

01 Oct 1992

State-Selective Capture in Collisions of Protons with Noble Gases

Michael Schulz

Missouri University of Science and Technology, schulz@mst.edu

D. M. Blankenship

S. W. Bross

A. D. Gaus

et. al. For a complete list of authors, see https://scholarsmine.mst.edu/phys_facwork/1299

Follow this and additional works at: https://scholarsmine.mst.edu/phys_facwork

 Part of the [Physics Commons](#)

Recommended Citation

M. Schulz et al., "State-Selective Capture in Collisions of Protons with Noble Gases," *Physical Review A*, vol. 46, no. 7, pp. 3870-3876, American Physical Society (APS), Oct 1992.

The definitive version is available at <https://doi.org/10.1103/PhysRevA.46.3870>

This Article - Journal is brought to you for free and open access by Scholars' Mine. It has been accepted for inclusion in Physics Faculty Research & Creative Works by an authorized administrator of Scholars' Mine. This work is protected by U. S. Copyright Law. Unauthorized use including reproduction for redistribution requires the permission of the copyright holder. For more information, please contact scholarsmine@mst.edu.

State-selective capture in collisions of protons with noble gases

M. Schulz, D. M. Blankenship, S. W. Bross, A. D. Gaus, T. J. Gay, W. Htwe, J. T. Park, and J. L. Peacher

Department of Physics, University of Missouri–Rolla, Rolla, Missouri 65401

and Laboratory for Atomic and Molecular Research, University of Missouri–Rolla, Rolla, Missouri 65401

(Received 8 April 1992)

We have measured coincidences between neutral H atoms and Lyman- α photons for collisions between 50-keV protons and noble gases as a function of the projectile scattering angle. The coincidences are dominated by capture to the $2p$ state of the projectile. While the total cross sections depend strongly on the target, the shape of the angular distribution of the differential cross sections was found to depend only weakly on that parameter. The data indicate that electrons are captured predominantly from the outermost shell of the target atom for the collision systems studied here.

PACS number(s): 34.70.+e, 34.50.Fa

I. INTRODUCTION

Electron capture in ion-atom collisions has been studied extensively since the first measurements in the 1920s [1–7]. Investigations have been performed both experimentally and theoretically for a large variety of collision systems and over a large range of collision energies. In spite of these continuous efforts our understanding of capture processes is still rather incomplete. One reason for this limited understanding is the fact that in a capture process, in contrast to an excitation process, for example, the nucleus with which the electron is associated changes, as does the electron's subsequent average linear momentum. The correct description of this change is a difficult task, involving the introduction of electron translational factors (see, e.g., Refs. [8–10]). Therefore, it is important to measure capture cross sections for many collision systems over a large range of projectile energies, scattering angles, and initial and final electron states so that detailed tests of theoretical calculations can be made.

Total capture cross sections have been measured at projectile energies ranging from as low as 0.1 eV/amu [11] for highly charged ions to as high as a few MeV/amu [e.g., [12]]. The agreement between experimental data and theoretical calculations varies depending on the projectile energy region and the model used in the calculation. However, there is no theoretical model available at this time which gives good agreement with measured data over a large range of projectile energies. State-selective capture has been studied especially at small collision energies using energy-gain spectroscopy (see, e.g., Refs. [13,14]). At intermediate projectile energies state-selective capture has been studied by detecting photons emitted after capture to an excited state in coincidence with the projectile which has captured one or more electrons [15,16]. These experiments provided a better understanding of the n - and l -state distribution of the captured electron. A large variety of data exist for differential cross sections for capture to all states (see, e.g., Refs. [17–20]) providing more sensitive tests of theoretical calculations than total cross sections.

Experimental studies of differential cross sections with

state-selective capture are still relatively rare. At low projectile energies differential cross sections were measured for capture to the $2p$ state of hydrogen for protons colliding with He [21] and Ar [22]. Furthermore, the polarization of Lyman- α radiation following capture to the $2p$ state in low-energy collisions of protons with He was measured as a function of scattering angle [23]. However, in this latter work no absolute cross sections were quoted. We have recently reported measured differential cross sections for capture to the $2p$ state of hydrogen for protons colliding with He at intermediate energies [24]. In this work we have extended these studies to investigate the heavier noble-gas targets Ne and Ar.

