Electrolitic Plasma Cell

Decades of experiences of preliminary evidences

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CONDITIONS

THIS PRESENTATION EXPLAINS EXPERIMENTAL EVIDENCES AND PRELIMINARY CONVERGENCES ON ANOMALIES EMERGING FROM TESTING **ON ELECTROLYTIC PLASMA CELL**

SUCH ANOMALIES ARE TWO TYPES: DETECTION OF NUCLEAR EVIDENCIES AND ENERGETIC ANOMALIES

DESPITE THE EVIDENCES, WHAT WE HAVE OBTAINED IT IS STILL AT A 7 **STAGE (SADLY) PRELIMINARY**

Cell configuration (last configuration)



Plasma cell during a run

Typical aspect during which the breakthrough of electrochemical behaviour is passed and the cell works in plasma mode



Steps to reach Plasma mode

< 100 V input



Hydrogen generation at cathode

 $2H_2O + 2e^- \rightarrow H_2 + 2OH^ 2H_3O^+ + 2e^- \rightarrow H_2 + 2H_2O$ 100 – 200 V input



Discharge (not Faradayan behaviour)

200 – 240 V input



stable plasma layer



conditions depending by cell's parameters > 250 V input



ambiente di plasma attivo

anomalies activity environment

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Funzione Lavoro per Elementi Metallici



plasma environment conditions (>200 V)





On cathodic superface:

- cracks on tungsteno bulk
- monoatomic hydrogen ions locally
- electronic high density
- electric field very high



ignition conditions



Data measurement system



EXPERIMENTAL RESULTS

or

nuclear phenomena
calorimetric anomalies

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In proximity of the cathodic surface:

- a) CR39 detector
- b) dosimeter's scheme assembling
- c) dosimeter



Misura dei neutroni



$CR39 = C_{12}H_{18}O_7$

 $(CH_2=CH-CH_2-O-CO-O-CH_2CH_2-O-CH_2CH_2-O-CO-O-CH_2-CH=CH_2)$ polymer sensitive at α (alpha) emission

boron contained into the H_3BO_3 has got an

natural isotopic distribution :

- $20\% \longrightarrow {}^{10}B$ sensitive at thermal neutrons (E @ 0,025 eV)
- 80% → ¹¹B

¹⁰B + n $\longrightarrow \alpha$ [1.47 MeV] + ⁷Li + γ (93.6%)

 $^{10}B + n \longrightarrow \alpha [2.79 \text{ MeV}] + ^{7}\text{Li}$ (6.4%)

Energetic range for CR39 vs α (alpha) particle's energy

 $\alpha \longrightarrow [0.04 \text{ MeV}; 4 \text{ MeV}]$



BORIC ACID = H_3BO_3

Dosimeter's calibration

A collection of 20 sample s containing CR39 + H_3BO_3 were delivered to the "National Institute of Metrology in Ionizing Radiations" (Casaccia ENEA).

Such samples were **exposed at a calibrated source of thermal neutrons, emitting a flux**

1.2·10² N/mm²·s (0,12 μS/s)

The source is composed by six sources based on Am-Be reaction, covered by graphite and polyethilene .

The exposition was done in 'single blind'. We don't know the sample exposed and the exposition time.





Dosimeter's calibration

In order to obtain a calibration and a reference for compare the measurements done through this dosimetric system, the 20 samples were divided into six groups, exposed at thermal neutron flux according:

- 1. group
- 2. group
- 3. group
- 4. group
- 5. group
- 6. group

- NOT exposed (blank)

- 1' exposure time
- 5' exposure time
- 20' exposure time
- 40' exposure time
- 60' exposure time



Dosimeter's calibration



After etching, the sample is analized through electronic magnifier and a specific software to counts the tracks on CR39 plate after the etching process

Thermostatic chamber for CR39

Dosimeters' calibration (the gauge)



nple code	de Average trac Exposure time density		Group average track	
	(min.)	(tracks/mm ²)	density (tracks/m m ²)	
514	0	5	5	
276	1	5		
284	1	10	7	
678	1	6		
550	5	10		
310	5	12	9	
142	5	8		
117	5	8		
121	20	18	28	
120	20	37		
113	40	23		
311	40	36	32	
143	40	36		
262		40		
321	60	41	47	
565	60 60	53		
620) 60	55		

