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# **Towards Nuclear Spin Polarization of DT**

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## ABSTRACT

Recently, we have proposed an experiment to test the persistence of the polarization in a fusion process, using a powerful laser hitting a polarized HD target. In the present contribution, we first examine new possibilities towards this goal, in view of recent developments of new polarized D samples presented in this conference. We then advocate that sizeable amounts of polarized DT fuel could be obtained by the Nuclear Dynamic Polarization (DNP) of DT molecules, if a successful protocol for the DNP of the HD analog isotopic molecule could be established. We suggest that a revival of an early HD DNP experiment performed more than forty years ago, should be undertaken, taking advantage of detailed polarized HD properties revealed by thoroughly repeated static polarization of HD targets during the last ten years.

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### 1. Introduction

It is accepted that the parallel polarization of D and T nuclei should increase their reactivity when used as fuel material in fusion processes induced either by magnetic or by inertial confinement, because the fusion reaction goes mainly through the excitation of a resonant  ${}^{5}He \ 3/2^{+}$  intermediate state. In addition, the reaction products would be emitted with a  $sin^{2}\theta$  angular distribution, by reference to the polarization axis. This can be very useful to reduce damage or activation of costly equipments. The question is to know if the polarization will persist in dense and hot plasmas as anticipated from theoretical considerations, both for Magnetic Confinement Fusion (MCF) [1] and Inertial Confinement Fusion (ICF) [2]. The persistence of the polarization in a fusion process could be tested using a powerful laser hitting a polarized *HD* target. The polarized *deuterons* accelerated in the plasma generated by the laser can fuse, according to the reaction:  $(D + D \rightarrow {}^{3}He + n)$ . The angular distribution of the emitted *neutrons* and the change in the corresponding total cross section are signatures to estimate the polarization persistence [3].

In a more recent paper [4], we have discussed the experimental difficulties in using such a polarized HD target produced by the static polarization method "*Brute Force*" (BF) which is costly and takes long time, in addition to the sophisticated cryogenic holding equipment necessary to maintain the target on a laser beam line, as well as the lack of a clean flat surface of the solid HD on which to adjust the laser focusing. The main difficulty comes from the fact that the polarized material must be contained in a thin Mylar bag, offering to the laser only a small window-less surface. But so far, the BF method was almost the only one to get rather pure polarized targets either by using hyper-polarized molecules from Atomic Beam Sources (ABS) [5] or highly nuclear-spin-polarized deuterium atoms from UV photodissociation [6,7]. We shall briefely discuss below, how the corresponding experiments could be done.

#### 2. New polarized D2 samples

Polarized atoms from an ABS can recombine to molecules on a cold surface inside a dedicated apparatus [5]. During this process, it is expected that a large fraction of the initial polarization (close to 100 %) could be preserved. This would allow to produce a thin coating of solid  $D_2$ , which could be used as windowless target instead of the bulky *HD* target in our initial proposal [3]. Depending on the ABS intensity and the efficiency of the whole solidification process, a 1cm<sup>2</sup> target of a few µm thickness (enough for a laser-*D2* interaction) could be produced within hours, to be compared with months of production for a polarized *HD* target by BF with final lower *D* polarization rates [8].

Such a target would allow to repeat the Pretzler experiment [9] in quasi identical conditions except for the polarization of D nuclei. It should be noted that in Ref. [9], a prepulse of the laser is used to to generate a preplasma with a scale length of 30  $\mu$ m when the main pulse arrives after 300 ps. The authors note that without such a prepulse, no DD fusion where observed, so that the fusion reactions take place inside the preplasma.

This suggests that an other kind of polarized target could also be used. It has been shown in this symposium [7] how laser can be used to create polarized deuterium samples at nearly atmospheric pressure ( $10^{19}$  cm<sup>-3</sup>). The exact experimental conditions of [9] ( $n_e \sim 10^{21}$  cm<sup>-3</sup>) are not readily realized, but intensive numerical simulations might tailor the proper choice of the various experimental parameters to get the best fusion rate. It is likely that with modern ultrashort lasers, even with moderate pulse energies, some fusion will occur allowing to perform a significant exeptiment within a reasonable time.

It is worth noting that the initial idea was that in the plasma created by the laser, polarized deuterons would be heated enough to undergo mutual thermal fusion. However, it is unlikely that thermal fusion alone could account for the observed rate of neutron production as seen in previous exepriments. More likely, some deuterons are accelerated in the plasma by the laser plasma interaction and those accelerated ions act as a beam to generate nuclear reactions with colder deuterons in the plasma. Thermal fusion dominance would probably require much more energetic laser pulses!