II. EXPERIMENTAL METHOD

The experimental apparatus is shown schematically in Fig. 1. The experiment was performed with the 200-kV accelerator which is part of the University of

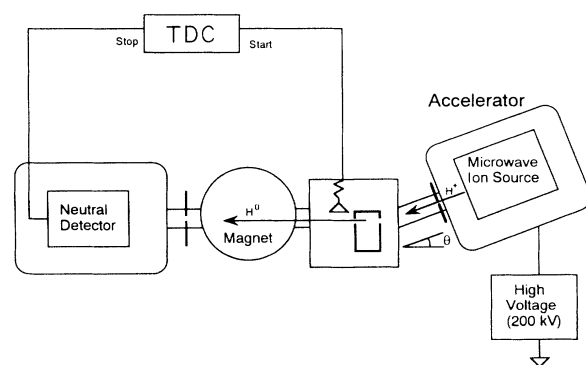


FIG. 1. Experimental setup. The scattering angle of the projectiles was set by rotating the accelerator around the center of the target chamber. The photons emitted by the projectiles after capture to the $2p$ state were detected by a channeltron detector mounted downstream from the target cell in order to avoid detection of photons emitted by the target atoms. The neutralized projectiles were measured in coincidence with the photons.

Missouri–Rolla ion energy-loss spectrometer [25]. Protons were produced with a recently developed 2.45-GHz microwave ion source which is similar to one described earlier [26], and accelerated to an energy of 50 keV before being focused into a gas cell. A movable slit in front of the target chamber was used to collimate the beam to a size of approximately $0.15 \times 0.15 \text{ mm}^2$. The beam intensity going through the target cell was typically 0.5 nA. Protons were separated from the neutralized projectiles magnetically. The H atoms passed through the 0° port of the separation magnet and were detected by a focused mesh electron multiplier. A movable slit in front of this detector was used to define the beam trajectory. The scattering angle was set by rotating the accelerator around the center of the gas cell [27]. The accelerator angle could be set with an accuracy of $3.3 \mu\text{rad}$. However, the angular resolution of particles detected with the neutral detector is determined by the geometry of the trajectory-defining slits. From the measured angular distribution of the incident beam, which was determined with the target gas removed from the scattering chamber and the switching magnet turned off, we estimate that our experiment has an angular resolution of approximately 0.2 mrad (see Fig. 2).

The experiment was performed with He, Ne, and Ar gas targets. The target region was differentially pumped and had an interaction length of about 1 cm. Target pressures were typically 4 mTorr for Ar, 10 mTorr for Ne, and 50 mTorr for He. The pressure dependence of the neutral-atom count rate was linear at these pressures ensuring single-collision conditions. They were monitored by a capacitance manometer and kept constant to within 5% by a microprocessor-based pressure controller [28]. Pressures below 2×10^{-6} Torr in the surrounding vacuum and a few times 10^{-7} Torr in the rest of the beam line could be maintained with these target pressures.

Emitted photons resulting from the collision were detected by a channeltron detector which was mounted downstream from the target cell. With this geometry, photons emitted by the target atoms were not detected. The entrance of the channeltron was covered with a 1-

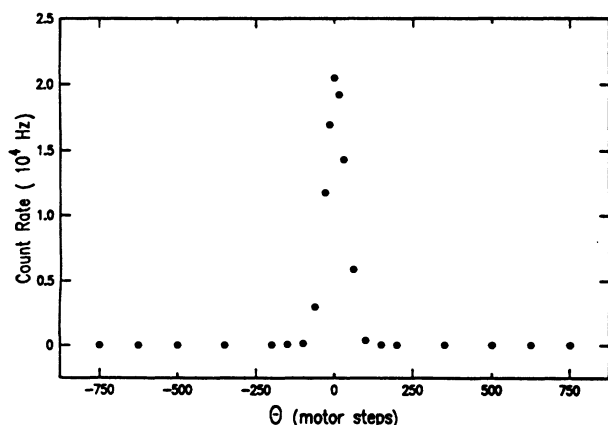


FIG. 2. Angular distribution of the incident proton beam with no target gas. One motor step corresponds to $3.3 \mu\text{rad}$.

mm-thick MgF_2 window. This kept electrons from being detected and also discriminated against photons with wavelengths shorter than 1140 \AA due to absorption in the MgF_2 window. Photons with wavelengths longer than $\approx 1400 \text{ \AA}$ were discriminated against due to the low-energy cutoff of the channeltron response. Therefore, the photon detectors were sensitive to H Lyman- α photons ($\lambda = 1216 \text{ \AA}$), but not to Lyman- β photons ($\lambda = 1026 \text{ \AA}$) or transitions from higher- n states.

The neutral-atom- and photon-detector signals were measured in coincidence. The signal from the photon detector provided the start signal for a time-to-digital converter (TDC) and the neutral-atom detector provided the stop signal. The TDC signals and the total count rates of both detectors were recorded with a computer-aided measurement-and-control based data-acquisition system. The total photon count rate was used to normalize the coincidence count rate (see Sec. III). Typically data were taken for five scattering angles in each run. The collection time could be varied for each angle. For small angles the collection time was typically 20 sec in each scan and a few minutes for large angles.

