Development of Regenerable, High-Capacity Boron Nitrogen Hydrides For Hydrogen Storage

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Overview

Timeline

- Project start 3/15/2005
- Project end 11/30/2008
- Percent complete 10

Budget

- Total project funding
 - DOE share \$ 1.6M
 - Contractor share \$ 0.4M
- Funding received in FY04 0
- Funding for FY05 \$ 0.2M
- Funding for FY06 \$ 0.15M

Barriers

- A Storage system and fuel cost
- B Gravimetric/volumetric hydrogen storage density
- R Cost effective regeneration processes

Partners

- State Scientific Research Center of Russian Federation (GNIIChTEOS)
- Internatix Corporation



Objectives

Overall – Develop a boron-nitrogen hydride-based hydrogen storage system to meet U.S. DOE's 2015 hydrogen density and cost targets.

Hydrogen Release – Develop an efficient on-board heating system to extract most of the hydrogen from ammonia-borane (NH₃BH₃).

Regeneration – Develop an off-board, energy efficient, integrated process for regenerating spent materials (~ BN) utilizing commodity hydrogen and chemicals.



Technical Barriers Addressed

- A Storage system and fuel cost (2015 targets: System cost \$67/ kg H₂; Fuel \$2 to 3/GGE (~kg H₂)
- B Gravimetric/volumetric hydrogen storage density (2015 targets: 0.09 kg H₂/kg and 0.081 kg H₂/L)
- C Energy efficiency
- D Durability
- E Refueling Time
- G System Life-Cycle Assessment
- R Cost effective regeneration processes
- S By-Product/Spent Material removal
- T Thermal Management for hydrogen extraction



Phase I Program Approach

- Task 1 Regeneration Demonstrate individual chemical steps for converting BN to ammonia-borane.
- Task 2 Hydrogen Release Ammonia-borane heating approaches for maximum hydrogen yield.
- Task 3 Recycle Spent Products Regeneration of ammonia-borane decomposition products.
- Task 4 Preliminary Storage System Design Develop a preliminary prototype (1 kg hydrogen capacity) design with >9 wt% hydrogen capacity.
- Task 5 Feasibility Determine technical and economic feasibility.



Ammonia-borane Properties

- Molecular Formula NH₃BH₃
- White crystalline solid stable in ambient air
- Hydrogen Content 19.6% by weight
- Density 0.74 g/cm3
- Heat of formation - 42.54 kcal/mole
- Melting temperature ~ 105°C
- Solubility at 20 °C is ~ 33.6 g in 100 g of water.
- Water solutions of Ammonia-borane are also very stable during long term storage.



Ammonia-borane for Transportation

- On-Demand Decomposition by Direct Heating.
- Hydrogen combustion or electrical heating powered by Fuel Cell - Net material-based Hydrogen Density > 17% by wt with complete hydrogen release.
- Spent Boron Nitride must be converted to Ammoniaborane (AB) Off-board in a central processing facility utilizing commodity hydrogen.
- Develop AB Distribution / BN Recycling Network.





Aminoborane in close energy transfer cycle



Technical Accomplishments

- Task 1 Regeneration Initiated BN chlorination studies at laboratory scale – literature search for process conditions, catalysis, reactor materials, product separation and analytical techniques.
- Task 2 Hydrogen Release Conducted controlled ammonia-borane heating studies. Demonstrated release of 85% of available hydrogen in < 30 seconds; confirmed with elemental analysis of the decomposition residue.

Material-based hydrogen density achieved ~ 16.5 wt%

Task 4 – Preliminary Storage System Design Identified components for the prototype design.



