# Photo induced collisions with laser cooled He\* atoms

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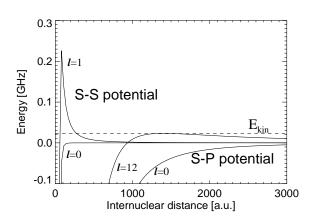
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**Abstract.** This paper presents an experimental investigation of cold collisions between metastable Helium atoms in an optical trap at 1mK. Penning (PI) and associative (AI) ionization reactions are distinguished using a mass spectrometer and studied under influence of near resonant laser light. Sensitive behavior of the ion rate is observed when the laser is tuned close to resonance. Experimental findings are well described, on an absolute scale, by a semi-classical model we have developed for optical collisions and by a modified Julienne-Vigué model.

**PACS.** 32.80.Pj Optical cooling of atoms; trapping – 33.80.Eh Autoionization, photoionization, and photodetachment – 34.50.Rk Laser-modified scattering and reactions

#### 1 Introduction

The ability to trap and manipulate neutral atoms by means of laser light have created new fascinating environments for studies of atomic interactions at low temperatures [1, 2]. With the invention of the magneto-optical trap (MOT), dense and cold samples of atoms are now produced routinely and studied under controlled conditions. Collision dynamics of cold atoms have recently attracted considerable attention experimentally as well as theoretically [3–9]. When a near resonant light field is introduced into the cold atom cloud the rate coefficient for collisional processes can be dramatically modified [9]. As two slowly moving atoms approach they form a quasi molecule which can be excited during the collision. If the molecular complex is excited to a repulsive energy state the atoms will be pushed apart thus preventing them from reaching small internuclear distances where reactions may take place. This effect is called optical shielding of collisions and has been observed in several systems [6, 10]. In the case of a light field detuned below the atomic resonance, excitation to an attractive potential curve will take place. Note, that the interaction potential will depend critically on the electronic state of the atoms involved. When the two atoms are in the ground state we have a van der Waals interaction which scales as  $1/R^6$ . With one atom in an excited P-state, the potential energy arises from an induced dipole-dipole interaction scaling as  $1/R^3$ . Since the excited state potential reduces the centrifugal barrier more partial waves will contribute in the collision process, exemplified for the He\*-He\* system in Fig. 1. Typically only zero or one partial wave come in play for colliding atoms in the ground state at mK temperatures while usually 5–10 partial waves will contribute for the excited state potential.



**Fig. 1.** Lowering of the centrifugal barrier in the excited state for the He\*-He\* system. The ground state S-S potential scales as  $1/R^6$  and has a centrifugal barrier, which is already for  $\ell=1$  too high to overcome. The excited state S-P potential scales as  $1/R^3$ , which for a temperature of 1 mK allows 12 partial waves to reach short internuclear distances.

Interaction with the light field, leading to excitation, spontaneous emission and stimulated emission of the quasi molecule, is an intrinsic part of the cold collision dynamics and offers additional room for influencing the outcome of a collision event. Here the lifetime of the atom in question is important as well as the power and polarization of the light used to probe the collision. This situation is in sharp contrast to collisions with laser prepared atoms at thermal or higher temperatures. Here light is used to excite an independent, isolated atom which then collides with another particle.

The present experiment addresses optical collisions between cold metastable triplet atoms in an optical trap at

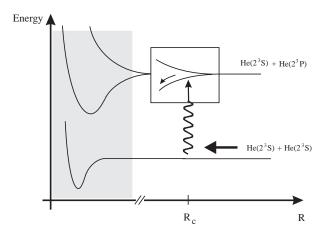


Fig. 2. The quasi molecule formed by two metastable atoms in the S-S ground state is excited to the S-P state at the Condon point  $R_c$ , where the laser frequency equals the dipole-dipole interactions energy. If the laser detuning is negative, as shown in figure, the molecular S-P complex is accelerated towards small internuclear distances where ionization reactions may take place, indicated by the shaded box.

