

The Nuclear Transmutation Induced by the Electrolysis
Using Tungsten, Rhenium and Gold Electrodes
(Mainly Plasma Electrolysis with Tungsten and Rhenium)

Tadayoshi Ohmori and Tadahiko Mizuno

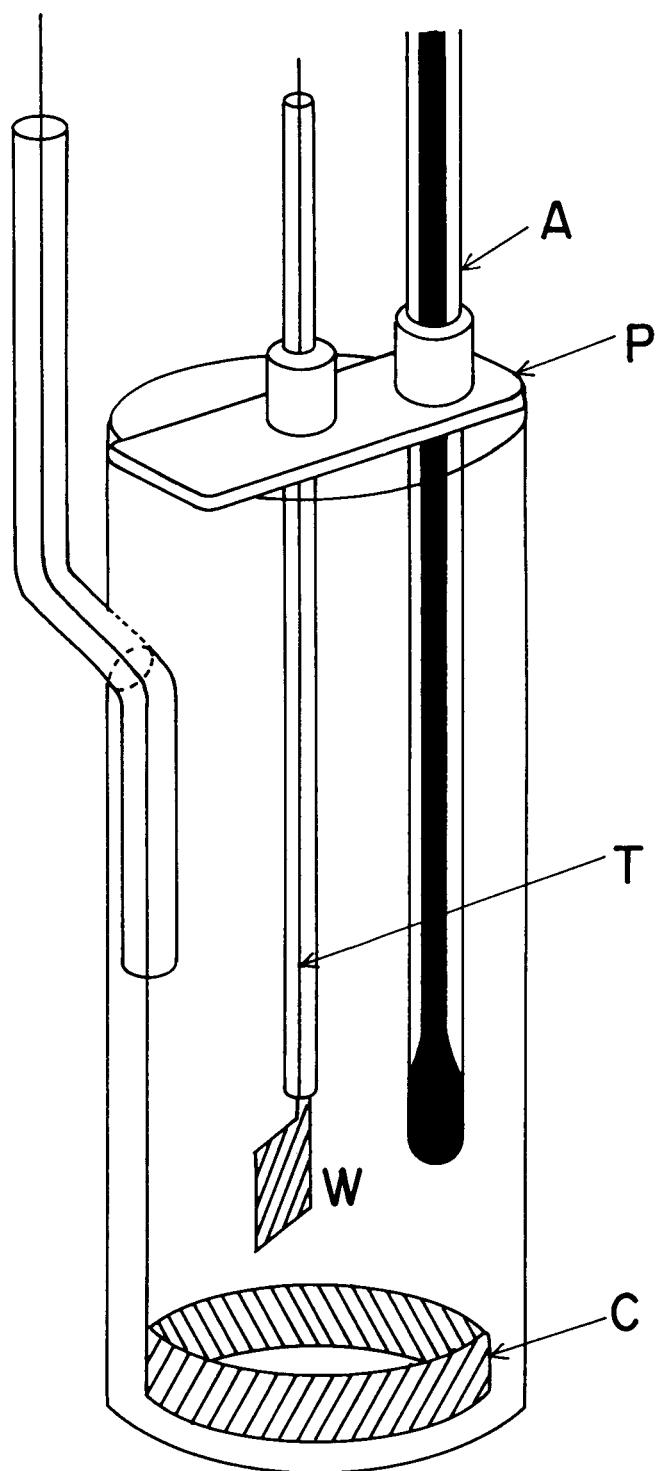
Hokkaido University

DEVELOPMENT

1. Nuclear transmutation in Au / H₂O electrolysis system.
2. Nuclear transmutation in W, Re / H₂O plasma electrolysis system.
3. Nuclear transmutation in Re / D₂O plasma electrolysis system.

RESULTS

1. Excess heat evolution ranging 50 – 150 W.
2. Detection of new elements with anomalous isotopic distribution.
3. Anomalous topography of the electrode surface after electrolysis.