Dosimeters' calibration (the gauge)

average trend





operative conditions





Dosimeter after a run very close at cathode



Tracks comparison (CR39)

average trend vs exposed



IMPORTANT NOTES

- The neutron detection method based on CR-39 nuclear track detectors, coupled with a boron converter, has demonstrated neutron generation by plasma discharge in an electrolytic cell with alkaline solution.
- A significant number of tracks were revealed by the CR-39 detector samples 7 positioned in close proximity to the plasma discharge, next to the tungsten cathode of the electrolytic cell.
- the blank detectors show no tracks, if positioned far from electrolytic cell. 7

NUCLEAR TRANSMUTATIONS

ICP analysis of electrolithic solution – 07/2012



methods



- Test done using the same electrode (broken in 2 parts) on two tests,
- The first one in a plasma test
- The second one in a simple electrolysis test (same electric energy for both run).
- After each test, the cathode was dissolved in solution using electrolytic

ICP analysis of electrolithic solution (I)-07/2012



ICP analysis of electrolithic solution (I)– 07/2012



Several important increasements of composition, in terms of percentage and in terms of quantities, but also important decreasements, of chemical species (like indium, but also nikel, rhenium, osmium, hafnium, platinum, gold, mercure, germanium, arsenicum and selenium) not comparable to the increasing of new chemical elements after plasma action.

analisi ICP della soluzione elettrolitica (II) – 06/2012

I valori di concentrazione espressa in parti per milione degli elementi richiesti sono riportati nella tabella seguente.

	Re	Os	Au	Pt
A [ppm]	0.13	0.26	2.59	0.47
Std dev	0.0145	0.0232	0.0215	0.00275
B [ppm]	0.004 (tracce)	0.06	Inf. limiti di detezione	Inf. limiti di detezio
Std dev	0.0023	0.0299		

Oro e platino nel campione "B" risultano inferiori al limite di rilevabilità dello strumento.

ICP analysis independent, performed by an important Italian research institution showed similar abnormalities in the composition of the solution that emerged after the step of plasma prolonged.



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ALMA UNIVERSITAS TAURINENSIS



Università degli Studi di Torino DIPARTIMENTO DI SCIENZE DELLA TERRA

ENERGETIC ANOMALIES

Previous experimental campaign





All previous measurements were affected by errors related to the underestimation of the input energy due to the inadequacy of the instruments in the detection of all components induced plasma, mainly in the high frequencies





The cell was equipped with a stabilized power supply and noise filtration system. The cell and its connections are enclosed in a Faraday cage.



INSIEME FILTRI PASSA-BASSO 50 KHz - 20 MHz



(blu = 50kHz, black = 500kHz, green = 20MHz, red = risultante)

filters d = risultante)





All the measurements performed so far with this system are constrained from having a flow rate of the cooling fluid set by the circulation pump and the thermal exchange between the fluids of measurement through the glass. These two features represent two substantial limitations:

The last correction to be made to the measuring system cell is the implementation of the measuring system calorimetric which represents a major limitation in the analysis of the cell to the low response speed

impossibility of control of the T cell (thus the plasma)

calorimetric measurement times with <u>'shift' with respect to an</u> <u>electrical stimulus</u> due to the characteristics of the glass

Despite the limitations imposed (obligation to long-term tests, no control on the T cell, no check on the flow rate) which determine the scatter of the calorimetric anomalies of short duration, they are measured anomalies that require the need for system analysis.



Future

- The first preliminary measurements and the first results are NOT definitive. Its are just PRELIMINARY evidencies for a long and deep investigation.
- From the point of view calorimetric the way is already marked and characterization of the plasma will be performed through the steps already described relative to electrical measurements, and calorimetric.



NOT definitive. Its are ation. ed and e steps already stric.

GRAZIE