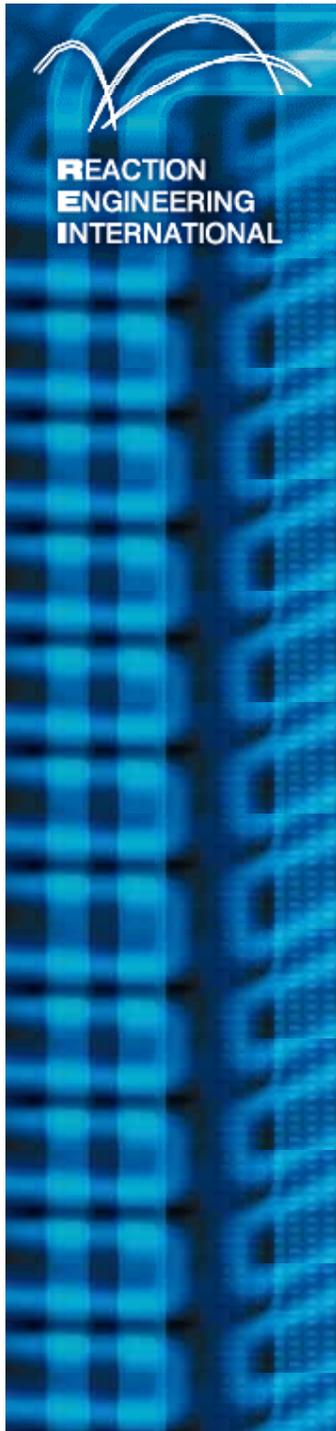


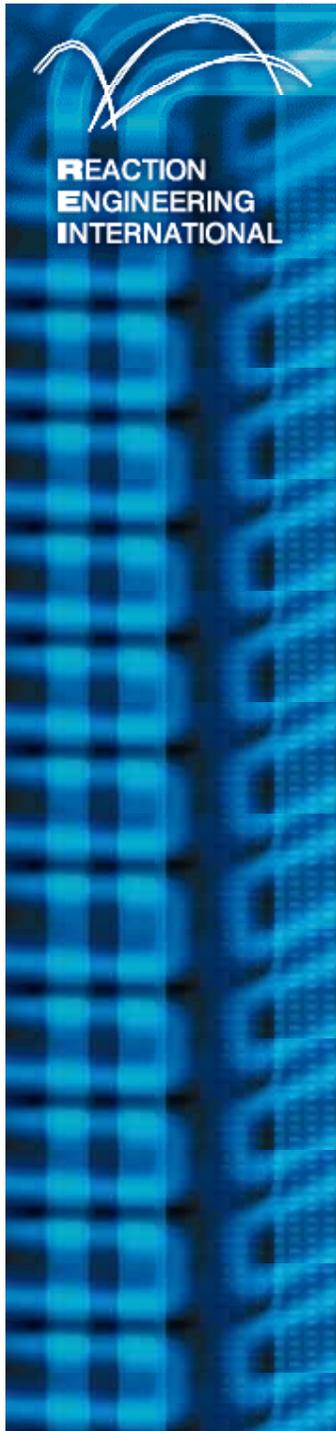
# Oxidation of Mercury Across SCR Catalysts in Coal-Fired Power Plants

C. L. Senior and T. Linjewile  
Reaction Engineering International



# Project Objectives

- Gather data on the behavior of mercury across SCR catalysts
  - Better understanding of Hg behavior
  - New model
- Measurements at power plant burning blend of bituminous, subbituminous coal
- Slipstream reactor with six catalysts
  - One blank honeycomb
  - Three commercial honeycomb catalysts
  - Two commercial plate catalysts



# Project Organization

- Slipstream reactor built under catalyst deactivation program (DOE- NETL)
- Mercury testing carried out under separate program (DOE – NETL)
- Additional support from EPRI and Argillon GmbH
- Field test support from AEP



