Electron Spectroscopy at Storage Rings



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Scientific goals:

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- I reaction mechanisms excitation and decay channels (impact parameter dependence) correlated processes: (R)TE (sub shell population), DR (crossed beam)
 - few electron systems in strong fields (high Z) Rydberg states, meta stable configurations
 - Rydberg states, meta stable configurations Auger decay lines \rightarrow QED contributions, FS-splitting, Breit-terms
- interplay of nuclear and atomic structures: conversion electrons from H-, He-, Li- like ions after Coulomb excitation
- precision measurement of 1s binding energy conversion coefficients \rightarrow e- wave function at nucleus
 - HFS : nuclear magnetic moments from excited nuclei ($\tau \le 1$ ns)





<u>Synthetic spectrum</u> of 857 eV electrons from internal conversion of H-like Gd⁽³⁺ projectiles. Ep* denotes the cusp peak at projectile velocity. Ee* and Ee* correspond to the kinematical shifts (forward - backward) observed at zero degree.



Example of Coster-Kronig transitions $(1s^2pnl) \Rightarrow 1s^2ss + c (n = 6...*)$ from O⁺ projectiles after electron capture from He in slow collisions measured at 0^o. Such states can be studied at storage rings for high atomic numbers where the stretching of line-spacing is amplified due to the high v_{μ} Besides collisional generation-mechanisms (capture, cecitation ...) information is obtained on QED, relativistic and magnetic electron interactions, similar to DR – experiments. Here Auger transitions are studied in situ without background from charge changing collisions or from disturbing magnetic and electric cooler fields.



<u>The hyperfine splitting</u> of 1s-conversion electrons can be observed if an electron spectrometer with high resolution ($\Delta E E < 6x 10^{-5}$) is applied. It gives access to HFS of excited nuclear states with life times below nano seconds allowing determinations of magnetic momenta and nuclear g-factors. The intensities -besides the statistical weights- reflect state-dependent nuclear life-times within the spectrometer acceptance length.



Instrumental Method: zero degree electron spectrometry:

- I a dispersion free magnetic transport system (270° deflector, fig.1) transports electrons from the (gas) target in a forward cone out of the beam line. Large momentum (ΔP/P ~ 2,5/1) and angle (Δθ = ± 12°, Δϑ = ±8°) acceptances enable the study of a large band of electron energies with high efficiency (fig.2).
 II energy measurements are performed by
 - energy measurements are performed by 1) a window-less SI-detector covering the complete energy band with moderate energy resolution or by

 a high-resolution iron-free magnetic analyzer (r^{-1/2}-field) with high dispersion.



Ions in storage rings are dressed with Rydberg electrons via different interactions with cooler- electrons. A certain class of n,l states are stable against field ionisation (quenching) in the ring magnets with life times $\tau > 0.2 \, \mu$ s. They can be measured at the experimental area free from photon and electron noise of the cooler, providing they fall into the ring q-acceptance. Collisions with target atoms transfer Rydberg electrons to the cusp – continuum or generate (via isolated core excitations like $1s^22s \rightarrow 1s^22p_1$, ...) specific Auger decay lines which are fully received from the forward electron spectrometer. Alternatively photo ionisation and excitations are release these upper bound electrons more distinctively into the analyzer. The system represents a versatile tool for investigation of cooler induced recombination processes as well as for precise spectroscopic studies.