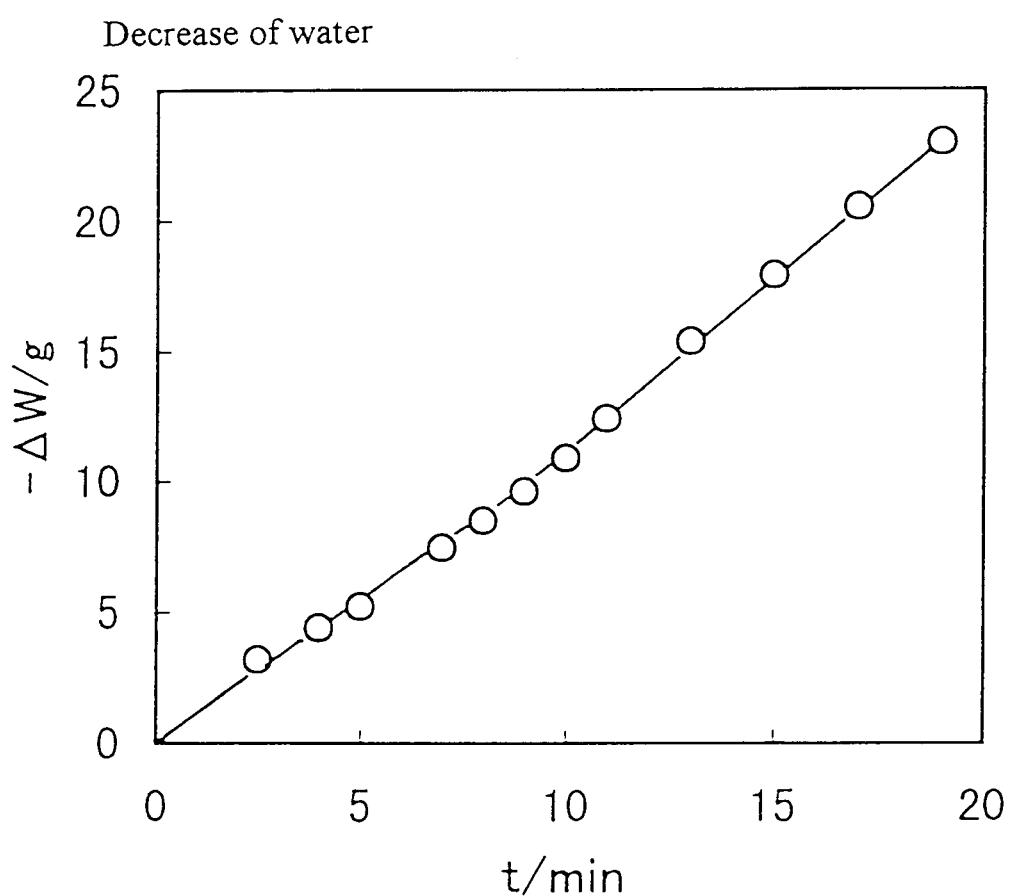
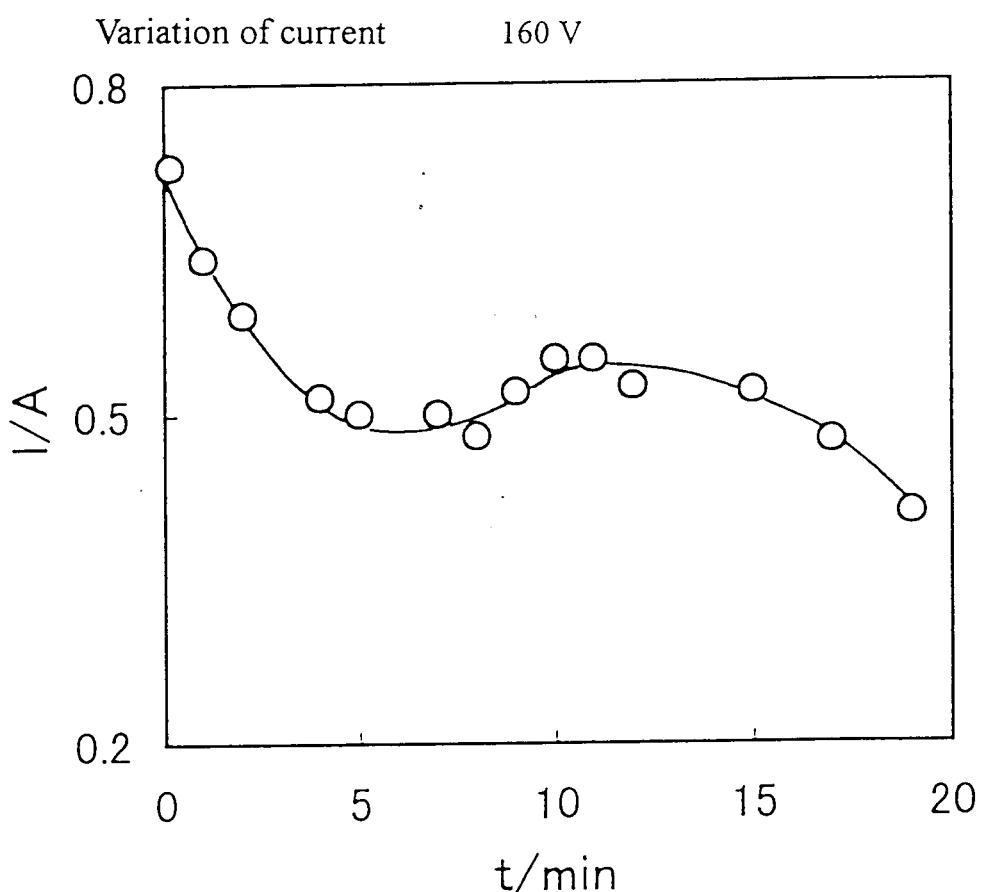
WORKING ELECTRODE UNDER AN INCANDESCENT CONDITION

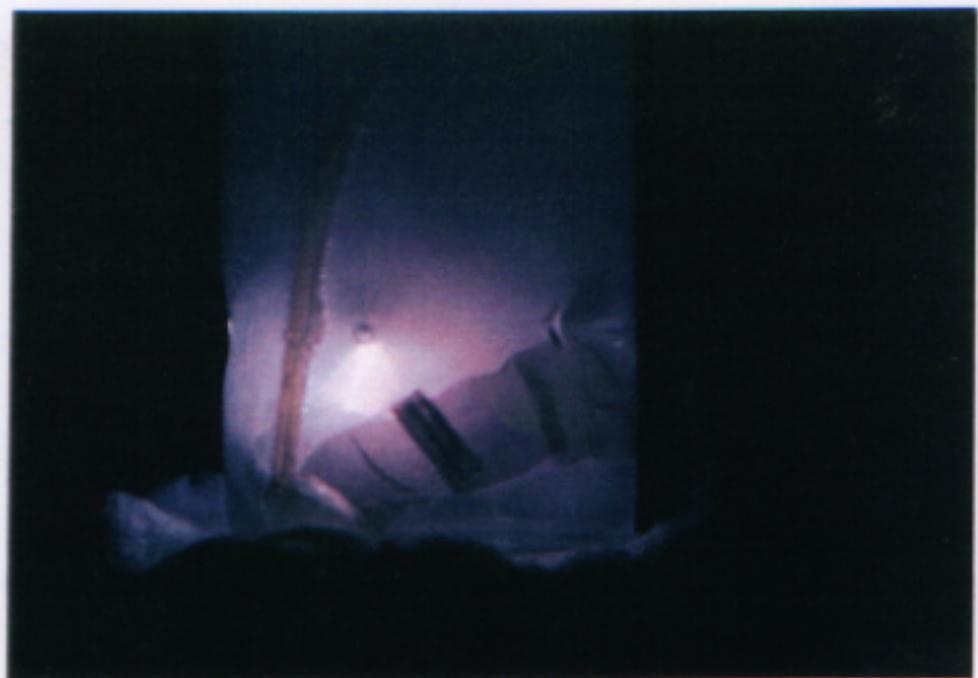


When electrolyzed under this condition, the working electrode was suddenly heated incandescently and reddish-purple glow emerged in the solution around the electrode. A typical incandescent situation of the electrode is shown in this picture. This condition was continued during the electrolysis.

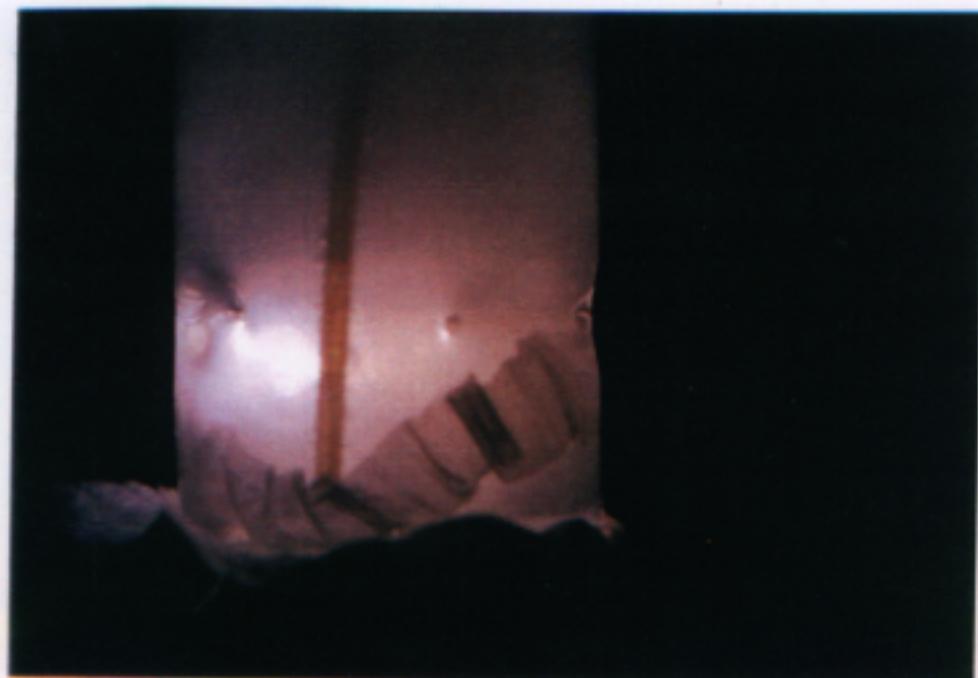
Perhaps, the glow is caused by plasma emerged from the W electrode. It is surprising that the plasma is formed even in such a liquid phase. The temperature of the solution was 85°C just before the ignition. After the ignition, the temperature rapidly increased and reached its boiling point within 80 sec. This phenomenon was easily reproduced. We measure the temperature using both thermocouple and alcohol thermometer. After the ignition, the measurement by the thermocouple becomes impossible owing to the strong emission of the electromagnetic wave, so that we used alcohol thermometer.

