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On the characterisation of a Bragg spectrometer with X-rays from an ECR source

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Abstract

Narrow X-ray lines from helium-like argon emitted from a dedicated ECR source have been used to determine the response function of a Bragg crystal spectrometer equipped with large area spherically bent silicon (111) or quartz (10 $\bar{1}$) crystals. The measured spectra are compared with simulated ones created by a ray-tracing code

based on the expected theoretical crystal's rocking curve and the geometry of the experimental set-up.

Key words: X-ray spectroscopy, Exotic atoms, ECR source

PACS: 29.30.Kv, 36.10.-k, 29.25.Ni

1 Introduction

In an ongoing experiment at the Paul Scherrer Institute (Switzerland) the strong interaction broadening of the ground state level of pionic hydrogen of about 1 eV should be determined with an accuracy of less than 10 meV by measuring the line shape of X-ray transitions feeding the atomic 1s state [1,2,3]. The energies of the pionic hydrogen X-rays in question are 2.436, 2.886 and 3.043 keV for the 2p-1s, 3p-1s and 4p-1s transitions, respectively. The experiment uses a Bragg spectrometer in Johann mounting [4] and was equipped during recent measurements with spherically bent silicon or quartz crystals having a diameter of 100 mm and a thickness of 0.3 mm or 0.2 mm, respectively. The curvature radius R_C of the bent crystals is $R_C = 2982.5 \pm 0.6$ mm. For these parameters bent crystal theory predicts a negligible influence of the bending process on the crystal's rocking curve [5]. Imperfections in the crystal material itself and defects produced during the fabrication could lead to deviations which should be determined experimentally. In order to meet the requirements imposed by the pionic hydrogen experiment a determination of the spectrometer's response function requires at least 20000 events in the cali-

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bration line. Moreover the measurement should be performed under conditions equivalent to the real experiment.

In earlier experiments a rough energy calibration was done by using X-rays from geometrically extended fluorescence targets excited by means of X-ray tubes [6]. A determination of the response function is out of reach with this method as the natural line widths are much broader than the resolution of the spectrometer. In addition the shape of these lines is distorted by satellite transitions.

Alternating the pionic hydrogen measurement in regular intervals with calibration experiments using narrow X-ray lines from other exotic atoms would be ideal. The measured low count rates of about 100/hour, however, render such an approach unfeasible.

The use of X-rays emitted from highly ionized ions therefore offers an appropriate solution albeit in a remote measurement. It makes use of the fact that for low to medium Z atoms the E1 X-rays from hydrogen-like ions as well as M1 X-rays from helium-like ions have natural line widths negligibly small compared to the expected resolution of the Bragg crystals. To give an example, the lifetime of the M1 transition of helium-like argon used mostly in the present study is $0.21 \mu\text{s}$ only [7]. The required X-rays are emitted in copious quantities from ECR or EBIT sources. ECR sources are preferable as the X-rays are emitted from a geometrically extended region important for a Johann set-up whereas EBIT sources produce X-rays in the region of the very compressed electron beam only, which would require an observation perpendicular to the electron beam cutting down the accepted intensity considerably. Decisive, however, is the low kinetic energy of the ions in ECR sources on

the level of less than 1eV only, which had been determined at different ECR sources with high resolution optical spectroscopy [8,9]. From this a Doppler broadening of less than 40 meV can be expected for transitions around 3 keV energy in helium-like argon. In principle EBIT sources could be tuned to low kinetic energies of the captured ions by reducing the voltage of the drift tube, which, however, results in a drastic reduction in intensity [10].

The intention of this paper is much different from earlier publications using crystal spectrometers in combination with ECR sources. Those mostly aimed at a determination of plasma properties leading to a deeper understanding of the processes governing the physics of these sources [11,12,13]. Here, for the first time the characterisation of a crystal spectrometer with a narrow X-ray line from an almost completely ionized atom is described. For this purpose the transition from helium-like argon at 3.104 keV [7] close in energy to the 4p-1s transition in pionic hydrogen was used.

2 Experimental set-up

The experiment as depicted in Figure 1 can be subdivided into three parts:

- The Electron-Cyclotron Resonance Ion Trap (ECRIT) consists out of a superconducting split coil magnet, which together with special iron inserts, provides the mirror field configuration, an AECR-U style permanent hexapole magnet and a 6.4 GHz power regulated microwave emitter [14]. The mirror field parameters provide one of the highest mirror ratios for ECR sources with a value of 4.3 over the length of the plasma chamber. The hexapole is cooled at the front pieces and on the inner radius by a

forced flow of demineralised water. The plasma chamber is formed by a 0.4 mm thick stainless steel tube of inner diameter of 85 mm and a length of 265 mm axially limited by copper inserts. At the position of the hexapole gap the stainless steel tube is perforated by a series of 2.5 mm diameter holes allowing for radial pumping in addition to axial pumping. The microwave HF power is introduced directly to the plasma chamber with waveguides exhibiting a small angle to the axis. In this way the path of the X-rays was at no point cut by any obstacle which could distort the shape of the response function.

