EXCESS HEAT IN ELECTROLYSIS EXPERIMENTS AT ENERGETICS TECHNOLOGIES

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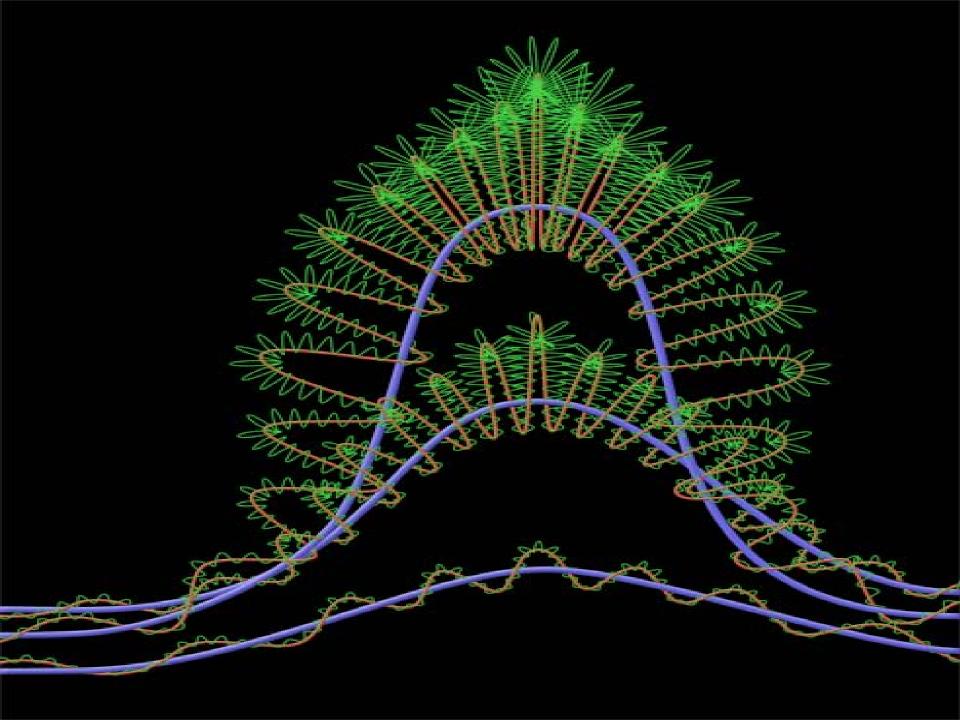
- 1. Presentation objective
- 2. SuperWaves[©] used for driving the cell
- 3. Review of glow discharge experiments
- 4. Description of ET electrolytic cells
- 5. Pd Cathode and its pretreatment
- 6. Excess heat obtained
- 7. Excess tritium
- 8. Material analysis

PRESENTATION OBJECTIVES

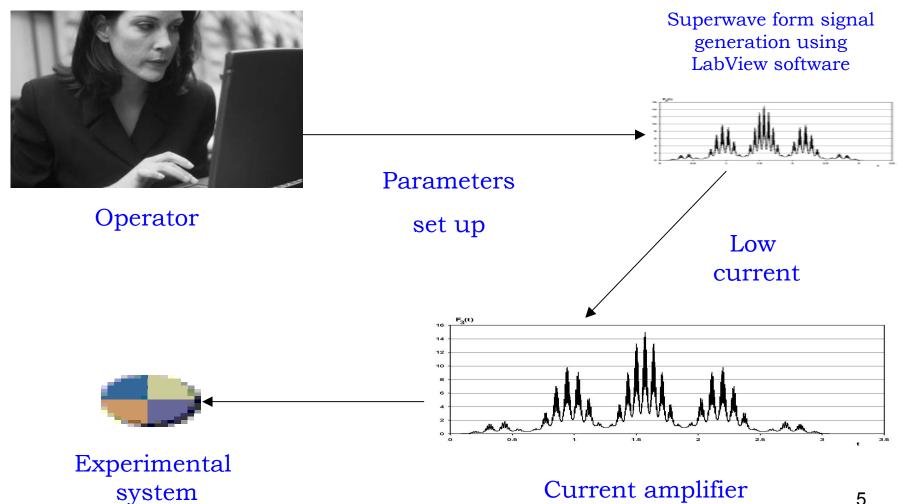
- 1. Review of SuperWave[©] principles
- 2. Review of Glow Discharge Experiments
- 3. Description of 3 ET electrolytic cell experiments that resulted in significant excess heat generation:

Experiment #	56	64a	64b
Cycle № Loading time (s) Excess heat (EH) (%) Duration of EH (h) Excess energy (EE) (MJ) Specific* EH (W/g Pd)	4 80 80 300 3.1 11	1 5 2500 17 1.1 71	2 16 1500 80 4.6 62
Specific* EE (KeV/Pd atom)	13.5	4.8	20 (24.8)

 $^{^*}$ - pertaining to effective part of cathode ($6 \times 0.7 \text{ cm}$)

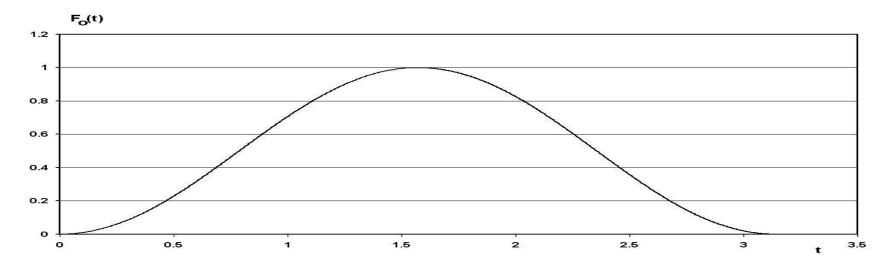


EC are driving by SuperWaves

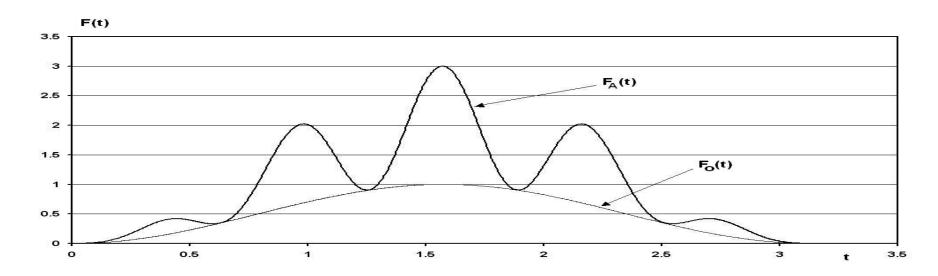


SuperWaves formation principles

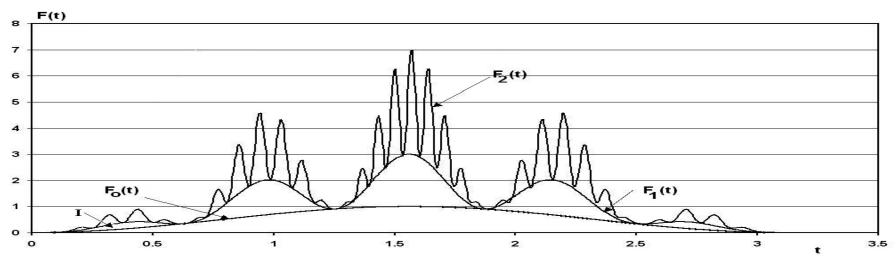
$$F_0(t) = A_0 \sin^2(\omega_0 t)$$



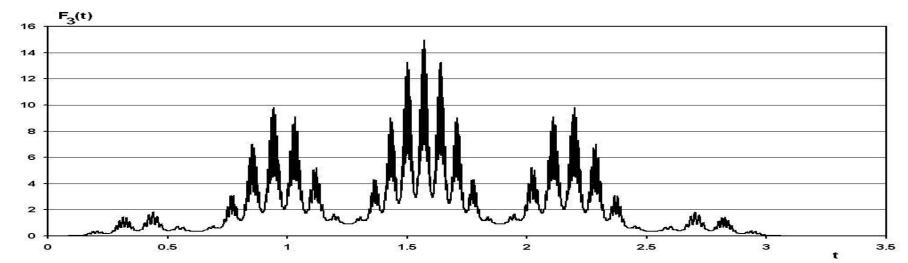
$$F_1(t) = A_0 \sin^2(\omega_0 t) (1 + A_1 \sin^2(\omega_1 t))$$