# Project Team

REI: Planning/analysis, slipstream reactor operation

- Connie Senior, Temi Linjewile, Darren Shino, Dave Swensen

URS: Mercury measurement and analysis

- Carl Richardson, Mandi Richardson, Tom Mahalek

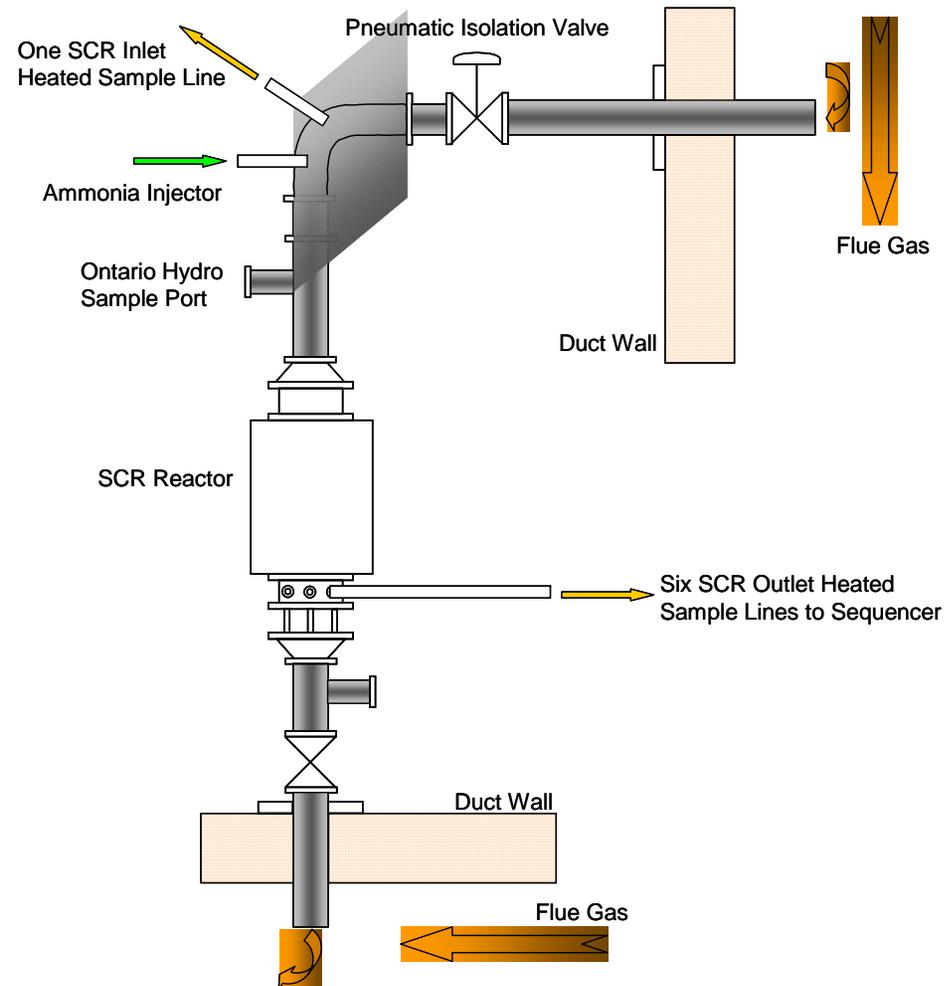
AEP: Field test support and program review

- Steve Pfeister, Steve Batie
- Gary Spitznogle, Aimee Toole

Program review

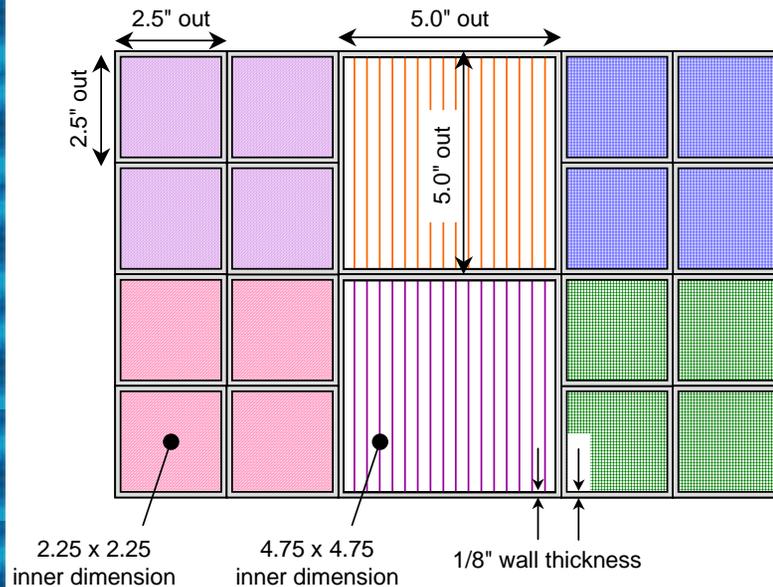
- José Figueroa, Bruce Lani, Lynn Brickett (DOE-NETL)
- Chuck Dene (EPRI)
- Jeanette Bock (Argillon GmbH)

# Multi-catalyst Slipstream Reactor



# Catalyst Dimensions

Chamber:	1 (Blank)	2	3	4	6	5
Catalyst type:	Monolith	Monolith	Plate	Plate	Monolith	Monolith
Chamber porosity:	58.7%	70.0%	85.0%	86.9%	70.0%	68.3%
Length of catalyst in chamber (inch):	24.40	21.50	39.25	43.25	20.06	19.75



- Five commercial catalysts
- One blank cordierite honeycomb

# Testing Summary

- AEP Rockport:
  - Two 1300 MW<sub>e</sub> B&W opposed-wall, supercritical boilers
  - Testing on Unit 1 across air preheater
  - Burn a subbituminous-bituminous blend
- Two test series (March and August)
- Measurements
  - Coal, economizer ash, ESP ash composition
  - Ontario Hydro measurements at inlet to slipstream
  - SCEM measurements at inlet/outlet of catalyst chambers
  - NO<sub>x</sub> and O<sub>2</sub> at inlet/outlet of catalyst chambers
  - Carbon trap and acid gas measurement at inlet of catalyst

