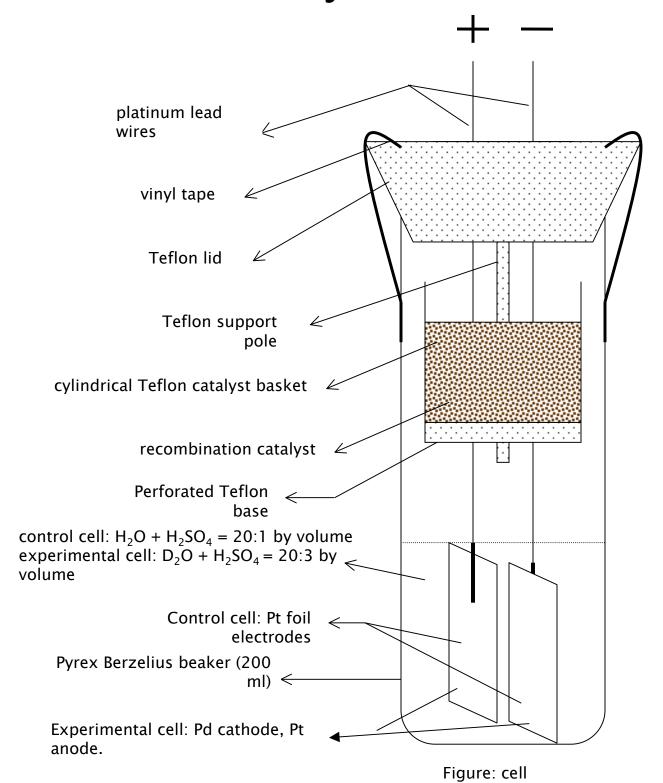
ICCF11 TUTORIAL

MARSEILLES, FRANCE, 10-31-04

SEARCH FOR OPTIMUM CONDITIONS
TO PRODUCE EXCESS HEAT FROM
THE ELECTROLYSIS OF HEAVY
WATER WITH A PALLADIUM CATHODE

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Electrolysis Cell



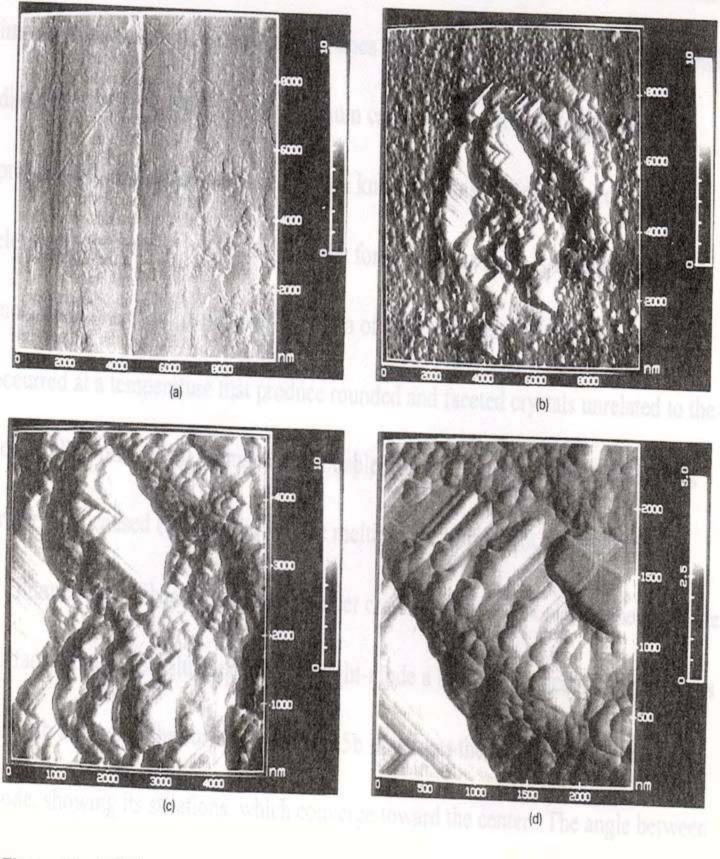


Figure 13. AFM images of a) Pd foil surface after cold rolling (the vertical lines resulted from contact with the steel rolls); b) Pd cathode surface after electrolysis for 12 minutes; c) and d) are enlargements of surface pit in (b) to show the rounded and faceted features.

