

# Excess Heat Triggered by Different Current in a D/Pd Gas-loading System

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**Abstract--** In order to study the relationship between the triggering current, deuterium pressure and the excess heat, a series of experiments were made in a D/Pd gas-loading system. By comparing the system constants ( $k=\Delta T/\Delta P$ ) in both nitrogen and deuterium atmosphere we found an optimum current (8A) and a deuterium pressure ( $9\times 10^4$  Pa) in which the system could release a maximum excess power (more than 80W). The reproducibility was 16/16 and the excess energy released in the longest experiment was about 300MJ within 40 days, which was corresponding to  $10^4$ eV for each palladium atom. Analysis of the palladium surface with a scanning electron microscopy (SEM) and an energy dispersive spectrometer (EDS) revealed that some new surface topographical feature with concentrations of unexpected elements (such as Ag, Sn, Pb and Ca) appeared after the current triggering. The results implied that the excess heat might come from a nuclear transmutation.

**Key words--** current triggering, deuterium pressure, D/Pd gas-loading system, excess heat

## I. INTRODUCTION

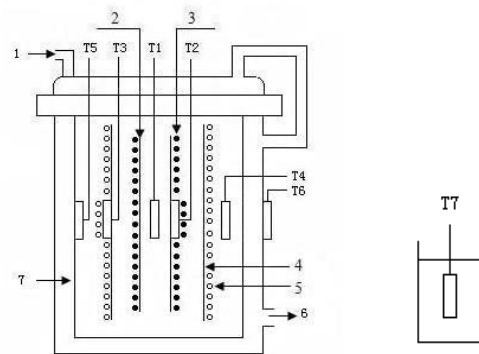
S. Focardi et al [1] reported on the existence of a large excess heat production observed in a H/Ni gas-loading system. A. Rossi [2] developed the method and enlarged the phenomenon into 1MW power generation device afterwards. In recent days few of people knew Rossi's secret catalyzer and many had skeptics and questions on the demonstration reactor. With a curiosity to that event a D/Pd gas-loading system was chosen and a series of similar experiments were made in order to prove if the evidence could be true and if the excess heat could be also occurred in other different system.

## II. EXPERIMENTAL

### A. Materials and Apparatus

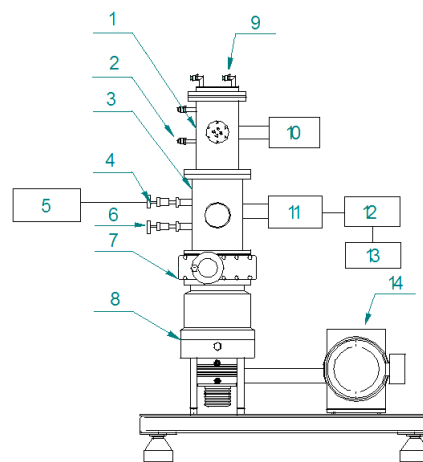
Figures 1 and 2 gave a schematic of the experimental system. Figure 1 was a reaction chamber that made up of stainless steel with a double-jacket structure, where the circulating water could flow through. It has internal dimension of diameter =100mm and high = 240mm with useful capacity of about 1.9 L. Two Pd wires were 99.98% in purity (made by General Research Institute of Nonferrous Metals, Beijing), one of them was for being triggered with the dimension of 0.5mm in diameter and 210cm in length ( $V=4.1\times 10^{-1}\text{cm}^3$ ). And the other was for triggering with the same size in diameter and 400cm in

length ( $V=7.854\times 10^{-1}\text{cm}^3$ ). In order to monitor the temperature at different positions continuously, Seven Pt100 resistor thermometers were placed inside and outside the reaction chamber. The chamber structure as shown in figure 1:



where, 1 was the inlet of circulating water; 2,4 ceramic tube; 3 Pd wire for triggering; 5 Pd wire for being triggered; 6 outlet of circulating water; 7 double-jacket;  $T_1\sim T_7$  Pt100 resistor thermometers were used for taking the temperatures in different position:  $T_1$  the center of the chamber inside the ceramic tube;  $T_2$  wound on triggering Pd wire;  $T_3$  wound on being triggered Pd wire;  $T_4$  inside the chamber between inner wall and outer wall of the ceramic tube;  $T_5$  inner wall of the chamber;  $T_6$  outside wall of the chamber;  $T_7$  ambient temperature.

The apparatus connected to the reaction chamber was shown as in figure 2:



where 1 was the chamber; 2 the inlet and outlet for circulation water around the chamber; 3 transition chamber; 4 D<sub>2</sub> needle valve; 5 D<sub>2</sub> generator; 6 air-released valve; 7 gate valve; 8 turbo-molecular pump; 9 the inlet and outlet mouth for water circulation on the top; 10 DC power supply; 11 vacuum gauge; 12 Keithley 2700 multifunction data-inquisition meter; 13 a computer for data recording and controlling; 14 mechanical pump.

