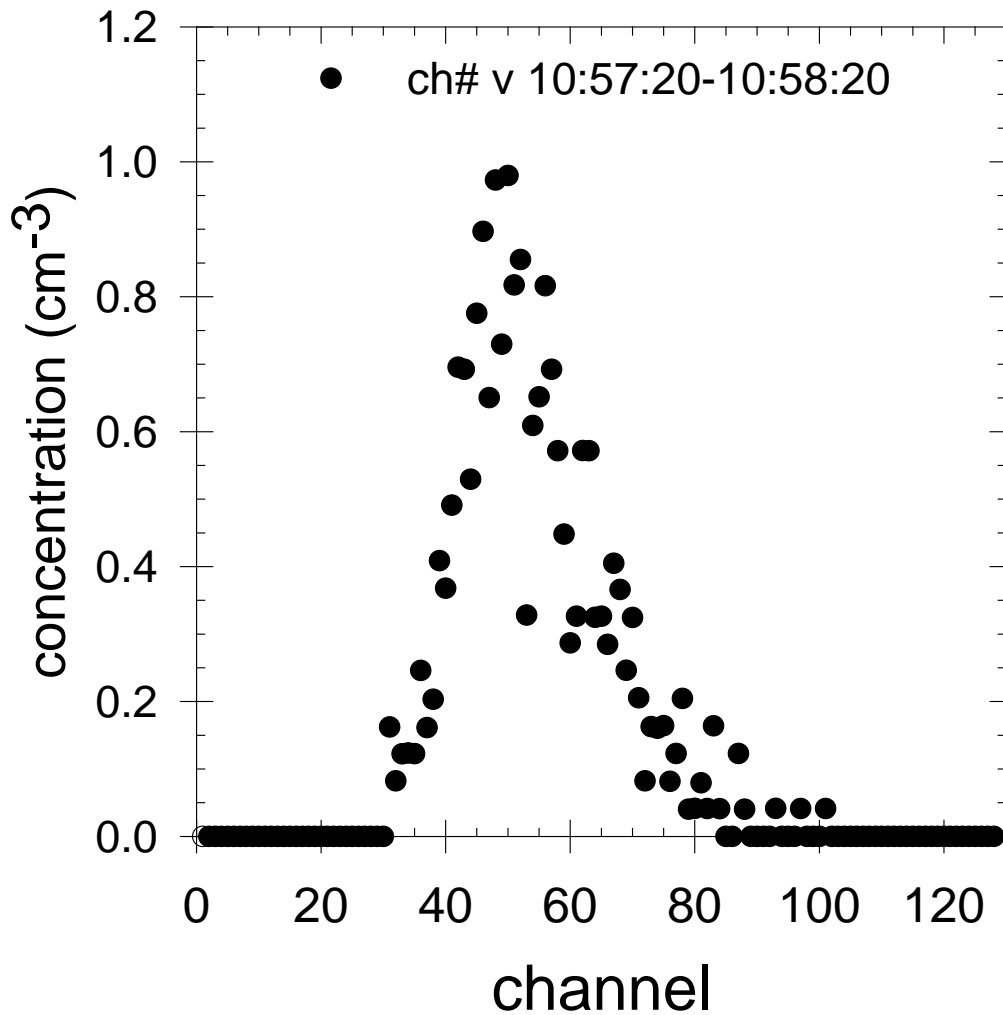
Calibration is done with nuclei of known composition (e.g., NaCl) and size (differential mobility analyzer—DMA—electrostatic classifier--EC).

Assumes that all CCN with the same  $S_c$  regardless of composition (or size) produce the same droplet sizes. Calibration holds only if all chamber parameters (i.e., flows and temperatures) remain constant.

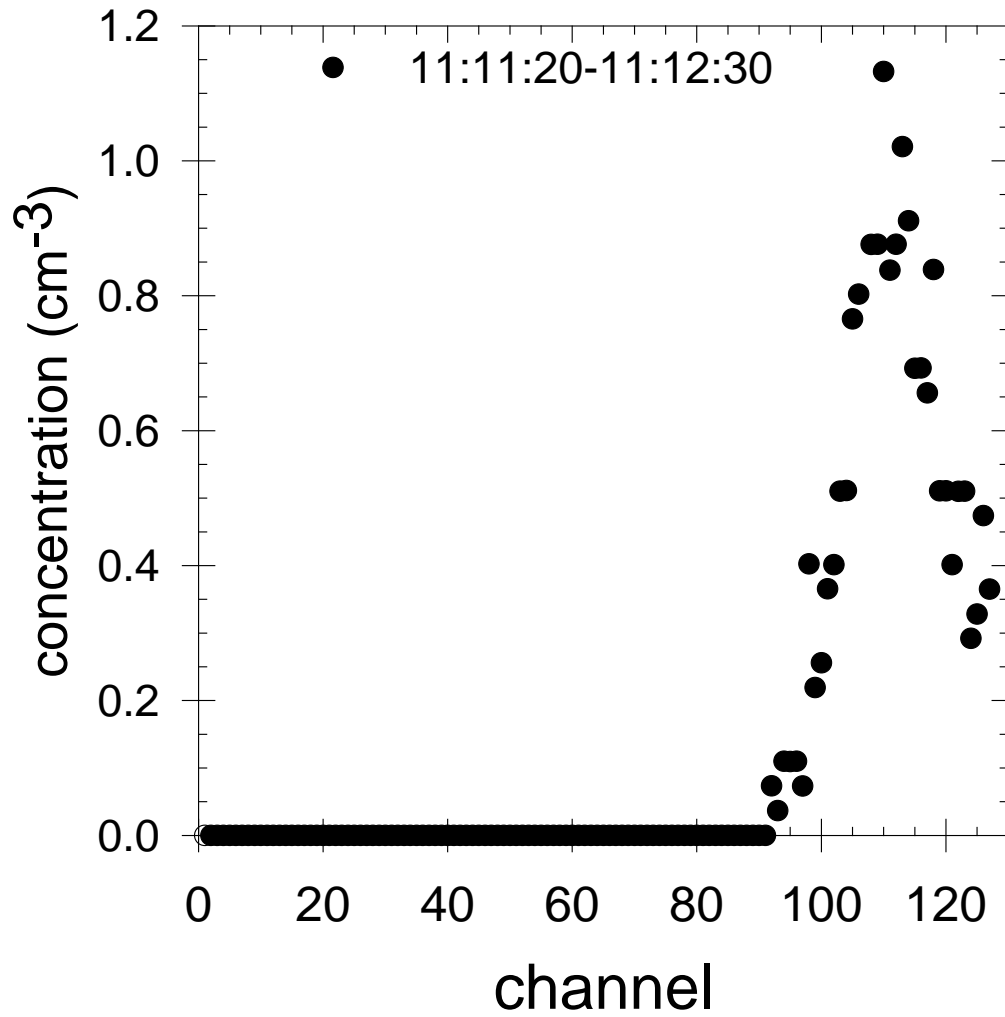
$S_c$  inversely proportional to number of soluble ions. Traditionally CCN plots are cumulative because clouds act cumulatively on the aerosol— all nuclei with  $S_c < \text{cloud } S$  produce “activated” cloud droplets.

Also previous CCN instruments had too few data points to produce a differential spectrum. DRI CCN spectrometers have enough data points to produce differential spectra.

**July 25, MASE**  
**NaCl calibration**  
**27 nm ch-53 0.79%**

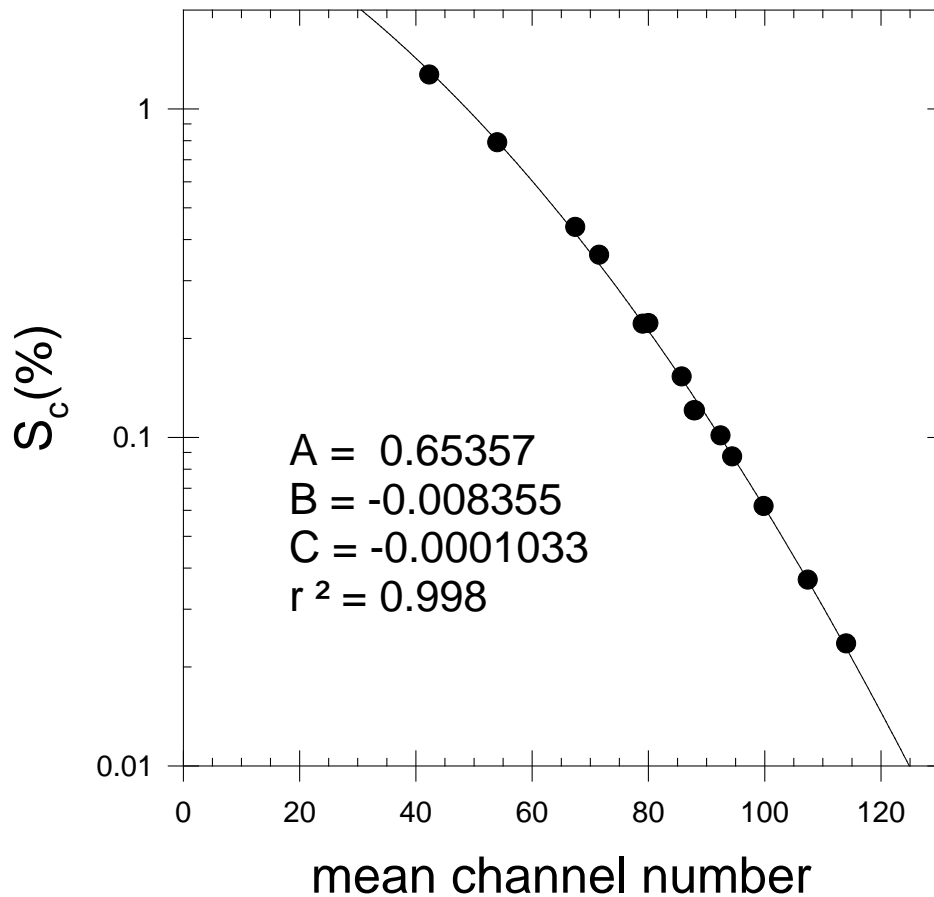


**July 25, MASE**  
**NaCl calibration**  
**280 nm ch-114 0.02%**



July 25, 2005, 1051-1231  
MASE, Sacramento, 6 holes open old sf meter  
new spec. OPC 50106  
LV = 6.66  
HV = 450, gain 8.0, offset 0.2  
sf = 1.58-1.59v  
DT = 0,1.5,3,5.5,7,8  
TM = 25C  
EC 10 lpm NaCl calibration curve

● mnn v Sc10nc  
— Plot 3 Repr



Extension the traditional CCN (Aitken) range below 0.1% to Large Nuclei, is necessary because a large proportion of CCN have  $S < 0.1\%$ . This is needed because:

- Many clouds have  $S < 0.1\%$
- Static closure—aerosol and CCN
- Dynamic closure—CCN and cloud droplets.
- Large nuclei are embryos for precipitation.
- Droplet spectral width.
- Interface with giant nuclei.
- CCN sizes.



# Why two spectrometers

1. Redundancy
2. different S ranges in each instrument with different optimal S validity; check in overlapping range.
3. In flight calibrations
4. Volatility
5. Size versus  $S_c$

Spectrometry is foiled by instrumental broadening, coincidence and vapor depletion.

Broadening is more of a problem for steep spectra.

Coincidence is more of a problem for higher concentrations.

Vapor depletion—high concentrations.

S range 1-0.02% (i.e., 101-100.02% RH) cloud S range.

Higher  $S_c$  for smaller (less hygroscopic) particles.

Fundamental limitations of CCN spectral range.

1. **Instrumental broadening**—same  $S_c$  particles not producing identical signals.

Crossover between channels is more of a problem with steeper spectra.

Improvements have reduced this problem.

2. **Coincidence**—two droplets simultaneously within detection volume artificially convert two small (high  $S_c$ ) droplets to one large (low  $S_c$ ) signal.

Causes overestimates of low  $S_c$  particles.

Higher concentrations exacerbate coincidence.

Lower sample flow rates reduce coincidence.

### 3. Vapor Depletion

high concentrations—competition for water vapor reduces droplet sizes

lower sample flow, but may cause diffusion losses

Not a problem in clean air.

Spectra more challenging for lower part of S range.

Lower  $S_c$  (larger particle) concentrations are generally lower.

Thus broadening and coincidence of high concentrations of high  $S_c$  particles can obscure measurements of low  $S_c$  particle concentrations.

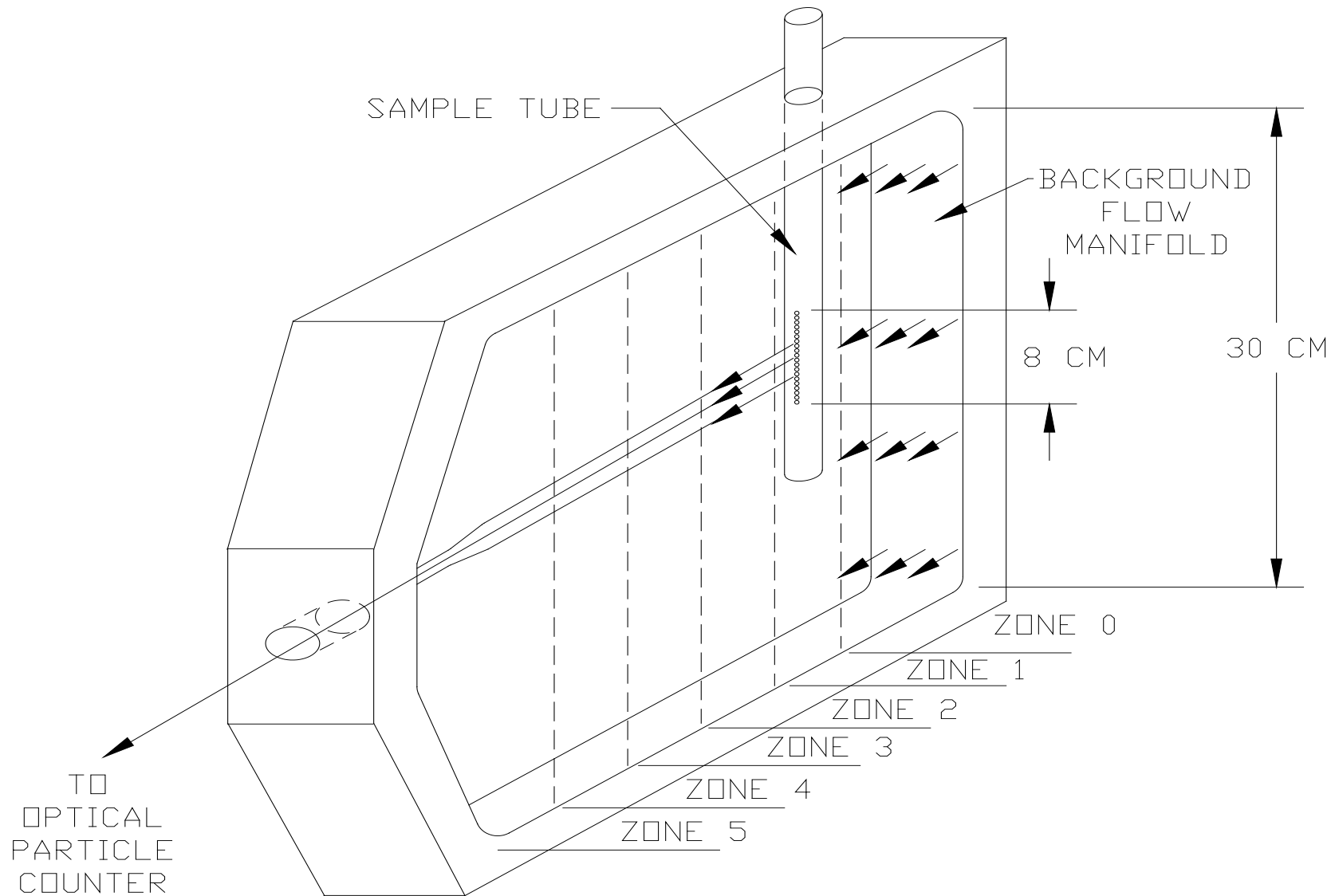
Low  $S_c$  measurements by an instrument operating only at low S (not measuring the high  $S_c$  particles) should be more accurate; they are not obscured by higher  $S_c$  particles.

Therefore, comparisons of low  $S_c$  measurements by instruments operating over different S ranges can confirm operation at high S.

