

# Heavy-Duty NOx Emissions Control: Reformer-Assisted vs. Plasma- Facilitated Lean NOx Catalysis

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# Acknowledgements

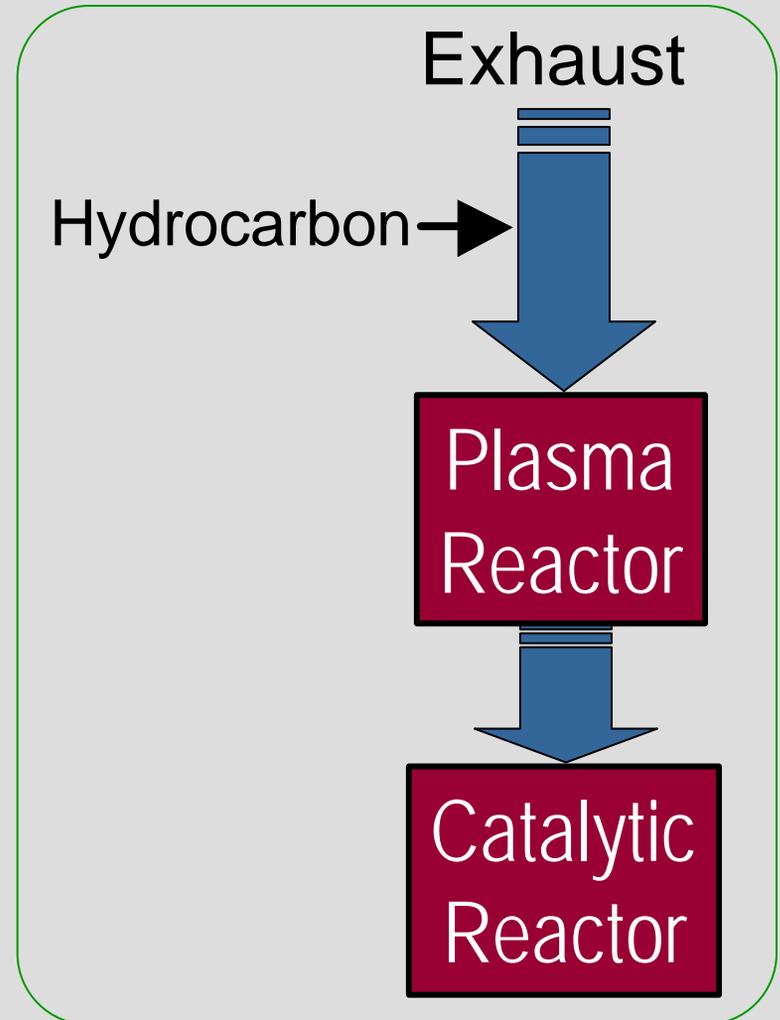
- ▶ K. Rappe, R. Rozmiarek, D. Mendoza – PNNL
- ▶ J. Hoard, C. Peden – LEP NTP CRADA
- ▶ G. Singh, K. Stork, DOE-OFCVT

# Outline

- ▶ Background
  - Flowsheets
  - Motivation for examination reformers
  - Suitable reformation approaches
  - Current work
- ▶ Data on reformer-assisted lean NO<sub>x</sub> catalysis
  - HC speciation using plasma reformer w/C<sub>3</sub>H<sub>6</sub>
  - Ag<sub>x</sub>O/γ-Al<sub>2</sub>O<sub>3</sub>
  - Ba/zeolite-Y ® Ag<sub>x</sub>O/γ-Al<sub>2</sub>O<sub>3</sub>
- ▶ Comparison between PFC and RAC
- ▶ Summary and future plans