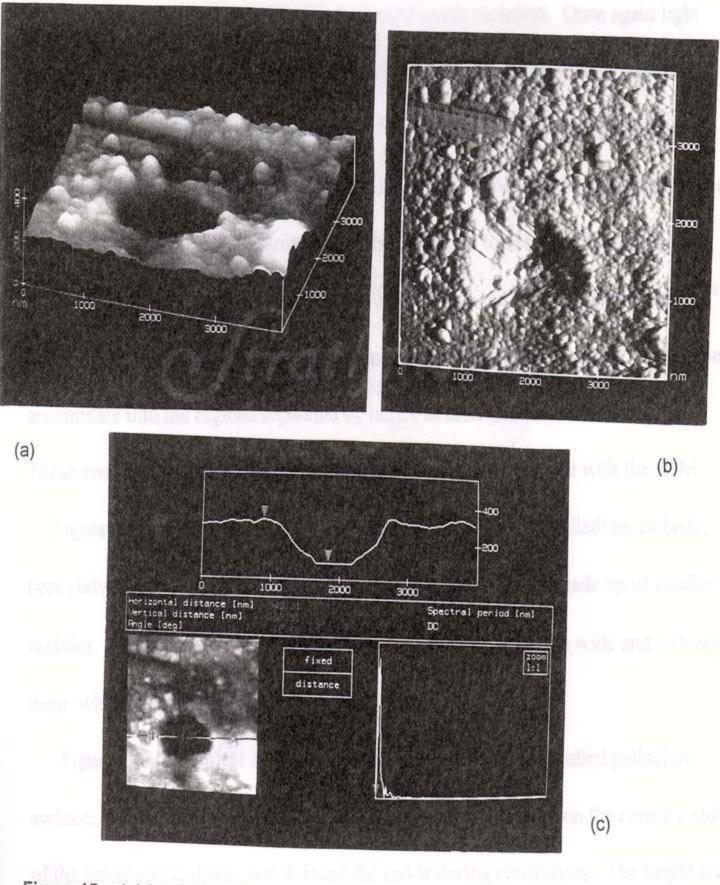
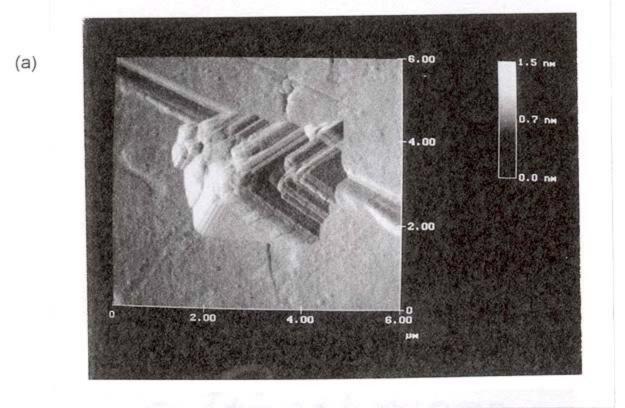


Figure 15. a) A bowl-shaped cavity in Pd cathode, illustrated a) in the height-mode b) in the force-mode, showing striations that converge into its center, and c) contour of the crater. The crater drops > 280 nm vertically from its rim.



(b)

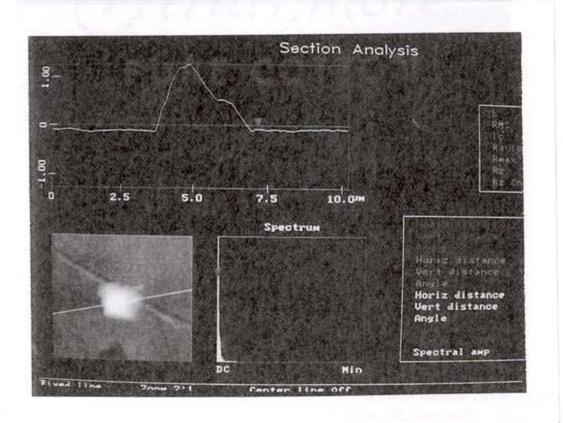
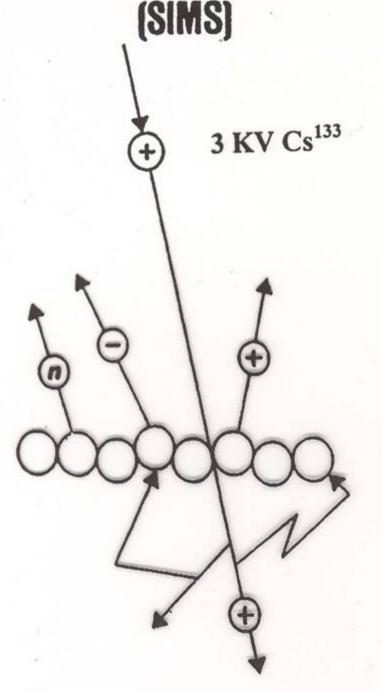


Figure 22. An AFM force-mode image a) of the palladium cathode side facing away from the anode, electrolyzed for a total of 105 seconds. The asperity seen here appears to be from melting and recrystallization. The angles of the facets suggest that these are traces of $\{111\}$ planes. The section analysis b) indicates that this feature rises about 1 μ m above the surrounding surface.

SECONDARY ION MASS SPECTROSCOPY



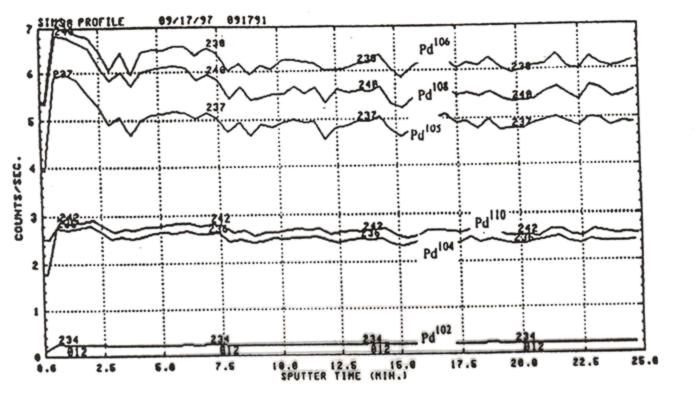
Beam scanned
1.4 mm × 1 mm
1 Å/sec removed
from Pd surface

Cathode

Figure 29. SIMS is a procedure that bombards a small area of the Pd surface with a primary beam (Cs133). Surface atomic layers are sputtered off, and the mass of the ions from the resulting sputtered particles are analyzed using a mass spectrometer to provide elemental and isotopic identification.