1 mK. The atoms may be in the  $\text{He}(2^3\text{S})$  ground state or the  $\text{He}(2^3\text{P})$  excited state and will spontaneously Penning ionize if their internuclear distance becomes sufficiently small

$$\text{He}(2^3\text{S}) + \text{He}(2^3\text{S}, 2^3\text{P}) \rightarrow \begin{cases} \text{He}^+ + \text{He}(1^1\text{S}) + e^- \\ \text{He}_2^+ + e^- \end{cases}$$
 (1)

Depending on the energy transferred to the electron the ion products may end up as an atomic ion  $(\mathrm{He^+})$  or a molecular ion  $(\mathrm{He_2^+})$ . Monitoring this ion yield as a function of the light field frequency provides detailed information on the long-range interaction between two slowly colliding metastable atoms. This is illustrated in Fig. 2.

A red detuned photon transfers the He\*-He\* complex to the attractive part of the excited state potential at the Condon point  $R_c$ . Here it is accelerated towards small internuclear distances where ionization may occur. Inevitably spontaneous decay of the S-P complex will reduce the chance of reaching small distances on the excited state potential. Depending on the absolute value of the detuning, i.e., the location of  $R_c$ , the chance of decaying varies. This can be controlled by adjusting the frequency of the photons.

The He\* collision system possesses several attractive features compared to other systems investigated so far. Having no nuclear spin the He atoms do not suffer from the presence of hyperfine structure. This facilitates considerably interpretation of data and offers the possibility of accurate and detailed modeling on an absolute scale.

Yet another property of the He\* system is interesting. Two distances characterize an optical collisions:  $R_{\tau} = v\tau$ , the average distance traveled by an atom during one lifetime, and  $R_{\lambda} = \lambda/2\pi$ , the characteristic length where retardation effects become important. Although  $R_{\lambda}$  is in the same order of magnitude for all systems studied so far,

**Table 1.** Critical distances  $R_{\tau}$  of molecular excitation that have been accessed in optical collision experiments. Reactions are studied for molecular excitations in the He\*+He\* system far beyond the point of  $R_{\lambda}$ .

| system                              | $\lambda$ [nm] | $\tau$ [ns] | $\langle v \rangle$ [m/s] | $R_{	au}$ [Å] | $R_{\lambda}$ [Å] | Ref.    |
|-------------------------------------|----------------|-------------|---------------------------|---------------|-------------------|---------|
| He*+He* Na+Na Kr*+Kr* Rb+Rb Xe*+Xe* | 1083           | 98          | 2.0                       | 2744          | 1724              | [14]    |
|                                     | 589            | 16          | 0.3                       | 65            | 940               | [3,7,8] |
|                                     | 811            | 31          | 0.2                       | 87            | 1290              | [5]     |
|                                     | 780            | 27          | 0.1                       | 45            | 1240              | [4]     |
|                                     | 882            | 33          | 0.1                       | 42            | 1404              | [6]     |

 $R_{\tau}$  is two orders of magnitude larger for the He\*+He\* system compared to the other systems, as seen from Table 1. Therefore we expect the He\* system to be different from other systems and, in addition, retardation effects to become important.

### 2 Experimental setup

The experimental setup consists of three main sections separated by differential pumping stages; a metastable helium source, a Zeeman slower, and a magneto-optical trap, shown schematically in Fig. 3. Metastable atoms are produced in a high pressure DC discharge expanding through a small 0.5 mm nozzle hole. To reduce the initial atom velocity the source unit is cooled with liquid nitrogen and operated at a low discharge power (700 V, 3 mA). Under these conditions we have a  $He(2^3S_1)$  output yield of  $10^{14}$  ${\rm sr}^{-1}{\rm s}^{-1}$  with a mean beam velocity of 900 m/s. Before the metastable atoms can be loaded into the MOT they must be slowed down sufficiently. In the Zeeman slower the atoms are decelerated by a counter propagating laser beam tuned close to the  $He(2^3S_1)$ -  $He(2^3P_2)$  transition. A spatial varying magnetic field along the beam axis provides a Zeeman detuning which keeps the atoms in resonance through the slower unit. By optimizing the current in the Zeeman coils, and the laser detuning, final velocities in the range of 50 – 100 m/s can be obtained. Even lower velocities are attainable but at the expense of a significant transverse expansion of the beam. Ultimately the beam will fan out and this results in a poor beam intensity at the trap position. In the MOT section the atoms are further decelerated and finally trapped. The MOT is composed of three pairs of counter propagating laser beams crossing at right angles in the center of a quadrupole magnetic field. The quadrupole field is generated by two circular coils mounted in an anti-Helmholtz configuration. Typical field gradients are 7.5 G/cm in the plane of the coils and 15 G/cm along the center axis perpendicular to the coil plane.