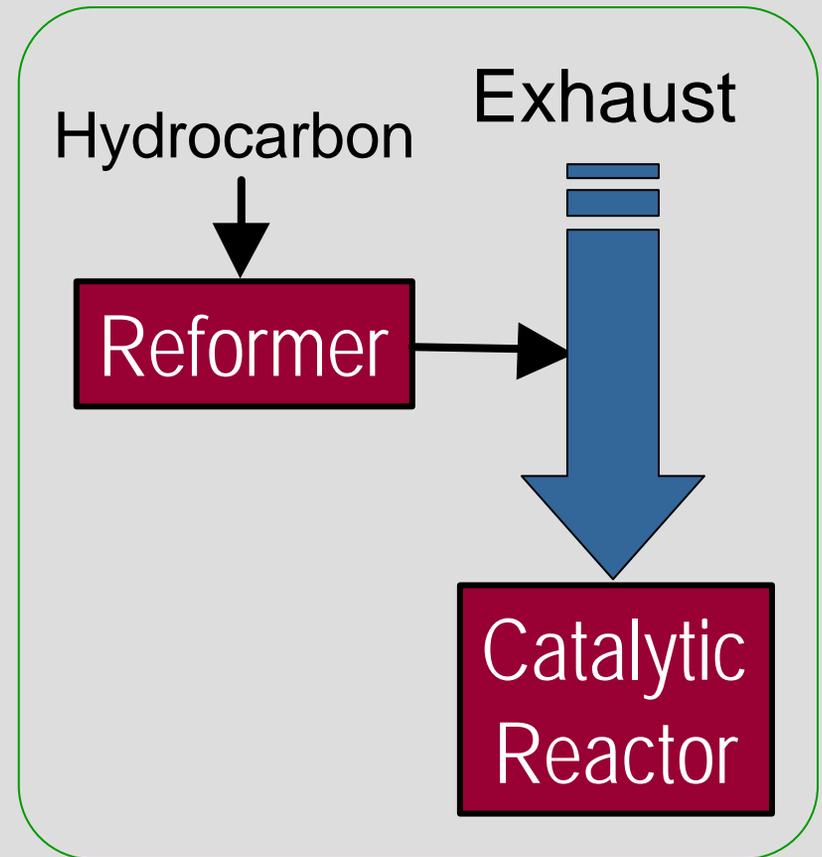
# Plasma-Facilitated Lean NO<sub>x</sub>

- ▶ Step 1. Exhaust passed through NTP
  - Selective oxidation of NO to NO<sub>2</sub>
  - Partial oxidation of hydrocarbon
- ▶ Step 2. NTP-treated exhaust passed over lean NO<sub>x</sub> catalyst
  - LD – zeolites; HD – alumina
  - NO & NO<sub>2</sub> reduced to N<sub>2</sub>
  - Consumption of hydrocarbons



