A new path toward gravity experiments with antihydrogen

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Abstract

We propose to use a 13 keV antiproton beam passing through a dense cloud of positronium (Ps) atoms to produce an \( H^+ \) “beam”. These ions can be slowed down and captured by a trap. The process involves two reactions with large cross-sections under the same experimental conditions. These reactions are the interaction of \( \bar{p} \) with Ps to produce \( H \) and the e\(^+\) capture by \( H \) reacting on Ps to produce \( H^+ \). Once decelerated with an electrostatic field and captured in a trap, the \( H^+ \) ions could be cooled and the e\(^+\) removed with a laser to perform a measurement of the gravitational acceleration of neutral antimatter in the gravity field of the Earth.

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1. Introduction

The measurement of the gravitational mass of antiparticles has motivated physicists since over three decades. It is interesting to note that general relativity would not contradict antiparticles to “fall up” in the gravity field of the earth [1]. Experiments to test such ideas have been proposed on positrons and antiprotons [2], but have never succeeded. Several experiments at CERN are now producing neutral antimatter in the form of antihydrogen or antiprotonic helium atoms [3,4] for CPT tests. The trapping of neutral antihydrogen atoms is the next step for these experiments. Some proposals to measure the fall of these atoms have been presented [5]. The possibility to measure the free fall of positronium has also been studied [6].

Recently, it has been proposed [7,8] to measure the gravity acceleration of antimatter using the \( H^+ \) ion. The advantage of this ion is that it can be cooled down to 13\( \mu \)K, a temperature suitable for a gravity experiment. The authors do not investigate in detail the \( H^+ \) production scheme but suggest the interaction of Ps Rydberg atoms with \( \bar{H} \) atoms [9,10]. In this article, we concentrate on the \( H^+ \)
production scheme and propose a “beam to cloud” experimental configuration instead of a trapped system. This configuration allows the isolation of $\text{H}^+$ from the production region.

The positronium cloud, hereafter referred to as “the target” is not contained: it is made of positronium atoms emitted from an aluminum crystal surface, which is bombarded by a flux of positrons. The atoms are emitted in a direction normal to the surface with a few mrad angular spread [11]. Their energy is \( \sim 2 \text{ eV} \) and their speed is \( \sim 1 \text{ mm/ns} \). The density of the Ps target is proportional to the positron flux. As explained further, an amount of order \( 10^{11} \) positrons is needed to get the required density for the Ps target. However, the short Ps lifetime requires this amount to be delivered in a few nanoseconds while the highest foreseen rates of slow positrons are at best \( 10^{14} \text{ s}^{-1} \). The positrons have then to be accumulated in the crystal vicinity and accelerated toward the crystal. To counteract the effects of space charge, the positrons are accumulated in a small neutral \( \text{e}^-\text{e}^+ \) “plasma” for a short time. The plasma size is \( \sim 3 \text{ mm}^2 \times 1 \text{ cm} \). Then an electrostatic field accelerates the positrons towards an aluminum crystal, where they are converted into positronium atoms.

A 13 keV antiproton beam is guided parallel to the crystal at a distance of \( \sim 150 \mu\text{m} \). The target length is crossed in 6 ns by the 13 keV antiprotons.

The flux of positronium atoms is maintained as long as the flux of positrons toward the crystal is kept. It is the flux of positronium atoms, which constitutes the “target”. The overall layout for such a gravity experiment is presented in Fig. 1.

Apart from the gravity experiment, the technique to accumulate positrons in the plasma volume described in this article could prove useful as a first step toward the realization of a positronium [12] BEC, a 511 keV laser [13] and the observation of the antimatter molecule Ps$^2$. If Greaves–Surko traps [14–16] are available with a capacity to hold \( 10^{12} \) positrons and provided that the neutral plasma can be held during the time required to empty the trap, one can produce a density of positronium of \( \sim 0.3 \times 10^{14} \text{ cm}^{-3} \). An experiment to observe the stimulated annihilation process would then become feasible.