A detailed picture of the optical setup is given in Fig 4. Both slowing and trapping lasers are operated on the  $\mathrm{He}(2^3\mathrm{S}_1)$ - $\mathrm{He}(2^3\mathrm{P}_2)$  transition of metastable helium. The 1083 nm laser light required for this transition is provided by two SDL single mode diode lasers, each capable of giving an

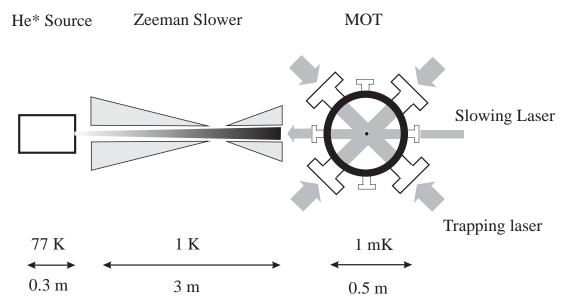


Fig. 3. Schematic view of the experimental setup from above. A beam of metastable atoms is produced in a DC discharge. After being decelerated in a Zeeman slower by a counter propagating laser beam the atoms are loaded into a magneto-optical trap. Here the atoms are further cooled and studied.

output power of 50 mW. For trapping, we use a single laser beam. Two telescopes optimize circular beam shape and diameter while a Glan-Thomson polarizer, followed by a  $\lambda/4$ -plate, ensures the right circular polarization state of the light before it enters the trap region. First the trap beam traverses the symmetry plane of the MOT coils, then mirrored out of the plane and directed along the center axis of the coils, and finally reflected back by a mirror placed under the MOT chamber (not shown). A  $\lambda/2$ -plate changes the helicity of the laser light just before it is mirrored out of the symmetry plane while a  $\lambda/4$ -plate on top of the back reflecting mirror ensures correct polarization state for the back going light beam.

To acquire an optimal loading rate of the MOT it is operated at a large negative detuning of  $15\Gamma$  (natural line width  $\Gamma=1.6$  MHz), a high trap laser power of  $50~I_0$  (saturation intensity  $I_0=0.16$  mW/cm²), and with a large trap beam diameter of about 35 mm (full width at  $1/e^2$ -height). The latter increases the capture velocity and capture area of the MOT. With a background pressure of  $3\times 10^{-9}$  mbar we typically trap about  $10^5$  atoms in a volume of 4 mm³ yielding an average density of about  $10^7$  atoms/cm³. The lifetime is typically 3–4 s and the decay of the MOT, after the load beam is shut off, is a pure exponential.

For diagnostics of the trapped atom cloud we have installed a mass spectrometer and a micro channel plate, both mounted in the symmetry plane of the two circular coils but opposite to each other, as shown in Fig. 4. A plate mounted in front of the mass spectrometer is biased by a small negative voltage (typical -150 V) in order to extract ions originating from the MOT volume. The ions may be formed in collisions between slow helium atoms in the MOT [see Eq. (1)] or in Penning ionization with background gas atoms. In particular, mass selective detection

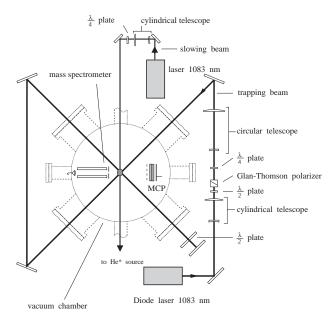
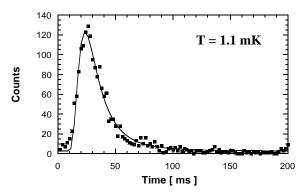


Fig. 4. Detailed picture of the optical setup used for trapping and slowing. Two diode laser systems are used, one for initial slowing of the atoms and the second for trapping of the atoms. A mass spectrometer and a microchannel plate is mounted close to the MOT center for diagnostics of the cold atoms.