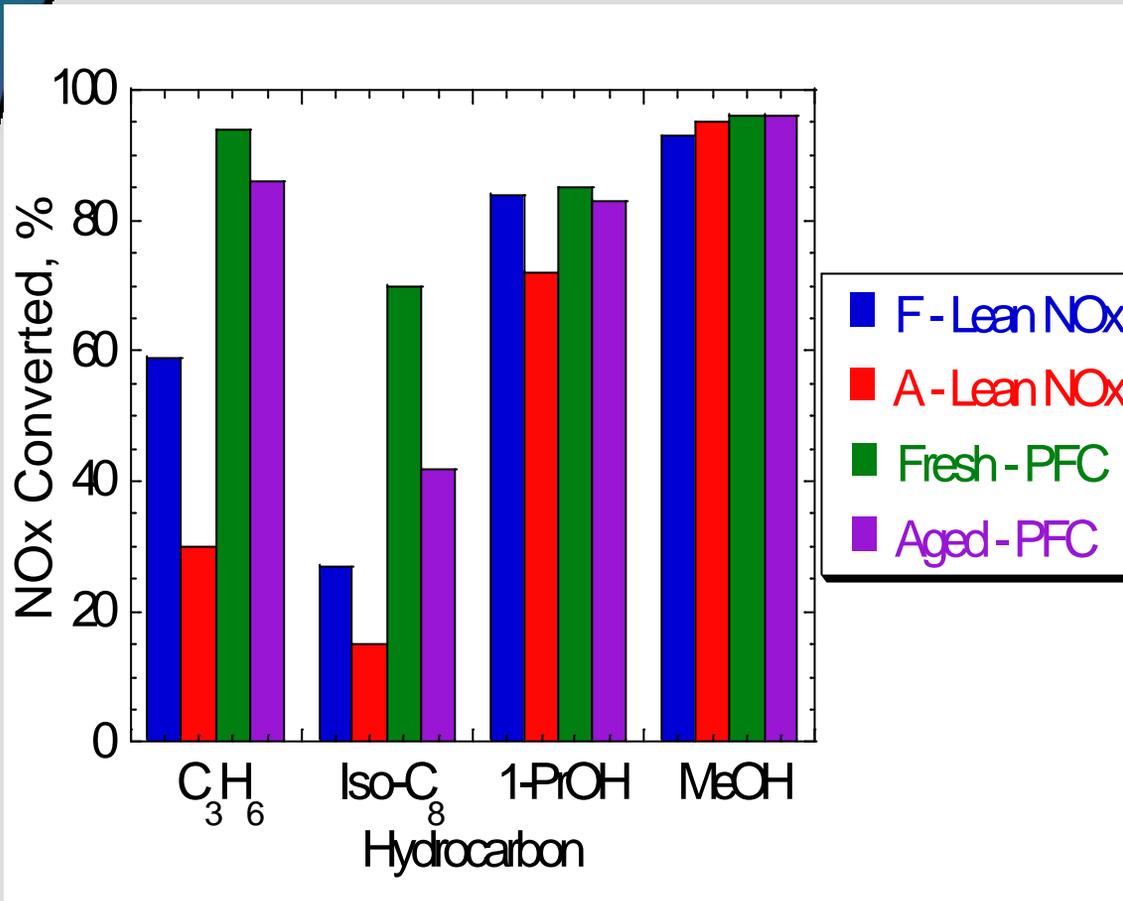
# Reformer-Assisted Lean NOx

- ▶ Step 1. Hydrocarbon is treated in a side stream
  - Partial oxidation of hydrocarbon
- ▶ Step 2. Treated HC injected in to the exhaust prior to lean NOx catalyst
  - NO reduced to N<sub>2</sub>
  - Consumption of hydrocarbons
  - Active catalyst should be similar to PFC



# Motivation for RAC

350°C:  $\text{In}_x\text{O}/\gamma\text{-Al}_2\text{O}_3$

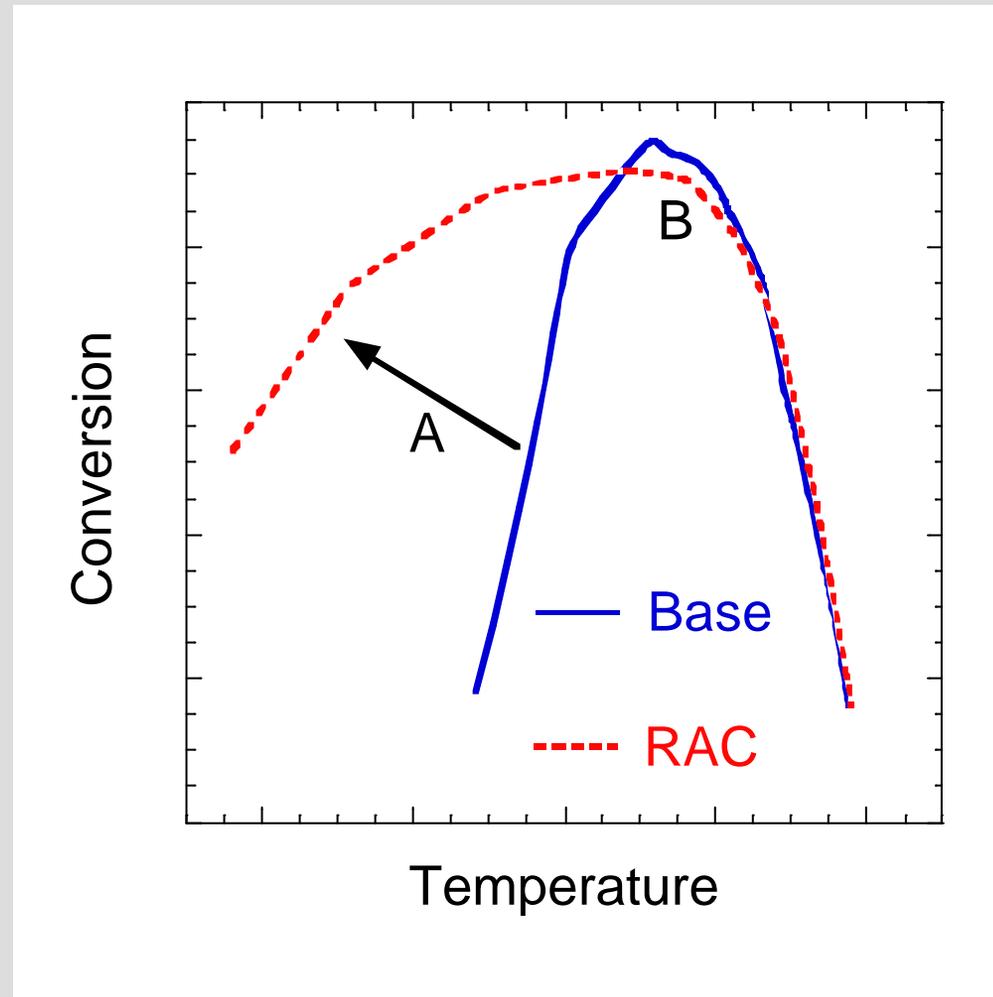


- ▶ Oxygenates result in higher activity on lean NOx catalysts
- ▶ Temperature range for high catalyst activity broadened
- ▶ Sulfur tolerance with oxygenates is better
- ▶ Potential to reduce energy cost (fuel penalty) and reduce the size of the system as a whole

Equal C<sub>1</sub>:NOx basis. Aging in 30 ppm SO<sub>2</sub>, 7% H<sub>2</sub>O, 500 ppm NO, 7% O<sub>2</sub> for 250 Hrs.

