

Hydrogen, anti-hydrogen beams production in a long linear vacuum pipe (submitted to Nucl. Inst. Meth.)

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This paper presents the possible production of hydrogen and anti-hydrogen from p^\pm and e^\mp beams merged in a long vacuum chamber (about 2500 m). In such a configuration with a beam kinetic energy for the p^\pm around 100 KeV, the two key parameters are the density of the e^\mp beam and the temperature of the merged beams. The production rate is computed for several parameter sets and it is shown that it is possible to recombine $10^6 p^\pm$ with nearly 100% efficiency. Possible experiments using this configuration are presented, including the fine structure constant measurement, the anti-hydrogen spectroscopy and the comparison of the gravitational mass for hydrogen and antihydrogen. PACS numbers: 25.43.+t (antiproton and antihydrogen), 06.20.Jr (fine structure constant, gravitational mass), 41.75.Lx (crystalline beams)

1. Introduction

Several experiments using storage rings or electromagnetic traps have been proposed to produce and store anti-hydrogen [2,19]. In both cases it is expected that the anti-hydrogen atoms are produced at very low rates. The configuration proposed here is a long vacuum pipe inside which the particles travel only once and the anti-hydrogen is produced by bursts of about 10^6 atoms. The topics presented and discussed are the following:

- the configuration and the different sets of beam hypothesis,
- the computation of the production rate and its application to the different sets,
- the focussing of the anti-atom or atom beam at the exit of the pipe,
- the experiments' configurations.

In the last section the production of p^- is briefly discussed. The hydrogen atom is referred to as H while the anti-hydrogen atom is referred to as \bar{H} .

2. Configuration and hypotheses

The general set-up is shown in figure 1. It is derived from the existing experiments made at ion storage rings where ion beams are merged with

electrons in an electron cooler [5]. The recombination region defined by the length of the cooler is approximately 1 meter and the unused ions are recycled in the storage ring and re-used at the next path through the cooler. In the long pipe configuration, the ions are not recycled but the interaction time, defined by the duration of the two beams superposition, is about 2500 larger. This longer time allows for a recombination of all the p^- up to 10^6 particles (see section 4). The point where the beams are injected will be referred to as the *entry* and the other pipe extremity as the *exit*.

The other features of ion storage ring experiments are identical here: the e^\mp beam has a diameter of approximately $1 \sim 2$ cm and the cooled p^\pm beam collapses into a cylinder of approximately 2 mm. The e^\mp beam is kept within this diameter along the 2500 m by a weak magnetic field.

As the experiments on the radiative recombination into hydrogen atoms of $e^- p^+$ have shown, the reaction can be enhanced by a laser [4,5]. Both cases, with and without laser induced radiative recombination, will be considered. When there is a laser, the light is injected from the exit of the pipe. The photon energy seen in the rest frame of the p^\pm is then Doppler shifted.

The atom production rate is computed for several sets of parameters, including some very dif-

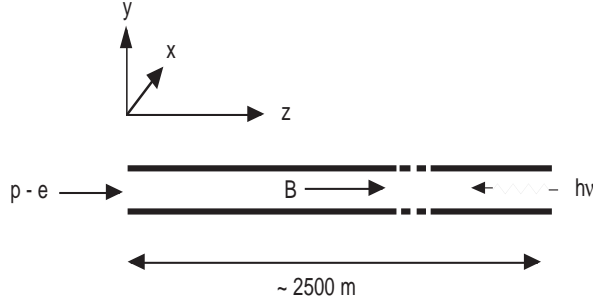


Figure 1. Merged beams configuration: the two beams are injected and superposed on one side and the laser light is injected at the other side. A weak magnetic field parallel to the pipe is used to guide the e^\mp beam.

ferent from the suggested ones: these sets have been chosen to compare the results with past experiments. Each set contains:

- the kinetic energy of the p^\pm : $K(p^\pm)$,
- the corresponding kinetic energy for the e^\mp to have the same velocity: $K(e^\mp)$,
- the velocity: V ,
- the relativistic coefficients: β, γ ,
- the time of flight across the pipe: τ ,
- the Doppler factor: $\sqrt{\frac{1+\beta}{1-\beta}}$.

Tables 1 and 2 correspond to a long time of flight in order to produce beams of $10^5 \sim 10^6 \text{ } \dot{H}$ per burst.

Table 3 corresponds to the kinetic energy of the p^- beam produced at CERN with the Anti-proton Decelerator (AD). Table 4 shows the kinetic energy required to have a Doppler shift factor of 5 suited for the boosted spectroscopy experiment to be presented in a later section.

The experiments on the radiative recombination of e^-p^+ into hydrogen atoms performed at

Table 1

Set 1	
p^\pm	50 KeV
e^\mp	27.23085 eV
γ	1.00005328943
β	0.0103232912
V	3094844.8551 $m.s^{-1}$
τ	0.808 10^{-3} s
$\sqrt{\frac{1+\beta}{1-\beta}}$	1.01037713079

Table 2

Set 2 represent the suggested target parameters.

p^\pm	100 KeV
e^\mp	54.46170 eV
γ	1.00010657887
β	0.0145987557
V	4376596.86545 $m.s^{-1}$
τ	0.57 10^{-3} s
$\sqrt{\frac{1+\beta}{1-\beta}}$	1.01470689052

Table 3

Set 3: $K(p^-)$ produced at CERN with the AD.

p^\pm	5 MeV
e^\mp	2.72308505 KeV
γ	1.00532894336
β	0.102826529
V	30826617.9578 $m.s^{-1}$
τ	81.1 10^{-6} s
$\sqrt{\frac{1+\beta}{1-\beta}}$	1.10870342938

Table 4

Set 4: boosted spectroscopy experiment.

p^\pm	1.501235696 GeV
e^\mp	817.598496 KeV
γ	2.6
β	0.923076923
V	276731499.692 $m.s^{-1}$
τ	9.03 10^{-6} s
$\sqrt{\frac{1+\beta}{1-\beta}}$	5

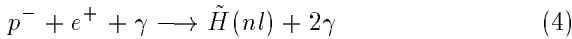
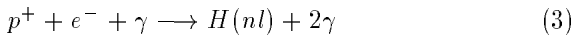
the TSR ring in Heidelberg [5] and with the MEIBE [4] apparatus at the University of Western Ontario used p^+ beams with a kinetic energy of 21 MeV and 330 KeV respectively. These values are within the range defined by the parameter sets chosen here. These experiments have shown experimental evidence that laser induced radiative recombination works and is well described by the theoretical formulae.

The main differences are:

- the interaction length which is 1.5m at TSR and 8.6 cm at MEIBE
- the beam temperature as it shall be discussed in the next section.

3. The recombination rate

The two reactions involved are the radiative recombination (RR = 1 & 2) and the laser induced radiative recombination (LIRR = 3 & 4).



The expected production rates of H and \tilde{H} being the same, in the following computation, only the case of H is considered.

The notation is the following: $H(nl)$ is the atom in the state with principal quantum number n and orbital quantum number l . The energy of the ground level ($n = 1$) is the ionization energy, $E_0 = -13.6 \text{ eV}$. The photon energy E_γ matches the difference between the kinetic energy of e^- in the rest frame of p^+ and the energy of level n :

$$E_\gamma = h\nu = \frac{1}{2}mv_e^2 + \frac{|E_0|}{n^2} \quad (5)$$

In this formula we have only taken into account the principal quantum number because the spread in photon energy is dominated by the

spread in e^- velocity. The separation between states sharing the same principal quantum number and having different orbital quantum numbers l is negligible in this case.

