# Observation of macroscopic current and thermal anomalies, at high temperature, by hetero-structures on thin and long Constantan wires under H<sub>2</sub> gas

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#### Abstract

Since 2011, we introduced in LENR Research field the use of a Constantan alloy to absorb and adsorb proper amounts of  $H_2$  or  $D_2$  (concentrated and/or mixed with noble gases of low thermal conductivity) and to generate thermal anomalies even at low temperatures (>200°C wire temperature). Based on this idea, we developed a reactor with a core of sub-micrometric layered Constantan wires that produced measurable excess power and showed result with some reproducibility. During the years, we modified this base configuration with the purpose of improving both the reproducibility and the Anomalous Heat Effect (AHE). We used fiberglass sheaths for ensuring electrical insulation and found out, by chance, that this material even improves the performance of the reactor. In the most recent configuration, we studied the effects of the addition of Fe nanolayers to Constantan wires and of several small knots along their extension, resulting in a larger excess power growing with wire temperature. Finally, we detected a new anomalous electric effect, consisting in the generation of a spontaneous voltage between the ends of a floating wire in the reactor, enhanced and stabilized by Fe presence.

*Keywords*: calorimeter, Constantan, sub-micrometric surfaces, H<sub>2</sub> adsorption/absorption, iron, second law of thermodynamics, LENR.

# 1. Constantan alloys as H2-dissociation catalyzers

Our investigations concerning the ability of metals such as Palladium (Pd) and Nickel (Ni) to absorb  $D_2$  and  $H_2$ , in connection with anomalous heat generation at high temperatures received a new impulse with the introduction of the Constantan alloys in the LENR research field in 2011.

Our original idea was to individuate a low-cost material able to replace the very expensive (and mechanically weak) Pd in LENR experiments.

	ΔE (eV)
Ni <sub>0.3750</sub> - Cu <sub>0.6250</sub>	+3.16
Ni <sub>0.6250</sub> - Cu <sub>0.3750</sub>	+2.86
Ni <sub>0.8125</sub> - Cu <sub>0.1875</sub>	+2.10
Ni	+1.74
Ni <sub>0.1825</sub> - Cu <sub>0.8175</sub>	+1.57
Ag <sub>0.8125</sub> - Pd <sub>0.1875</sub>	+0.57
Ag <sub>0.625</sub> - Pd <sub>0.375</sub>	+0.51
Ag <sub>0.1875</sub> - Pd <sub>0.8125</sub>	+0.51
Pd	+0.42
Cu	-1.11
Ag	-1.42

Table 1: Catalytic power of different metals and alloys with respect to the reaction  $H_2 \rightarrow 2H$ , computed in Density Functional Theory [14].

We pointed our interest to the family of (Copper-Nickel) Constantan alloys as materials that could fit our purposes because of their ability to dissociate molecular Hydrogen [1]. In particular, we selected a low-cost commercial material called ISOTAN44, with atomic composition  $Cu_{55}Ni_{44}Mn_1$  (Isabellenhütte Heusler, Germany). Together with a measurable H<sub>2</sub>-diffusion coefficient at high temperature, this material offers good mechanical resistance against the aging effects of the thermal cycles and the H<sub>2</sub> absorption/desorption. Moreover, it has very large values of (calculated) catalytic power with respect to hydrogen dissociation, showed in Table 1.

We demonstrated experimentally that Constantan at nano/micrometric size and at low temperatures (T >120 °C, in comparison with about 2000 °C for Tungsten) is able to catalyze the dissociation reaction  $H_2 \rightarrow 2H$  and absorb/adsorb (at least) atomic Hydrogen even inside the

bulk of the lattice, apart surface.

For our experimental set-up we employed Constantan wires of length L = 100 cm and diameter  $\Phi = 0.1 - 0.2$  mm (Figure 1). To increase their effective surface available for catalytic processes, the wires were subjected to specific thermal/electric treatments that created sub-micrometric and multilayered structures at the surface and deeper in the bulk (Figure 2). The sub-micrometric structures were just created by oxidation, with a threshold temperature of 600 °C in free air. Such structures are vaguely similar to hetero-structures.



View Mad. 397.6 µm Det SE Date(m/dy): 05/17/11 100 µm NEXT - LNF - INFN Figure 2: SEM image of the cross section of the constantan wire as provided by Isabellenhütte Heusler. The plastic cover of the wire is visible as the light area rounding the wire.