RELATIVE ABUNDANCES AND WEIGHTS OF NATURALLY OCCURING ISOTOPES

Elem.	AMU	Isotopic Comp. (at.%)
Pd ¹⁰²	101:905634	1.020
Pd ¹⁰⁴	103.904029	11.14
Pd ¹⁰⁵	104.905079	22.33
Pd ¹⁰⁶	105.903478	27.33
Pd 108	107.903895	26.46
Pd110	109.905167	11.72
Ag 107	106.905092	51.839
Ag 109	108.904757	48.161
Cd100	105.906461	1.25
Cd ¹⁰⁸	107.904176	0.89
Cd110	109.903005	12.49



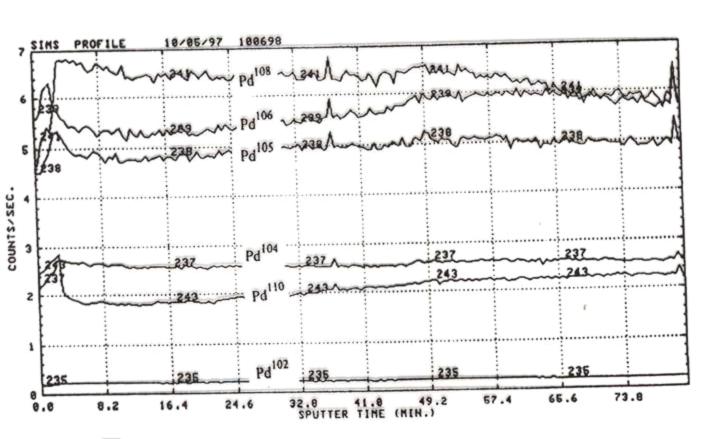


Figure 1. SIMS profile of the six palladium isotopes for a region of the palladium sample not electrolyzed (top), and a region of the six minute heavy water electrolyzed sample (bottom). The latter shows isotopic inversions of Pd¹⁰⁸ with Pd¹⁰⁶, which merge after ~65 minutes sputtering. Also Pd¹¹⁰ and Pd¹⁰⁴ are inverted.

Energy Dispersive X-Ray Spectrometer

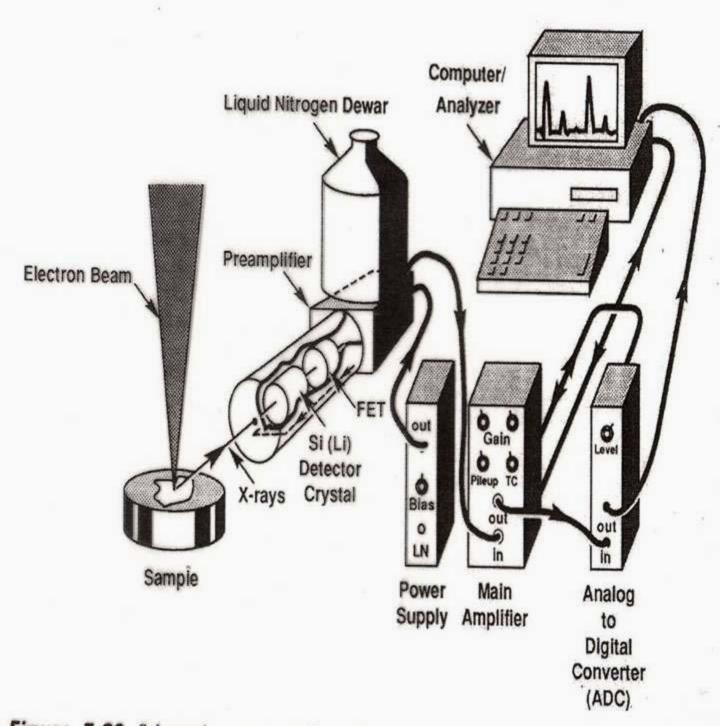
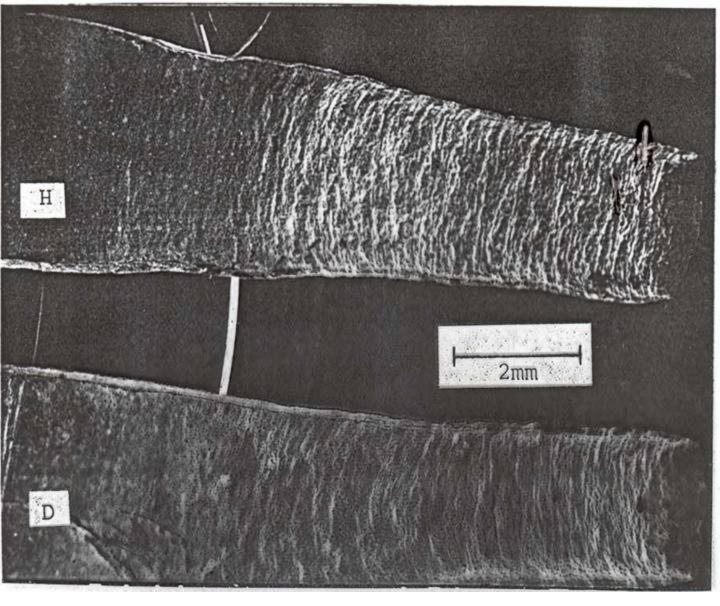


Figure 5.20. Schematic representation of an energy-dispersive spectrometer and its associated electronics.