# What to Expect From RAC?

- ▶ A) Major benefit is broadening the active temperature window due to higher reactivity of oxygenates over catalyst at lower temperatures
- ▶ B) May be a slight loss in maximum level of reduction due to lower  $C_1:NO_x$  over catalyst



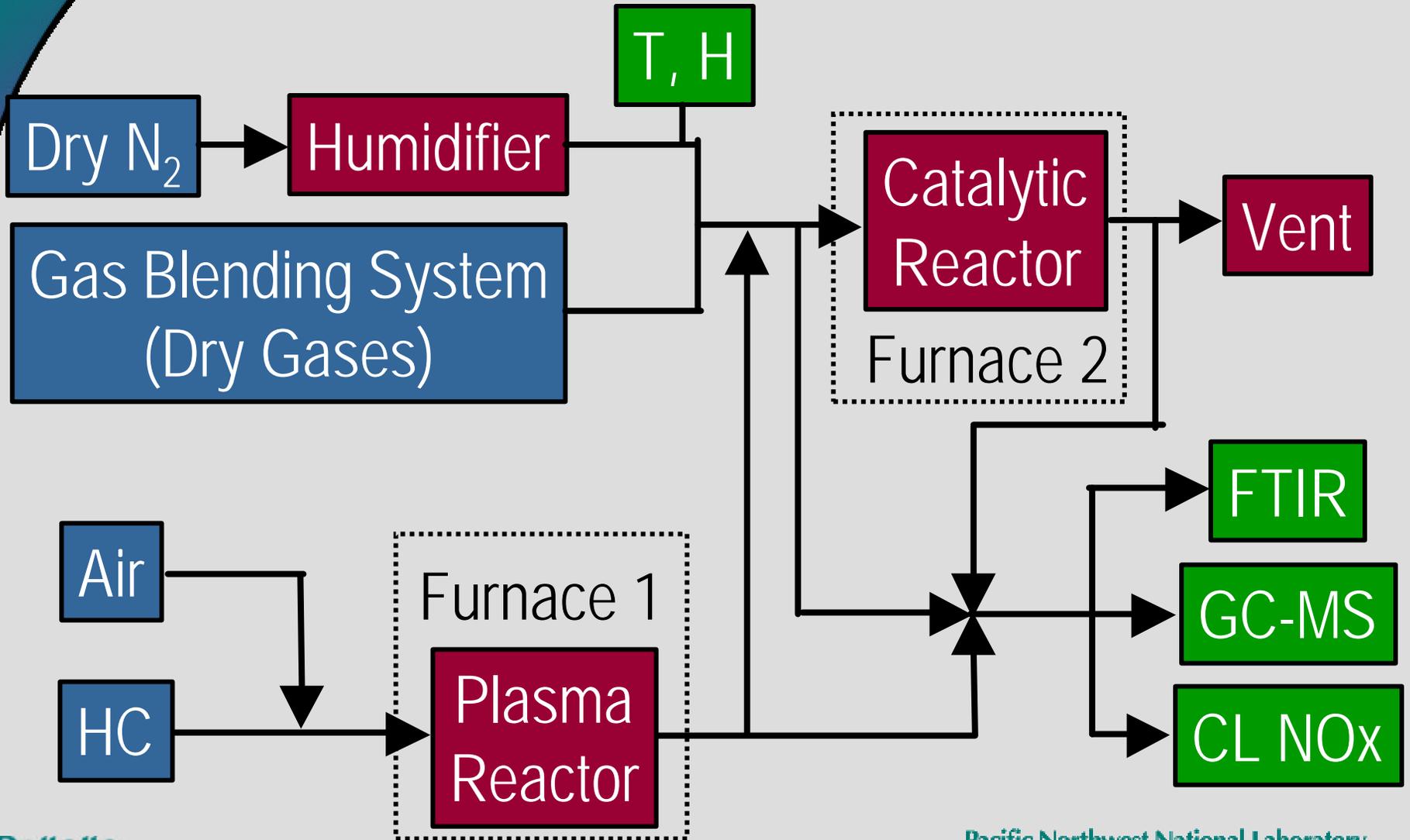
# Reformer Options

- ▶ NTP alone
- ▶ NTP-catalyst combinations
- ▶ Thermal catalytic reformers
  - Partial oxidation of alkanes
  - Reform fuel to  $H_2/CO$  and use synthesis chemistry to obtain desired reducing agent