SEM MAG: 58.18 kx Det: SE Vac: HIVac Date(m/d/y): 04/20/12 1 µm NEXT - LNF - INFN Figure 1: SEM image of the constantan wire surface at micrometric scale after repeated high-power pulse treatments.

The treatment includes electric high peak power pulses (20 kVA/g of material) with a rise time  $T_r < 1 \mu s$ , corresponding to a current density J > 50 kA/cm<sup>2</sup> even neglecting skin effects. Such pulses induce extremely fast thermal treatments (warming $\rightarrow$ cooling) and shock waves. A rough evaluation (by fast photo-camera) of light color emitted from the wire revealed a surface temperature even larger than 1000 °C in some specific tests. At the end of the process we obtained glassy materials at the surface.

The treatment just described produces sub-micrometric geometries, a sort of chaotic mixture of Ni, NiO, Cu, CuO,  $Ni_xCu_yO_z$ , reducing/avoiding, at the same time, the usual (deleterious) self-sintering processes due to the high temperatures. SEM observations revealed that the wires so treated had a large number (up to 700 in some samples) of multilayered structures with thickness of 20 -100 nm.

Our treatment was inspired by "*Melt Spinning and Quenching*" metallurgical process, largely used by Prof. Yoshiaki Arata and his Collaborators (Osaka, Tohoku Universities-Japan) to produce *nanomaterials* (Pd, or  $Pd_xNi_y$ , both dispersed into a matrix of  $ZrO_2$  at 65% concentration) for his Solid State Fusion devices under interaction with pressurized (up to 60 atm)  $D_2/H_2$  at 150-300 °C. In the quenching process their cooling rate was over 100000 K/s.

### 2. First generation experiments: the introduction of Constantan wires

The dissipation reactor we realized to perform the tests with the new generation Constantan wires consisted of a mica support on a central Stainless Steel (SS) tube ensuring electrical insulation, on which two wires, one active (the surface-modified constantan wire) and the other supposed inert (Ni-Cr control wire), were parallel and helicoidally rounded. The Ni-Cr wire (because intrinsically stable against oxidation or other stresses) was used to give power to the reactor itself (indirect heating). The core of the reactor was contained inside a borosilicate Schott Duran glass tube with 3 mm thick wall. Temperatures at the external glass wall and inside the reactor were detected by means of several Type K, SS-



Figure 4: Time behavior of the measured quantities during the first loading by  $H_2/Ar$  mixture: T of the external glass (green), T of the internal mica support (red),  $R/R_0$  of the inert wire (blue),  $R/R_0$  of the active wire.



Figure 3: Temperature of the external glass wall (red), the internal SS tube (blue), the internal mica support (orange) and the ambient (green) as a function of time.

screened (diameter 1.5 mm) thermocouples [1].

Calibrations were made by noble gases (He, Ar) with different powers applied to the inert wire.

The first test was conducted in an atmosphere of  $H_2/Ar$  mixture in the ratio 75/25 at 7 bar of total pressure. The power input was 48 W. Figure 3 shows the behavior of the measured quantities with time. In green color we display the temperature of the external borosilicate wall, while in red the temperature of the mica inside the reactor. The key monitor parameter is the ratio  $R/R_0$  between the resistance of the wire at a given temperature T and that at room temperature  $T_0$ . For Constantan, this ratio, because of Hydrogen interaction, decreases with increasing temperature and time. In blue we indicate  $R/R_0$  for the inert (Ni-Cr) wire and in orange  $R/R_0$  for the active (Constantan) wire. We just recall that one of the key characteristic of Constantan is its excellent resistivity stability ( $\pm 1\%$ ) from -75 up to 500 °C.

We observed that when temperature inside the reactor reached 120 °C,  $R/R_0$  of the active wire dropped down to 0.92 in about 2500 s and to 0.88 in 10<sup>5</sup> s. Correlated with this decrease of the resistance, we observed also an increase of the excess power at the output of the reactor. Correspondingly, all temperatures related to the reactor grew. External glass temperature increased from about 140 °C to 150 °C; SS tube temperature from 168 °C to 190°C and gas temperature from 170 °C to 195 °C. These variations could not be directly related to room temperature. Instead, room temperature instabilities somehow helped the anomalous heat generation, introducing non-equilibrium conditions. In fact, after long time room temperature went back to its initial value, while heat production continued increasing.