An extraction voltage of 2 kV had been routinely applied at the side opposite to the crystal spectrometer. The measurement of the current drawn served as a control for a stable operation.

A reference pressure (without plasma) of $1.7 \cdot 10^{-7}$ mbar was achieved. Gas filling was supplied radially by UHV precision leak valves through the gaps in the open structure hexapole. The gas composition was routinely surveyed with a quadrupole mass spectrometer. For an optimised plasma source a drastic increase of the number of energetic electrons was discovered which required the use of a cleaning magnet installed at a distance of one meter in front of the crystal.

- A silicon(111) as well as a quartz($10\bar{1}$) crystal were investigated, which had been recently applied for measuring pionic hydrogen transitions [15]. The Bragg angles Θ_B , corresponding to the M1 transition of helium-like argon with an energy of 3.104 keV, are $\Theta_B = 36.68^\circ$ for the quartz and $\Theta_B = 39.57^\circ$ for the silicon crystal. The crystals were installed at a distance of 2330 mm from the centre of the ECRIT resulting in a position of the plasma about 500 mm outside the Rowland circle.
- A CCD pixel detector with a pixel size $40 \mu\text{m} \times 40 \mu\text{m}$ and an energy

resolution of 140 eV at 3 keV was used to detect the X-rays [18]. The detector consisted out of six chips with 600 x 600 pixels each resulting in a total height of 72 mm and a width of 48 mm. The distance of the CCD detector from the crystal could be changed remotely over a length of 86 mm without breaking the vacuum.

The CCD detector and the following electronics were protected against light as well as the HF stray field by a 30 μm thick Beryllium window installed in the vacuum tube in front of the CCD cryostat.

3 Tuning the ECRIT

In a first step with low HF power and minimum field values around 1.4 kG the proper working of the whole set-up was checked with X-rays of low ionisation degree. In consequence the HF power was gradually increased and with the Bragg angles set to the proper values for a simultaneous observation of the well separable X-rays from both Ar^{10+} and Ar^{11+} an optimum for the HF power was found at about 450 Watt. The turn-around time from introducing a change of parameters to the evaluation of a measurement was about one minute. Going to Bragg angles valid for higher ionisation degrees the strength of the mirror field as well as the argon pressure were optimised for a maximum intensity of X-rays from highly ionised atoms. In addition a gas mixture Ar/O_2 with a mixing ratio of 1:9 was found to be necessary to achieve this goal [16]. The working pressure was $1.4 \cdot 10^{-6}$ mbar. After a total tuning time of 10 hours the M1 transition $^3S_1-^1S_0$ at 3104 eV in Ar^{16+} could be observed as shown in Figure 2. The typical illumination time of the CCD chips of 1 min before readout resulted in a high probability of double hits per pixel even in the M1

peak. In order to reduce the intensity and especially in order to improve the peak/background ratio an aluminium collimator was inserted at a distance of 185 mm from the centre of the plasma into the direction of the crystal. It left a free hole with the dimensions 16 mm(horizontal) x 10 mm(vertical), which was wide enough not to influence the measurement of the response function.

A 6.4 GHz emitter proved to be sufficient to provide a highly intensive source of M1 X-rays from Ar^{16+} ions in the special hybrid magnetic structure used with radial pumping together with the gas mixing technique. After turning the system on, it worked in a stable and reproducible way after about one hour warm up time. The observation of the X-rays alone was sufficient for a successful tuning in a short time thus making the usual momentum analysis of extracted ions superfluous. There was no measurable effect of the applied extraction voltage on the intensities of the X-rays for values between 0 and 6 kV, which is the maximum achievable voltage.

4 Response function of the crystal spectrometer

With the ECR source a number of 20000 events was reached for the narrow M1 transition of helium-like Argon in about 30 minutes time to be compared with a number of 5000 counts reached after 40 hours with X-rays from pionic methane. A total of about 10 hours was needed, however, to determine the spectrometer's response function in sufficient detail including changes of the distance CCD detector-crystal (focal scans) and changing apertures in front of the crystals. The measurements were confronted with theoretical predictions using the following procedure:

- The XOP program package [17] was used as a theoretical reference for the present measurements. It provided a theoretical rocking curve for a flat ideal crystal.
- The theoretical rocking curve was used as input for a Monte Carlo X-ray tracking routine. With it the Bragg reflection of a spherically bent crystal was calculated for the geometrical conditions of the special set-up parameters like source diameter, detector position and aperture at the crystal.
- The output of the tracking routine can be considered to be the theoretical limit of the response function which under ideal circumstances would reproduce the data. In order to allow for additional broadening this response function is folded with a Gaussian distribution. Its width was determined with a least square fitting of the data using the program package MINUIT from the CERN program library. The statistics of the Monte Carlo response function was chosen high enough not to contribute to the error of the fit.