The RR rate in a given state level nl is given by:

$$r(n) = n_e N_p \frac{\tau}{\gamma} \alpha(nl) \quad (6)$$

Where:

- n_e is the e^- beam density,
- N_p is the number of p^+ in the beam,
- τ/γ is the time of flight seen in the p^+ rest frame,
- $\alpha(nl)$ is the recombination coefficient.

The recombination coefficient is computed from the e^- velocity distribution in the rest frame of the p^+ and from the recombination cross section:

$$\alpha(nl) = \int_0^\infty v_e f(v_e) \sigma_{rec}(nl) dv_e \quad (7)$$

The velocity distribution in the beams follows a Maxwell distribution and with the notations used in [2] we have:

$$f(v) = \frac{\exp[-(\frac{v_\perp}{\Delta_\perp})^2] \exp[-(\frac{v_\parallel}{\Delta_\parallel})^2]}{\Delta_\perp^2 \Delta_\parallel \pi \sqrt{\pi}} v^2 \quad (8)$$

$$v_\perp = v^2 \sin(\theta)^2 \quad v_\parallel = v \cos(\theta)$$

Where \perp corresponds to the velocity projection in the transversal plane (x, y) and \parallel corresponds to the velocity projection along the z axis. The temperatures in the transversal and longitudinal directions will be noted T_\perp and T_\parallel . The temperatures correspond to kinetic energy distributions and will be estimated in energy units (meV) rather than Kelvins. The use of "kinetic energy" is reserved for single particles or for target beam energy.

Introducing the $erf(z)$ function, equation 7 can be rewritten in the form¹:

¹see [2] p256 eq 11 and 13

$$\alpha(nl) = \frac{2}{\Delta_{\perp}^2 \Delta_{\parallel}} \int_0^{\infty} \sigma_{rec}(nl) v^3 \exp\left(-\left(\frac{v_{\perp}}{\Delta_{\perp}}\right)^2\right) \frac{erf(z)}{z} dv \quad (9)$$

Where z is defined by:

$$z = v \sqrt{\frac{1}{\Delta_{\parallel}^2} - \frac{1}{\Delta_{\perp}^2}} \quad (10)$$

In the following we will suppose that the beams have a "flattened" distribution: $\Delta_{\parallel} \ll \Delta_{\perp}$.

With this hypothesis, equation 10 becomes:

$$\alpha(nl) = \frac{2}{\Delta_{\perp}^2} \int_0^{\infty} \sigma_{rec}(nl) v^2 \exp\left(-\left(\frac{v_{\perp}}{\Delta_{\perp}}\right)^2\right) dv \quad (11)$$

This integral is computed analytically in the appendix.

The LIRR process is similar to the stimulated emission of a photon with the difference that while stimulated emission occurs between two discrete quantum states, LIRR occurs between the continuum and a discrete state [2]. The same analogy can be drawn between the RR process and spontaneous emission. In the same way that spontaneous emission and stimulated emission probabilities are related by the Einstein relations, similar relations exist between RR and LIRR. If we define the gain $G(n)$ by the ratio of the LIRR and RR processes for a recombination into the atom with energy level n then:

$$G(n) = \frac{\sigma(LIRR)}{\sigma(RR)} = \frac{Pc^2}{F\Delta\nu 8\pi h\nu^3} \quad (12)$$

where:

- P is the laser power in Watts,
- F is the laser beam area cross-section
- ν is the laser frequency

- $\Delta\nu$ is computed from equation 5 and is dominated by the e^- kinetic energy spread (the atomic level width is negligible).

Therefore the LIRR rate is obtained from the RR rate computation.

It shall be noted that the LIRR process is limited by the photoionization when the photon flux increases: the atom, created by the LIRR process, is immediately destroyed by photoionization because both processes are induced by photons with the same wavelength.

The maximum photon flux Φ before saturation due to photoionization is given by:

$$\Phi = t_{min} \times \sigma_{phot}(n, \nu) \quad (13)$$

Where:

- t_{min} is the shorter of the laser pulse time length and the target state (nl) life time,
- $\sigma_{phot}(n, \nu)$ is the photoionization cross section for the atom at energy level n and with photons of frequency ν .

The TSR experiment has observed LIRR with the target state level $n = 2$ while the MEIBE experiment has observed LIRR with the target states $n = 11$ and $n = 12$. Depending on the target state, typical gains with conventional lasers are in the range (50, 2000). It shall be noted that the experiment at TSR, due to a miss-alignment of the two beams, had an electrostatic field which enhanced the photoionization cross section [8–11], therefore limiting the gain. This effect does not appear when the beams are well aligned. In the configuration presented here, the e^{\mp} beam will produce an electric field which is zero on the beam axis and reaches its maximum on the edge at a radius of approximately 1 cm. In order to avoid the enhancement of the photoionization cross section, the laser light shall be concentrated within a diameter of 2 ~ 3 mm. As the RR rate will show, a gain of 50 is sufficient here.

4. Numerical results for the different sets

The target energy levels of the atom after recombination and their life time are given in table 5.

Table 5
12 first energy levels and life times of the (anti) hydrogen atom.

n	$E(eV)$	$\lambda(nm)$	$\tau_0(ns)$
1	-13.6	91.91	∞
2	-3.4	367.65	2.1
3	-1.5	833.30	10.2
4	-0.85	1470.60	33.5
5	-0.54	2314.80	88
6	-0.37	3308.80	196
7	-0.28	4503.60	81.2
8	-0.21	5882.40	108.3
9	-0.168	7444.80	173.5
10	-0.136	9191.10	238.3
11	-0.11	11121.30	317.6
12	-0.09	13235.20	412.7

The origin of the energy scale corresponds to the system of free particles $e + p$. The recombination emits a photon whose wavelength is in the 3rd column of table 5. After the recombination into the state (nl) , the atom radiates photon(s) to reach the ground level. The life time of the atom [1] in the excited level (nl) is the 4th column in table 5.

The target beam temperatures for this experiment are below 100 meV, therefore the values in column 2 show the excited states up to level 12. Above this level, excited states can be re-ionized by collisions and the number of atoms produced results from the competition between recombination and ionization. These states will not be taken into account for the estimation of the production rate. Hence the rates are slightly underestimated (less than a factor 2). The numerical results of the computation of the recombination coefficient $\alpha(nl)$ are presented in tables 6, 7 and 8 in units of $10^{-12} cm^3.s^{-1}$. The transversal beam temperature T_{\perp} in meV is shown on top of each column.

