Development and commissioning of ion-optical elements for ion and antiproton beams with energies up to 5 keV

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ABSTRACT: In nuclear and atomic physics experiments, charged ion beams often need to be guided from the ion production to the experimental site. In the PUMA experiment, an ion source beamline was developed, which can be operated with up to 5 keV beam energy at a base pressure of 10^{-9} mbar or better. In this paper, a low-energy pulsed drift tube for beam energy modification, a hybrid einzel lens assembly for beam focusing and steering and an iris shutter assembly for separating beamline sections with different vacuum requirements are described with their design principles and performances.

KEYWORDS: Beam Optics

1 Introduction

The PUMA (antiProton Unstable Matter Annihilation) experiment aims at studying the neutron-toproton ratio in the density tail of stable and exotic nuclei, indicative of nuclear phenomena like halo nuclei and neutron skins by use of low-energy antiprotons [1, 2]. The first goal of the experiment is to characterize the neutron-to-proton ratio of stable isotopes at the Antiproton Decelerator (AD) at CERN in a Penning-Malmberg trap by mixing them with antiprotons. The ions of interest have to obey strict requirements with respect to the beam emittance to allow for an efficient injection into the PUMA Penning trap, and to the isotopic purity for the subsequent annihilations with antiprotons. Thus, a dedicated, versatile ion source beamline was designed to create, transport and shape the beam from the ion source to the PUMA Penning trap.



Figure 1. The ion source beamline of the PUMA experiment consists of four sections: ion production, mass separation, ion cooling and stacking and transfer to the antiproton beamline. The beamline includes einzel lens and steerer assemblies (ELSA), pulsed drift tubes (PDT) and iris shutter apertures (IS). After preparation, the beam is guided towards the PUMA Penning trap.

2 Experimental setup

In figure 1, a schematic view of the beamline is shown. The ions are produced in bunches with energies of up to 5 keV in an electron impact plasma ion source (IQE 12/38, SPECS Surface Nano Analysis GmbH). The beam is then purified in a multi-reflection time-of-flight mass separator (MR-ToF MS), capable of achieving a mass resolving power $m/\Delta m$ of approximately 10⁵ [3]. Further downstream, the bunches are cooled and accumulated in a quadrupole radio-frequency cooler-buncher (RFQcb) [5] which uses the buffer-gas cooling technique [4].

During normal operation, the desired ion species is produced with a kinetic energy of 3 keV. The most critical part in terms of beam transportation is the injection and ejection of the beam into and out of the RFQcb, whose injection cone has an inner diameter of 2 mm. Furthermore, for a successful trapping in the biased RFQcb, the ions have to be decelerated from an energy of about 3 keV to 0.3 keV while keeping the beam spot below 2 mm.

In order to meet these requirements, dedicated ion-optical elements were designed to be installed in between the sections of the beamline: an electrostatic einzel lens in combination with a segmented steerer for steering and focusing, as well as a pulsed drift tube (PDT) for modifying the kinetic energy of the ion beam. Additionally, the strict vacuum requirements (UHV, $< 10^{-9}$ mbar) at the end of the ion transfer section have to be respected. Thus, a separation of the vacuum sections with adjustable apertures based on iris shutters was implemented. In this work, the three beamline elements are presented in detail.

3 Simulation method

The design of the ion-optical elements was determined based on simulations with SIMION[®]2020, a software package used to calculate electric and magnetic fields based on a given configuration of electrodes and magnets by solving the Laplace equation [7]. Extensive descriptions on the

simulation methods used in SIMION[®] can be found, e.g., in reference [8]. All electrodes were defined in a 1 mm grid and refined with a convergence objective of 10^{-7} V.

In the simulation, ${}^{40}\text{Ar}^+$ ions were considered following a Gaussian angular distribution with a FWHM of 1.5° and a Gaussian spatial distribution of 1 mm, corresponding to a transverse FWHM emittance of 8.3π mm mrad, an initial kinetic energy of $3000 \pm 10 \text{ eV}$ FWHM and a uniformally distributed bunch length of 300 ns, which corresponds to a length of 3.61 cm for the initial kinetic energy. The ions were created directly after the ion source.