Before and after each run the angular distribution of the total neutral-atom rate (without the coincidence requirement) and of the incident beam was measured. The coincidence count rates as a function of scattering angle were later corrected for the angular spread of the incident beam (see Sec. III).

III. DATA ANALYSIS

In Fig. 3 we show a coincidence time spectrum for 50-keV protons colliding with Ar at an accelerator angle of 0.1 mrad. A sharp coincidence peak at about 700 nsec with a full width at half maximum of approximately 10 nsec can be observed on top of a flat random background. This width we mainly attribute to the uncertainty of the projectile's time of flight due to accelerator-voltage fluctuations and to the charge-collecting time in the detectors. The true coincidences are dominated by capture to the $2p$ state. Capture to higher-lying states is negligible because of the long transition time for the cascade transition that would have to precede the emission of a Lyman- α photon.

The number of true coincidences were normalized to the total photon rate at each angle. The angular distribution of these normalized coincidence rates is affected by the divergence of the incident beam and were therefore deconvoluted with the measured incident beam profile. This was important because scattering angles smaller than the beam divergence were studied. The deconvolution procedure has been described in detail by Park *et al.* [27].

In order to obtain absolute differential cross sections the coincidence count rates normalized to the total photon count rates were integrated over the scattering angle and then normalized to the total cross sections for capture to the $2p$ state. For He the total $2p$ capture cross sections were taken from a fit to measured data reported previously for a number of projectile energies by different authors [29]. In the case of Ne and Ar the total $2p$ cap-

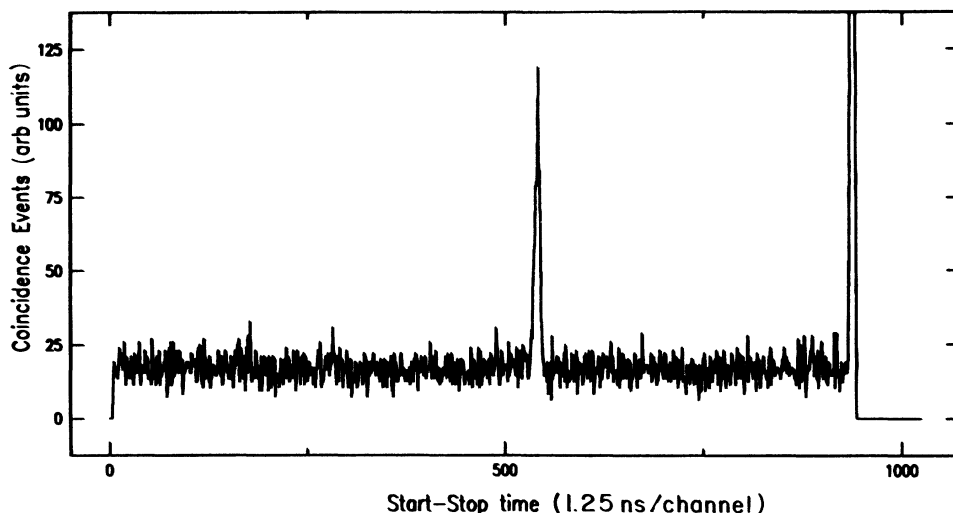


FIG. 3. Coincidence time spectrum taken at an angle of 0.1 mrad for protons colliding with Ar. The acquisition time for this spectrum was approximately 10 min.

ture cross sections were taken from measured data by Hughes *et al.* [15]. The change in photon count rate as a function of the target observed in our experiment is consistent with these previously measured data.

In principle, the determination of a $2p$ capture cross section from the coincidence count rate requires

knowledge of the angular distribution of the emitted photons. It has been shown [21,30] that for electric dipole radiation the coincidence count rate dN/dt as a function of the projectile scattering angle Θ_n is related to the differential cross sections for capture to the $2p_0$ and $2p_{\pm 1}$ substates $(d\sigma/d\Omega)_0$ and $(d\sigma/d\Omega)_1$ by

$$\frac{dN(\Theta_n)}{dt} = A \left\{ 7 \left[\frac{d\sigma}{d\Omega} \right]_0 + 11 \left[\frac{d\sigma}{d\Omega} \right]_1 - 3 \left[\frac{d\sigma}{d\Omega} \right]_1 \sin^2 \Theta_\gamma \cos[2(\Phi_n - \Phi_\gamma)] - 3 \left[\left[\frac{d\sigma}{d\Omega} \right]_0 - \left[\frac{d\sigma}{d\Omega} \right]_1 \right] \cos^2 \Theta_\gamma + 3\sqrt{2} \operatorname{Re} \langle a_0 a_1^* \rangle \sin(2\Theta_\gamma) \cos(\Phi_n - \Phi_\gamma) \right\}. \quad (1)$$