If traps with such capacity are not available, an alternative path for a stimulated annihilation observation could be to use the neutral plasma in an MCEO trap [17]. The positronium atoms produced by the 3 body reaction are in Rydberg states. It takes a few \( \mu\text{s} \) for them to reach the ground state. There, the magnetic field couples the triplet state to the singlet state, which has a lifetime of 0.125 ns. Therefore, the lifetime of the positronium produced inside the plasma is dominated by the decay time toward the ground state. Using an infrared laser onto the Ps$^*$ gas, one can re-ionize some atoms while preventing the decay into states with a low level of excitation. Hence, the lifetime can be increased and, in principle, longer trains of positrons could be filled into the plasma volume.

![Fig. 1. Atoms/ions production scheme.](image)
This configuration is also well suited for spectroscopy experiments where atoms containing a positron, or a positronium atom \([18,10]\) are produced by the interaction of atoms/ions with the target. The incoming antiproton beam is then replaced by an ion or atom beam.

The path described in this article to produce an antimatter ion beam is the following:

- the accumulation of positrons in a neutral \(e^-e^+\) plasma,
- the separation of \(e^-\) and \(e^+\) by an electrostatic field and the interaction of the \(e^+\) with an aluminum crystal to produce a positronium cloud, referred to as the target,
- the interaction of 13 keV (anti)protons with the target to produce the atoms and the ions as a beam.

The beam-to-cloud configuration presented in this article is made of several devices, which were developed separately for various applications:

- the 10 MeV \(e^-\) beam on a thin foil to produce an intense \(e^+\) source,
- the buffer gas and Greaves–Surko trap,
- the MCEO trap,
- the Charge Focusing Aluminum Converter (CFAC).

The present study takes the parameters of these devices as they appear in the literature and shows that with little modifications, these devices can be assembled to produce an \(\overline{\Pi}^+\) ion beam.

The steps involved in the target production are discussed in the next section. The atom and ion production rates are presented in Section 3. Several technical features are discussed in Section 4.

2. The positronium target

The positronium target is obtained by accelerating positrons from an \(e^-e^+\) plasma toward an aluminum crystal. The positrons hit the crystal with a kinetic energy above 40 eV to avoid elastic and specular reflection \([19]\).

We describe a way to produce the neutral plasma and the subsequent extraction and focalisation of the positrons from the plasma onto the Al crystal in the following.

We foresee two modes of operation for the creation of the neutral plasma:

- the slow loading mode, where the positrons are extracted continuously from the buffer gas section of the Greaves–Surko trap, which cools them to room temperature (≈25 meV) (the loading time is a few seconds),
- the fast loading mode where the positrons are stored and cooled to 2 meV in a Greaves–Surko trap \([16]\) and extracted in \(\sim10\mu s\). This fast extraction heats the positron beam. This mode of operation requires less confinement time for the neutral plasma and may allow higher plasma densities.

The electrons are provided by a buffer gas or magnetic trap depending on the desired temperature. Since electrons do not annihilate on the container walls and are easy to produce, the final neutral plasma temperature is tuned by setting the electron beam temperature.

We assume that the positrons are produced continuously through the interaction of an intense beam of 10 MeV electrons with a thin tungsten foil \([20]\) with a rate, after solid Neon moderation, of \(10^{11}s^{-1}\). The positrons are cooled to room temperature (≈25 meV) in a buffer gas. In the slow loading mode, the positrons are taken at the exit of the buffer gas section, while in the fast loading mode they are first accumulated in a Greaves–Surko trap \([14–16]\), where they are stored and cooled to \(\sim2\) meV. The low temperature of the positrons is a feature of the Greaves–Surko trap: the strong magnetic field is produced by a supraconducting magnet and the positrons are in thermal equilibrium with the magnet container at a temperature of 2 meV. Such a low temperature enhances the 3-body reaction which absorbs the positrons if the neutral plasma were to be stored in this trap.

The preferred device for accumulating the \(e^+\) in a neutral plasma is the MCEO trap \([17]\), which does not require a very low temperature such as
the one reached in a Greaves–Surko trap. Therefore, the heating of the positrons can be accommodated and is beneficial to accumulate enough positrons in the neutral plasma.

The positrons are guided by a 100 G magnetic field, from the Greaves–Surko trap (or the buffer gas), to the neutral plasma. The injection pipe is shielded to minimize its leaking field.