Pd cathodes 0.35mm thick, after electrolysis for about 400 hours

H - electrolyzed in H₂O - H₂SO₄ D - electrolyzed in D₂O - H₂SO₄

- 0.1 to 0.2 watts excess power produced by D₂O cell compared with H₂O cell during first 300 hours of operation
- 0.1 to 0.2 watts excess power produced by H₂O cell compared with D₂O cell during final 100 hours of operation

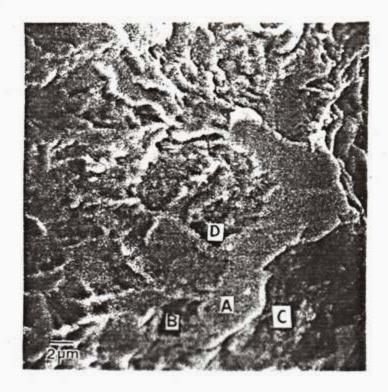
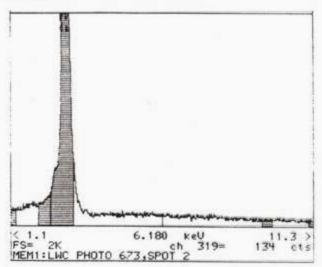


Fig. 2. Enlargement of lower right corner of H₂O cathode in Fig. 1.

the electrolyte where it occurs due to slow dissolution of the Pt anode. Au, nowever, is not expected to arise from a pure Pt anode. Nor is it expected to occur inhomogeneously as an impurity in Pd because Au and Pd are completely miscible in the solid state³.



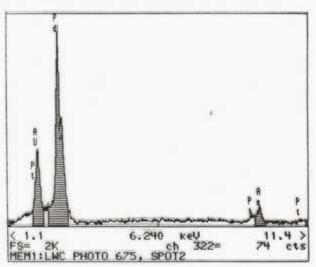


Fig. 3. EDS spectrum from region A of the H₂O cathode shown in Fig. 2.

Fig. 4. EDS spectrum from region B of the H₂O cathode shown in Fig. 2.

Assume that the following fusion reaction occurs:

$$D^+ + D \rightarrow {}^{3}He (0.82 \text{ MeV}) + n(2.45 \text{ MeV})$$

Subsequently, the neutron produced in this reaction is absorbed by a Pt atom on the surface of the Pd electrode. The energy of the neutron is converted to heat, and the Pt atom decays to Au by the following reaction:

$$196$$
Pt + n = 197 Pt = 197 Au

197Pt is an unstable isotope, with a half life of 80 min. It yields an orbital electron, thus becoming 197Au (stable).