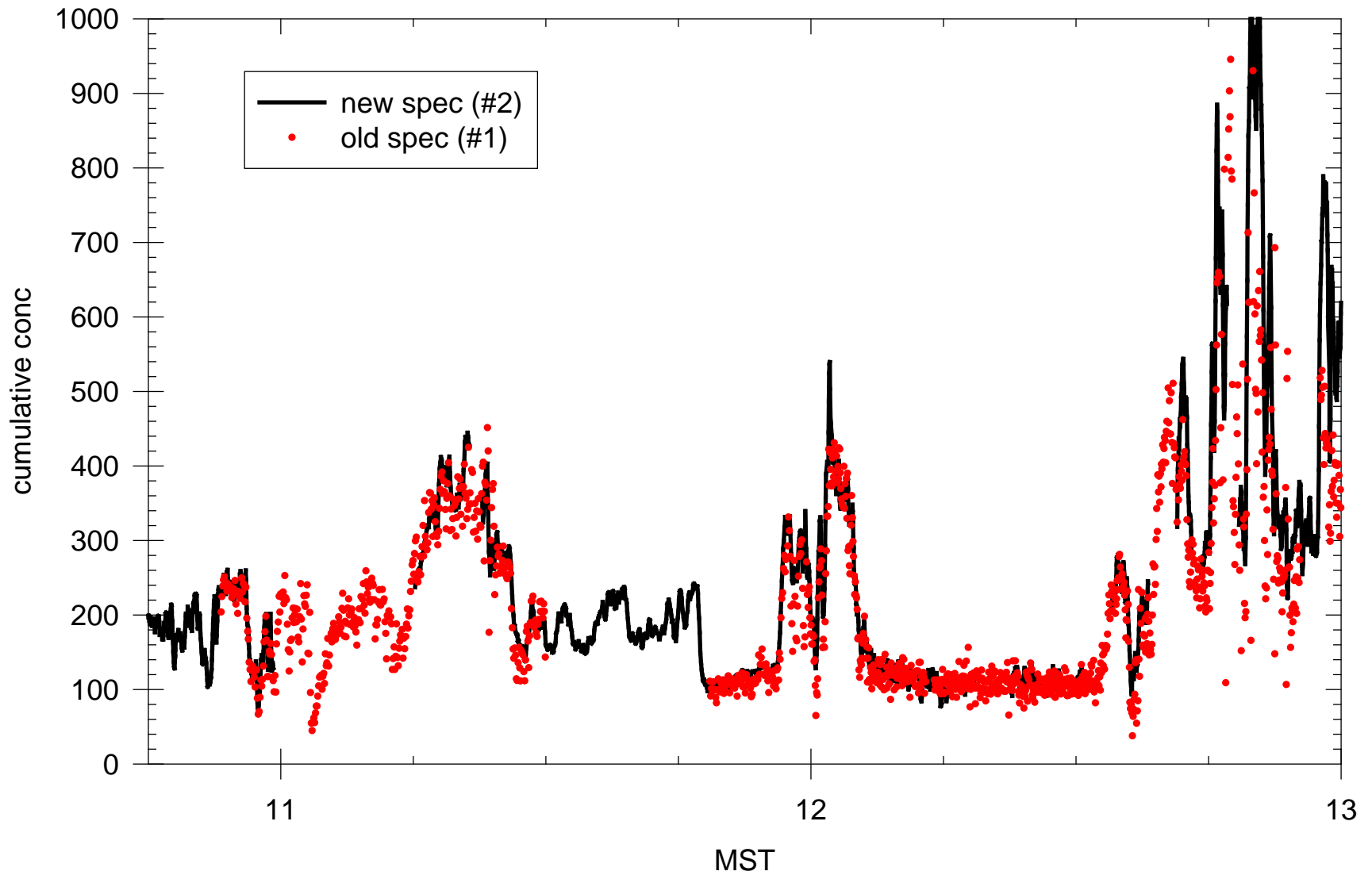
Agreement suggests validity of the challenging low  $S_c$  measurements by the instrument operating at high S.

If lower S part of range is accurate then higher S part of range is also probably accurate.

1. Ambient sampling—1s continuous
2. Different or similar  $S$  ranges?  
Different—better test, greater  $S$  range—0.01-2% $S$   
Similar—redundancy and processing
3. Volatility—one remains on ambient other switches back and forth every minute or so to normalize changes. Greater editing job—remove transitions and separate ambient from processed. Change temps from 20C to 1000C takes ~30 min.
4. Size- $S_c$  –difficult in clean air—constant alt., won't work at high altitude—one continues to monitor ambient, at least one minute per size, one minute transition so at least 10 minutes for 5 sizes. Also calibrations another 10 min.
5. Size-  $S_c$  squared—test Kohler theory—no ambient
6. Volatility and size- $S_c$  together? Very slow and difficult.

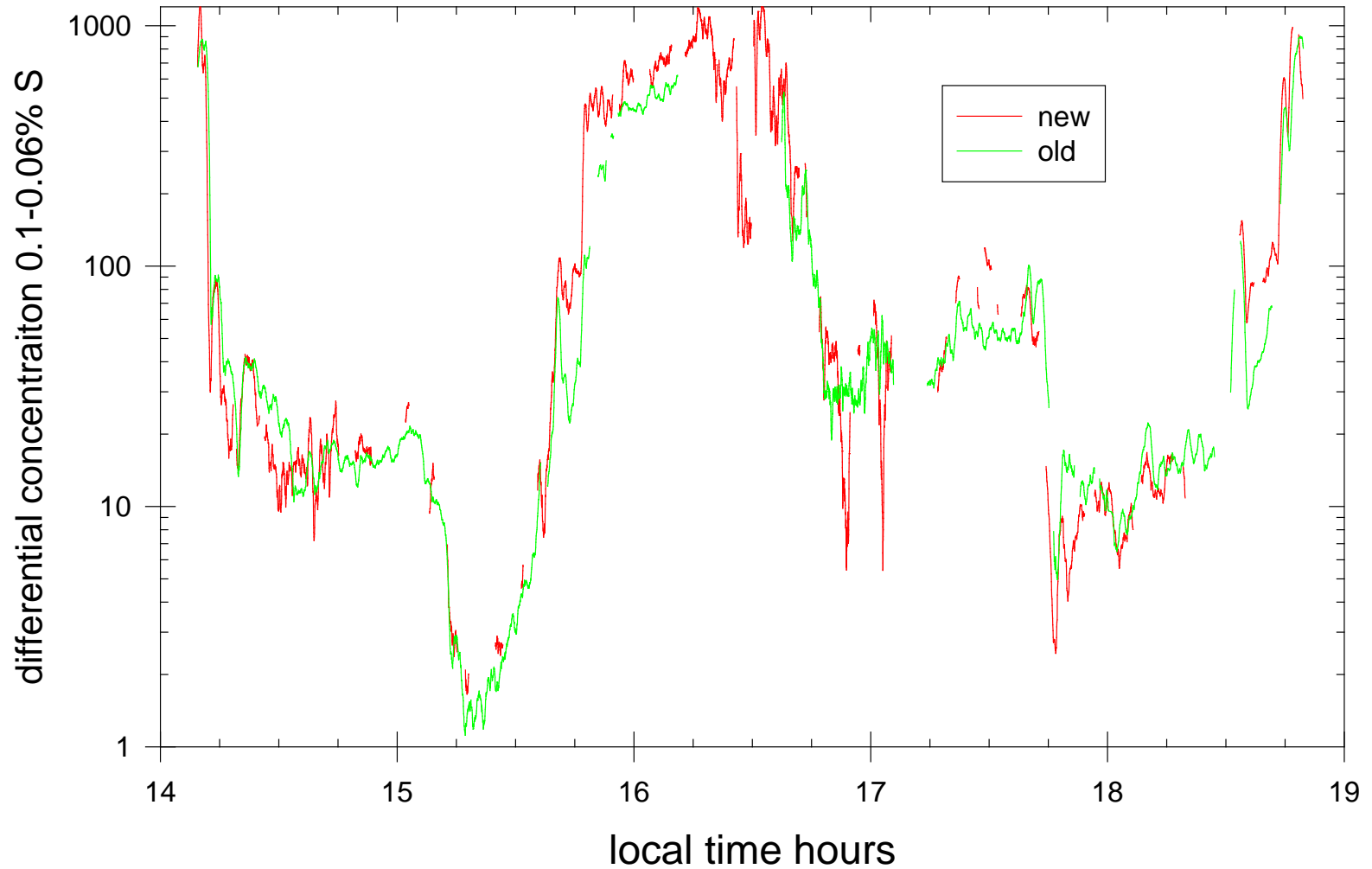


November 20, 2007 ICE-L  
0.30% S

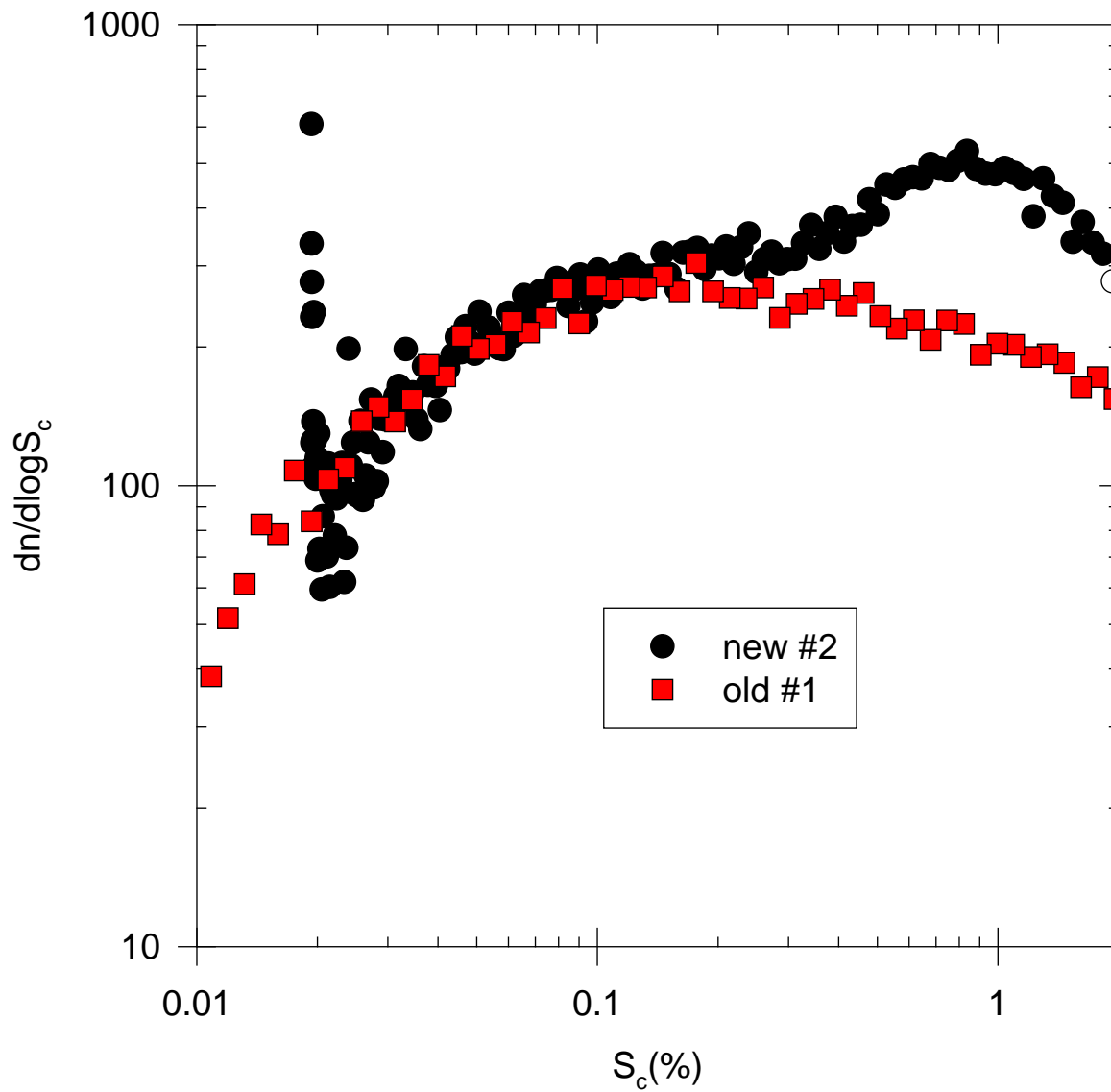




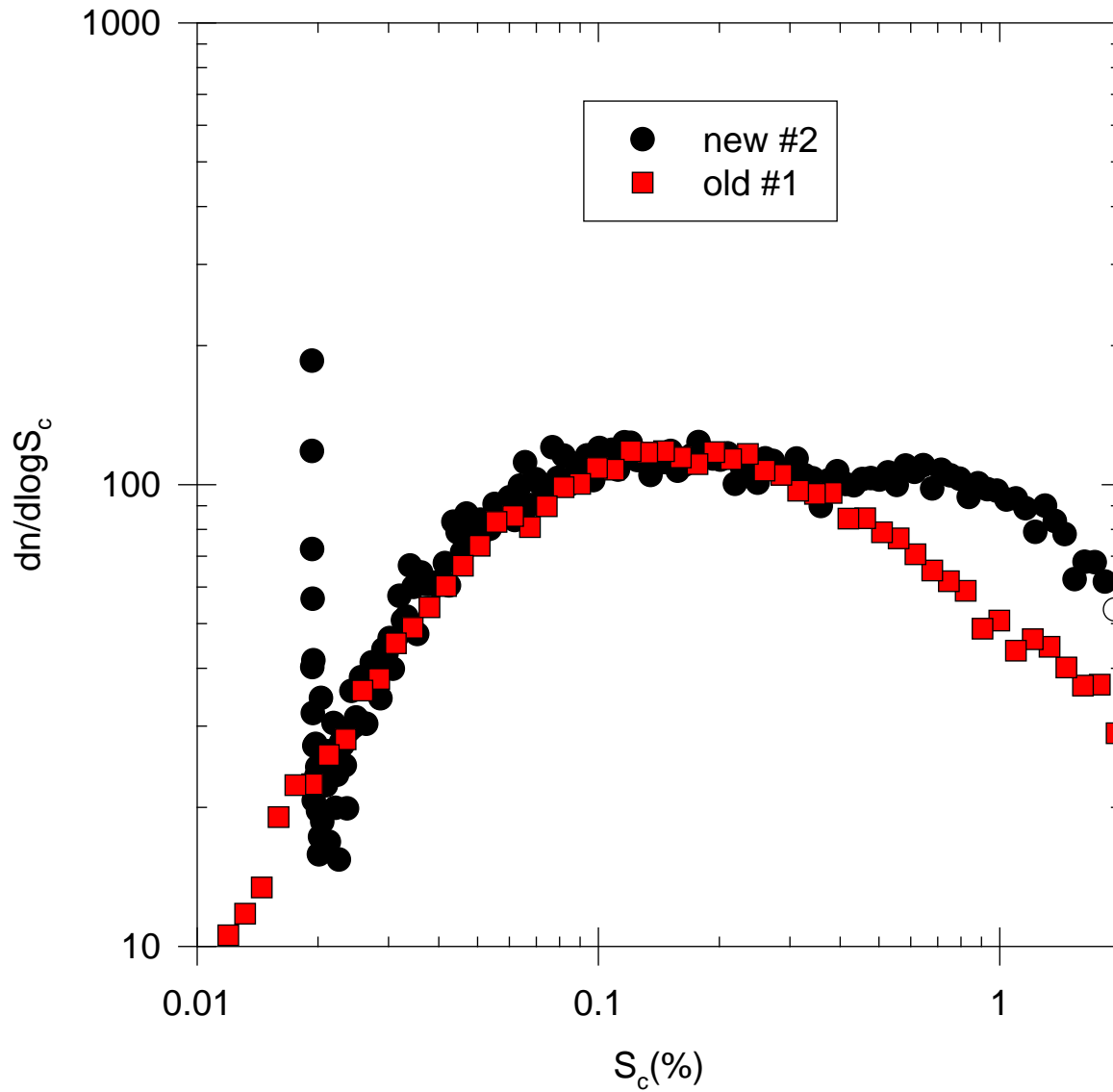
# Nov 10, 2003, AIRS-2, Cleveland comparison running average of 19 records