# Coal Properties

Date	3/28/03	4/1/03	4/2/03
<b>(As Received):</b>			
Carbon	50.67	51.80	51.75
Hydrogen	3.51	3.64	3.46
Oxygen	10.89	11.04	11.18
Nitrogen	0.76	0.78	0.75
Sulfur	0.32	0.30	0.37
Ash	5.12	5.99	6.10
Moisture	28.74	26.45	26.39
HHV	8,723	8,989	8,989
<b>(Dry Basis):</b>			
Hg, ug/g	0.088	0.118	0.091
Cl, ug/g	120	160	200
SO <sub>2</sub> , lb/MBtu	0.74	0.67	0.82
Hg, lb/TBtu	10.10	13.13	10.13
Hg, ug/dnm <sup>3</sup> (5%O <sub>2</sub> )	8.02	10.82	8.46

- Coal blend – mostly subbituminous
- Higher Cl than typical subbituminous
- 8-10 μg/dnm<sup>3</sup> Hg (gas-phase equivalent)
- Ash contains ~6 wt% Fe<sub>2</sub>O<sub>3</sub>, ~16 wt% CaO

# Flue Gas Composition

(Inlet to Slipstream Reactor)

## Calculated from coal:

HCl 6-12 ppm (5% O<sub>2</sub>)

SO<sub>2</sub> 275-325 ppm (5% O<sub>2</sub>)

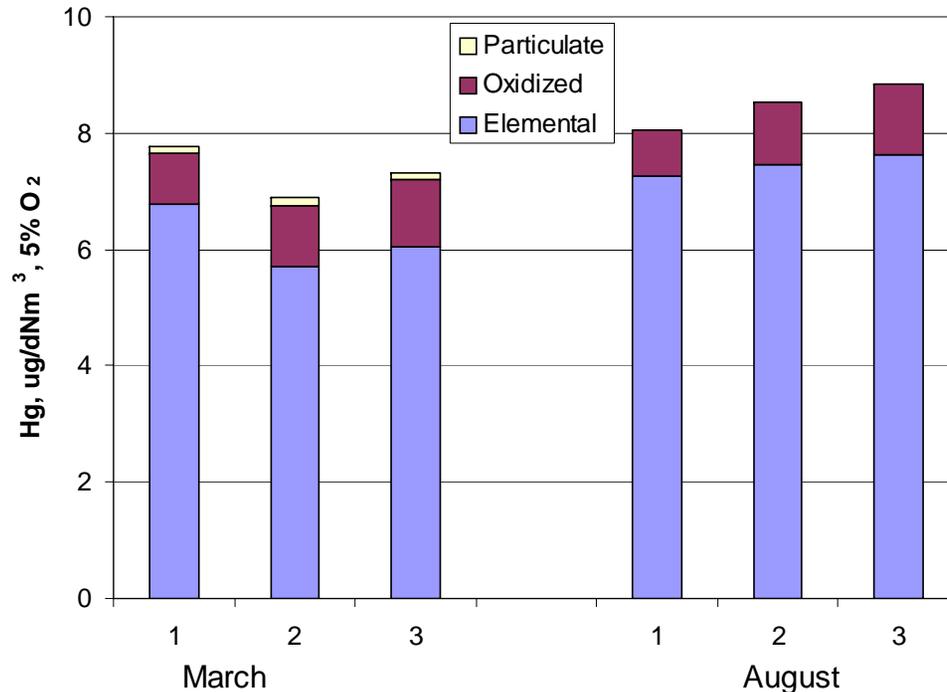
## Measured:

NO<sub>x</sub> 300-350 ppm (5% O<sub>2</sub>)

Total Hg 7-9 ug/dNm<sup>3</sup> (5% O<sub>2</sub>)



# Ontario Hydro Data



- Hg concentration in OH ash higher than in ESP fly ash BUT fraction of Hg in particulate very low
- 80-90% elemental Hg at inlet to catalysts