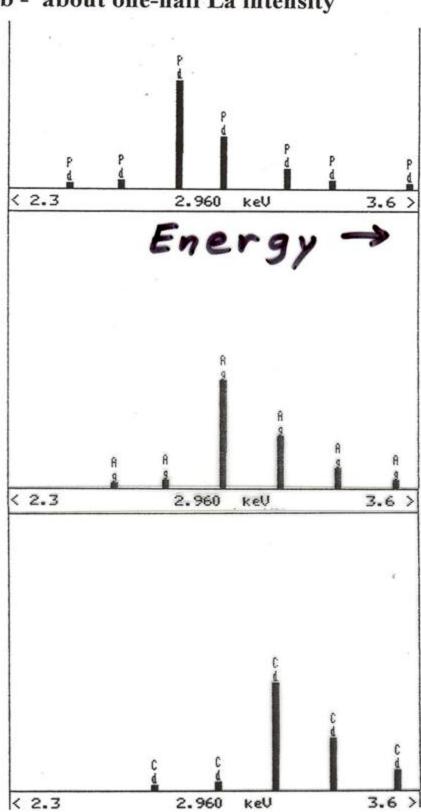
20k 728 <u>10 ym</u>

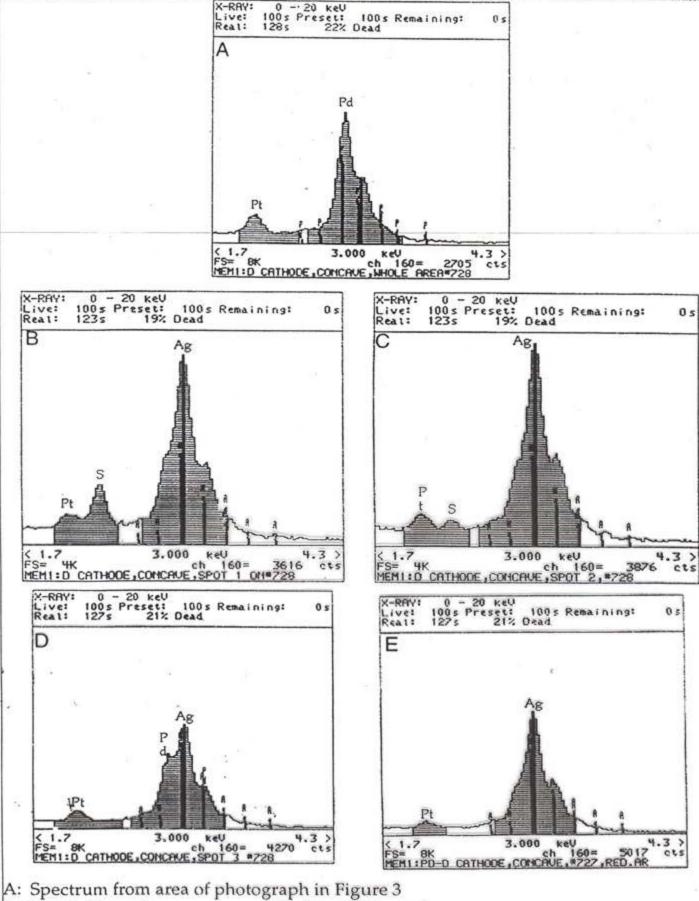
Characteristic Peaks for Pd, Ag, and Cd

La - most intense

Lb - about one-half La intensity

1 Intensity





B: Spot 1, Figure 4

C: Spot 2, Figure 4

D: Spot 3, Figure 4

E: Spot 4, Figure 4

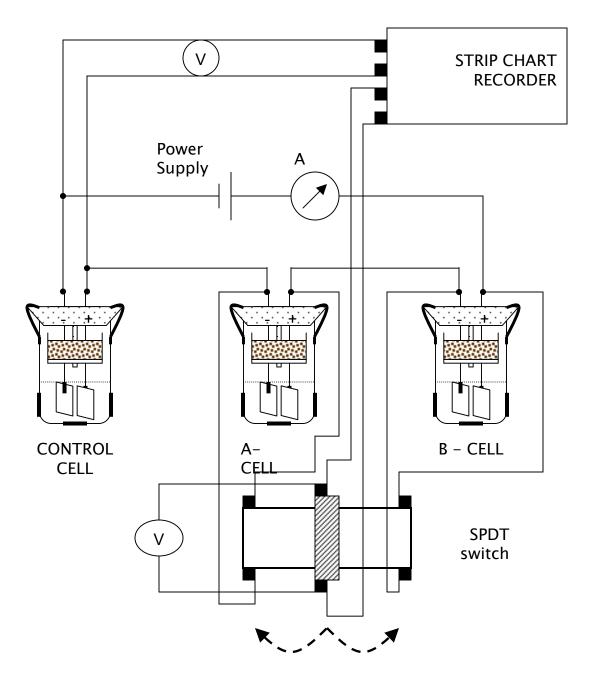


Figure: Circuit diagram. A Single-pole-double-switch (SPDT) switch was used to switch the voltmeter reading between the two experimental cells.

'Represents a K-type thermocouple used for temperature measurement. All thermocouples were attached to a STP-36CJC-102-02A board manufactured by Kiethley. The board was connected to a computer which was used to read and paste temperature values in a MS Excel worksheet.

Equations for excess heat

At steady state:

Power IN = Power OUT – Anomalous power

$$IV_{1} = k(T_{1} - T_{A}) + \frac{dH_{1}}{dt}$$

$$IV_{2} = k(T_{2} - T_{A}) + \frac{dH_{2}}{dt} - \frac{dH_{xs}}{dt}$$

I = constant current

 V_1 = control cell voltage

V₂ = experimental cell voltage

k = a constant

 T_1 = control cell temperature

 T_2 = experimental cell temperature

T_A = ambient temperature

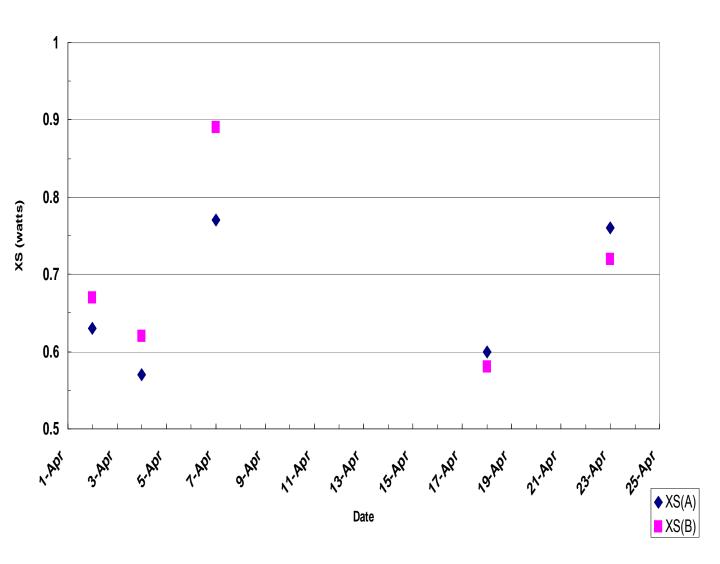
dH₁/dt = power lost from control cell due to evaporation

dH₂/dt = power lost from experimental cell due to evaporation

 $dH_{xs}/dt = excess thermal power$

Excess power data for 2 cells with Pd cathodes and heavy water electrolyte in comparison with a control cell which had a Pt cathode and light water electrolyte.

Current = 3A,Current density = 0.48 A/cm-sq. Precision of measurement is estimated at +/- 0.1W



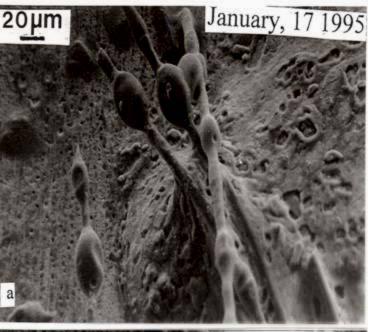
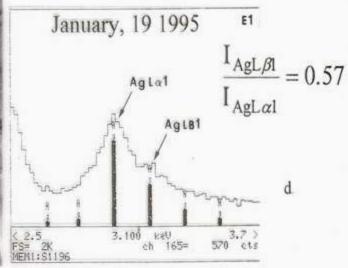
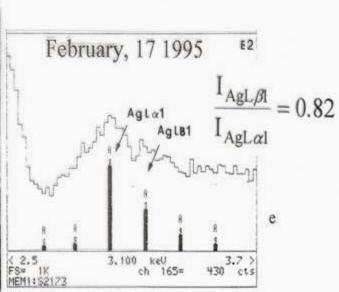