# Ongoing Work of Interest

- ▶ First record: Engelhard USP #6176078
- ▶ PNNL-Caterpillar CRADA (Aardahl): Plasma-based oxygenate production
- ▶ GM (Schmieg): Thermal catalytic oxygenate production (LEP CRADA with PNNL)
- ▶ LANL (Borup) POx for olefin production from fuel
- ▶ Catalytica, NOxTech: Fuel reformers/converters for aftertreatment
- ▶ Potential sources of syngas
  - DOE-EERE Hydrogen Program: POx-LANL; Autothermal-ANL; SMR-PNNL
  - MIT (Bromberg): Plasmatrons
  - Seimens (Hammer), several others: Plasma SMR
  - Delphi (Fisher), Univ. Minn. (Schmidt): Millisecond POx

# Bench Scale RAC System



# Stream Compositions

## ▶ Catalytic Reactor

- Simulated exhaust composed of 2% water, 8% CO<sub>2</sub>, 9% O<sub>2</sub>, 500 ppm NO, and 300 ppm CO.
- Steady-state data taken: constant reactor temperature and reformer conditions for a given point.
- 3 grams of catalyst and 3 SLM flow.
  - Ag<sub>x</sub>O/γ-alumina
  - Ba/zeolite Y ® Ag<sub>x</sub>O/γ-alumina

## ▶ Reformer

- Feed streams are house air and propene.
- Propene rate fed to the reformer maintains constant propene to NO<sub>x</sub> ratio of 4 in the catalytic reactor. Values reported are feed ratios only, not actual HC:NO<sub>x</sub> levels in the catalytic reactor.
- 1:1 to 15:1 C-to-O ratios examined
- Reformer held at 500°C and 6 kJ/L

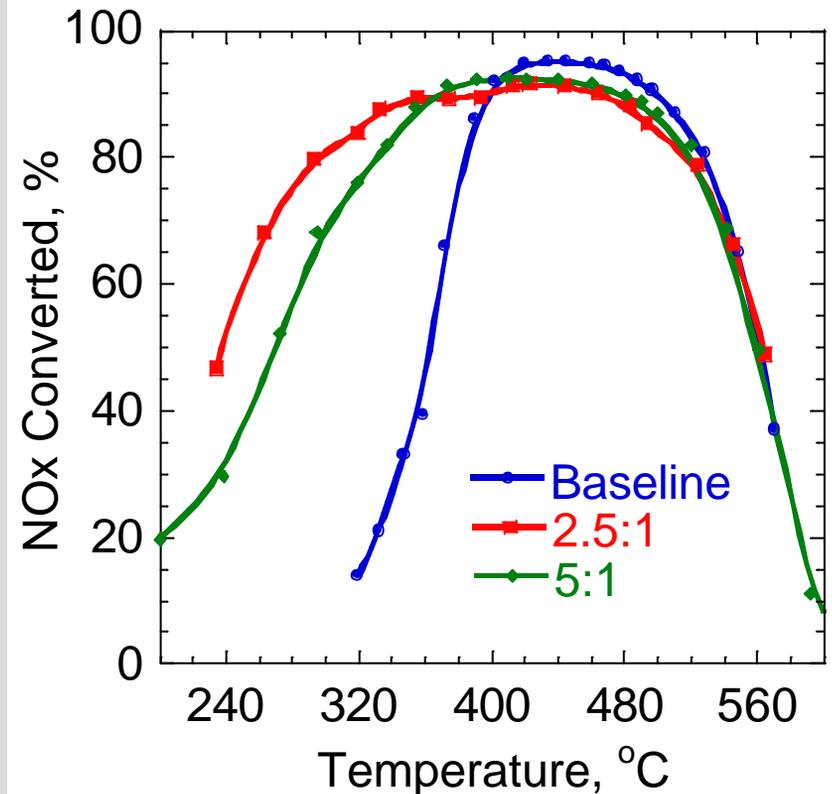
# Reformer Chemistry for C<sub>3</sub>H<sub>6</sub>

- ▶ Higher energy density produces more oxygenates
- ▶ Observed compounds include **acetaldehyde**, propylene oxide, **propanal**, acetone, **2-propenal**, **2-propanol**, **formaldehyde**, **methanol**, **ethanol**, **1-propanol**, **2-propen-1-ol**, 1-hydroxy-2-propanone, carboxylic acids
- ▶ Higher power shifts alcohol production to somewhat higher C:O ratios
- ▶ Single pass propene conversion is less than 20% for conditions used.
- ▶ Limited CO<sub>2</sub> production at high C:O ratio. Selectivity deteriorates heavily below C:O = 2. Complete C balance still needed.