Decrease of water in solution after incandescent condition





Wave length of potassium: 404.72, 404.41 nm



Wave length of hydrogen: 636.28, 486. 13, 434.05, 410.17 nm

Wave length of oxygen: 700.22, 645.61, 645.46, 645.37,
436.83 nm

Method of Excess Energy measurement

$$E = W_{\text{vap}} + W_{\text{sol}} + W_{\text{cell}} + W_{\text{wall}} - W_{\text{in}}$$

W_{vap} : heat necessary to vaporize the water of solution.

W_{sol} : heat necessary to rise the solution temperature up to 100°C.

W_{cell} : heat necessary to rise the cell container up to 100°C.

W_{wall} : heat effused through the wall of the cell container during the electrolysis.

W_{in} : input energy (current x input voltage)

Excess Heat Evolved in the H₂O Electrolysis on Tungsten Electrodes

Electrode	Solution	Conc. M	Volume ml	Time min	Hin W	Hout W	Excess H W	Efficiency %
W	Na ₂ SO ₄	0.5	170	30	107	213	106	199
		0.5	170	30	112	196	84	175
		0.5	170	30	104	191	87	184
	Na ₂ CO ₃	0.5	150	30	99	162	63	164
		0.5	150	6	101	224	123	222
	NaNO ₃	0.5	150	23	101	188	87	186
		0.5	150	15	112	197	85	176
	NaClO ₄	0.5	140	15	104	199	95	191
		0.5	120	15	96	174	78	181
	K ₂ CO ₃	0.5	120	15	77	181	104	235
		0.5	170	30	86	196	110	228
	KNO ₃	0.5	170	30	85	174	89	205
		0.5	150	30	85	156	71	184
	Rb ₂ CO ₃	0.5	150	30	71	132	61	186
		0.25	170	30	93	185	92	199
	Cs ₂ CO ₃	0.25	170	30	94	228	134	243
		0.25	150	30	93	162	69	174
	Ba(NO ₃) ₂	0.25	120	15	69	143	74	207
		0.25	150	30	125	195	70	156
	Ba(ClO ₄) ₂	0.25	150	30	71	155	84	218
		0.25	150	15	99	163	64	165
		0.25	150	15	94	205	111	218
					70	147	77	210
					78	173	95	222
					83	188	105	227

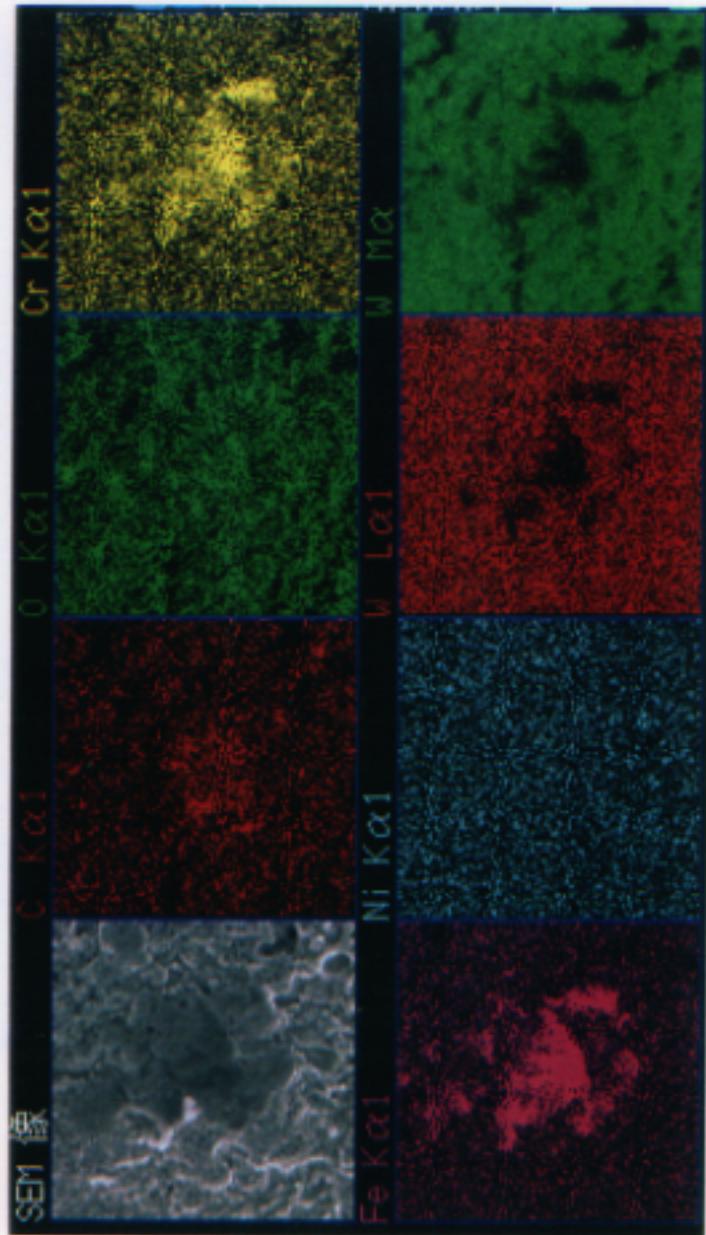
Excess Heats Evolved in the Electrolysis on
Rhenium Electrode