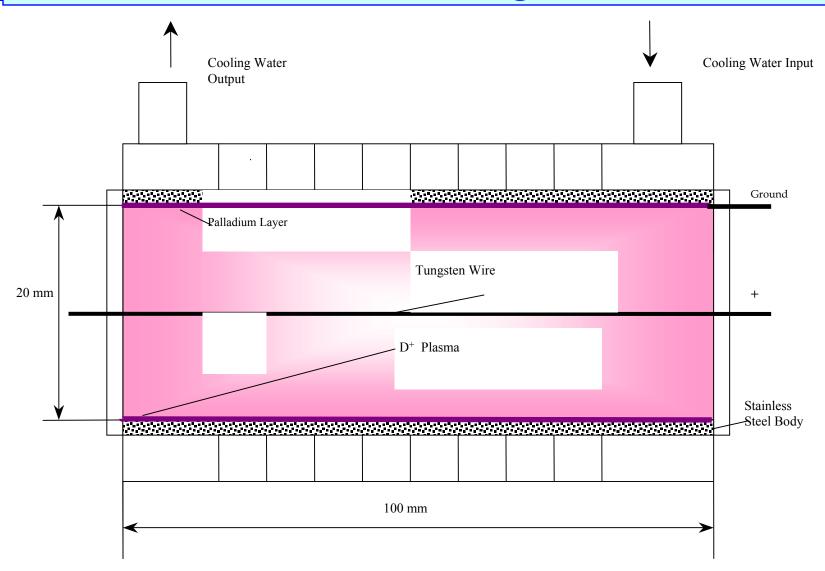
$$F_{2}(t) = A_{0} \sin^{2}(\omega_{0}t)[1 + A_{1} \sin^{2}(\omega_{1}t)(1 + A_{2} \sin^{2}(\omega_{1}t))]$$



 $F_3(t) = A_0 \sin^2(\omega_0 t)[1 + A_1 \sin^2(\omega_1 t)(1 + A_2 \sin^2(\omega_2 t)(1 + A_3 \sin^2(\omega_3 t)))]$



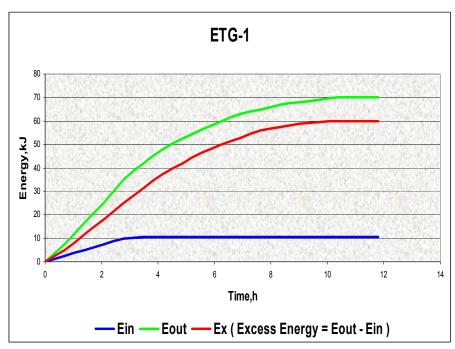
Glow Discharge Cell

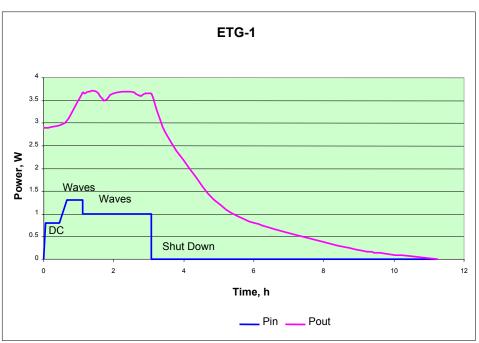


Cell Assembly



Experimental Results with thin Palladium film (about 1 μm) on Stainless Steel Body

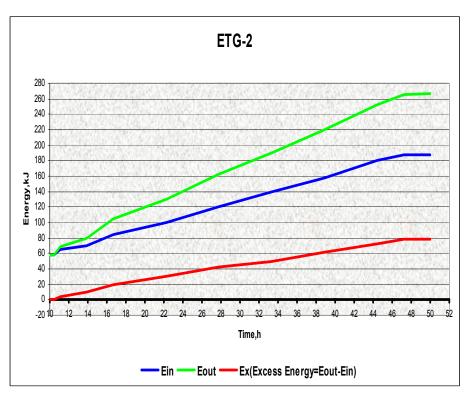


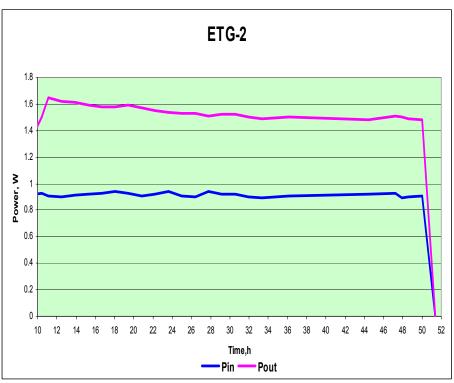


Maximum Excess Power = 3.7 x Input Power

Total Excess Energy = $6.7 \times 10^{-2} \times 10^{-2$

Experimental Results with thick Palladium foil (100 μm)

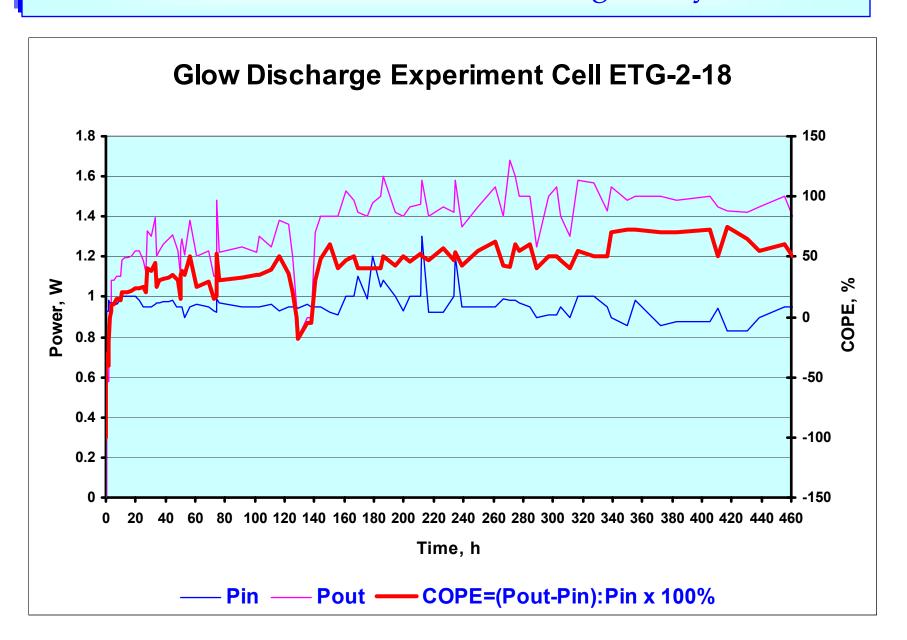




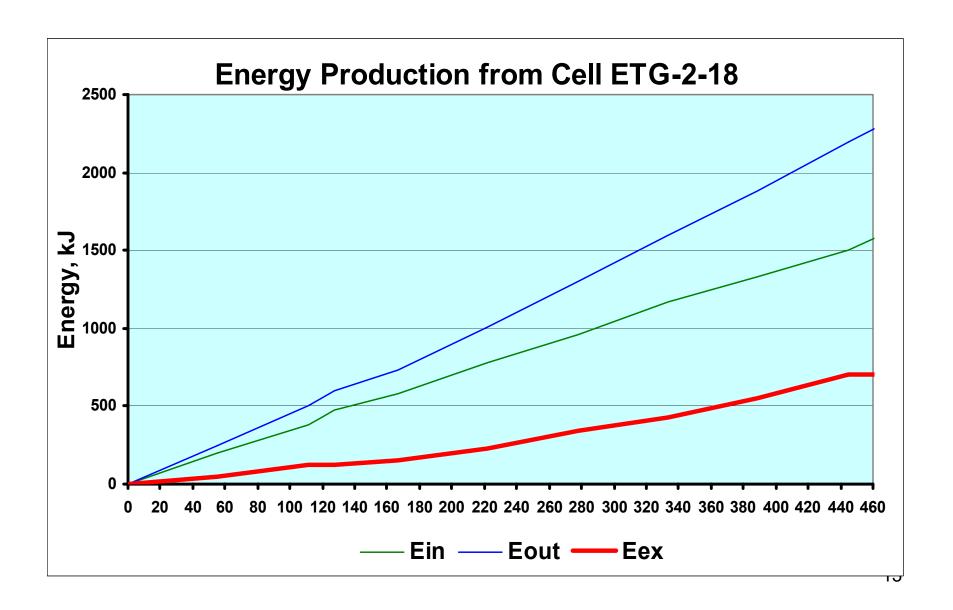
Maximum Excess Power = 0.8 x Input Power