On the occasion of the National Instruments Annual Meeting in USA (NI-Week, August 2012, Austin-TX-USA), the reactor was disassembled from Frascati Laboratories-Italy, shipped and there reassembled, working in a public demo for three days. We also observed the highest values of maximum excess power for this experimental set-up: about 21 W with indirect heating (power applied on the inert wire) and about 25 W with direct heating of the active wire against an input power of 48 W. Just after the NI Week, the reactor was again shipped to South Korea for our participation at the ICCF17 Conference at Daejeon. Even there, overcoming the new strict-rules conditions of the air travel from USA (because of safety rules, no gas inside, apart from air, and/or no vacuum/pressure conditions were allowed), the reactor functioned properly with quite reproducible results:  $5 \div 10$  W of AHE with 50 W of electric input power.

### 3. Second generation experiments: the addition of glass

The successive series of experiments exhibited an unsatisfactory overall reproducibility. Using SEM/EDS/ICPMS analyses, we found out that the first batches (produced before 1970) of raw material we used in our experiments had a composition different from those later employed. Analyses revealed a Fe contamination in the order of 1000-5000 ppm, and locally up to 10000 ppm.

Because of a budget cut in the late 2013, we were forced by external events to redesign and reschedule our experiments with the purpose to study again and more deeply some of the most interesting effects obtained in the past and, if possible, to increase the AHE.



Figure 5: Data comparison between tests conducted with a small and a large amount of glass in the reactor. In red we display external glass temperature, in blue internal SS tube temperature after subtracting ambient temperature. Under the same conditions, temperature is higher when more glass is employed.

In the new experimental set-up we modified the geometrical disposition of the wires inside the reactor [2]. The wires were no longer two, but three: a 500-layers Constantan wire, a 2-layer Constantan wire and a Pt wire for control and monitoring purposes. They were inserted inside fiberglass sheaths (L = 100 cm,  $\Phi_{ext} = 1$  mm, produced by SIGI, Italy and France), with each fiber having a mean diameter of 5 µm, and closely braided together. The braid was then twisted around the central SS support that was likewise covered with a fiberglass sleeve with internal diameter 12 mm. As before, the core was inserted into the thick-wall borosilicate glass tube previously used.

Since February 2013, all the sheaths were embedded in a  $Sr(NO_3)_2$  solution and further decomposed in SrO by thermal treatment. Strontium is a material with a *low working function for electron emission*, similar to the Calcium oxide used by Yasuhiro Iwamura at Mitsubishi Heavy Industries Laboratories (Yokohama-Japan) since 1999. Electron emitter materials are empirically recognized to have "beneficent effects" on LENR reaction.

We calculated the emitted power using Stefan-Boltzmann law (emission proportional to  $T^4$ , in K) for the radiated energy from the glass wall and Newton's law (proportional to T) for the energy dissipated by convection.

During the tests several uncontrolled and unexpected phenomena occurred - included a spontaneous overheating that damaged the 500-layer wire - and the results were unclear. Therefore, we decided that the only way to clarify the role of the fiberglass in the process was to add a larger amount of glass. The SS tube, with the wires twisted around, was inserted inside two more glass hoses.

For the calculation of the heat excess, we imposed that the tests carried out with a smaller amount of glass were the threshold, i.e. the blank. Even with such a conservative constraint, the comparison between the two experiments showed that glass has intrinsic co-effects in the generation of anomalous heat. In Figure 5 we compare the temperatures at external wall and at internal SS tube in the two cases of small and large amount of glass. We note that, under the same conditions of pressure and input power, temperature in the second case is clearly larger than the corresponding values for the experiments with a lower glass content in the reactor.

## 4. Effect of fiberglass on H storage

The experiments we carried out clarified that borosilicate fiberglass that we used as electrical insulator plays a role in the generation of heat excess.

We individuate the mechanism responsible for it in the adsorption property of (borosilicate) glass, which was observed by Prof. Irving Langmuir (Nobel Laureate in Chemistry in 1932) since 1920 during his studies on Hydrogen dissociation at high temperatures (>2000°C, by W wires, at H<sub>2</sub> low pressures). The Hydrogen atoms, produced by the dissociation of molecular Hydrogen by means of the catalytic action of our sub-micrometric Constantan wires on the reaction (at quite low temperatures, about 200°C), are largely adsorbed onto the surface of the micrometric glass fibers, forming a thin film.