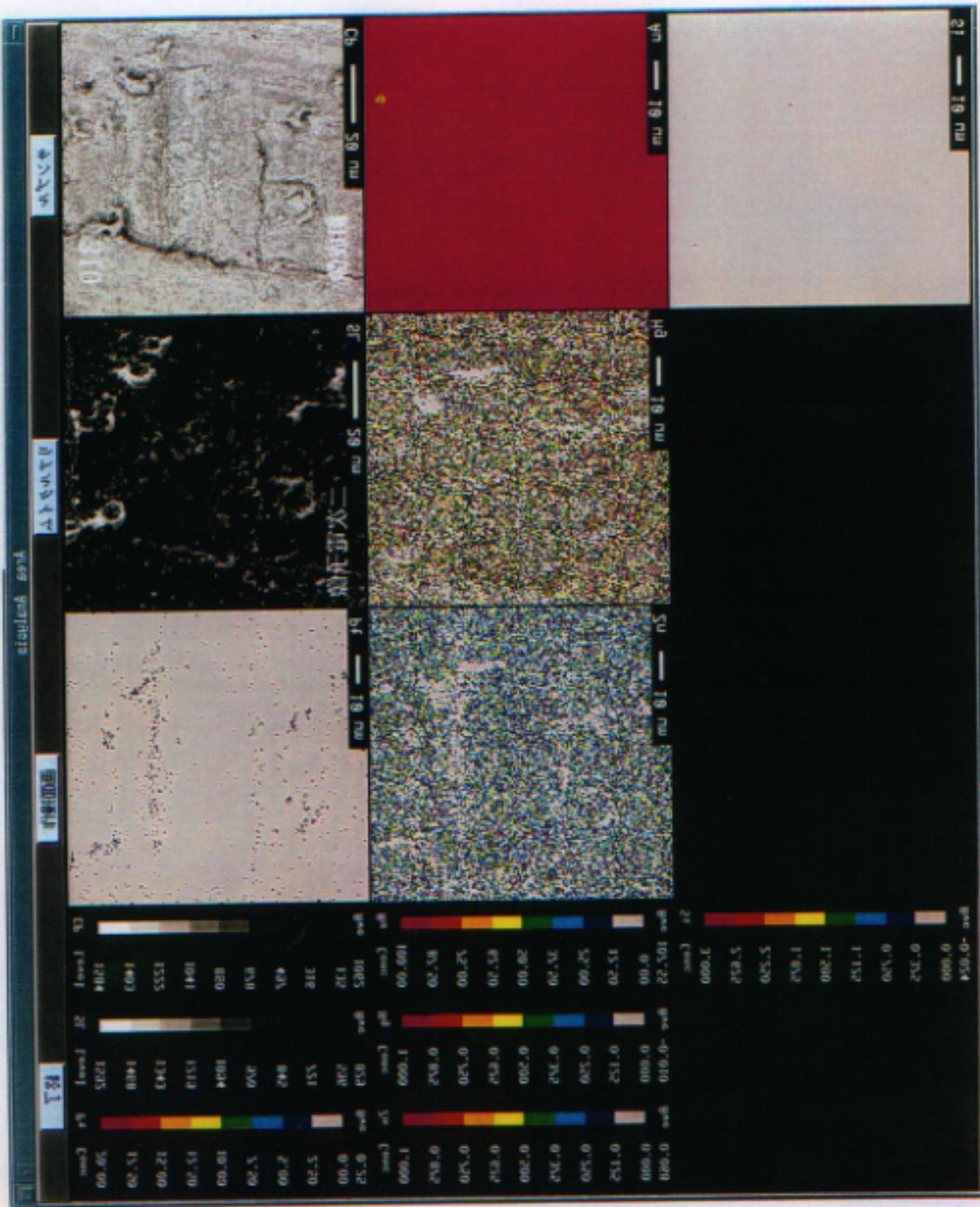
H₂O electrolysis

Electrode	Solution	Conc. M	Volume ml	Time min.	Hin W	Hout W	Excess H W	Efficiency %
Re	K ₂ CO ₃	0.5	150	16	86	137	51	159
		0.5	150	18	82	134	52	163
		0.5	175	7	88	160	72	182
		0.5	175	9	83	170	87	205
		0.5	175	19	85	131	46	154
		0.5	175	10	89	148	59	166

D₂O electrolysis

Electrode	Solution	Conc. M	Volume ml	Time min.	Hin W	Hout W	Excess H W	Efficiency %
Re	K ₂ CO ₃	0.5	150	15	65	58	123	211
		0.5	150	18	78	138	60	178
		0.5	150	14	95	155	60	163
		0.5	150	15	66	128	62	192

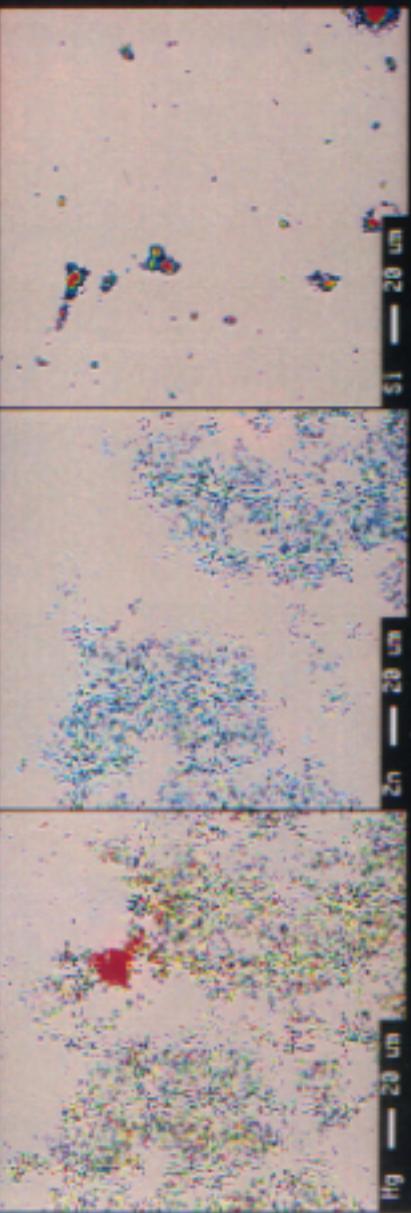
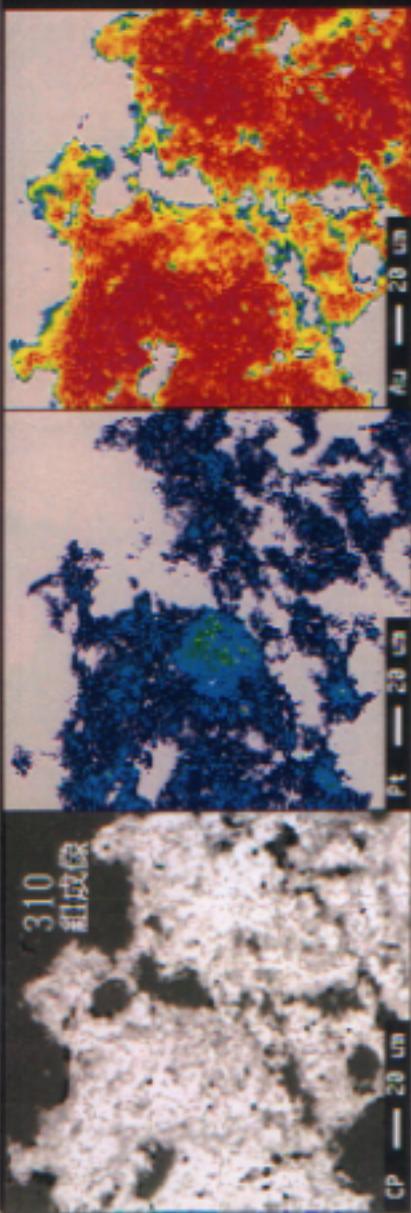