4 Einzel lens and steerer assembly

4.1 Description

The general design strategy for einzel lenses is, e.g., described in reference [9]. The einzel lens and steerer assembly (ELSA) is shown in Figure 2. It is contained in a CF160 tube of length 20.32 cm. Drawings and dimensions are provided in appendix A.1. The assembly consists of three highly polished aluminum cylinders. Aluminum was chosen as material due to its low weight, cost and low outgassing rates. The lens electrode is sandwiched between a four-fold segmented steerer electrode and a grounded electrode. The grounded electrode is connected directly to the flange. The steerer electrodes are separated from the focus electrode by alumina (Al₂O₃) split bushes (tectra, BGB-M4) to be able to supply high voltage to all electrodes. Their assembly is also shown in appendix A.1. The maximum applied voltage to the electrodes are \pm 5 kV DC, limited by the electrical feedthroughs. The ELSA is mounted on a customised double-sided CF160 flange, with five radially positioned CF-16 flanges, that are equipped with single-pin SHV-5 feedthroughs each (see appendix A.2). Two geometries are considered for the assembly, which are shown in figure 2, one with cylindrical and one with planar steerer electrodes. The ELSA is used successfully as a standard component in the PUMA ion source and antiproton beamline.

4.2 Length of focus electrode and radius of einzel lens assembly

The focal point position z_f and beam spot size r_f were determined in dependence of the applied voltage and the design parameters of the einzel lens assembly. z_f is measured from the center of the einzel lens segment. The beam spot r_f is the mean radius of the ion beam at z_f . Using only DC fields, the results for z_f and r_f can be scaled for different initial beam energies, thus simulations were only performed for one initial beam energy.

Figure 3 shows z_f and r_f in dependence of the applied voltage U for various radii r_e of the ELSA and lengths of the lens electrode l_e . A smaller r_e allows to reach smaller r_f for lower applied voltages. z_f and r_f saturate for $l_e > 70$ mm, which is short enough to contain the ELSA in the CF160 tube. The larger r_e , the higher is the required potential to reach small r_f . For ELSAs used in the case of the PUMA ion source beam line, a radius of 30 mm was chosen. This will allow to sufficiently focus the beam to pass orifices < 2 mm in the beam line with a z_f that corresponds to the distance between ELSA and orifices, using an applied voltage of approximately 2 kV.

4.3 Steerer design

The effect of the steerer shape was simulated for a cylindrical and planar geometry. A steering voltage $\pm V_s/2$ is applied to opposite steering electrodes, leading to a voltage difference V_s . The



Figure 2. The einzel lens is mounted on a customized double sided CF-160 flange and consists of 6 electrodes: a grounding element, a focus element and a four-fold segmented electrode for x-y steering of the beam. Two configurations of steerer plates were tested: cylindrical (left) and planar (right).

achieved angle in the horizontal or vertical plane was plotted for the applied voltage V_s . The results were fitted with a straight line:

$$\Theta_{\text{cyl}}(V_s) = (0.0074/\text{V} \cdot V_s - 0.0034)^\circ, \tag{4.1}$$

$$\Theta_{\text{pla}}(V_s) = (0.0091/\text{V} \cdot V_s - 0.0100)^{\circ}.$$
(4.2)

The planar geometry requires approximately 0.81 V_s to reach the same steering angles as the cylindrical geometry. In the case of the PUMA ion source beamline, a maximum V_s of ±1 kV can be applied, limited by the power supplies and cables used, leading to elevation and azimuthal angle $\Theta_{cyl,max} \approx 7.4^{\circ}$ and $\Theta_{pla,max} \approx 9.1^{\circ}$.

The value of r_f increases for a deflected beam compared to a non-deflected beam. This is furthermore influenced by the applied focus voltage U, e.g., when applying U = 1.8 kV in the planar steerer assembly, r_f increases by 34.8 %, while for U = 2.0 kV, r_f increases by 45.1 %. Further increasing U to 2.2 kV yields an increase of 21.2 %. The cylindrical design has a better r_f stability than the planar design, as can be seen in Fig. 4, especially when focusing diagonally.