Langmuir measured that the density of Hydrogen atoms adsorbed onto the surface of a glass bulb (presumably of type borosilicate) kept at low temperature (about 90 K) is of the order of  $10^{15}$  atoms/cm<sup>2</sup>. In his experiments, the dissociation was obtained through a hot Tungsten filament in H<sub>2</sub> atmosphere and the fraction of dissociated molecules depended on pressure and temperature: lower the pressure and higher the temperature, larger the fraction ([**3**], [**4**]).

In our experiments the effective surface of each sheath is larger than 1 m<sup>2</sup>. The total surface of the used fibers could be larger than 50 m<sup>2</sup>, corresponding at about  $10^{20} - 10^{21}$  adsorbed atoms according to Langmuir's relation.

The H atoms concentrated onto the glass surface by adsorption are much closer to each other than the Hydrogen atoms moving in the bulb as a gas, increasing so their probability of recombination in the molecular form. This reaction is largely exothermic with the release of 4.52 eV to the environment.

According to this reconstruction, the net effect of the introduction of glass in the reactor is the increase of the rate of the chemical reaction of Hydrogen recombination. Therefore, adsorption property of glass appears to be a cofactor in the generation of heat excess in addition to the main unknown LENR process of non-chemical origin.

We also suppose that adsorption may enhance absorption of atomic hydrogen in constantan lattice, supporting in this way the main LENR process of heat generation. Actually, the possibility that the presence of an H film closely surrounding the wire favor the diffusion of atomic hydrogen inside the metal is not to exclude. In conclusion, further investigations aimed to clarify the effects and the role of glass in such kind of phenomena are necessary.

# 5. The hypothesis of energy localization

In 1954 Enrico Fermi (Nobel Laureate in Physics, 1938) and his coworkers J. Pasta and S. Ulam with the help of M. Tsingou performed, at LANL-USA, the numerical simulation of a discrete nonlinear system using the computer MANIAC I [5]. The dynamical system they studied consisted of 64 one-dimensional oscillators coupled to the nearest-neighbors by nonlinear force terms. Fermi's expectation was that the excitation energy associated to a normal node would have distributed along all the normal modes after a sufficiently long time interval, that is the system would have evolved toward a state of energy equipartition. Surprisingly, energy remained localized to few modes and showed a sort of periodicity, returning most of it to its initial mode.

Prof. Brian Ahern (previously at DARPA-USA) has recently revisited the Fermi-Pasta-Ulam problem, proposing it as a candidate for explaining some of the complex phenomena occurring in LENR experiments [6]. While in most solids the atoms reside in parabolic potential wells and therefore undergo simple harmonic motion, generally at high frequency and low amplitude, in some materials atoms are subject to a non-parabolic potential, corresponding to nonlinear force terms, that leads to nonlinear vibrational modes of large amplitude and low frequency. Hence, Ahern proposes that energy localization may take place in structures with a small number of atoms. Both properties – countable number of atoms and nonlinear coupling – are possessed by nanoparticles of size 3-12 nm (the "right" dimension suggested by Y. Arata). In nanoparticles, a considerable fraction of atoms is located at the surface and is subject to nonlinear binding forces with the internal atoms.

The transition to the new regime would be triggered by a pulse of energy that the cluster can receive from thermomechanical oscillations of the surrounding medium or from electrical pulses. As a consequence, a very small number of atoms in the cluster acquire a significantly greater amount of energy than they would in conditions of thermal equilibrium. Locally, the large oscillations manifest as hot regions. A close similarity with the phenomenon of *oscillons* in granular media can be noticed [7]. We note that, in our experimental conditions (i.e. specially the effect of glass fiber sheaths), the, local, pulse of energy could be provided by the  $H+H->H_2$  recombination reaction. Energy localization can clarify catalysis processes, usually associated to the lowering of the activation energy of a reaction. According to the depicted scenario, few excited atoms, corresponding to a locally hot region, make their energy available for the activation of chemical reactions that could take place only at higher temperature if the cluster was absent. Ahern sustains that energy localization could also explain the initiation of LENR reactions in  $H_2/D_2$  saturated nanostructures. As a consequence, energy localization circumvents the second law of thermodynamics, since nanoparticles act as Maxwell's demons able to convert part of their thermal energy into valuable chemical potential.

The violation of the second law is apparent, because not applicable to ensembles with a small number of particles. We quote Maxwell's words [8]: "The truth of the second law is ... a statistical, not a mathematical, truth, for it depends on the fact that the bodies we deal with consist of millions of molecules... Hence the second law of thermodynamics is continually being violated, and that to a considerable extent, in any sufficiently small group of molecules belonging to a real body." Violations of the second law at nanoscales have also been observed experimentally as occasional short-time (less than two seconds) fluctuations around the thermal equilibrium state ([9], [10]).