Area Analysis

リバウンド

サンプル



EDS
Pt
Cu
Zn
Hg

20.0 kV
9.75eV/2A
20.00
25.4720
X: 0.90
Y: 0.80

EDS
Pt
Au
Hg
Zn
Hg
Zn
Pt

0.060
0.075
2.125
1.875
0.625
0.550
1.550
0.125
0.050
-0.224
-0.554
-0.013

統計分析

統計分析

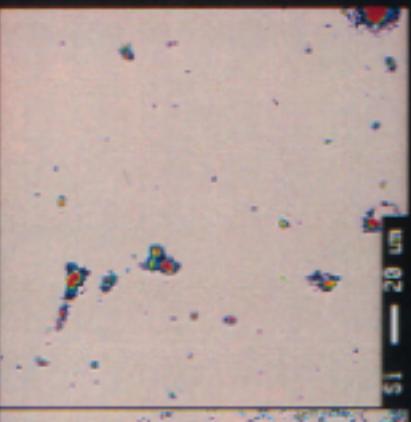
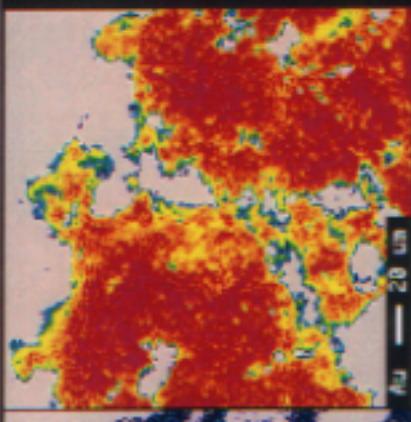
CP Level Pt Cu Au

1675
1762
1523
1355
1153
1019
837

60.69
52.59
45.69
37.59
30.69
22.59
15.69

60.69
67.59
75.69
62.59
50.69
37.59
25.69

12.59
0.09
51.21
2.125
1.875
0.625
0.550
0.125
0.050
-0.224
-0.554
-0.013



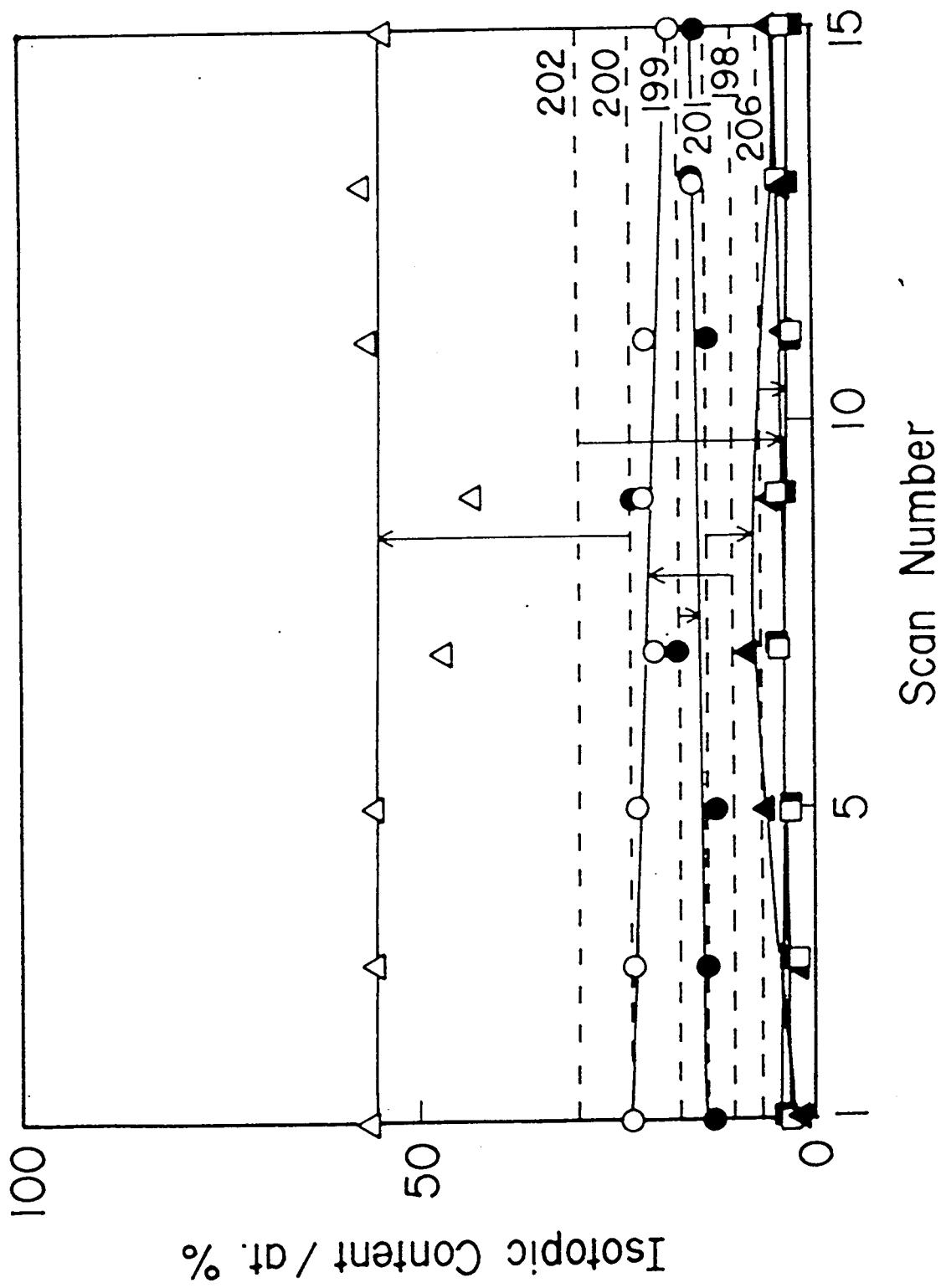
EDS
Pt
Cu
Zn
Hg

20.0 kV
9.75eV/2A
20.00
25.4720
X: 0.90
Y: 0.80

EDS
Pt
Au
Hg
Zn
Hg
Zn
Pt

0.060
0.075
2.125
1.875
0.625
0.550
1.550
0.125
0.050
-0.224
-0.554
-0.013

Isotopic Distribution of Hg Present on a gold electrode
After the H_2O Electrolysis (by SIMS)



