

Kidwell, D., et al. *Does Gas Loading Produce Anomalous Heat? (PowerPoint slides)*. in *15th International Conference on Condensed Matter Nuclear Science*. 2009. Rome, Italy: ENEA.

This is a PowerPoint presentation converted to Acrobat format. The original PowerPoint slides are here:

<http://lenr-canr.org/powerpoint/KidwellDdoesgasloa.ppt>

Here is the abstract for this paper:

Does Gas Loading Produce Anomalous Heat?

David A. Kidwell, Allison E. Rogers, Kenneth Grabowski, and David Knies
Chemistry Division, Naval Research Laboratory, Washington, DC 20375;
Materials Science and Technology Division, Naval Research Laboratory, Washington, DC 20375

Simple pressurization of nanosized palladium with deuterium appears to be a simpler and more rapid method to generate anomalous heat compared to electrolytic systems. A survey of the literature indicates that palladium particles less than 2 nm in size can obtain a Pd/D loading near one at modest deuterium pressure. In hundreds of reactions, we have routinely prepared palladium nanoparticles inside an aluminosilicate matrix and have found that these systems produce up to 8 fold more heat with deuterium compared to hydrogen. Furthermore, a characteristic signature of a pressurization reaction is its reversibility – the heat released upon pressurization should be absorbed upon evacuation. This reversibility is observed with hydrogen but not deuterium. Although we are still seeking conventional explanations for this excess heat, the anomalous heat does not appear to be explained by impurities in the deuterium gas nor other simple chemical or physical sources. The selection and preparation of the particles, the experimental set-up, and results will be discussed.

This is from ENEA. *Abstracts*. in *15th International Conference on Condensed Matter Nuclear Science*. 2009. Rome, Italy, p. 41.

<http://lenr-canr.org/acrobat/ENEAabstracts.pdf>

Does Gas Loading Produce Anomalous Heat?

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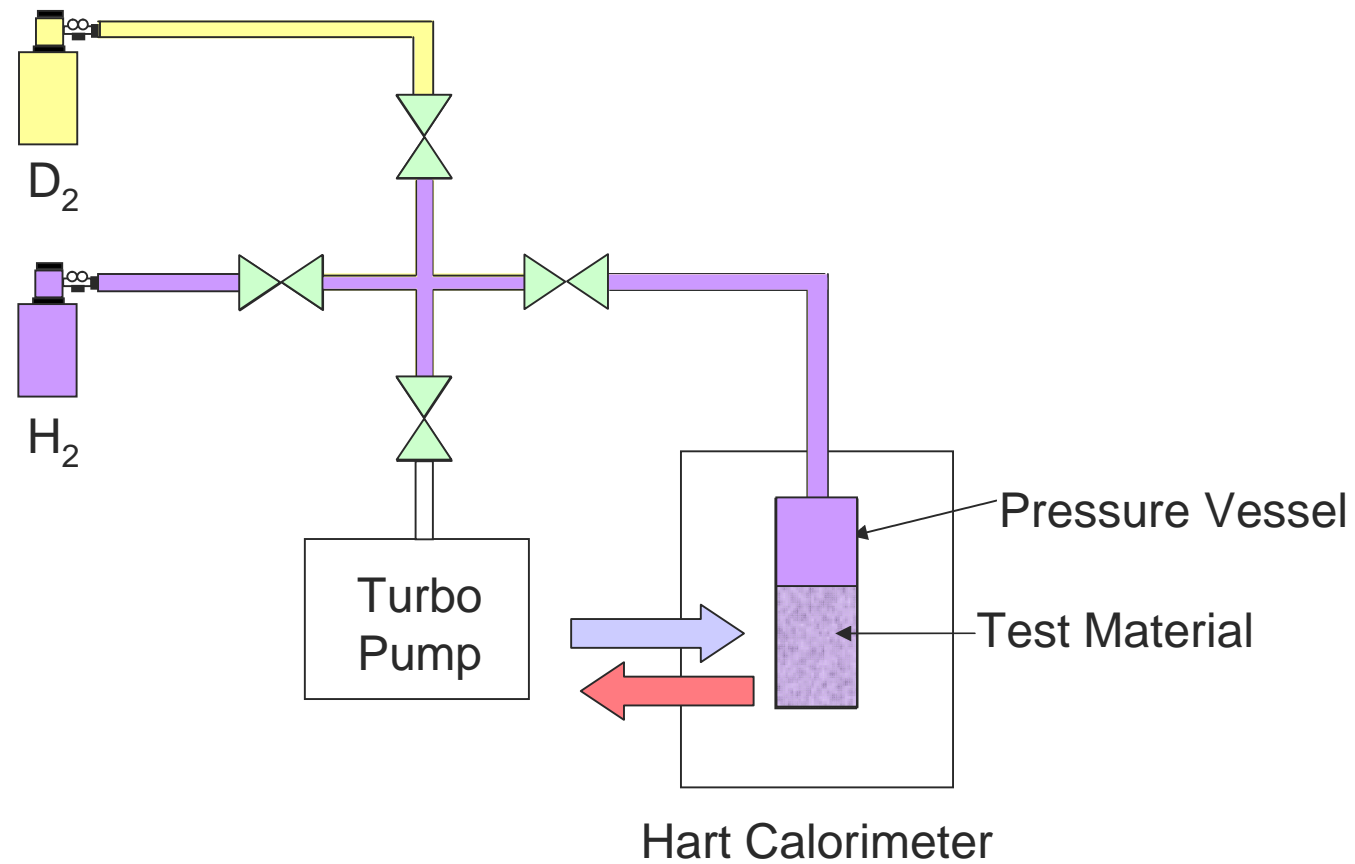
(202)767-3575

David.Kidwell@nrl.navy.mil

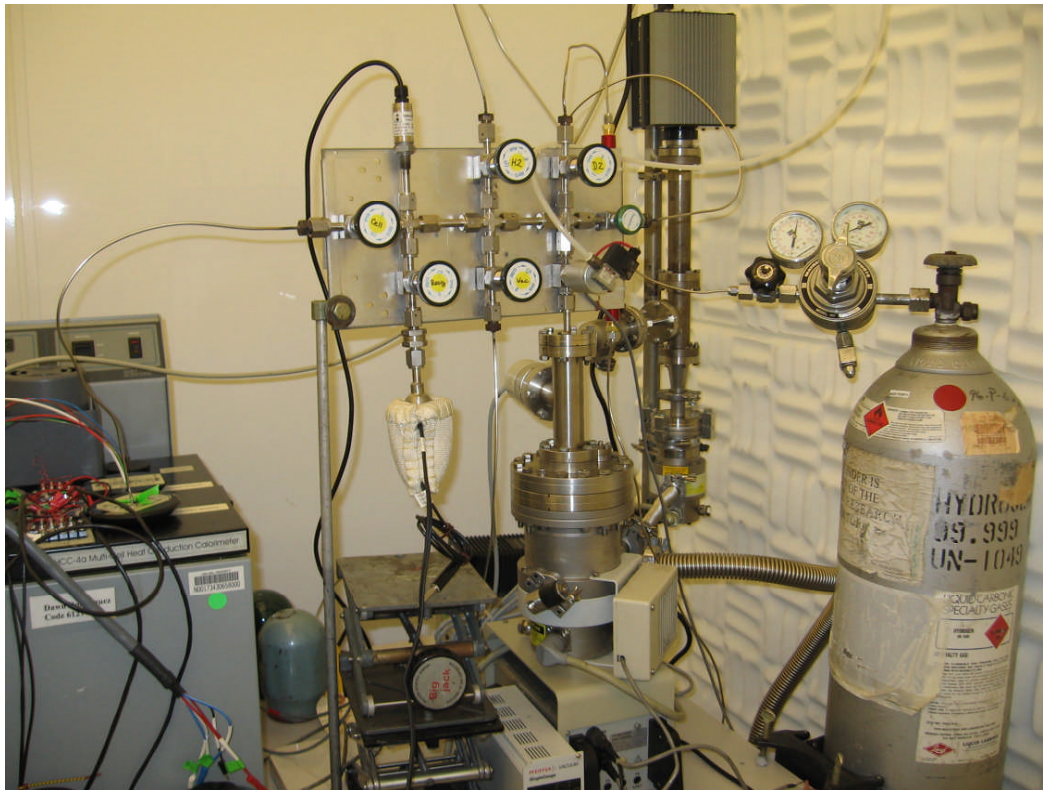
October 6, 2009



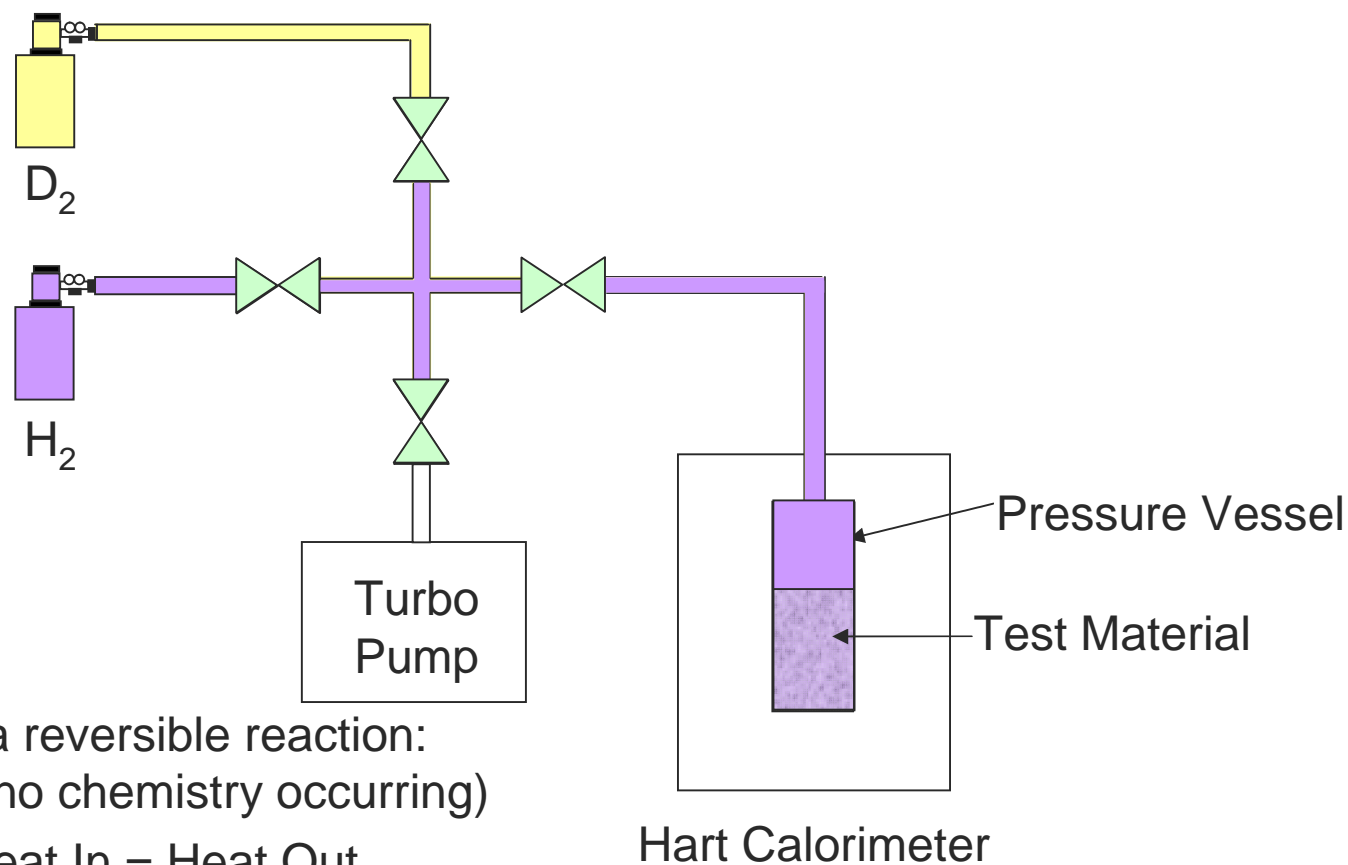
Basic gas loading experiment



Set-up in Hart calorimeter in Nanoscience Institute (NSI)