The transversal beam temperature of $T_{\perp} = 200 meV$ is included to be compared with the experimental value found at Novosibirsk [2,7]:

Table 6
 $\alpha(nl)$ in units of $10^{-12} cm^3.s^{-1}$

n	0.5 meV	1.0 meV	1.5 meV	2.0 meV
1	10.716	7.577	6.187	5.358
2	4.286	3.031	2.475	2.143
3	4.072	2.879	2.351	2.036
4	3.372	2.383	1.945	1.684
5	2.697	1.906	1.556	1.347
6	2.247	1.588	1.296	1.121
7	1.926	1.360	1.110	0.960
8	1.684	1.190	0.970	0.839
9	1.497	1.057	0.862	0.745
10	1.347	0.950	0.775	0.670
11	1.224	0.863	0.703	0.608
12	1.121	0.791	0.644	0.556
Total	36.189	25.575	20.874	18.067

Table 7
 $\alpha(nl)$ in units of $10^{-12} cm^3.s^{-1}$

n	25 meV	50 meV	75 meV
1	1.5155	1.0716	0.8749
2	0.8335	0.5894	0.4811
3	0.5759	0.4072	0.3325
4	0.4703	0.3281	0.2646
5	0.3734	0.2588	0.2075
6	0.3084	0.2122	0.1691
7	0.2617	0.1787	0.1415
8	0.2264	0.1534	0.1208
9	0.1988	0.1337	0.1046
10	0.1767	0.1178	0.0917
11	0.1584	0.1048	0.0811
12	0.1433	0.0940	0.0724
Total	6.2423	3.6502	2.9418

Table 8

 $\alpha(nl)$ in units of $10^{-12} \text{ cm}^3 \cdot \text{s}^{-1}$

n	100 meV	200 meV
1	0.7577	0.5358
2	0.4167	0.2143
3	0.2879	0.2036
4	0.2264	0.1534
5	0.1767	0.1178
6	0.1432	0.0940
7	0.1193	0.0771
8	0.1013	0.0646
9	0.0873	-
10	0.0761	-
11	0.0671	-
12	-	-
Total	2.4597	1.4606

$1.46 \cdot 10^{-12} \text{ cm}^3 \cdot \text{s}^{-1}$ versus $2.2 \cdot 10^{-12} \text{ cm}^3 \cdot \text{s}^{-1}$. The difference is due to the limitation of the sum over the 8 levels which are not affected by re-ionization at that temperature while all levels participate in the experimental value.

In equation 6, the hypothesis on the number of p^\pm and the density of the e^\mp beam is that the product $n_e N_p = 5 \cdot 10^{19} \text{ cm}^{-3}$ (this hypothesis is discussed in the next section).

Then, the recombination coefficient values together with times of flight τ and the relativistic boost γ tabulated in tables 1 2 3 4 allows the computation of the number of recombined atoms produced at each burst in the vacuum pipe via equation 6.

The numbers obtained in table 9 show that, with a laser to set the gain at 50, $10^6 p^-$ can be recombined at any beam temperature below 200 meV and a kinetic energy $K(p^\pm) = 100 \text{ KeV}$. If we define the apparatus efficiency as the proportion of recombined p^- , then the efficiency is 100 %: all the p^- produced are used. In the long vacuum pipe configuration, the p^- are not recycled like in a storage ring but nevertheless they are all used. Furthermore, to keep p^- in a storage ring a very good vacuum has to be maintained.

Table 9

Millions of recombined atoms per burst without laser gain

T_\perp	50 KeV	100 KeV
0.5 meV	1.46	1.03
1.0 meV	1.03	0.73
1.5 meV	0.84	0.59
2.0 meV	0.73	0.51
25 meV	0.252	0.178
50 meV	0.147	0.104
75 meV	0.119	0.084
100 meV	0.099	0.070
200 meV	0.059	0.042

Table 10

Millions of recombined atoms per burst without laser gain

T_\perp	5 MeV	1.5 GeV
0.5 meV	0.14	0.006
1.0 meV	0.10	0.004
1.5 meV	0.08	0.0035
2.0 meV	0.07	0.0031
25 meV	0.025	0.0011
50 meV	0.015	0.0006
75 meV	0.012	0.0005
100 meV	0.010	0.00042
200 meV	0.006	0.00025

At the AD storage ring at CERN, the limitation of the number of p^- produced at each burst is due to the vacuum. The AD vacuum is approximately 10^{-10} torr which corresponds to a residual gas density equal to that of the p^- beam ($\sim 2 \cdot 10^7$ particles, 2 mm diameter and 100 m length). It shall be noted that the AD was built with recycled magnets that cannot be heated to remove adsorbed particles while a vacuum of approximately 10^{-11} torr can be achieved for the system presented here. The total time spent in the AD by the p^- for deceleration and cooling is about 80 s. In the long vacuum pipe configuration, the time of flight is less than 10^{-3} s which allows much shorter beams without the limitation due to the residual gas. Another difference with storage rings, which continuously produce recombined atoms at low rate, is that the beams do not need to be in a stable state during the path in the long vacuum pipe: shorter dense beams which could not be kept in a storage ring can be packaged at the entry provided that their instability has a characteristic time larger than 10^{-3} s. The last electron cooling in the AD takes 10^6 loops, that is 6 s. The cooler length is approximately 10^{-2} of the total AD length, hence the transfer of heat between beams takes about $6 \cdot 10^{-2}$ s. In the long vacuum pipe, the beam heating due to the density will not have time to take place.

5. Focussing of the charged beams

Experimentally, there is no difficulty to produce e^- beams with a density of $5 \cdot 10^{13} \text{ cm}^{-3}$ and beams of p^+ with 10^6 particles. The situation for p^- and e^+ is more difficult: in order to be produced and collected, the p^- requires a p^+ primary beam above $\approx 12 \text{ GeV}$. The e^+ are easier to produce and bursts of 10^{13} particles have been achieved at SLAC where bursts with more particles are foreseen.

With these numbers, the density of e^+ is seven orders of magnitude larger than the density of p^- particles: the two superposed beams can be considered as "a pure e^+ beam". At a p^+ kinetic energy of 100 keV , the corresponding e^- kinetic energy is only 54.4 eV . There are two consequences:

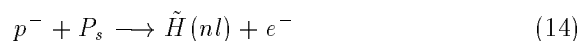
- the e^+ beam is destroyed by its internal

electric field,

- no focussing can be applied to compensate for this field.

To cure this problem, a third beam of e^- is superposed to compensate the space charge so that a weak axial magnetic field can guide the beams.

The superposition of an e^- beam with an e^+ beam at low temperature produces positronium atoms. There are three reactions to be considered:



Computations [34,35] confirmed by an experiment [36] have shown that in the temperatures range considered here, the cross section² of the reaction 14 is about $10 \pi a_0^2 \sim 10^{-15} \text{ cm}^2$. This is 10000 times larger than the RR cross section. Therefore, the positronium enhances the production of \tilde{H} .

The cross section of the reaction 15 is the same as the RR cross section of the e-p recombination provided that the ionization energy is replaced by the one of positronium. Hence this cross section is about 1/2 of the RR cross section:

$$\alpha_{e-e} = \frac{1}{2} \alpha_{e-p} \quad (17)$$

In the singlet spin state, the life time of positronium $P_s(1s)$ is 0.142 ns . In the triplet spin state, the life time is 142 ns . Therefore, the reaction 16 removes e^+ from the beam.

Since the positronium is about 10000 times more effective for producing \tilde{H} but lives approximately 100 ns , the positronium is equivalent to the presence of an e^+ particle during 1 ms , that is during the time of flight in the pipe.

Since the wavelength of the laser which sets the gain is tuned for the e-p reaction, if the beam of e^+ particles is neutralized by a beam of e^- , then most of the H/\tilde{H} production is done through the

²where $a_0 = 0.529 \cdot 10^{-10} \text{ m}$ is the Bohr radius

e-p channel. For instance, at a temperature of 10 meV , $\alpha_{e-p} \sim 10^{-11}\text{ cm}^2$, hence the probability for a p^\pm to be recombined during a 1 ms flight is approximatively:

$$\alpha_{e-p} \times n_e \times 10^{-3} = 10\% \quad (18)$$

And the probability for producing a P_s is approximatively 5%. With a gain set at $G = 10$ the production of H/\tilde{H} is 20 times more likely to happend through the e-p channel than through the P_s channel. The total number of e^\mp absorbed by the production of P_s is only 5% of the total. Therefore, the beam neutralization, while important for keeping the charged beams in the pipe, plays only a small role in the H/\tilde{H} production.