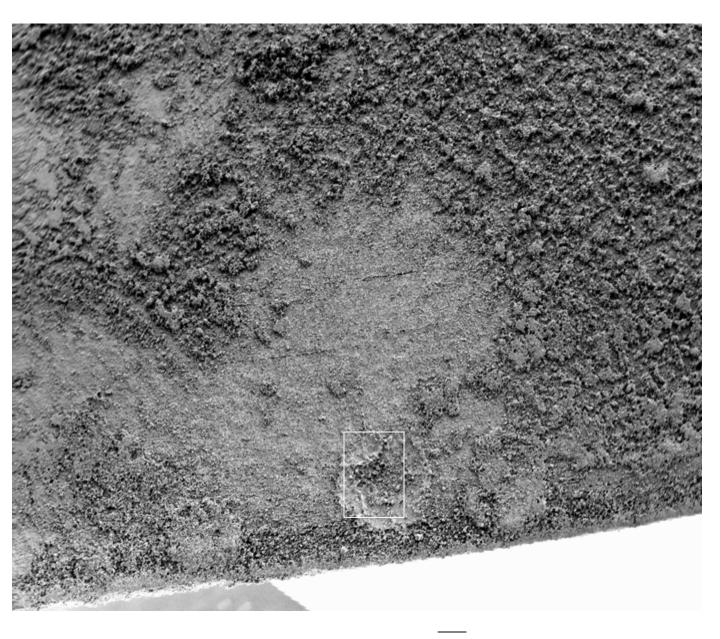
Fig. 5. Changes in morphology and EDS spectra of the features in the square shown in Fig. 4a. These changes occurred during storage at room temperature with no further electrolysis.