# Hg and Cl in ash

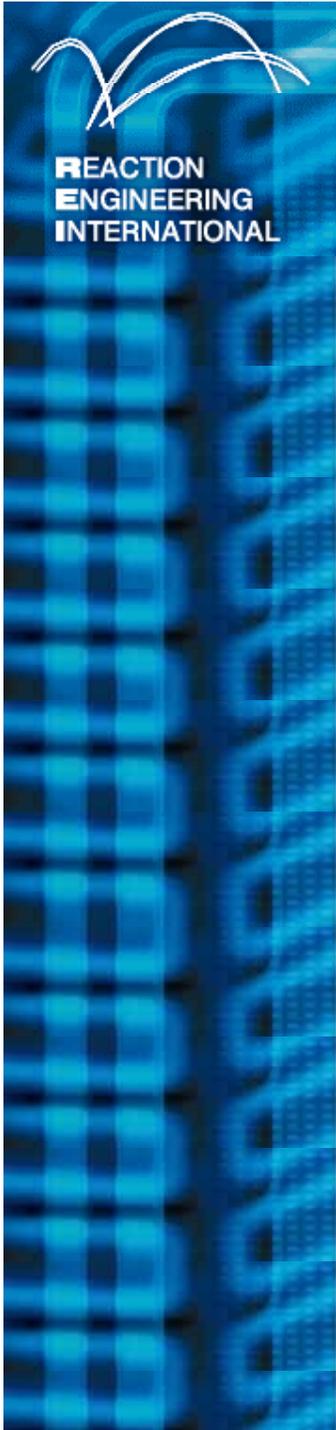
Date	LOI, wt%	Hg, ug/g	Cl, ug/g
<b>Economizer</b>			
3/28/03	0.08%	0.005	29
8/11/03	0.00%	0.005	<5
8/15/03	0.00%	0.000	<5
<b>ESP Hopper</b>			
3/28/03	0.31%	0.081	20
3/31/03	0.37%	0.118	25
4/1/03	0.31%	0.127	24
4/2/03	0.34%	0.101	27
8/7/03	0.06%	0.034	21
8/11/03	0.30%	0.050	21
8/15/03	0.13%	0.055	23

- Economizer ash has 10-20 times less Hg than ESP ash
- Little Cl on economizer ash
  - Expect most Cl in gas phase at inlet to slipstream
  - Not consistent with gas sampling

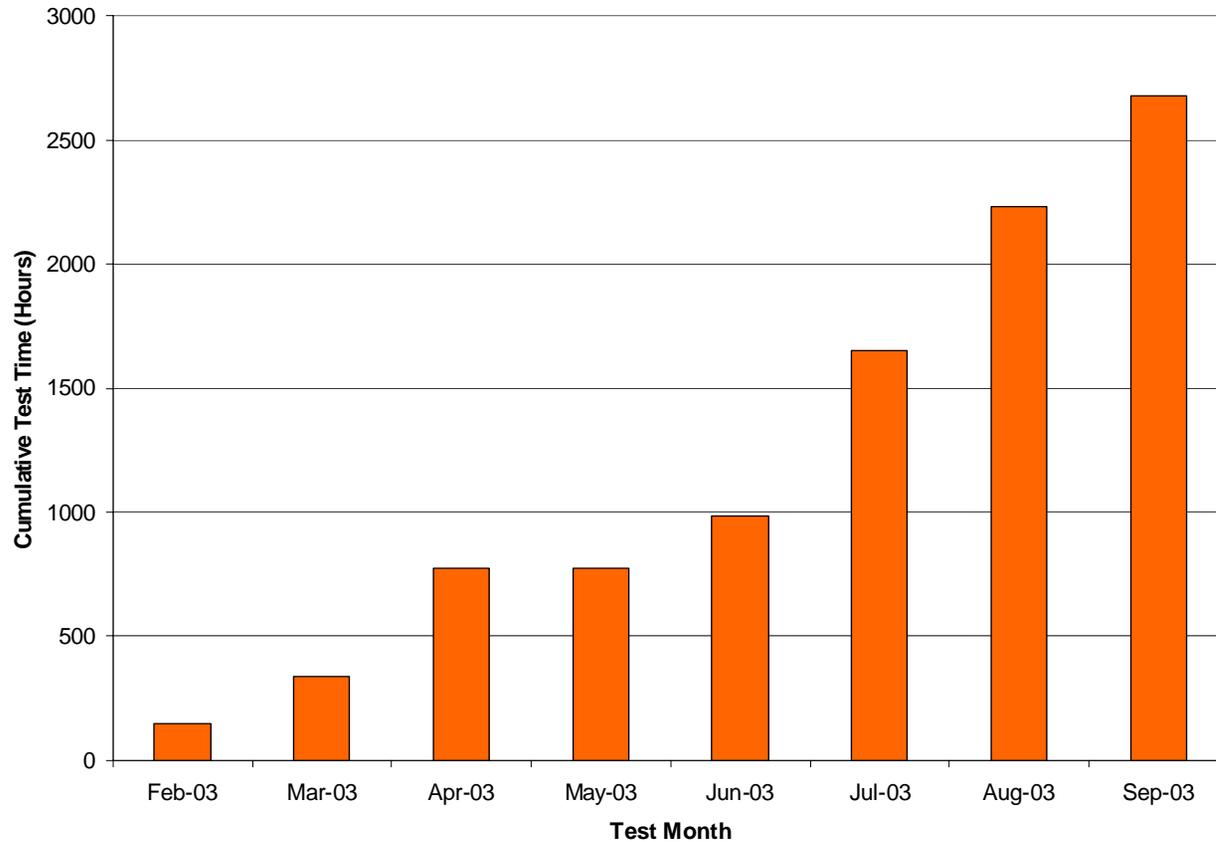
# Hg and Cl in ash

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8/15/03	0.13%	0.055	23

- Low LOI in ESP ash
  - Predominantly PRB
- ESP ash has very little Hg, ~0.5% of coal Hg (consistent with OH data)
- Cl content of ESP ash low ~1.5% of coal