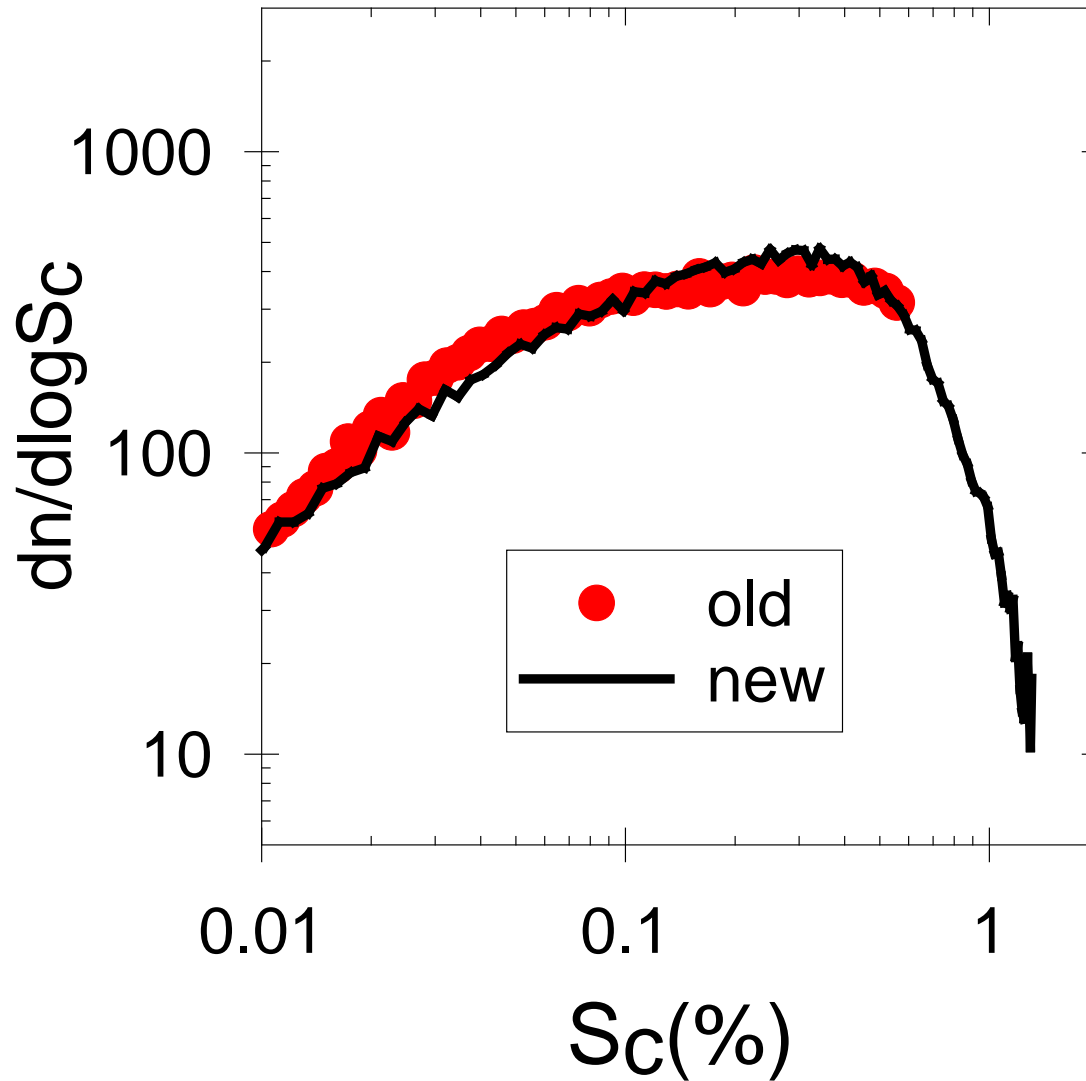
Nov 20, 2007; 1115:00-1117:00 MST



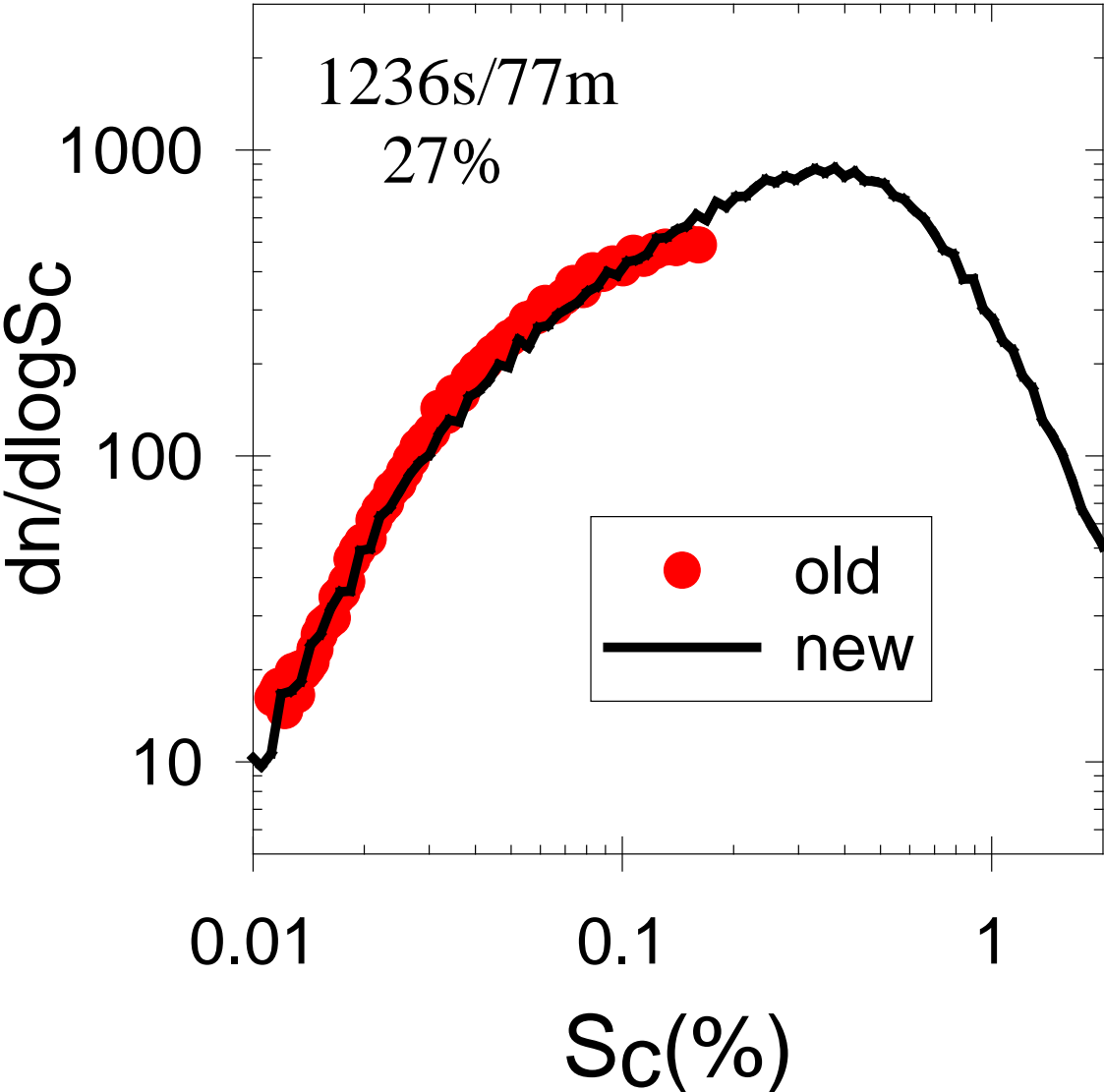
Nov 20, 2007; 1215:00-1230:00 MST



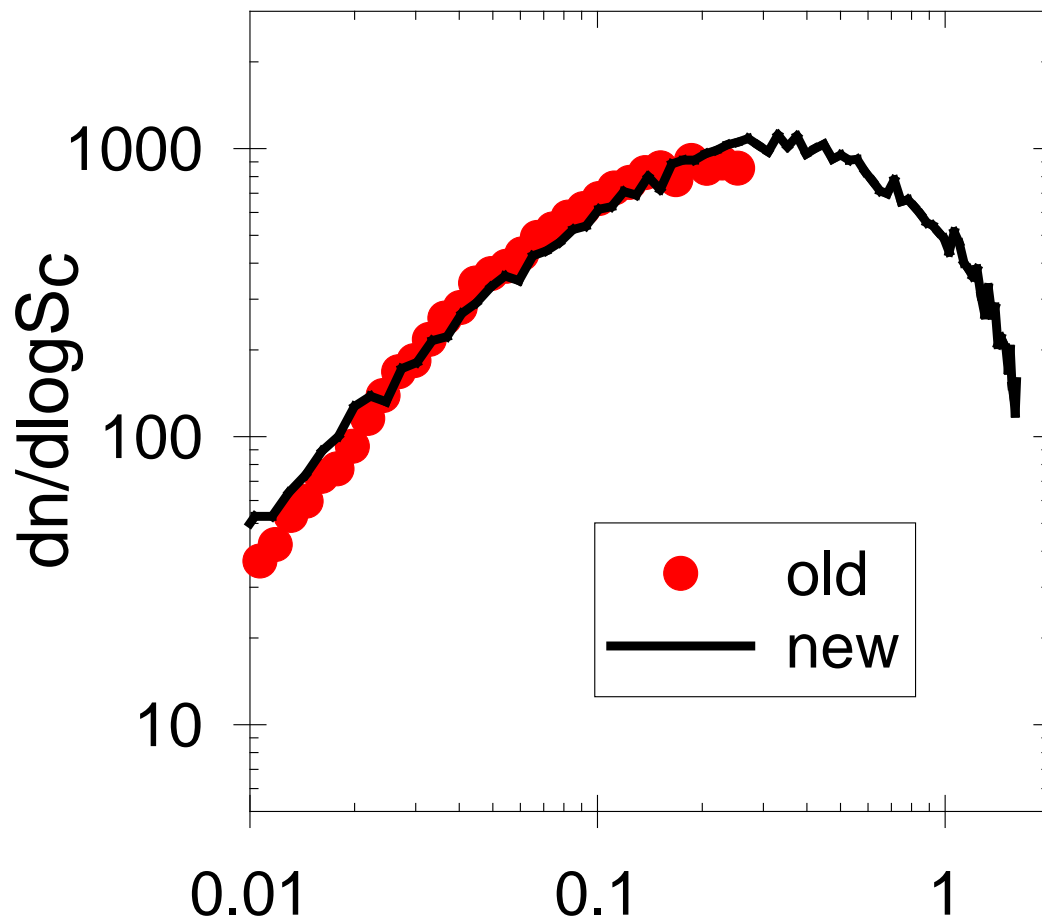
# July 22 below



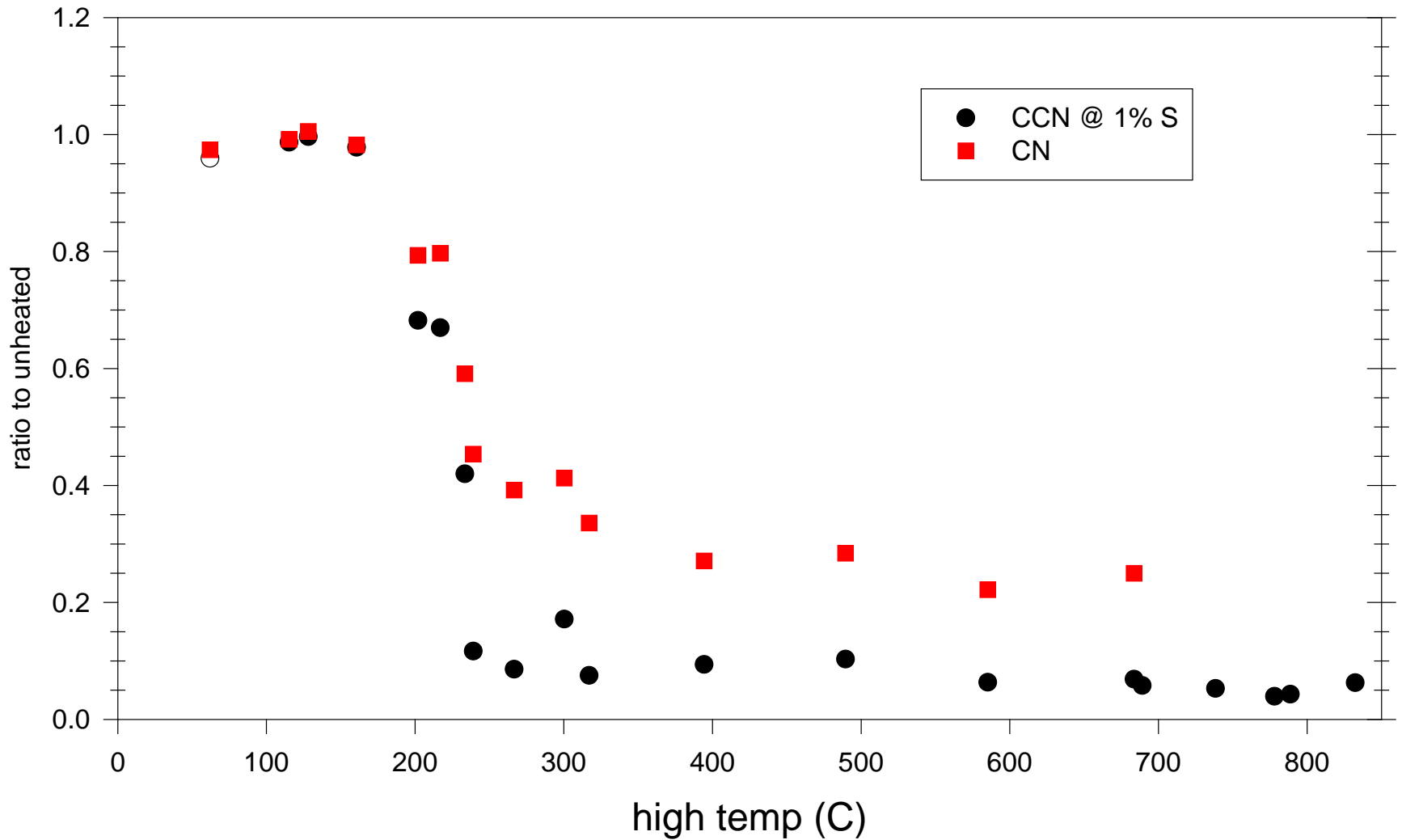
# July 18 below



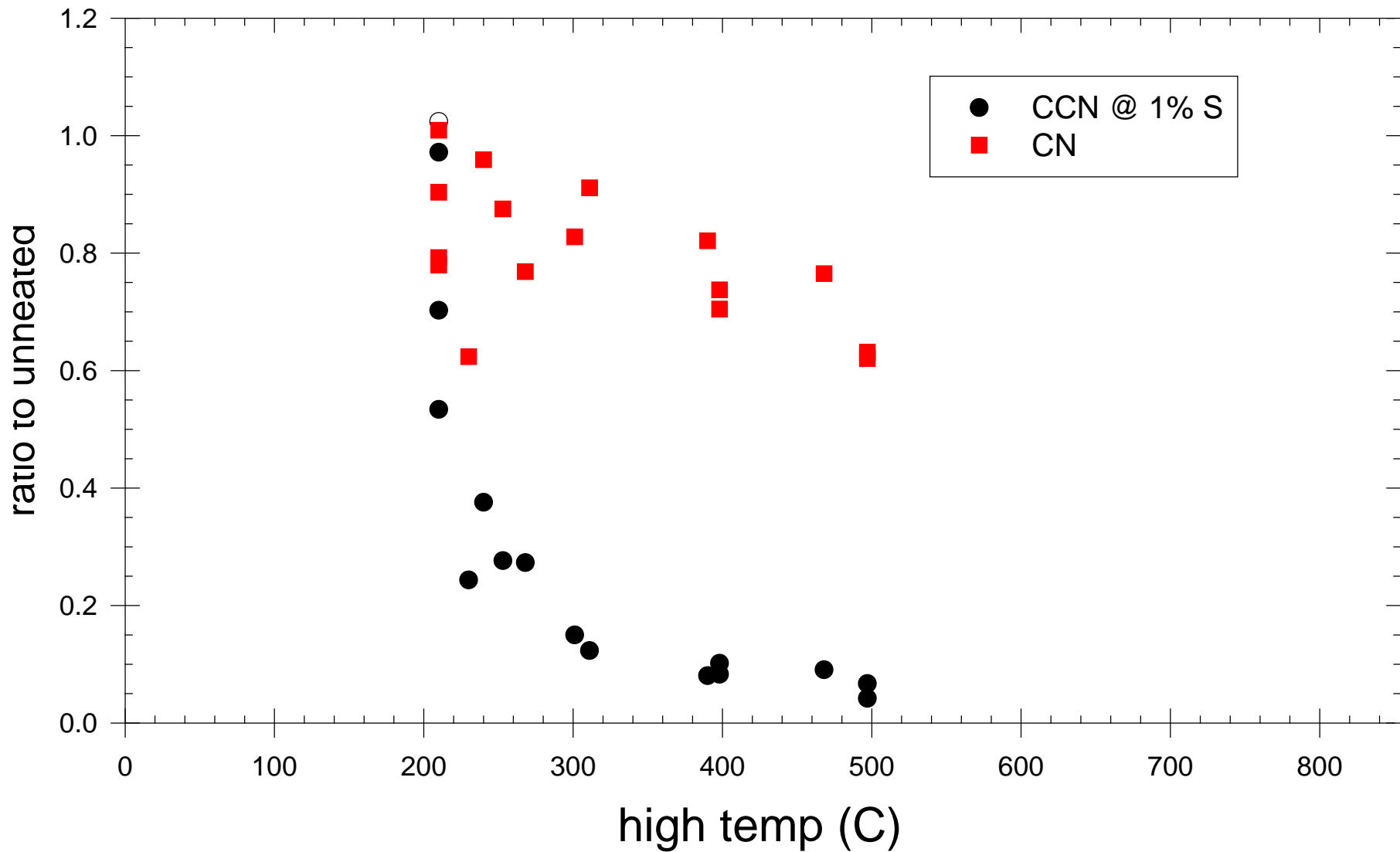
# July 17 above



Sept 2 2007 PASE RF11  
1231-1553 XST



July 22, 2005, 1042-1303  
MASE; near central California coast  
@ 955-976 mb





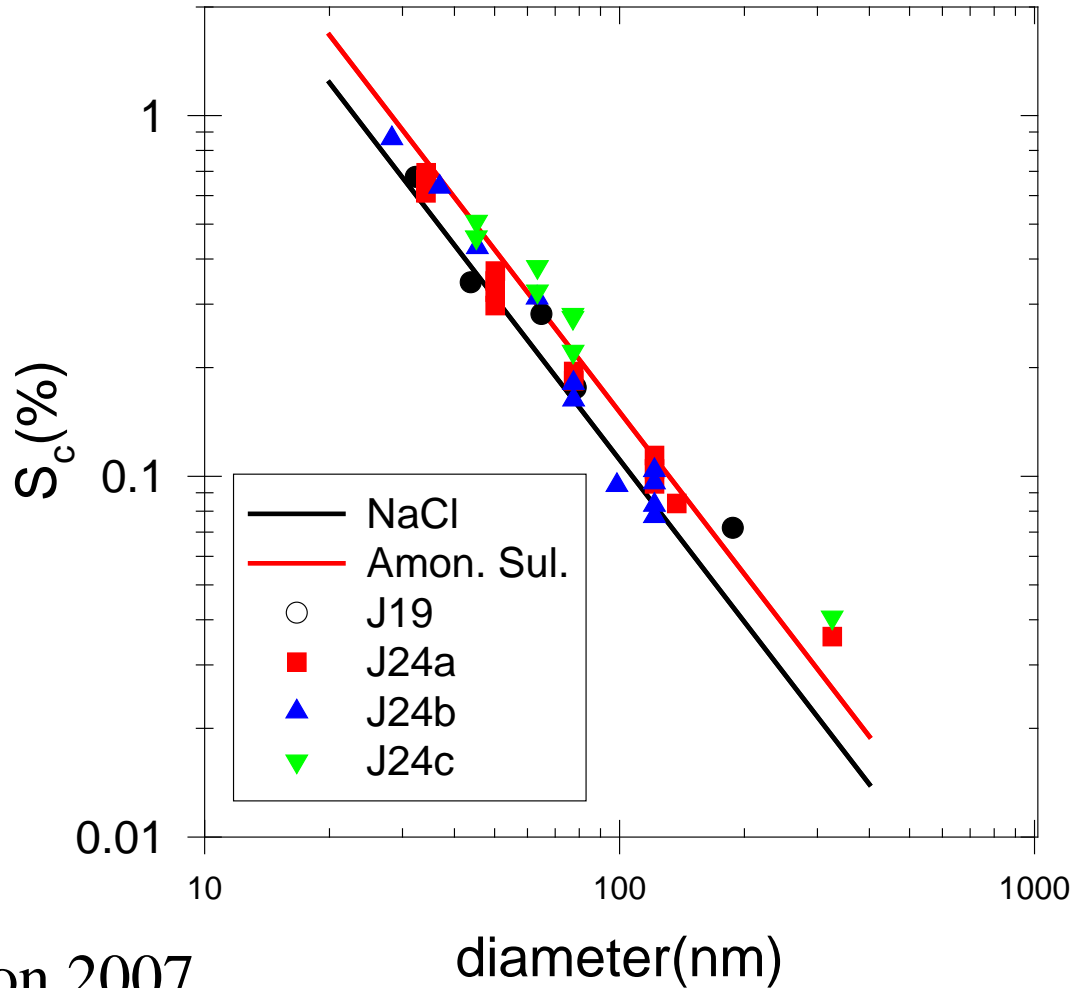
To compare different sizes or critical supersaturations ( $S_c$ ) Fitzgerald et al. (JAS 1982) introduced the hygroscopicity or solubility parameter,  $B$ , (non-dimensional).