Task 1 - Ammonia-borane regeneration

Reaction **Reaction Enthalpy**

 $2 \text{ BN} + 3 \text{ Cl}_2 \rightarrow 2 \text{ BCl}_3 + \text{N}_2$ -36.8 kcal/mole BN (1000 C)

 $2 \text{ BCl}_3 + 6 \text{ H}_2 \rightarrow \text{B}_2\text{H}_6 + 6 \text{ HCl} + 32.1 \text{ kcal/mole BCl}_3 (400 \text{ C})$

 $B_2H_6 + 2NH_3 \rightarrow 2 BH_3NH_3$

-20.9 kcal/mole BH_3NH_3 (30 C)

Overall Reaction

 $2 \text{ BN} + 3 \text{ Cl}_2 + 6 \text{ H}_2 + 2 \text{ NH}_3 \rightarrow 2 \text{ BH}_3 \text{ NH}_3 + \text{ N}_2 + 6 \text{ HCl}$

Overall Reaction Enthalpy ~ -25.6 kcal/mole BH₃NH₃ (exothermic)



Task 1 – Activities

- Literature review for conducting BN chlorination; process temperature ~ 1000 C; catalysis – group lb, VIIb and VIII element chlorides; reactor materials – quartz, alundum; product separation and analysis.
- Diborane synthesis by hydrogenation of boron chloride - the most challenging process step but is industrially practiced.
- Prior experience in synthesizing AB from diborane.
- Laboratory apparatus assembled for studying BN chlorination, product separation, and analysis of gas streams and liquid samples.



Task 2 - Hydrogen Release

- Two possible modes of supplying heat: direct hydrogen combustion, Resistive heating using fuel cell power.
- AB cartridge design: a) Fixed material cartridge with heat directed to progressive zones – simple compartmentalized cartridge design -- slower heating rate, intermediate products.
- AB cartridge design: b) dispensing in a fixed hot zone rapid heating rate, need separate product storage and solid powder/pellet dispensing system.



Task 2 – TGA-MS Activities

- TGA studies conducted with controlled heating patterns, ramp rate, hold times and final temperatures.
- All TGA studies experienced mechanical loss of sample due to melting and subsequent gas release through melt regardless of ramp rate.
- Maximum rate of weight loss was at 100 °C coincident with first stage hydrogen release.
- Exit gas analyzed by online Mass Spectrometer for species identification.



TGA Sample weight loss

TGA of Aminoborane





Rate of Weight Loss with Temperature





MS analysis of Gas Released - Hydrogen





MS analysis of Gas Released - Borazine

Ammonia Borane mass spec data / mass 80





Conclusions from TGA-MS studies

- First stage hydrogen release occurs at ~ 100 °C with a second stage release at ~140 °C. Hydrogen release at < 90 °C temperature below MS detection limit.</p>
- Melting of ammonia-borane also around 100 °C caused foaming of sample and mechanical weight loss in TGA studies.
- All of the detectable higher MW emissions occurred after reaching 175 °C, e.g. MW = 54 (B₂N₂H₄) and MW = 81 (B₃N₃H₆ - borazine).
- Melting of ammonia-borane and foaming as hydrogen is released – possible problem in progressive heating approach.



Task 2 – High Temp. Reactor Studies

- High temperature tubing reactor used for quantitative determination of hydrogen released as a function of reactor temperature and sample hold time.
- Reactor temperature was varied from 400 to 500 °C and sample hold time was varied from 15 sec to 10 minutes.
- Amount of gas generated was measured by direct water displacement.
- Ammonia-borane decomposition residue analyzed for elemental composition.



High Temperature Reactor Test Results

- Specific gas release of ~ 1980 cc/g ammonia-borane at 500 °C. (~ 85% of the maximum possible hydrogen).
- Material-based hydrogen density achieved ~ 16.5 wt%.
- Specific gas release at 500 °C ~ 10% greater than at 400 °C.
- Hold time of 30 seconds is adequate for gas release.
- Elemental analysis of the residue indicated composition of BN_{0.9}H – consistent with observed specific gas release.
- GC analysis of gas indicated hydrogen purity ~ 98.5%.
- The nature of impurities analyzed by Mass Spectrometry.



