

# **Collective Protection 2005**

## **Design of Catalytic Process for the Removal of CW Agents and TIC's**

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# **Introduction**

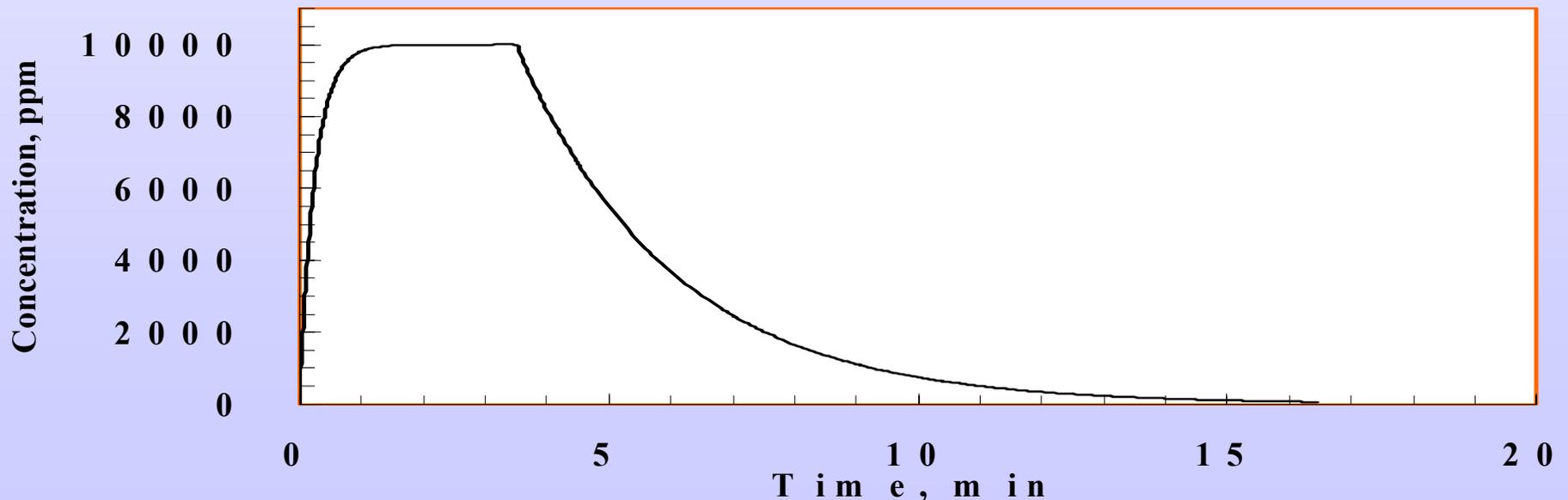
## **Catalytic Oxidation Technology is Well Suited for Military Air Purification Applications**

- 1. Technology Non-selective: Agents and TIC's reduced to CO<sub>2</sub>, H<sub>2</sub>O and haloacids**
- 2. Elevated Temperatures may Offer Biological Decontamination**
- 3. Unlimited Capacity for Agents and TIC's: Catalytic sites not consumed during reaction**
- 4. Technology lends itself well towards integration with many host applications**

# Introduction

## Catalytic Process versus Chemical Threat

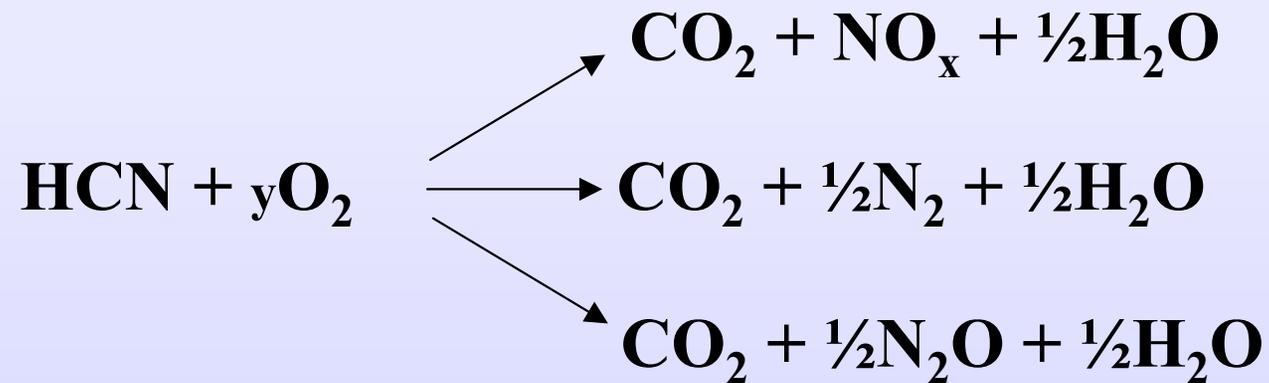
1. **Chemical threat; how catalytic process will be exposed to chemical agent, differs greatly with how gas-mask and CP filters are tested.**
2. **During a chemical attack, a filtration device can be exposed to a rapidly increasing, high concentration of chemical agent that will decay over time.**



# Introduction: Catalytic Response

- 1. Catalytic reaction rates can be highly non-linear in concentration. Catalyst must be operated at conditions sufficient to achieve high destruction efficiencies when challenged with high concentration of agent.**
- 2. When exposed to high concentration of chemical agent, the temperature within the catalyst will increase significantly in a short period of time.**
  - o Conversion will increase, as catalyst performance increases exponentially with increasing temperature.**
  - o For nitrogen-containing compounds (e.g HCN, NH<sub>3</sub>), product selectivity will shift from N<sub>2</sub> to NO<sub>x</sub> with increasing temperature.**
- 3. NO<sub>x</sub> (NO and NO<sub>2</sub>) is difficult to filter within the constraints of regeneration system. Therefore, catalytic process must be designed to minimize NO<sub>x</sub> formation during the destruction of nitrogen-containing compounds.**