Isotopic distribution of Si present in the deposits measured from 1 to 15 scans

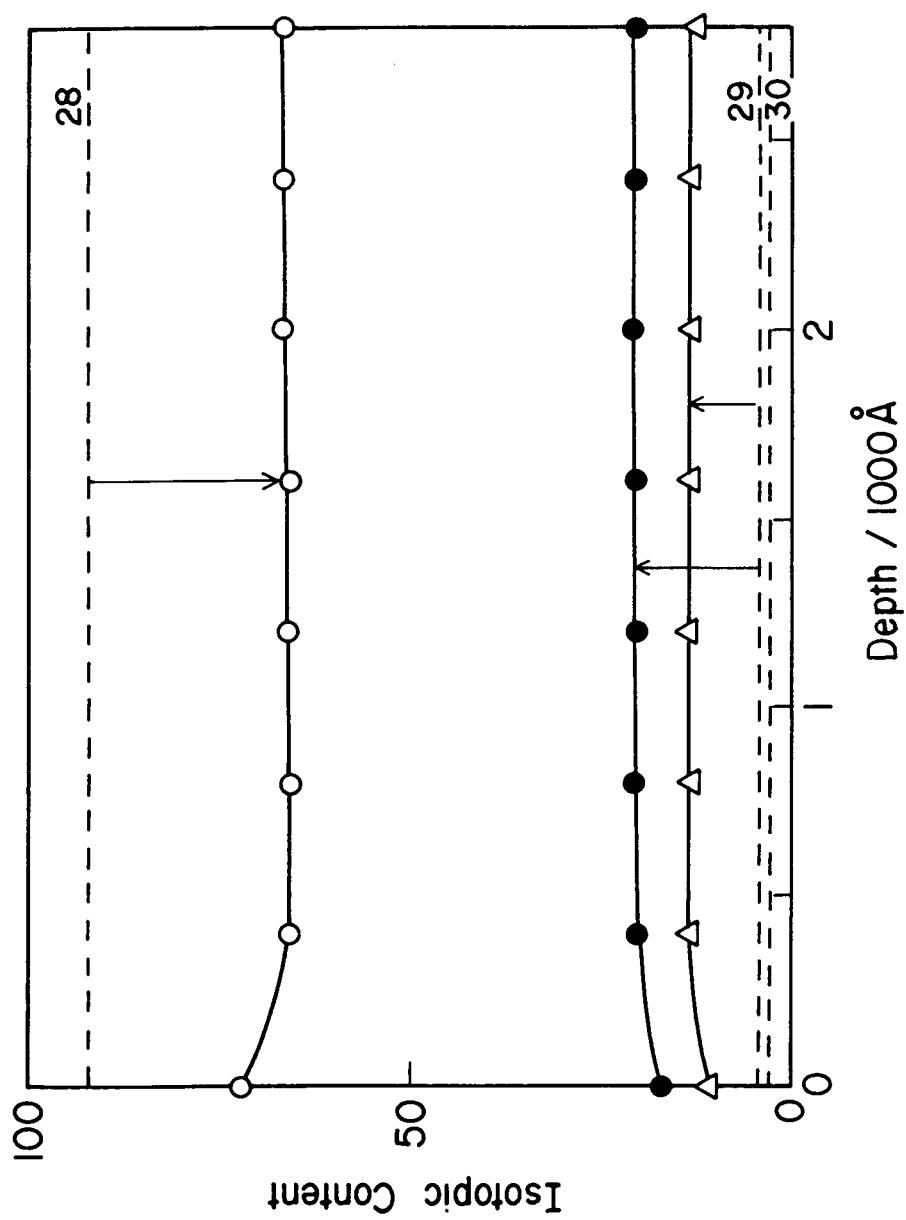
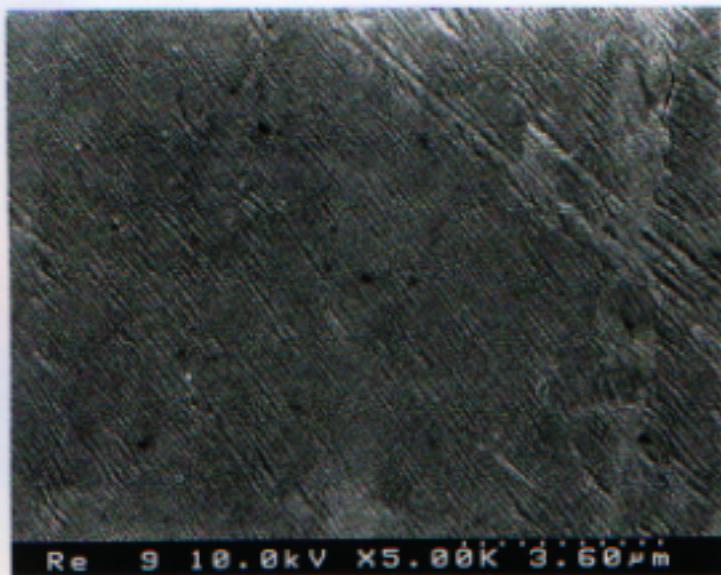
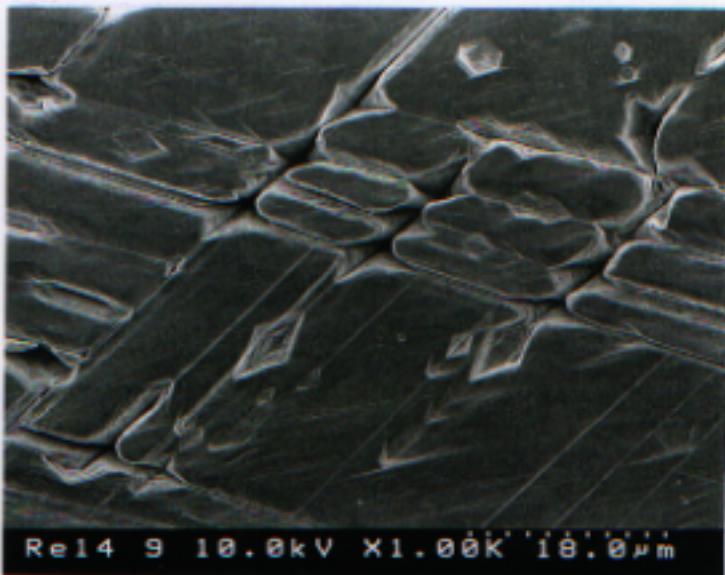
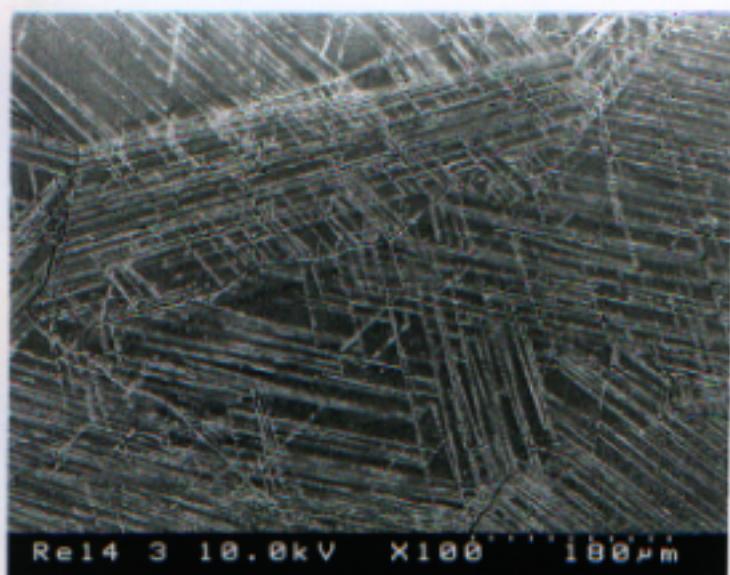
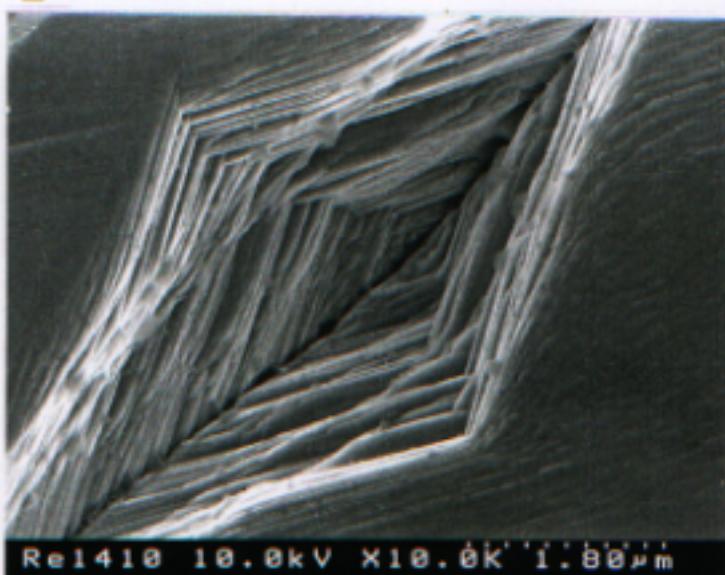
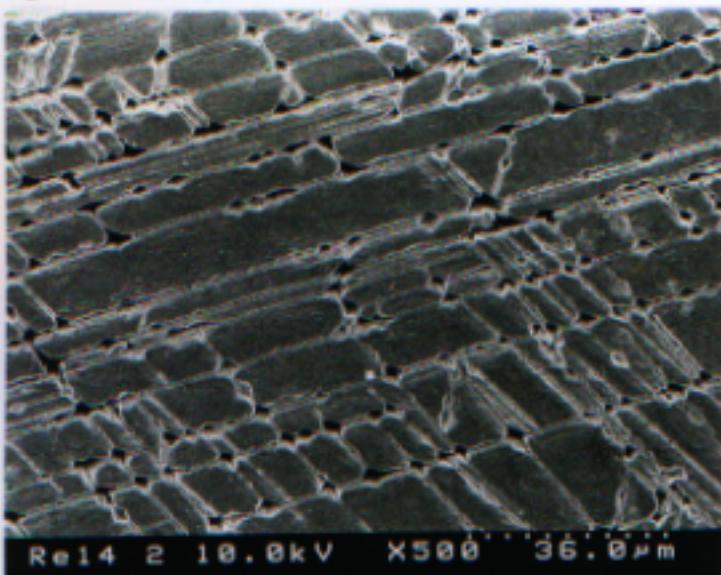