The original MCEO trap is a magnetic cusp with an octupolar electric field. It is made of several pairs of coils of 10 cm radius arranged symmetrically with respect to the center of the trap. The most central coils are located at 4 cm from the center. The other coils are separated from each other by 5 cm. The currents in each pair of symmetric coils are of opposite sign. The current amplitude can be varied for each pair.

It is designed to mix both positrons and antiprotons. It is to be implemented in the ASACUSA beam line at CERN under the name of Musashi trap. In our application this trap is meant to mix positrons with electrons. Therefore, a scaled down version has to be designed.

The scaled down version of the MCEO trap has the following parameters: the magnetic coils radius is 1.5 cm, the most central coils are at 1.2 cm from the center, the next coils are separated by 1.5 cm, the total current for each set of coils is 4 k Ampere-turns, the octupolar electrostatic field is set by a central electrode at 70 V and a length of 1.0 cm. The key feature of this trap is a null field region in its center (Fig. 2). Once a charged plasma is loaded, it acts as a potential well for the opposite charge. We use electrons to create the well for the positrons. The neutral plasma will be located at the center of the well. The $e^-$ and $e^+$ beams are injected along the $x$-axis. The octupolar electrostatic field of the MCEO trap is simulated using an analytical form [21].

A simple trap made of two coils acting as mirrors [22] is also considered, but the injection of the positrons is more difficult.

In order to extract the positrons from the neutral plasma and accelerate them toward the crystal, an electrostatic field is used. The electrode configuration is shown in Fig. 3.

The electrodes are arranged in two parallel planes. The cathode is made of Al crystal. Its surface is $100 \mu m \times 1 \text{ cm}$. The neutral plasma is located in the volume between the two planes. This configuration will be referred to as Charge Focusing Aluminum Converter or CFAC.

This configuration is inspired by a technology developed for high-energy physics detectors called the Micro-Strip Gas Chamber [23], or MSGC. The distances between electrodes and field magnitude are taken from this development.

In the CFAC, the electrostatic field of the MSGC is modified by introducing 4 wires to smooth the side field as shown in Fig. 4.

The CFAC was put in the center of the trap by the simulation described below in order to get an estimation of the positronium density and time distribution. The insertion of metallic conductors
in the center of the MCEO trap is unrealistic. An integrated design is proposed in the discussion section.

A fortran program was written to simulate:

- the stability of the neutral plasma during the 10 \( \mu \)s loading time,
- the acceleration by the CFAC electrostatic field and the time distribution of the positrons hitting the Al crystal.

2.1. The target simulation

In the MCEO trap the positrons are injected along its axis, parallel to the magnetic field which guides them.

2.1.1. The CFAC electrostatic field

The CFAC electrostatic field is computed using analytical formulae [23]. The gap of the CFAC, i.e. distance between the drift electrode and the cathode planes, is 1.5 cm. The cathode plane is located 1 cm from the \( x \)-axis and parallel to it. The drift electrode size is 1 cm \( \times \) 7 mm. The cathode, made of the Al(111) crystal, has a 100 \( \mu \)m width. The distance between the edge of the cathode and the next electrode in the cathode plane is 200 \( \mu \)m. The drift electrode plane being at 1600 V, the cathode is grounded and the bias voltage is 1550 V. The plasma temperature is 30 meV.

During the loading phase, the CFAC field is set “off”. The acceleration phase starts when the CFAC electrostatic field is set “on”, while the octupole electrostatic field and the cusp magnetic field are set “off”.

The positron capture and the emission of positronium atoms is a fast process, the delay of which on the nanosecond scale can be neglected [24]. Therefore, the time distribution of the positrons hitting the crystal is also the time distribution of the positronium atoms exiting the crystal surface (Figs. 5 and 6).

The positronium atoms exiting the crystal are in the ground state. The magnetic field couples the singlet and triplet states [25], reducing the Positronium lifetime to a minimum of 0.125 ns.