where A is a constant of proportionality. The angles (Φ_n, Θ_n) and $(\Phi_\gamma, \Theta_\gamma)$ are the azimuthal and polar angles of the scattered particle and emitted photon with respect to a coordinate system with its z axis in the direction of the incident proton. The detector geometry in our experiment is shown in Fig. 4. The term $\operatorname{Re} \langle a_0 a_1^* \rangle$ is the density-matrix element which describes the relative phase between the $m_l=0$ and ± 1 scattering amplitudes. For $\Phi_n=0^\circ$ and Θ_γ and $\Phi_\gamma=90^\circ$, the photon detector is mounted perpendicular to the scattering plane and Eq. (1) reduces to

$$\frac{dN(\Theta_n)}{dt} = 7A \left[\left[\frac{d\sigma}{d\Omega} \right]_0 + 2 \left[\frac{d\sigma}{d\Omega} \right]_1 \right]. \quad (2)$$

Since the collision plane contains the z axis the cross sections for populating the $m_l=+1$ and -1 states are equal. The differential $2p$ capture cross section is therefore given by

$$\left[\frac{d\sigma}{d\Omega} \right]_{2p} = \left[\frac{d\sigma}{d\Omega} \right]_0 + 2 \left[\frac{d\sigma}{d\Omega} \right]_1, \quad (3)$$

and under the conditions leading to Eq. (2) the coin-

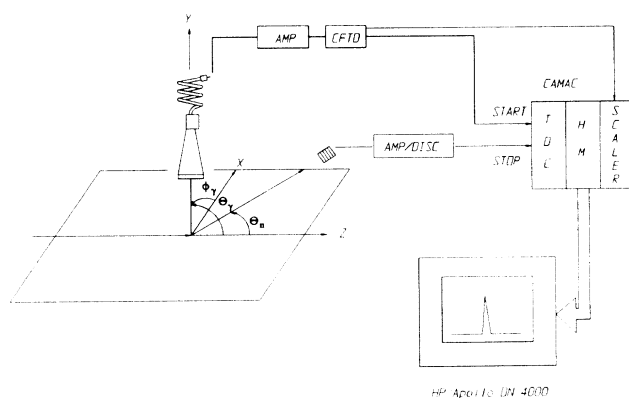


FIG. 4. Detector geometry and electronic coincidence setup. The photon detector was mounted perpendicular to the scattering plane defined by the incident and scattered beam axes. The scattering angle is Θ_n , and Θ_γ and Φ_γ are the polar and azimuthal angles of the photon detector with respect to the scattering plane. The acronyms CFTD, TDC, and HM stand for constant-fraction timing discriminator, time-to-digital converter, and histogramming memory, respectively.

cidence count rate is thus proportional to the differential $2p$ capture cross section. It should be noted that if any of the photons detected are emitted in a direction not perpendicular to the scattering plane the $2p$ capture cross section can only be determined if the relative population of the $m_l=0$ and ± 1 states are known. Furthermore, Eq. (2) does not apply to measurements of total cross sections where the scattered projectile is not detected since in that case the collision plane is no longer defined.

In our experiment the photon detector was mounted perpendicular to the scattering plane. However, due to the acceptance angle of the neutral-atom and the photon detectors, some photons were detected which were not emitted perpendicular to the scattering plane. Equation (1) therefore has to be integrated over the acceptance angle of both detectors and then takes the form

$$\frac{dN(\Theta_n)}{dt} = C \left[\left. \frac{d\sigma}{d\Omega} \right|_0 + 2D \left. \frac{d\sigma}{d\Omega} \right|_1 \right], \quad (4)$$

where C is another constant of proportionality and D , which depends on the scattering angle, is a factor resulting from the integration over the finite detector solid angles. The coincidence rate is now only proportional to the differential $2p$ capture cross section if $D=1$. The values of D calculated for our detector geometry are listed for some scattering angles in Table I. From this table one can see that except for 0° , D is equal to 1 to within less than 10% and that the coincidence count rate is thus proportional to the differential $2p$ capture cross section to a good approximation. In order to estimate the error introduced by the acceptance angles of the detectors, we rewrite (4) as

$$\frac{dN}{dt} = C \left[\left. \frac{d\sigma}{d\Omega} \right|_{2p} + 2(D-1) \left. \frac{d\sigma}{d\Omega} \right|_1 \right], \quad (5)$$

where $(d\sigma/d\Omega)_0$ obtained from Eq. (3) was substituted in (4). The second term on the right-hand side can be considered the error introduced by the acceptance angles of the detectors. According to measured data by Hippler *et al.* [31] $\sigma_1/\sigma_{2p} \approx 0.2$ for the He target. Assuming that these numbers are similar for the Ne and Ar targets and that they do not vary significantly with scattering angle, we estimate the relative error to be of the order of 4% at

TABLE I. Result of the integration of Eq. (1) over the acceptance angles of the projectile and the photon detectors. For $D=1$ the coincidence count rate is proportional to the differential $2p$ capture cross section.