MS analysis of HT Reactor Gas

Dycor System 2000

09/27/05 09:25:35 092705b 001 002.dat





MS analysis of Gas bubbled through water

Sample bubbled thru water

Dycor System 2000

09/27/05 17:21:33 sample H2O.dat



MS analysis of HT Reactor Gas

- MS analysis of the HT reactor gas indicated possible presence of cyclic B,N,H compounds (e.g. MW = 39 (B_2NH_3) , 54 $(B_2N_2H_4)$, 64 $(B_3N_2H_3)$, and 81 $(B_3N_3H_6)$ with successive addition of B/N atom).
- Need positive identification of individual species.
- These species appear to be stable even at 500 °C.
- Passing the gas through a water bubbler significantly reduced the concentration of these species by a factor of 2 to 3.



Task 4 - Preliminary Storage System design

- Components identified reagent/product storage chambers, heating system for the reactor chamber, solid feed and extraction system, buffer hydrogen tank, filter/purifier if needed, and pressure sensor-based control system.
- Solid feed with an adequate hold time in the reactor.
 Allowable design pressure and ammonia-borane feed rate/injection will determine volume of the reactor.
- Buffer tank capacity determined by the startup hydrogen requirements.
- Heat can be supplied by direct hydrogen combustion or electrical heating during startup. The hydrogen release process is expected to be self-sustaining after start-up.



Activities anticipated in FY06

- Task 1 Continue evaluation of process conditions and catalysts for maximizing product yield in each of the individual reaction steps starting from BN chlorination.
- Task 2 Continue evaluating direct and catalytic decomposition approaches for hydrogen release.
- Task 3 Determine additional considerations for starting with decomposition product instead of BN in Task 1, Evaluate catalytic regeneration of decomposition products.
- Task 4 Develop preliminary design of an on-board hydrogen storage and delivery system.
- Task 5 Estimate costs of regeneration process, stored hydrogen, and the hydrogen storage system.



Activities anticipated in FY07

- Task 1 Finalize process conditions and catalysts for maximizing product yield in each individual reaction steps.
- Task 2 Select a heating system and process conditions for maximum hydrogen yield.
- Task 3 Demonstrate regeneration of ammonia-borane decomposition products.
- Task 4 Complete the preliminary design of an on-board hydrogen storage and delivery system using the heating approach and conditions identified in Task 2.
- Task 5 Update estimates of costs of stored hydrogen and the storage system



Phase I Technical Milestones

- Synthesis of pure ammonia-borane from boron nitride at a laboratory scale.
- Demonstration of a process, suitable for on-board deployment, for extraction of pure hydrogen from AB.
- Demonstrate lab-scale synthesis of pure ammonia-borane starting from AB decomposition products.
- Design a prototype, 1 kg hydrogen capacity, on-board ABbased hydrogen storage system with > 9 wt% H₂ capacity.
- Determine technical and economic feasibility of ammoniaborane-based hydrogen storage system leading to a Go/No Go decision point for continuation to Phase II.



Project Summary

Relevance – Develop a hydrogen storage system to meet DOE's 2015 hydrogen density and cost targets.

- Approach Utilize thermal decomposition of ammoniaborane (19.6 wt% hydrogen) to produce hydrogen onboard. Regenerate spent materials chemically off-board.
- Technical Accomplishment Demonstrated release of 85% of available hydrogen in < 30 seconds. Materialbased hydrogen density achieved ~ 16.5 wt%. Identified a chemical route for regeneration of spent products.
- Future Activities Demonstrate efficient ammoniaborane decomposition and regeneration processes.

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Hydrogen Safety

The most significant hydrogen hazards:

- Handling toxic chemicals such as diborane, chlorine, hydrogen and ammonia during evaluation of aminoborane synthesis processes.
- Extreme conditions during synthesis processes, e.g. high and low temperatures (e.g. 1000 C to -200 C).
- Uncontrolled release of hydrogen during evaluation of approaches for hydrogen extraction from aminoborane.



Hydrogen Safety

Our approach to deal with these hazards:

- Safe laboratory practices during handling of chemicals and extreme process conditions eliminating exposure to TLV levels of toxic chemicals involved.
- Limit scale of experiments to assure safety during synthesis experiments, increasing the scale gradually as experience is gained in handling hazardous process conditions.
- Design hydrogen extraction experiments so as to eliminate any possibilities of uncontrolled release as well exposure of hydrogen to any open flame.
- Conduct all experiments in ventilated fume hood areas