In an experimental test with a MCEO trap and with electrons the radial field on the central electrode surface was 360 G [26]. This trap is designed to hold both positrons and antiprotons. When antiprotons are used the magnetic field is much stronger, but in the beam-to-cloud configuration the antiprotons cross the trap axis and are not kept. Hence, with a weak field we take an effective lifetime larger than 0.200 ns. This lifetime corresponds to a travel distance of 200 \( \mu \)m. The target volume is defined as a box parallel to the crystal, which starts at a distance of 100 \( \mu \)m. The target is 100 \( \mu \)m large, 1 cm long and 100 \( \mu \)m long.
thick, which gives a volume of $10^{-4}\text{cm}^3$. This target volume is centered at a distance of 150\,\mu m from the crystal surface.

An estimate of the target density is obtained by assuming all Ps atoms fly perpendicularly from the crystal surface [11]. The number of positrons stored in the plasma is $N_{e^+} = 10^{11}$. The positron flux from the plasma to the crystal has a duration of $\sim 10$\,ns. The Ps atoms spend only 0.100\,ns in the target volume; hence, the target density is given by

$$n_{Ps} = \frac{10^{11} \times 0.100}{10^{-4}} \varepsilon_{PS,\text{CFAC}}$$

$$= 0.25 \times 10^{13}\text{cm}^{-3}. \quad (1)$$

The target volume could also be counted starting from the crystal surface with some antiproton losses (antiprotons hitting the crystal).

The simulation of the CFAC gives an efficiency of $\varepsilon_{\text{CFAC}} = 0.5$ for a positron to reach the crystal. The positronium emission efficiency from the crystal surface is taken as $\varepsilon_{Ps} = 0.5$.

A correcting factor due to the positronium decay between the crystal and the target volume will reduce the density by less than a factor 2. This correction depends on the magnetic field strength.

### 2.2. Positron losses

When mixing $e^-$ and $e^+$ with a density above $\sim 10^9\text{cm}^{-3}$ and a temperature of a few meV, the plasma collapses into the positronium through a 3-body interaction:

$$e^- + e^+ + e^\pm \rightarrow \text{Ps}(n'l') + e^\pm. \quad (2)$$

The rate of other reactions, namely the radiative recombination and the direct annihilation, is negligible. This reaction and the subsequent positronium annihilation destroy the positrons.

The number of positrons in the neutral plasma as a function of time is given by

$$N_e = \sqrt{\frac{a}{\lambda}} \times \tanh(\sqrt{a\lambda}t) \quad (3)$$

$$\lim_{t\to\infty} N_e(t) = \sqrt{\frac{a}{\lambda}} \quad (4)$$

where $a$ is the injection rate and $\lambda$ is a constant related to the 3-body reaction.

The tables obtained in the appendix show that in the worst case, in $\sim 2$\,s and at a temperature equal or above 4\,meV, it is possible to accumulate $1$See computation in the appendix.
\[ 2 \times 10^{11} \text{e}^+ \text{ in the plasma. For less than } 10 \text{s of accumulation there is little variation with temperature above } 4 \text{meV. Once the injection of positrons ends and during } \sim 1 \text{s the loss of positrons is small and on the time scale of a few } \mu \text{s it can be neglected.}

In order to have a negligible loss by annihilation on the remaining gas, a vacuum environment is required with a residual gas partial pressure below \( \sim 10^{-9} \text{Torr} \).

3. Antihydrogen production

The charge exchange on positronium to produce the atom is a resonant reaction. The cross-section calculations \([27,29,30]\) have been confirmed by an experiment \([28,31]\).

Once the \( \text{H} \) atom is produced, it captures a positron to make the ion. There is a non-resonant channel, the 2-body recombination, and a resonant one, the charge exchange on positronium.

The choice of the antiproton kinetic energy is given by the overlap of the resonant cross-sections. In the region of interest, the antiprotons have a kinetic energy of \( \sim 13 \text{keV} \) (Fig. 7). The atom production cross-section obtained experimentally \([28]\) is \( 7.8 \pi a_0^2 \).\(^2\) At this kinetic energy, the computed cross-section \([32]\) for the ion production is \( \sim 0.05 \pi a_0^2 \) (Fig. 8).