Basic gas loading experiment



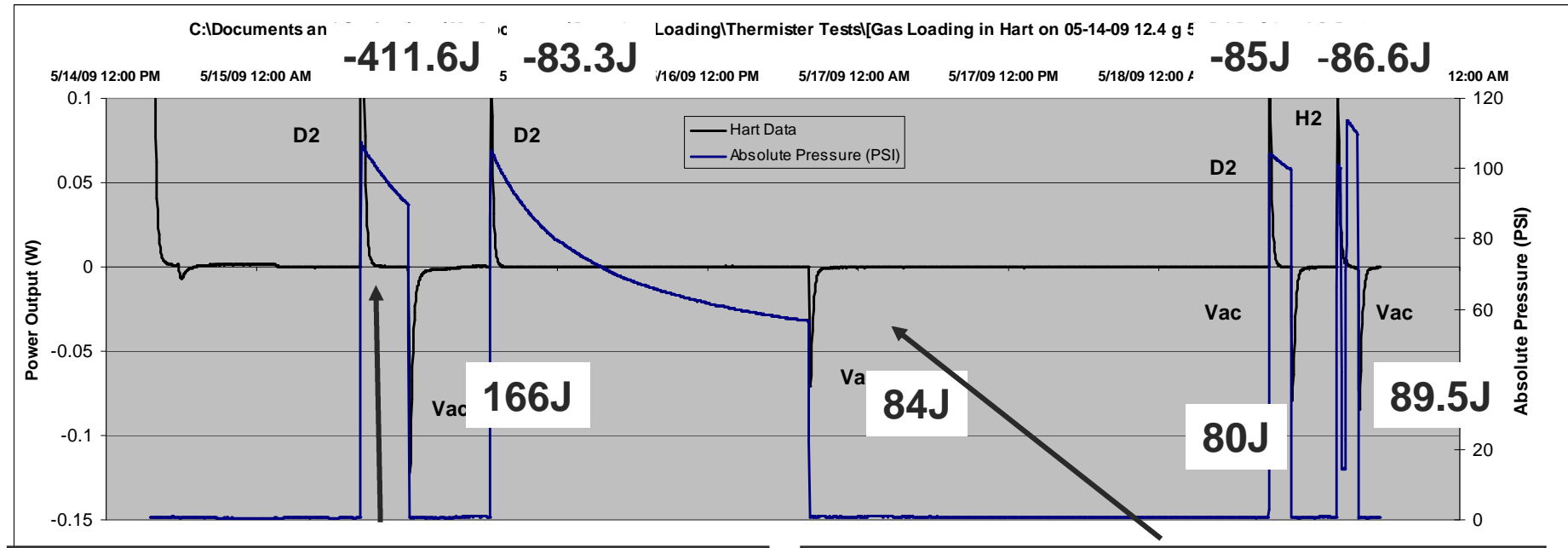
- For a reversible reaction:
(i.e. no chemistry occurring)

- Heat In = Heat Out
(Internal Control)

- $H_2 = D_2$
(External Control)



Loading using commercial 5% Pd/BaCO₃ - Control



Exothermic:

- Heat of Chemical Reaction
- Heat of D₂ uptake
- Heat of Pressurization (PV work)
- **Other Heat**

Endothermic:

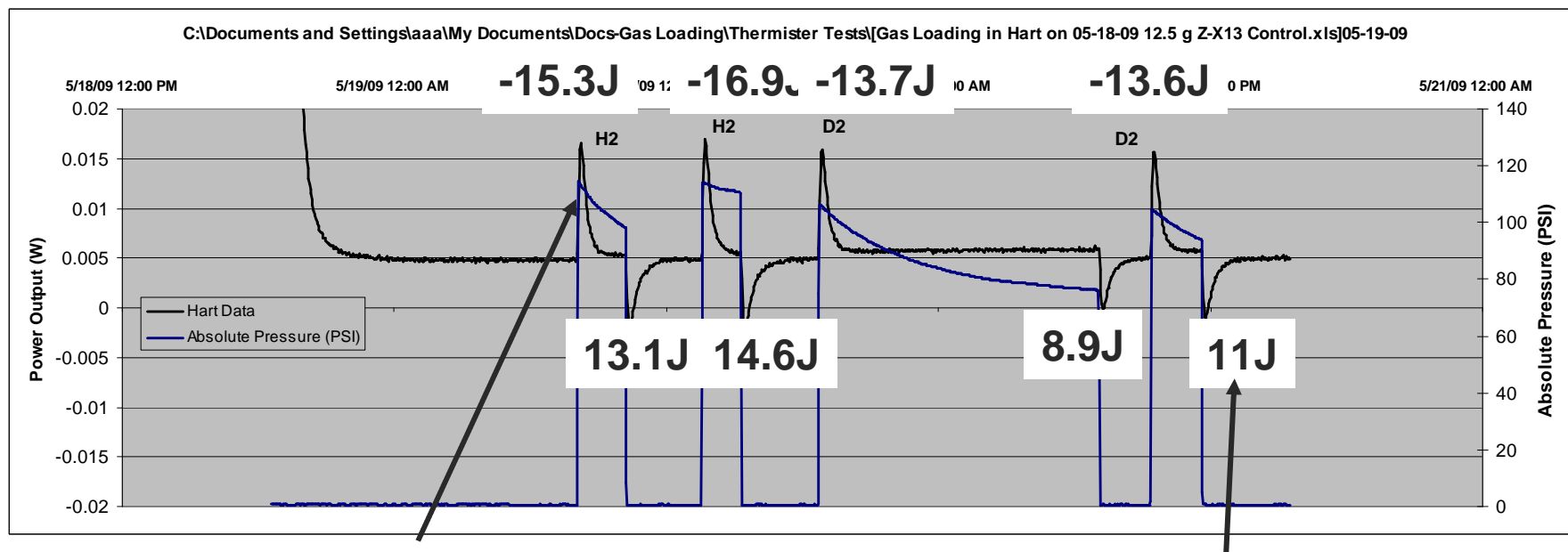
- Heat of D₂ release
- Heat of Evacuation (PV work)

■ Hart calorimeter - no excess heat is observed

- Heat In = Out; D₂ = H₂
- Internal and External Controls OK



Matrix without Pd - Control



Endothermic:

- Heat of Pressurization (PV work)

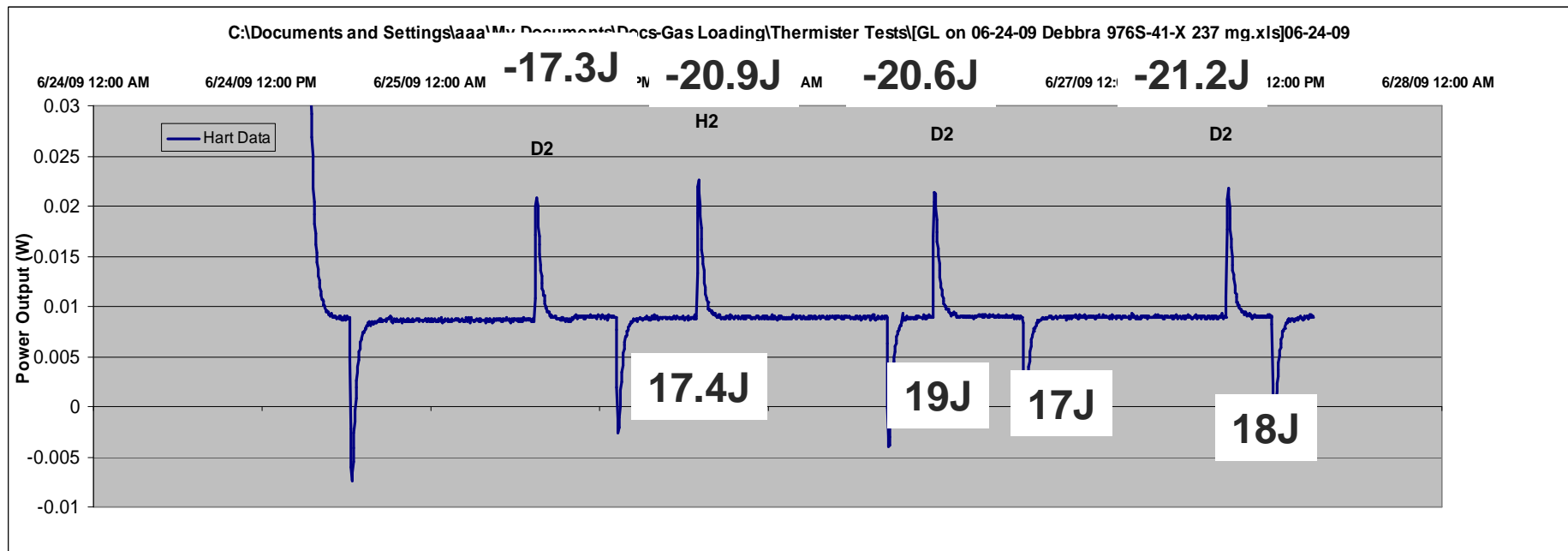
Endothermic:

- Heat of Evacuation (PV work)

- Hart calorimeter- no excess heat is observed
 - Heat In = Out; $D_2 = H_2$
 - Internal and External Controls OK



Loading using Pd/Graphite Paper



- Hart calorimeter- no excess heat is observed
 - Heat In = Out; $D_2 = H_2$
 - Internal and External Controls OK



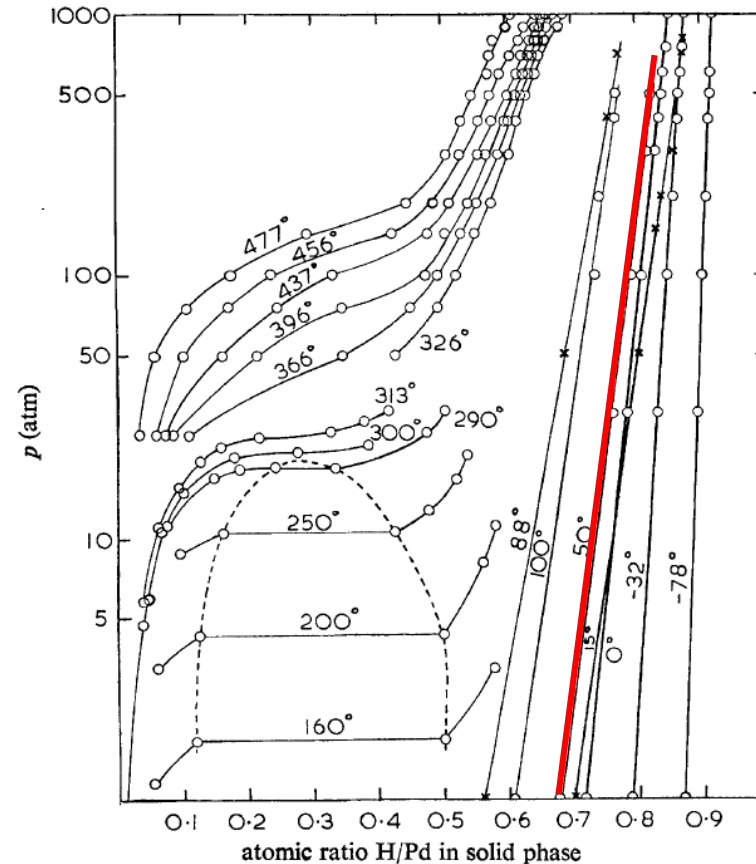
Tying Gas Loading to Electrolytic Loading

- Hydrogen diffusing into Palladium expands the lattice
- Because of this counter force, most Palladium takes-up 0.5-0.7 parts hydrogen
- Generally very high pressures are needed to get Palladium to uptake hydrogen beyond $\text{PdH}_{0.7}$ – est. 10^5 - 10^6 Atm to reach PdH_1
- In 1908, Paal and Gerum reported $\text{Pd:H}_{0.98}$ using Palladium black prepared with hydrazine (*Berichte*, **41** (1908) 818.)



Tying Gas Loading to Electrolytic Loading

- Hydrogen diffusivity
- Because of this, it can absorb hydrogen
- Generally very high hydrogen uptake
- In 1908, Paal and Lehn prepared palladium black



0.5-0.7

um to
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From: P.L. Levine and K.E. Weale, "The Palladium + Hydrogen Equilibrium at High Pressures and Temperatures", *Transactions of the Faraday Society*, **56** (1960) 357-362.