4.4 Angular deviation of the beam

The angular deviation describes the opening half-angle of the cone in which the ions are distributed around the central trajectory. It counteracts the focusing power of the einzel lens. Thus, z_f is



Figure 3. The length of the focus electrode l_e and the radius of the lens assembly r_e were varied in dependence of the applied focus voltage U (abscissa) in simulations to determine the impact on the focus point properties z_f and r_f . Positive and negative applied voltages are displayed on the same positive axis, indicated by different line styles. The dashed lines indicate a positive applied voltage, the dashed-dot lines a negative applied voltage.

increasing for higher angular deviations for a fixed applied potential U, see figure 5. Additionally, r_f degrades with higher angular deviation, which will lead to beam losses in low acceptance components. In case of inefficient transmission through a beam line, additional focusing elements might be necessary to compensate for a large angular deviation.

The angular deviation also has an effect on the behaviour of the beam after steering. Again, the planar geometry is more affected by an increase in angular beam deviation and will lead to a bigger r_f compared to the cylindrical geometry. For this reason, the cylindrical geometry was chosen for the ELSAs in the PUMA ion source beamline.

5 Pulsed drift tube

5.1 Description

Pulsed drift tubes have been designed for a large spectrum of applications, the earliest included drift tube acceleration of heavy ions [12]. In more recent applications, the concept of a pulsed drift tube is used, e.g., for the capture of ions in MR-ToF devices with an in-trap lift [13], bunch reacceleration after an RFQcb [14] or the HV deceleration of antiprotons for the injection in particle



Figure 4. Simulated r_f for steering angles Φ , Θ for (left) the cylindrical geometry and (right) the planar geometry for an applied focus voltage U = 2 kV. r_f is larger when steering the beam towards the edges of the steerers, and the effect is pronounced for the planar geometry.



Figure 5. Simulated focal point properties for different FWHM angular deviations of the beam cone for the PUMA ELSA in the cylindrical geometry. The larger the angular deviations, the larger is the minimum r_f that can be reached and the longer is z_f . The dashed lines indicate a positive applied voltage, the dashed-dot lines a negative applied voltage to the focus electrode.

traps [15, 16]. The dimensions of the low-energy PDT can be found in appendix A.3. The PDT assembly is mounted on the same electric feedthrough flange as the einzel lens and is contained in a CF160 tube. It consists of two cylindrical electrodes: a drift tube and an isolated electrode which are made of highly polished aluminum and separated by alumina split bushes. The drift tube is also separated with split bushes from the mounting flange to create a second gap between flange and drift tube. Thus, the drift segment is sandwiched between two isolated segments, which are usually grounded. An additional focusing effect can be achieved by applying high voltage to the isolated electrode.

5.2 Operation principle

A pulsed drift tube modifies the kinetic energy of traversing ions. A de- or accelerating field is created in the gap between the first grounded electrode and the drift tube. After the energy modification, the ions enter an ideally field-free drift region. While the ion bunch is fully contained in the tube, the applied voltage on the tube is switched to a lower or higher value respectively. Thus, upon leaving the PDT, the ions remain in their modified energy state. In the most simple case, the drift tube is switched to or from ground respectively, but other switch patterns can be implemented. Since the phase-space volume of the ion bunch is constant, decelerating the ions results in an increased spatial spread that has to be compensated by the use of ion-optical elements like an einzel lens. An acceleration will lead to a focusing effect.

A PDT can be characterized by relating the switch time t_s to the time-of-flight (ToF). If switched when the ion bunch is outside the drift tube, the energy state of the ions will remain unchanged after traversing the PDT. If the ion bunch is only partly contained in the tube, it will also only be partly affected by the change of field leading to a high energy spread of the resulting bunch. Only if the bunch is fully contained in the tube during switching, all ions can achieve the desired energy state. The longitudinal acceptance of the drift tube has to be appropriately chosen to match the bunch length of the incoming beam.

Accordingly to the description above, also the second gap after the drift electrode can be used for energy modifications. For a larger longitudinal acceptance, the bunch should traverse the tube always in the lower-energy state. Thus, the gap before the drift tube will be used for a deceleration and the gap after the drift tube for an acceleration. Other switch patterns could be implemented, e.g., switching to higher or lower values than ground respectively, which can have an impact on the transversal and longitudinal emittance.