Total Excess Energy = 0.8 x Input Energy

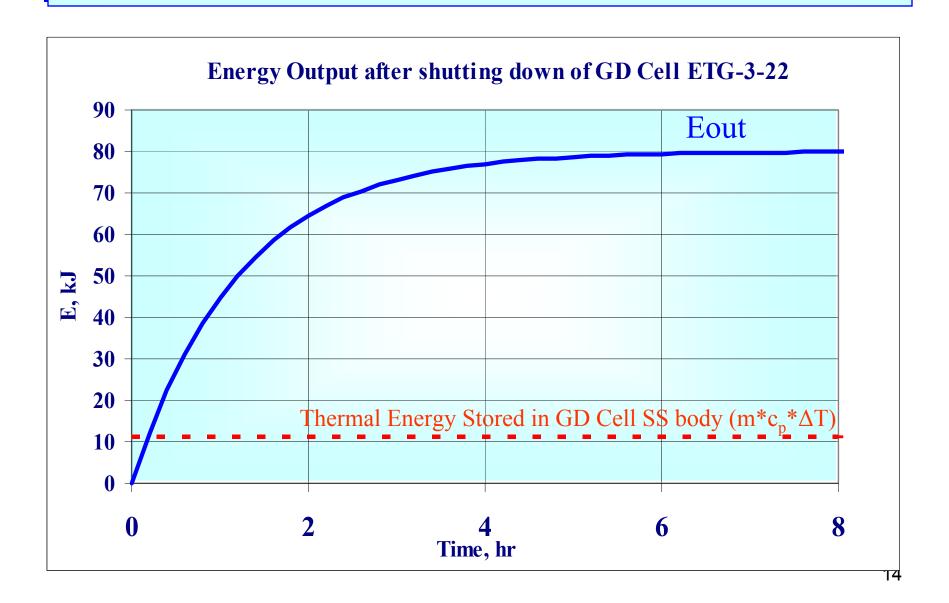
Excess Heat Generation during 20 days



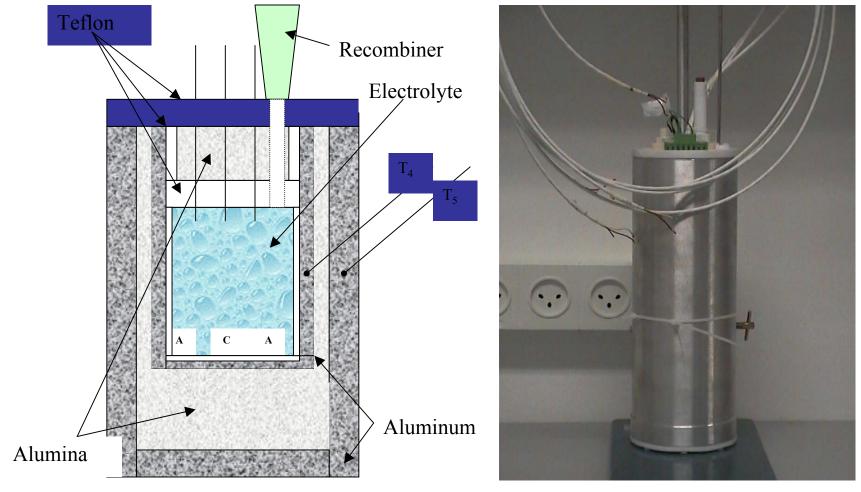
Excess Energy during 20 days



"Heat After Death"



ET Electrolytic Cell



 $0.1M \text{ LiOD in low tritium content } D_2O \text{ (230 ml)}$

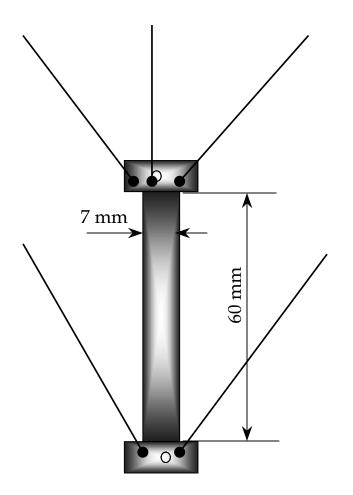
EC is inside a Teflon beaker that is placed inside an isoperibolic calorimeter

Electrolytic Cell



Three cells are immersed in a constant temperature water bath of $+2.5^{\circ}C \pm 0.25^{\circ}C$

Cathode & Pre-treatment



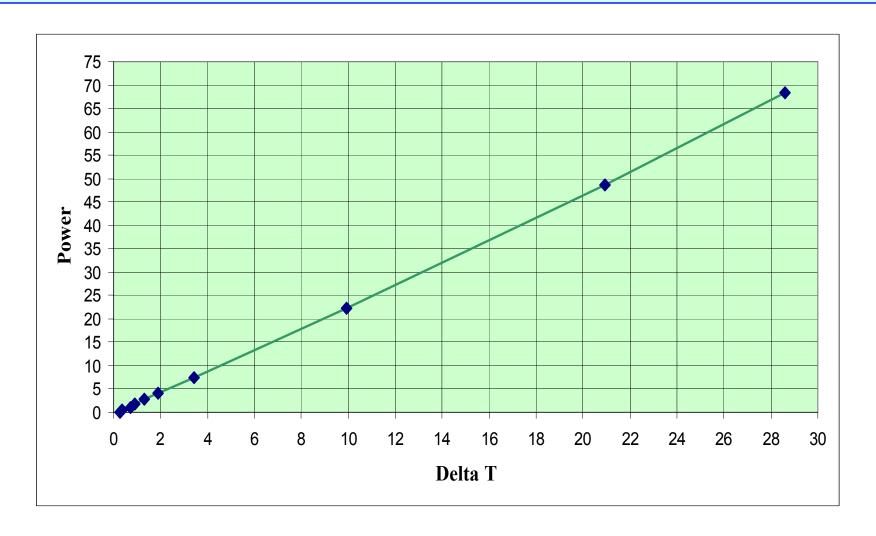
50 μm Pd foil, prepared by

Dr. Vittorio Violante (ENEA Frascatti, Italy)

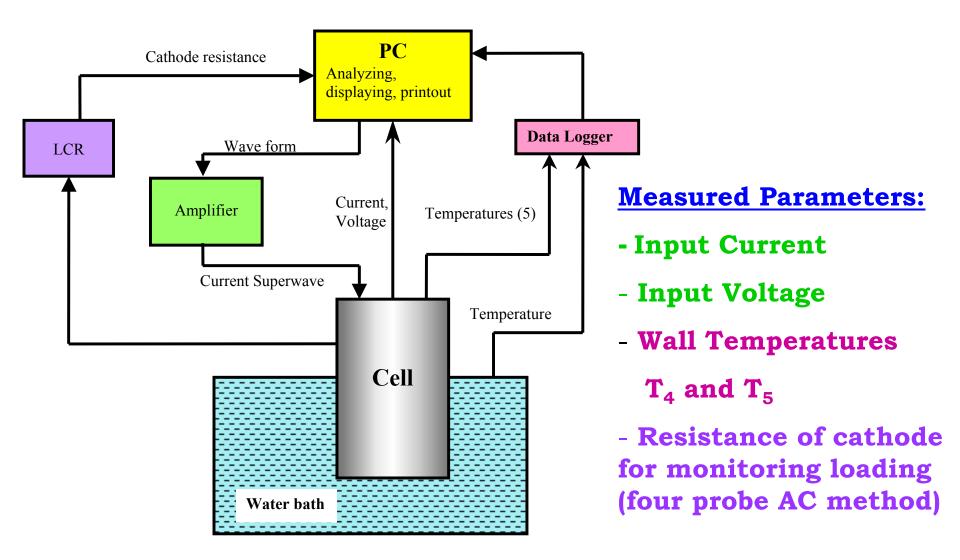
- Annealed at 870°C in vacuum for 1h
- Etched:
 - in Nitric Acid 65-67%; 1 min
 - in Aqua Regia 1:1 water solution; 1 min
- Rinsed:
 - D₂O four times
 - Ethanol 95% twice
 - Ethanol Absolute once
- Dried:

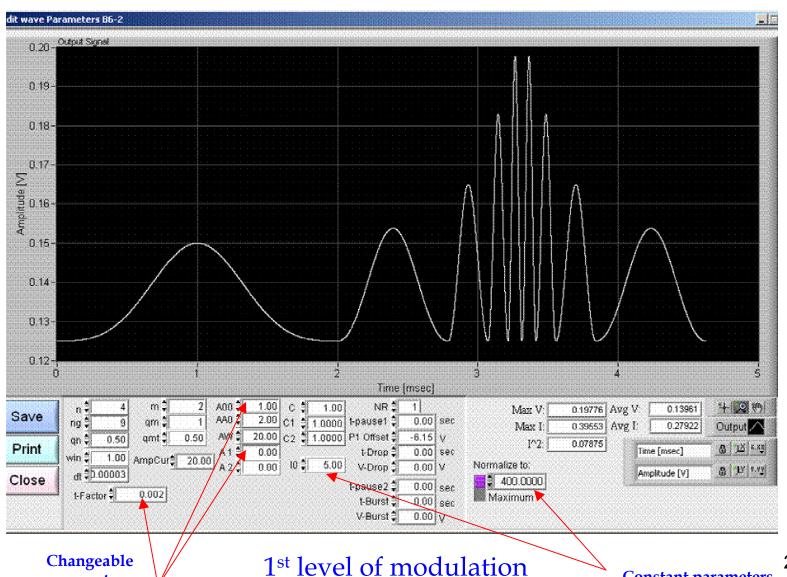
in vacuum at ambient temperature for 24 h

