

Clean Energy Research

Project III: Hydrogen Storage Using Chemical Hydrides

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STP38

This presentation does not contain any proprietary or confidential information

Overview

Timeline

- 10/31/2004-10/31/2006
- 55% complete

Budget

- Total project funding
 - \$559,875 DOE
 - \$134,250 contractor
- FY05 funding
 - \$365,000 DOE
 - \$91,250 contractor
- Funding for FY06
 - \$194,875 DOE
 - \$43,000 contractor

Barriers

- System Weight and Volume
- Thermal Management
- Discharging Rates

Partners

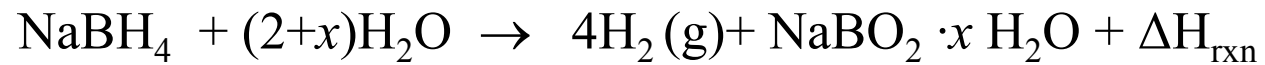
Millennium Cell, Inc

Eatontown NJ

Dr. Ying Wu, Director R&D

Objectives

- Develop hydrogen storage and delivery technology based on steam + chemical hydrides for automotive fuel cell applications



- Quantify/optimize steam + solid chemical hydride reaction kinetics as basis for production of hydrogen
- Compare experimental data to FreedomCAR targets
 - Mass efficiency of reaction (8MAA)
 - Hydrogen production rate / kinetic data (12MAA)
 - Analysis of water utilization of reactor and hydration characteristics of products (14MAA)
 - Prototype design development (16 MAA)
- Develop prototype of steam hydrolysis reactor

Approach

- Hydrolyze chemical hydrides with dry steam, rather than aqueous catalytic process
 - Chemically simple reaction with humid H₂ gas product
 - Minimal water inventory in the reactor
 - Autothermal integration: use reaction heat to produce steam
- Operate reactor at low temperatures (100 – 150°C) and pressures (~ atmospheric)
- Basic research to minimize water utilization and maximize H₂ delivery rate
- Apply research to design a prototype system

Critical Assumptions and Issues

- Unreacted water from steam hydrolysis can be recovered & recycled to the reaction
- Heat of reaction can be recovered
- Cost and energy requirements of manufacturing and recycling NaBH_4 are crucial; research by others is addressing this.

Hydrogen Safety

- The most significant hydrogen hazards associated with this project are:
 - High reactivity of solid chemical hydrides when exposed to humidified air
 - Instability of aqueous NaBH_4 solutions
 - Toxicity and corrosivity of basic NaBH_4 solutions
- Our approach to deal with these hazards:
 - Steam/solid reaction minimizes water inventory in the reactor
 - Reactant is stored and transported in dry form
 - Reaction products are nearly dry, not an aqueous basic solution

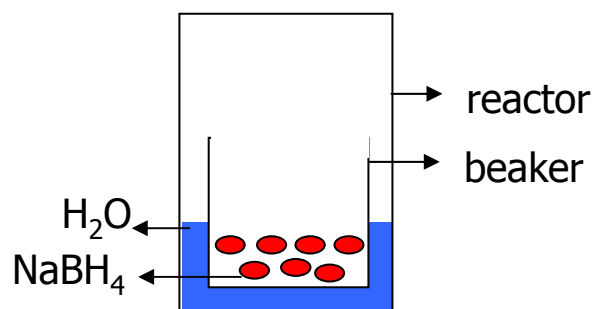
1. Characterization of Hydrolysis Reaction Products (Hydrated Metaborates)