5.3 Calculation of the ideal switch time

Consider a low-energy ion of mass *m* and charge *q* passing through a conductive concentric-to-axis cylinder with a gap of width *S* and inner diameter *D* with a kinetic energy T_0 in positive direction *z*. The first side of the cylinder has an applied voltage of V_0 and length l_0 , the other side V_1 and $2l_1$, see figure 6. Traversing the gap will modify the particles energy to T_1 . The ideal switch time, when the bunch is centered in the tube at l_1 , is calculated for two different assumptions of the field in the energy modification gap.



Figure 6. The ideal switch time can be calculated dependent on the dimensions and applied voltages to the PDT.

5.3.1 Plate capacitor

A step-wise calculation is performed for the drift time of the ions: first, until they reach the gap, second, passing through the gap and third, from the start of the drift tube to it's center. In a simplified view, an electric field of a plate capacitor is assumed inside the gap, the region before and after the gap is field-free. Thus, ions will drift for distances l_0 and l_1 with constant velocity outside the gap, while the electric field E in the gap and consequently the constant acceleration a along the z-axis compute to

$$\vec{E} = \frac{V_1 - V_0}{S} \vec{e_z} \Rightarrow a = \frac{q(V_1 - V_0)}{Sm}.$$
(5.1)

The ideal switch time computes then to

$$t_s = \frac{l_0 \sqrt{m}}{\sqrt{2T_0}} + \frac{S\sqrt{2m}}{\sqrt{T_1} + \sqrt{T_0}} + \frac{l_1 \sqrt{m}}{\sqrt{2T_1}}.$$
(5.2)

Considering no energy modification ($T_0 = T_1$) in the first gap, e.g., when the second gap is used for an energy modification, equation 5.2 simplifies to

$$t_{s,a} = \frac{\sqrt{m}}{\sqrt{2T_0}} (l_0 + S + l_1).$$
(5.3)

5.3.2 Axially aligned cylindrical electrodes

A more realistic potential in a gap between two axially aligned cylindrical electrodes was computed, e.g., in reference [17], from which follows the electrical field and consequently the acceleration, which both depend on the position z of the ion bunch, where z = 0 is in the center of the gap:

$$\vec{E(z)} = -\nabla\phi(z) = \frac{V_1 - V_0}{2S} (\tanh(w(z + S/2) + \tanh(w(z - S/2)))\vec{e_z} \Rightarrow \vec{a} = \frac{q|\vec{E}|}{m}, \quad (5.4)$$

where w = 2.64/D. By integration, the ideal switch time t_s can be determined

$$t_{s} = \int_{-(l_{0}+S/2)}^{l_{1}+S/2} \left(\frac{q(V_{1}-V_{0})}{wSm} \ln\left(\frac{\cosh(w(z+S/2))}{\cosh(w(z-S/2))}\right) + \frac{2T_{0}}{m} \right)^{-\frac{1}{2}} dz,$$
(5.5)

which can be solved numerically, and accounts for both acceleration and deceleration using first or second gap. Using the second gap, which implies $V_0 = V_1$, eq. (5.5) simplifies to eq. (5.3).



Figure 7. The resulting energy *E* and the energy spread ΔE of the deceleration as a plot of the switch time t_s depending on PDT radius r_d (left) and length l_d (right). A ⁴⁰Ar⁺ ion bunch with a bunch length of 300 ns and an emittance of 8.3 π mm mrad was simulated to determine the preferable dimensions.

5.4 Length of drift tube and radius

The PDT was characterized by simulating the deceleration of ${}^{40}\text{Ar}^+$ ions with a kinetic energy of 3 keV ions to 0.3 keV, which is required for ion trapping in the RFQcb of the PUMA ion source beamline. The ions have, as above, a bunch length of 300 ns and an emittance of 8.3π mm mrad. The length of the drift tube l_d and the radius of the PDT r_d have an impact on the energy modification efficiency and longitudinal acceptance as shown in 7. The larger r_d , the smaller is the switch-time plateau, on which the desired energy E with energy spread ΔE is reached. This is caused by the fringe field leaking further into the drift region for larger r_d . Thus, a radius of 20 mm was chosen considering longitudinal acceptance and plateau length. Consequently, $l_d = 150$ mm was chosen which is sufficient for accepting bunches with a bunch length < 1.4 µs in the case the deceleration from 3 keV ions to 0.3 keV.