allows a discrimination between Penning and associative ionization reactions, taking place between the cold atoms, as well as ionization with "hot" background particles. A grid mounted in front of the microchannel plate (MCP) can be biased negative (typical  $-400~\rm{V})$  when the total ion yield is measured, or positive (typical  $+100~\rm{V})$  when only metastable atoms are detected. When the MOT is operat-



**Fig. 5.** Time of flight distribution of cold metastable atoms after being released from the MOT. The temperature is found to be about 1 mK which corresponds to a mean velocity of 2 m/s.

ing, we observe a total ion yield of about 25 kc/s mainly due to Penning ionization with the background gas.

The temperature of the atom cloud was measured by a time of flight technique. Biasing the MCP grid by a positive voltage and releasing the atom cloud to expand freely in the gravitational field we could measure the flight time to the MCP located 70 mm from the MOT center. Fig. 5 shows the result of such a measurement. The time of flight distribution is peaked at about 30 ms corresponding to an average velocity of about 2 m/s. By fitting a Maxwell-Boltzmann distribution to the measured data we obtain a temperature of 1.1 mK in good agreement with prediction of standard Doppler theory using our MOT parameters [11]. In addition, we obtain an estimate for the total number of atoms in the MOT. Knowing the exact area of the MCP we can correct for the solid angle and obtain about  $10^5$  atoms, consistent with the total ion rate measurement. The MCP detection efficiency for a metastable helium atom is assumed to be 60%.

#### 3 Results and discussion

This section presents results on optical collision experiments with cold metastable triplet atoms at 1 mK. Collisions between the slow atoms in the trap and background atoms produces an ion rate

$$R = \alpha N + \frac{\beta}{2} \int n(\mathbf{r})^2 d\mathbf{r}$$
 (2)

where  $N = \int n(\mathbf{r})d\mathbf{r}$  is the total number of atoms in the trap,  $\alpha$  is characterizing the production of ions in collisions with background gas atoms and  $\beta$  the PI and AI rate as defined in Eq. (1). The  $\alpha$  coefficient is related to the cross section by

$$\alpha = \sum_{i} \sigma_{i} n_{bg}^{i} \bar{v}_{i} \tag{3}$$

with  $\sigma_i$  being the ionization cross section of metastable helium colliding with the background atoms of type i,  $n_{ba}^i$ 

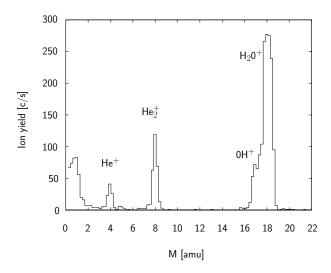


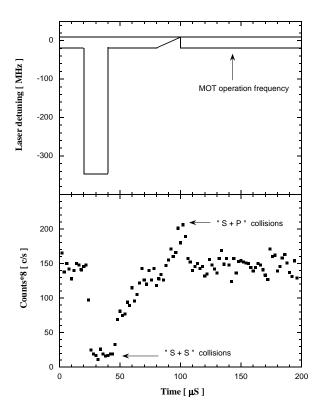
Fig. 6. Ion mass spectra from the MOT. The  $\mathrm{He}^+$  and  $\mathrm{He}_2^+$  peaks are produced in collisions between cold metastable atoms in the MOT. The  $\mathrm{H}_2\mathrm{O}^+$  is produced in Penning ionization with water molecules on trapped  $\mathrm{He}^*$  atoms.

the density and  $\bar{v}_i$  the average velocity of the background atoms. For the  $\beta$  coefficient we have

$$\beta = \sigma_* \bar{v}_r \tag{4}$$

where the  $\sigma_*$  is the cross section for PI and AI ionization and  $\bar{v}_r$  the average relative velocity between metastable atoms in the MOT. The density profile of the MOT is given by  $n(\mathbf{r})$  and is typically approximated by a Gaussian distribution function. In our case we estimated the mean radius of the MOT from our temperature measurement. Since the atoms are moving in a harmonic potential we have  $\frac{1}{2}k_BT = \frac{1}{2}\kappa\bar{x}$ , where T is the temperature in the MOT,  $\kappa$  the spring constant of the MOT and  $\bar{x}$  the average position. We measure the temperature and estimate the spring constant from our MOT model, which successfully predicted our measured temperature from the known MOT parameters [11]. This approach yields an average MOT density.