TABLE II
**Isotopic Contents of Product Elements Present on/in the Electrode within
 160 Å from the Electrode Surface and Tungsten of the Electrode Material**

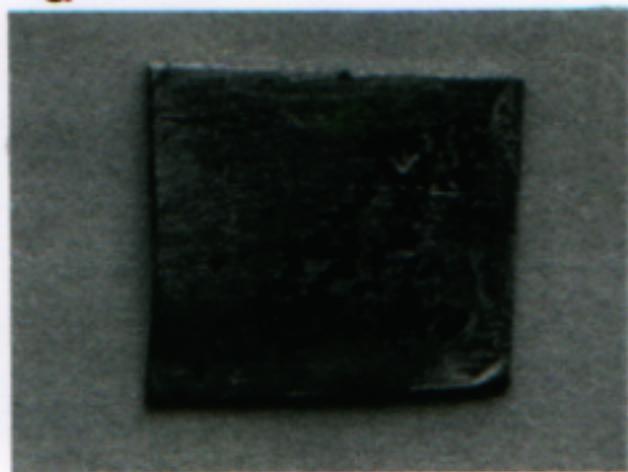
Element	Isotope	Isotopic Content at.%	Natural Isotopic Abundance at.%
Cr	^{50}Cr	6.1	4.31
	^{52}Cr	79.1	83.75
	^{53}Cr	12.4	9.55
Fe	^{56}Fe	91	91.66
	^{57}Fe	2.9	2.19
Ni	^{58}Ni	63	67.98
	^{60}Ni	30.8	26.23
Re	^{183}Re	48.4	37.1
	^{185}Re	51.6	62.9
Pb	^{206}Pb	38.5	25
	^{207}Pb	55	22
	^{208}Pb	6.5	52
W	^{182}W	25.3	26.3
	^{183}W	14.9	14.3
	^{184}W	30.4	30.6
	^{185}W	29	28.6
	$^{182}\text{W}^*$	26.6	26.3
	$^{183}\text{W}^*$	14.5	14.3
	$^{184}\text{W}^*$	30.4	30.6
	$^{186}\text{W}^*$	29	28.6

* Isotopic contents of W within 1700 - 2400 Å from the electrode surface.