Typical Calibration Curve for Electrolytic Cell Calorimeter

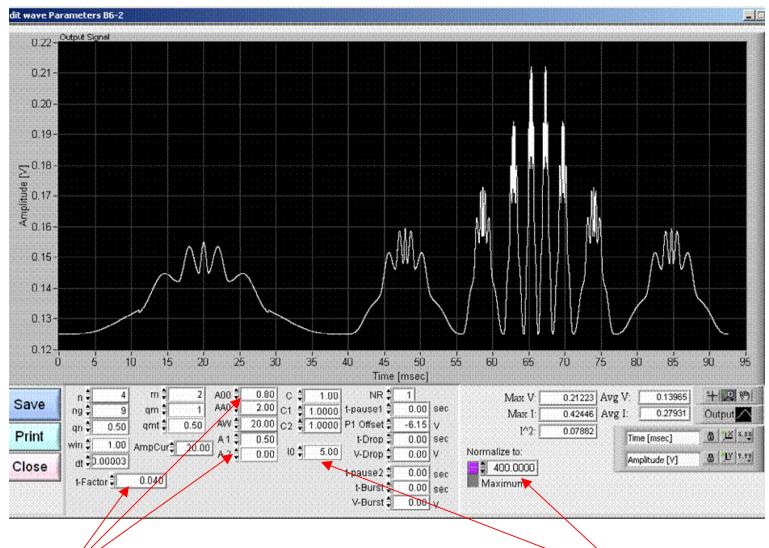


Block diagram



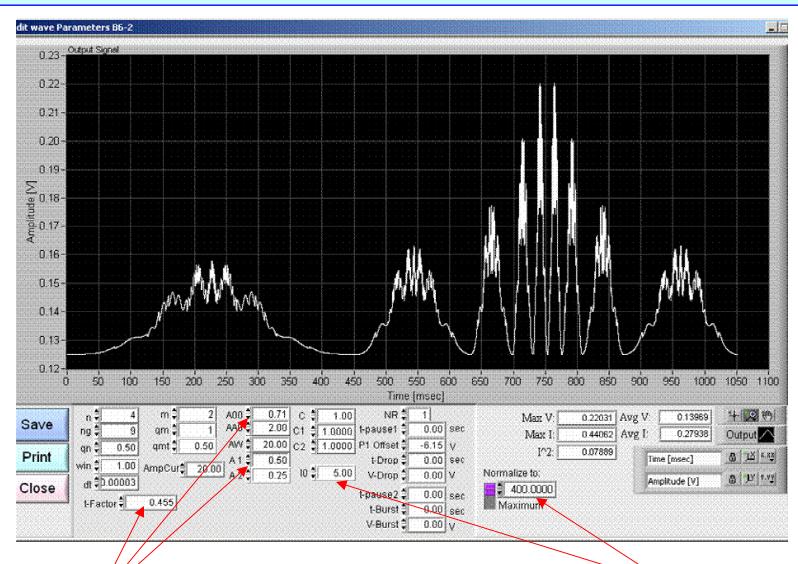


parameters



Changeable parameters

2nd level of modulation

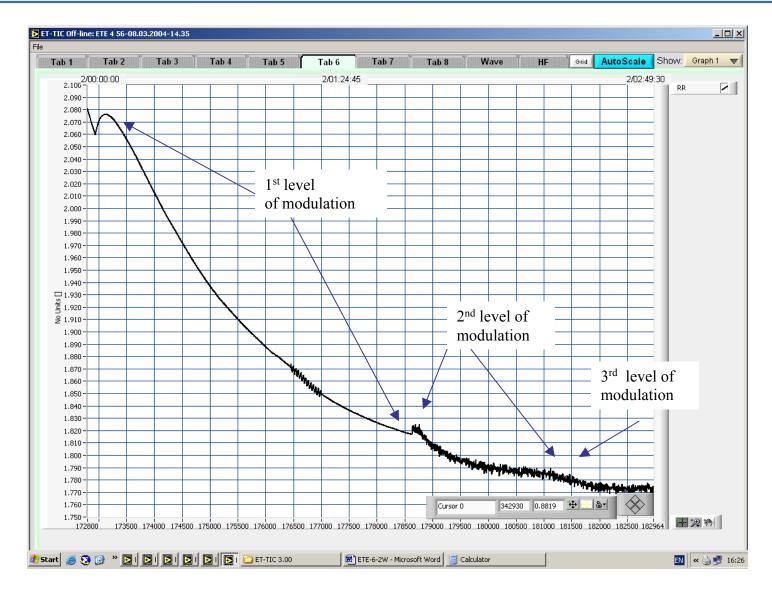


Changeable parameters

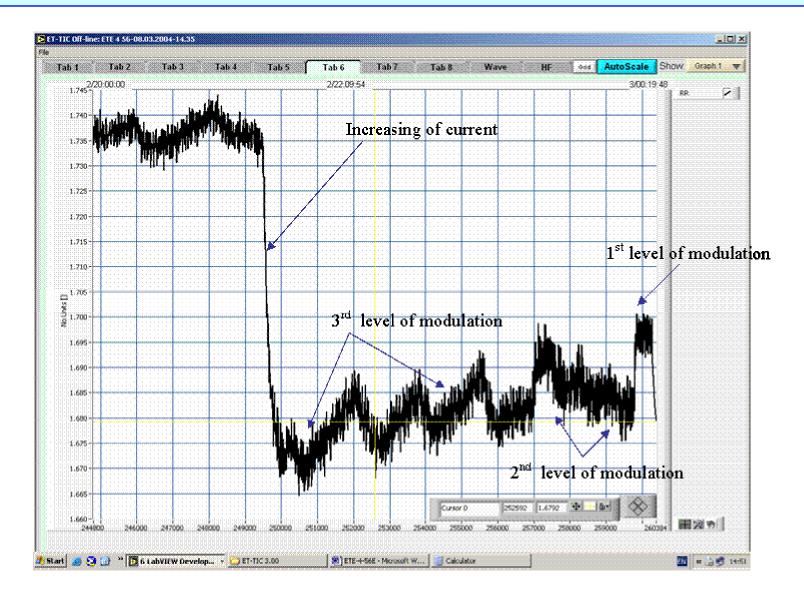
3rd level of modulation



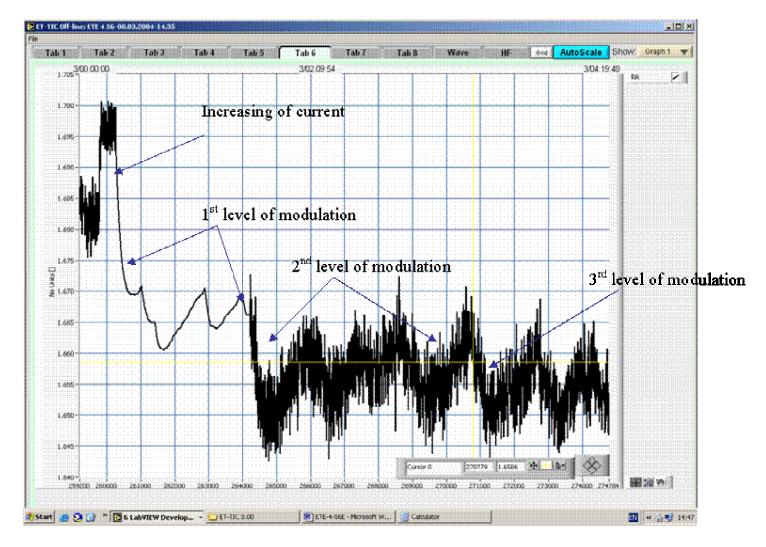
1.