Literature Results – Particle Size

Certain size particles (< 2 nm) take-up hydrogen 1:1

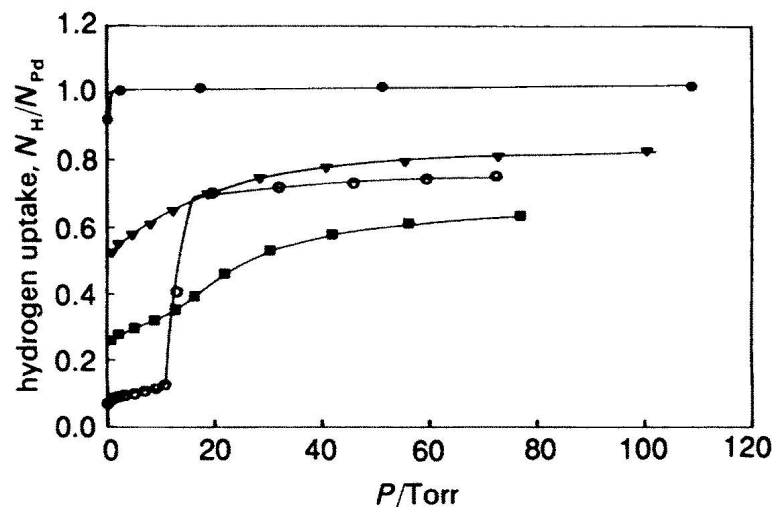


Fig. 1 Uptake isotherms for sorption of hydrogen on Pd/Al₂O₃ sample at 298 K: ○, 2% Pd(N)/Al₂O₃; ■, 2% Pd(Cl)/Al₂O₃; ▽, 1% Pd(Cl)/Al₂O₃; ●, 0.5% Pd(Cl)/Al₂O₃

From: Shu-Chin Chou, Shu-Hui Lin, and Chuin-Tih Yeh, "Isosteric Heat of Sorption of Dihydrogen on Alumina-supported Palladium", *J. Chem. Soc. Faraday Trans.*, **91**(1995) 949-951.

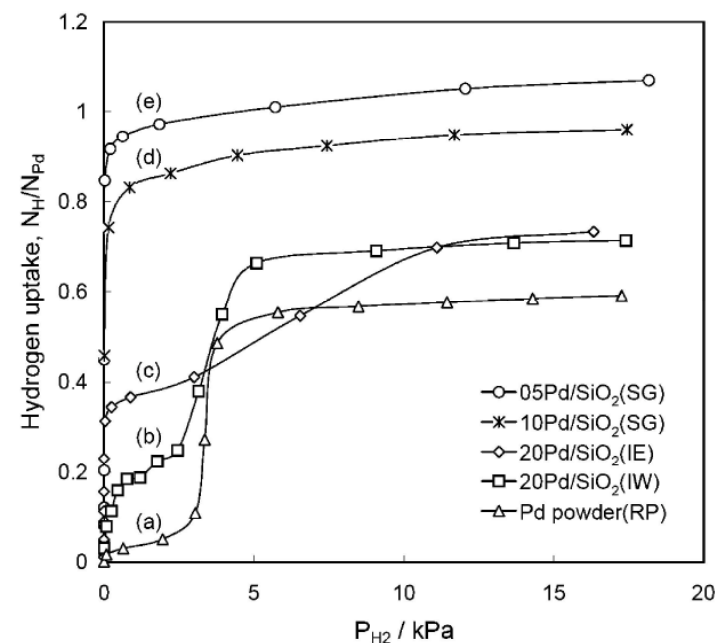


Figure 1. Adsorption isotherms of hydrogen uptake at 313 K for different palladium samples: (a) Pd powder (RP); (b) 20Pd/SiO₂ (IW); (c) 20Pd/SiO₂ (IE); (d) 10Pd/SiO₂ (SG); and (e) 05Pd/SiO₂ (SG).

From: Sheng-Yang Huang, Chin-Da Huang, Boh-Tze Chang, and Chuin-Tih Yeh, "Chemical Activity of Palladium Clusters: Sorption of Hydrogen", *J. Phys. Chem. B*, **110** (2006) 21783-21787.



Literature Results – Particle Size

Certain size particles (< 2 nm) take-up hydrogen 1:1

Preparation	Estimated Particle Size (nm)	Heat of Hydrogen Adsorption (kJ/mole)	Ratio H:Pd @ 0.2 Atm
Pd Powder	9	94	0.55
1.86% Pd/SiO ₂ (IW)	~4	92	0.68
10% Pd/SiO ₂ (SG)	1.1	131	0.9
5% Pd/SiO ₂ (SG)	1	183	1.05

Data from: Sheng-Yang Huang, Chin-Da Huang, Boh-Tze Chang, and Chuin-Tih Yeh, "Chemical Activity of Palladium Clusters: Sorption of Hydrogen", *J. Phys. Chem. B*, **110** (2006) 21783-21787.

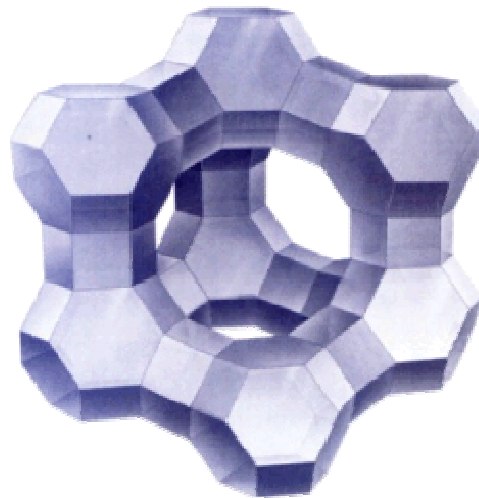


Hypothesis for Excess Heat

- Particle size is important
 - <2 nm allows rapid loading to Pd:D 1:1
 - Need isolation to keep from sintering – used Zeolites



Type A Zeolite



Type X Zeolite

From: http://www.molecularsieve.org/Zeolite_Molecular_Sieve.htm

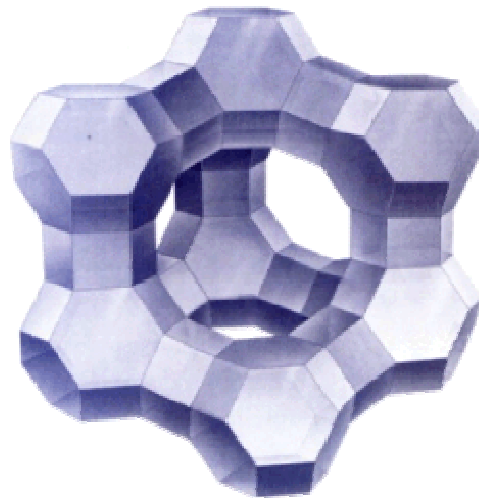


Hypothesis for Excess Heat

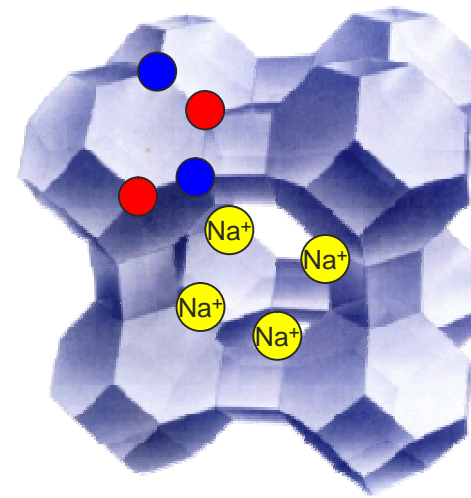
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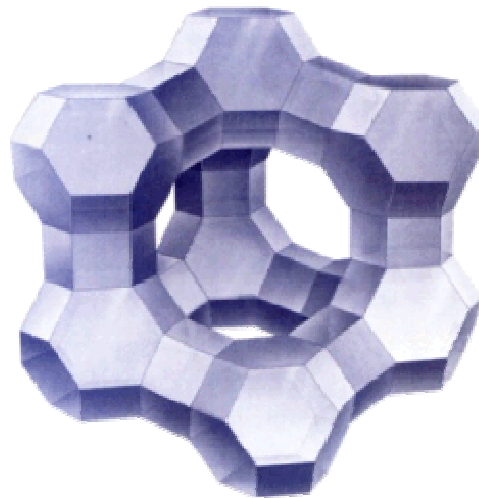


Hypothesis for Excess Heat

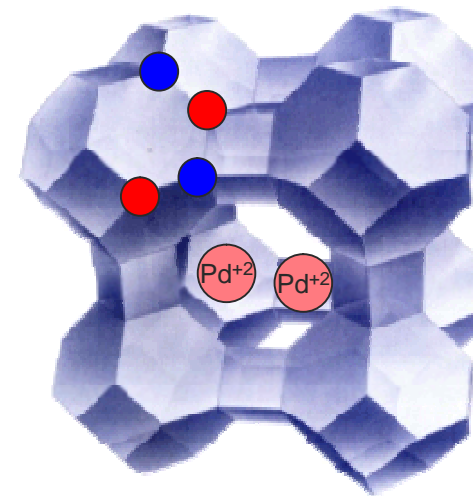
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Type A Zeolite