6. Focussing of the anti-atom or atom beam

The recombined atoms are produced all along the pipe and, depending on the transversal beam temperature, they may or may not need guiding devices to reach the exit without interacting with the pipe. Table 11 shows the average transversal distance at the pipe exit (in cm) for the atoms created at the entry: it gives the radius of the cone inside which the atoms would be after a 2500 m flight, if no focussing device is installed.

Since the atoms are on the ground level, and since there is no laser available with a wave length of 121 nm it is not possible to use Doppler shifted stimulated emission of the one-photon $1s-2p$ transition for transferring transversal quanta of momentum to focus the beam.

Therefore it is proposed to use the hyperfine (hfs) structure of the $1s$ energy level. This transition is used in MASER experiments where few atoms in the ground level excite a spherical cavity tuned on the hfs transition (1.420 GHz) which, in turn, produces stimulated emission of the other atoms, generating the measured electric signal. The focussing device is a MASER operated in reverse mode: the electric signal is not extracted but injected in the cavity. The cavity is excited in the TE101 mode at a frequency Doppler shifted so that the atoms with transverse velocity transfer transverse momentum to the electromagnetic

Table 11
Atom beam cone radius (without focussing) at the exit in cm

T_\perp	50 KeV	100 KeV	5 MeV	1.5 GeV
0.1 meV	11.2	7.9	1.1	0.1
0.2 meV	15.8	11.2	1.6	0.2
0.5 meV	25.0	17.6	2.5	0.3
1.0 meV	35.4	24.9	3.5	0.4
2.0 meV	50.0	35.3	5.0	0.6
5.0 meV	79.1	55.8	7.9	0.9
10 meV	111.8	78.9	11.2	1.2
20 meV	158.2	111.6	15.9	1.8
50 meV	250.1	176.4	25.1	2.8
100 meV	353.7	249.5	35.5	3.9
150 meV	433.1	305.6	43.5	4.8
200 meV	500.1	352.8	50.2	5.6

field in the cavity by stimulated emission. Several shapes can be selected for the cavity: spherical, cylindrical, rectangular. For MASER experiments the spherical shape is often selected and will be taken as practical example. The mode to be excited, TE101, is shown in figure 2. For this mode a cavity made of copper has the following parameters:

- the sphere radius a ,
- the electrical resistance R_s ,
- the quality coefficient Q ,
- the frequency width $\Delta\nu$.

This mode is the superposition of spherical waves travelling from the z axis to the edge and from the edge to the z axis. The cavity is tuned with a Doppler shift so that the atoms with transverse momentum are within the resonance while the other ones are de-tuned. This is possible due to the width of the mode which is slightly larger than the Doppler shift for an atom with kinetic energy of 200 meV . The transversal velocity and the corresponding Doppler shift for different temperatures is presented in table 13.

One can see in figure 3 that, when the cavity tune is shifted by $29328 Hz$, atoms with no transversal velocity are de-tuned.

The crossing of the cavity for a beam with kinetic energy of $100 KeV$ takes $96 ns$ during which the field in the cavity oscillates 136.3 times. By shifting the beam target energy to $99.56 KeV$ the number of oscillations is exactly 136 therefore the charged particles moving together with the atoms will not receive or lose energy during the crossing. Furthermore, the electric field on the z axis is $E = 0 V.m^{-1}$ and the p^- beam is only $2 mm$ in diameter while the e^+ beam is $1 cm$ in diameter. At a distance of $1 cm$ in the transversal plane, the field is only $\approx 5 \cdot 10^{-3} \times E_{max}$ and the e^+ are guided by the axial magnetic field both inside the cavity (oscillating field) and outside. Hence, crossing the cavity shall create only negligible perturbations of the charged beams.

The focussing power of the cavity is limited by the 136 field oscillations. Each oscillation corresponds to one stimulated emission and each hfs photon has an energy of $5.87 \cdot 10^{-6} eV$, therefore the transversal kinetic energy of the atom is decreased by $136 \times 5.87 \cdot 10^{-6} eV = 0.798 meV$. This gives the focussing power of a single cavity and allows the computation of the number of cavities needed to remove the transversal kinetic energy. These numbers are summarized in table 14 which contains for each kinetic energy, the number of stimulated emissions and the corresponding number of cavities.

The electromagnetic power required to force the atoms' spin to oscillate 136 times is less than a few Watts. This is due to the life time of the upper hfs energy level which is very long: $\tau \sim 1s$. If A and B are the Einstein coefficients for spontaneous emission and absorption then:

$$A = \frac{1}{\tau} = 1 \quad B = \frac{A \lambda^3}{8 \pi h} = 5.6 \cdot 10^{29} \quad (19)$$

Therefore, the spontaneous emission is negligible and the number of transitions per unit time is given by:

$$dN = -N B \rho_\nu dt \quad (20)$$

Table 12
Spherical cavity parameters for the mode TE101

ν_{hfs}	$1420405751.768 Hz$
$a = \frac{\lambda_{hfs}}{1.39}$	$15.18 cm$
$R_s = 2.6 \cdot 10^{-17} \sqrt{\nu_{hfs}}$	$9.8 \cdot 10^{-3} \Omega$
$Q = \frac{1}{R_s} \sqrt{\frac{\mu_0}{\epsilon_0}}$	38473.508
$\Delta \nu = \frac{\nu_{hfs}}{Q}$	$36919.06 Hz$

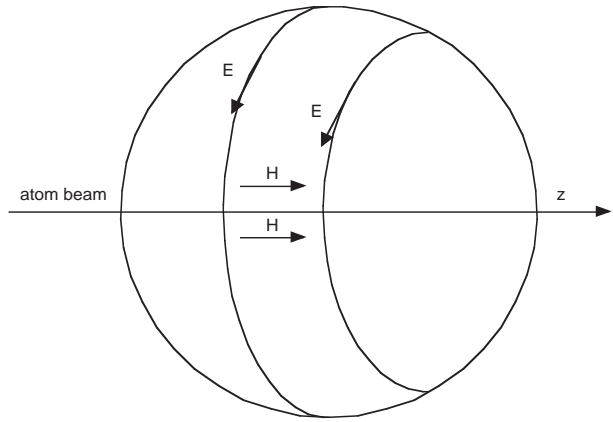


Figure 2. Field in a spherical cavity excited in the TE101 mode: the beam crosses the cavity along the z axis. Atoms with transverse momentum recoil from the stimulated emission of hfs photons.

The energy density in the cavity is:

$$\rho_\nu = \frac{1}{2} \epsilon_0 E(\nu)^2 + \frac{1}{2} \mu_0 H(\nu)^2 \quad (21)$$

For a field of $1 V/m$, this leads to a transition characteristic time:

$$\frac{1}{B \rho_\nu} = \frac{1}{10^{-12} \times 10^{29}} = 10^{-17} s \quad (22)$$

In other words, even with a weak field of about $1 V/m$ the atoms spin follows the field oscillations.