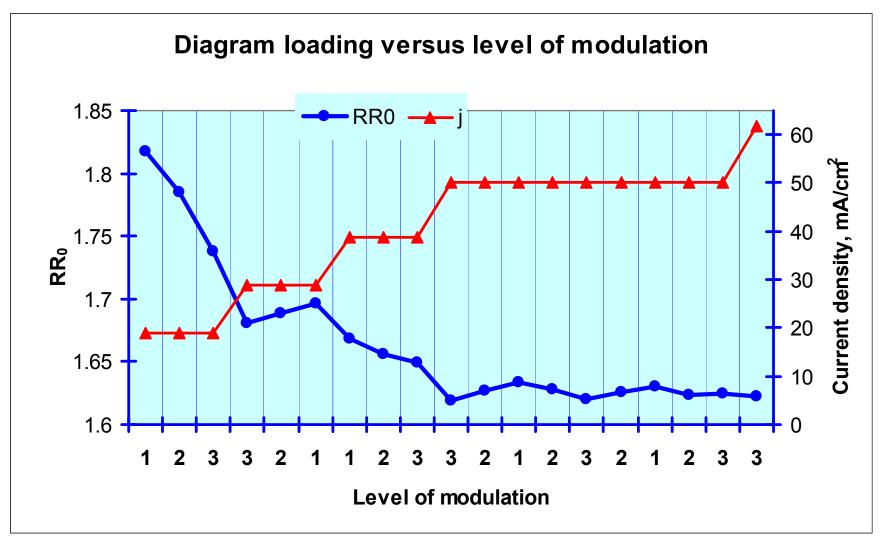


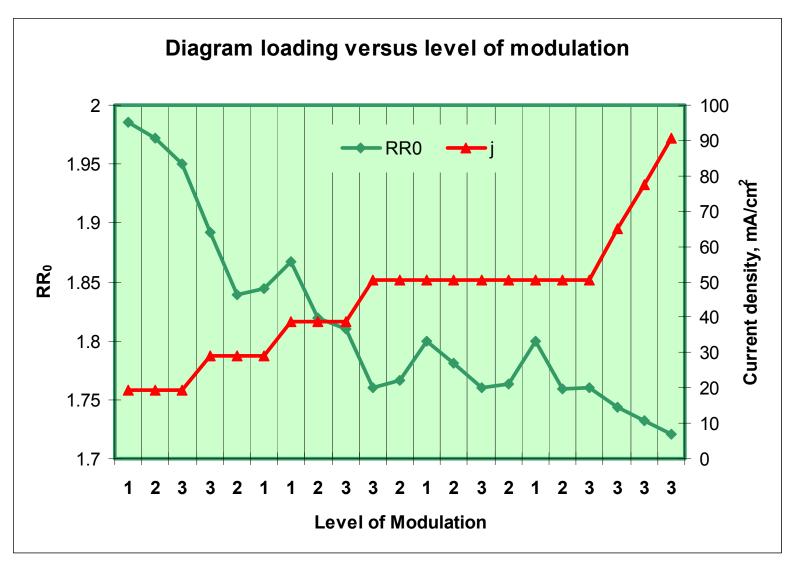
2.