# RAC Results: $\text{Ag}_x\text{O}/\text{g-Al}_2\text{O}_3$

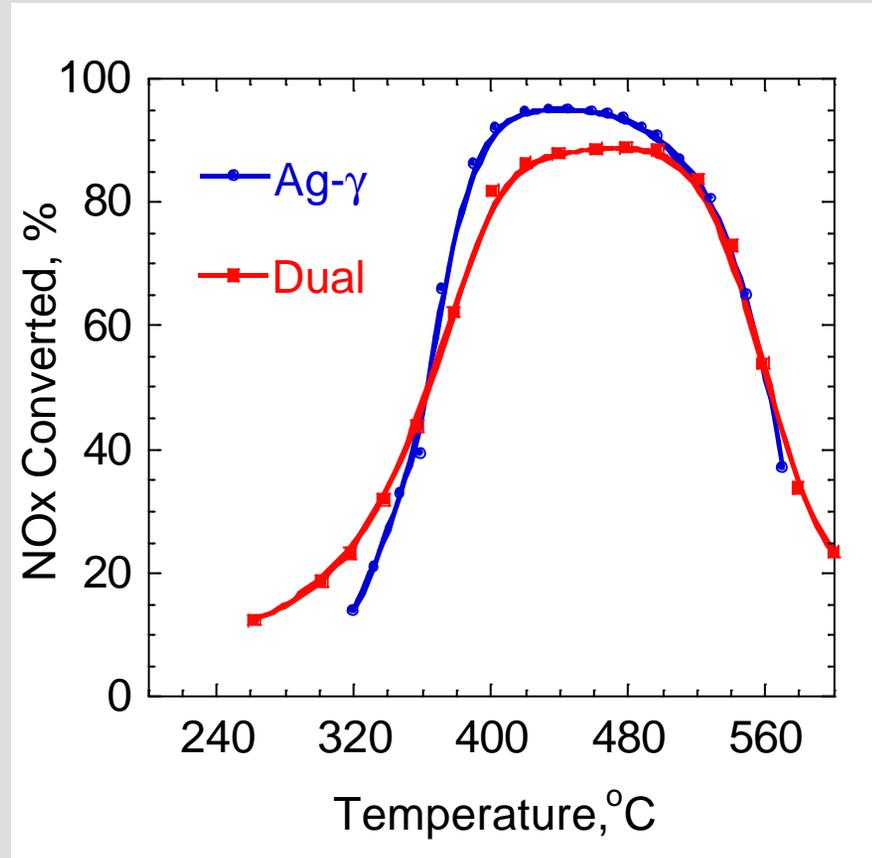
Reformer Operation: 500°C, 6 kJ/L

- ▶ Baseline is equivalent to lean NO<sub>x</sub> catalysis – no reformation
- ▶ As expected, temperature window broadened with slight loss in peak NO<sub>x</sub> reduction efficiency
- ▶ SV = 22000 1/hr



# Lean NOx: Dual System vs Single Catalyst

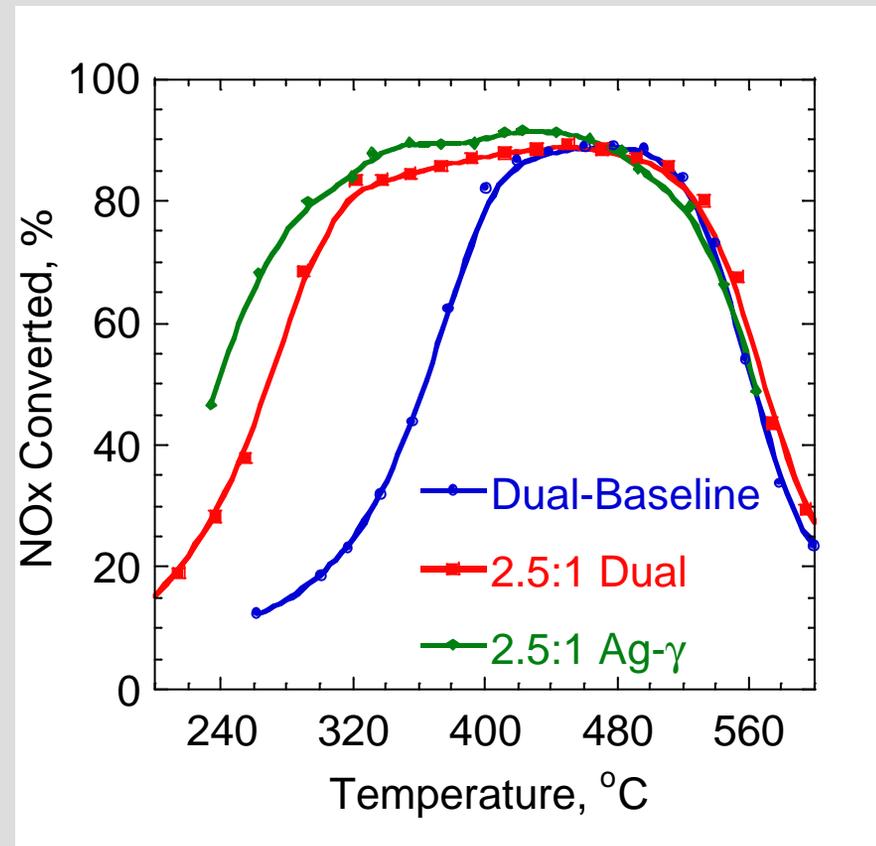
- ▶  $\text{Ag-}\gamma = \text{Ag}_x\text{O}/\gamma\text{-Al}_2\text{O}_3$
- ▶ Dual catalyst represents Ba/zeolite Y followed by Ag- $\gamma$  (SV = 28000 1/hr; SV= 44000 based on Ag- $\gamma$  alone)
- ▶ Total amount of catalyst by weight is the same for each case
- ▶ Drop in maximum efficiency due to SV effect. Ba/zeolite Y is not active with propene feed