7. Experimental configurations

The long vacuum pipe configuration brings specific capabilities which are difficult to achieve oth-

Table 13
Velocity and Doppler shift

T_{\perp}	$V \text{ m.s}^{-1}$	$\Delta \nu_{hfs} \text{ Hz}$
0.1 meV	138.41	655.8
0.2 meV	195.74	927.4
0.5 meV	309.50	1466.4
1.0 meV	437.70	2073.8
2.0 meV	619.00	2932.8
5.0 meV	978.71	4637.1
10 meV	1384.11	6557.9
20 meV	1957.43	9274.2
50 meV	3094.97	14663.9
100 meV	4376.95	20738.0
150 meV	5360.64	25398.7
200 meV	6189.94	29328.0

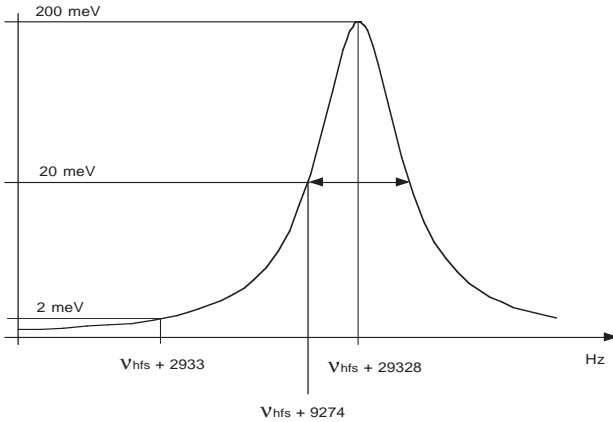


Figure 3. Doppler shift of the atoms according to their transversal kinetic energy

Table 14
Focussing power needed as a function of temperature

T_{\perp}	$nb \ h\nu_{hfs}$	$nb \ cavities$
100 meV	17023	125
50 meV	8511	63
20 meV	3405	25
10 meV	1702	16
5 meV	851	6
2 meV	340	3
1 meV	170	2
0.5 meV	85	1
0.2 meV	34	1
0.1 meV	17	1

erwise. They will be illustrated for three experiments:

- the Lamb shift measurement,
- the boosted spectroscopy,
- the gravitational mass measurement.

7.1. The Lamb shift measurement

Precise measurements of the Lamb shift in atomic hydrogen were obtained using the technique of the Separated Oscillatory Field [12–17] for the level $n = 2$.

Extensive reviews on this technique have been published. These measurements are among the most stringent tests of QED. In the SOF experiment, the difficulty is to produce a cold beam of atomic hydrogen with a kinetic energy in the 100 keV range from which atoms in the state $2S_{1/2}(F = 0)$ are extracted and used for the interferometry. These beams are produced with a small proton accelerator followed by recombination into atoms through charge exchange collisions with nitrogen gas. The two advantages of using e^-p^+ recombination instead of charge exchange are:

- the number of atoms with energy level $n = 2$ can be enhanced with laser induced gain as in the TSR experiment,

Table 15

Number of recombined atoms in the $2S$ state per burst without laser gain

T_{\perp}	50 KeV	100 KeV
0.5 meV	64939	45811
1.0 meV	45917	32392
1.5 meV	37493	26449
2.0 meV	32469	22905
25 meV	9184	6479
50 meV	6494	4581
75 meV	5302	3740
100 meV	4592	3239
200 meV	3247	2290

- the beam temperature can be lower (1 meV corresponds to 11.6 K).

The first feature could be used in a storage ring but it is counter-balanced by the very small rate of atoms produced. The long vacuum pipe configuration does not have this limitation. In the case of p^+ , the number of atoms is only limited by the acceleration to 100 keV. Contrary to the charge exchange process, there are no losses in the nitrogen gas. In the case of p^- tables 15 and 16 show the number of atoms produced in the $2S$ state per burst: these numbers can be enhanced by a factor ≈ 50 with a laser. Finally the long vacuum pipe configuration allows the measurement to be made on matter and anti-matter produced in the same conditions: the relative comparison of the Lamb shift is free from systematic effects related to the anti-matter production.

7.2. The boosted spectroscopy

The possibility of producing H and \bar{H} beams at a kinetic energy of 1.5 GeV suggests a new type of spectroscopy. This kinetic energy corresponds to table 4. Let's consider the configuration shown in figure 4.

The light frequency in the laboratory reference frame and in the beam reference frame are referred to as follows:

- ν_0 is the frequency of the light, emitted by the laser, in the laboratory reference frame,
- ν_1 is the frequency of the light in the atom

Table 16

Number of recombined atoms in the $2S$ state per burst without laser gain

T_{\perp}	5 MeV	1.5 GeV
0.5 meV	6518	279
1.0 meV	4609	197
1.5 meV	3763	161
2.0 meV	3259	140
25 meV	922	39
50 meV	652	28
75 meV	532	23
100 meV	461	20
200 meV	326	14

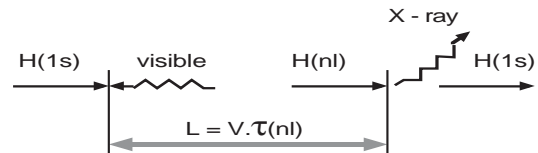


Figure 4. Source and echo in boosted spectroscopy

reference frame,

- ν_2 is the frequency of the light emitted by the atom, in the laboratory reference frame.

These frequencies are related by:

$$\nu_2 = \sqrt{\frac{1+\beta}{1-\beta}} \times \nu_1 \quad (23)$$

$$\nu_1 = \sqrt{\frac{1+\beta}{1-\beta}} \times \nu_0 \quad (24)$$

$$\nu_2 = \frac{1+\beta}{1-\beta} \times \nu_0 \quad (25)$$

The large Doppler factor (≈ 5) allows the use of common lasers with wave lengths in the visible range $\lambda \approx 500 \text{ nm}$ to excite any state from the ground state. Conversely, the light re-emitted from the atom in the transition to the ground state is seen in the laboratory reference frame as X-rays with $\lambda \approx 25 \text{ nm}$. In the following, the excitation light at $\lambda \approx 500 \text{ nm}$ is called the *source*, while the X-rays with $\lambda \approx 25 \text{ nm}$ are called the *echo*. Two measurements are available:

- the energy level of an excited state,
- the life time of the excited state.

To measure the energy level of an excited state, the source wave length is carefully tuned around the transition while the intensity of the echo is recorded. A fit on the intensity curve is made to locate the maximum and the corresponding wave-length of the source gives the energy of the level.

To measure the life time, the point where the laser is focused and the atoms are excited is scanned until the intensity of the echo is maximum: the time of flight of the atoms between this point and the detection region is equal to the life time of the excited states multiplied by the relativist boost $\gamma = 2.6$.