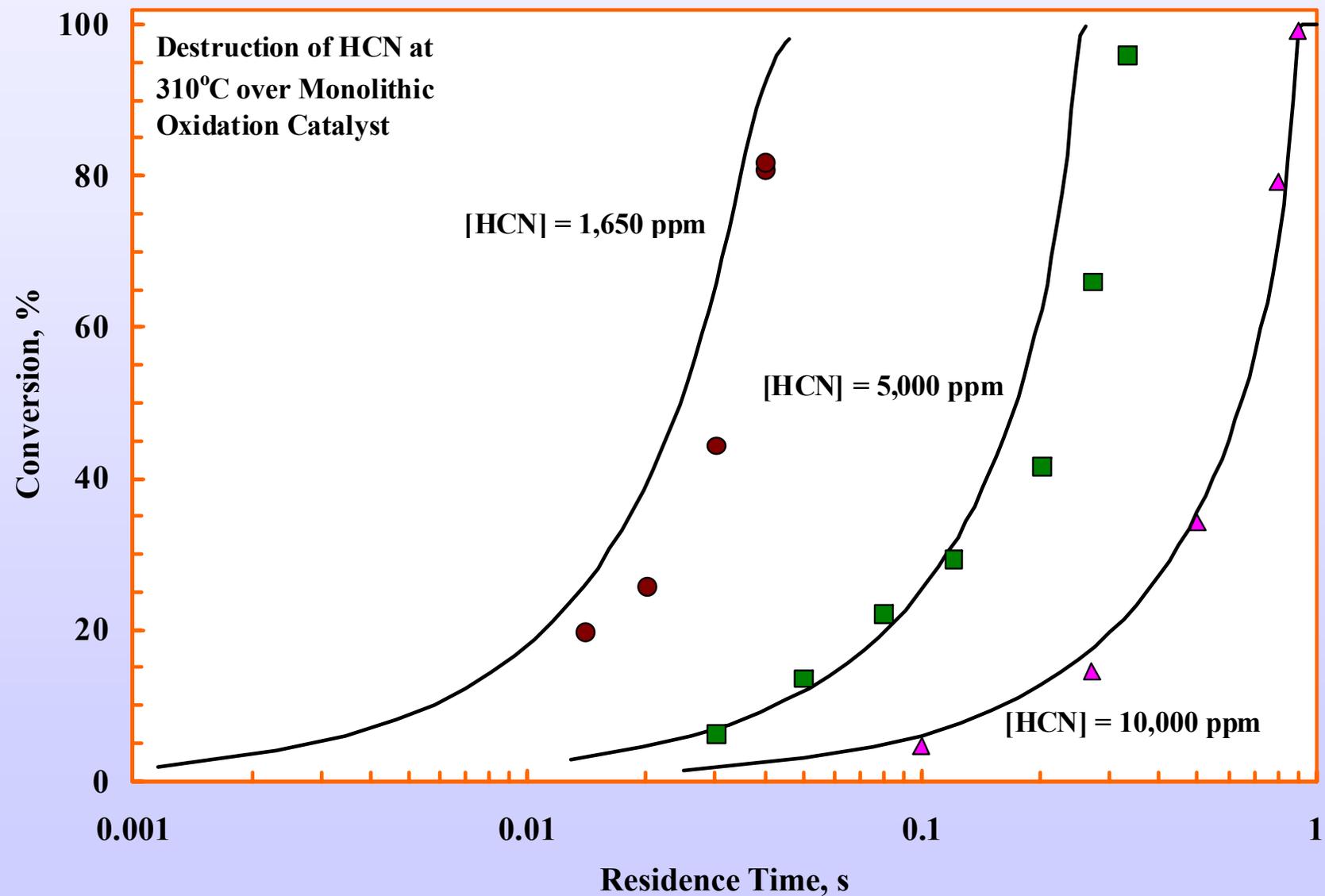
## Catalytic Destruction of Nitrogen-Containing Compounds



### Requirements:

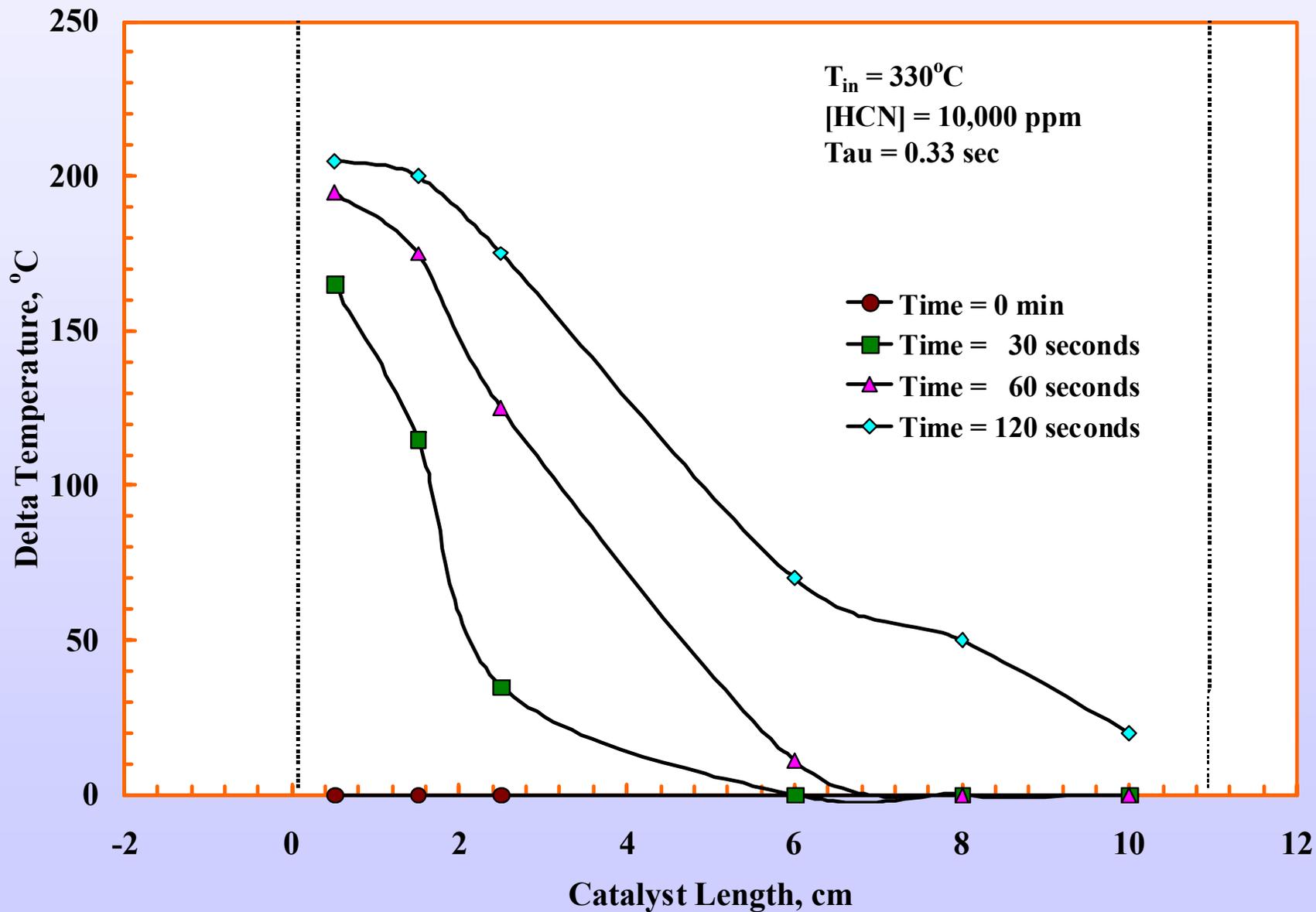
- NO:** Less than 30 mg/m<sup>3</sup>
- NO<sub>2</sub>:** Less than 5 mg/m<sup>3</sup>
- N<sub>2</sub>O:** Less than 60 mg/m<sup>3</sup>