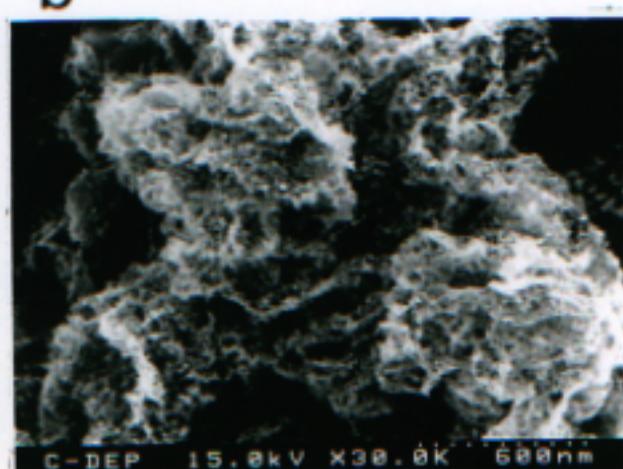
A**D****B****E****C**

Structures of the deposits from the craters of the Au electrode formed during the electrolysis

a



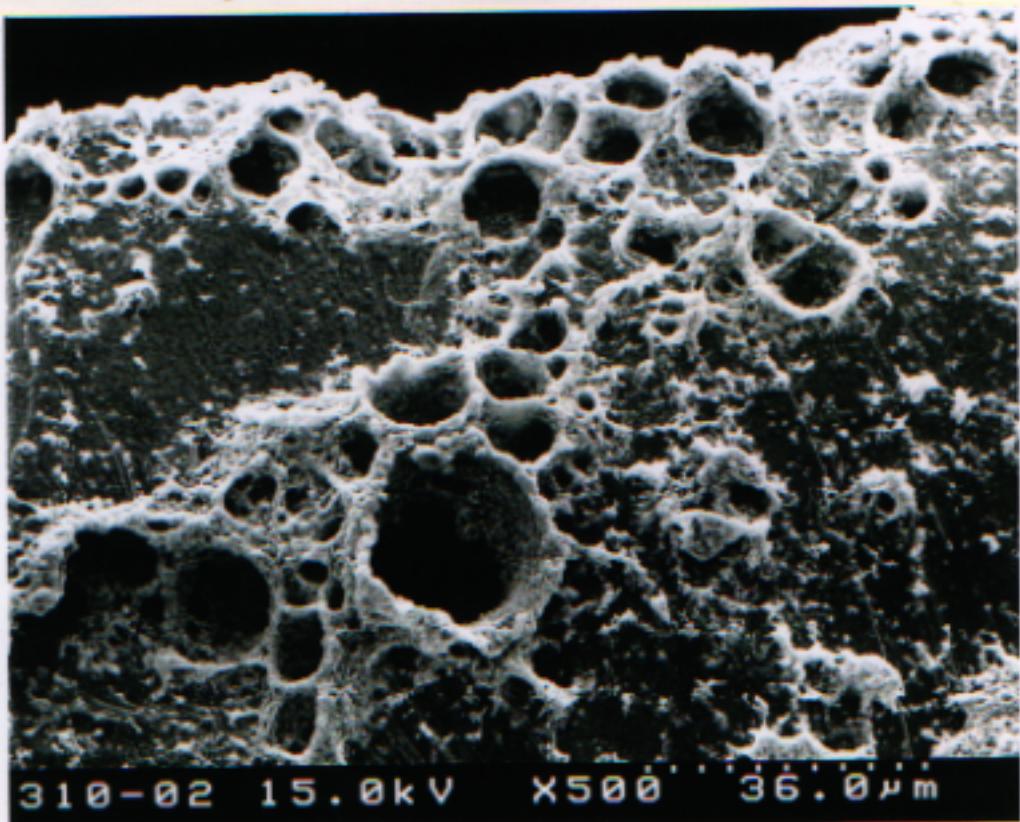
b



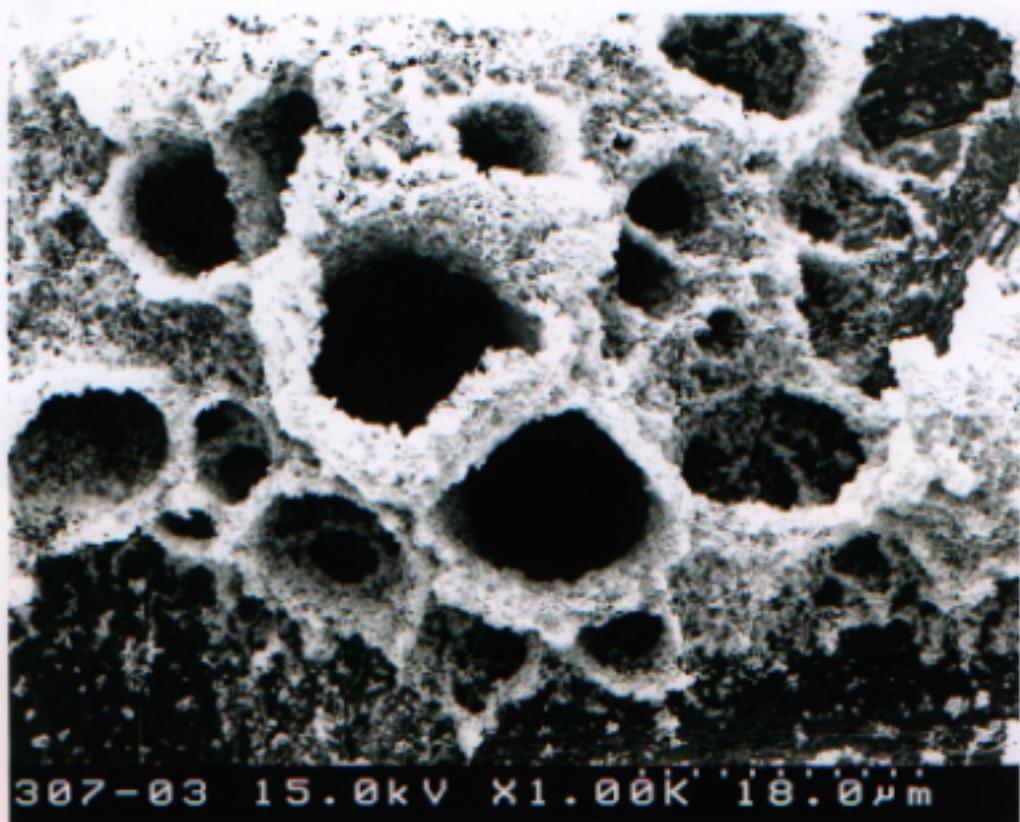
C-DEP 15.0kV X30.0K 600nm

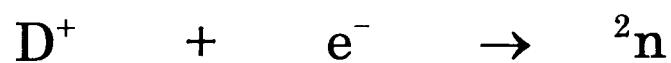
A

The surface of the Au electrode after the electrolysis at a current density of 0.75 A/cm^2



B





Fe



Si

Re Fe



C

Where the energy producing neutron come from? From vacuum ?

Summary

- Excess heat was confirmed during the plasma electrolysis of tungsten electrode.
- Many elements were detected on the electrode and in the electrolyte for the case of excess heat generated experiment.
- Experimental results can be well explained by the photo fission mechanism.