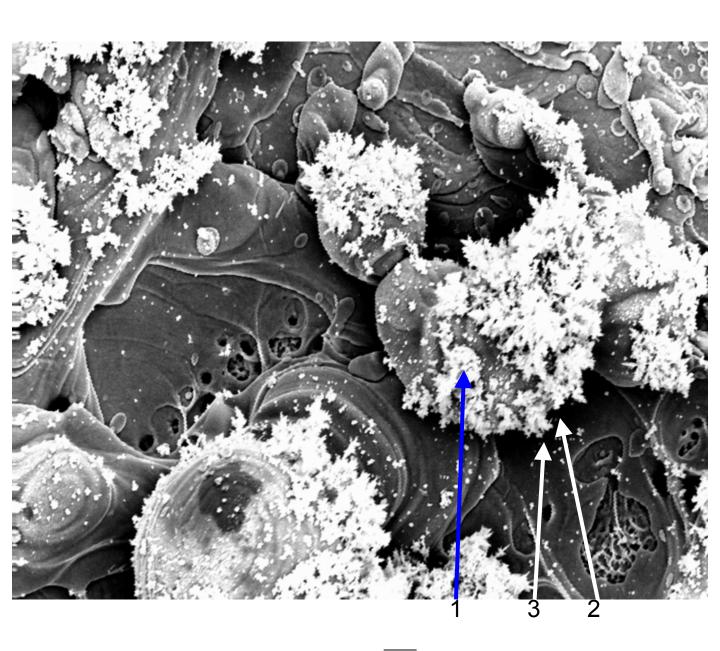




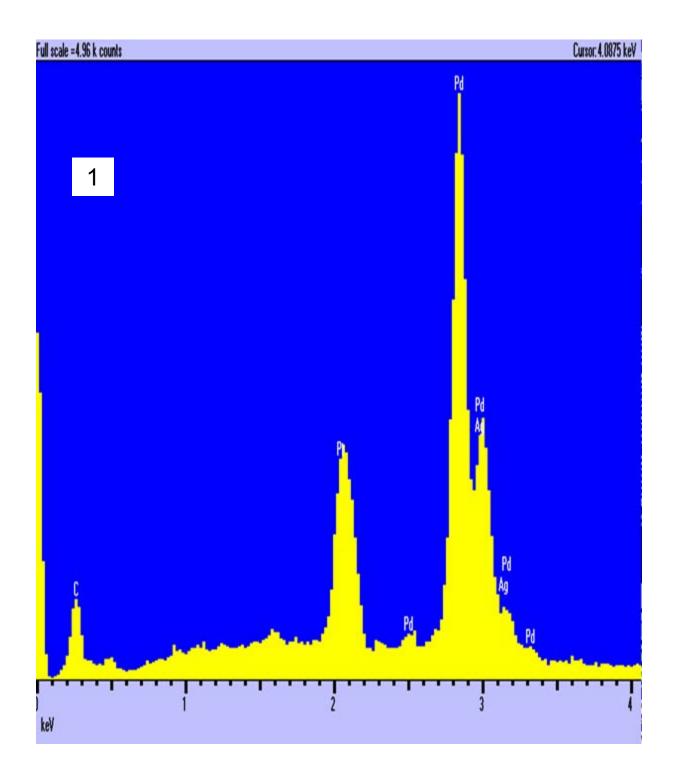




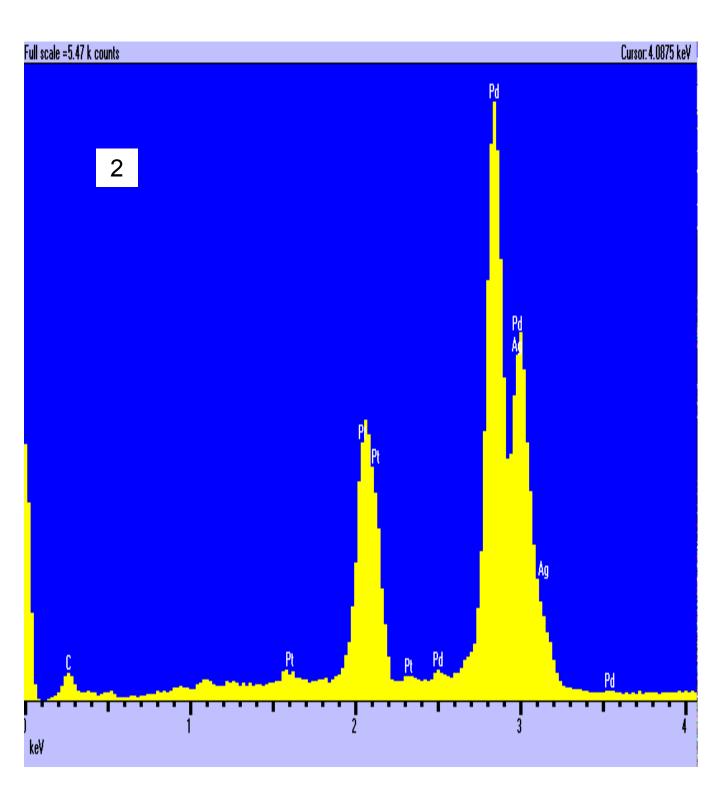
0.58mm



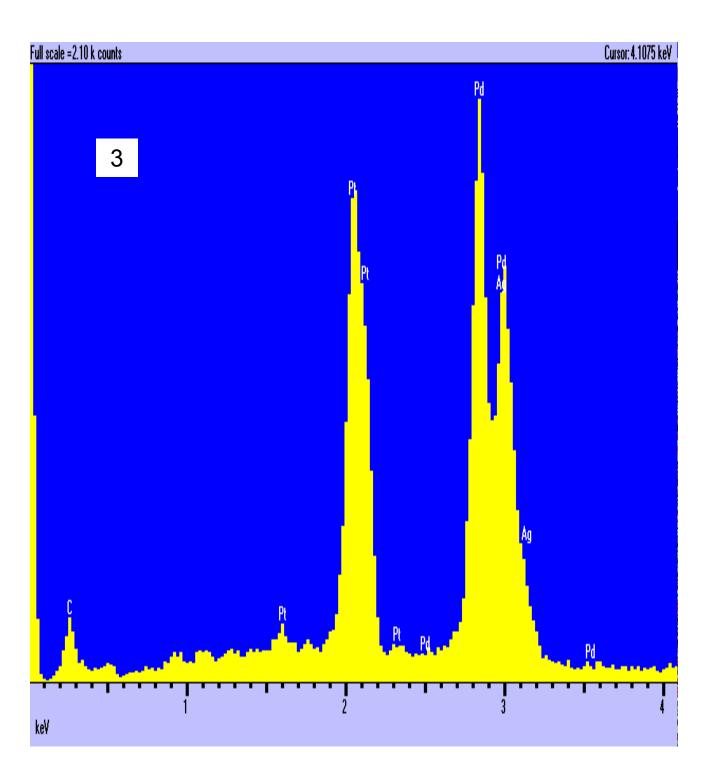
10µm



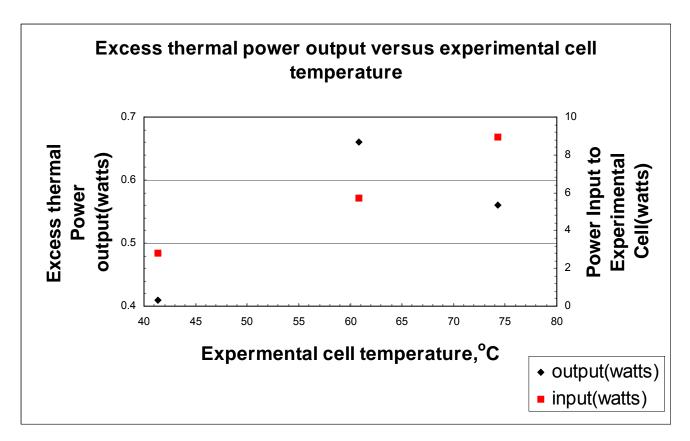
Palladium $L\beta/L\alpha = 0.4$ \rightarrow Elemental Silver = 0%



Palladium $L\beta/L\alpha = 0.6$ \rightarrow Elemental Silver = 7%



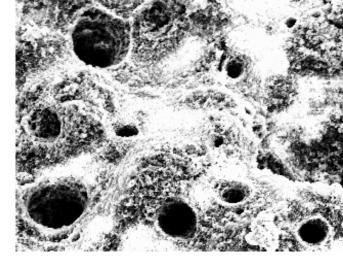
Palladium $L\beta/L\alpha = 0.74$ \rightarrow Elemental Silver = 10%



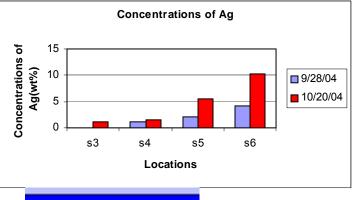
Excess thermal power output versus experimental cell temperature. Current density was about 0.5A/cm² for all three experiments. Pd cathode thickness was 0.05mm.