# Effects of Concentration of Reaction Rate



Rossin, 1995

# Catalyst Thermal Response during Pulse of HCN



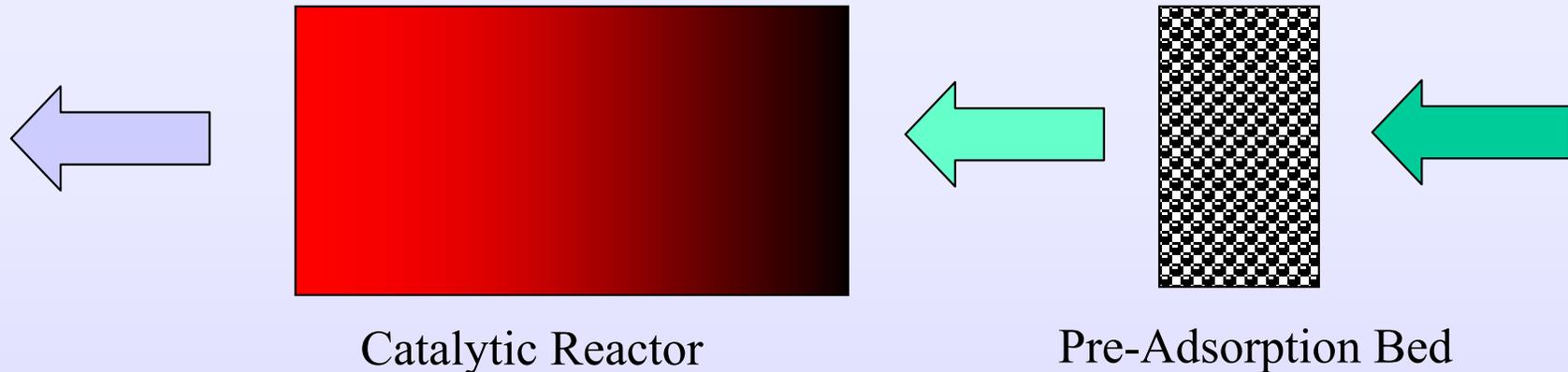
Rossin, 1995

# Objective

Evaluate the use of an adsorption bed up-stream of the catalytic reactor to minimize  $\text{NO}_x$  formation

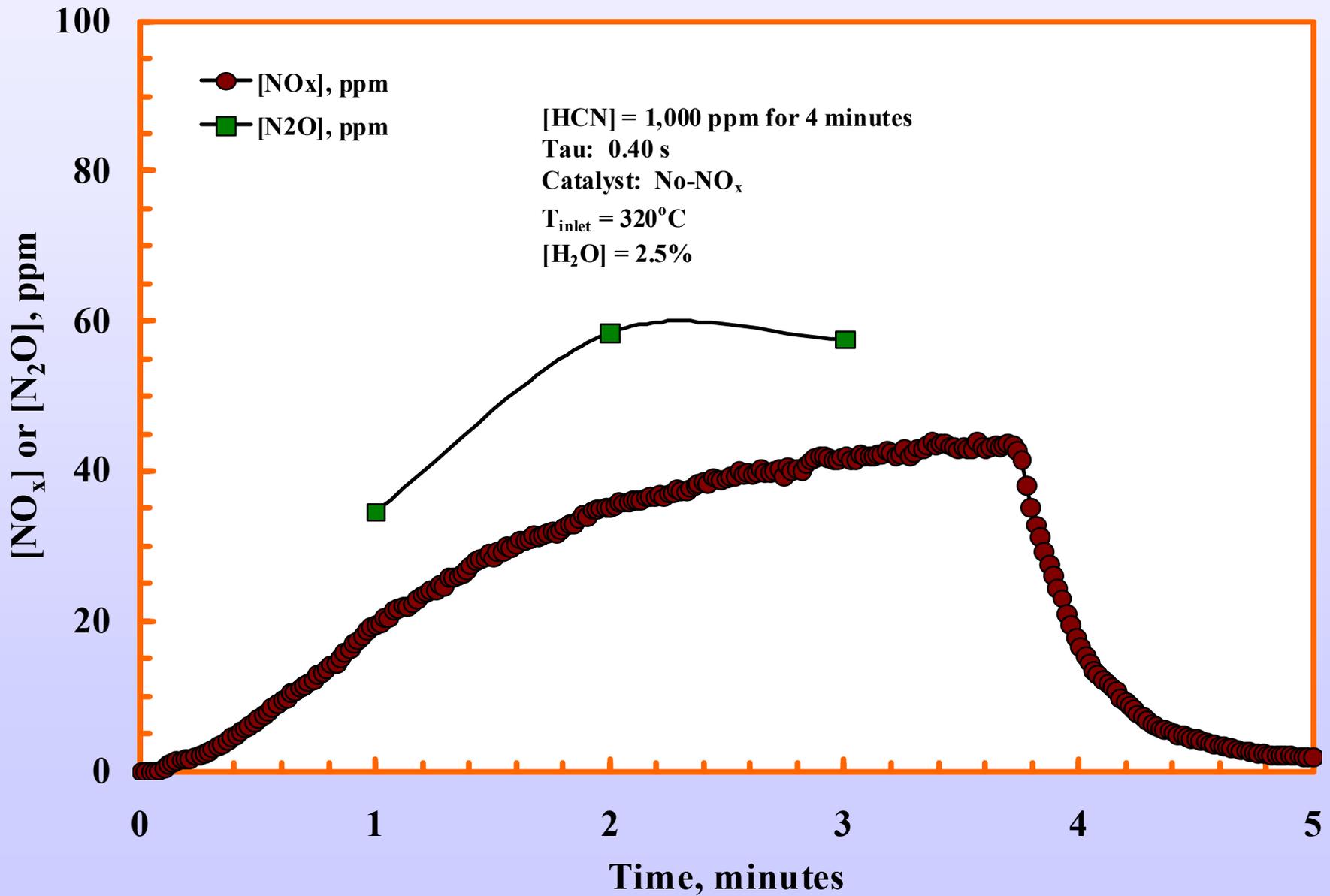
- o Adsorption bed expected to dampen high concentration spikes in chemical
- o Chemical will be delivered to the catalyst at lower concentrations over longer time.
  - Favorable reaction kinetics
  - Greater thermal management
- o  $\text{NO}_x$  formation will be reduced via lower catalyst temperature and lower chemical concentration.