Figure 6 shows results of a mass spectrum taken when the trap was in operation. Two remarkable features are observed. Firstly, two clear peaks appear at mass 4 amu and 8 amu which we identify with the formation of He<sup>+</sup> and He<sub>2</sub><sup>+</sup> ions produced in collisions between trapped helium atoms. They cannot be produced in collisions with the background gas or the loading beam, since the density is several orders lower compared to the MOT density. The two peaks have different height indicating different reaction rates of the PI and AI process defined in Eq. (1). Unfortunately the ions are not detected with the same efficiencies. The He<sub>2</sub><sup>+</sup> ion is produced with a very low velocity and is detected with a high efficiency. The He<sup>+</sup> ion is produced with a significant higher velocity due to the dissociation of the molecular ion. This reduces the extraction efficiency of the He<sup>+</sup> ions. In order to correct for this we simulated the extraction of the ions by calculating the



**Fig. 7.** Optical modulation of the  $\text{He}_2^+$  ion yield. The laser is first detuned far below resonance, then ramped to resonance and finally returned to the normal trapping frequency. Notice the dramatic effect when the laser is shifted off resonance and only atoms in the ground state is present. Close to the  $\text{He}(2^3\text{S}_1)$ -  $\text{He}(2^3\text{P}_2)$  resonance a fraction of the cold atoms are in the excited state enhancing the ion production significantly.

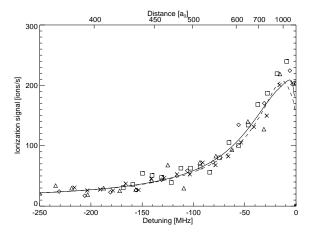
trajectories for different initial velocities, obtained using the potential in [12]. This gave correction factors for the measured ion yield. Secondly, the large peak at 18 amu is associated with Penning ionization with water on cold He\* atoms. This peak is due to a cross section of this process of more than 70 Ų [13]. Using the mass spectrometer we are able to distinguish collisions taking place in the MOT, between slow metastable atoms, and collisions with the background atoms. In the following we will concentrate on intra MOT collisions, i.e., the peaks with mass 4 amu and 8 amu.

In the mass spectrum shown above we have a mixture of atoms with about 80% in the ground state  $(2^3P_1)$  and 20% in the excited state  $(2^3P_2)$ . It is desirable to separate the signal contributions arising from collisions between atoms in the ground state and the excited state. Figure 7 presents results from an experiment in which we detected  $He_2^+$  ions. The MOT laser is detuned far below resonance in a short time interval and then returned to normal trapping operation. The time interval is chosen to be of the order of 20  $\mu$ s, which is long compared to the natural lifetime of 100 ns but short compared to the expansion time of the atom cloud. The scan cycle is repeatedly integrated until satisfactory statistics is obtained. We observe

a dramatic drop in the He<sub>2</sub><sup>+</sup> ion production as the laser is shifted off resonance. If the laser is detuned far below the atomic transition frequency only ground state  $He(2^3S)$ atoms are present. The ion production thus takes place via interaction on the potentials  $1^1\Sigma_g^+$ ,  $1^3\Sigma_u^+$  and  $1^5\Sigma_g^+$ , connecting asymptotically to He(2<sup>3</sup>S)-He(2<sup>3</sup>S) manifold. The quintet state, however, will not contribute due to violation of the total spin conservation in the ionization process. Detailed investigations, based on the theoretical model presented in the next section, show that even for detunings of 350 MHz, as used in the present experiment, the ion rate cannot be considered originating from pure S-S collisions. About half of the ion signal at 350 MHz appear to be due to S-P interactions. When the laser is shifted to resonance the ion yield increases strongly. Now the MOT contains, in addition to atoms in ground state a fraction of atoms in the excited 2<sup>3</sup>P<sub>2</sub> state. Considering the high powers used for the trapping laser we assume the number of excited atoms to be 50% on resonance. The significant enhancement of the ion yield at resonance we ascribe to an increased ion rate when atoms react via the excited state  $He(2^3S)$ - $He(2^3P_2)$  potential. The reason is twofold. Firstly, the attractive nature of the S-P dipole-dipole interaction "captures" more partial waves compared to the S-S potential (see Fig. 1). The number of contributing partial waves for the S-P potential is about 10 while only the l=0 wave contributes for the S-S potential at 1 mK. Secondly, the lifetime of the S-P complex is very long so once excited, even at large internuclear distances, it will survive until it reaches small distances where ionization occurs. This is in contrast to all other systems investigated. The characteristic distance,  $R_{\tau}$ , traveled for these systems is, as seen from Table 1, more than one order of magnitude less compared to the He\* system. As the laser frequency is tuned close to resonance the ion rate drops rapidly for these systems since the excited quasi molecules do not live long enough to reach small internuclear distances.