The catalytic action of nanostructures at the surface of our surface-modified Constantan wires can partially explain the thermal anomalies we found in our experiments, as they enhance  $H_2$  dissociation even at temperatures T <<800 °C.

If further confirmed, the energy localization would be the mechanism for coordinating the casual thermal motion of particles in nanometric regions of matter into localized high-energy oscillations.

### 6. Spontaneous voltage generation

In our experimental set-up we can control only two wires at the same time. For this reason one of the Constantan wires is not connected to the Data Acquisition System (PIXIe, NI) and left unconnected and floating. However, we measure periodically its resistance by means of a general-purpose multimeter to evaluate the amount of adsorbed Hydrogen. When the wire is heated in the presence of Hydrogen, its resistance ratio can decrease up to 0.7.

On June 25, 2014, we noted just by chance that the floating Constantan wire generated by itself a macroscopic voltage of the order of hundreds of microvolts that resulted function of many parameters: temperature, type of gas, pressure, resistance ratio. The highest measured values were about 1400  $\mu$ V for the tension and 120  $\mu$ A for the current with duration of only few hours, while stable outputs were about half.

We remind that the spontaneous tension cannot be ascribed to the usual Seebeck effect, because we consider only one single wire and not a junction of two different materials as in the thermocouples.

### 7. New third generation experiments: the effect of Iron

In previous experiments we found out that Iron contamination in the wires had a positive influence in terms of stability and excess power of the reaction. Hence, we decided to develop a procedure to add Iron of nanometric size to the surface of constantan wires and even deeper into the bulk for some microns during our thermal/electric treatments for the preparation of the nanolayers.

From a chemical point of view, Fe is characterized by a solubility of Hydrogen in lattice that increases largely with temperature: in 100 g of material, from 0.37 cc at 400 °C to 7 cc at about 1000 °C. To fully exploit such Fe property, the reactor had to reach higher temperatures than we previously obtained. For this reason, we added another sheath over the usual borosilicate glass one, made of alumina (Al<sub>2</sub>O<sub>3</sub>), with a higher melting temperature ( $T_{max} = 1200$  °C).

To increase wire temperature at constant input power, we used a mixture of  $H_2$  with other low thermal conductivity noble gases (Ar, Xe). The effectiveness of such mixture was verified by comparing the results about heat excesses.



Figure 6: Tests performed in  $H_2$  atmosphere for the configurations with knots with (red) and without (blue) Fe addition. The difference of power between the two configurations is displayed in black.



Figure 7: Tests in gas mixture atmosphere of Xe (1.4 bar) and  $H_2$  (1.7 bar). The temperature of the SS internal support is showed in magenta.

With the new procedure, fully developed at INFN-LNF, we produced 20-40 Fe thin layers, similar to multiple heterostructures, at wire surface. Moreover, we introduced further non-equilibrium conditions in the wire by making several (up to 20) knots with hole diameter 0.3-2 mm. These knots represent local geometrical variations of the path of the current flowing in the wire and are crossed by the lines of the magnetic field they generate. Because of the large currents (up to 2.3A) we inject in the wire, they are place of large thermal non-homogeneities. The tests we carried out with all different combinations of elements (wire with/without Iron/knots) showed that the new configuration (Iron plus knots) offers measurable advantages in terms of Hydrogen absorption and Anomalous Heat Excess.

As usual, we calculated very conservative estimations of the excess power, imposing as blank the corresponding result obtained with the configuration with no Iron in the wire. The best result (by  $H_2/Xe$  mixture) was 15 W against an input power of 100 W. The local temperature of the wire was estimated, through the Pt wire, in around 800 °C.

In Figure 6 we report the results of the tests carried out in Hydrogen atmosphere.  $T_{ext} - T_{amb}$  is plotted as a function of the input power in the case of standard Constantan wire (blue) and Fe-added constantan wire (red). In black we indicate the excess power of the configuration with Iron with respect to that without it. Clearly, excess power grows with temperature. This behavior was confirmed by the experiments with Xe/H<sub>2</sub> atmosphere, where larger wire temperatures were reached (Figure 7). Wire temperature was estimated through SS-support temperature  $T_{ss}$  measured by the thermocouples. For comparison, while at an input power of 90 W excess power is about 7.5 W by pure H<sub>2</sub> atmosphere, this raises up to 15 W by gas mixture.