5.5 Deceleration of ions

In figure 9 a ToF spectrum for an increasing switch time t_s is displayed for the simulation (left) and the measurement (right). The results are normalized to total counts per scan step. The measurement was performed with PDT 1 in section I of the ion source beamline, figure 1. Ions with the same properties as in the simulation are produced in the ion source and their ToF is measured after a deceleration from 3 keV to 0.3 keV. The horizontal lines in red (eq. (5.2)) and blue (eq. (5.5)) show the results of the calculation of the ideal switch time for the dimensions of the beamline. **Simulation** In the simulation an energy of 322.4 ± 5.2 eV was reached with a switch plateau of approximately 1.4 ± 0.1 µs. Applying 2.7 kV to the PDT, one would expect to reach a final kinetic energy of 300 eV, but the deviation can be explained by the behaviour of the PDT switch. The switch pattern of the PDT switch (Behlke HTS 81-06-GSM) was recorded and implemented in the simulations (see figure 10). The initial measured voltage applied to the PDT was 2705.72 V. Switching induces significant noise during and shortly after the switch down, afterwards a damped oscillation converges to 28.42 V, leading to a final energy of 322.7 eV.

Measurement In the measurement, the ⁴⁰Ar⁺ ToF spectra show a similar behavior as in the simulation. The switch plateau length is determined to be $1.6 \pm 0.1 \,\mu\text{s}$. The measured ToF spectrum is shifted on the ToF axis, because the ion production and thus the trigger for the data recording takes place at an unknown position further inside the ion source while in the simulations the ejection point of the ion source was considered as the starting position. An energy analyzer was used in between the PDT and the ToF detector to estimate the resulting energy. Three grids with a transmission of 92 – 95 % are axially aligned, and a potential is applied to the central grid to form a blocking potential. The measured ion count rate as a function of the blocking potential can be fitted with the cumulative distribution function (CDF) of a skew normal distribution, which allows to determine the mean energy μ and the standard deviation σ . For further visualization, the difference in ion counts between two points is plotted together with the corresponding skew normal PDF using the parameters from the CDF, as seen in figure 8. The skew normal distribution function and $\Phi(\alpha x)$ as the standard normal probability density distribution function and $\Phi(\alpha x)$ as the standard normal cumulative distribution function with skewness parameter α .

In this way, the mean energy was determined to be 284.0 ± 0.4 eV. This is about 38 eV below the expected value, explaining the deviations in ToF difference of simulation and measurement. Recording the switch pattern again after the measurement, it was observed that by small modifications of the switch setup, e.g., the electrical connection, the maximum applied current or the type of grounding, the voltage when switching to 0 V varies by several tens of volts. This is effect is less pronounced when applying a non-zero voltage to the switch channel, and could probably be prevented by grounding the channel instead of connecting it to a power supply.

Furthermore, several deceleration schemes were tested with the setup to modify the ion energy from 3 keV to 0.3 keV or 1 keV. The results are summarized in table 1. Acceleration schemes were successfully tested for a re-acceleration after the RFQcb to 4 keV but are not presented in the table due to the voltage limitation on the energy analyzer.

5.6 Longitudinal acceptance

In figure 11 (left), the mean ion energy is plotted as a function of t_s for uniform initial bunch lengths of 0.4, 0.8 and 1.2 µs. For increasing bunch lengths, larger parts of the bunch are in the fringe field or partly outside the drift tube when switching at non-ideal switch times, and thus the switch plateau for minimal final energy and energy spread decreases. The initial energy spread is 10 eV, which is improved for some t_s for all tested bunch widths.