We have measured the ratio of  $\mathrm{He_2^+/He^+}$  on and off resonance. Corrected for the transmission efficiency we find, on resonance  $0.16\pm0.02$  and off resonance  $0.03\pm0.003$ , where the error bars refer to statistical uncertainties. The change in this ratio is a clear evidence, that the reaction on- and off-resonance takes place via different reaction channels.

To explore further the dynamics of the collision at long internuclear distances we measured the  $\mathrm{He}_2^+$  ion yield as a function of the trap laser frequency. In a time interval of about 50  $\mu\mathrm{s}$  the trap laser is linearly ramped from resonance to -300 MHz, while the  $\mathrm{He}_2^+$  ion signal is monitored. Figure 8 shows results of several such measurements. Near resonance the ion signal increases considerably and then drops gradually to the S-S rate as the laser detuning increases. At larger detunings, ( $\leq -50$  MHz), the excitation of isolated  $\mathrm{He}^*$  atoms is completely negligible but we still observe a relatively high ion rate. This is attributed to resonant excitation of transient molecules, i.e., when two atoms approach to an internuclear distance close to the Condon point at which the laser frequency equals the dipole-dipole interaction energy.



**Fig. 8.** The frequency dependence of the  $\mathrm{He}_2^+$  ion yield. The solid line represents the our semi-classical model calculation and the dashed lined is predictions of a modified JV-model. Both are in very good agreement with the absolute rate measured.

## 4 Theoretical modeling

To describe the data as shown in Fig. 8 we have developed a model based on semi-classical considerations. In this model we describe the excitation in the region of the Condon point  $R_C = (C_3/\hbar\delta)^{1/3}$  within the dressed atom picture, in which a crossing between the potentials for ground and excited state of the quasi molecule arises. The flux of atoms crossing the Condon radius is given by the classical distribution function [16]  $F = 4\pi R_c^2 n^2 v_{\rm rad}$ , with  $v_{\rm rad}$  the radial velocity. The excitation probability at the Condon radius is calculated using the Landau-Zener expression [17]

$$P_{ex} = 1 - \exp(-\pi\Lambda),\tag{5}$$

with the Landau-Zener parameter  $\Lambda$  given by

$$\Lambda = \frac{2\hbar\Omega^2}{\alpha v} \tag{6}$$

Here  $\Omega$  is the Rabi-frequency and v the velocity of the atom. Furthermore, the parameter  $\alpha$  is the slope of the interaction potential at the Condon point, which in this case is given by

$$\alpha = \left| \frac{dV}{dR} \right|_{R} = \frac{3C_3}{R_c^4}.$$
 (7)

After excitation the two atoms are attracted towards each other due to the induced dipole-dipole interaction and they have a probability of Penning ionization at small internuclear distances. However, there are two constraints. Firstly, the quasi molecule has to reach small internuclear distances, where the ionization takes place. Secondly, the flight time may not be too large, since the lifetime of the quasi molecule in the excited state is in the same order as the atomic lifetime. Therefore the molecule can deexcite before reaching small internuclear distances by spontaneous emission.