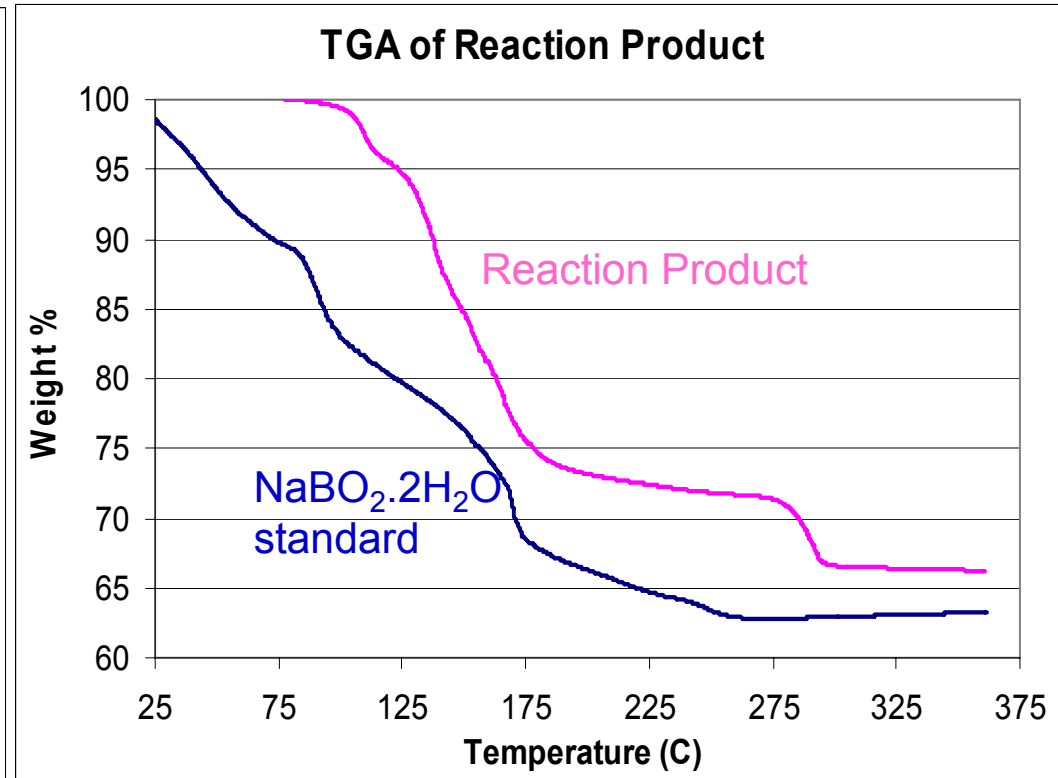
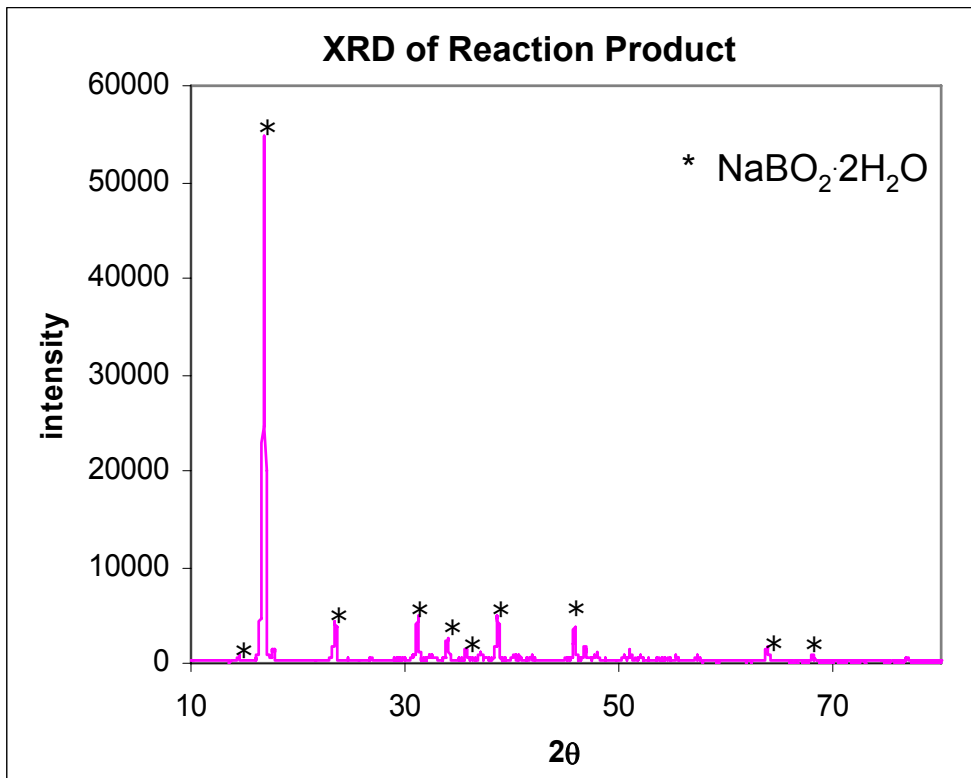
NaBH_4 was placed in the small beaker inside an isolated vessel as shown at left. Hydrated metaborates, with unknown degree of hydration x , were generated by controlled exposure of NaBH_4 to H_2O vapor.

$\text{NaBO}_2 \cdot x\text{H}_2\text{O}$ were characterized using TGA and XRD; analyses were repeated for standard samples of $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ and $\text{NaBO}_2 \cdot 4\text{H}_2\text{O}$.

Results are shown on the next two slides.



1. Characterization of Hydrolysis Products after Controlled Exposure to Steam

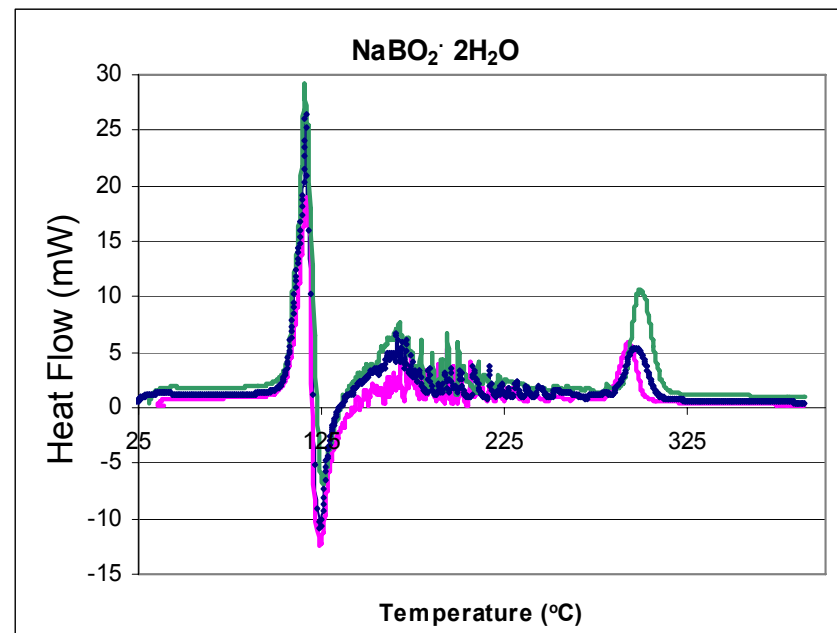
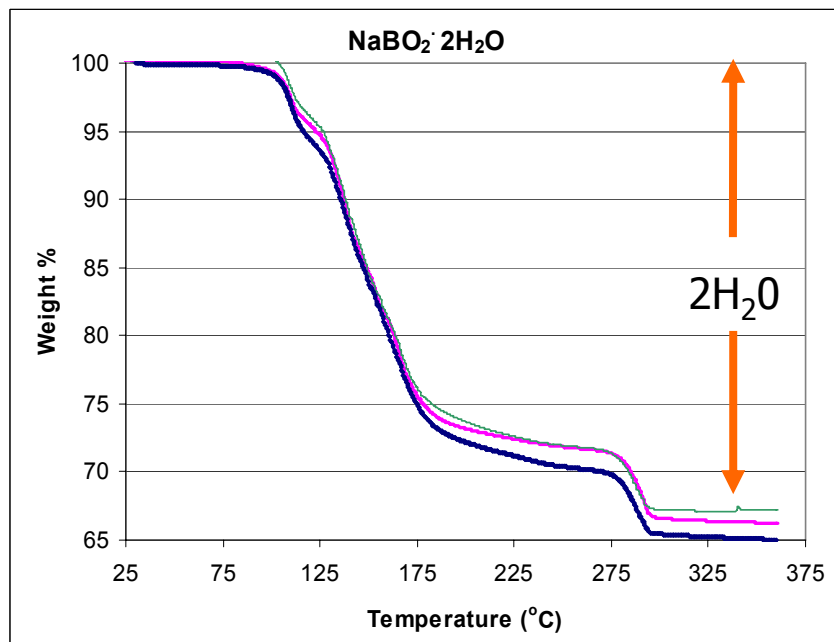