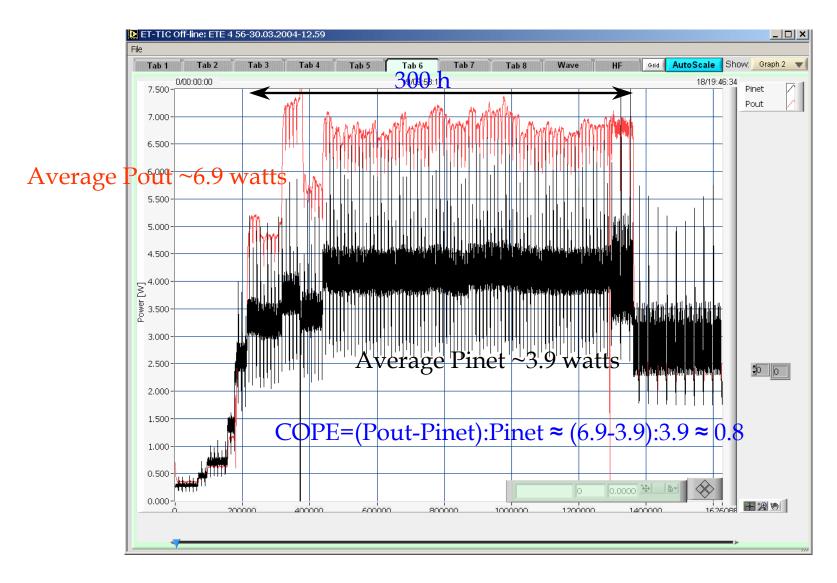




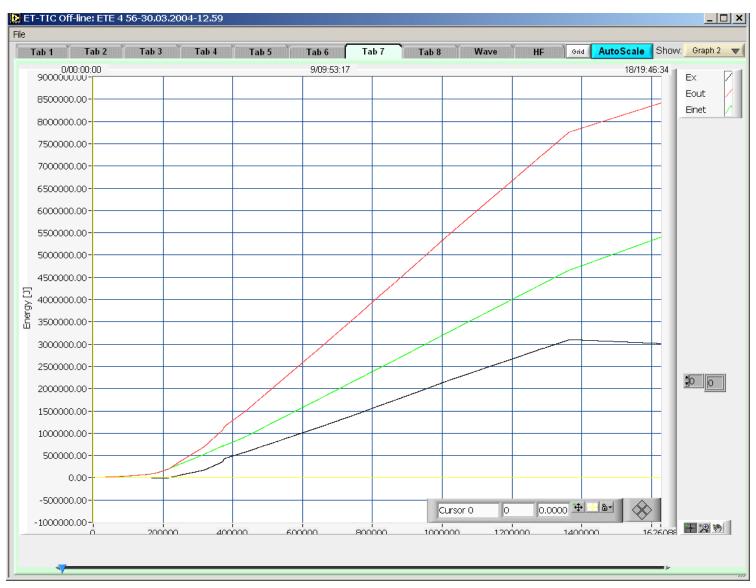




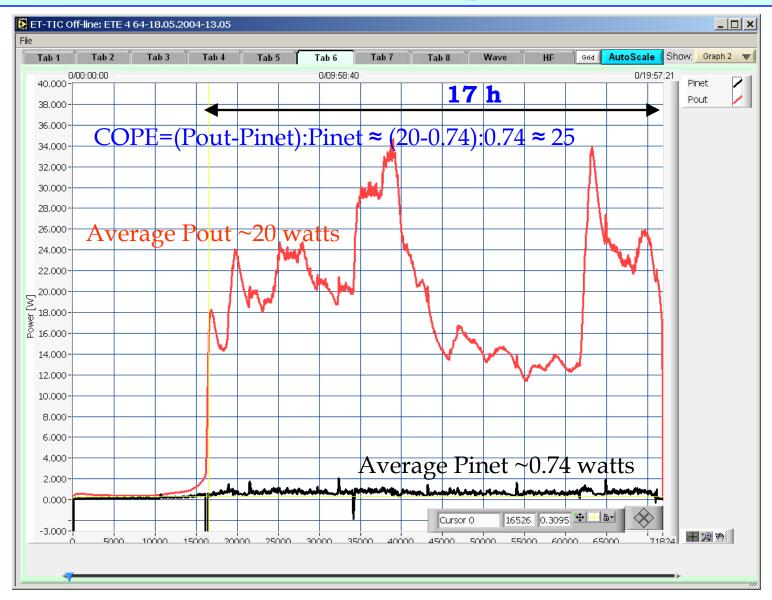
Excess Power; Exp. # 56



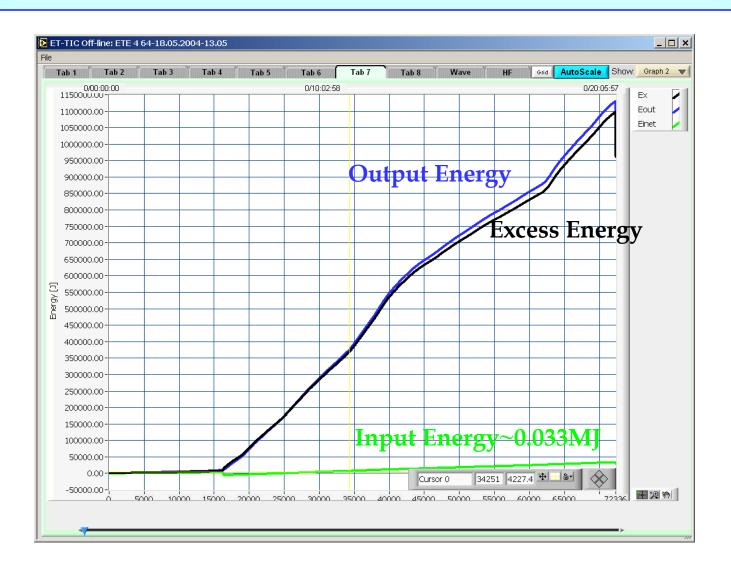
Excess Energy; Exp. # 56



Excess Power; Exp. # 64a



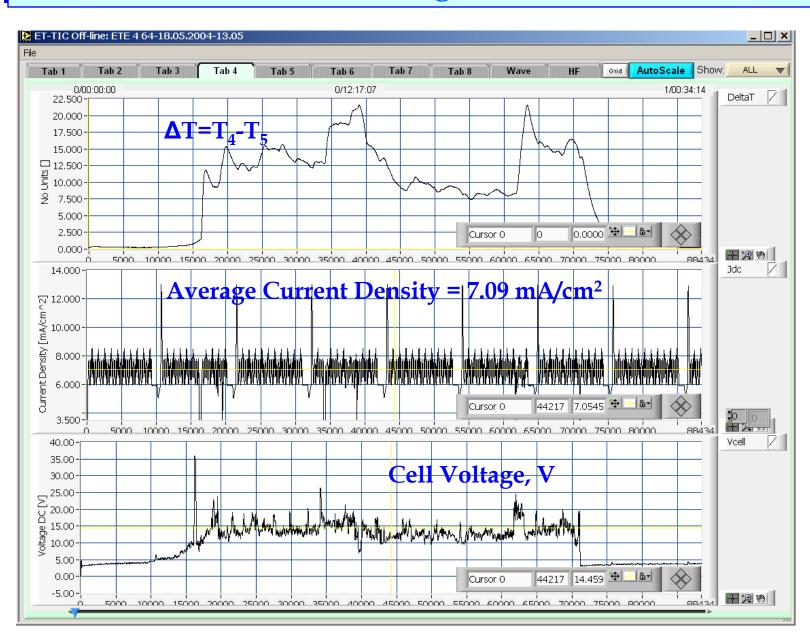
Excess Energy; Exp. # 64a



Temperature Evolution, Exp. # 64a



Current & Voltage; EXP. # 64a

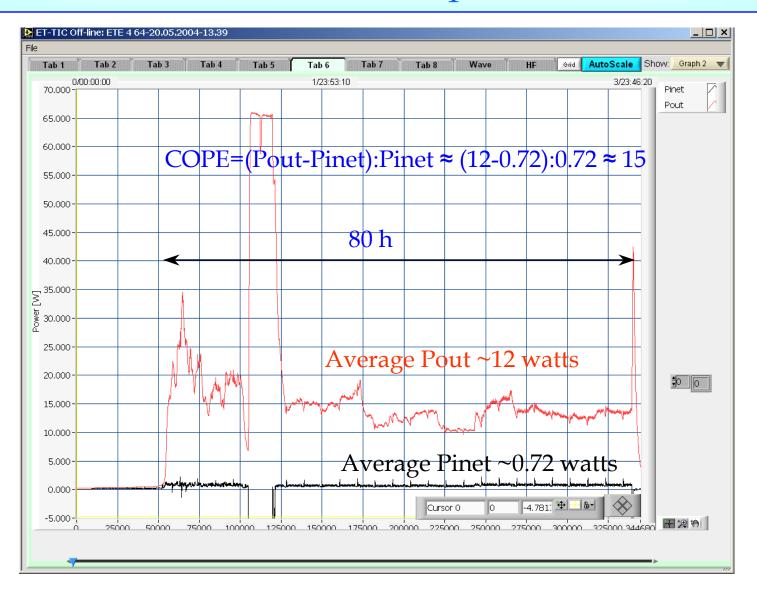


R/Ro & Power; EXP. # 64a

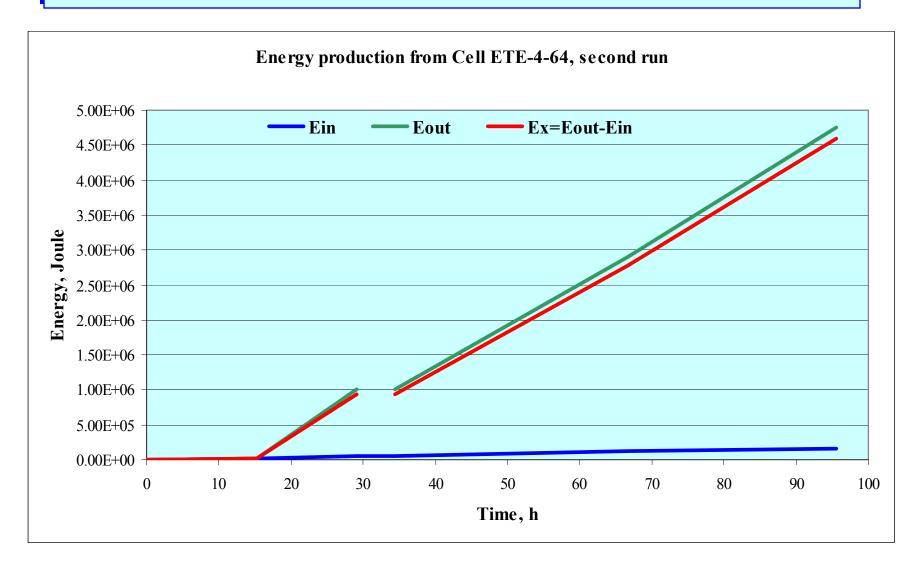


Loading is relatively low (~0.8)

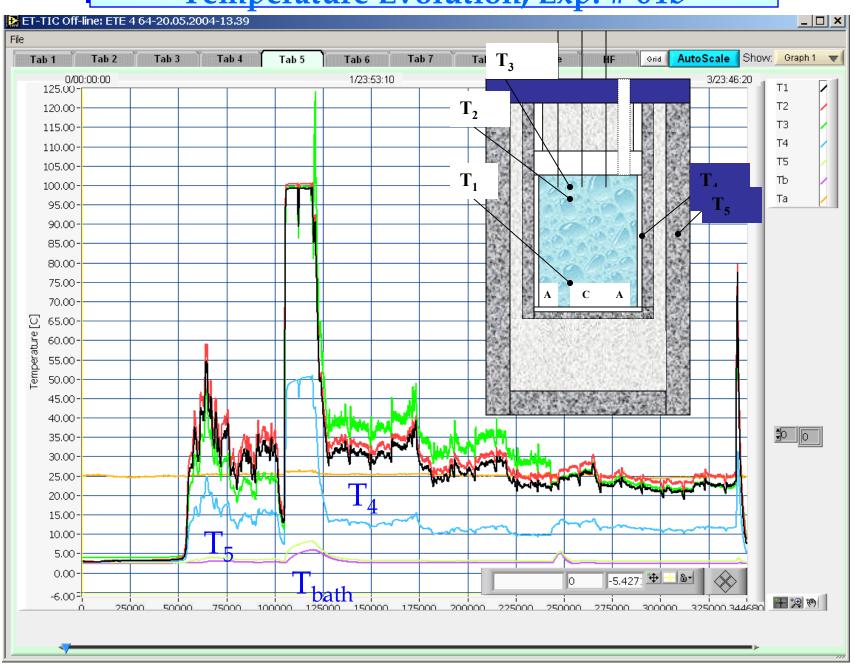
Excess Power; Exp. # 64b



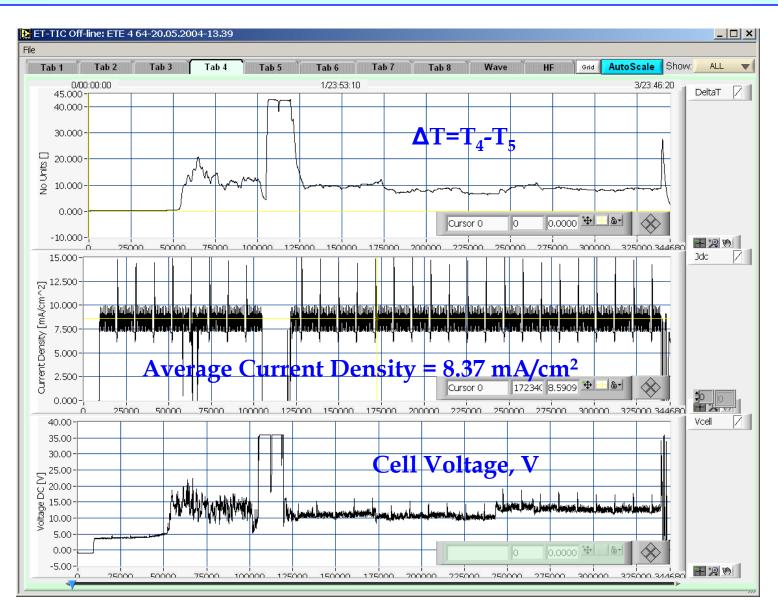
Excess Energy; Exp. # 64b



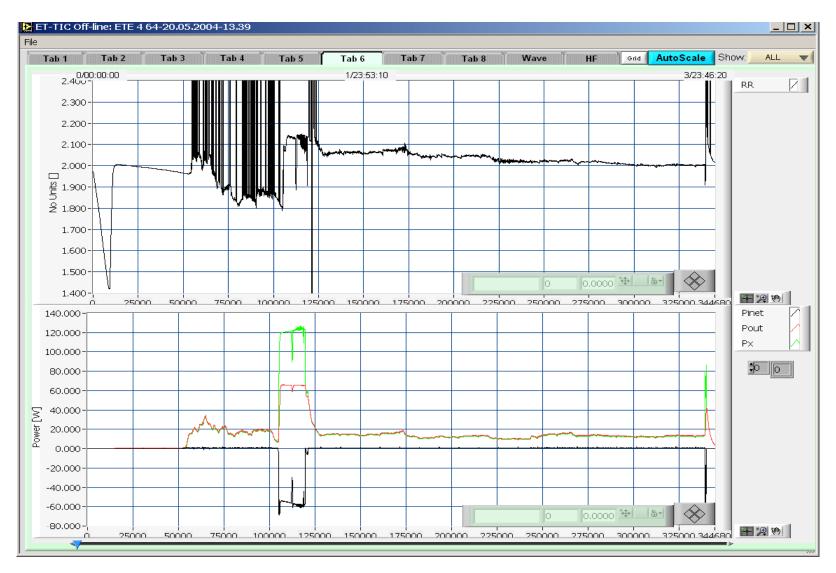
Temperature Evolution, Exp. # 64b



ΔT, Current Density & Voltage; EXP. # 64b



R/Ro & Power; Exp. # 64b



Excess Tritium in # 64

- Tritium concentration measured at end of #64 analysis ~ 250% of reference
- ~625 cm 3 of D_2O has been added to make-up for evaporation; initial inventory was 230 cm 3
- Assuming TDO/D₂O evaporation rate ~ 1.0
- Estimated T effective concentration ~ 750%
- This amount of tritium corresponds to <1J negligible as compared with the 5.7MJ excess energy generated.