### B. Calibration

Two palladium wires for triggering and being triggered were placed in the reaction chamber respectively. At the beginning some natural nitrogen gas was introduced into the chamber by four steps of  $P_{N_2}=20, 1 \times 10^4, 5 \times 10^4$  and  $9 \times 10^4$  Pa. In each step, an initial current of 1A and at a regular increase ( $\Delta I=1A$ ) were passed through the triggering Pd wire. The maximum current was 8A. Then the relation between the temperature increases and different input power was obtained. The same process was done exactly when the nitrogen was replaced by deuterium. The promising result was occurred in  $P_{N_2/D_2}=9 \times 10^4$  Pa and calibration were shown in figure 3 and 4:

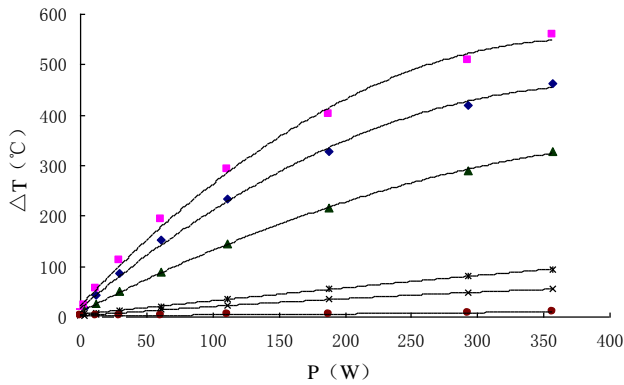


Fig. 3 Calibration curves: temperature (relative to the T7) vs. power under N<sub>2</sub> environment at  $9 \times 10^4$  Pa. (T1(◆), T2(■), T3(▲), T4(×), T5(\* ) and T6(●))

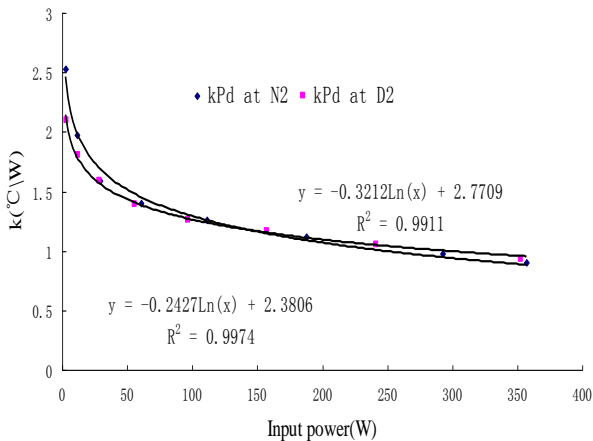


Fig. 4 The fitting curve of input power and  $k$  under N<sub>2</sub>(◆),D<sub>2</sub>(■) environment at  $9 \times 10^4$  Pa

### C. Current triggering

From the data, the correlation between input power and  $k=\Delta T/\Delta P$  [3] on being triggered Pd wire in N<sub>2</sub> and D<sub>2</sub> were fitted. The fitting curves were as shown as in figure 4. From it we can see that the value of  $k$  in N<sub>2</sub> atmosphere was higher than that in the D<sub>2</sub> when input power was less than 150W. But when the input power was more than 150W, the value of  $k$  in D<sub>2</sub> was higher than that in the N<sub>2</sub> at high temperature. And with the input power increasing, the difference between  $k$  in D<sub>2</sub> and  $k$  in N<sub>2</sub> became larger. However, taking into account the specific circumstances of laboratory equipment and the palladium wire, the triggering current of 8.0A and  $P_{D_2}=9 \times 10^4$  Pa were chosen in the whole experiment. The triggering result was shown as in Figure 5:

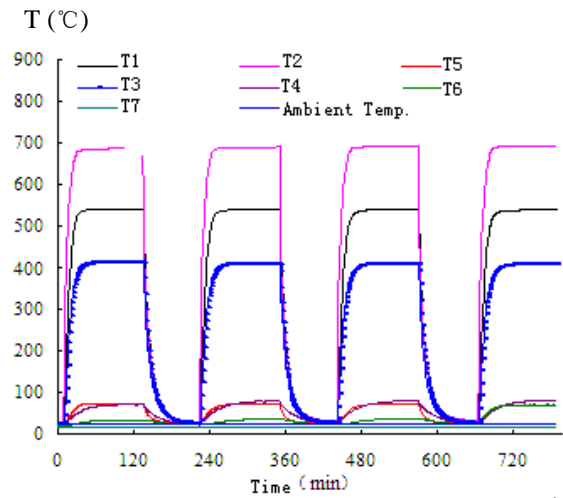


Fig.5 Current triggering results at 8A in  $P_{D_2}=9 \times 10^4$  Pa

The drawing in figure 5 was only the first four times' triggering in our experiment. The total was as many as 16 times. There were 80 90 W excess power in these experiment. The longest one was lasted for 40 days. It looked like the experiment could keep going on as long as possible.