XRD of reaction product corresponds to $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$

TGA of reaction product is consistent with $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ standard

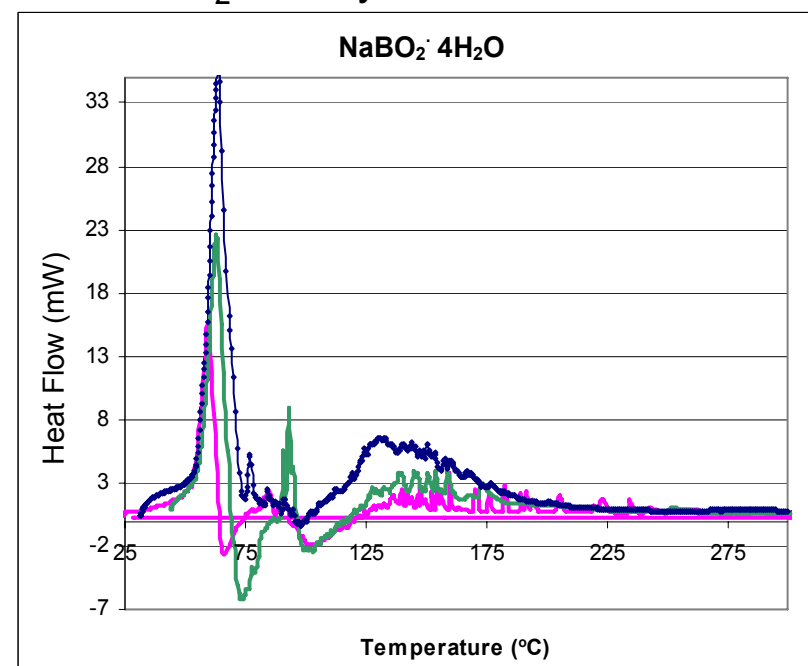
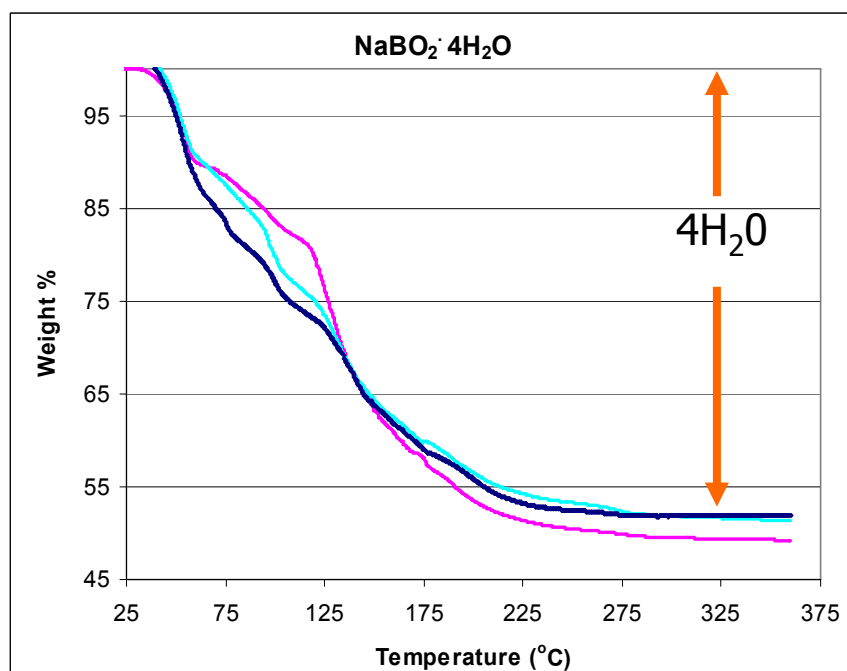
$\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ was the primary product of the hydrolysis reaction. This translates to a decrease in the potential gravimetric and volumetric efficiencies of 33%