$B = 1.23$  for NaCl

$B = 0.70$  for ammonium sulfate

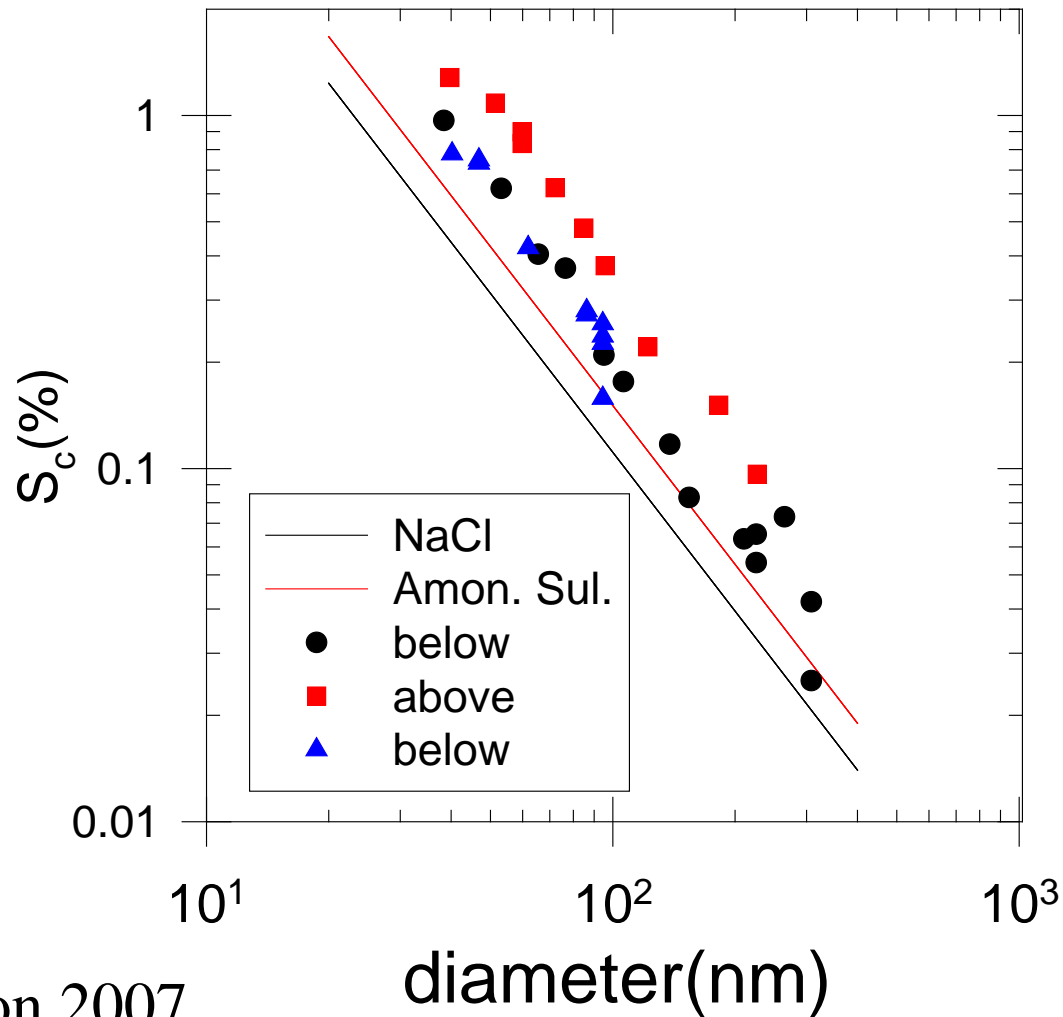
lower for less hygroscopic substances.

**Particle critical Supersaturation ( $S_c$ )  
versus dry particle diameter  
January 19 and 24, 2005  
RICO Eastern Caribbean near Antigua  
low altitude**



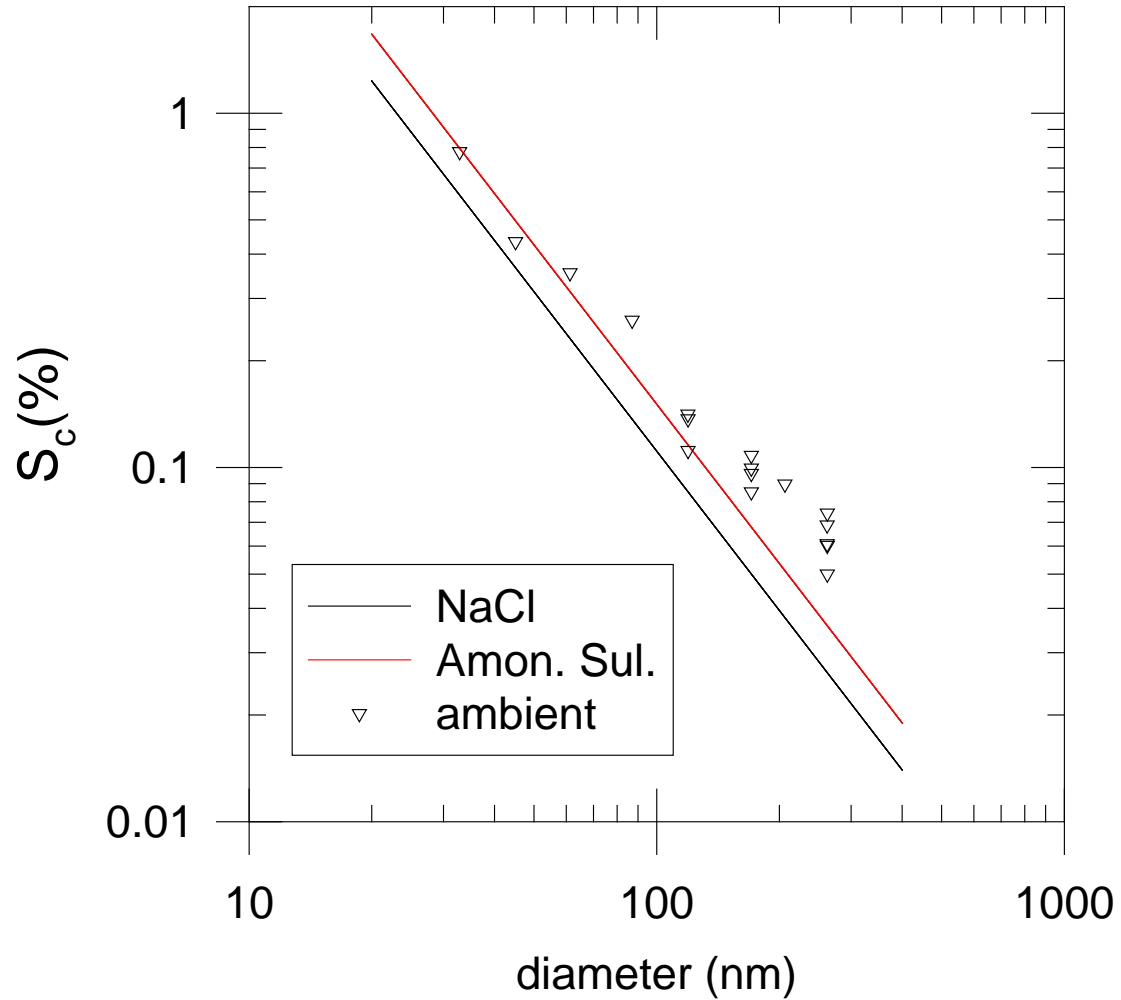
From Hudson 2007

**Particle critical supersaturation ( $S_c$ )  
versus dry diameter**  
July 25, 2005  
MASE off the Central California Coast  
below and above stratus cloud layer

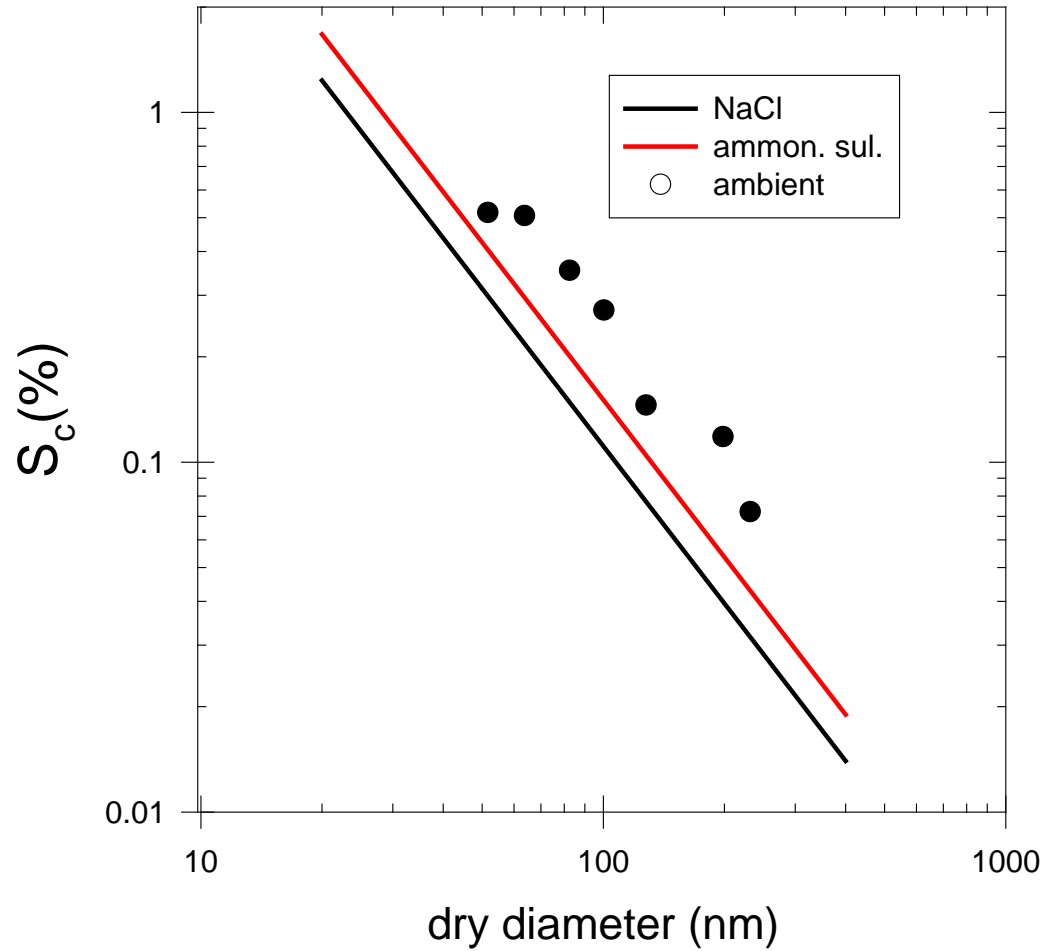


From Hudson 2007

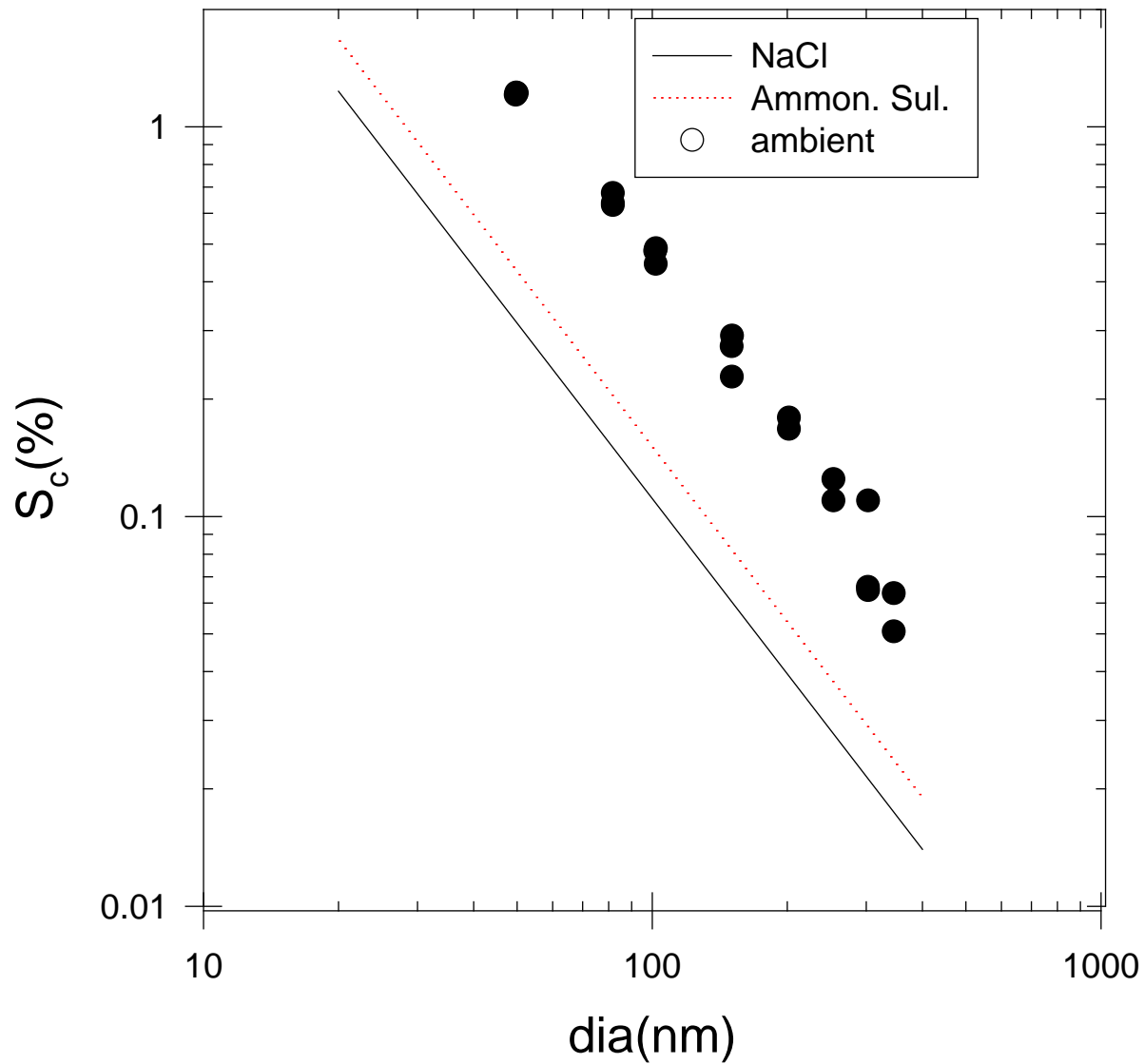
**Particle critical Supersaturation ( $S_c$ )  
versus dry diameter  
September 2, 2007 1417-1637 local time  
PASE near Christmas Island  
low altitude**



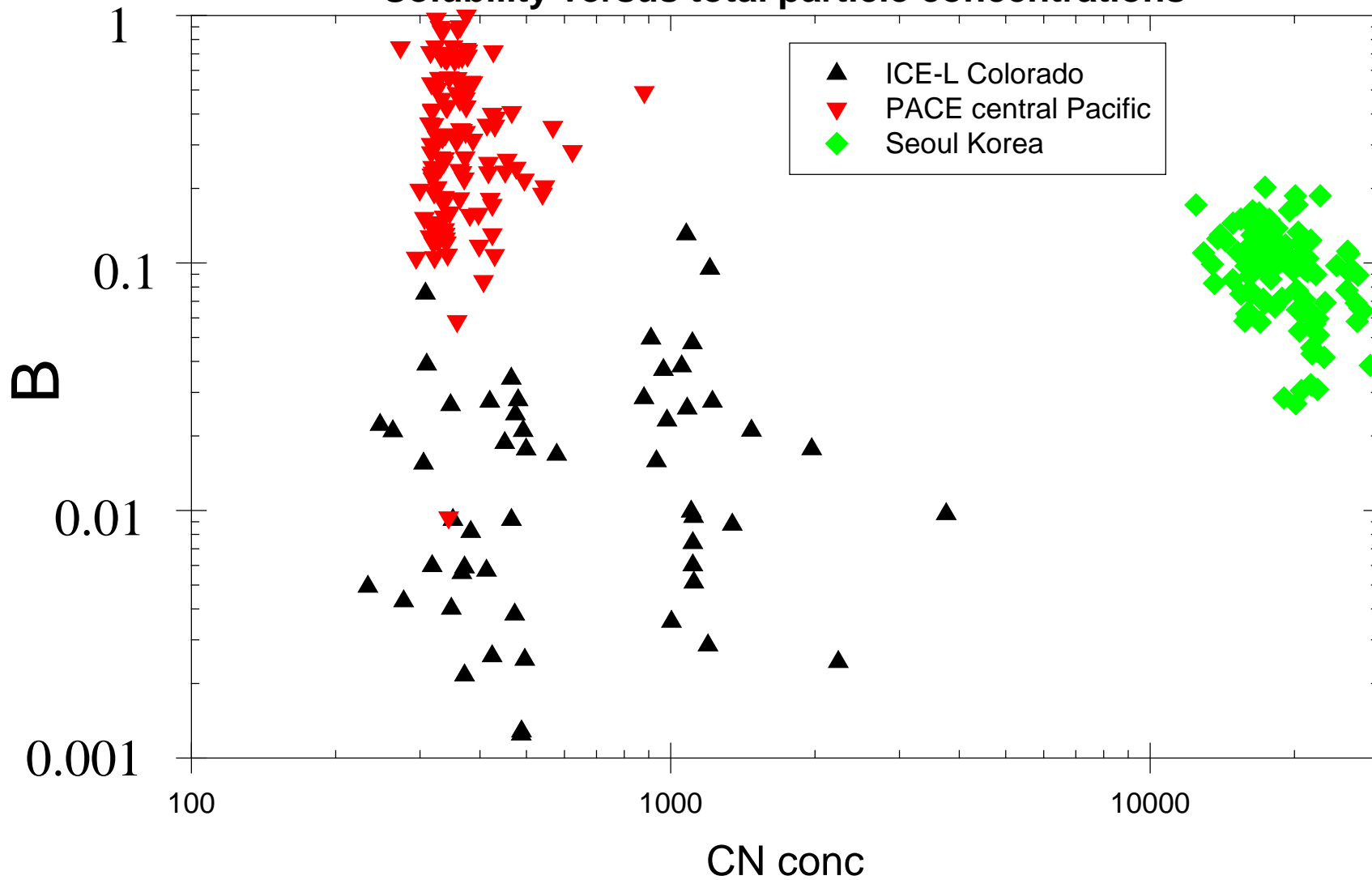
Aug 15, 2008 1121-1136  
Twin Otter, POST  
off central California coast  
size- $S_c$  ambient