Type X Zeolite

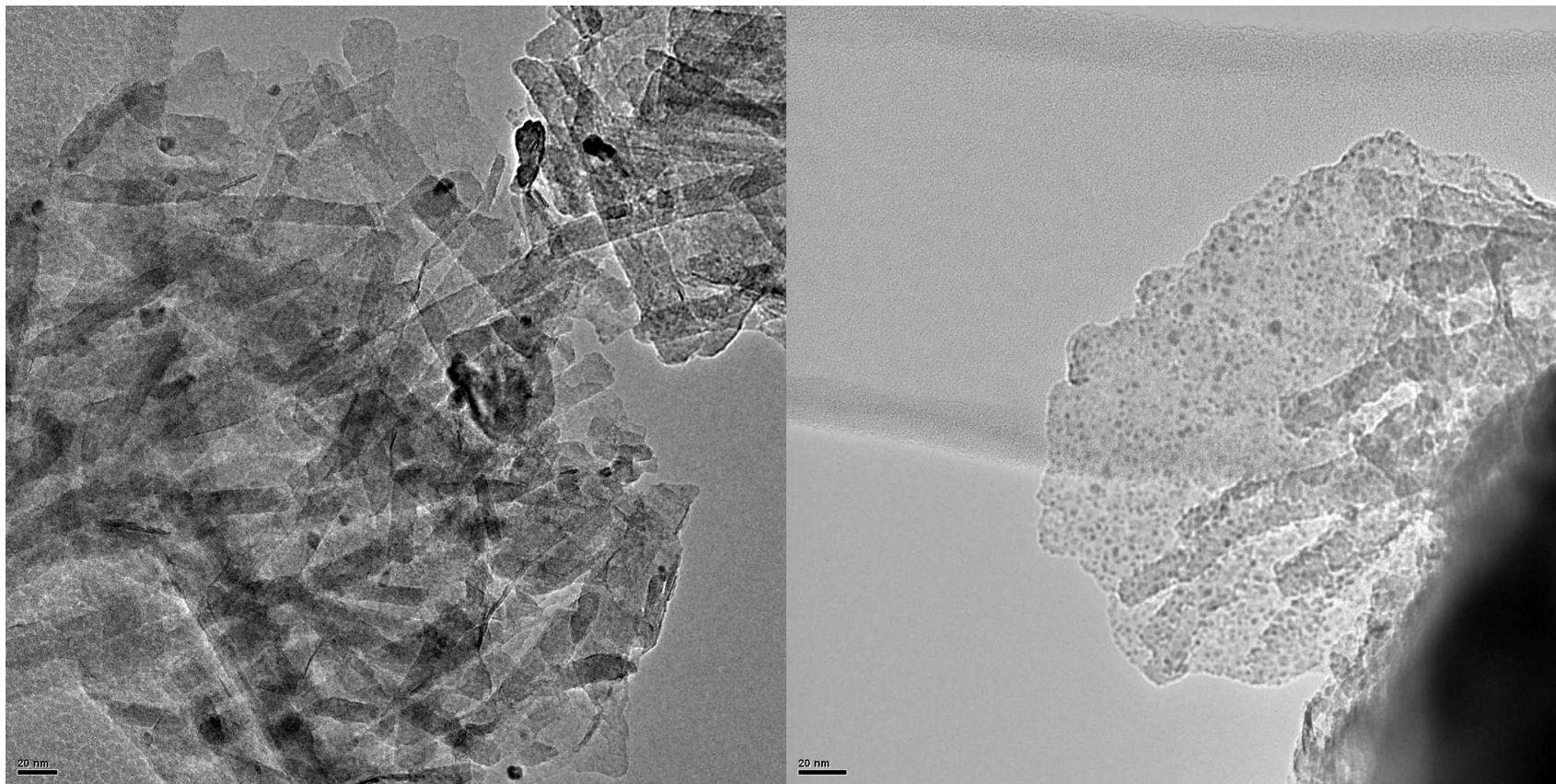


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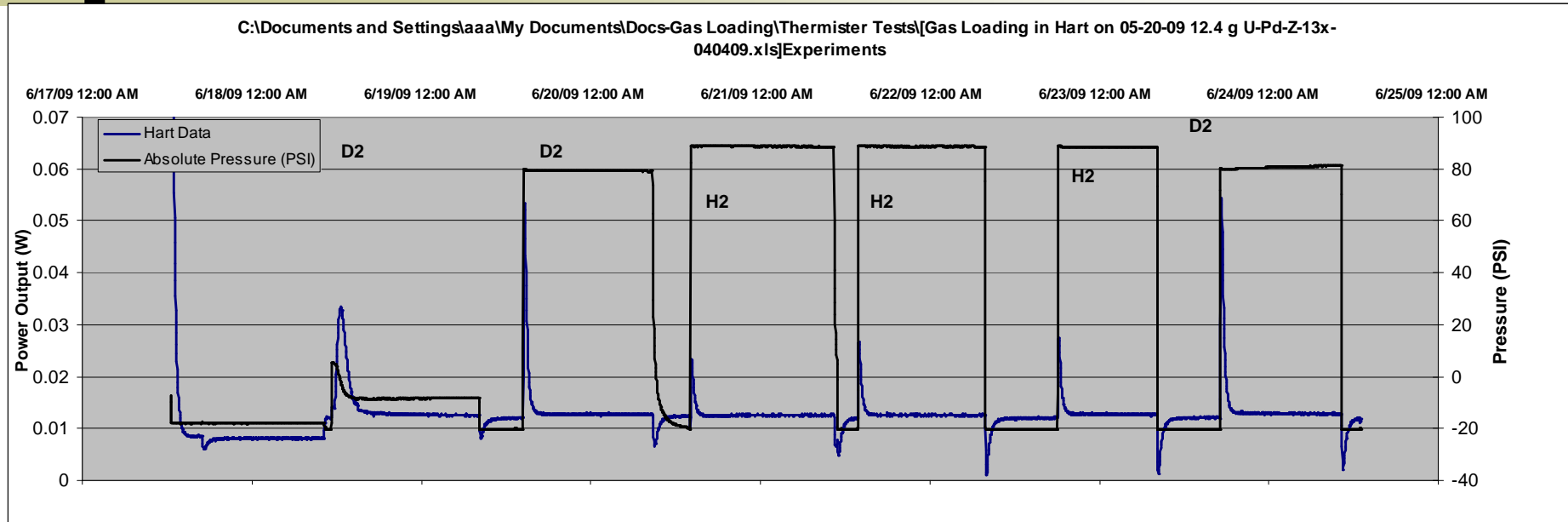


Characterization of Particle Size

- Not readily visible via TEM due to matrix



Matrix with Pd – 0.5% Pd on NRL Matrix

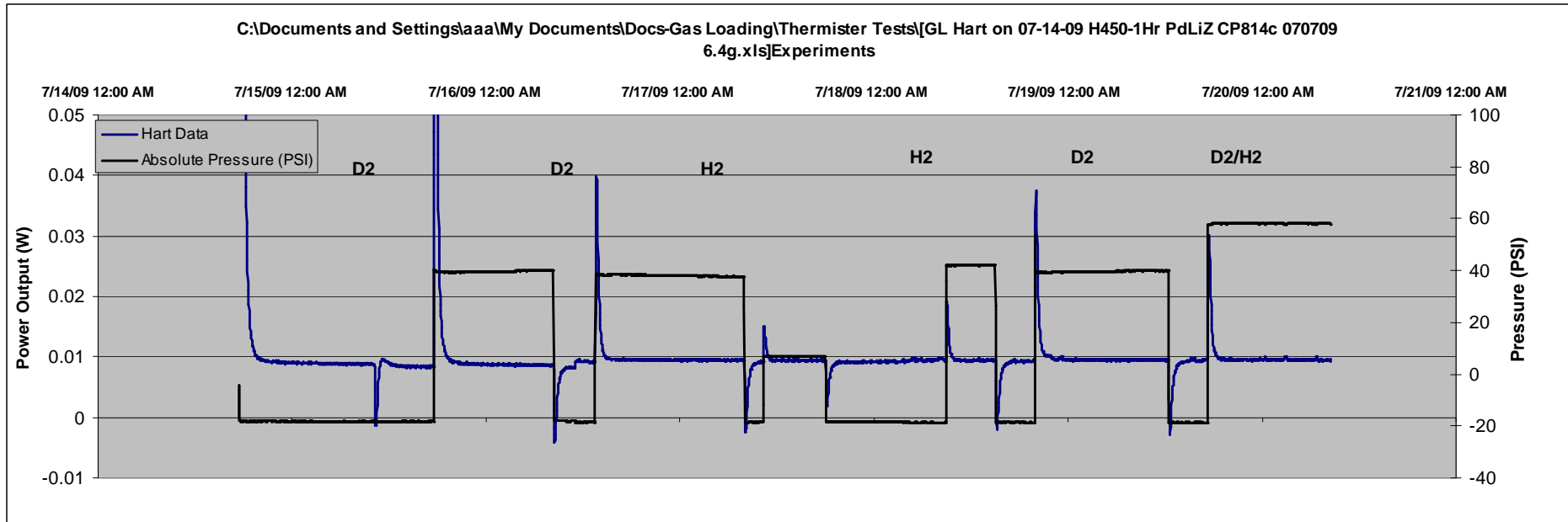


Heat from 2 nd D2 pressurization	-77.6
Endothermic heat from 2 nd D2 evacuation	19.7
Heat 3 rd H2 pressurization	-26
Endothermic heat from 3 rd H2 evacuation	31.5

- Hart calorimeter- no excess heat is observed
 - Heat In \neq Out; D₂ \neq H₂
 - Internal and External Controls NOT OK for D₂ but OK for H₂



Matrix with Pd – 1% Pd on NRL Matrix

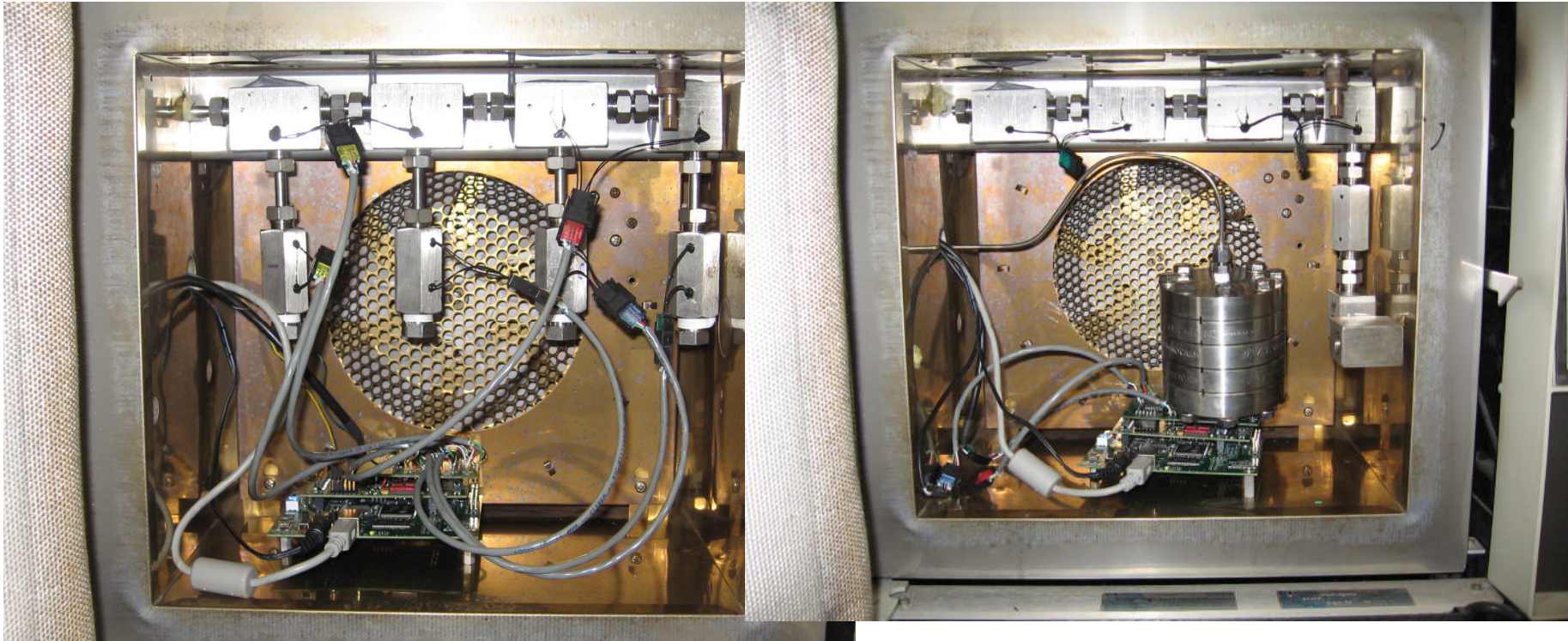


Measured Heat Deuterium pressurization at 17:35 on 07/15/09	-154.7
Measured Heat Deuterium removal at 8:21 on 07/16/09	22.4
Measured Heat of Deuterium pressurization at 13:48 on 07/16/09	-43.1
Measured Heat of Deuterium removal at 7:58 on 07/17/09	20.9
Measured Heat Hydrogen pressurization at 10:17 on 07/17/09	-7.2
Measured Heat of Hydrogen removal at 18:02 on 07/17/09	13.5
Measured Heat Hydrogen pressurization at 8:54 on 07/18/09	-12.9
Measured Heat Hydrogen removal at 15:04 on 07/18/09	20
Measured Heat Deuterium pressurization at 19:54 on 07/18/09	-46.5
Measured Heat Deuterium removal at 12:29 on 07/19/09	20.1
Measured Heat Hydrogen/Deuterium Mix pressurization at 17:54 on 07/19/09	-28.8