1. Characterization of Borate Standards

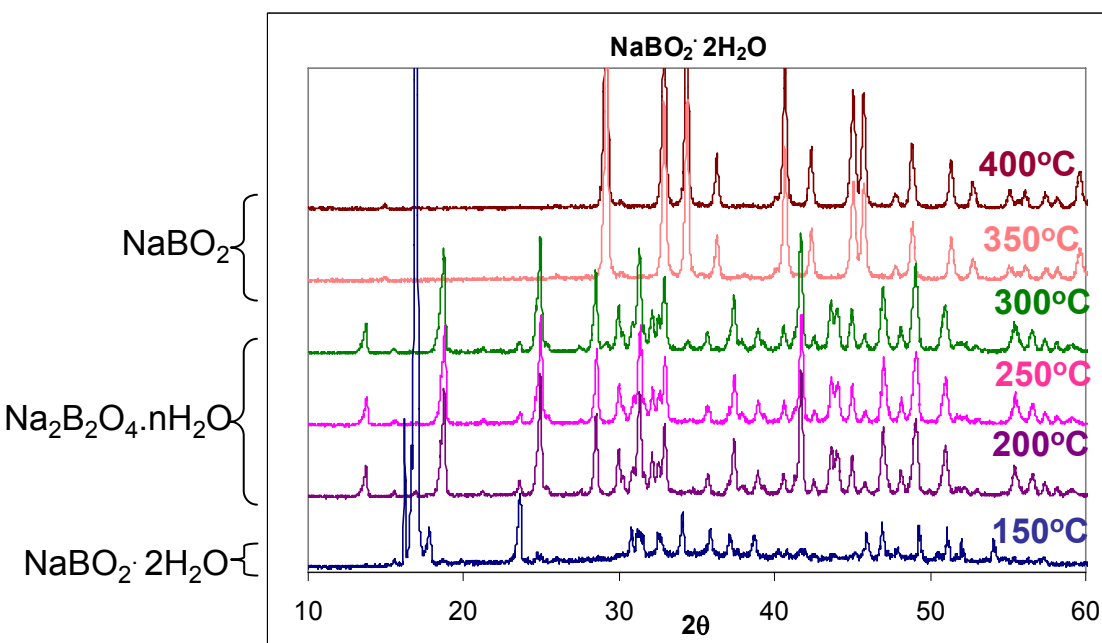
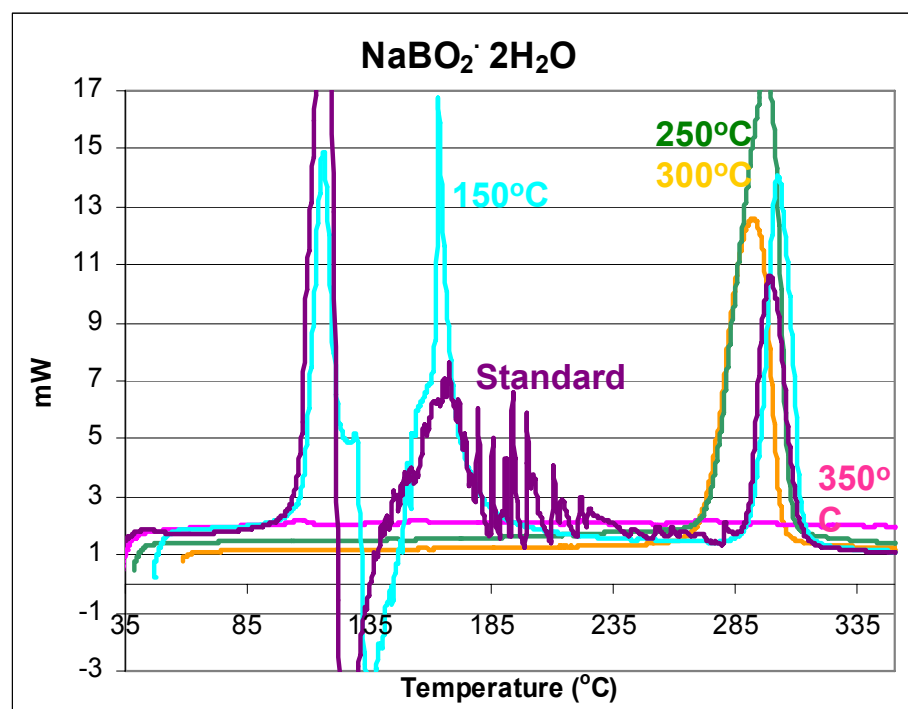
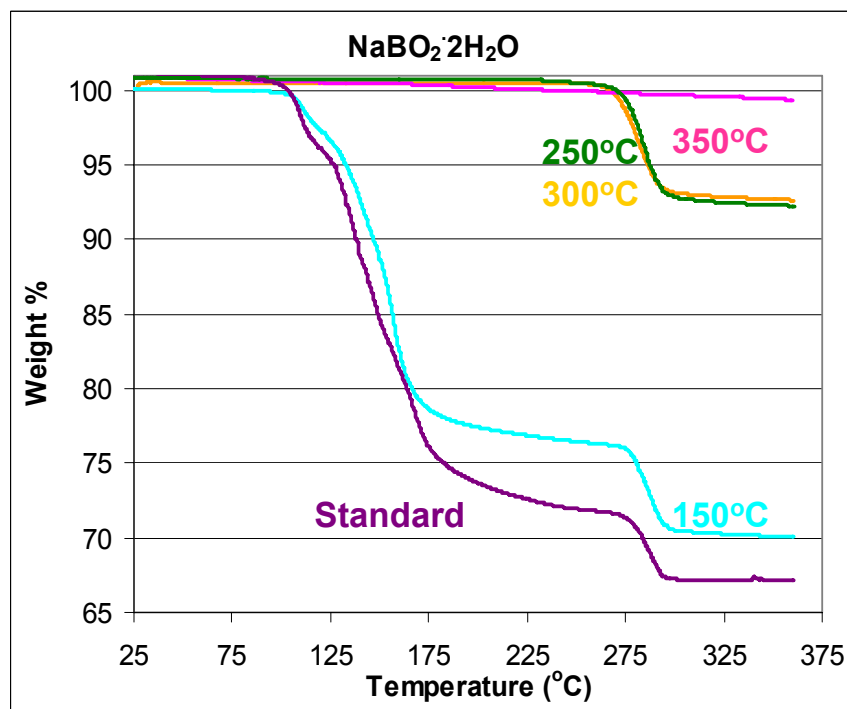


NaBO₂·2H₂O lost all its water (35%) before 400°C
NaBO₂·4H₂O lost all its water (52%) before 400°C

NaBO₂·2H₂O shows crystal transformation at 280 °C
NaBO₂·4H₂O is less reproducible due to larger amount of H₂O in crystal structure