On the right side of figure 11, the resulting bunch lengths are plotted with respect to t_s . Measurement (markers) and simulation (dashed lines) can be compared. For the measurements, the bunch length could be changed using a pulsed deflector electrode in front of the ion source to chop the continuous



Figure 8. The beam energy distribution was measured using an energy analyser grid. By fitting the cumulative distribution function (CDF) the mean energy and its standard deviation could be determined using a skew normal distribution fit. In the background the difference between two measurement points is plotted together with the skew-now PDT to further visualize the shape of the distribution.



Figure 9. Plot of the ion ToF spectrum as a function of the switch time t_s of the PDT for a deceleration from 3 keV to 0.3 keV, comparing the simulation (left) to the measurement (right). The intensity is color coded and normalized to the total count rate per slice.



Figure 10. A measurement of the output voltage of the PDT switch over the switch time. When switching from 2.7 kV to ground, a damped oscillation is visible over a time of about 0.5 µs until the applied potential converges at 28.42 V. This increases the energy that is reached after deceleration.

Switch pattern	E [eV]	$\sigma_E [eV]$	Skewness α	TOF FWHM [ns]
2700 V to 0 V	284.0 ± 0.4	4.1 ± 0.2	-0.83 ± 0.12	294 ± 51
0 V to $-2700 V$	322.9 ± 0.8	7.4 ± 0.5	-0.9 ± 0.1	277 ± 12
2500V to $-200V$	327.6 ± 2.1	9.3 ± 1.3	0.3 ± 0.2	302 ± 34
200 V to $-2500 V$	317.8 ± 0.6	6.3 ± 0.4	-0.7 ± 0.1	264 ± 16
2000 V to 0 V	988.4 ± 0.9	8.8 ± 0.5	-1 ± 0.1	279 ± 32
0 V to $-2000 V$	1019.5 ± 0.6	6.0 ± 0.4	-0.9 ± 0.1	279 ± 18
1000V to $-1000V$	1015.1 ± 0.7	5.9 ± 0.5	-0.7 ± 0.2	278 ± 21

 Table 1. Measured ion bunch parameters after deceleration with the PDT for different switch patterns.

beam, resulting in a uniform bunch shape. Since the shape of the ion bunch distribution after switching can be non-uniform and non-Gaussian, depending on t_s the bunch length was determined independently of the ion distribution within the ion bunch for a better comparison. For this, the width is defined by the difference of the ToF positions at which the ion distribution falls below and rises above 10% of the distributions maximum value. The simulations describe the measurement results well, only the bunch length of the bunches located in the fringe field while switching is slightly overestimated. It can be concluded that by choosing an appropriate t_s , the bunch length degradation can be minimized. This is especially important in cases where long distances have to be covered after an energy modification.



Figure 11. The energy *E* and energy spread ΔE (left) and bunch length Δt (right) depending on the switch time t_s of the PDT for different bunch lengths 0.4, 0.8 and 1.2 μ s for the deceleration of ⁴⁰Ar⁺ ions from 3 keV to 0.3 keV.

6 Iris shutter assembly

6.1 Description

The dimensions of the iris shutter assembly, which is made of a C-276 alloy (SAHM, 95032), can be found in appendix A.4. Figure 12 shows a photograph of the assembly. The iris shutter is mounted on a customized double-sided CF160 flange with two radial ports, one of which holds a micrometer precision linear feedthrough (VAb, NC16-100) that allows to regulate the opening of the iris aperture. The micrometer feedthrough is connected to the iris via a mechanical lever. With this, the iris aperture can be opened (30 mm diameter) or closed with 48 rough adjustment steps, which follow an almost linear dependency. Therefore, the iris diameter can be varied in steps of 625 μ m or better. In the case of the PUMA ion source beamline the shutters are meant to decouple the vacuum regions with different pressure of a beamline by reducing the conductance between the sections while still allowing a focused beam to pass through.

The iris apertures can additionally be used to optimize the overlap of ion beam velocity axis with the central axis, used for beamline tuning. The same technique could also be considered for laser beam overlap. By gradually closing it, the beams centrality and its size of focus can be checked, as described, e.g., in reference [18]. Furthermore, if the iris shutter is insulated from the flange by a thin teflon layer, one can electrically connect it to the second radial port and read the current of the impinging beam.