We also verified the presence of a spontaneous voltage in the non-powered wire in the reactor. In wires having several knots with hole diameter smaller than 1 mm and filled with Iron at nanometric size, we observed in gas mixture atmosphere currents up to 150  $\mu$ A and voltages up to 1900  $\mu$ V, stable over long times. We compared current generation for different wires with the same number of knots and measured values about 2.2 times larger with Iron addiction. In particular, we ascertained as extremely important for maximizing the effect the local presence of Iron in the knots.

### 8. Conclusions

The observation of a single-wire spontaneous voltage generation in our experimental set-up, apart the aimed increase of AHE amount (using Fe in the most recent specific case), represents one of the main developments of our recent research activity. The phenomenon consists in the generation of a voltage between the extremities of the non-powered wire in the reactor.

The effect was first revealed almost casually in the second-generation reactor with a large amount of glass. Even more, the successive addition of Iron nanolayers and small knots in the wire (third generation reactor) increased and stabilized the magnitude of the generated voltage (up to 1900  $\mu$ V) and current (up to 150  $\mu$ A).

As concerns the anomalous heat production, Iron has a beneficial effect because of its solubility of  $H_2$  in the lattice largely increasing with temperature, while the knots are meant to create further non-equilibrium conditions inside the reactor. With such new wire features, we obtained larger excess powers.

From our observations and experiences we deduce that anomalous heat excess and spontaneous voltage generation are somehow related to the following conditions, or at least to some of them:

- Temperature as large as possible, obviously avoiding material sintering.
- Large Hydrogen absorption/adsorption by means of catalytic materials, i.e. nanomaterials.
- Hydrogen flux as large as possible from regions at high concentration to regions at lower concentration.
- Presence of elements with Hydrogen concentration increasing with temperature, like Fe.
- Presence of non-equilibrium conditions as large as possible: this is the main condition for getting any type of thermal or electrical anomaly.

We highlight that wires with good performances in terms of heat excess showed remarkable values of spontaneous voltage. One key aspect to be clarified is the role of Strontium (Sr) in the generation of the spontaneous voltage. Supposing that the effect is due to some motion of electrons from the powered-wire toward the non-powered one, an electron-emitting material like Sr might give a large contribution to it. Unfortunately, we performed no measurement of wire voltage for the old set-up with no Sr deposited on the glass sheaths and targeted tests are therefore needed.

As regards Iron, we found out that the used fiberglass sheaths (by SIGI) may contain a small fraction of Iron (< 2%) and Iron oxide (< 1%). Hence, a certain amount of Iron was already present in the second-generation experiments.

In this article we have also formulated some hypotheses about the mechanisms leading to thermal excesses in our experiments. We have focused on Hydrogen dissociation/recombination reactions, individuating three combined processes:

- The catalytic properties of Constantan help molecular Hydrogen dissociation.
- The produced atomic hydrogen is partly adsorbed at fiberglass surface. Adsorption favors Hydrogen recombination (exothermic chemical reaction) but could also promote hydrogen absorption into the metallic lattice.
- If demonstrated, energy localization at nanoscales would enhance Hydrogen dissociation by converting thermal mechanical energy in valuable chemical potential.

The next step of our activity is aimed to deeply investigate spontaneous voltage generation and possibly identify its nature and the variables that are related to this anomalous effect as well as to get its maximization.

\*\* Addendum: after the ICCF19 Conference, in May 2015, we become aware that Prof. Leif Holmlid (Göteborg University, Sweden) suggested the use of Iron, at nanosize, as cofactor in its Deuterium catalyzer (so-called "*hydrogen transfer catalyst*"), to increase the amount of the so-called, ultra-dense, Rydberg matter in its specific reactor by Inertial Confinement Fusion (ICF) processes. He aimed, and demonstrated experimentally possible by high quality and sophisticated charge particle measurements, to produce very large excess power (energy gain over 1000) just using Deuterium gas (made ultra-dense by a proper catalyst) as initial fuel and a table-top Laser as stimulator for the fusion processes. His main ideas are resumed in the recent Patent application EP 2680271A1 [11].

Moreover, Prof. Friedwardt Winterberg (Nevada University, USA), the well-known expert in ICF at International level, suggested the use of a magnetic field to increase the amount of Rydberg matter in Holmlid's experiments [12]. We just observe that our knots (carrying a sufficiently large current), filled with Fe and/or Fe oxides at nanosize, can represent a simple/first step, experimental set-up, to match some of such requirements.

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