1. Thermal Dehydration of $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ Standard



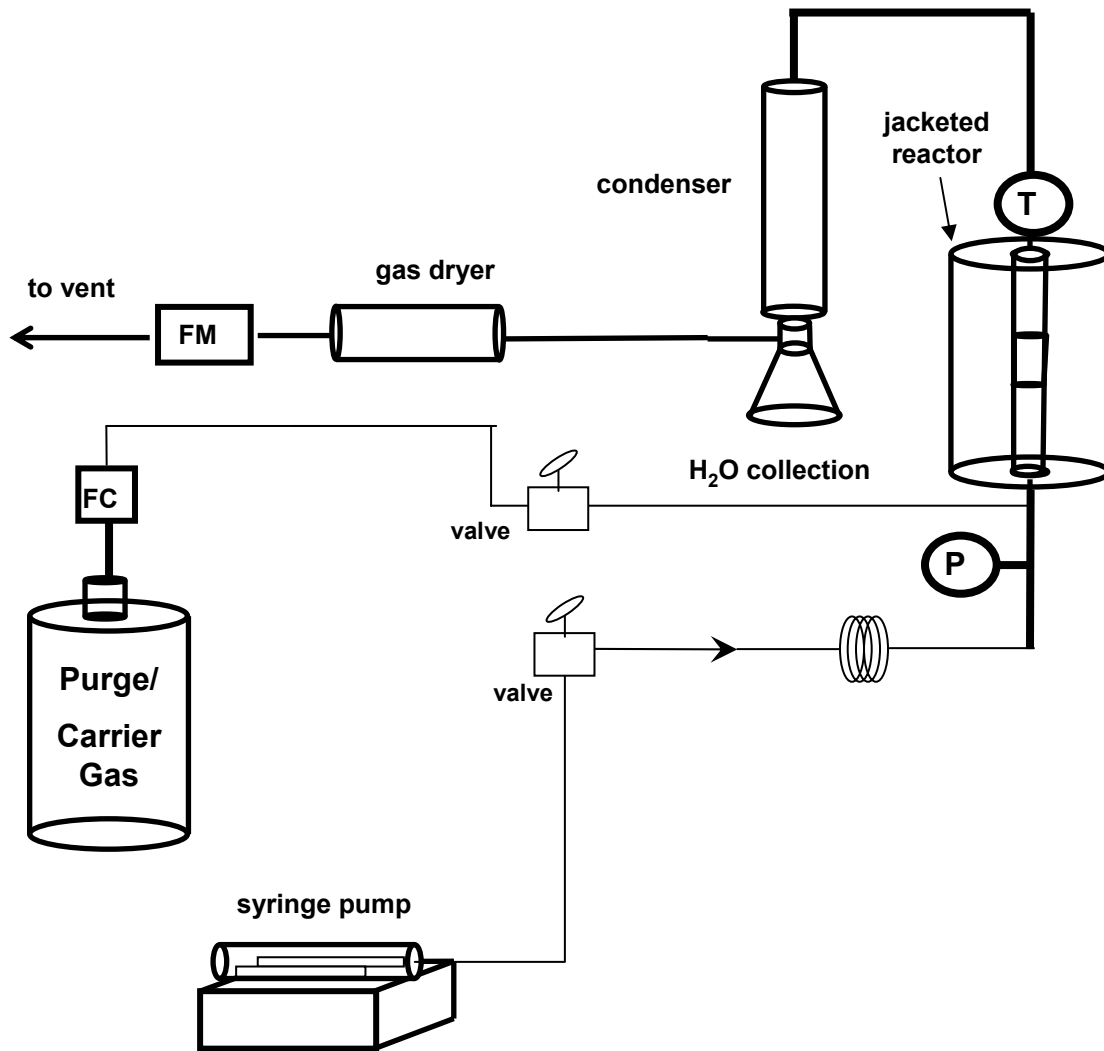
TGA: $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ dehydrates in stages below 350°C

DSC: Peak at 280°C disappears after crystal structure transformation to NaBO_2 occurs

XRD: $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ changes crystal structure with water loss

$\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ transforms into NaBO_2 above 300°C