Angle (mrad)	D
0.0	0.79
0.2	0.91
0.4	0.98
0.6	0.99
0.8	0.995
1.0	0.995
1.5	1
2.0	1

0.2 mrad. At larger scattering angles the error will even be smaller since the deviation of D from 1 is systematically decreasing with increasing scattering angle.

IV. RESULTS AND DISCUSSION

In Fig. 5 the differential cross sections for capture with simultaneous emission of a photon are plotted as a function of the laboratory scattering angle for He, Ne, and Ar, respectively. The error bars show statistical errors only. The systematic errors are mainly determined by the errors of the total $2p$ capture cross sections to which we normalized our relative differential cross sections integrated over the scattering angle. The errors reported for the measured total $2p$ capture cross sections are 30% for He [29] and 50% for Ne and Ar [15].

While the total $2p$ capture cross sections depend strongly on the target ($1.9 \times 10^{-18} \text{ cm}^2$ for He, $3.4 \times 10^{-18} \text{ cm}^2$ for Ne, and $1.1 \times 10^{-17} \text{ cm}^2$ for Ar), only

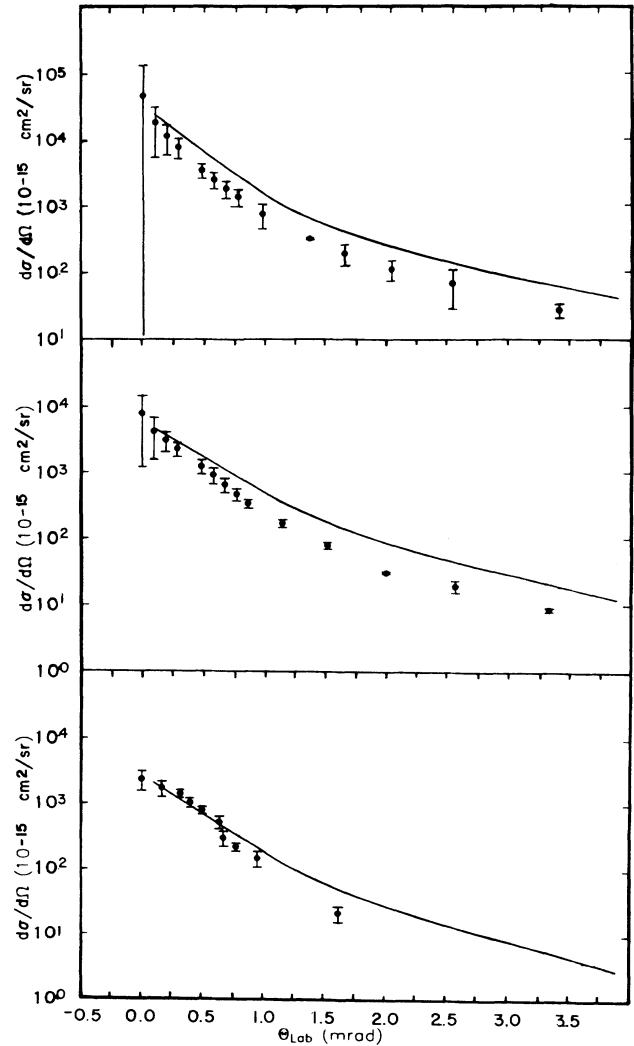


FIG. 5. Differential cross sections in the laboratory frame for capture to the $2p$ state of hydrogen from Ar (top), Ne (center), and He (bottom).

small differences are observed in the shapes of the differential cross sections. There is a noticeable difference only at very small angles where the data for He are somewhat flatter than those for Ne and Ar. In an attempt to understand the similarity in the shape of the differential cross sections, we have calculated impact-parameter-dependent $2p$ capture probabilities $P(b)$ from the differential cross sections. $P(b)$ is related to the differential cross section by

$$P(b) = \left[\frac{d\sigma}{d\Omega} \right] \sin\Theta / \left[b \frac{db}{d\Theta} \right]. \quad (6)$$