# Catalytic Oxidation Process

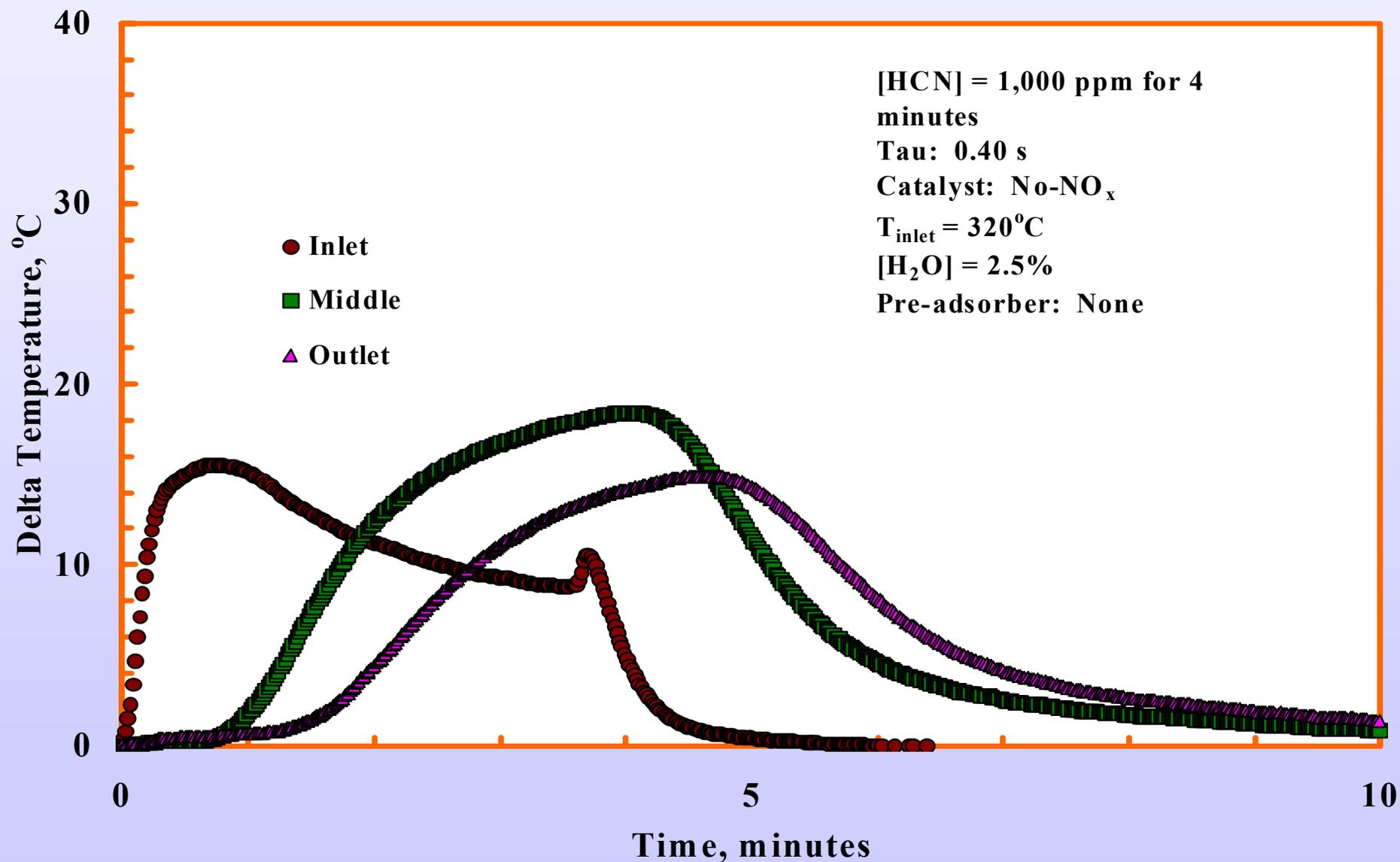


1. Expose catalytic process to pulse of chemical
2. Record catalyst temperature in real time
3. Monitor  $\text{NO}_x$  in real time using  $\text{NO-NO}_x$  analyzer
4. Monitor concentration of  $\text{N}_2\text{O}$ ,  $\text{CO}_2$  and chemical in effluent stream using grab bags and discrete on-line analysis.