In the first measurement, the precision is limited by the velocity spread of the atoms:

$$\frac{\Delta\nu_2}{\nu_2} = \frac{\Delta\nu_0}{\nu_0} + \frac{2\Delta\beta}{1-\beta^2} \quad (26)$$

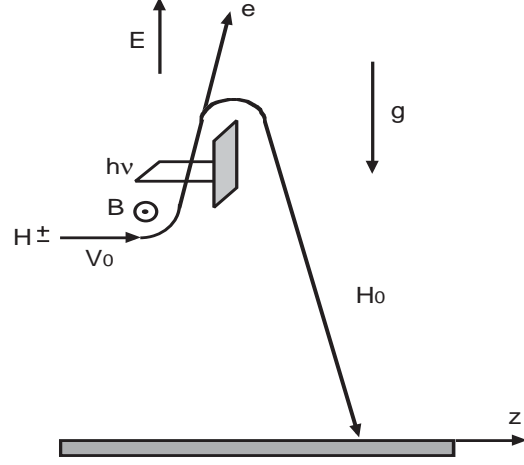


Figure 5. Gravitational mass experiment

If we take the same velocity spread as in the TSR experiment then $\Delta\beta = 2 \cdot 10^{-4}$. Inserting it into equation 26 and considering that the excited states of the hydrogen are at $\approx 10 \text{ eV}$ the error on the energy level is given by:

$$\Delta h\nu_2 = \Delta h\nu_0 + 27 \text{ meV} \quad (27)$$

In the second measurement, if the maximum is located with 1 cm precision, then since the atom velocity is $2.7 \cdot 10^8 \text{ m.s}^{-1}$ the error is $\approx 0.037 \text{ ns}$.

In table 10 the number of anti-hydrogen atoms produced at a temperature of 25 meV is 1100, but this number can be easily enhanced by a factor ≈ 20 with a laser induced gain. These anti-hydrogen atoms are collimated in a cone of radius equal to 1.8 cm according to table 11. With these parameters there is no difficulty in detecting the X-rays of the echo signal. Furthermore, due to the boost, 80% of the echo light is emitted in the forward direction of the beam in the laboratory reference frame. Simple detectors made of scintillators leading to photomultipliers can collect the signal: no unconventional technology is required.

7.3. The gravitational mass measurement

The anti-matter mass measurements performed until now are performed on elementary charged particles trapped in a magnetic field: they are inertial mass measurements [21]. Experiments are underway, [20,22] where p^- and e^+ plasma are cooled, and then merged, to produce \tilde{H} nearly at rest in the laboratory frame. It might be possible then, to perform gravitational mass experiments. The long vacuum pipe configuration offers another path to perform this experiment using the \tilde{H}^+ ion. Since the recombination of p^- and e^+ takes place inside an e^+ beam, some of the \tilde{H} atoms produced capture an e^+ to form the \tilde{H}^+ ion. The \tilde{H}^+ ion can then be slowed down to a velocity of a few $cm.s^{-1}$ and manipulated before the e^+ is removed by a laser pulse. The acceleration and manipulation of the ion has been performed [23] in the past up to $800 MeV$ and there is no difficulty in decelerating, manipulating and removing the e^+ .

The experience is shown in figure 5. The notations are as follows:

- V_0 is the ion velocity,
- B is the magnetic field to tune the parabolic flight,
- g is the gravitational field,
- E is a weak electrostatic field that is used to remove the e^+ ,
- $h\nu$ is the laser light used to separate \tilde{H} from e^+
- \tilde{H}_0 is the atom in the ground state,
- z is the distance before \tilde{H} reaches the detector.

The magnetic field and the initial velocity allows to tune the parabolic flight. The e^+ is separated from the \tilde{H} by a laser with a wave length slightly below the binding energy ($0.75 eV$) and a mirror so that the separation is made through a two photon process. In such a process, the ion can absorb the two photons in three different ways:

- the two photons come from the laser,

Table 17

$$\sigma(e^+ + \tilde{H} \rightarrow \tilde{H}^+) = \sigma(e^- + H \rightarrow H^-)$$

e^+ kinetic energy (eV)	cross-section $\times 10^{-22} cm^2$
0.135	0.456
0.406	0.570
0.677	0.583
1.354	0.547
2.708	0.455
5.416	0.315
8.124	0.300
10.832	0.270

- the two photons come from the mirror,
- one photon comes from each side.

Therefore there can be momentum transfer from each of the sides or no transfer at all giving three separated signals along the z axis. These signals allow to monitor the effect of the laser and to avoid systematic effects due to momentum transfer during the separation.

The magnetic field, the velocity and the distance are sufficient to make the gravitational mass measurement but another parameter can be added to obtain redundancy. The laser can be pulsed to define the starting time of the flight and the time of flight can be detected when the atom reaches the detector.

Such a configuration allows to make a free fall experiment, atom per atom without interference with residual electric fields since the atoms are neutral.

The radiative attachment cross-section is known theoretically and experimentally with precision [18] due to its important role in astrophysics observations and its values are shown in table 17. This cross-section tends to zero when the kinetic energy of the e^+ decreases toward zero. The kinetic energy is taken in the reference frame of the atom. To choose a suitable kinetic energy for e^+ , one has to take into account the affinity $e-H$ which is equal to $0.75 eV$. Above this energy, the ions start to be dismantled by e^+ col-

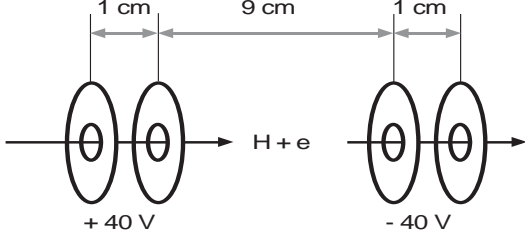


Figure 6. One pair of electrostatic cells for ion production

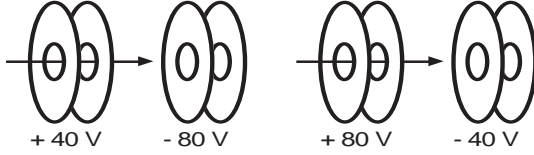


Figure 7. Stack of electrostatic cells for ion production

lisions. The chosen compromise is to operate at a kinetic energy of 0.4 eV . This energy is achieved by pairs of electrostatic cells of 1 cm length with an electric potential of 40 V as shown in figure 6. The cells are separated by a space without field, the length of which is 9 cm . It is in that space that the ion production takes place. Since the kinetic energy gained in one cell is compensated in the next one, the e^+ kinetic energy is unchanged after crossing one pair of cells.

If several pairs of cells are stacked, then to avoid cumulative delay between the e^+ beam and the \tilde{H} beam, the direction of the electric field is alternated, hence the internal cells have an electric potential of 80 Volts as shown in figure 7.

The number of ions produced between a pair of cells is computed with the following approximation: the kinetic energy of e^+ is constant and

equal to 0.4 eV (the spread of kinetic energy in the e^+ beam is few meV and it is neglected).

The parameters are:

- velocity of the e^+ in the \tilde{H} reference frame:
 $V = 375107 \text{ m.s}^{-1}$,
- active length of the cell: $L = 9 \text{ cm}$,
- density of e^+ : $n_e = 5 \cdot 10^{13} \text{ cm}^{-3}$,
- number of \tilde{H} : $N_{\tilde{H}} = 10^6$,
- the cross-section: $Q = 0.570 \cdot 10^{-22} \text{ cm}^2$,
- the relativistic boost $\gamma = 1$.

The number of ions produced in each cell is given by:

$$N(\tilde{H}^+) = n_e N_{\tilde{H}} \frac{L}{V} Q = 0.02565 \quad (28)$$

Over a length of 100 m the number of cells is 1000. Then the number of ions \tilde{H}^+ produced per burst is: $N(\tilde{H}^+) = 25.65$.

8. The production of p^-

The hypotheses for the production of antiproton beams are based on the the Antiproton Decelerator (AD) operated at CERN [24,25]. The main characteristics of the AD are presented now.