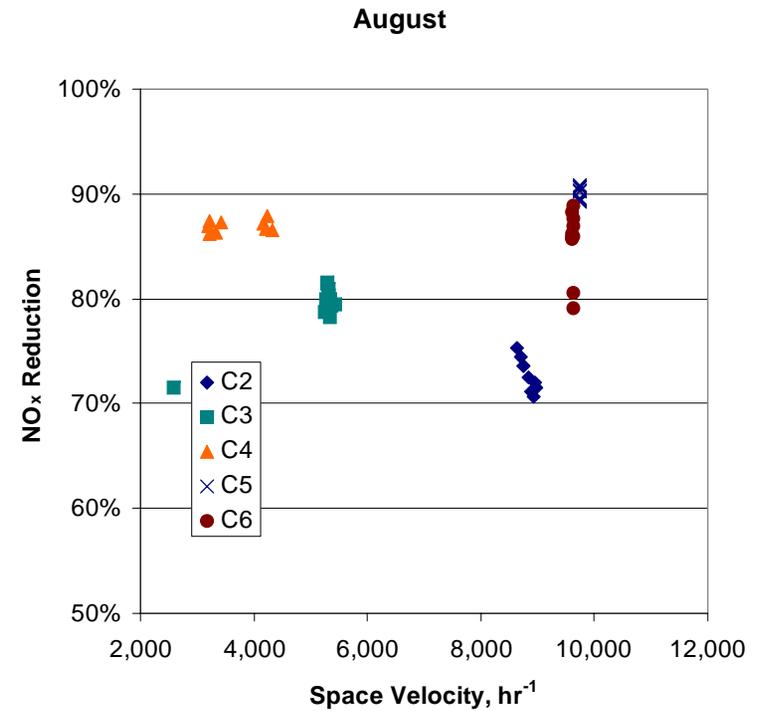
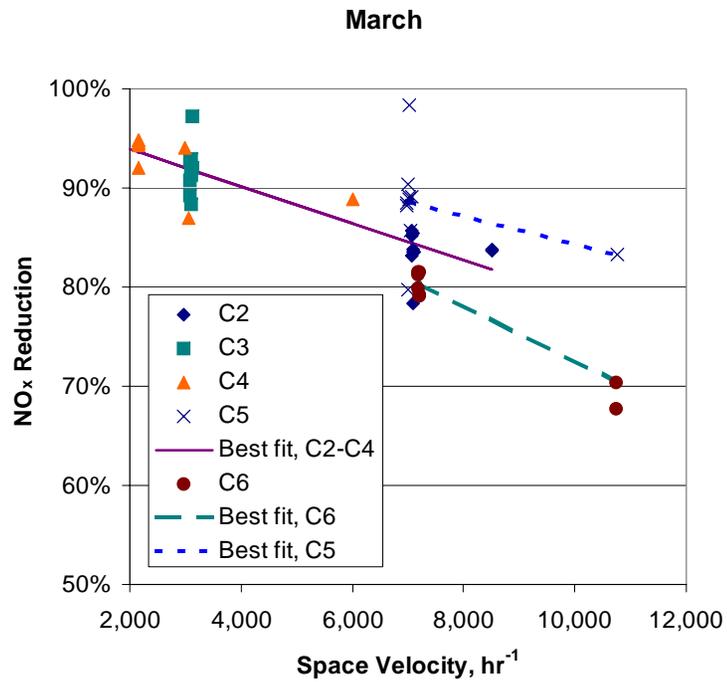