# RAC for Dual Catalyst System

- ▶ Reformulation shows improvement from lean NO<sub>x</sub> baseline.
- ▶ Dual catalyst not better than Ag- $\gamma$  alone.
- ▶ Likely that Ba/zeolite Y is not active with reformat as reducing agent.
- ▶ Drop in NO<sub>x</sub> conversion in dual catalyst from Ag- $\gamma$  due to SV effect.

Reformer Operation: 500°C, 6 kJ/L

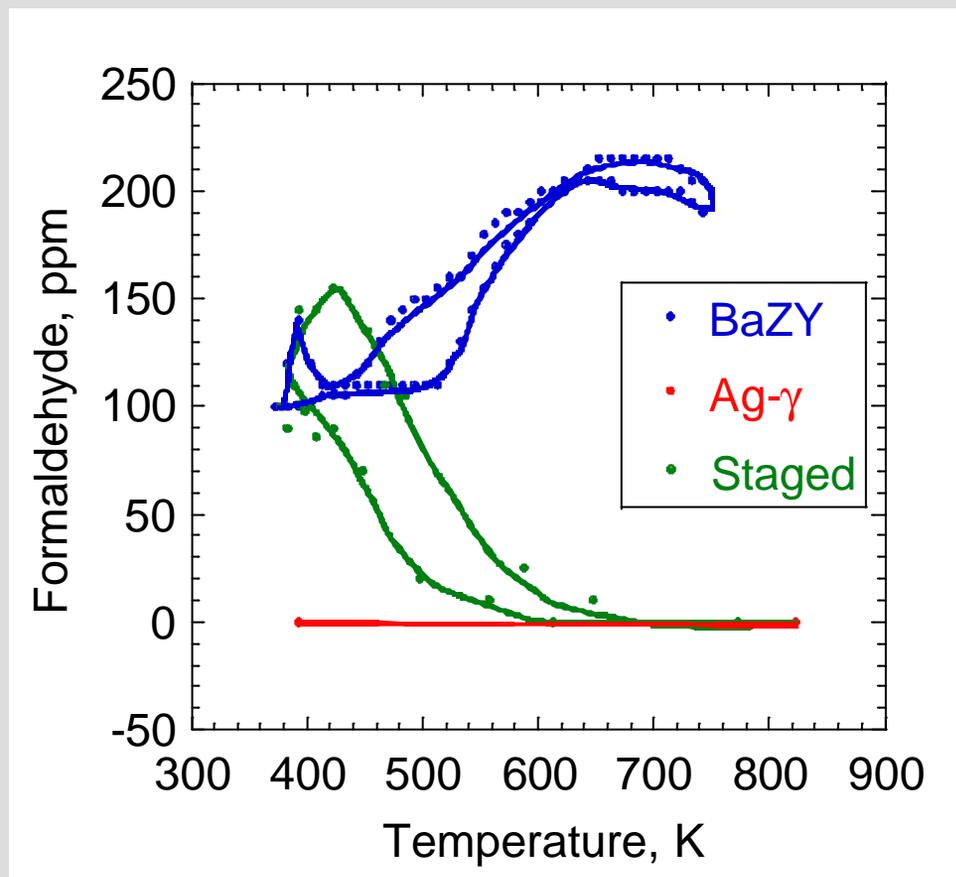


# Impact of Oxygenate Speciation

RAC shows positive results for  $\text{Ag}_x\text{O}/\gamma\text{-Al}_2\text{O}_3$ , but not for the staged catalyst system with Ba/zeolite Y.