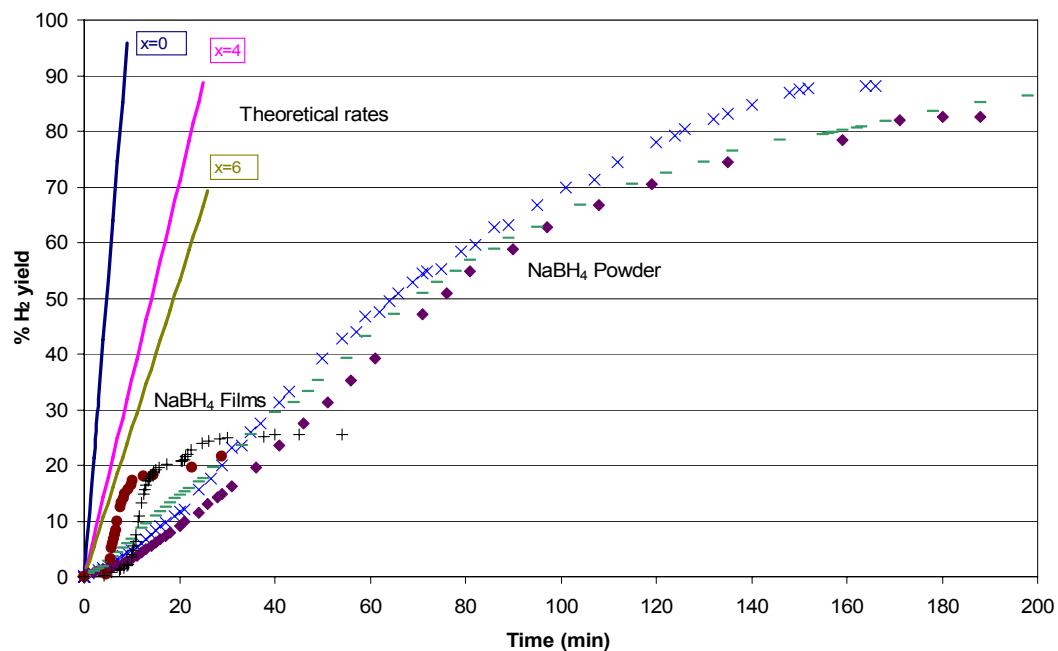
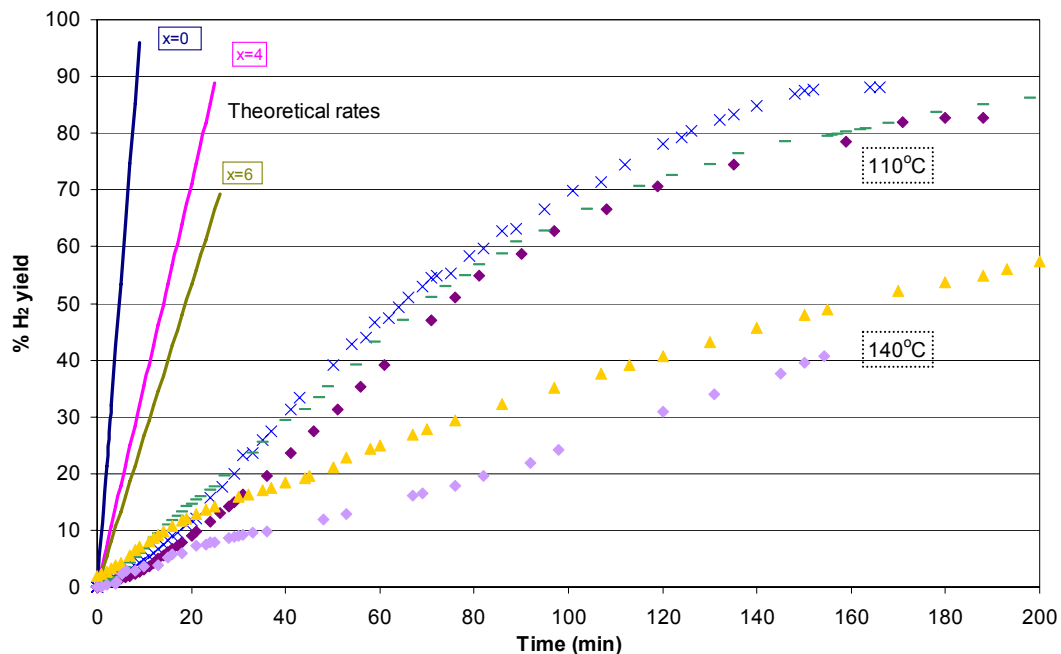
2. Steam Hydrolysis Kinetics: Apparatus and Procedure



- Heated packed bed reactor in upflow configuration
 - FY05 used tilted orientation
- N₂ sweep gas is used for inert blanketing
- Product gas flow rate is measured with digital flow meters.
 - FY05 used H₂O displacement
- Water in the product stream is condensed and quantified
- Initial $T_{rxr} = 110^{\circ}\text{C}$
- Liquid water flow rate = 0.1 mL/min

2. Results from FY05

- Good yields, low rates at 110°C
- Poor yield and rates at 140°C
- Thin films give higher initial rates
- Decreased yields are attributable to insufficient reactant contact



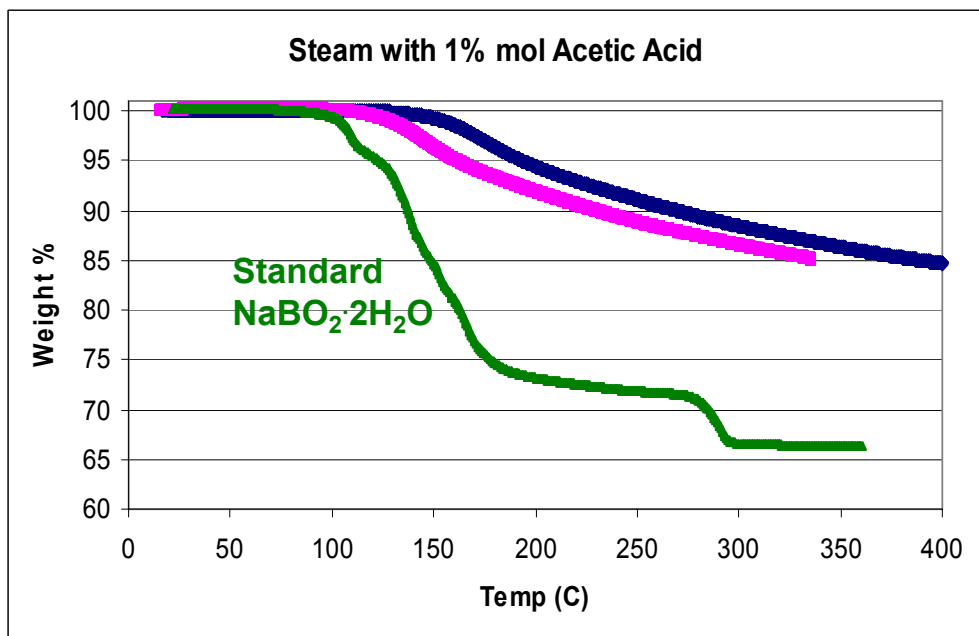
Description	T (°C)	Max Slope (mol/kgNaBH ₄ *min)	% of Theo. H ₂ Yield
NaBH ₄ powder	110	1.018	88.2
NaBH ₄ powder	110	0.870	82.7
NaBH ₄ powder	110	0.825	94.0
NaBH ₄ powder	140	0.628	67.1
NaBH ₄ powder	140	0.327	40.8
NaBH ₄ on glass beads	110	4.42	19.7
NaBH ₄ on glass beads	110	5.14	22.9
Theoretical Rates	x=0	11.1	100
	x=4	3.70	100
	x=6	2.78	100