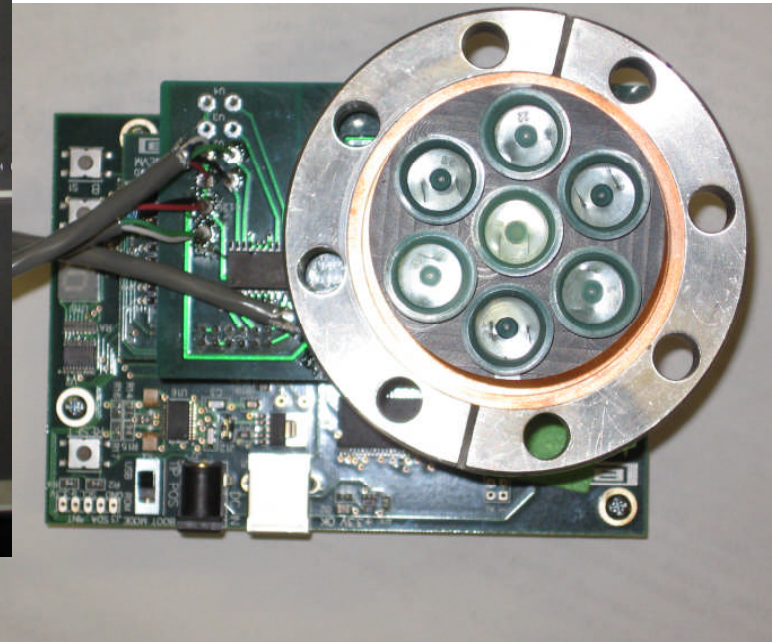
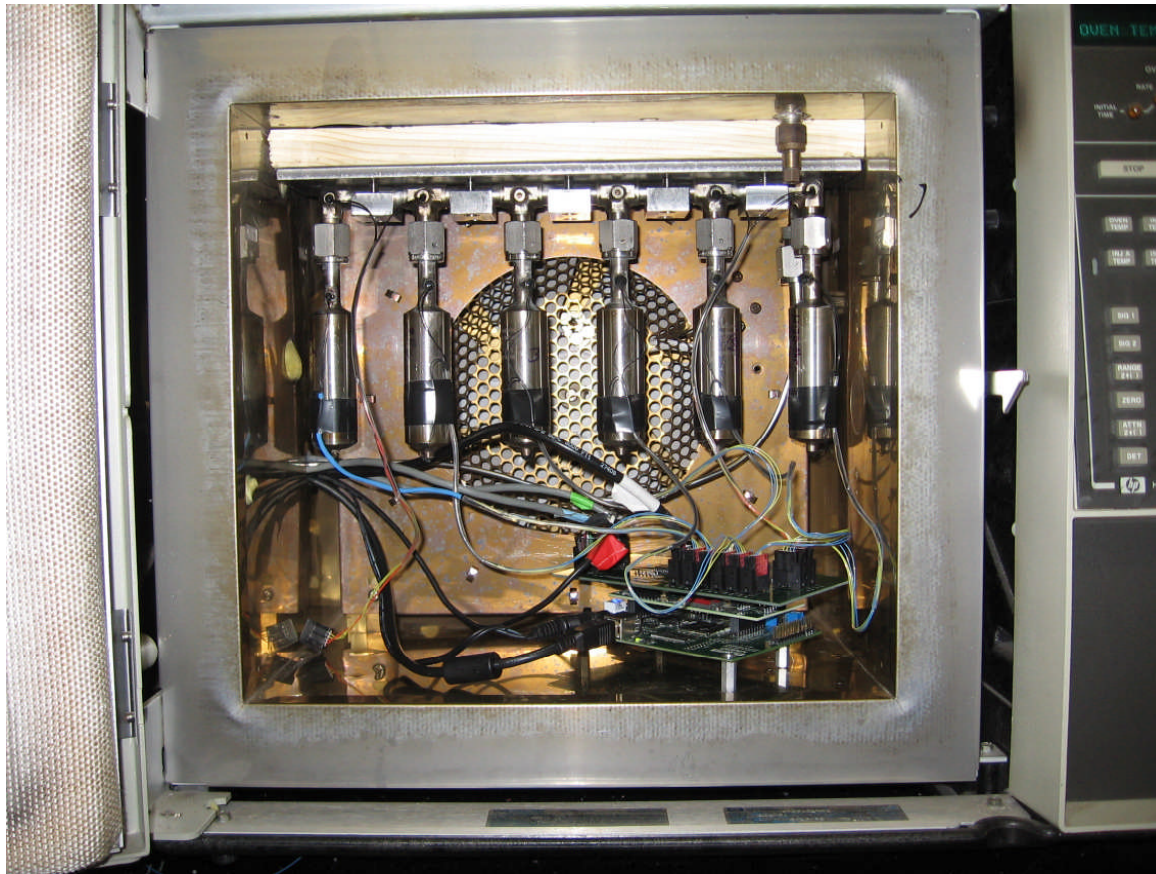
- Hart calorimeter- no excess heat is observed
 - Heat In \neq Out; $D_2 \neq H_2$
 - Internal and External Controls NOT OK for D_2 but OK for H_2



Examples of set-up in GC Oven

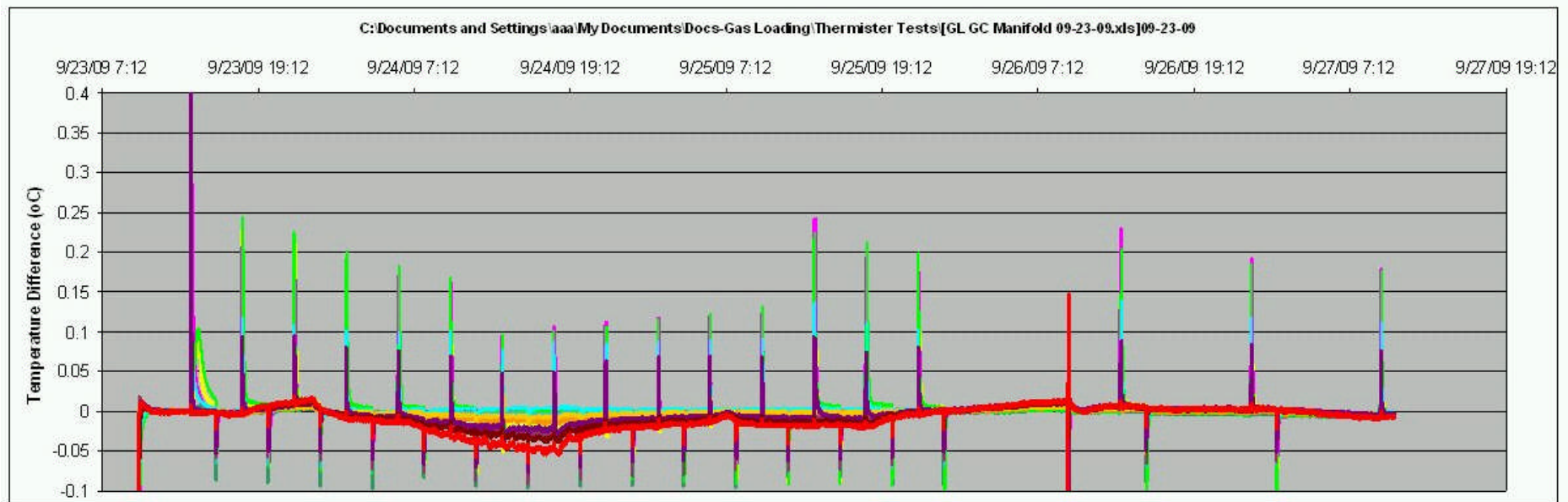


Examples of set-up in GC Oven



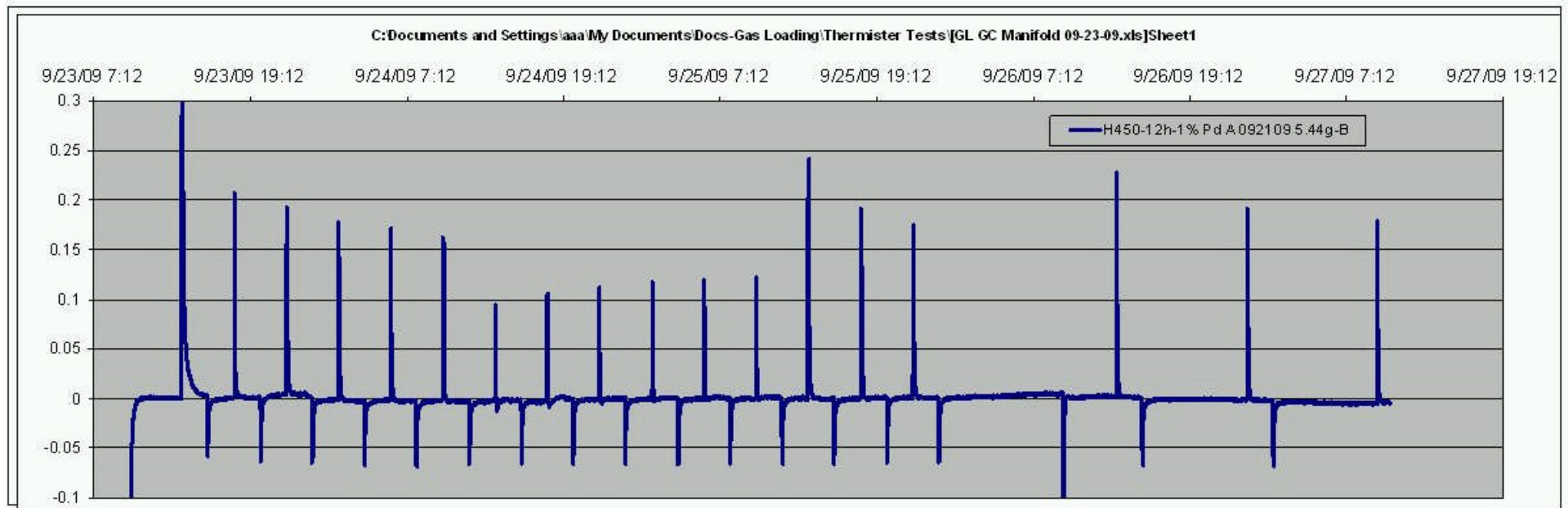
Closer Look of a Run in the GC

Various matrices



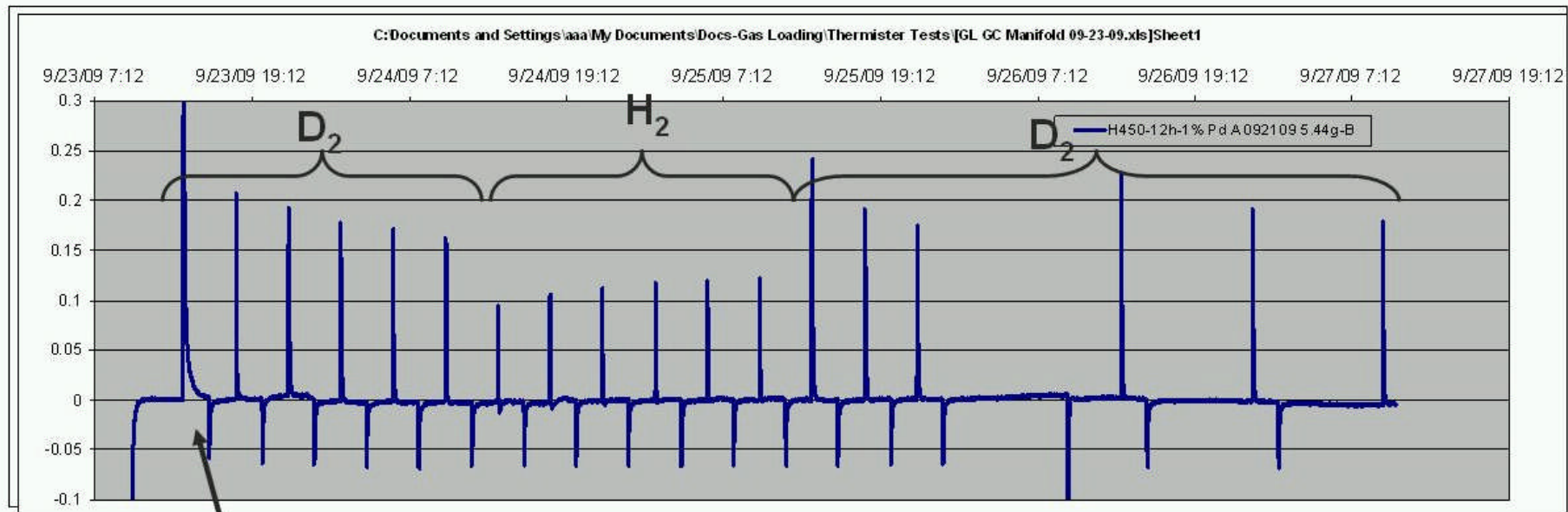
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Closer Look of a Run in the GC