**Hypothesis:** Oxygenates produced in the plasma can participate in the deNOx chemistry over alumina, but not the zeolite.

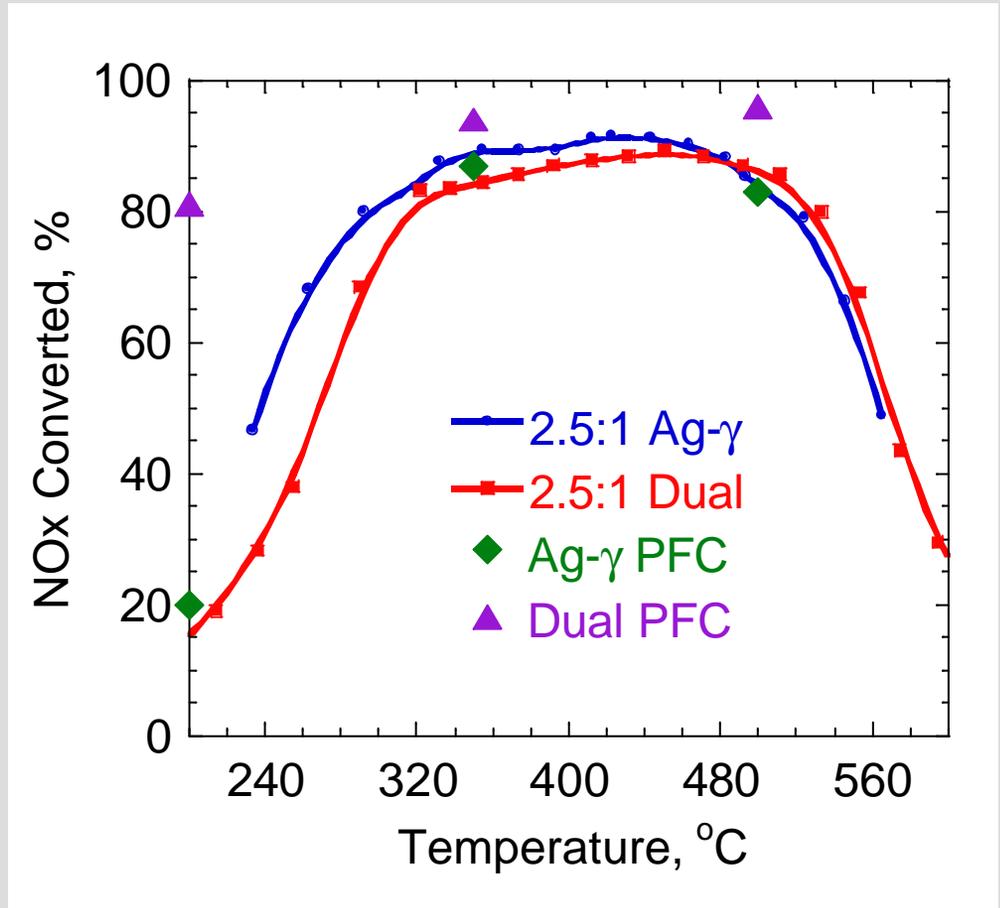
Observed before with formaldehyde.



Data taken by J. Hoard (Ford Research Labs)

# Comparison: PFC vs RAC

- ▶ In the case of Ag- $\gamma$ , RAC is equivalent to PFC (40 J/L).
- ▶ Still advantages for the dual catalyst system in using PFC.



# Summary

- ▶ Reformer-assisted lean NO<sub>x</sub> catalysis (RAC) is equivalent or better than plasma-facilitated catalysis (PFC) over Ag<sub>x</sub>O/γ-Al<sub>2</sub>O<sub>3</sub>.
- ▶ The configuration of RAC used here (air:HC) was not successful in promoting NO<sub>x</sub> reduction over Ba/zeolite Y. Other species in exhaust may be key to make ample amounts of acetaldehyde.

# Outlook

- ▶ Currently looking at larger alkanes (dodecane and hexadecane) in the bench system
  - Fuel vaporizer required
  - Line heating an issue
  - Coking an issue?
- ▶ Catalytic NTP reforming could benefit
  - System is simple
  - Chemistry is not so simple
- ▶ Provided continued improvement examined, RAC will be examined at full scale during transient PFC testing at Caterpillar Tech. Center in Spring-Summer 2004.