## III. RESULTS AND DISCUSSIONS

### A. Calculation for Excess Heat

The first purpose of this experiment was to investigate whether the excess heat could also appear in D/Pd gas-loading system by current triggering as in that in Focardi's H/Ni system. The second was whether the excess energy obtained in the system could be higher than chemical heat triggered by different electric current when deuterium atoms were charged into metal palladium lattice. Let  $P_i$  be the electrical power supplied into the reaction chamber. At the thermal equilibrium the difference between the  $T_6$  and  $T_7$  could be measured. If  $k$  is the system heat equilibrium constant which was determined by the rise of temperature with a unit power input into the system, the following relation must be verified:

$$P_i = k(T_6 - T_7) \quad (1)$$

If some kind of physic-chemical process occurs into the system and palladium wire is the related enthalpy variation, the following relation holds:

$$P_i + \sum_{i=1}^n \frac{dH_i}{dt} = C \frac{dT_5}{dt} + k(T_6 - T_7) \quad (2)$$

where  $C$  is the heat capacity. Equation (2) is valid if the characteristic times of the heat exchanges inside the reaction chamber are shorter than  $\tau = C/k$  (chamber time constant). If all the physic-chemical processes last for a limited period before stopping, equation (2) reduces to equation (1). In the case for which one process excess heat power ( $P_o$ ) does not stop and  $P_o = dH_j/dt \rightarrow \text{constant}$ , on assuming quasi-stationary conditions, that is, the temperatures of the chamber remain practically constant during a few chamber time constant. The excess heat power can be calculated by equation (3).

$$P_o = P_i - \frac{T_6 - T_7}{k} \quad (3)$$

According to equation (3) the excess heat power could be calculated as in Table 1:

TABLE I  
8A CURRENT TRIGGERING RESULTS IN  $P_{D_2}=9 \times 10^4 \text{ Pa}$

Triggering	$P_{\text{input}}$ (W)	Loading radio	Equilibrium temperature ( $^{\circ}\text{C}$ )	Excess heat power (W)
0	-	0.094	23.959 $\pm 0.003$	-
1	416.875 $\pm 0.011$	0.018	409.138 $\pm 0.022$	90.629 $\pm 0.002$
2	416.689 $\pm 0.015$	0.029	407.166 $\pm 0.037$	88.380 $\pm 0.015$
3	416.778 $\pm 0.019$	0.092	406.408 $\pm 0.156$	87.399 $\pm 0.019$
4	416.71 $\pm 0.004$	0.032	405.532 $\pm 0.022$	88.217 $\pm 0.004$

The longest triggering experiment lasted for 40 days. The average excess heat power was more than 87W, the total excess heat energy was about 300MJ, which was corresponding to  $1.8 \times 10^4 \text{ eV/atom Pd}$ . Apparently it was more than the energy for each Pd atom released in a chemical process.

### B. SEM and EDS Analysis

The SEM image of original Pd wire before triggering was shown in figure 6. From it many scratches on Pd wire surface could be seen due to its industrial production. The SEM of Pd wire charged by deuterium for 5 times was given in figure 7. Where many small cavities on Pd wire surface occurred. Comparison to the figure 6 and 7, Pd wire had an apparently morphological difference in the surface before and after deuterium loading and current triggering. Because a certain number of deuterium atoms went into the Pd lattice, many cavities formed on

the surface. These cavities or channels, made it easy that deuterium atoms charged into the Pd lattice. This is the reason why deuterium charging repeatedly into Pd became more easily afterwards.

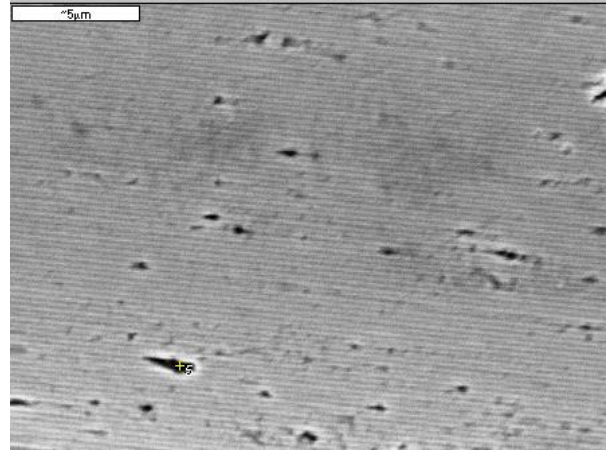


Fig. 6 SEM of palladium wire before triggering (400 $\times$ )