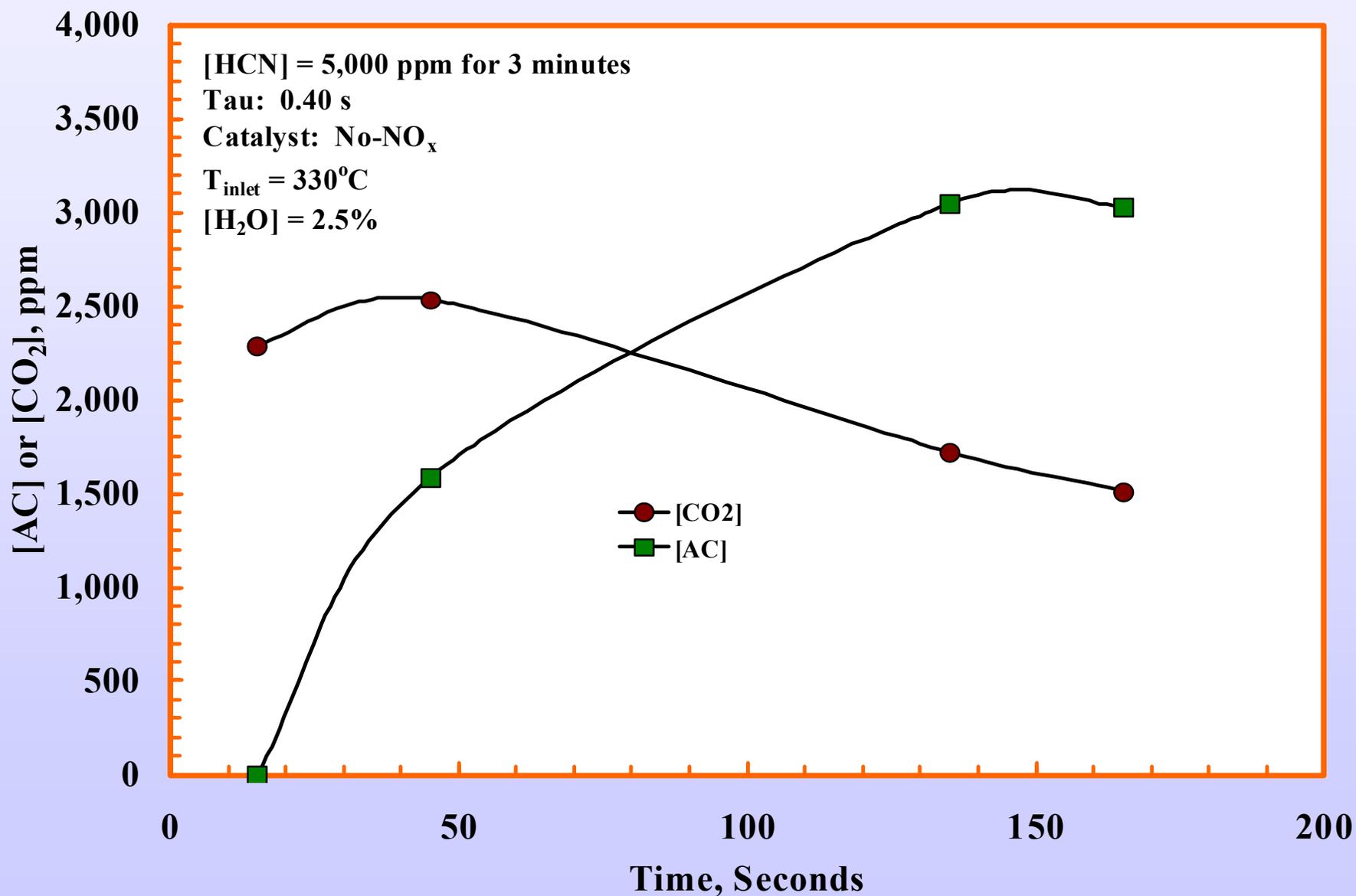
# No Pre-Adsorber, Low Concentration



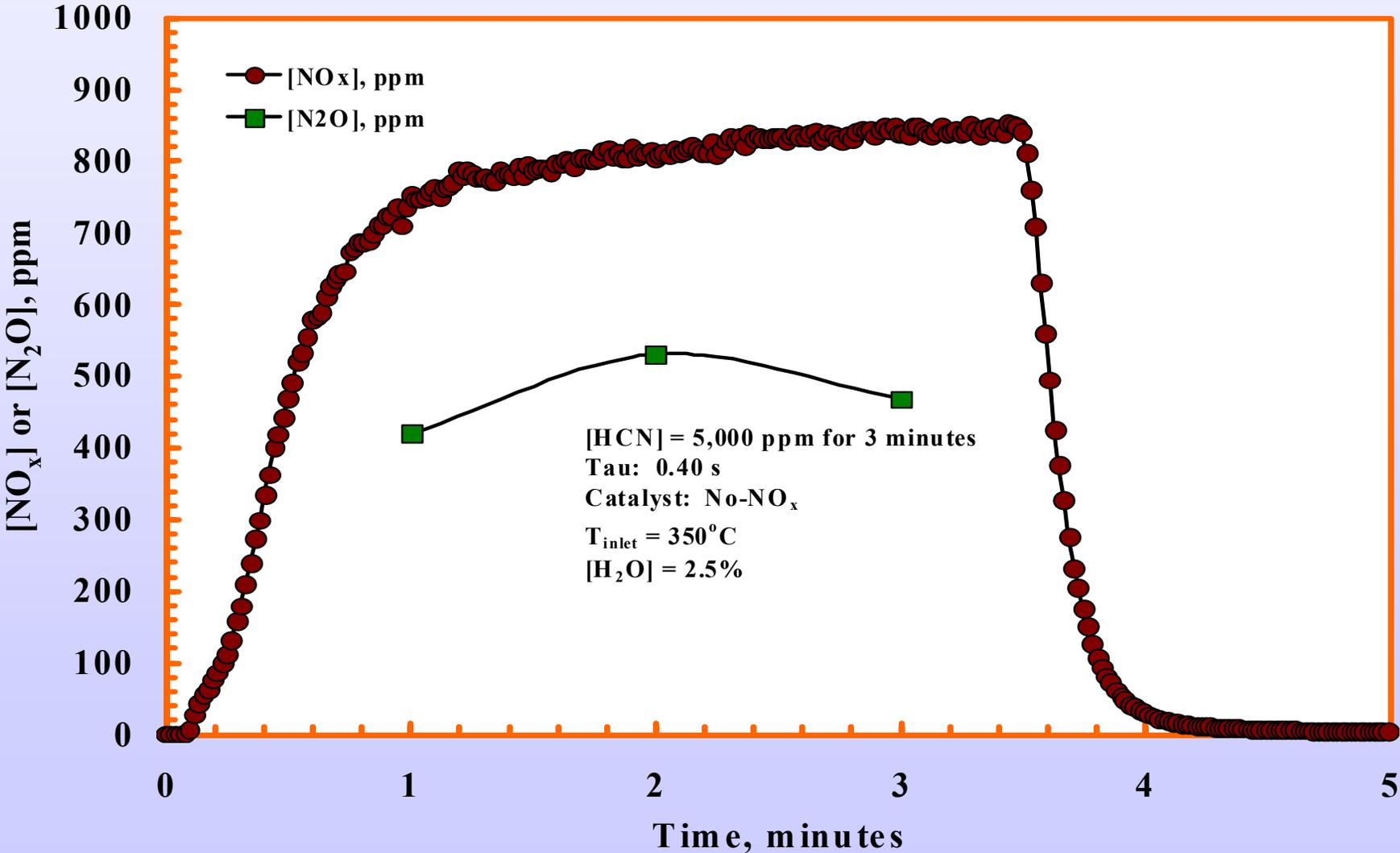
## No Pre-Adsorber, Low Concentration



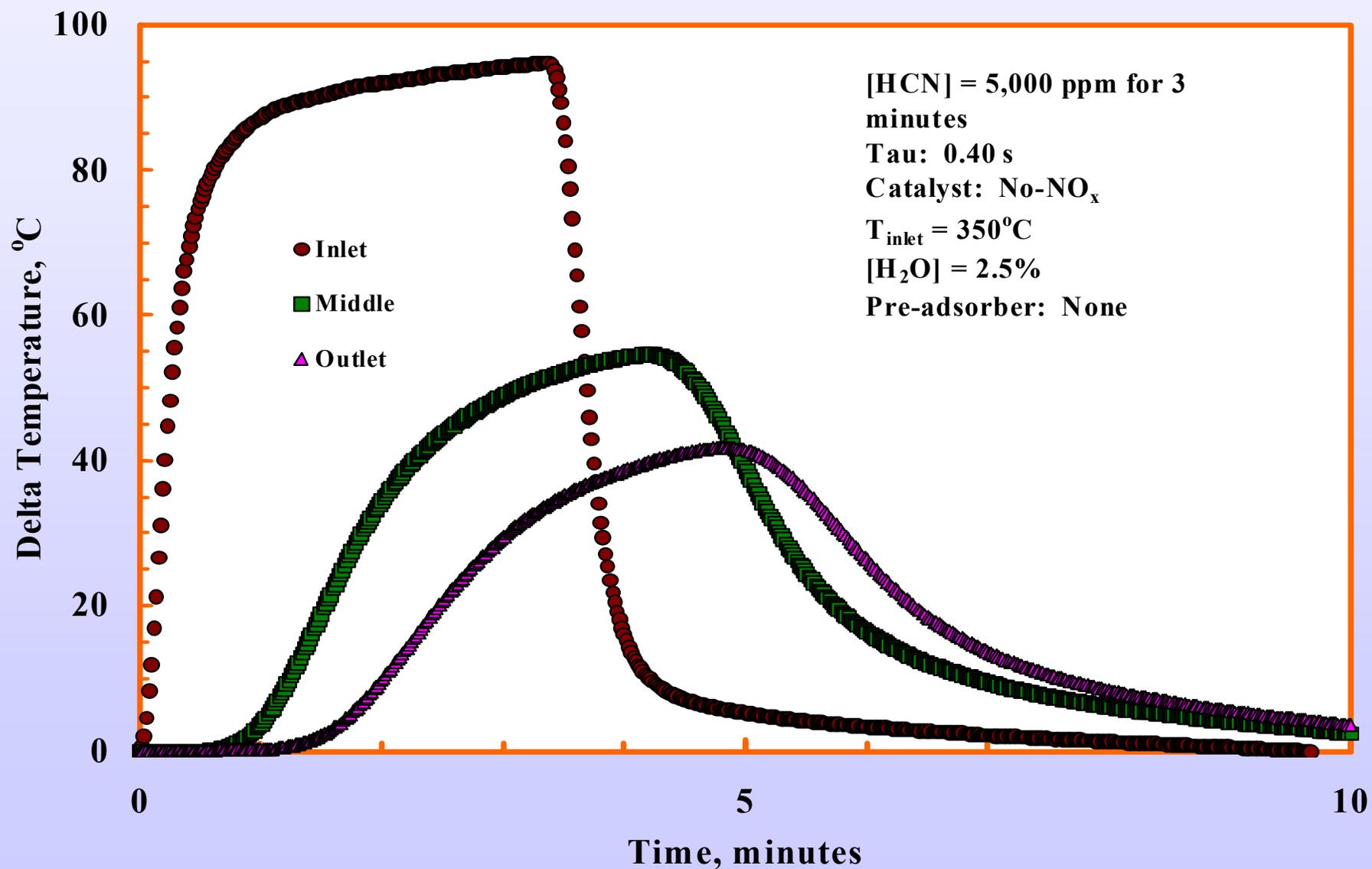
## No Pre-Adsorber, Insufficient Temperature