The impact parameter b and $db/d\Theta$ were calculated using a model potential for the target nucleus fit to the Hartree-Fock potential, in which screening is accounted for by a superposition of a number of exponential screening functions [32]. It should be noted that by calculating the impact parameter using only screened nuclear potentials, binary collisions of the projectile with the target electrons are neglected. On the other hand, at large collision distances the scattering angle may be strongly affected by such binary collisions with the target electrons. Here, we use the impact-parameter dependence only to make some qualitative arguments. The $P(b)$'s ob-

tained for the three targets using Eq. (6) are shown in Fig. 6. For all targets a pronounced maximum can be seen at relatively large impact parameters. If one extrapolates the small-impact-parameter wing of these maxima to zero-impact parameter, it appears that the $2p$ capture probability is essentially zero for very close collisions for the He and Ne targets and is strongly reduced for the Ar target compared to far collisions.

At a given impact parameter, the projectile predominantly penetrates the electron cloud for a certain n state. A comparison of the impact-parameter-dependent probabilities with the spatial distribution of the electrons of a given n state an electron is captured from. The curves in Fig. 6 show the square of the product of the radial wave function with the radius $|r\phi(r)|^2$ for the individual nl terms of the ground state and for their sum. For the wave functions the parametrized representation given by Clementi and Roetti [33] was used. It can be seen that for all targets the maximum in $P(b)$ is close to the maximum in $|r\phi(r)|^2$ for the valence electrons and that $P(b)$ is small in the region where $|r\phi(r)|^2$ maximizes for the

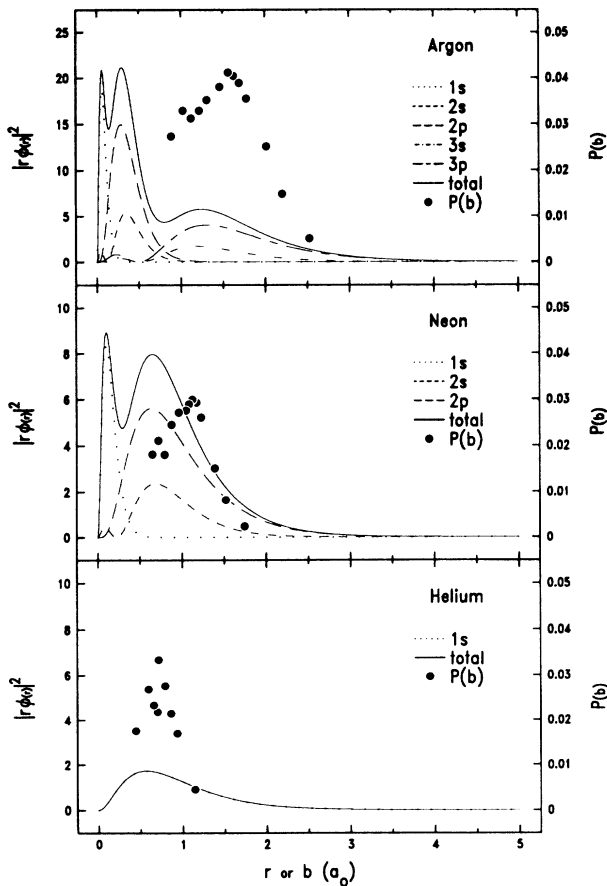


FIG. 6. Impact-parameter-dependent $2p$ capture probabilities for Ar (top), Ne (center), and He (bottom). Also shown is the radial distribution of the target electrons for various nl states.