The (anti)hydrogen production rate normalized to the (anti)proton flux is

\[ n_{\text{Ps}} \times \sigma_{\text{Ps}} \times v_p. \tag{5} \]

The (anti)proton speed at 13 keV is \( v_p = 1.58 \times 10^8 \text{cm} \text{s}^{-1} \). Therefore, the normalized rate is \( 0.28 \times 10^6 \text{s}^{-1} \). The target length is 1 cm; hence, the crossing time by the (anti)proton beam is 6.3 ns and the number of antihydrogen atoms produced by each antiproton crossing the target is 0.0017.

The (anti)hydrogen ion production rate normalized to the antihydrogen flux is

\[ n_{\text{Ps}} \times \sigma_{\text{H}^+} \times v_{\text{H}^+}. \tag{6} \]

The normalized rate is \( 0.18 \times 10^4 \text{s}^{-1} \). After convolution for the \( \text{H}^+ \) production in the same 1 cm

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\(^2\)Here \( a_0 \) is the Bohr radius and \( \pi a_0^2 = 0.880 \times 10^{-16} \text{cm}^2 \).
long target, the number of ions produced for each antiproton crossing the target is $10^{-8}$. In the Ps density computation, only the fast component of Ps emitted by the crystal was taken into account: if we consider also the slow component, this number becomes $10^{-7}$.

At the edge of the crystal and at a distance of 150 μm, the CFAC electrostatic field component transverse to the antiproton path is ~0.12 MV/m. Therefore, an antiproton aiming at the edge of the crystal will be pulled transversely by 240 μm on a distance of 1 cm. The transversal electrostatic field, which is null on the axis and grows toward the edges of the target, will cause a maximum pull of ~0.5 mm and a beam divergence of ~50 mrad.

4. Discussion

In the integration of the devices (CFAC, MCEO trap, buffer-gas, Greaves–Surko trap) into a single experiment there is room for changing their parameters: gains from tuning the device parameters and losses are difficult to simulate and have to be established experimentally. When changing the parameters in the simulation these gains/losses are within a factor 10.

Still, there are many technical questions that are not addressed by the simulation. We shall now list some of them.

The field in the buffer gas trap is 1.5 kG and 5 T in the Greaves–Surko trap: a set of iron plates and auxiliary fields shall be designed to reach the 100 G guiding solenoidal field [16].

Positron plasmas with a density of $10^{10} e^+ \text{ cm}^{-3}$ at meV temperatures have been stored in a Greaves–Surko trap: a set of iron plates and auxiliary fields shall be designed to reach the 100 G guiding solenoidal field [16].

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For antihydrogen creation, the very large $e^\pm$ plasma densities discussed above allow to transform the antiprotons with a relatively high efficiency. But antiprotons are difficult to produce. However, in order to test the process one can use protons and an existing trap: the lower plasma density being compensated by a larger number of protons (say $10^8 p$ per beam pulse).

Another aspect of this path for a gravity experiment is that the reactions involved are in all steps charge symmetric: by switching from protons to antiprotons on the time scale of 1 h, it is possible to measure $g$ in the same gravitational field with little or no tide effect due the movement of the moon and other masses.

The integration of the CFAC into an MCEO trap requires some modification. We foresee cutting the central electrode in 4 (or more) sections separated by a thin insulator (Fig. 9). In the trapping mode all the sections are set at the same potential acting as a single electrode. In the CFAC mode the potentials are set to produce an electrical field similar to the one in Fig. 3.
The H$^+$ ion beam is needed for the gravity experiment, but a large number of H atoms are also produced in the beam. Using a laser to stimulate the transition toward the $n = 2$ level, one gets a beam in the $10–20$ keV range. Then the Separated Oscillatory Field method [35–37] allows to measure the $2S_{1/2}$–$2P_{3/2}$ fine structure (Tables 1–3).

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We wish to express our sincere thanks to all the people from different fields with whom we had fruitful discussions: P.K. Biswas, G. Chardin, C. Guyot, B. Mansoulié, C. Surko, R. Greaves, A.P. Mills, J.M. Rax, Y. Yamazaki. One of us is