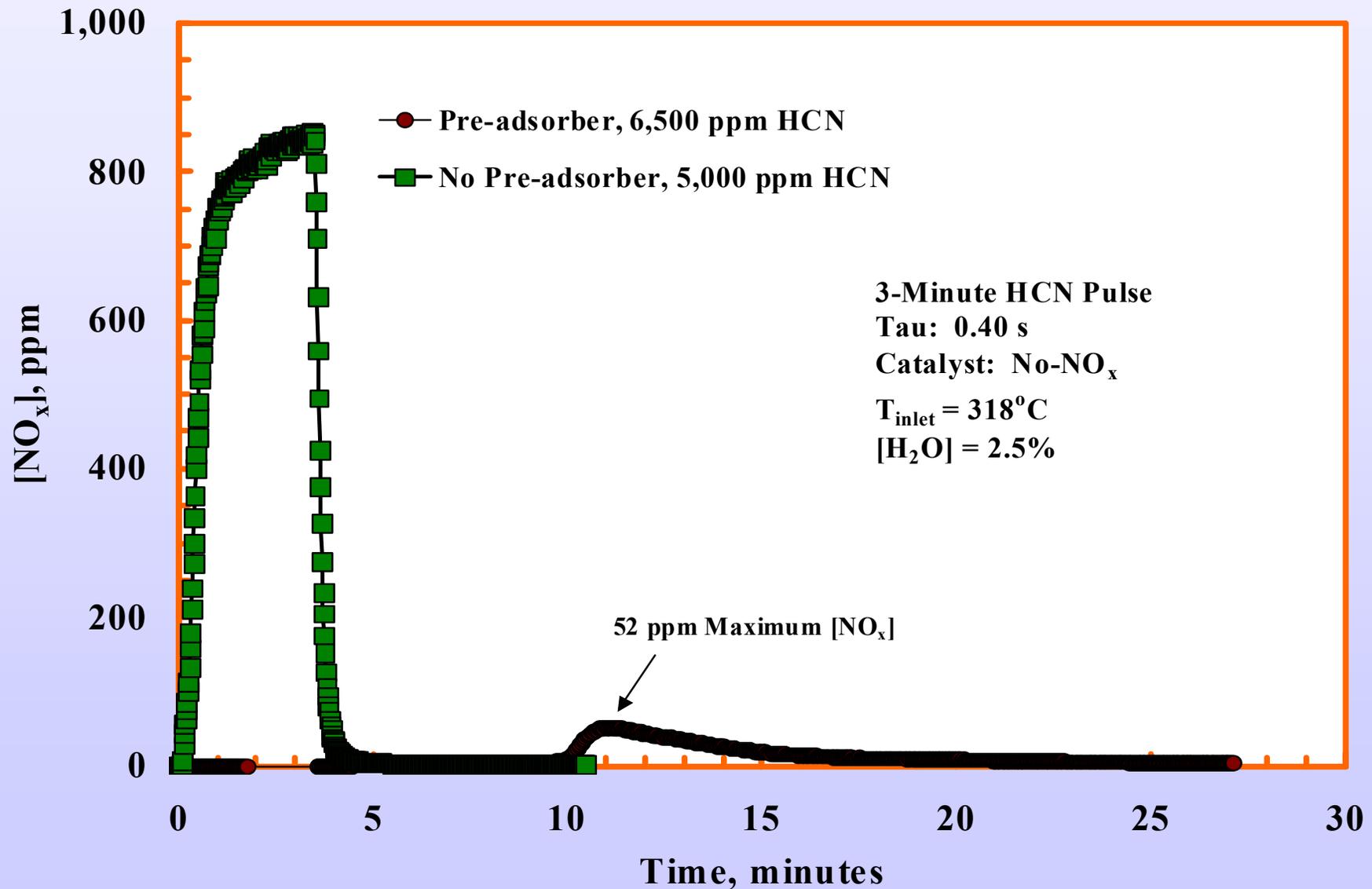
# No Pre-adsorber, Increased Temperature



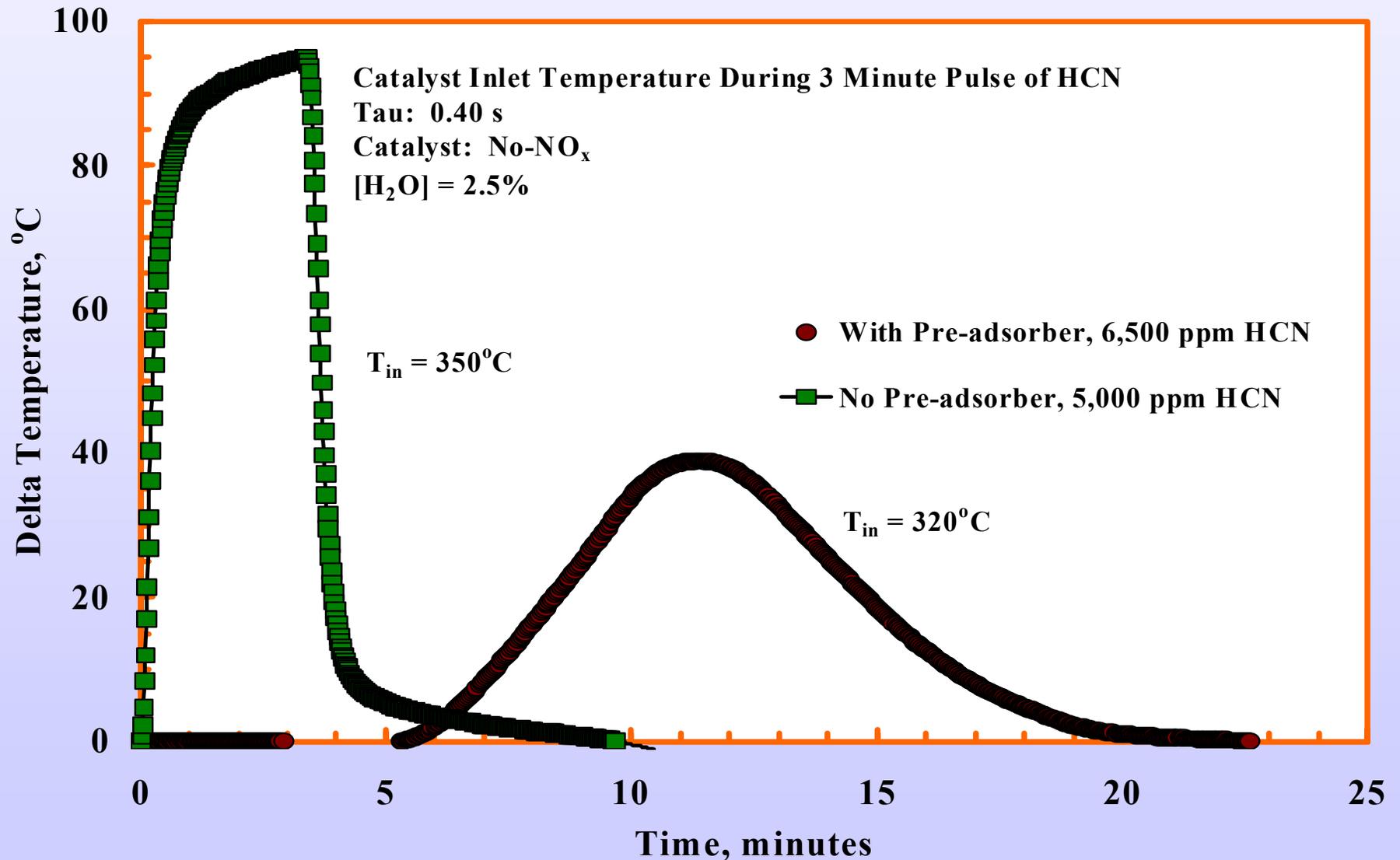
## No Pre-adsorber, Increased Temperature