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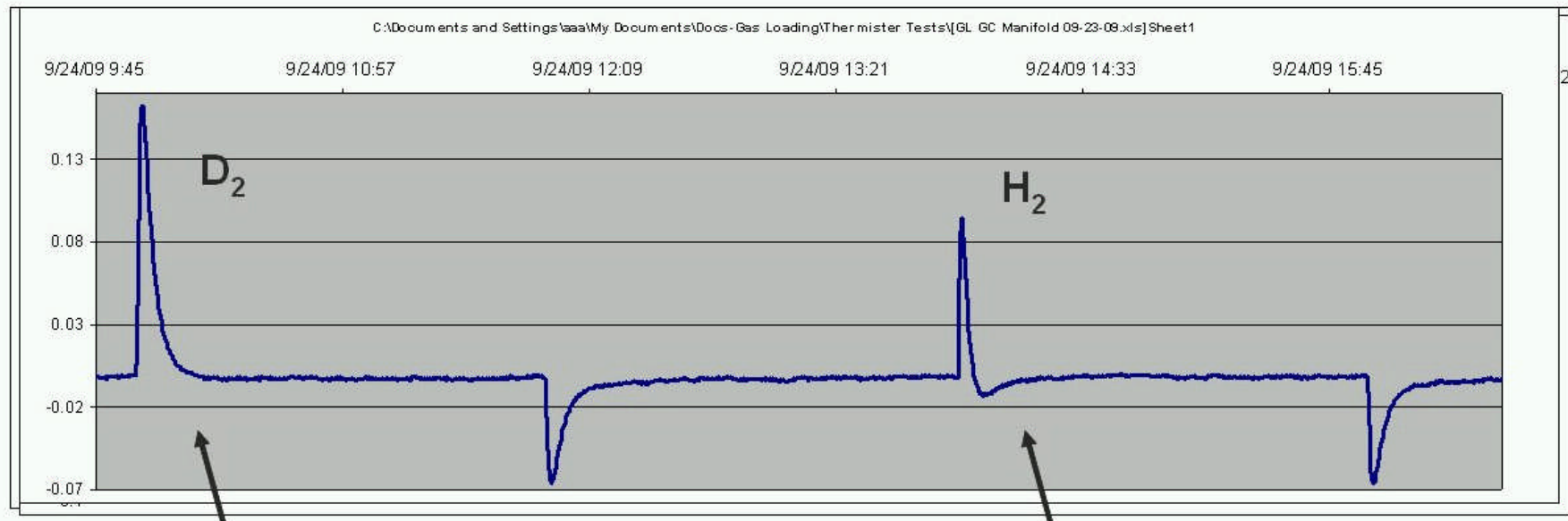


Heat of Chemical Reduction
Heat of D_2 uptake
Heat of Pressurization (PV work)
Other Heat



Closer Look of a Run in the GC

Various matrices



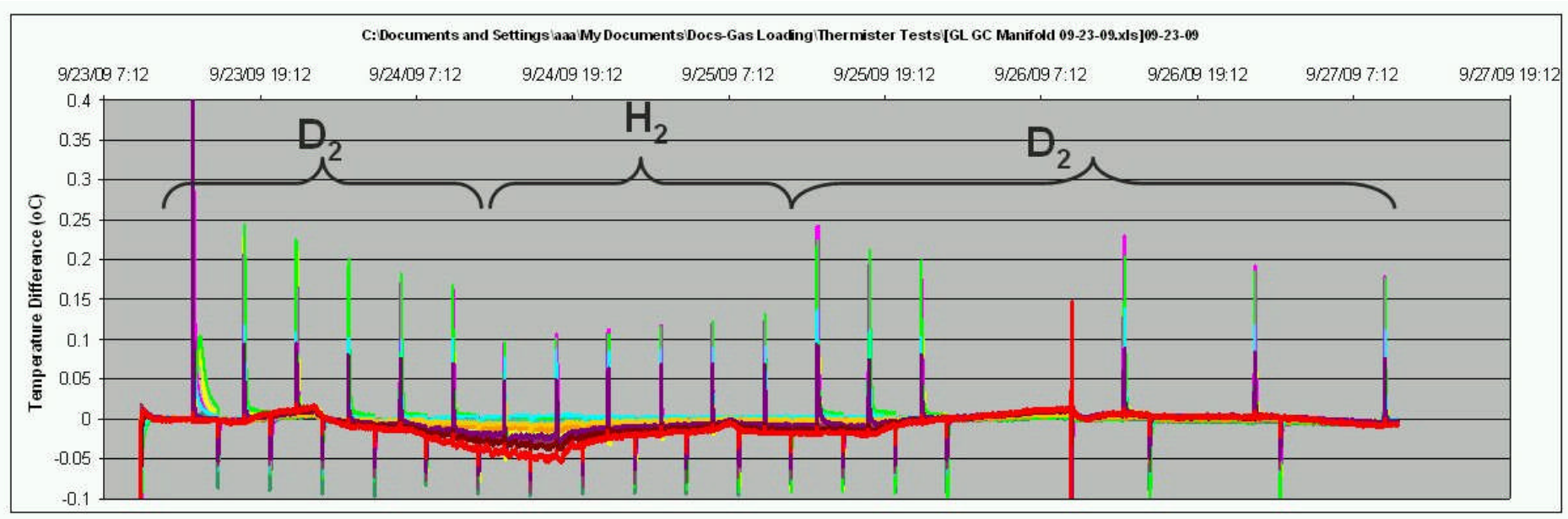
■ Results from GC:

- Excess other heat is observed
- Heat In \neq Out for D_2 but equal for H_2 , $H_2 \neq D_2$



Closer Look of a Run in the GC

Various matrices



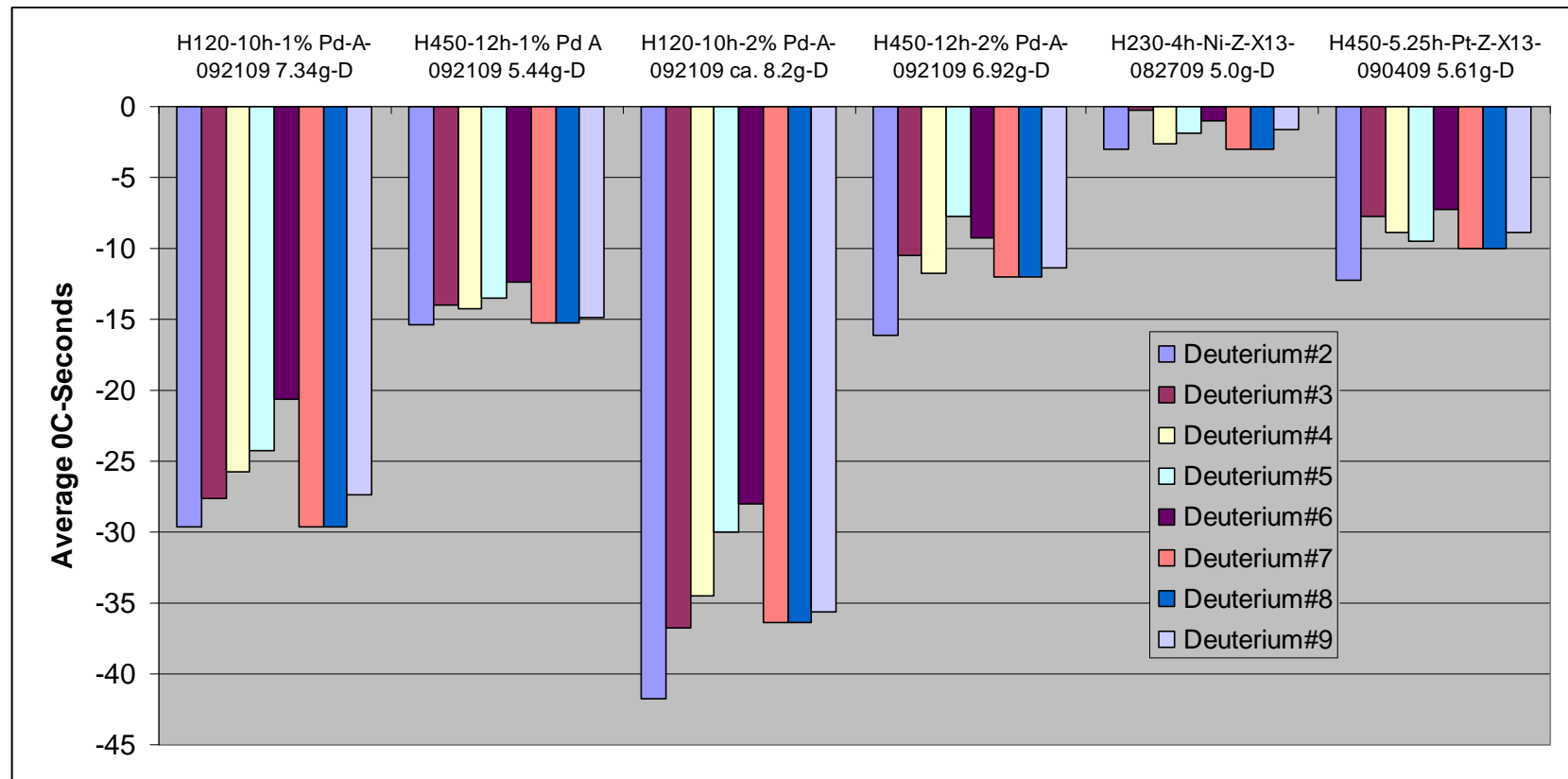
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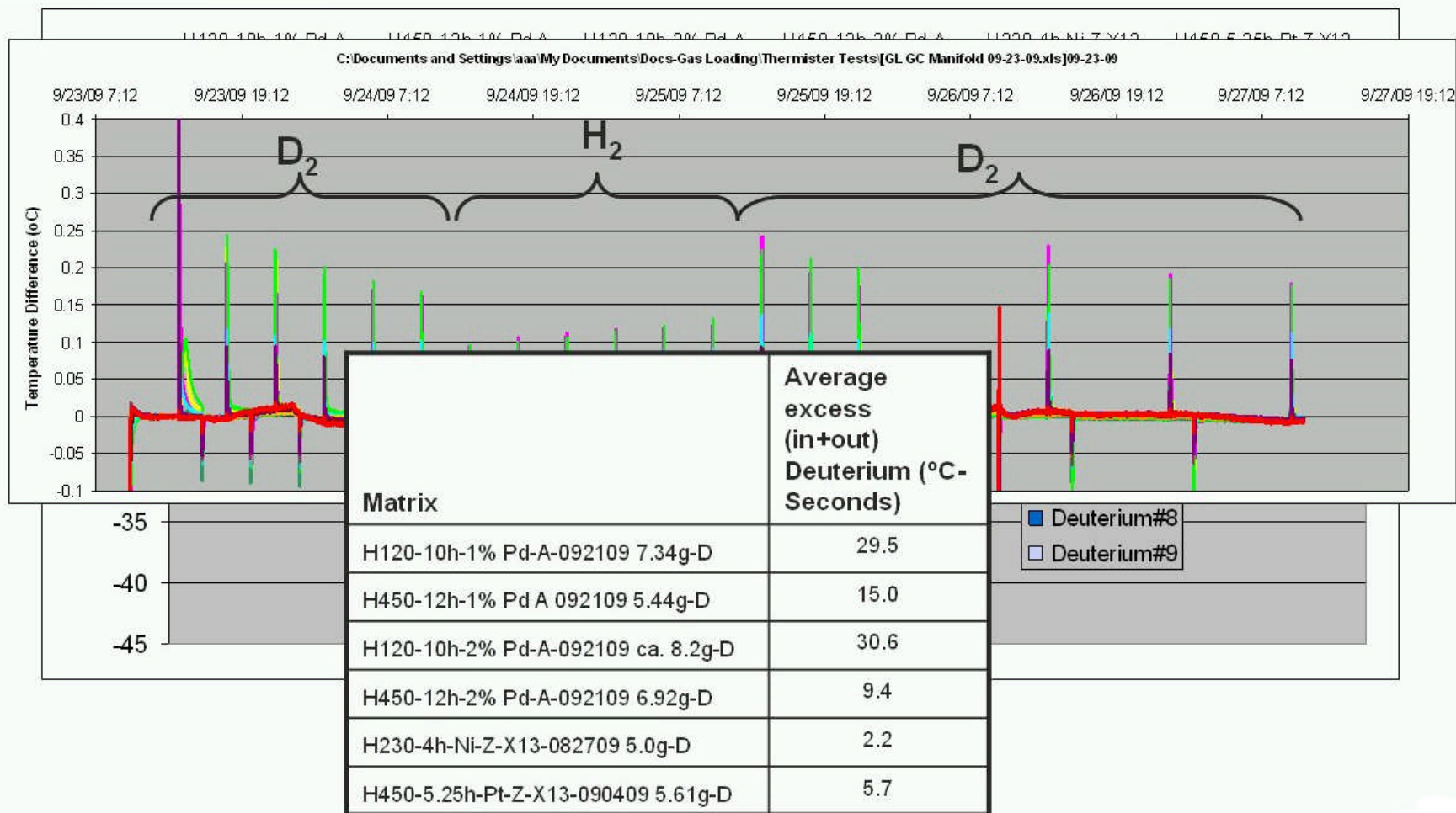
Closer Look of a Run in the GC

Reproducibility of Depressurizations



Closer Look of a Run in the GC

Reproducibility of Depressurizations





But
Is there an explanation for the
“other” heat repeatedly observed?