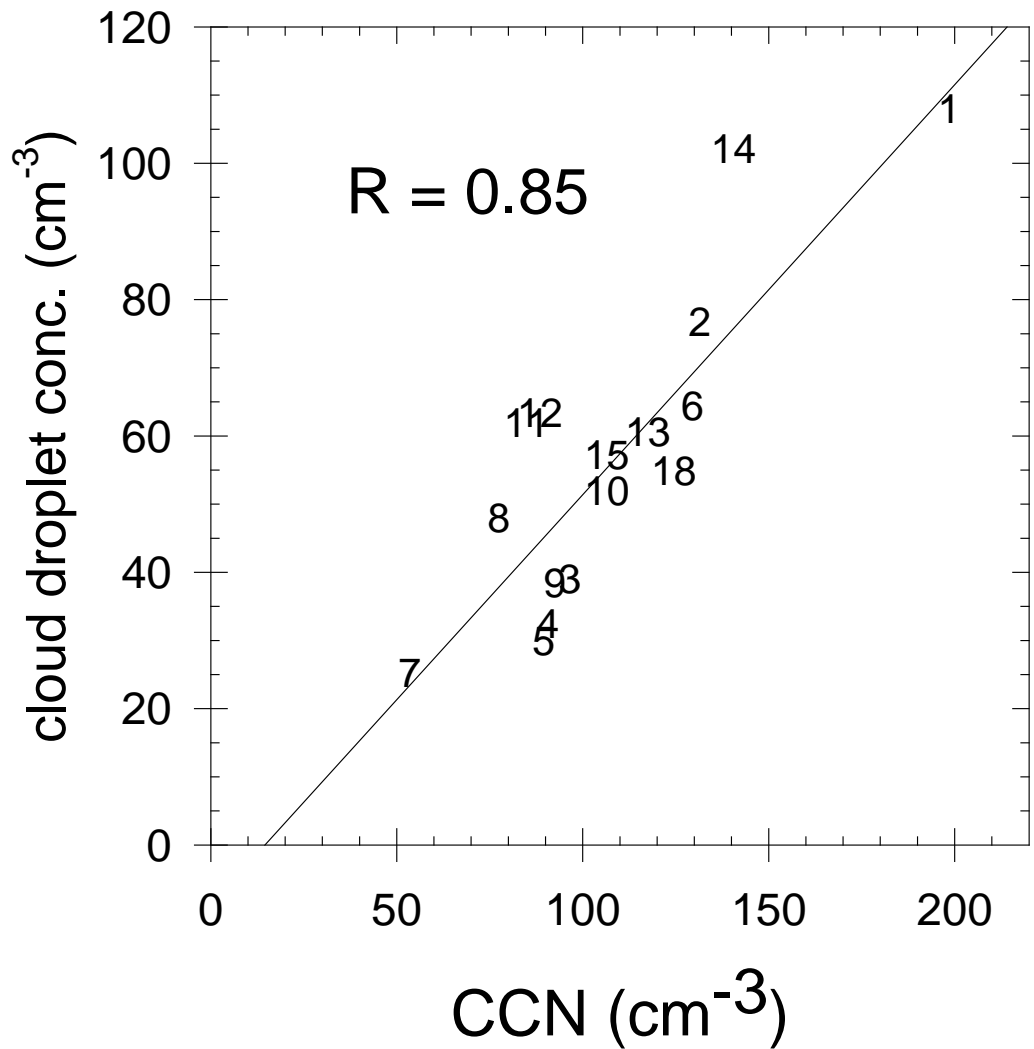


**Oct 21, 2006, 1854-1941**  
**Yonsei, Seoul, Korea**



# Solubility versus total particle concentrations

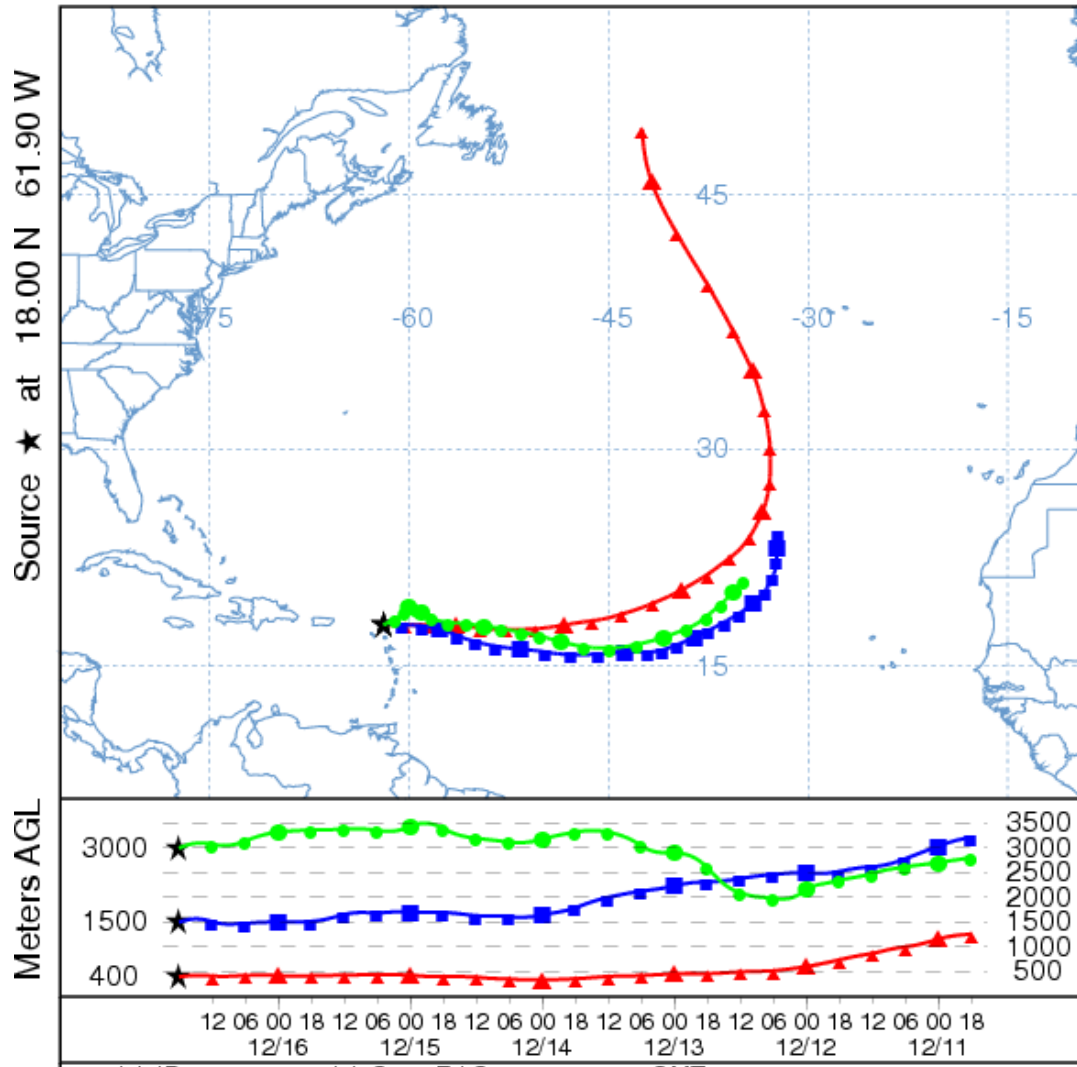




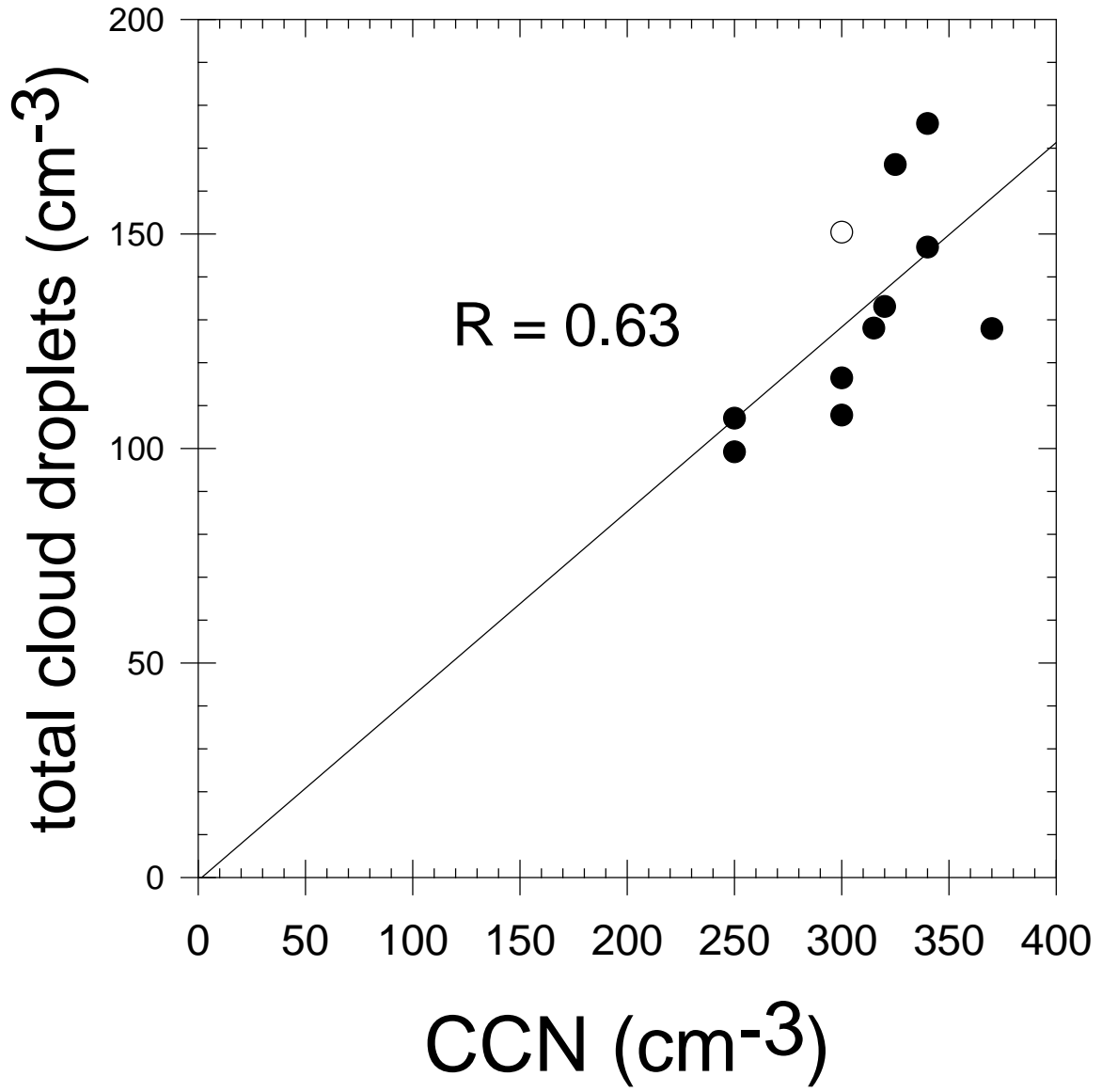
RICO flight ave  
Cloud droplet conc  
For LWC > 0.1gm<sup>-3</sup>  
Versus CCN conc



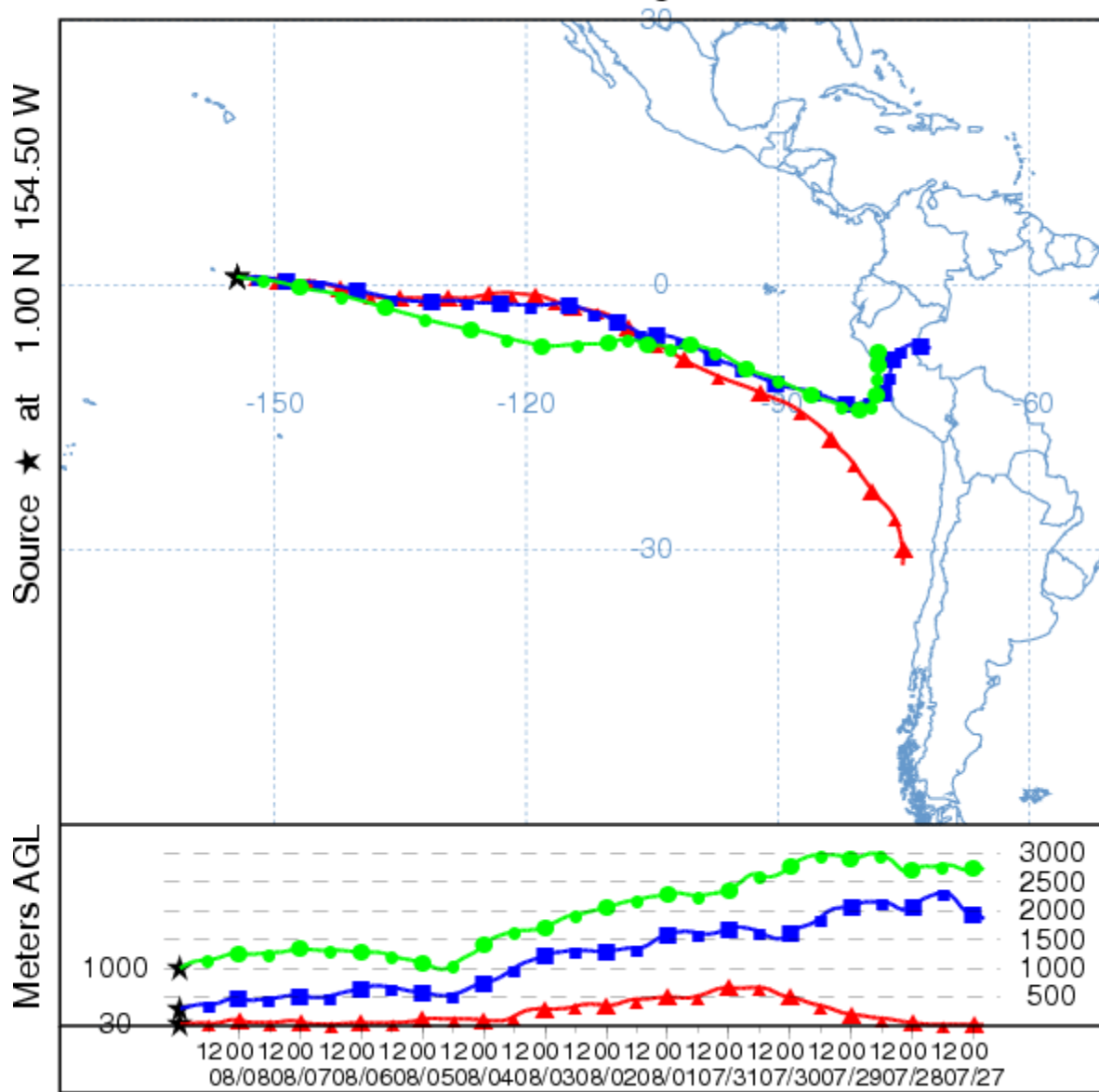
NOAA HYSPLIT MODEL  
Backward trajectories ending at 18 UTC 16 Dec 04  
FNL Meteorological Data



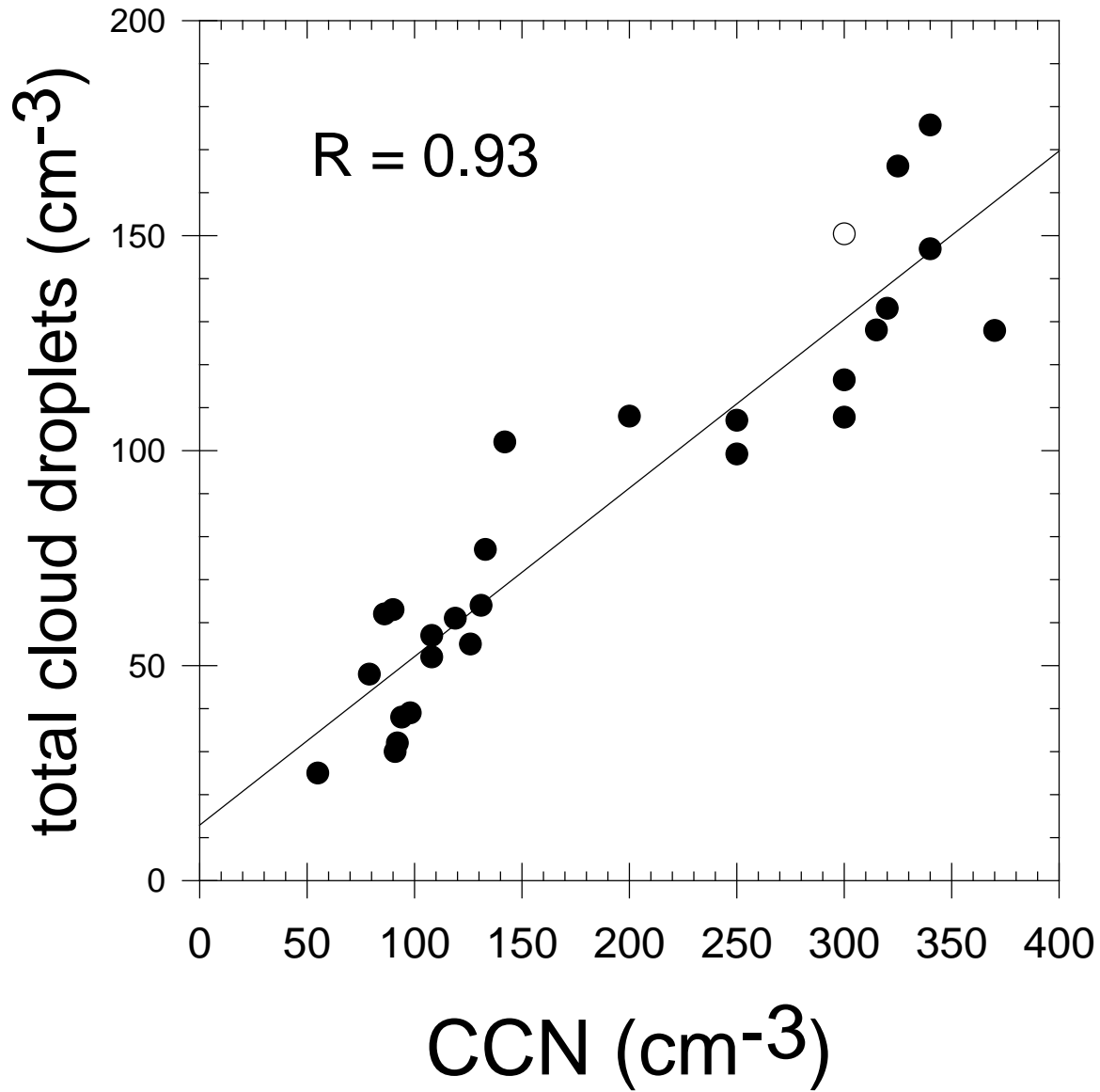
**PASE**  
**for LWC > 0.1 g m<sup>-3</sup>**