2. Representative Results, FY 06

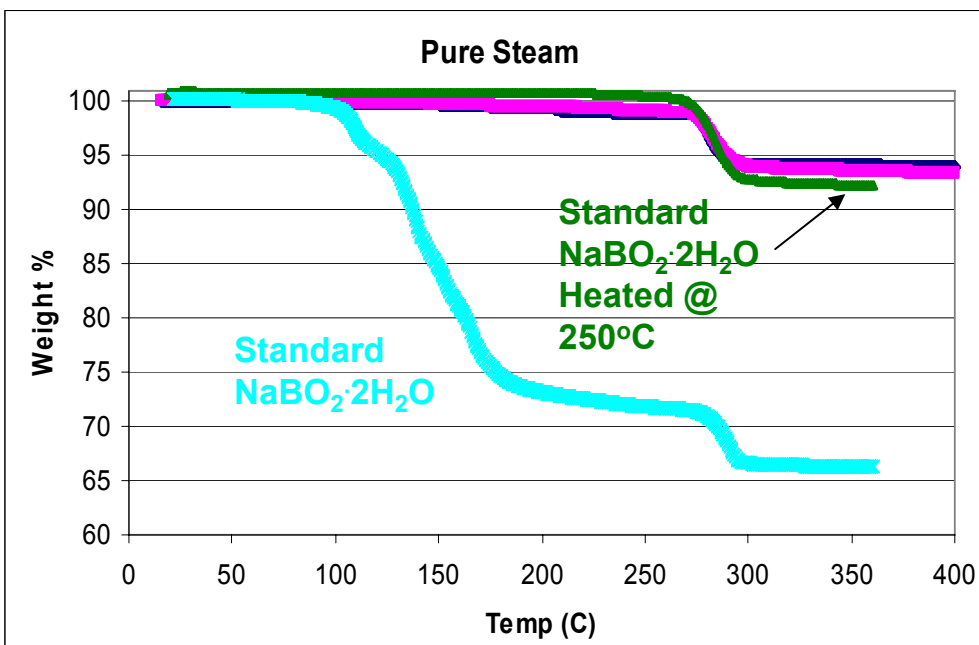
Condition of Reactant Stream	Initial Rate (mol/kg*min)	Theoretical H ₂ Yield %	H ₂ O Collected (% of amt fed)	Gravimetric Efficiency, % (mass H ₂ /(mass NaBH ₄ + H ₂ O))
Pure Steam	0.843	72.5%	53%	1.34
Pure Steam	0.892	85.4%	92.5%	7.31
Pure Steam	0.906	90.3%	87.5%	5.01
1mol% HAc	0.790	92.4%	75%	3.0
1mol% HAc	1.044	92.7%	90%	6.6
15 mol% MeOH	0.629	72.9%	90%	4.85

Gravimetric efficiency assumes that condensed water can be recovered and recycled.

2. TGA of Products from Reactor



Reaction products allowed to dry for several hours under inert N_2 atmosphere prior to characterization analysis



Reaction with acetic acid vapor solution generates product other than $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$

Reaction with pure steam yields product similar to $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ that was preheated to 250°C

3. ^{11}B NMR Analysis

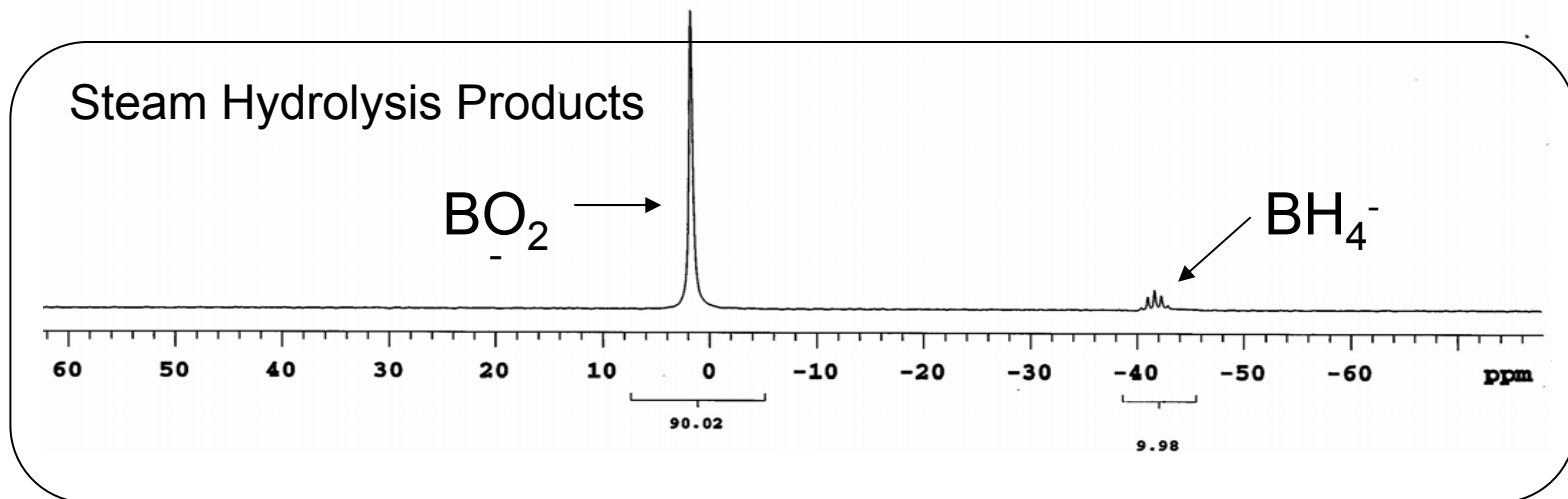
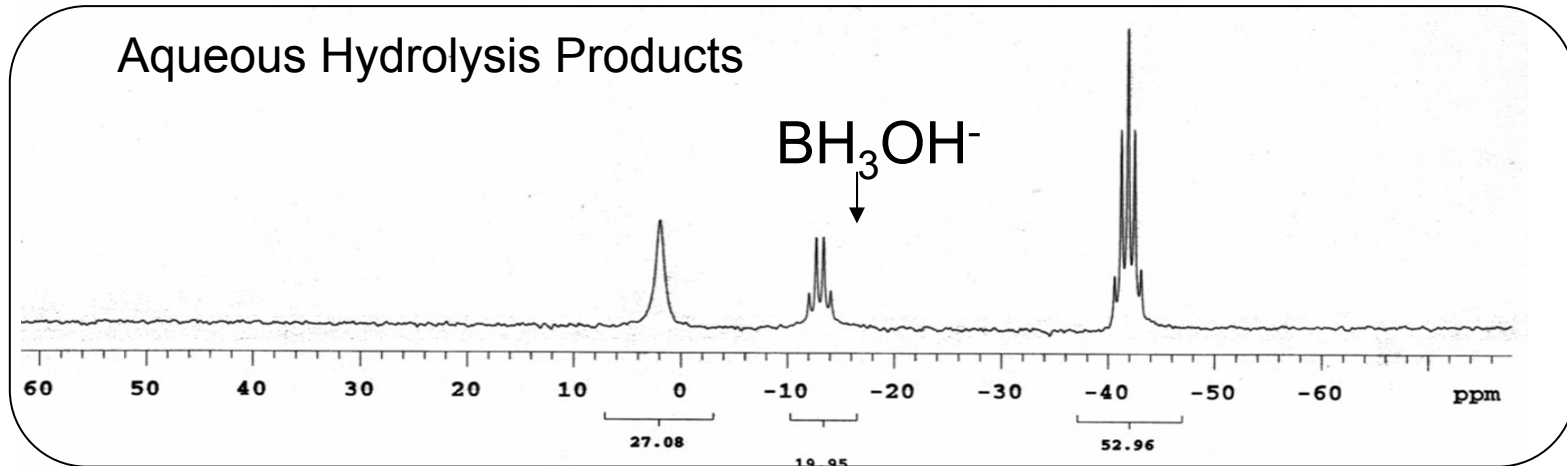
Objectives