The parameters of the primary beam of p^+ are:

- momentum: $26 \text{ GeV}/c$,
- diameter: 1 mm ,
- number of particles per pulse: $1.5 \cdot 10^{13}$

This beam interacts with an iridium target to produce p^- which are collected in a magnetic horn [26]. The interaction produces $5 \cdot 10^7 p^-$ with a momentum of 3.5 GeV . The limit to avoid the target destruction is $6 \cdot 10^{14} p^+$ per GeV/c which corresponds to $2.5 \cdot 10^{13}$ for the CERN p^+ beam. After being collected, the p^- are alternatively cooled and decelerated. There are two stochastic cooling steps and two electron cooling steps. Finally the p^- beam is delivered to the experiments with a momentum of $100 \text{ MeV}/c$ (kinetic energy of 5 MeV) and a momentum spread of

$\Delta p/p \approx 5 \cdot 10^{-4}$. The AD delivers pulses of approximately $3 \cdot 10^7 p^-$.

After the AD, the ASACUSA experiment [27] uses an inverted Linac which further reduces the kinetic energy to approximately $63 \pm 60 \text{ keV}$. This last deceleration has a yield of 40%.

The main limitation of the AD comes from the fact that it cannot be heated to remove adsorbed particles and the vacuum is 10^{-10} torr . There is no use to further cool down the p^- beam because the collisions with the residual gas heat the beam. The beam temperature is approximately 200 meV .

On ion storage ring experiments, electrons with a temperature of $10 \sim 20 \text{ meV}$ are obtained with inhomogeneous magnetic fields. These electrons can be used to further cool down a p^- beam, provided that there are no other effects which heat the beam.

At low temperature, below few meV , beams are expected to reach a crystalline state [31] where the relative positions of the particles in the beam is constant. Studies [29] have shown that in a machine where the particles are focussed with an alternating gradient quadrupole system (4-AG), the intrabeam scattering and the space-charge tune shift prevent the crystalline state to be created. Conversely, it is predicted [30] that if alternating gradient octupoles (8-AG) are used these limitations can be removed. In the case of e^\mp , an alternative design was proposed where the particles are guided along a strong magnetic field: a machine of this type, called LEPTA [32,33], is being built at the Joint Institute for Nuclear Research in Dubna, Russia. The first experimental results are expected in 2002.

From the AD parameters and from the study of an anti-proton accumulation and deceleration ring at KEK in Japan [28], an extrapolation shows that for the production of beams with $10^5 \sim 10^6 p^-$, a primary beam of $3 \cdot 10^{13} p^+$ at $12 \sim 14 \text{ GeV}$ is required. This beam would be produced in two steps:

- a first accelerator ring up to 4 GeV to produce relativistic p^+ ,
- a second accelerator to boost the energy to $12 \sim 14 \text{ GeV}$.

No technical novelty is required for the second accelerator.

At $12 \sim 14 \text{ GeV}$, the p^- are produced with a momentum of $2 \sim 3 \text{ GeV}/c$ and the collection and deceleration scheme would be similar to that of the AD. But the cooling/deceleration would be extended further to create crystalline beams. The first accelerator shall also be used as the decelerator. In order to reach lower temperature, it shall be an octupole machine where p^- beams can reach the crystalline state. The e^+ would be cooled and stored in a small ring (length of $\approx 20 \text{ m}$) similar to LEPTA.

9. Conclusion: target parameters

The experimental configuration presented in this article allows to produce atoms of antihydrogen in the form of beams. The computations lead to the following target parameters for the proposed experiments:

- p^- kinetic energy: $K(p^-) = 100 \text{ keV}$,
- number of p^- particles per beam: $N_p = 10^6$,
- density of the e^+ particles: $5 \cdot 10^{13}$,
- temperature of the p^- and e^+ beams: $T \leq 5 \text{ meV}$.

These parameters can be relaxed by increasing the gain from 10 to 500: the density of e^\mp can be relaxed by 1 or 2 order of magnitude, the kinetic energy of the p^\pm can be increased up to $5 \sim 100 \text{ MeV}$ and the temperature up to 100 meV .

For the boosted spectroscopy experiment the target kinetic energy is 1.5 GeV .

10. Acknowledgments

I wish to thanks D. Möhl and S. Maury from the CERN AD group who have guided me in the comprehension of the Antiproton Decelerator operation and who have suggested to look into crystalline beams. I wish to thanks also I. N. Meshkov for explaining and presenting me the status of electron cooling. I am grateful to Chi-Yu Hu who carefully presented the use of positronium

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11. Appendix

The recombination coefficient $\alpha(nl)$ is given by:

$$\alpha(nl) = \frac{2}{\Delta_{\perp}^2} \int_0^{\infty} \sigma_{rec}(nl) v^2 \exp\left(-\left(\frac{v_{\perp}}{\Delta_{\perp}}\right)^2\right) dv \quad (29)$$

For $n = 1$ the cross-section $\sigma_{rec}(1)$ can be approximated [2] by:

$$\sigma_{rec}(1) = \frac{E_0}{E_e} 1.676 \cdot 10^{-22} \text{ cm}^2 \quad (30)$$

Where E_0 is the ionization energy, 13.6 eV and E_e is the e^- kinetic energy.

For $n \leq 3$ the cross-sections are given by:

$$\sigma_{rec}(2) = 0.550 \times \sigma_{rec}(1) \quad (31)$$

$$\sigma_{rec}(2s) = 0.15 \times \sigma_{rec}(1) \quad (32)$$

$$\sigma_{rec}(2p) = 0.40 \times \sigma_{rec}(1) \quad (33)$$

$$\sigma_{rec}(3) = 0.380 \times \sigma_{rec}(1) \quad (34)$$

$$\sigma_{rec}(3s) = 0.05 \times \sigma_{rec}(1) \quad (35)$$

$$\sigma_{rec}(3p) = 0.153 \times \sigma_{rec}(1) \quad (36)$$

$$\sigma_{rec}(3d) = 0.177 \times \sigma_{rec}(1) \quad (37)$$

Therefore the recombination coefficients for $n \leq 3$ are obtained from the value of the integral:

$$I_1 = \int_0^{\infty} \frac{2}{\Delta_{\perp}^2} \cdot \frac{v^2 E_0}{\frac{1}{2} m_e v^2} \cdot \exp\left(-\left(\frac{v}{\Delta_{\perp}}\right)^2\right) dv \quad (38)$$

$$I_1 = \int_0^{\infty} \frac{4 E_0}{\Delta_{\perp}^2 m_e} \cdot \exp\left(-\left(\frac{v}{\Delta_{\perp}}\right)^2\right) dv \quad (39)$$

$$x = \frac{v}{\Delta_{\perp}} \quad dv = \Delta_{\perp} dx \quad (40)$$

$$I_1 = \frac{4 E_0}{\Delta_{\perp} m_e} \int_0^{\infty} e^{-x^2} dx \quad (41)$$

$$t = x^2 \quad dx = \frac{dt}{2\sqrt{t}} \quad (42)$$

$$I_1 = \frac{4 E_0}{\Delta_{\perp} m_e} \int_0^{\infty} e^{-t} \frac{dt}{2\sqrt{t}} \quad (43)$$

$$\int_0^{\infty} e^{-t} \frac{dt}{2\sqrt{t}} = \frac{1}{2} \Gamma\left(\frac{1}{2}\right) = \frac{\sqrt{\pi}}{2} \quad (44)$$