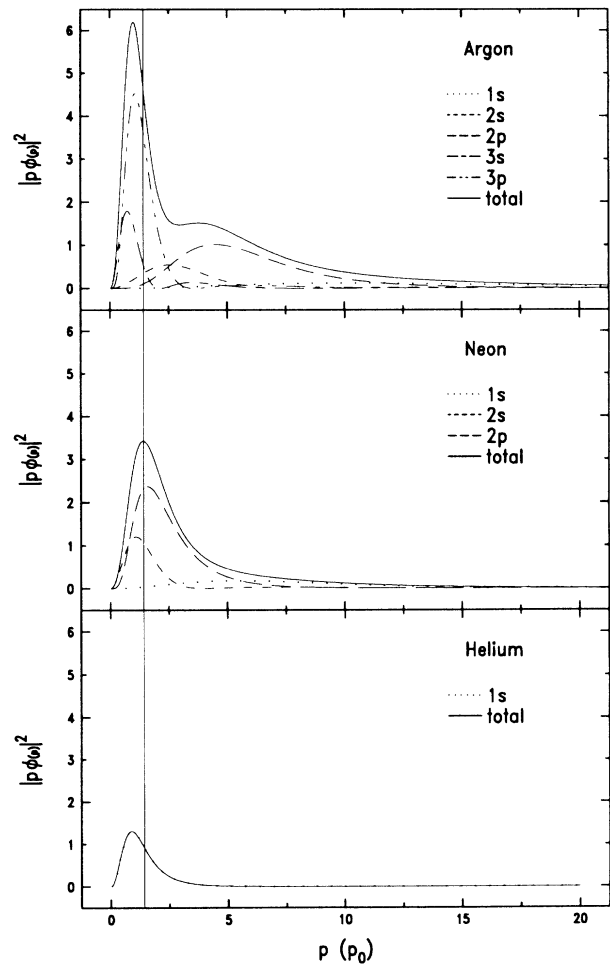


FIG. 7. Momentum distribution of the target electrons for Ar (top), Ne (center), and He (bottom). The vertical line indicates the reduced projectile momentum, which is the electron mass times the proton velocity.

inner shells. In the case of the Ar target there may be some contributions to $P(b)$ when the L shell or even the K shell is penetrated if the data are extrapolated to small-impact parameters.

From the radial wave functions alone one cannot understand why the $P(b)$'s are strongly reduced for close collisions where the inner shells are penetrated. However, a capture process does not only require a spatial overlap between the projectile and the target electron, but also an overlap of the projectile velocity with the velocity distribution of the target electrons. In Fig. 7 the square of the product of the electron momentum with the momentum wave function, which is simply the Fourier transform of the radial wave function, is plotted versus the electron momentum. The vertical line indicates the reduced projectile momentum (projectile velocity times electron mass). Again, as for the various radial wave functions, there is a strong overlap in momentum space with the valence electrons and almost zero overlap with the inner shells. Only the Ar target has a small overlap of the reduced projectile momentum with the momentum distribution of the $2s$ state. This lack of overlap in momentum space with the inner shells, which are only penetrated in close collisions, is a possible explanation for the small $2p$ capture probabilities at small-impact parameters. Also, the nonzero overlap with the $2s$ state of Ar may explain why the extrapolated probabilities do not fall as quickly at small-impact parameters for the Ar target as they do with the He and Ne targets. We are therefore led to conclude that capture to the $2p$ state is dominated by the valence electrons of the target with possibly some small contributions from the $2s$ state for the Ar target.

The full curves in Fig. 5 show a classical-trajectory Monte Carlo calculation by Olson and Lundy [34] for Ne and Ar and by Schultz *et al.* [35] for He. There are currently no other calculations available to the best of our knowledge. The CTMC technique has been described in detail by, e.g., Olson and Salop [36]. In this calculation only capture from the valence shell of the target atom was taken into account, which is consistent with

our conclusion above from our data. For the He target the agreement between the calculation and the data is excellent both in absolute magnitude and in the angular shape. Only at the largest measured scattering angle is there significant discrepancy. For the Ne and the Ar target there is good agreement with the data in the angular shape. Even though in absolute magnitude the calculation is somewhat higher in both cases the discrepancy is within the systematic error in the data.

V. CONCLUSIONS

We have studied capture to the $2p$ state of hydrogen for protons colliding with He, Ne, and Ar targets at a projectile energy of 50 keV by measuring coincidences between neutralized projectiles and Lyman- α photons emitted by the hydrogen atoms. While the total cross sections depend strongly on the target the angular distribution of the differential cross sections do not differ significantly for the targets studied here. From a comparison of the impact-parameter-dependent probabilities extracted from the measured differential cross sections with the spatial distribution of the target electrons we conclude that capture to the $2p$ state is dominated by the valence electrons of the target for the collision systems studied here. The small overlap of the reduced projectile momentum with the momentum distribution of the inner-shell target electrons explains why capture from an inner shell is unlikely even for a close collision. The dominance of the valence electrons in $2p$ capture, in turn, explains the similarity in the angular distribution of the differential cross sections for different target Z 's. The very good agreement of our data with a calculation which only takes capture from the valence shell into account supports our conclusion.

ACKNOWLEDGMENTS

We would like to thank Dr. R. E. Olson for making his calculations available to us prior to publication. This work was supported by the National Science Foundation.