In order to evaluate these constraints we have calculated classical trajectories for the quasi molecule. The trajectory of an atom, crossing the Condon radius with an initial velocity  $\mathbf{v}$ , is calculated and its "fate" is indicated with the parameter w. If the atom reaches small internuclear distances, we have w=1, whereas we have w=0, if not. From the calculation we extract the time interval  $\tau_0$  between the excitation at  $R_c$  and the arrival at small internuclear distance. The survival  $S(\tau_0)$  of the atom in the excited state is described by an exponential decay  $S(\tau_0) = \exp(-\Gamma \tau_0)$ . If we combine all four steps (flux inwards, excitation, reaching close distances, in the excited state) we arrive at our final expression

$$R_{ion} = V \int_{-\pi/2}^{\pi/2} F(R_c) P_{ex}(R_c) w(R_c) S(\tau_0) \sin \theta \ d\theta/2,$$
(8)

where the integration over the angle  $\theta$  runs over all angles of the initial velocity  $\mathbf{v}$  with respect to the internuclear axis and V denotes the MOT volume.

Our model becomes identical to the model of Gallagher and Pritchard [16] in the case of low excitation rate and large detuning. However, we extend on it by including a Landau-Zener excitation probability instead of a quasistatic approximation, which does not produce correct results for high excitation rates. Furthermore, we explicitly calculate the trajectory of the atom, in order to determine, if under the initial conditions the atom can reach small internuclear distances. Our model becomes identical to the model proposed by Julienne and Vigué [18], when we replace their quasi-static excitation rate with the Landau-Zener expression. However, the model of Julienne and Vigué is based on a quantum-mechanical description of the dynamics, whereas we rely on a classical description.

The results of our calculations is shown in Fig. 8 together with the measured data. The agreement between theory and experiment is satisfactory. In the calculation we have taken into account, that for the He\*-He\* system we have, due to the fine structure interaction, 10 attractive curves from which 9 can contribute to the ionization. For each of these curves we have carried out the calculation and finally performed the statistical weighted sum. Furthermore, we have averaged the calculation over the distribution of incoming velocities, which is determined by the temperature (1 mK) of the trap. There are some deviations at small detunings, but these are to be expected, since at small detunings the two dressed states come close together and the quasi molecule starts no longer from the lowest state. Then we expect that describing the excitation rate with the Landau-Zener expression becomes inaccurate. Also indicated in the figure by a dashed line the result of the Julienne-Vigué model, where we have replaced their quasi-static expression with the Landau-Zener expression. The agreement between the two models is excellent, indicating that both a classical and quantummechanical description of the dynamics work for our sys-

Apart from the detuning dependence of the ion rate the theoretical model also predict the ionization rates on an

absolute scale. For the maximum S-P rate, close to resonance, we obtain a rate coefficient of  $4 \times 10^{-9}$  cm<sup>3</sup>/s. Using our estimated values for the number of atoms and the volume of the trap our experimental results agrees within a factor of two with this value, which is reasonable given the uncertainty in the experimental parameters. However, this ionization rate is more than one order of magnitude different compared to the value published by Bardou et al. [14]. One possible reason for this discrepancy is that in their experiment the total loss rate, by binary collisions between metastable atoms in the MOT, is measured, whereas we measure those intra-MOT processes, that produces ions. Processes like radiative escape, which lead to trap loss, but do not produce ions will contribute to the rate measured by Bardou et al., but not to the rate measured in our experiment.

#### 5 Conclusion and outlook

We have studied collisions between slow metastable triplet He atoms in an optical trap. Characteristic for this system is its ability to ionize at short distances to atomic He<sup>+</sup> or molecular He<sub>2</sub><sup>+</sup> ions. Using a mass spectrometer we have been able to distinguish these two fundamental reaction channels. Significant enhancement of the ion rate is observed for both channels, when laser light is tuned close to the  $He(2^3S_1)$ -  $He(2^3P_2)$  resonance. This increase takes place as a result of the relative long distance the helium atoms can travel in the excited state. We modeled our results using an intuitive semi-classical model and obtained good agreement between theory and experiment. We also compared our model to a more elaborate quantum mechanical model by Julienne and Vigué [18] and found reasonable agreement for our system. This indicates that the dynamically aspect of the optical collision is well described by both models.

Future experiments will concentrate on optical shielding effects where the laser is tuned to the blue of the atomic resonance. Here the He\* system takes a prominent place due to its simple level structure.

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