Finally:

$$\alpha(1) = \frac{2 E_0 \sqrt{\pi}}{\Delta_{\perp} m_e} \times 1.676 \cdot 10^{-26} \text{ m}^3 \cdot \text{s}^{-1} \quad (45)$$

For $n > 3$ the cross-section $\sigma_{rec}(n)$ can be approximated [2] by:

$$\sigma_{rec}(n, E_e) = \frac{E_0^2 \times 2.11 \cdot 10^{-22} \text{ cm}^2}{n E_e (E_0 + n^2 E_e)} \quad (46)$$

This expression can be split into two parts:

$$\frac{1}{E_e (E_0 + n^2 E_e)} = \frac{A}{E_e} + \frac{B}{E_0 + n^2 E_e} \quad (47)$$

with

$$A = \frac{1}{E_0} \quad B = -\frac{n^2}{E_0} \quad (48)$$

The recombination coefficients for $n > 3$ are obtained from the value of the integral:

$$\alpha(n) = \int_0^{\infty} \frac{2}{\Delta_{\perp}^2} \sigma_{rec}(n, E_e) v^2 \exp\left(-\left(\frac{v}{\Delta_{\perp}}\right)^2\right) dv \quad (49)$$

$$E_e = \frac{1}{2}mv^2 \quad dv = \frac{1}{\sqrt{2m}} \frac{dE_e}{\sqrt{E_e}} \quad (50)$$

$$\alpha(n) = \frac{2}{\Delta_{\perp}^2} \times \frac{2}{m} \times \frac{1}{\sqrt{2m}} \times \int_0^{\infty} \sigma_{rec}(n, E_e) \sqrt{E_e} \exp\left(-\frac{2E_e}{m\Delta_{\perp}^2}\right) dE_e \quad (51)$$

$$x = \frac{2E_e}{m\Delta_{\perp}^2} \quad dE_e = \frac{m\Delta_{\perp}^2}{2} dx \quad (52)$$

$$\alpha(n) = \Delta_{\perp} \int_0^{\infty} \sigma_{rec}(n, x) \sqrt{x} e^{-x} dx \quad (53)$$

$$\alpha(n) = 2.11 \cdot 10^{-22} \text{ cm}^2 \times \frac{\Delta_{\perp} E_0^2}{n} \times I_n \quad (54)$$

And $I_n = I_A + I_B$ with:

$$I_A = \frac{2A}{m\Delta_{\perp}^2} \int_0^{\infty} \frac{1}{\sqrt{x}} e^{-x} dx \quad (55)$$

$$I_A = \frac{2A}{m\Delta_{\perp}^2} \Gamma\left(\frac{1}{2}\right) = \frac{2\sqrt{\pi}}{mE_0\Delta_{\perp}^2} \quad (56)$$

$$I_B = B \int_0^{\infty} \frac{1}{E_0 + \frac{n^2 m \Delta_{\perp}^2 x}{2}} \sqrt{x} e^{-x} dx \quad (57)$$

$$u = E_0 + \frac{m n^2 \Delta_{\perp}^2}{2} x \quad (58)$$

$$x = (u - E_0) \times \frac{2}{m n^2 \Delta_{\perp}^2} \quad (59)$$

$$dx = \frac{2}{m n^2 \Delta_{\perp}^2} du \quad (60)$$

$$I_B = B \cdot \frac{2}{m n^2 \Delta_{\perp}^2} \cdot \sqrt{\frac{2}{m}} \cdot \frac{1}{n \Delta_{\perp}} \times \int_0^{\infty} \frac{\sqrt{u - E_0}}{u} \exp\left(-\frac{2(u - E_0)}{m n^2 \Delta_{\perp}^2}\right) du \quad (61)$$

$$t = u - E_0 \quad a = \frac{2}{m n^2 \Delta_{\perp}^2} \quad (62)$$

$$I_B = B \cdot \frac{2}{m n^2 \Delta_{\perp}^2} \cdot \sqrt{\frac{2}{m}} \cdot \frac{1}{n \Delta_{\perp}} \times \int_0^{\infty} \frac{\sqrt{t}}{E_0 + t} \cdot e^{-at} dt \quad (63)$$

$$u = \sqrt{t} \quad dt = 2u du \quad (64)$$

$$\int_0^{\infty} \frac{\sqrt{t}}{E_0 + t} e^{-at} dt = \int_0^{\infty} \frac{2u^2}{E_0 + u^2} e^{-au^2} du \quad (65)$$

$$= 2 \int_0^{\infty} e^{-au^2} du - 2 \int_0^{\infty} \frac{E_0}{E_0 + u^2} e^{-au^2} du \quad (66)$$

Using the formula 7.4.11, 6.1.1 and 6.1.8 from the "Handbook of Mathematical Functions" [3], one gets:

$$\int_0^{\infty} \frac{E_0}{E_0 + u^2} \cdot e^{-au^2} du = \frac{\pi}{2} \cdot \sqrt{E_0} \cdot e^{aE_0} \cdot \text{erfc}(\sqrt{aE_0}) \quad (67)$$

$$\int_0^{\infty} e^{-au^2} du = \int_0^{\infty} \frac{1}{\sqrt{t}} e^{-at} dt = \frac{1}{2\sqrt{a}} \cdot \Gamma\left(\frac{1}{2}\right) = \frac{1}{2} \sqrt{\frac{\pi}{a}} \quad (68)$$

Grouping the two integrals and replacing B by its value leads to:

$$I_B = -\frac{n^2}{E_0} \cdot a \cdot \sqrt{\frac{2}{m}} \cdot \frac{1}{n \Delta_{\perp}} \times \quad (69)$$

$$\left(\sqrt{\frac{\pi}{a}} - \pi \sqrt{E_0} e^{aE_0} \operatorname{erfc}(\sqrt{aE_0}) \right)$$

Summing I_A and I_B and replacing a by its value in the factorization gives a cancellation of the two largest terms:

$$\frac{\Delta_{\perp} E_0^2}{n} \times I_n = \frac{2\sqrt{\pi} E_0}{n m \Delta_{\perp}} - \frac{2\sqrt{\pi} E_0}{n m \Delta_{\perp}} + \quad (70)$$

$$\frac{2\sqrt{2} \pi E_0 \sqrt{E_0}}{n^2 m \sqrt{m} \Delta_{\perp}^2} \cdot e^{aE_0} \cdot \operatorname{erfc}(\sqrt{aE_0})$$

Finally combining equations 54 and 70 gives the recombination coefficients for $n > 3$:

$$\alpha(n) = 2.11 \cdot 10^{-26} m^2 \times \quad (71)$$

$$\frac{2\sqrt{2} \pi E_0 \sqrt{E_0}}{n^2 m \sqrt{m} \Delta_{\perp}^2} \cdot e^{aE_0} \cdot \operatorname{erfc}(\sqrt{aE_0})$$

It shall be noted that this formula involves the multiplication of a very small number with a very large one:

$$\operatorname{erfc}(\sqrt{aE_0}) \sim e^{-374} \quad e^{aE_0} \sim e^{+370} \quad (72)$$

Therefore, to perform the numerical computations in this paper the Mathematica software was used requiring a precision of 40 digits. Alternatively, serie expansions can be used for these functions ([3] 7.1.2.8), provided that terms of the same order are multiplied together before performing the summation.

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