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**Table 1**

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**Table 3**

| $\sqrt{a/2}$, maximum number of e$^+$ that can be accumulated in the 30 mm$^3$ neutral plasma with an injection rate of $a = 10^{11}$ s$^{-1}$ for $A = 0.76$ ($B = 0$) and $A = 0.07$ ($B = \infty$) |
|-----------|------|------|------|------|------|------|
| $T$ (meV) | 2    | 4    | 10   | 25   | 50        | 100     |
| $A = 0.76$ | $1.81 \times 10^{11}$ | $8.60 \times 10^{11}$ | $6.76 \times 10^{12}$ | $5.31 \times 10^{13}$ | $2.53 \times 10^{14}$ | $1.20 \times 10^{15}$ |
| $A = 0.07$ | $5.95 \times 10^{11}$ | $2.83 \times 10^{12}$ | $2.23 \times 10^{13}$ | $1.75 \times 10^{14}$ | $8.32 \times 10^{14}$ | $3.96 \times 10^{15}$ |
grateful to Chi-Yu Hu [29] who carefully presented the use of positronium to produce antihydrogen at the poster session of the ICPEAC 2001 conference.

Appendix

The technique described in this article can be used for experiments on positronium BEC and a 511 keV laser, because it could reach a positronium density of \( \sim 10^{14} \) or \( \sim 10^{15} \text{ cm}^{-3} \). The appendix gives a numerical estimation of the density as a function of time and temperature.

The maximum number of positrons available to interact with an aluminum crystal is the result of the equilibrium between the injection rate and the 3-body reaction rate. This reaction (see Eq. (2)) and the subsequent positronium annihilation destroy the positrons. The time evolution of the number of positrons in the plasma is given by the injection rate and the 3-body reaction rate [38]:

\[
dN_e = a \, dt - r_3 \, dt,
\]

where \( N_e \) is the number of \( e^+ \) in the plasma volume, \( a \) is the injection rate and \( r_3 \) is the 3-body reaction rate given by

\[
r_3 = A \times n_e^2 \times v_e \times b^5,
\]

where the electron thermal velocity \( v_e = \sqrt{kT/m_e} \) and the impact parameter \( b = e^2/kT \) are numerically \( v_e (\text{cm} \, \text{s}^{-1}) = 4.19 \times 10^7 \sqrt{T} (\text{eV}) \) and \( b (\text{cm} \, \text{eV}^{-1}) = 1.44 \times 10^{-7}/T (\text{eV}) \), resulting in

\[
r_3(s^{-1}) = 2.6 \times 10^{-27} A \times n_e^2 / T^{4.5},
\]

where \( A \) is a parameter which varies with the magnetic field, \( n_e \) is the density in \( \text{cm}^{-3} \) and \( T \) is the plasma temperature in eV.

The neutral plasma volume is \( \sim 30 \text{ mm}^3 \).

Let \( V \) be the plasma volume and let us define \( \lambda \) as

\[\lambda = 2.6 \times 10^{-27} A/(V^2 T^{4.5}),\]

so that \( r_3 = \lambda N_e^2 \).

With the initial condition \( N_e(t = 0) = 0 \), we get

\[dN_e = (a - \lambda N_e^2) \, dt,\]

which leads to

\[N_e = \sqrt{a/\lambda} \times \tanh(\sqrt{a/\lambda} \cdot t), \]

\[\lim_{t \to \infty} N_e(t) = \sqrt{a/\lambda}.
\]

For the two extreme situations, \( B = 0 \) and \( B = \infty \), parameter \( A \) is, respectively, equal to 0.76 and 0.07.

In the computations the ion was supposed to be much heavier than the electron and therefore its trajectory in the field was neglected [38]. Recently, a computation was made with a proton, taking into account its trajectory [39] for fields of a few tesla: the variation of parameter \( A \) compared to the infinite field value was less than a factor of 2. Here, the ion is the positron, which is much lighter, and the magnetic field is a few gauss only. Therefore, the positron behaves as a heavier ion in a stronger field. The expected behavior is then an intermediate one, between the above computation with zero field and with infinite field.

A computation involving only electrons and positrons predicts a 3-body recombination rate twice greater than that for protons and electrons [40].

The computed number of positrons accumulated inside the neutral plasma for several temperatures is tabulated for both values of \( A \).

When the injection in the plasma ends, the evolution is given by the rate of the 3-body reaction:

\[dN_e = -r_3 \, dt = -\lambda N_e^2 \, dt\]

\[N_e(t) = \frac{N_0}{1 + N_0 \lambda t},\]

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