6.2 Pressure between vacuum sections

In the PUMA ion source beamline, four iris apertures are installed, see Figure 1. Argon buffer gas is injected in the RFQcb section (section III), leading to a vacuum from $1 \cdot 10^{-5}$ mbar to $5 \cdot 10^{-5}$ mbar. Especially the antiproton beam line has to be shielded from a vacuum contamination to keep the



Figure 12. A photograph of the iris shutter assembly. The iris shutter can be positioned between two beamline sections to lower the pass-through area for residual gas particles. It can be opened and closed from outside the vacuum using a the linear feedthrough.

vacuum < 10^{-9} mbar at the quadrupole bender junction. In the following, the impact of the apertures on the vacuum in sections I and II (see figure 1) is demonstrated.

In figure 13, three argon buffer gas flow rates were considered: $5 \cdot 10^{-4}$ mbar 1 s⁻¹, $1 \cdot 10^{-3}$ mbar 1 s⁻¹, $2 \cdot 10^{-3}$ mbar 1 s⁻¹. Each section is differentially pumped with turbomolecular pumps. The pressure was measured for different iris aperture diameters d_{I} , d_{II} and d_{III} with full range and cold cathode gauges at the side flanges. In the left figure, the diameter of Iris I is varied while Iris II is kept open. Reducing the opening of Iris I results in a decrease of the pressure by one order of magnitude. When also closing Iris II, a similar drop in pressure can be observed in section II (right). Meanwhile, the pressure in section I decreased by another factor two, when keeping Iris I at a diameter of 3.45 mm. Gas-flow simulations with the software MolFlow+ [19] were carried out to test the effect of using a stack of multiple iris shutters in a row. However, the non-evacuated region between the shutters acts as buffer zone for higher pressure and overall deteriorates the vacuum level instead of improving it. If the vacuum has to be further improved, an implementation of ion getter pumps and the use of baking is required.

6.3 Outgassing rates

The iris shutters were baked up to temperatures of 250 °C in a test campaign to determine their outgassing rates. 48 hours after the end of a 72 hour bake-out campaign, the iris shutter had a hydrogen degassing rate of less than $1.1 \cdot 10^{-9}$ mbarls⁻¹s. Furthermore, elements with atomic mass units of 53, 64, 66 and 68 are present after the bakeout, but with a factor 1000 less than hydrogen. [20].



Figure 13. The measured pressure in section I, II and III depending on the opening diameter d_i of the iris-type apertures. On the left site, the Iris I is varied while Iris II is kept open. The impact of Iris II is shown on the right side, while Iris I is kept on 3.45 mm. Three buffer gas flow rates were considered: $5 \cdot 10^{-4} \text{ mbar } 1 \text{ s}^{-1}$ (red), $1 \cdot 10^{-3} \text{ mbar } 1 \text{ s}^{-1}$ (blue), $2 \cdot 10^{-3} \text{ mbar } 1 \text{ s}^{-1}$ (green).

7 Conclusion

For the PUMA ion source beamline, operating with voltages up to 5 kV, three ion-optical components have been designed for an adequate transport of the beam towards the PUMA antiproton beamline, as well as for improving vacuum levels between the buffer gas insertion point for the RFQcb and vacuum-sensitive parts of the beamline. The comparision to simple models and simulations have been shown to be predictive and could be reliably used for a setup design.

The einzel lens and steerer assembly successfully focuses and steers the beam for the injection into the RFQcb, which has the smallest acceptance in the beamline. Furthermore, the PDT was used to decelerate ions to energies as low as 0.3 keV, while allowing to minimize the energy spread and bunch length by choosing appropriate switch times. Re-acceleration from 0.3 keV to 4 keV after the RFQcb was tested accordingly. The iris shutters significantly limit the impact of the RFQcb buffer-gas injection on the other vacuum sections by one order of magnitude per section shielded with the iris shutter assembly

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Data Availability Statement

Data are available on request from the authors.

A Drawings

A.1 Einzel lens and steerer assembly



A.2 Mounting flange



A.3 Pulsed drift tube





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