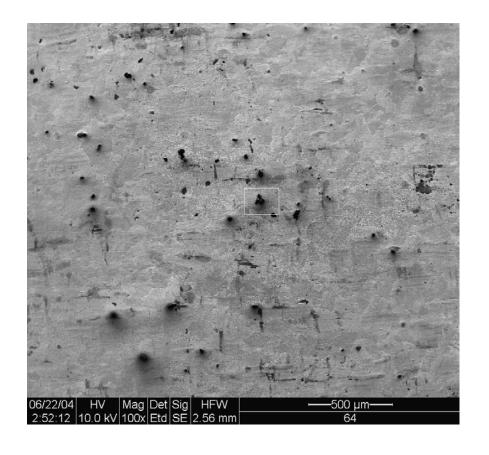
Material Analysis

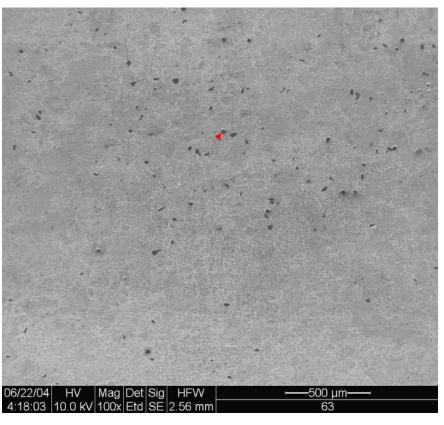
Diagnostics:

- Auger Electron Spectrometry
- Scanning Electron Microscopy-Energy Dispersive
 Spectrometry SEM-EDS
- Transmission Electron Microscopy (TEM)
- Secondary Ion Mass Spectrometry (SIMS)

Material Analysis # 64 vs. 63; SEM-EDS

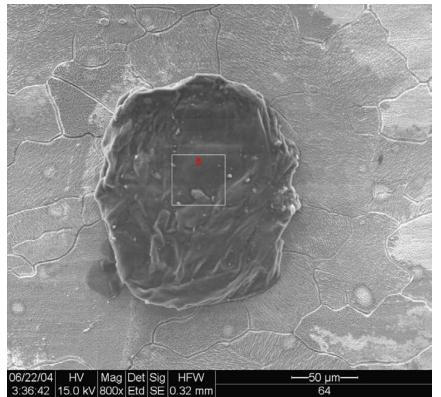
Surface of Pd foil after rolling and annealing at 870°C: sample #64 sample #63



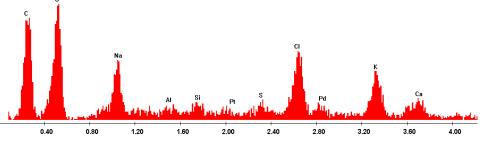


View of Typical Black Spot on # 64 and its composition

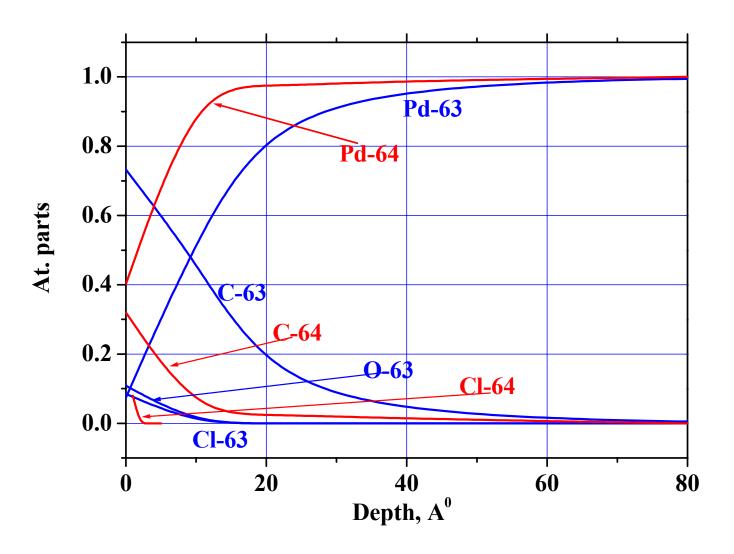
SEM-EDS



Elemen	t Wt %	At %
$\overline{\mathbf{C}}$	35.77	52.48
O	26.19	28.84
Na	4.92	3.77
A1	0.43	0.28
Si	1.05	0.66
Pt	0.39	0.04
S	1.44	0.79
Cl	10.68	5.31
Pd	2.55	0.42
K	11.07	4.99
Ca	5.52	2.43
Total	100.00	100.00

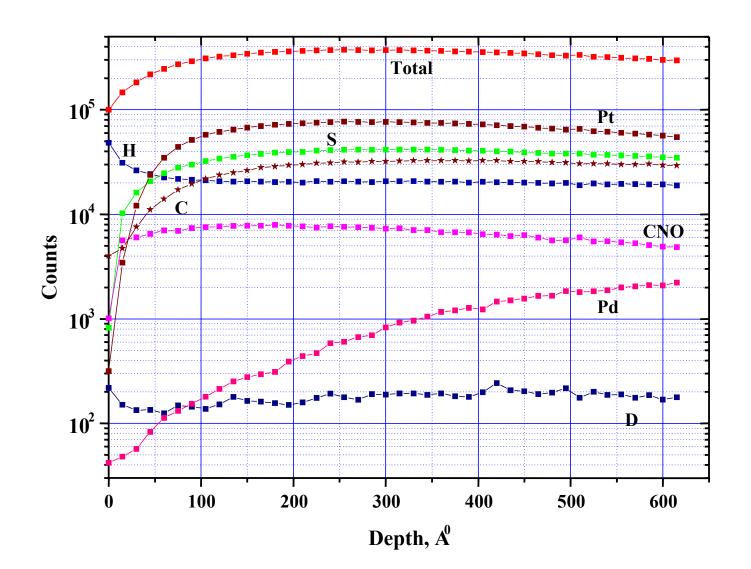


AES profiles of Pd, C, O and Cl of Pd foils #63 and #64

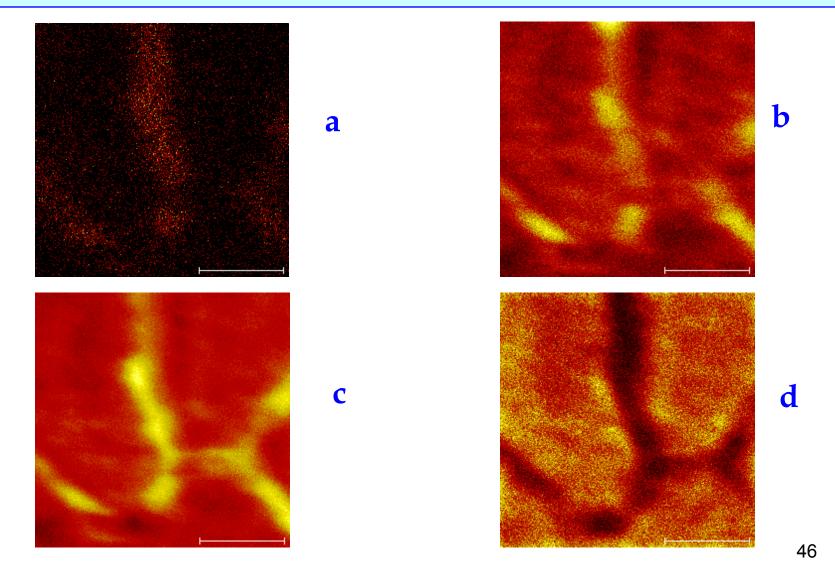


After etching in Aqua Regia

Sample 64. SIMS depth profiling.