- Characterize solid reaction products from steam hydrolysis; compare to liquid-phase reaction intermediates
 - Detect BO_2^- and unreacted BH_4^- in solid products
- Analyze condensate from product stream
 - Detect carry-over of boron-containing species (if any)
- Use these results to help interpret steam hydrolysis reaction pathway and differences from aqueous hydrolysis pathway

Procedure

- 1mL of sample dissolved in 0.1M NaOH in 5mm NMR tubes
- Analyzed using a Varian Mercury/VX 400 with a $\text{BF}_3 \cdot \text{O}(\text{C}_2\text{H}_5)_2$ reference

3. ^{11}B NMR Analysis



BH_3OH^- intermediate does not appear in analysis of steam reaction indicating a different reaction mechanism than the aqueous reaction. Understanding the steam hydrolysis mechanism may lead to strategies to reduce water usage and lower system weight.

4. Commercial / Industrial Application Development

- Steam hydrolysis of NaBH_4 is the basis of a start-up company DEnergy LLC.
- 3 invention disclosures have been filed with the University of South Carolina.
- Prototype design is currently being tested and developed for preliminary portable power applications
 - Gateway approach to automotive technologies
- More information is available at <http://ip.research.sc.edu/>

5. Future Research Directions

FY06

- Steam/solid NaBH₄ system
 - Measure intrinsic kinetic rate of reaction under different operating temperatures and partial pressures of steam
 - Determine hydration characteristics of products in order to improve gravimetric efficiency and understand kinetic limitations
- Investigate additional solid hydride systems
 - Evaluate additional hydrides and mixtures to meet FreedomCAR requirements
- Submit description of prototype system design
 - Design will be based on laboratory-scale experiments
 - Design will be evaluated according to FreedomCar targets such as mass and volumetric efficiency and startup dynamics

Summary

- Yields approach 100% with NaBH_4 in powder form with pure steam, and also with addition of acetic acid and MeOH
- Acetic acid and MeOH additives give consistent high H_2 yields
- Reaction product, $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$, corresponds to an excess hydration factor of 2 ($x=2$)
 - Potential limitation on gravimetric/volumetric efficiency
- Hydration and crystal structure of $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ are affected by reaction temperature and post-treatment.
 - Complete dehydration of $\text{NaBO}_2 \cdot 2\text{H}_2\text{O}$ occurs at $T < 350^\circ\text{C}$

Publications and Presentations

1. Michael A. Matthews, Thomas A. Davis, and Eyma Y. Marrero-Alfonso, “Hydrogen storage in chemical hydrides”. **ACS National Meeting**, Philadelphia. (Aug. 2004)
2. Michael A. Matthews, Thomas A. Davis, and Eyma Y. Marrero-Alfonso, “Production of hydrogen from chemical hydrides via hydrolysis with steam”. **AIChE Annual Meeting**, Austin. (Nov 2004)
3. Steam Hydrolysis of Chemical Hydrides: Meeting the Challenge of Hydrogen Storage. Marrero-Alfonso, E.Y., Gray, J., Matthews, M.A., Davis, T.A. 230th ACS National Meeting, Washington, DC, Aug 28-Sept 1, 2005.
4. New method for hydrolysis of chemical hydrides. Marrero-Alfonso, E.Y., Gray, J.R., Davis, T.H., and Matthews, M.A. Second International Conference on Green and Sustainable Chemistry, and Ninth Annual Green Chemistry and Engineering Conference, Washington D.C. June 20-24, 2005.
5. J.R. Gray, E.Y. Marrero, T.A. Davis, and M.A. Matthews, “Steam Hydrolysis of Chemical Hydrides : Meeting the Challenge of Hydrogen Storage”. **42nd Power Sources Conference** Philadelphia, Pennsylvania. June 12-15, 2006.