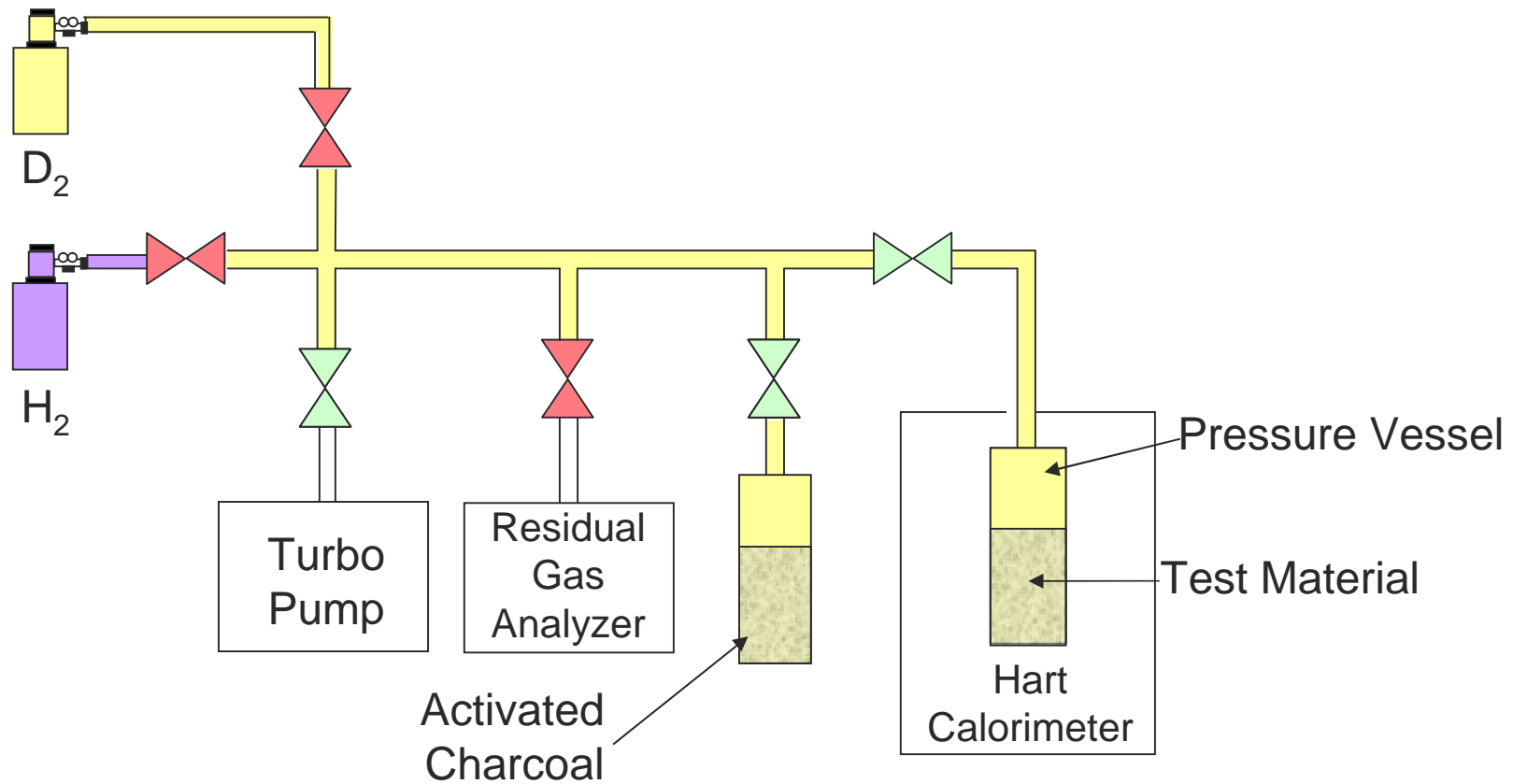
Possible Sources of Heat #1

Chemical energy in the deuterium but not in the hydrogen

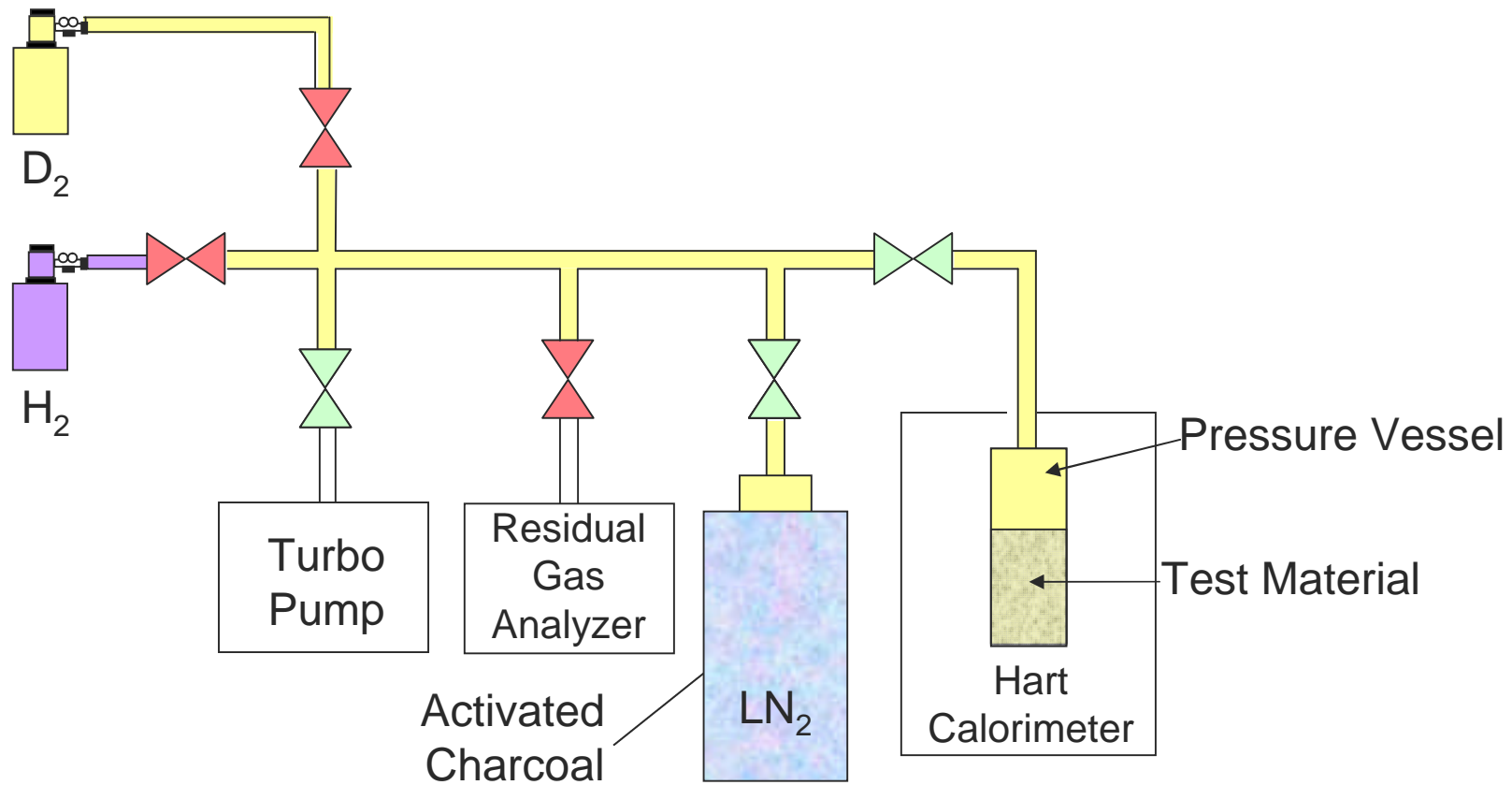
- Example O₂
- Argue against:
 - Increase pressure – no proportionate increase in heat
 - Other palladium particles show no effect
 - Hypothetically could be dependant on particle size but this is not supported chemically



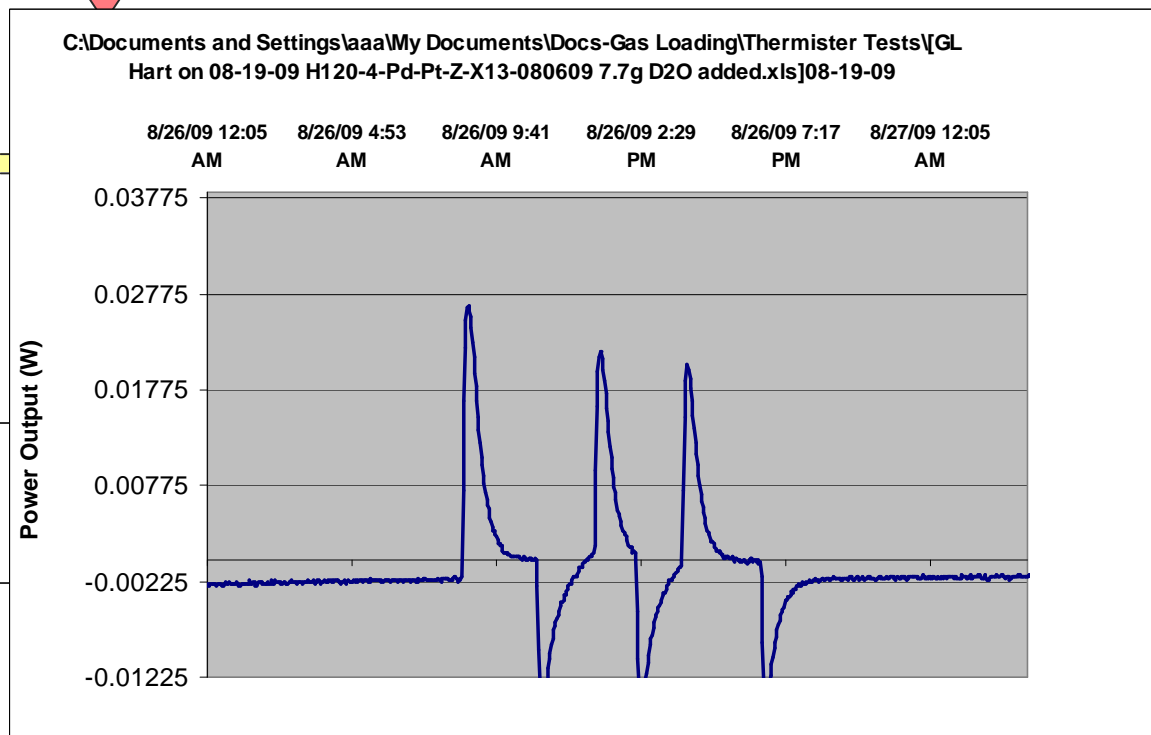
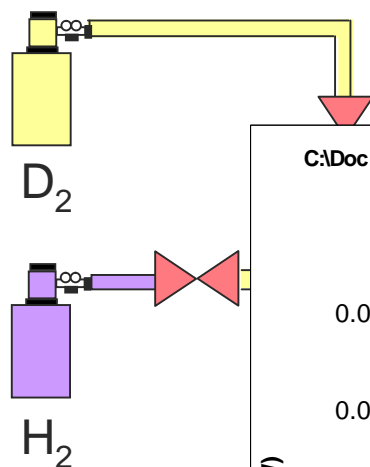
Recycling of Fill Gas