# Operating Experience

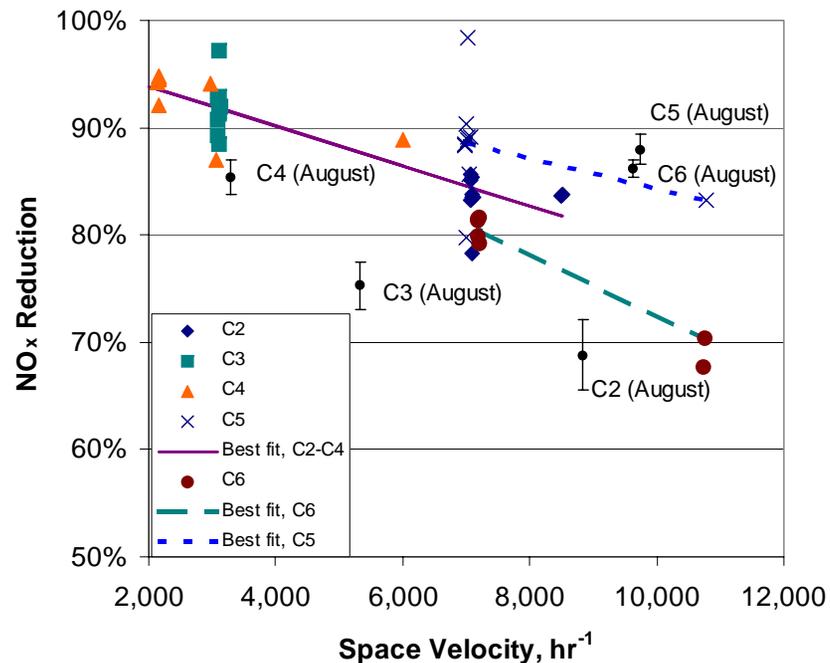


~2700 hours of cumulative flue gas exposure

# NO<sub>x</sub> Reduction



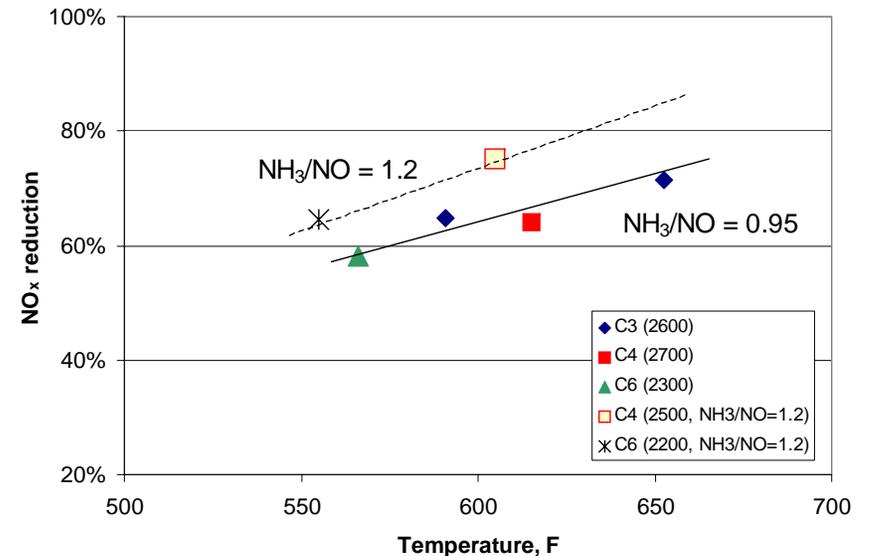
# Change in $\text{NO}_x$ Activity



- Activity decreased for catalysts C2, C3, C4
- Activity about the same for C5
- C6 appears higher?
  - Extrapolation of data to different temperature range

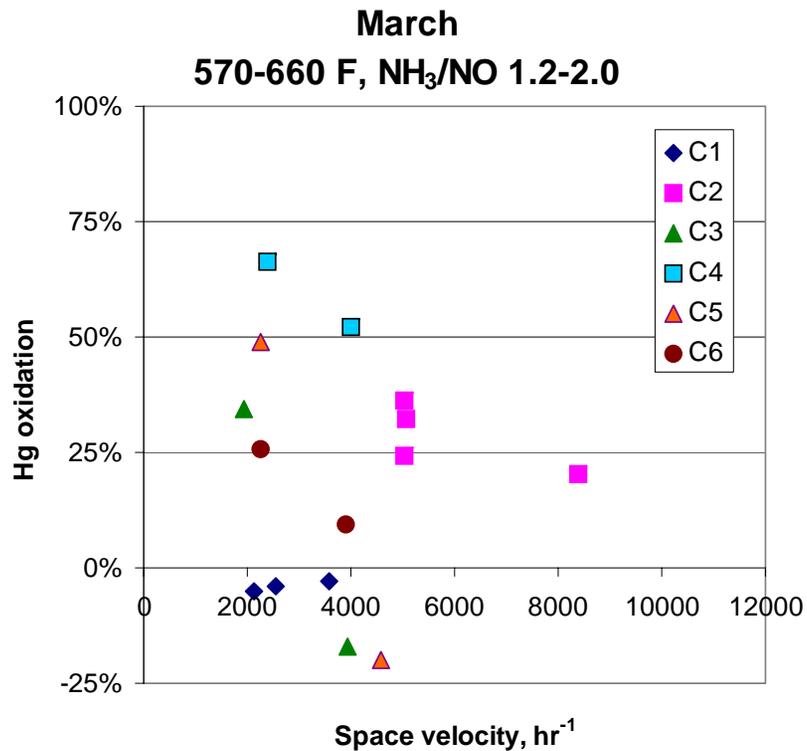
# Hg Testing: Temperatures

- Temperatures for August testing (500-550 F) lower for than March testing (570-660 F)
- $\text{NO}_x$  reduction decreases at lower T, as expected
- Hg effect?
  - Other lab, slipstream data<sup>1</sup> suggests Hg oxidation increases as temperature decreases