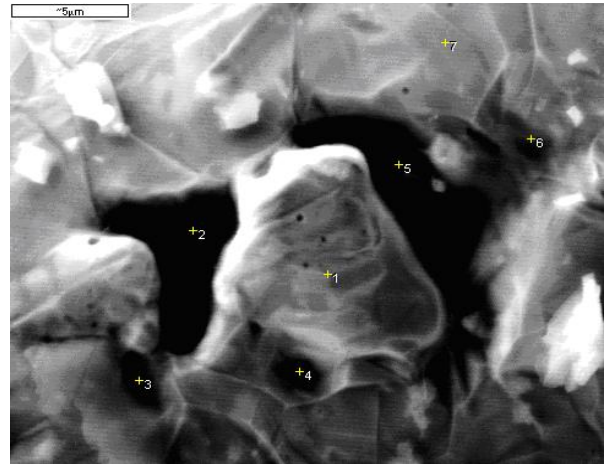


Fig. 7 SEM of palladium wire after triggering (400 $\times$ )

The original Pd wire was analysed with an EDS (figure 8). The result indicated that the purity of sample was of nearly 100%. The EDS analysis of palladium wire after being triggered was shown in figure 9. It is very clear that new elements of Ag, Pb, Sn and Ca were appeared after the triggering process. And it is necessary to invested how they did form on the palladium surface or inside the body. These elements might be produced during some transmutation process.

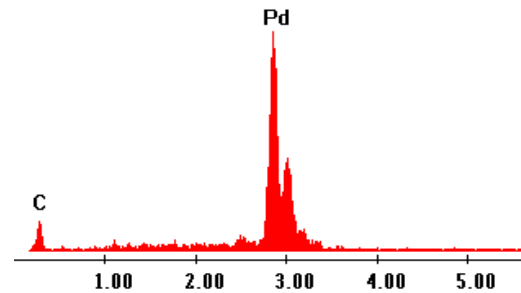
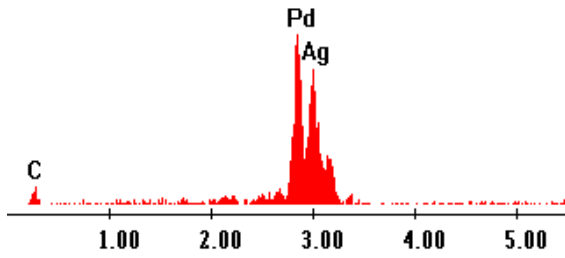


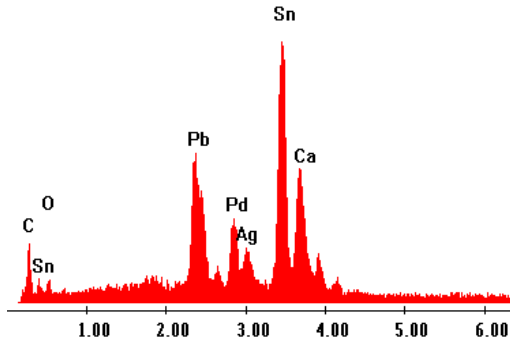
Fig. 8 EDS of Pd origin sample before triggering

## REFERENCES

- [1] S. Focardi, V. Gabbani, V. Montalbano, "Large excess heat production in Ni-H systems," *J. IL Nuovo Cimento*, vol. A111, pp. 1233-1242, November 1998.
- [2] Information on <http://ecatnews.com/?p=1160> /2011/10/30
- [3] J.Tian et al, "'Excess Heat" and "Heat after Death" in a Gas-loading Hydrogen/Palladium System," The 9th International Conference on Cold Fusion, Condensed Matter Nuclear Science (ICCF-9, 2002), Beijing (China), May 2002.



(a) Element Ag was found (point 2)



(b) Element Pb, Sn, Ca were found (point 5)

Fig. 9 EDS of Pd sample after triggering

## IV. CONCLUSIONS

Being triggered by current large amount of excess heat could be produced in D/Pd gas-loading system as same as in H/Ni system. Under the condition of bellowing an atmosphere, the higher of deuterium pressure, the larger of triggering current, the better the triggering effect. By this way even larger excess heat could be obtained under some other conditions (such as 2atms, 500 °C). The reproducibility of our experiments was good as 16/16 in the conditions of 8A triggering current and  $9 \times 10^4$  Pa deuterium pressure. 300MJ excess energy was obtained within forty days, which was corresponding to the energy of  $1.8 \times 10^4$  eV for each Pd atom. This is clear that gratitude was much higher than the energy for each Pd atom released in a chemical process. And on the surface of Pd after being triggered some new elements were found by SEM and EDS analysis. This phenomenon is need further studying and its origin might come from a nuclear transmutation process.

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