Recycling of Fill Gas



Recycling of Fill Gas



re Vessel

terial

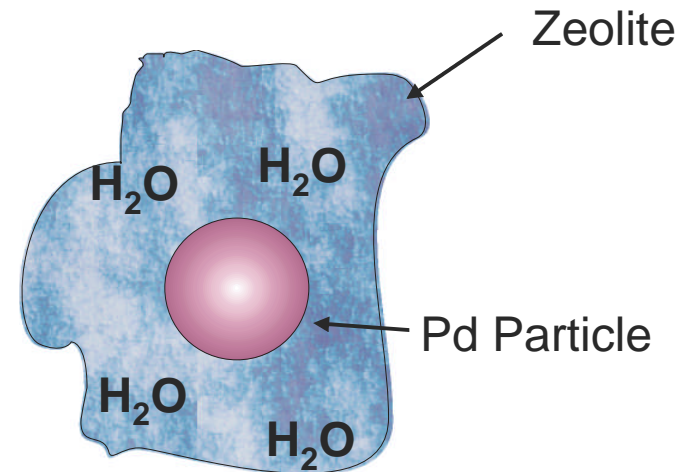
Initial Pressurization to 125 PSI	-48.9J
Recycle #1 to 122 PSI	-38.7J
Recycle #2 to 120 PSI	-31.3J



Possible Sources of Heat #2

Mechanism consistent with some observations

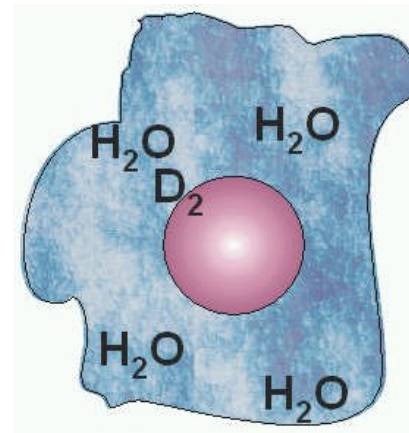
- Chemical exchange of D_2 with H_2O



Possible Sources of Heat #2

Mechanism consistent with some observations

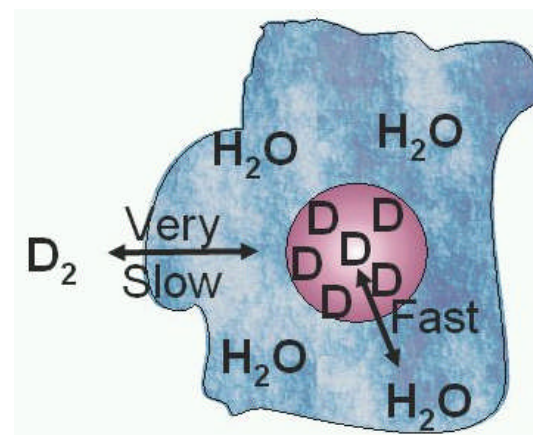
- Chemical exchange of D_2 with H_2O



Possible Sources of Heat #2

Mechanism consistent with some observations

- Chemical exchange of D_2 with H_2O



- Heat produced but no signature of H-D exchange in gas phase
 - Signature only upon removal of residual gas in Pd particle
 - Observed – D_2 gives some HD and H_2 gives some HD
- Calculations suggest for the amount of Pd present and going to D_2O (worse case)
 - Ignoring the spillover effect, only 7J heat available
- Using D_2O saturated Zeolite should see less heat with H_2
 - Observed – can be endothermic

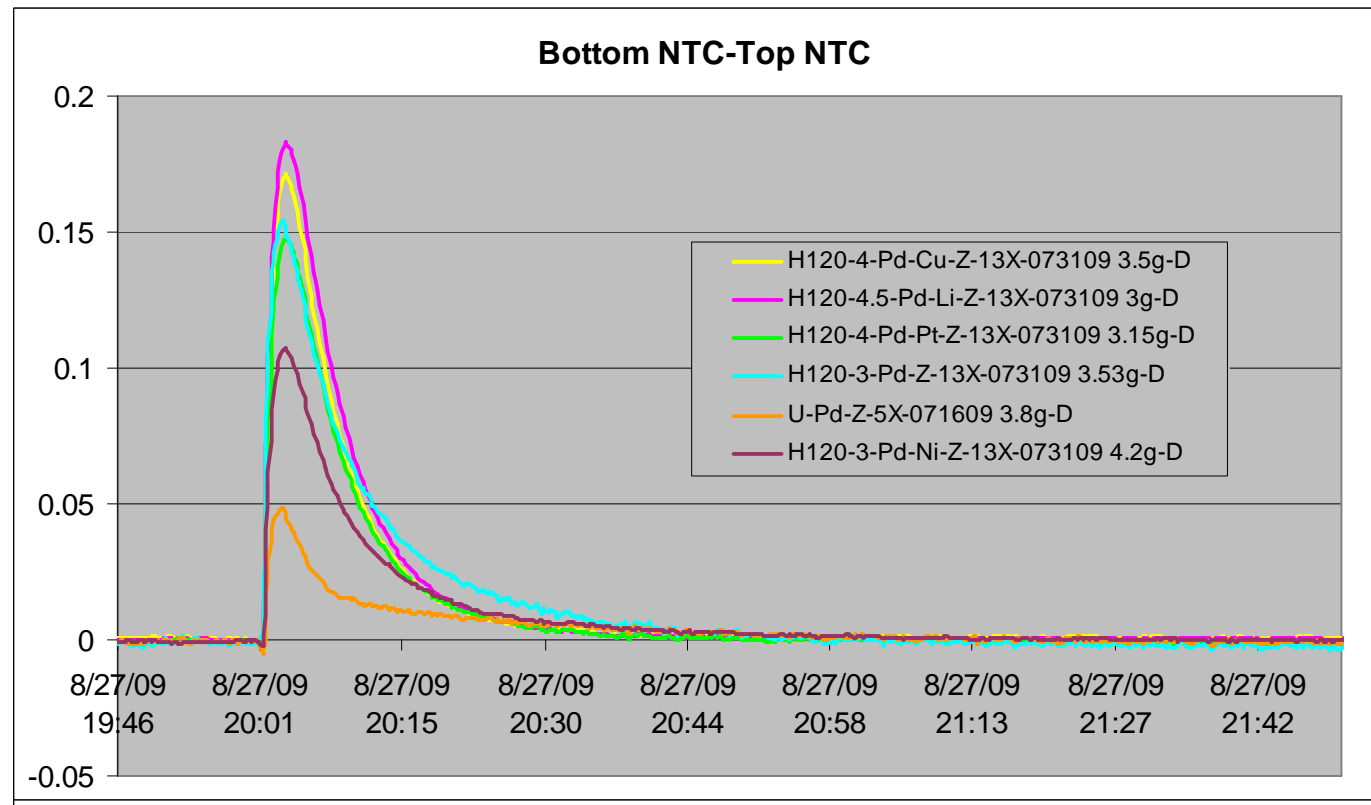


Possible Sources of Heat #2

Some observations

Chemical exchange of D₂ with H₂O

- D₂ + H₂O → HD + HDO -4.13 kJ/mole
- D₂ + H₂O → HD + D₂O -8.71 kJ/mole



GL GC Manifold 08-25-09 at 70C then 29C.xls

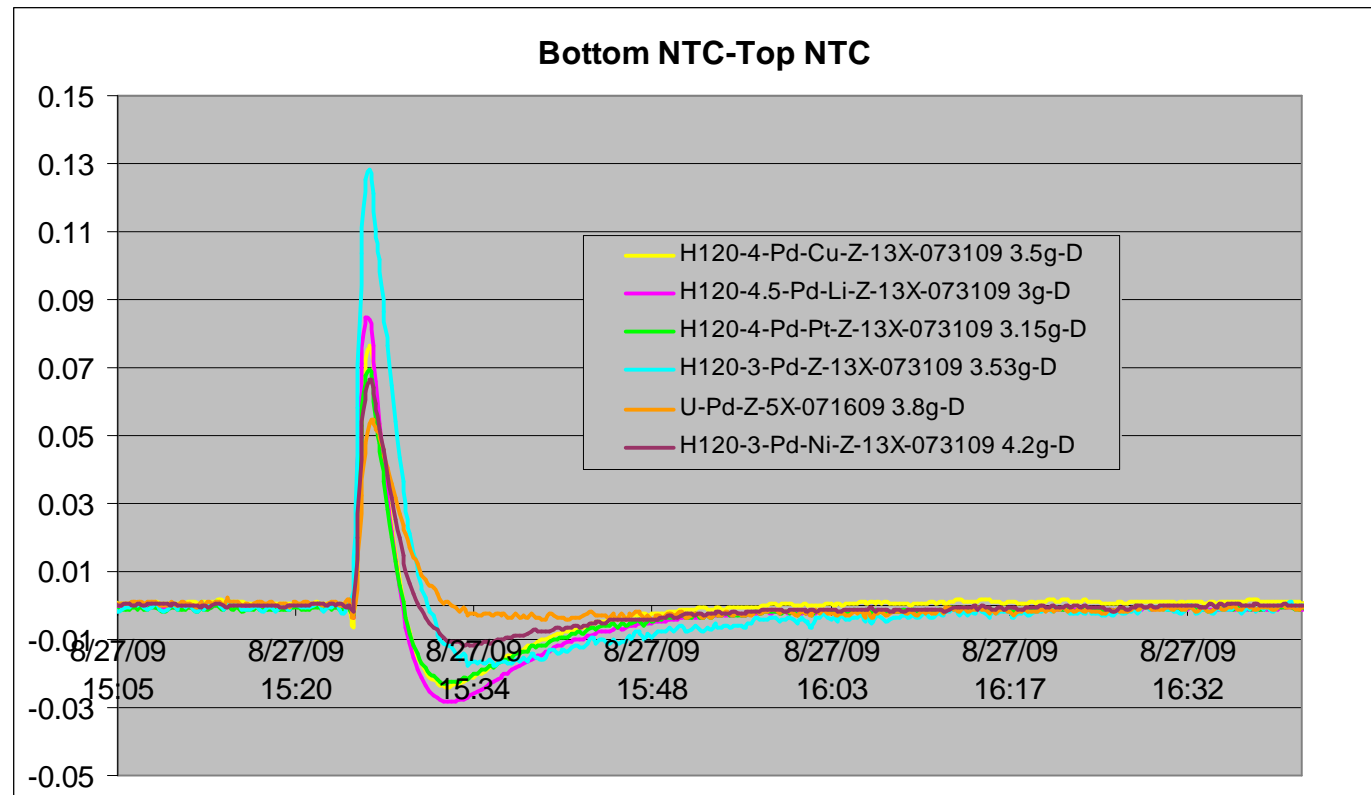
ICCF-15



Possible Sources of Heat #2

Some observations

- Chemical exchange of D₂ with H₂O
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GL GC Manifold 08-25-09 at 70C then 29C.xls



Why Consider D-H Exchange as a Possible Source of Heat?

- Chemical exchange of D₂ with H₂O
 - $D_2 + H_2O \rightarrow HD + HDO$ -4.13 kJ/mole
 - $D_2 + H_2O \rightarrow HD + D_2O$ -8.71 kJ/mole
- Can explain:
 - Internal controls – In <> Out
 - Why D₂ (exothermic) <> H₂ (endothermic)
- Effect still present with some “dry” matrices
 - Make water in initial reactions
 - $PdO + D_2 \rightarrow PdDx + D_2O$
 - $D_2O + Matrix \rightarrow Hydrated\ Matrix$
- Many pressurization/depressurization cycles decreases contribution



Conclusions

- Preparing Pd nano-particles inside Zeolites is reproducible and easy
 - Zeolites appear to provide Pd particles of the correct dimensions
- Interesting heat pattern was measured by two independent principles
 - Many iterations with unexpected heat present
 - Heat in presence of Deuterium but not Hydrogen
- Chemical effect due to Hydrogen-Deuterium exchange may account for some of the anomalous heat
 - Hydrogen may not be best control
 - Requires all the species present – Zeolite, Pd catalyst, water. D_2/H_2
 - But simple calculations do not account for all the heat observed
- Although chemistry of Zeolites is complex, because of the ease of preparation, further exploration of these systems is warranted



Acknowledgements

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Questions

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