Pd-0.05mm-8mm-36mm—cv-0.15kx-Sep.28

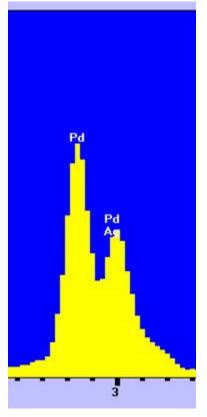


Pd-0.05mm-8mm-36mm—cv-0.15kx-Oct.20



Pd Aq

Pd-0.05mm-8mm-36mm—cv-0.15kx, Sep.28-s6 (Pd Lb/La=0.57)

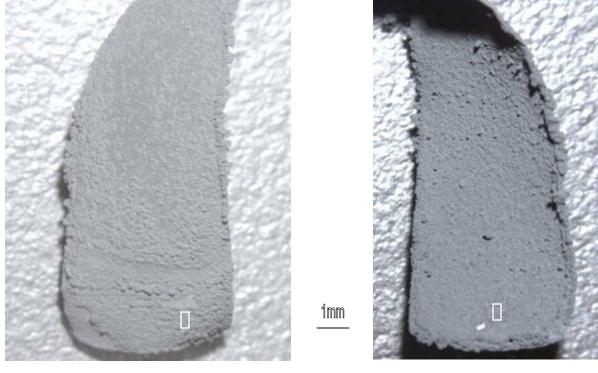


Pd-0.05mm-8mm-36mm—cv-0.15kx, Oct.20-s6 (Pd Lb/La=0.63)

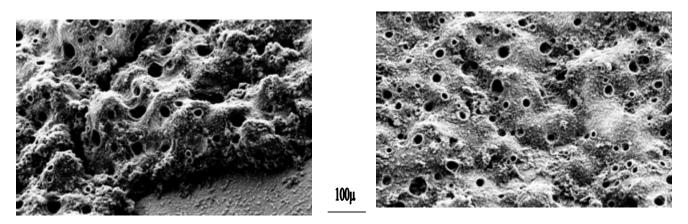


Pd-0.05mm-8mm-36mm Sample

1. Image

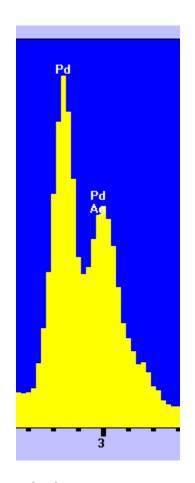


Pd-0.05mm-8mm-36mm-cv Pd-0.05mm-8mm-36mm-cc Fig.1.1 Image of sample Pd-0.05mm-8mm-36mm(small magnification)

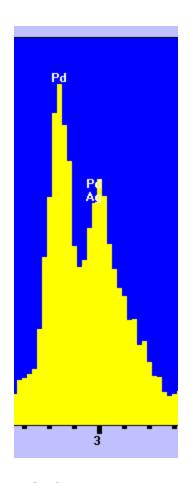


Pd-0.05mm-8mm-36mm-cv-0.15kx Pd-0.05mm-8mm-36mm-cc-0.15kx Fig.1.2 Image of sample Pd-0.05mm-8mm-36mm (big magnification)

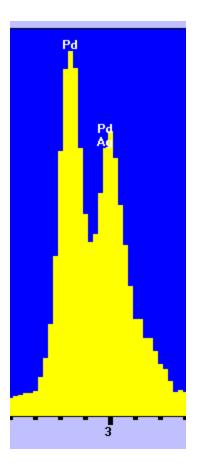
2. Spectrums



EDS of Pd-0.05mm-8mm-36mm convex, area s11, which has largest silver concentration, 4.02% (Pd Lb/La=0.63)



EDS of Pd-0.05mm-8mm-36mm convex, area s14, which has silver concentration, 1.9% (Pd Lb/La=0.73)



EDS of Pd-0.05mm-8mm-36mm concave, area s10, which has largest silver concentration, 9.56% (Pd Lb/La=0.78)

Figure: EDS of sample Pd-0.05mm-8mm-36mm

3. Concentrations

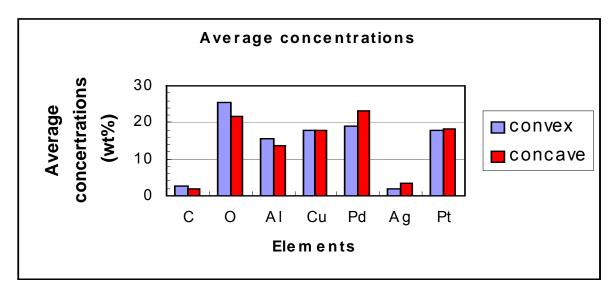


Figure: All concentrations of sample Pd-0.05mm-8mm-36mm

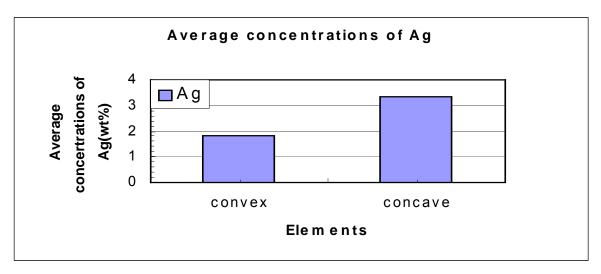


Figure: Ag concentrations of sample Pd-0.05mm-8mm-36mm

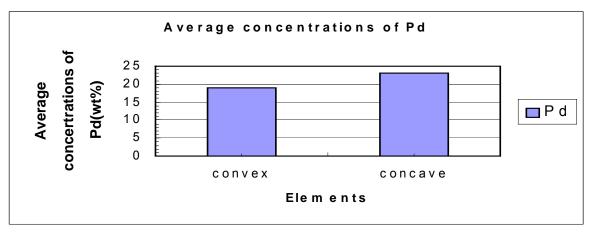


Figure: Pd concentrations of sample Pd-0.05mm-8mm-36mm