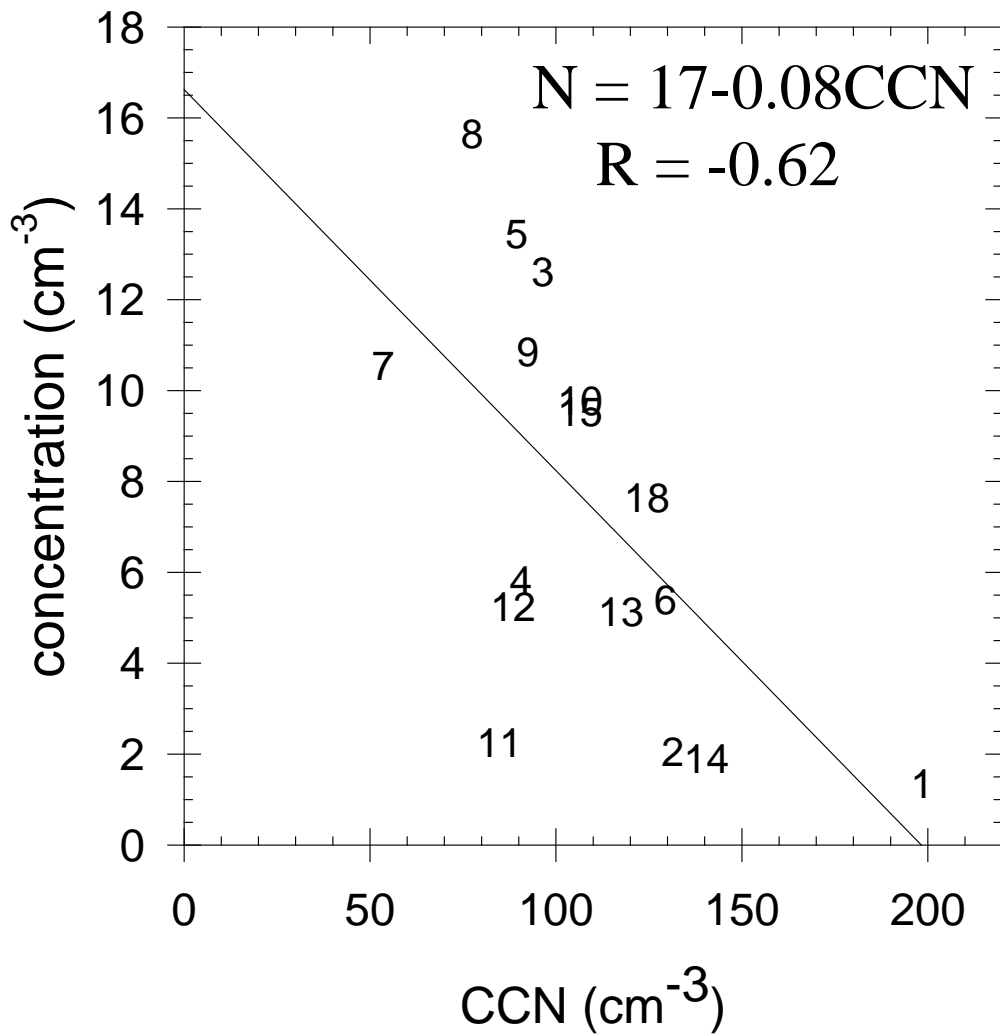


NOAA HYSPLIT MODEL  
Backward trajectories ending at 23 UTC 08 Aug 07  
CDC1 Meteorological Data

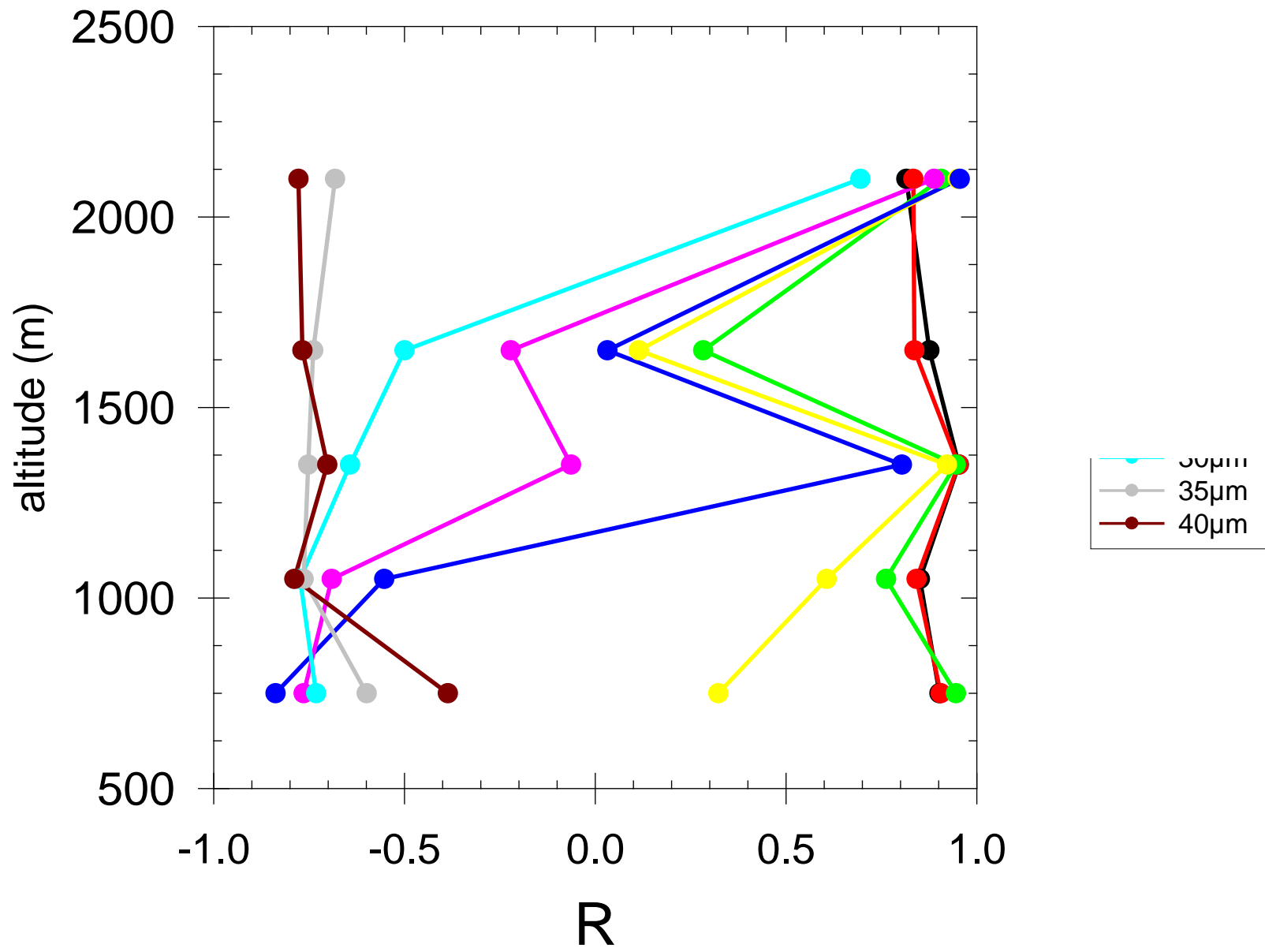


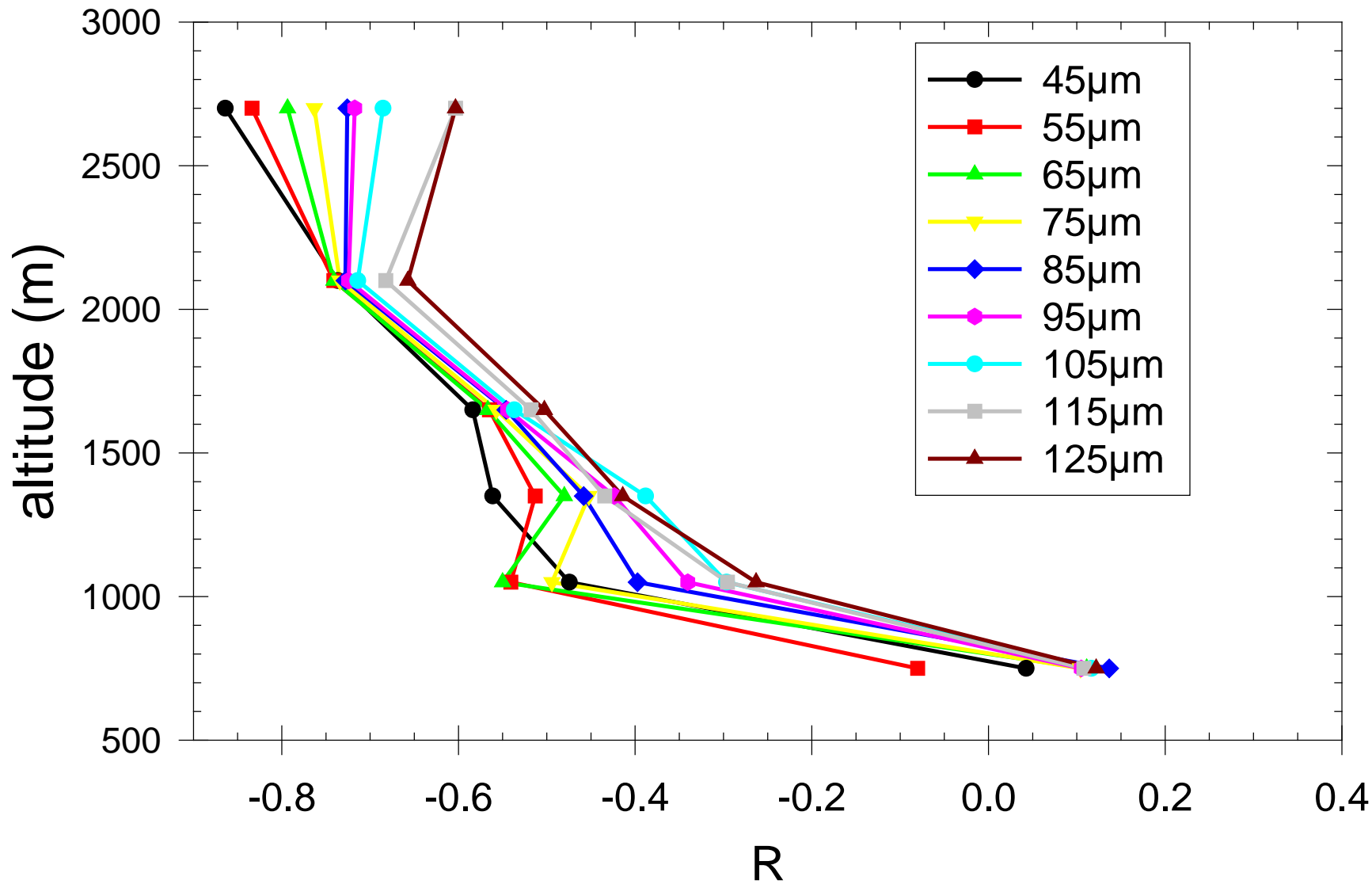
**RICO and PASE**  
**for LWC > 0.1 g m<sup>-3</sup>**

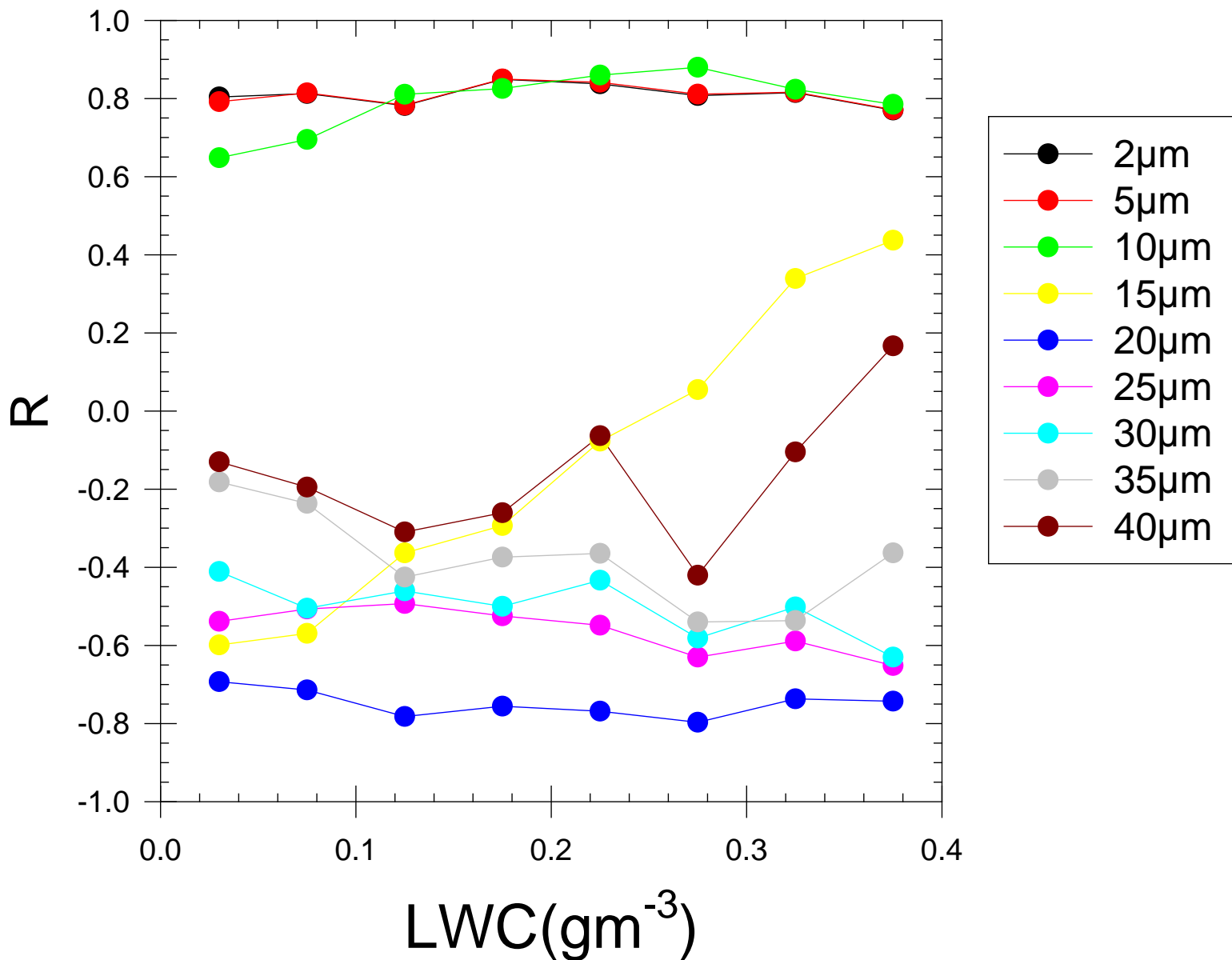




600-900m altitude and 16 flights  
with FSSP LWC > 0.1 gm<sup>-3</sup>  
droplets larger than 20 μm  
diameter. Note smaller  
droplet concentration range.