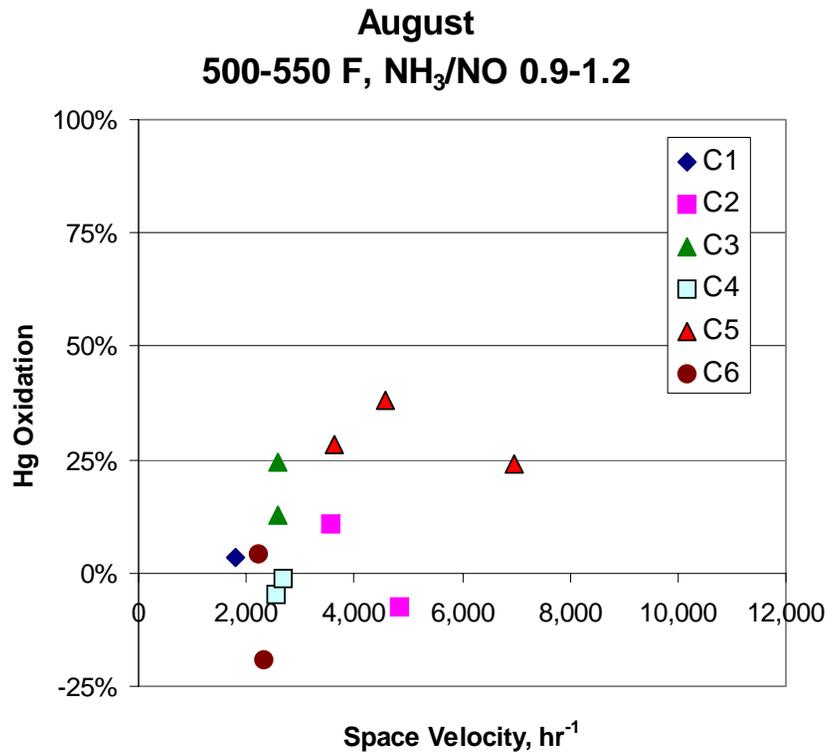
<sup>1</sup>Richardson, et al., AQIII Conference

# Oxidation of $\text{Hg}^0$



- Blank (C1) does not show oxidation
- March data in same range as previous pilot-scale data

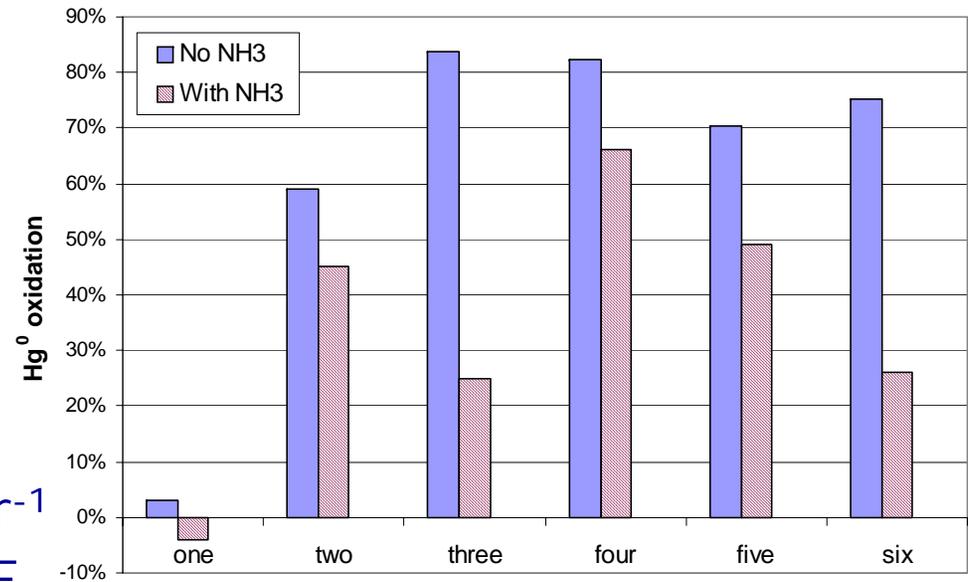
# Oxidation of $\text{Hg}^0$



- ▶ Blank (C1) does not show oxidation
- ▶ August data show some decrease in oxidation relative to March