#### 3. Nuclear spin polarization of solid Hydrogen

The Hydrogen family comprises 6 members: 3 homonuclear ones  $(H_2, D_2 \text{ and } T_2)$ , 3 heteronuclear ones (HD, HT and DT); among them 3 are radioactive (HT, DT and  $T_2$ ). Due to symmetry restrictions of the wave function, the polarization of homonuclear molecules is complicated by metastables states producing at low temperature 2 coexisting configurations with different polarizabilities. As an example, the polarization of  $H_2$  at low temperature in its ortho metastable configuration (2 nuclear spins aligned) could be attempted, however the equilibrium configuration at low temperature is the para one (2 nuclear spins antiparallel), so that in a few hours, by decay of the ortho to the para form, the  $H_2$  molecules lose their polarization. Therefore, in the last 10 years, the BF static polarization of HD has been pursued, essentially because the hetero HD, being formed of a fermion and a boson, has no symmetry restriction of its total wave function [8]. However, to reach sizable polarization rates by BF, it is necessary to go down to very low temperatures and high magnetic fields, namely dilution refrigerators and supraconducting coils which require heavy equipments. For example, at 10 mK and 17 Tesla, the proton polarization can reach ~ 90%, while the deuteron vector polarization is only  $\sim 33\%$  due to the lower deuteron magnetic moment. Reaching low temperatures requires some time and allowing the contaminant homonuclear species decay to their magnetically inactive forms takes more than a month. Accordingly, even at moderate low temperatures (4 K) and weak magnetic fields (1 T), relaxation times can become very long [8].

The *DT* has the most interesting Hydrogen configuration, because it contains both the *D* and *T* nuclei necessary for nuclear fusion. Unfortunately, the BF method cannot be applied to *DT*, because the heat generated by the Tritium radioactivity does not allow to reach very low temperatures. The power of dilution refrigerators at 10 mK, is in the range of  $\mu$ W, while the Tritium radioactivity produces a heat in the range of W/mol. For the DNP, temperatures around 1-4 K are sufficient and refrigerators in the range of a few W powers can be built. Since *HD* and *DT* have very similar magnetic properties, it is convenient to optimize the DNP protocol on *HD*. So far, the results onbtained by the DNP of *HD* were not encouraging. The low polarization rates (less than 5%) observed, could have been limited by significant amounts of  $H_2$  and  $D_2$  impurities [10]. Fig. 1 shows a schematic picture of the DNP process. The high polarization of electrons is transferred to the nuclear spins by RF transitions. The key points to get high polarization rates are: i) well resolved states so that RF transitions are connected to well defined

hyperfine states [11], ii) electron spin-lattice relaxation times  $T_{1e}$  much shorter than the nuclear relaxation ones  $T_{1N}$ . It turns out that for Hydrogen,  $T_{1e}$  is unusually long.

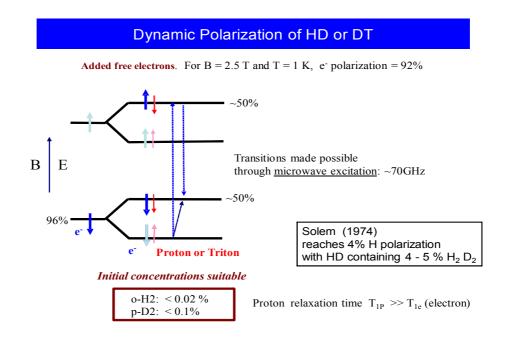


Fig. 1 : Sketch of the DNP process, according to the solid state effect. The polarization of unpaired electrons, is transferred to the nuclei by RF transition between hyperfine states. If the  $T_{1e}$  is much shorter than the  $T_{1N}$ , electrons undergo spin flip rapidly, while nuclear spins keep their orientation. By repeating this process, nearly all the electron polarization can be transferred to the nuclei.

To shorten  $T_{1e}$  Solem [10] added to the *HD* a small fraction (10<sup>-3</sup>) of paramagnetic  $O_2$ , which shortened drastically  $T_{1e}$  but at the same time broadened the hyperfine states, as shown by the resulting Electron Paramagnetic Resonance (EPR) spectra measured by Solem himself. For pure *HD*, the EPR sectra show well resolved hyperfine states, both for proton and deuteron after addind  $O_2$ , the states linked to the protons are significantly broadened, while those from the deuterons are severely smeared. No wonder that the deuteron polarization could hardly be build up! Pure *HD* samples are now available with extra long  $T_{1N}$ . As shown in Ref. [12],  $T_{1N}$  can be increased almost at will by using higly distilled *HD* gas and ageing it to let the ortho- $H_2$  and para- $D_2$  impurities decay to their magnetic inactive configurations. Compared to the  $T_{1e}$  in the range of seconds as mentionned by Solem,  $T_{1N}$  in the range of hours have been achieved [12]. A suitable density of unpaired electrons could be generated by intensive irradiation. Recent exposition of polarized *HD* samples to electrons have shown some polarization loss but no significant reduction of  $T_{1N}$  [13]. It is obvious that the Solem investigation of the DNP of *HD* should be repeated, using recent expertise on the *HD* properties. An optimum DNP protocol could be established and extended to  $D_2$  and *DT* for polarized fusion experiments.

#### 4. Conclusion

It has been shown that the experiment initially proposed to test the persistence of the polarization in a fusion process, could be done in much better experimental conditions, using presently available polarized window-less  $D_2$  samples. Although the related polarization

methods from ABS or UV dissociation have a great potential, it is not clear yet how they can produce enough polarized fuel for fusion.

A fusion reactor like ITER, with a power of 500 MW would need at least 1 mol/hour of DT, assuming 100% efficiency, which will not be by far the case! Therefore, it is highly desirable to find a way to polarize sizeable amounts of DT. It has been advocated that the DNP protocol of solid HD, if successful after repeating early experiments in presently much better conditions, could be extended to DT and even  $D_2$ . It should be noted that sizeable quantities of polarized material are already produced by DNP for nuclear physics experiments: the COMPASS polarized target at CERN is able to polarize 50 mol of material within a single polarization cycle [14].

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