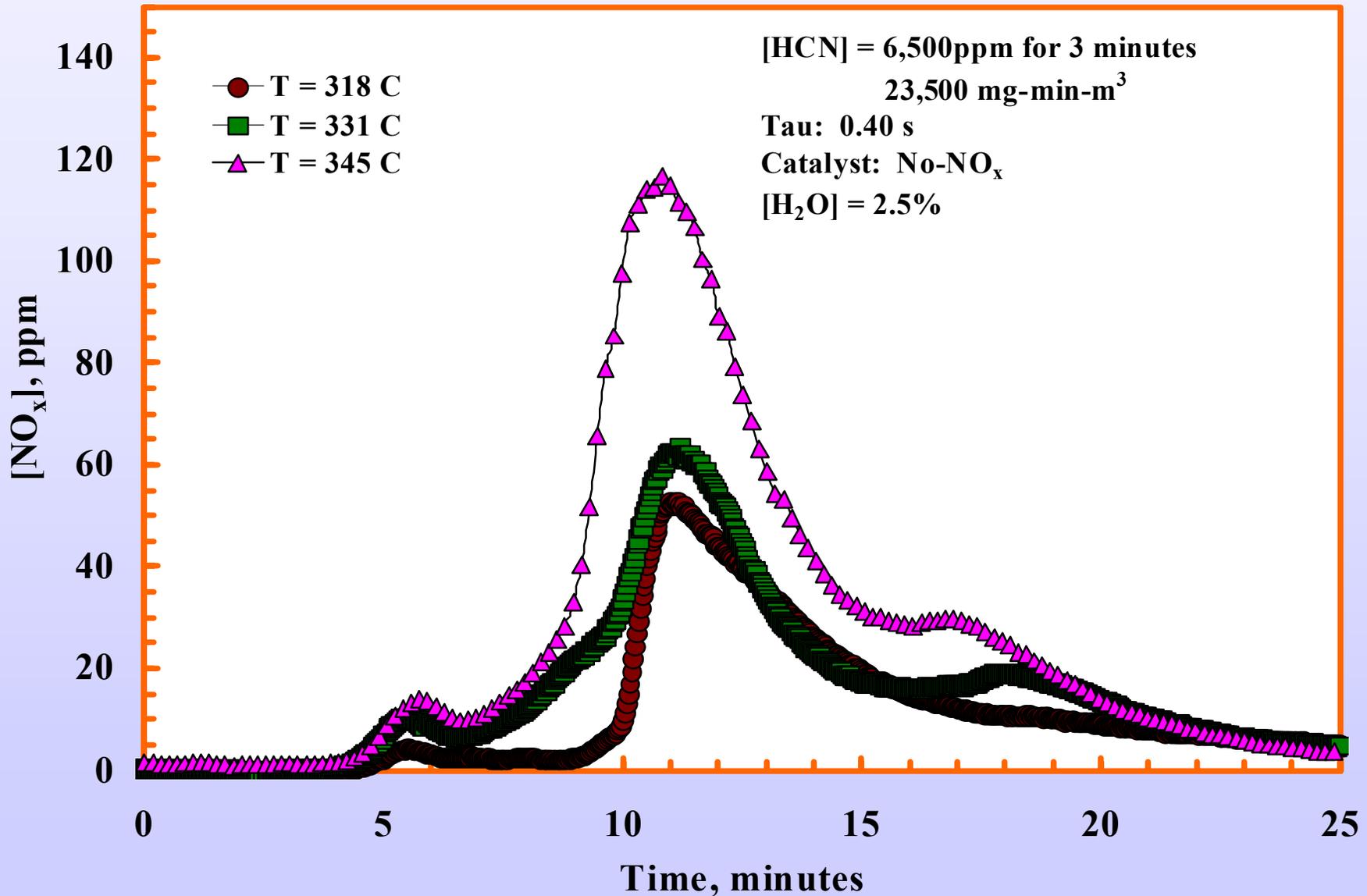
# Use of Pre-adsorber for Destruction of HCN