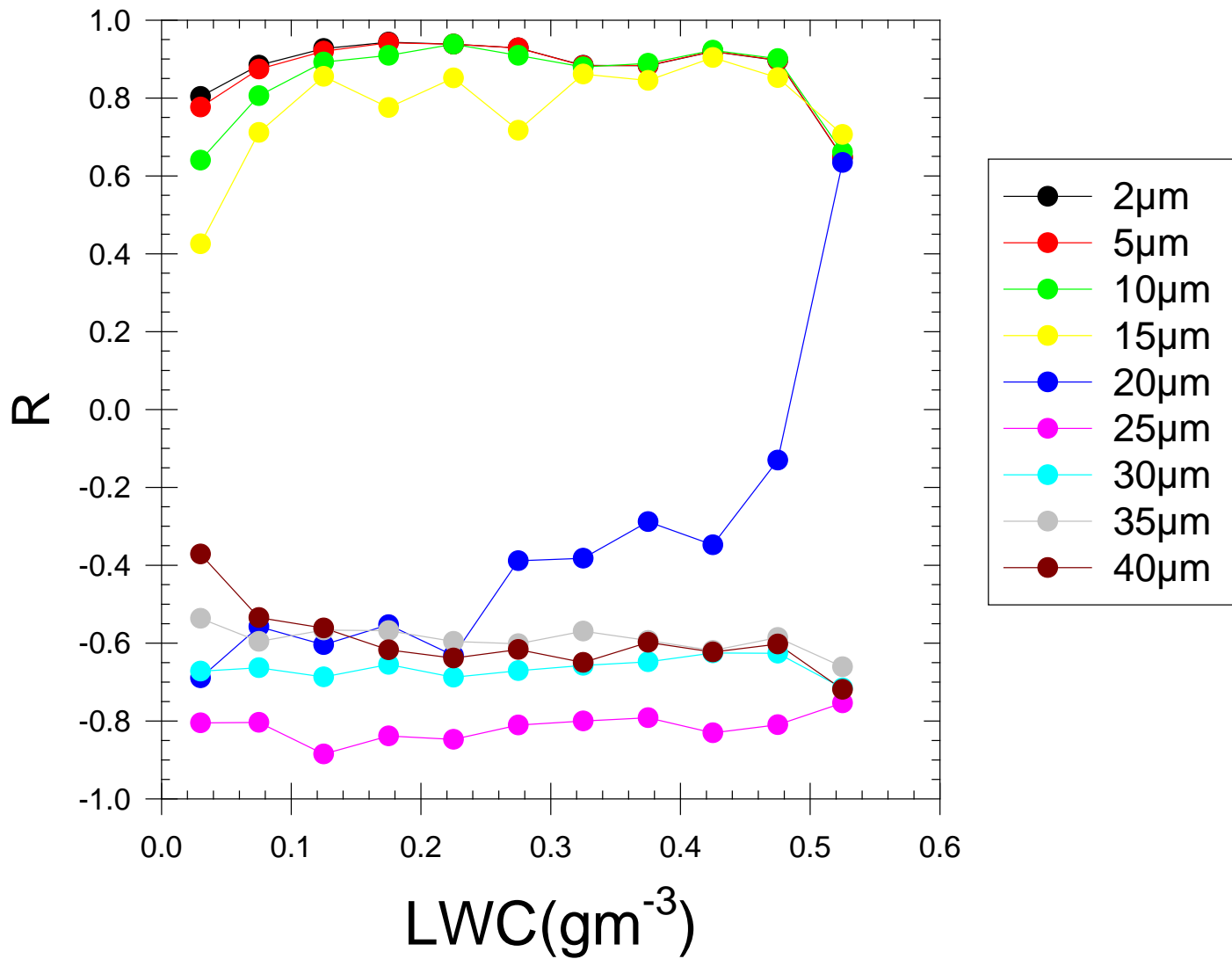




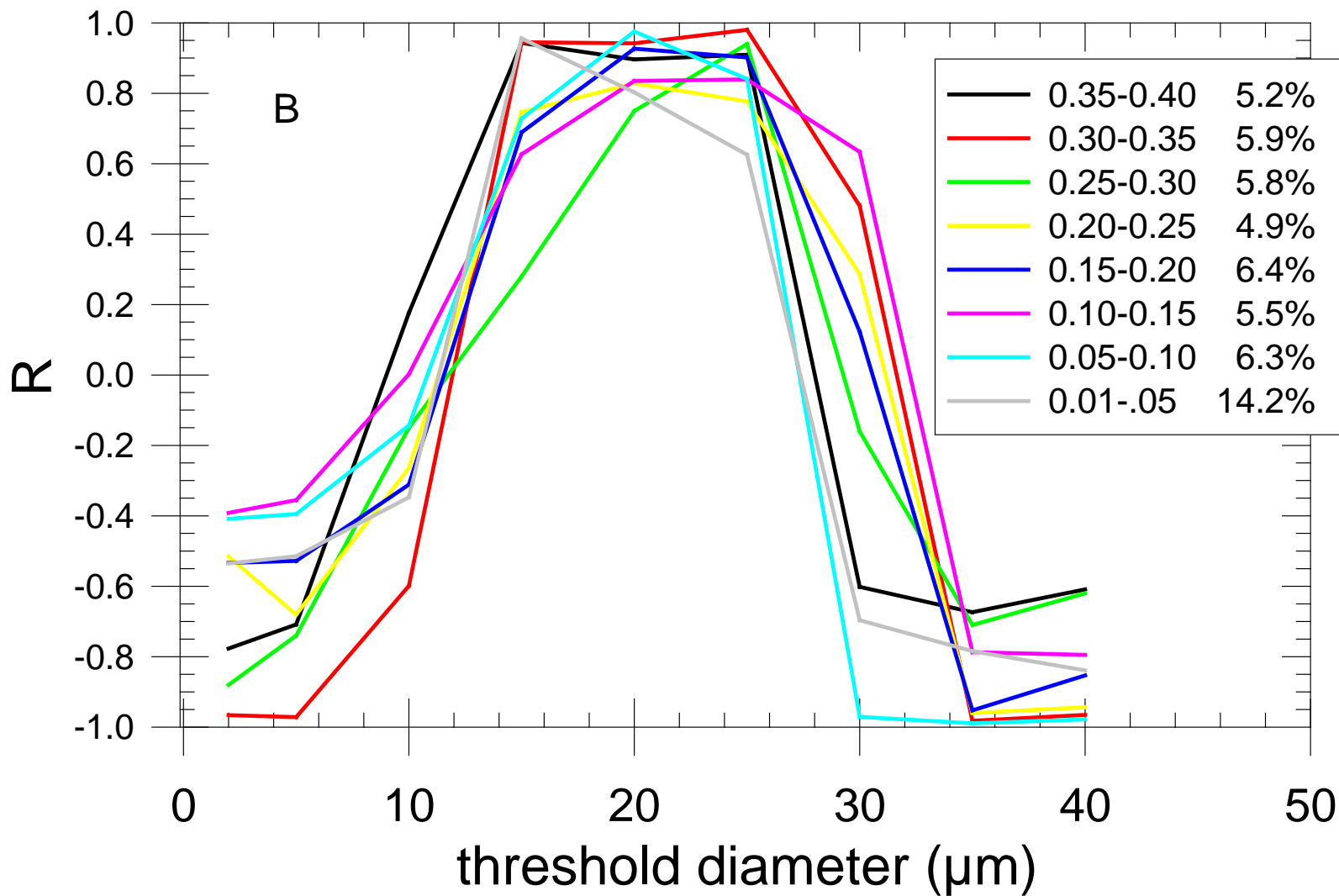


600-900 m altitude



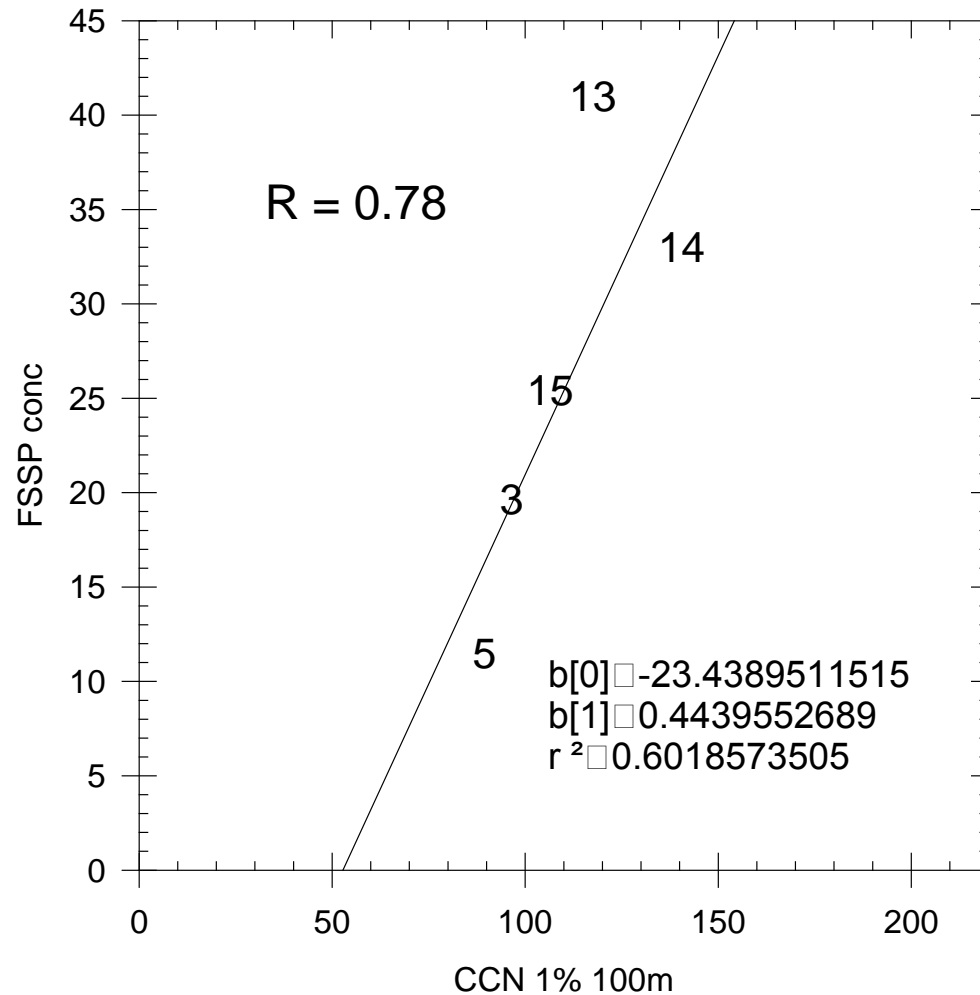


900-1200m altitude range.

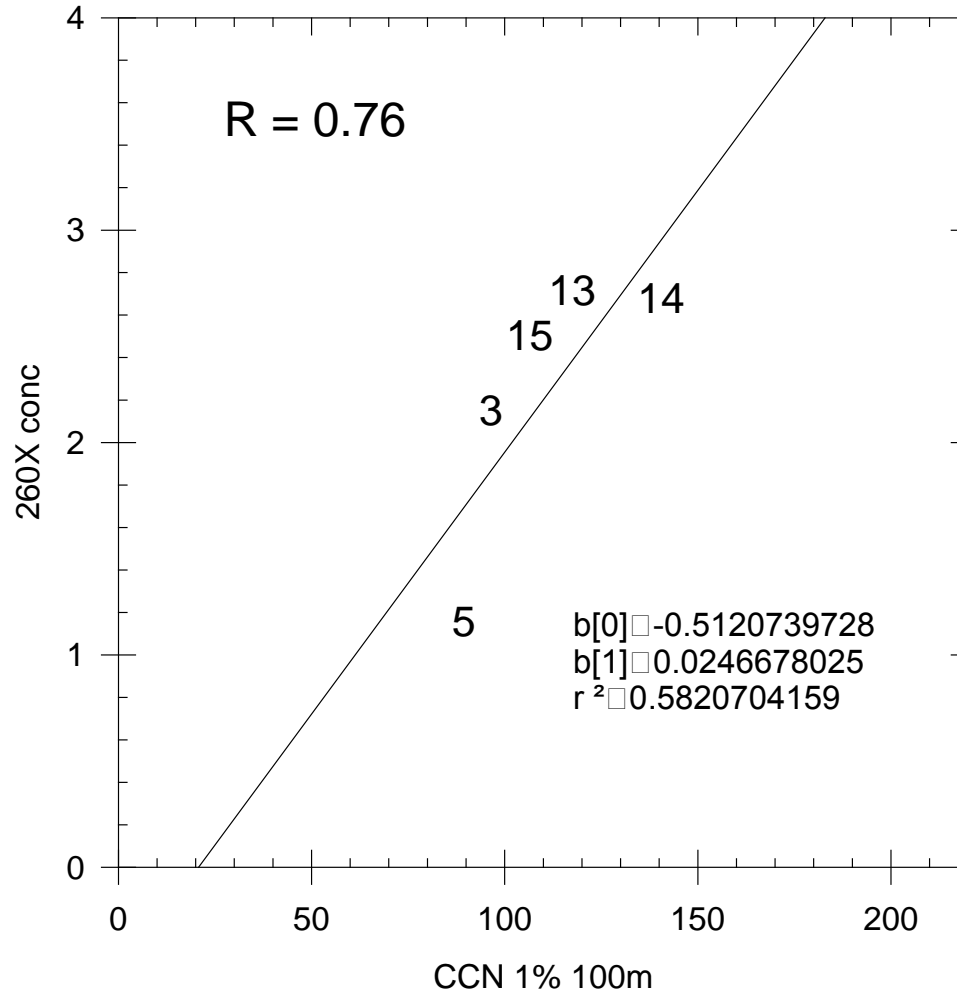


2400-3000m

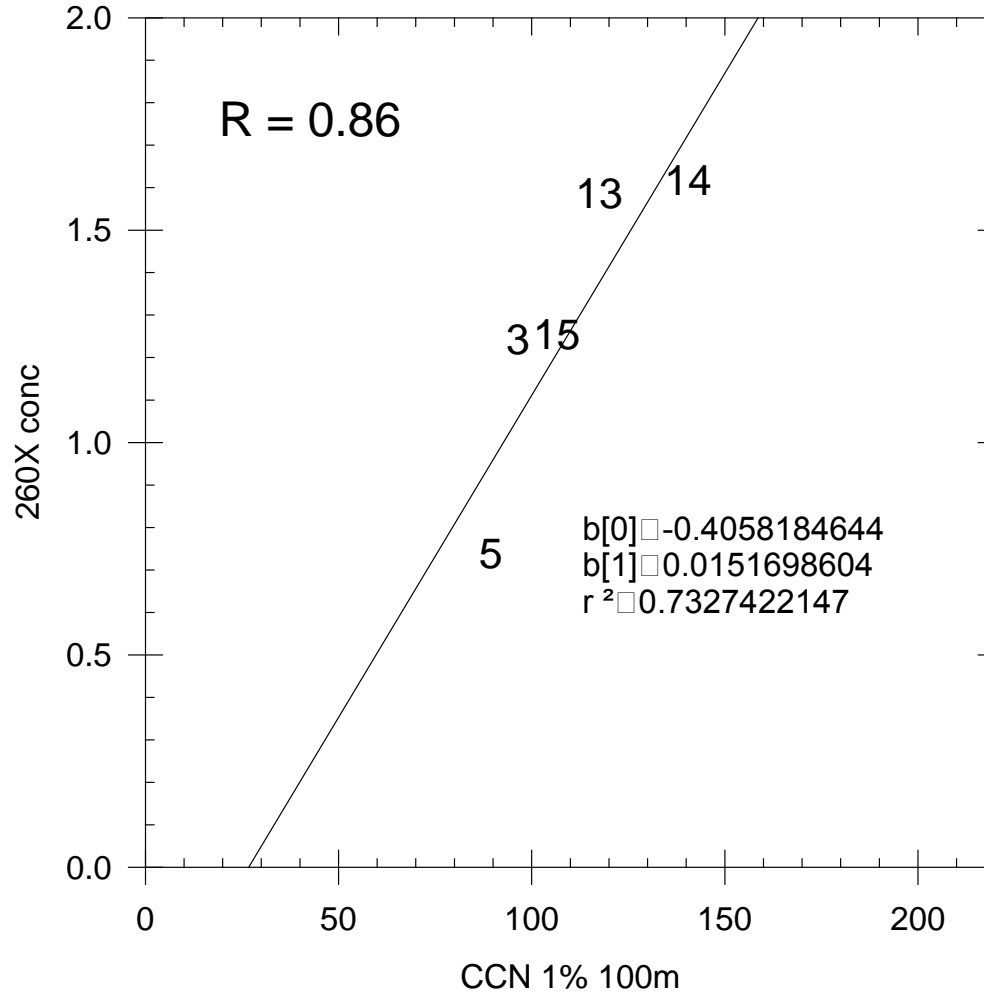
**RICO 2004-05, Antigua**  
**droplet concentration for all parcels with**  
**FSSP conc > 1, 3000m and up altitude**  
**FSSP conc all**  
**for 5 flights**



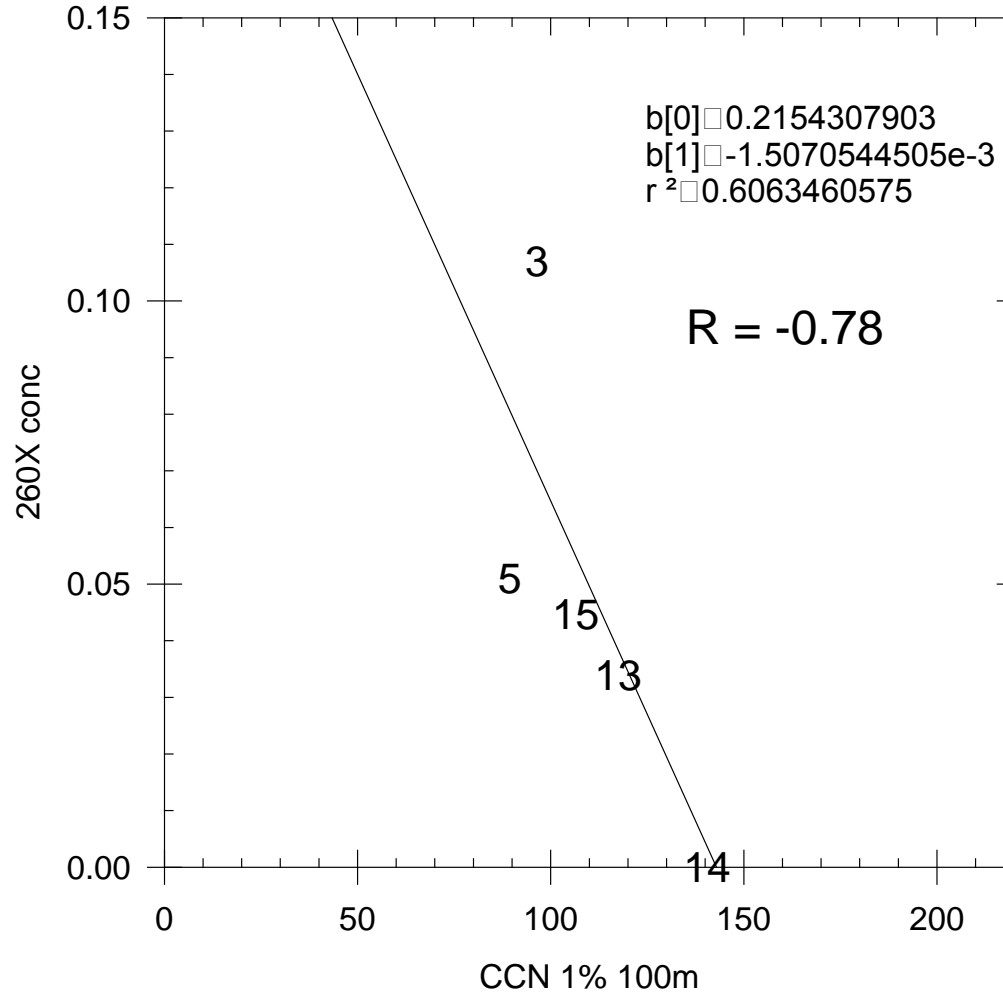
**RICO 2004-05, Antigua  
droplet concentration for all parcels with  
FSSP conc > 1, 3000m and up altitude  
260X conc > 295  $\mu\text{m}$   
for 5 flights**