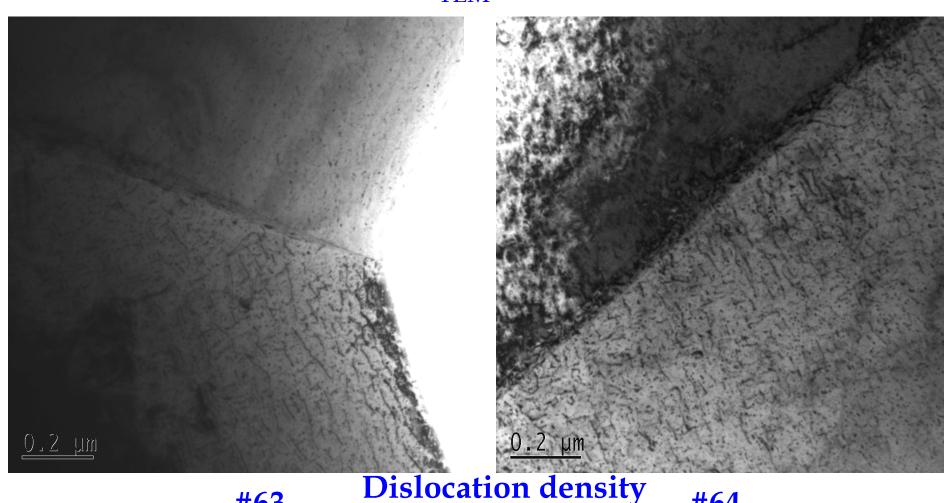


Spatial distribution of selected isotopes measured by SIMS in #64 a – D, b - 32 S, c - total isotopes content, d - 195 Pt. Scale bar - 30 μm .



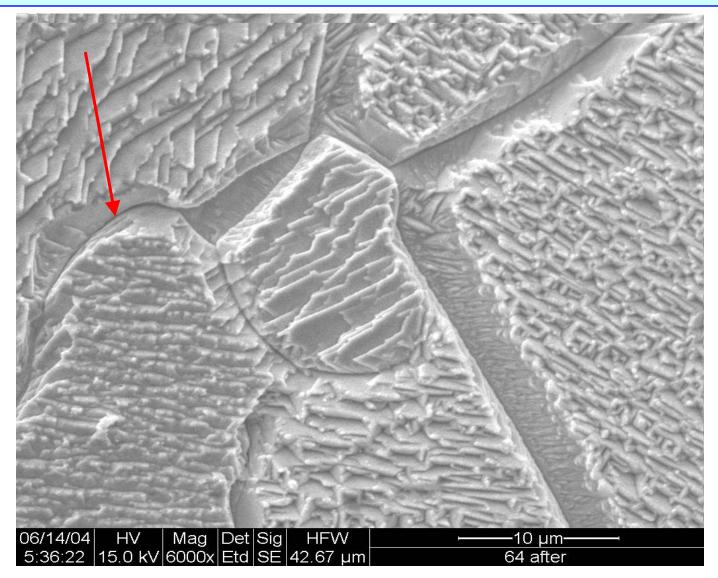
The dislocations in Pd foils after annealing.

TEM

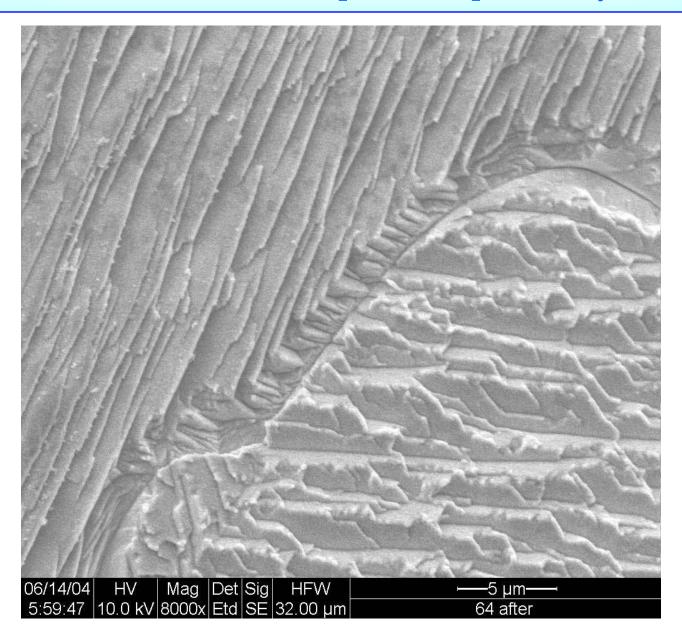


#63 3x10¹⁰, cm⁻² #64 6x10¹⁰, cm⁻²

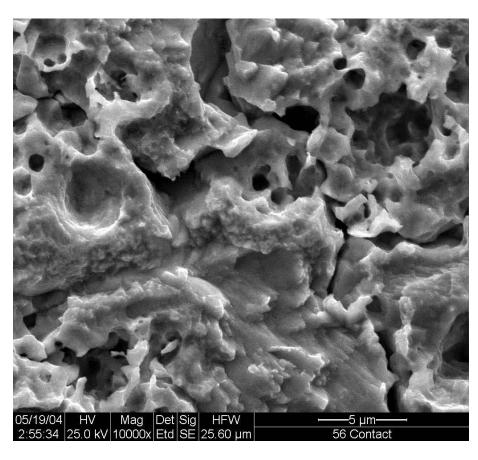
Plastic deformation of Pd foil caused by D2 absorption. Planes of sliding (111) are visible. Sample #64

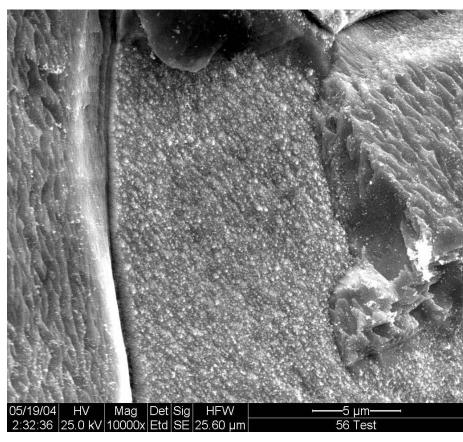


The area that is shown in the previous picture by an arrow.



Sample #56

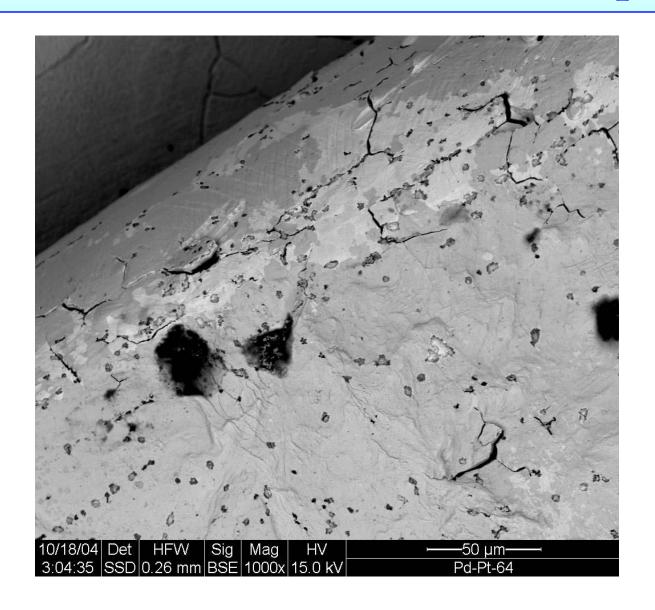




Contact area

Working area

Brittle destruction of Pt Wire Sample #64



Summary of Material Analysis

- Pd surface is covered at least with two types of impurities. The first one is a lubricant used at a rolling process with Pd during plastic deformation of the metal. The second one is a result of adsorption of air components by Pd surface.
- The lubricant stains are of various sizes and configurations and present on surface of all samples before and after the electrolysis.
- Annealing at 850^o does not fully remove the lubricant's components from Pd surface.
- The density of dislocations and the average size of grains in sample # 64 are twice higher, than in the reference sample.
- Nuclear reaction product can not be detected on surface zone due to high concentration of impurities. No He measurement has been attempted.

SUMMARY

- Significant amount of excess heat has been generated in 3 experiments using 2 Pd foils.
- Dardik's modified SuperWaves have been used for current drive in all the three experiments.
- The average current density was relatively low: < 10 mA/cm²
- The maximum excess power was **2500%**. This range of excess power is suitable for commercial applications (although the operating temperatures were too low for such applications).
- The maximum excess energy generated with a single Pd foil is **5.7 MJ**.
- This corresponds to a specific energy of 24.8 KeV per Pd atom.
- The highest average power density obtained is ~70 W/g Pd (versus 20 to 50 W/g U in commercial fission reactors).

SUMMARY (cont.)

- Significant increase in the tritium concentration in the electrolyte has been observed. However, the amount of tritium produced is negligible as compared with the excess energy generated.
- No measurement of He has been attempted.
- The Pd cathode surface was contaminated by what appears to be lubricant from the roller used for pre-treatment as well as impurities adsorbed from the air.