-
- [1] E. Rutherford, *Philos. Mag.* **47**, 277 (1924).
 - [2] J. R. Oppenheimer, *Phys. Rev.* **31**, 349 (1928).
 - [3] H. C. Brinkmann and H. A. Kramers, *Proc. Acad. Sci. Amsterdam* **33**, 973 (1930).
 - [4] N. Bohr and J. Lindhard, *K. Dan. Vidensk. Selsk. Mat.—Fys. Medd.* **28**, No. 7 (1954), and references therein.
 - [5] H. B. Gilbody, *Adv. At. Mol. Phys.* **15**, 293 (1979).
 - [6] R. E. Olson and A. S. Schlachter, *Comments At. Mol. Phys.* **18**, 237 (1986).
 - [7] R. A. Mapleton, *Theory of Charge Exchange* (Wiley-Interscience, New York, 1972).
 - [8] M. Kimura and C. D. Lin, *Phys. Rev. A* **34**, 176 (1986).
 - [9] M. Kimura and N. F. Lane, *Advances in Atomic and Molecular Physics*, edited by D. Bates and B. Bederson (Academic, New York, 1990), Vol. 26, p. 79.
 - [10] B. H. Bransden and D. P. Dewangan, *Advances in Atomic and Molecular Physics*, Vol. 25, p. 343.
 - [11] C. Havener, in *Physics of Electronic and Atomic Collisions*, edited by W. R. MacGillivray, I. E. McCarthy, and M. C. Standage (Hilger, Bristol, 1992), p. 381.
 - [12] W. Schwab, G. B. Baptista, E. Justiniano, R. Schuch, H. Vogt, and E. W. Weber, *J. Phys. B* **20**, 2825 (1987).
 - [13] C. Schmeissner, C. L. Cocke, R. Mann, and W. Meyerhoff, *Phys. Rev. A* **30**, 1661 (1984).
 - [14] C. Biedermann *et al.*, *Phys. Rev. A* **43**, 4075 (1991).
 - [15] R. H. Hughes, E. D. Stokes, Song-Sik Choe, and T. J. King, *Phys. Rev. A* **4**, 1453 (1971).
 - [16] M. B. Shah, J. Geddes, and H. B. Gilbody, *J. Phys. B* **13**, 4049 (1980).
 - [17] P. J. Martin, K. Arnett, D. M. Blankenship, T. J. Kvale, J. L. Peacher, E. Redd, V. C. Sutcliffe, J. T. Park, C. D. Lin, and J. H. McGuire, *Phys. Rev. A* **23**, 2858 (1981).
 - [18] T. R. Bratton, C. L. Cocke, and J. R. Macdonald, *J. Phys. B* **10**, L517 (1977).

- [19] E. Horsdal-Pedersen, C. L. Cocke, and M. Stöckli, *Phys. Rev. Lett.* **50**, 1910 (1983).
- [20] H. Vogt, R. Schuch, E. Justiniano, M. Schulz, and W. Schwab, *Phys. Rev. Lett.* **57**, 2256 (1986).
- [21] R. H. McKnight and D. H. Jaecks, *Phys. Rev. A* **4**, 2281 (1971).
- [22] R. Hippler, M. Faust, R. Wolf, H. Kleinpoppen, and H. O. Lutz, *Phys. Rev. A* **31**, 1399 (1985).
- [23] R. Hippler, M. Faust, R. Wolf, H. Kleinpoppen, and H. O. Lutz, *Phys. Rev. A* **36**, 4644 (1987).
- [24] D. G. Seely, S. W. Bross, A. D. Gaus, J. W. Edwards, D. R. Schultz, T. J. Gay, J. T. Park, and J. L. Peacher, *Phys. Rev. A* **45**, R1287 (1992).
- [25] J. T. Park, *Advances in Atomic and Molecular Physics*, edited by D. Bates and B. Bederson (Academic, New York, 1983), Vol. 19, p. 67.
- [26] S. R. Walther, K. N. Leung, and W. B. Kunkel, *Rev. Sci. Instrum.* **57**, 1531 (1986).
- [27] J. T. Park, J. M. George, J. L. Peacher, and J. E. Aldag, *Phys. Rev. A* **18**, 48 (1978).
- [28] E. Redd and J. T. Park, *Rev. Sci. Instrum.* **55**, 119 (1984).
- [29] Oak Ridge National Laboratory Report No. 6086/V1, 1990 (unpublished).
- [30] J. Macek and D. H. Jaecks, *Phys. Rev. A* **4**, 2288 (1971).
- [31] R. Hippler, W. Harbich, M. Faust, H. O. Lutz, and L. J. Dubé, *J. Phys. B* **19**, 1507 (1986).
- [32] H. L. Cox and R. A. Bonham, *J. Chem. Phys.* **47**, 2599 (1967).
- [33] E. Clementi and C. Roetti, *At. Data Nucl. Data Tables* **14**, 177 (1974).
- [34] R. E. Olson and C. Lundy (private communication).
- [35] D. R. Schultz, C. O. Reinhold, R. E. Olson, and D. G. Seely, *Phys. Rev. A* **46**, 275 (1992).
- [36] R. E. Olson and A. Salop, *Phys. Rev. A* **16**, 531 (1977).