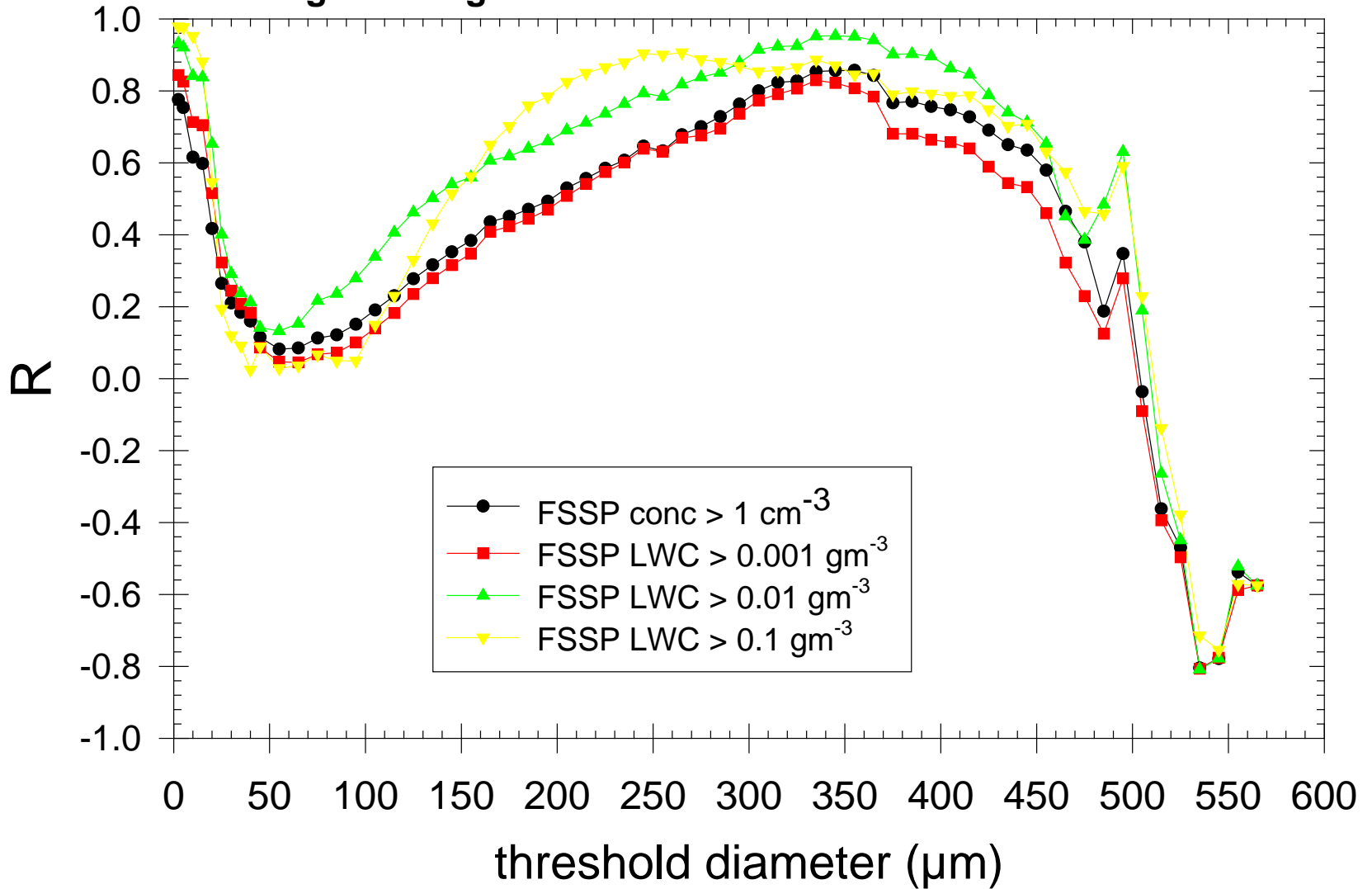
**RICO 2004-05, Antigua**  
**droplet concentration for all parcels with**  
**FSSP conc > 1, 3000m and up altitude**  
**260X conc > 245  $\mu\text{m}$**   
**for 5 flights**



**RICO 2004-05, Antigua  
droplet concentration for all parcels with  
FSSP conc > 1, 3000m and up altitude  
260X conc > 545  $\mu\text{m}$   
for 5 flights**



**Correlation coefficients (R) between CCN @ 1% concentrations at 100m altitude versus cumulative cloud droplet and drizzle drop concentrations within 3000+m altitude during the same five flights using various criteria**



3 CCN spectrometers

#1 (old) rectangular

S range 0.02-0.4%

#2 (new) rectangular

S range 0.1-1.5%

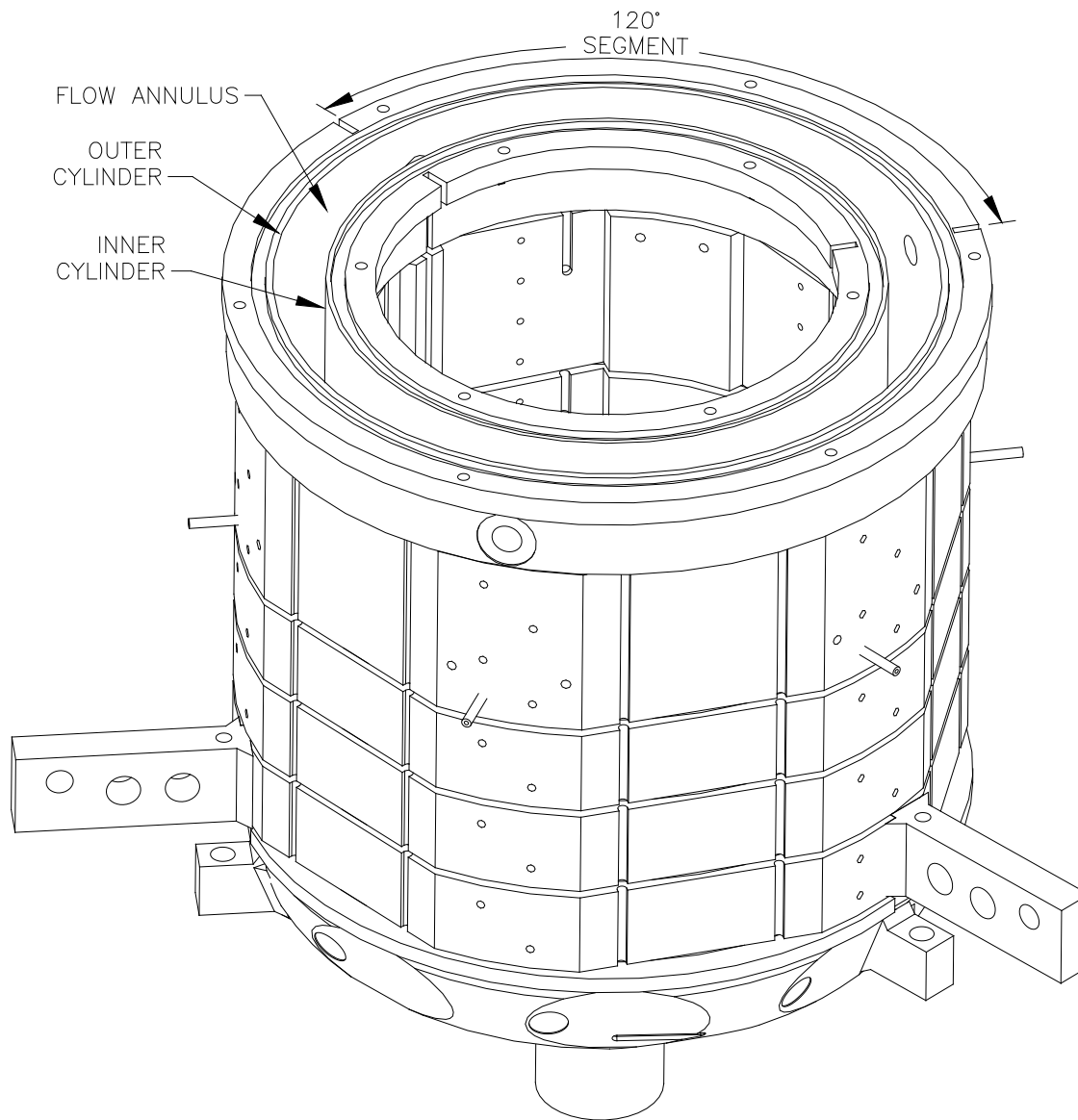
#3 double cylinder

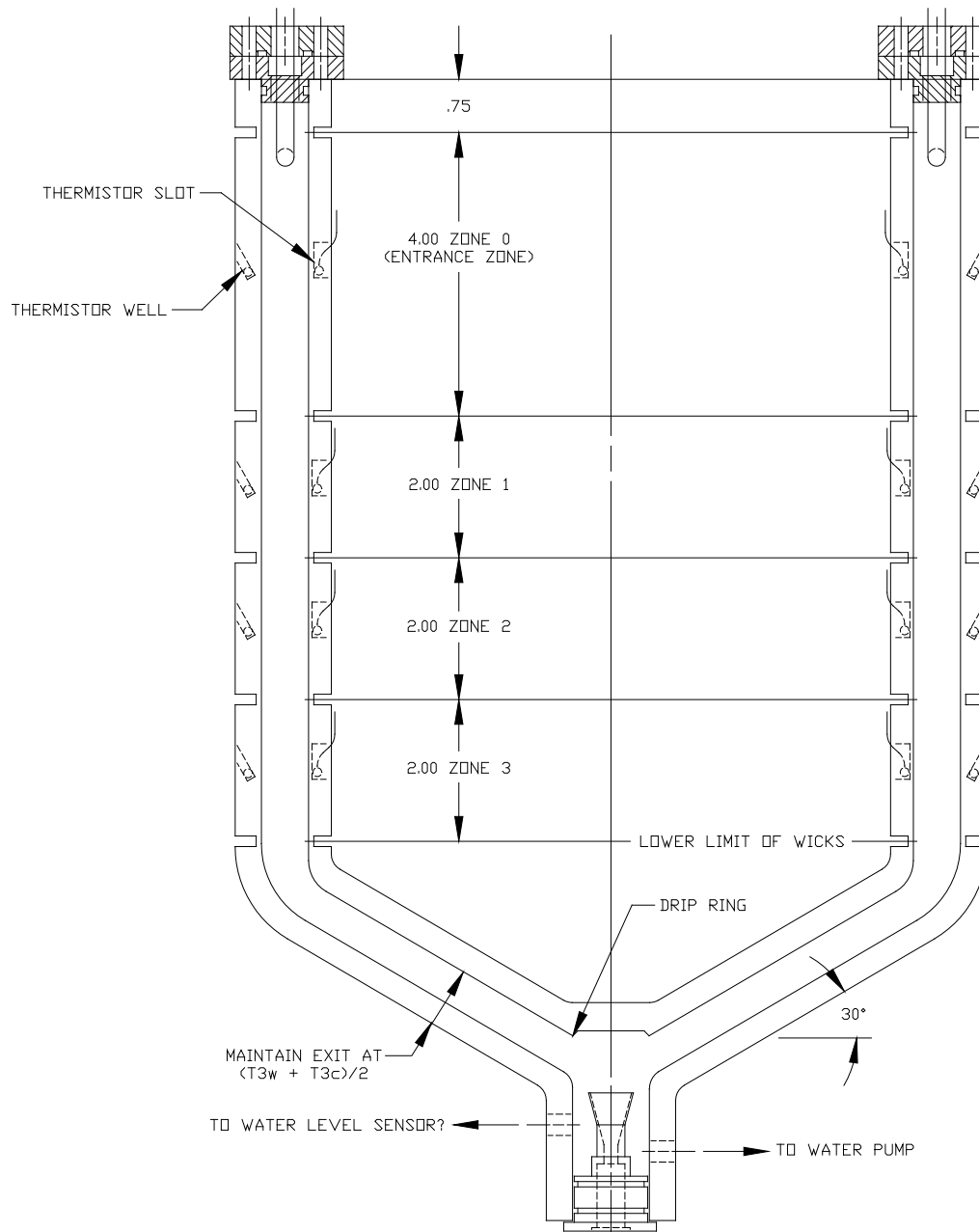
first aircraft deployment—ICE-L

Dec 07



1. Simultaneous high time resolution CCN spectra
2. Differential CCN spectra
3. Supersaturation (S) range extended below 0.1%.
4. Agreement between instruments





## DRI CCN Spectrometer #3

Cylindrical rather than rectangular

All sample has same path to OPC

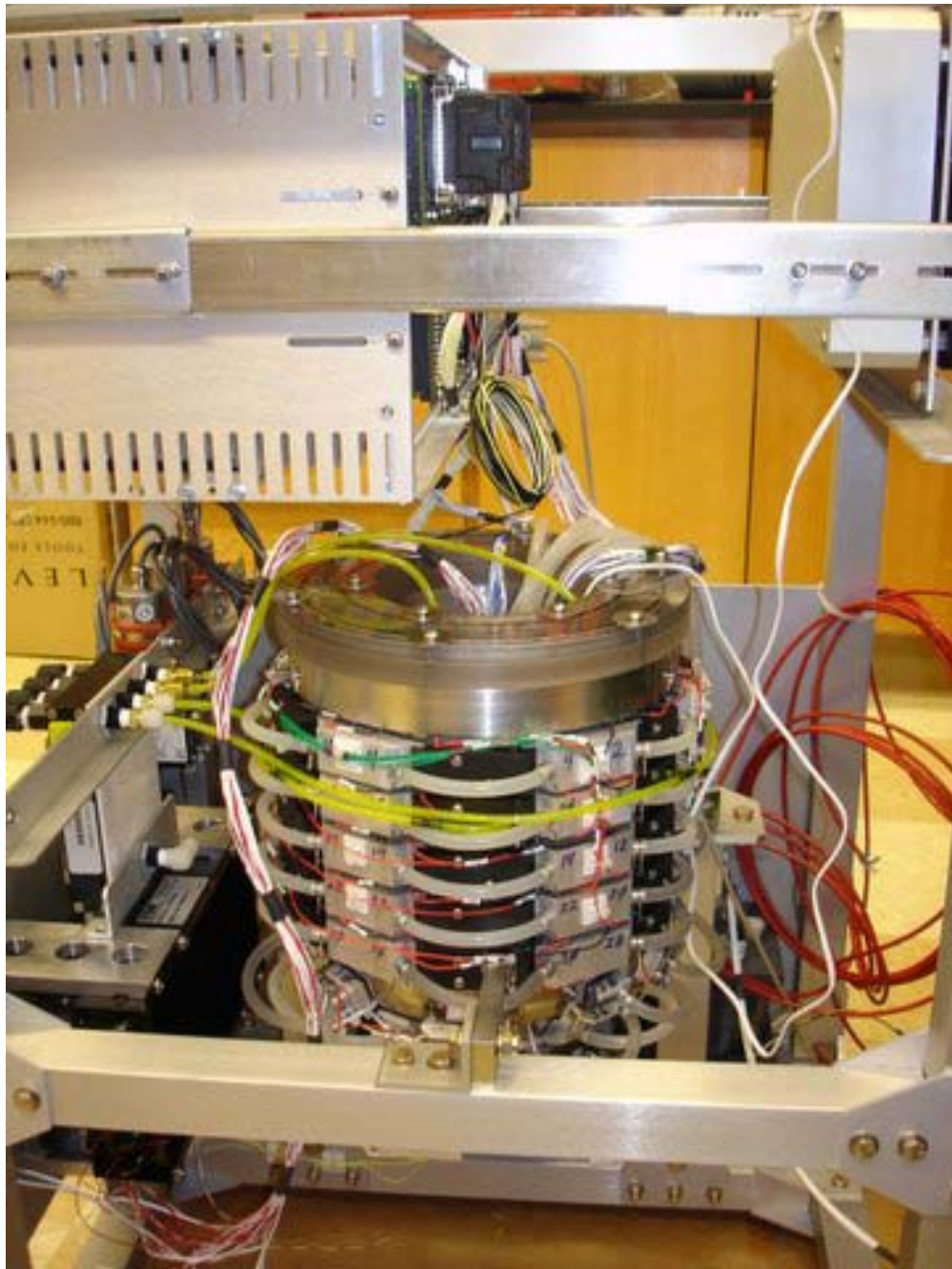
—should have less instrument broadening.

No edge/wall effects so sample can fill entire area between plates instead of confined to central area to avoid wall effects

—allows higher sample flow for the same chamber width/circumference.

Split azimuthally—provides two parallel chambers with separate sample flows with different S ranges—two chambers in one.

Has been aircraft tested on 5 C-130 flights in ICE-L in Dec. 07. Operated successfully.







But

Does not produce a narrower spectrum of calibration aerosol.  
Although convection is not a problem up to at least 12C delT  
in rectangular chamber it becomes a problem in cylindrical at 5C.  
Works at low S and does provide higher sample flow  
—better stats for low conc large particles.  
Try to stabilize and enhance S by decreasing the mean T  
downward (along the flow).  
But this decreases S.  
So try increasing T downward like DMT.  
Works but going against convection  
So should try inverting chamber (flow upward)  
Stable against convection  
Are streamwise and transverse  
(between plates) S additive?  
To produce higher S with  
limited del T.  
Is streamwise S enough? Will spectrum  
really be narrower?

Cylindrical or rectangular?