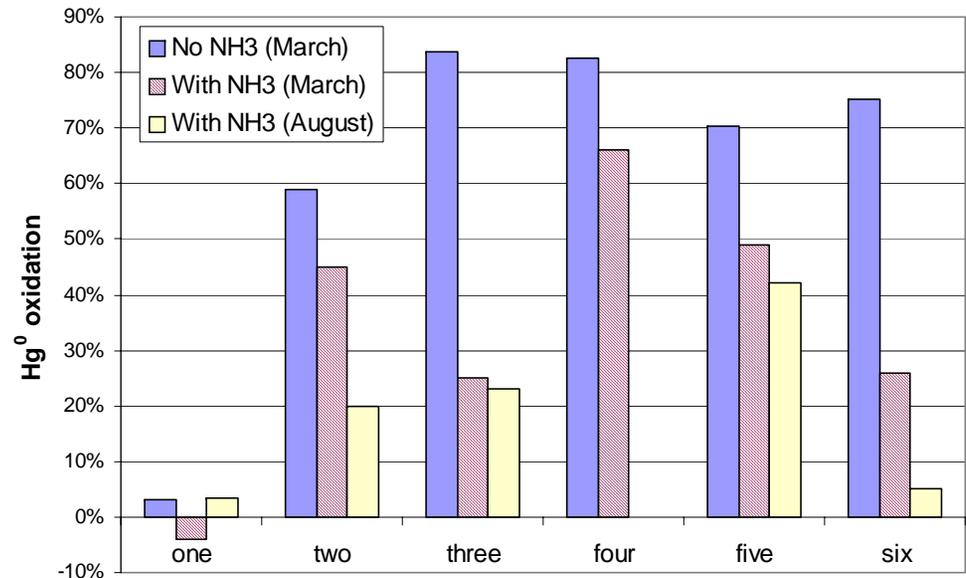
# Effect of Ammonia

- No ammonia vs. excess ammonia ( $\text{NH}_3/\text{NO} \sim 2$ )
- March:
  - SV  $\sim 2,500 \text{ hr}^{-1}$
  - T  $\sim 610\text{-}630 \text{ F}$
- Oxidation decreased in presence of ammonia
- No effect of blank monolith (C1)

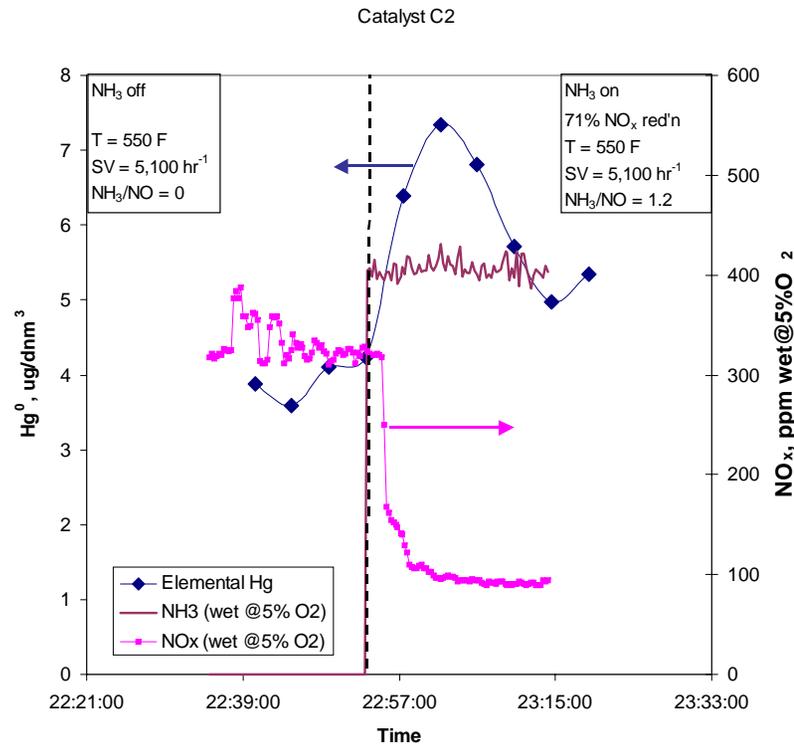


# Effect of Aging

- August:
  - SV ~ 2,500 hr<sup>-1</sup>
  - T ~ 500-550 F
  - NH<sub>3</sub>/NO ~ 0.9-1.2
- Two catalysts (C3 and C5 show little effect of aging on oxidation
- Other catalysts had decrease in oxidation
- No effect of blank monolith (C1)



# Transients



**Elemental mercury and NO<sub>x</sub> as a function of time for catalyst C2; T=550 F, SV=5,100 hr<sup>-1</sup>.**

- After ammonia turned off, concentration of elemental mercury initially increased, and then dropped
- Less oxidation with ammonia
- Time scale on the order of 10-20 minutes

# Conclusions

- Blank monolith showed no oxidation at 300 hours (March) or 2,200 hours (August)
- Oxidation of mercury increased without ammonia present
- Catalysts C3 (plate) and C5 (monolith) showed comparable oxidation in March and August
  - no loss of activity toward elemental mercury after ~2000 hours of exposure to flue gas.
- Other commercial catalysts exhibited less oxidation in August test series
- Hg oxidation not always correlated with NO<sub>x</sub> reduction
- Transient experiments: Hg desorption when ammonia turned on