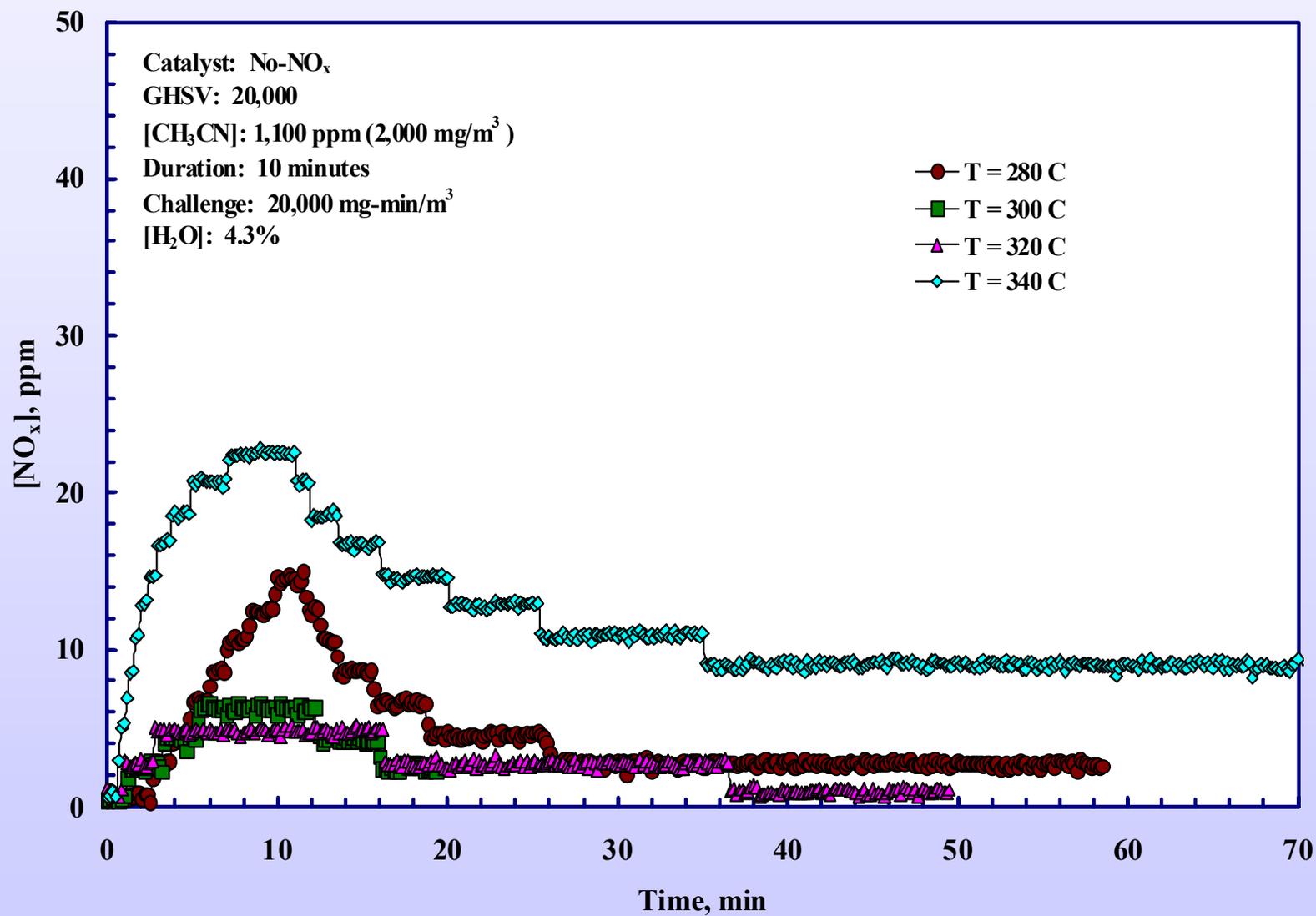
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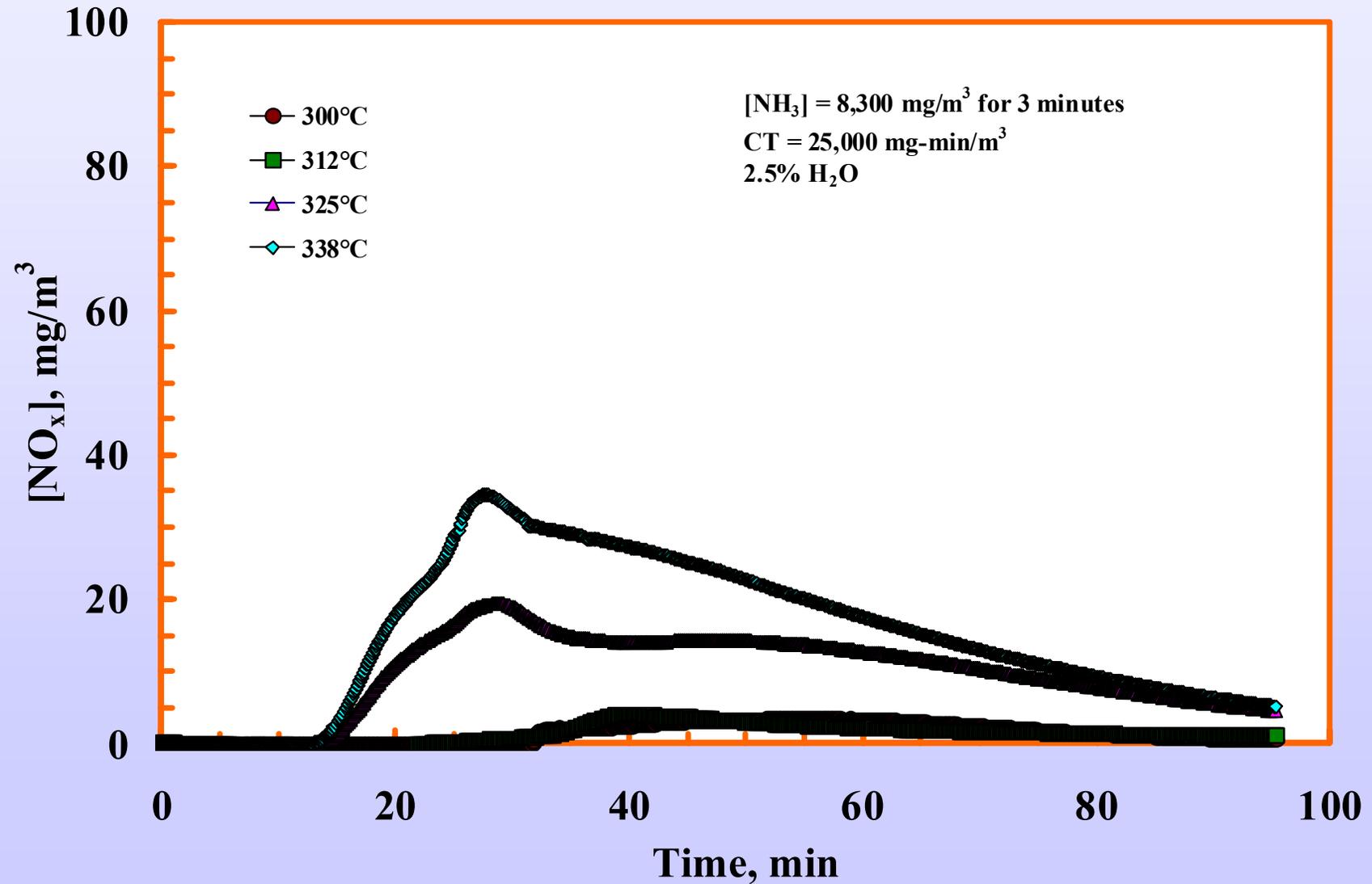
# Use of Pre-adsorber for Destruction of HCN



# Results: Destruction of Acetonitrile using pre-adsorption



## Results: Destruction of NH<sub>3</sub> using pre-adsorption



# Conclusions

- 1.  $\text{NO}_x$  generation during destruction of nitrogen-containing compounds can be greatly minimized using a pre-adsorber.**
- 2. Use of a pre-adsorption bed to minimize concentration excursions greatly improves catalyst performances**
  - o Greatly reduced  $\text{NO}_x$  formation**
  - o Improved thermal management**
- 3. In the case of HCN, not able to meet low  $\text{NO}_x$  requirements. Identification of improved adsorption materials should allow for meeting this objective.**