Two types of changes in the experimental set-up were investigated with this procedure. At first spectra were taken for different distances of the CCD detector from the crystals (focal scans). In a second step with the CCD detector located at the focal position different apertures were mounted in front of the crystals.

The focal scans served the two purposes:

- to verify the optimum focal position, which is given by $R_C \sin \Theta_B$. For this position the width of the response function should be at minimum. The minimum was searched for by moving the CCD in steps of 2-3 mm in direction to and from the crystal.
- to measure the line shape variations as a function of the crystal-detector

distance which served as a check of the validity of X-ray tracking calculations.

An example of a spectrum far off focus is shown in Figure 3 for the quartz crystal with the CCD detector shifted by 12.5 mm from the focal position into the direction of the crystal. The fit allowing for an additional Gaussian broadening reproduces the measured spectrum in detail.

In a second step the measured response function was compared with the simulated response function, as function of the crystal's reflection surface. This was achieved by applying apertures in front of the crystal and allowed the study of possible irregularities in the response function due to crystal material imperfections and/or deviations from the spherical shape. Four different apertures were used: two circular ones with openings of diameter 40 mm and 60 mm, respectively and two rectangular ones with horizontal openings of 40 mm and 60 mm, respectively, and a free diameter of 95 mm elsewhere. In table 1 the results for the theoretical response function are confronted with the measurement. Both lineshapes were fitted with a single Gaussian and result in the values for the FWHM depicted in columns 2 and 3. The rocking curve width for a plane crystal, as obtained by XOP, fitted by a single Gaussian resulted in FWHM values of $106.6 \mu\text{rad}$ for silicon and $97.4 \mu\text{rad}$ for quartz.

The difference between the FWHM of the spectrometer's response function and the crystal's rocking curve reflects expected geometrical contributions mainly due to the horizontal extension of the crystal (Johann broadening). In order to cope with the difference of the measured response function to the theoretical one, a least square fitting of the data was done with the width of a Gaussian folded with the theoretical response function as fit parameter. The

result of this fit is shown in the fourth column and is considered to be the main result of the present investigation.

The Si crystal shows an additional Gaussian contribution of $34.1(1.8) \mu\text{rad}$ as an average of all apertures chosen. This can be interpreted as an overall deviation of the crystal geometry from spherical shape eventually caused by the production process or by a failure in the crystal material. For quartz the three smaller apertures show an additional broadening on average of about $24.2(1.8) \mu\text{rad}$, but opening the aperture to 60 mm rectangular results in a considerable worsening. This points to an irregularity in the crystal material or its mounting in the upper and/or lower part of the reflecting surface and will be investigated in a future measurement for this special crystal.

The different dispersions of $3.756 \text{ meV}/\mu\text{rad}$ for Si and $4.166 \text{ meV}/\mu\text{rad}$ for quartz result in an energy resolution of about $478(6) \text{ meV}$ for the Si crystal and $506(8) \text{ meV}$ for the quartz crystal, both equipped with a 60 mm horizontal aperture. Going to 60 mm circular aperture these values reduce to $456(8) \text{ meV}$ in the case of Si and to $459(6) \text{ meV}$ in the case of quartz.

The result of a measurement for a circular aperture with a free diameter of 60 mm is shown in Figure 4 for the quartz crystal together with the corresponding fit.

In terms of energy resolution the values for the additional Gaussian correspond to $100(8) \text{ meV}$ for quartz and to $128(6) \text{ meV}$ for Si. Being different they cannot be caused by a common Doppler broadening caused by the motion of the argon ions. Moreover it should be recalled that from optical measurements an upper limit for the Doppler broadening of about 40 meV had been obtained [8,9].

Compared to the total width of the response functions the Doppler broadening can be neglected and the present accuracy is sufficient for the determination of the strong interaction width in pionic hydrogen. As the response function has been measured at 3.104 keV only, its shape has to be extrapolated to the lower energies of the different pionic hydrogen transitions. If one interpretes the additional Gaussian broadening as resulting from a geometrical deformation only this extrapolation is straightforward and the Gaussian broadening can be folded into the rocking curve before the X-ray tracking is performed.

5 Conclusion and Outlook

The characterisation of two large area spherically bent crystals was performed with M1 X-rays from helium-like argon with two orders of magnitude higher statistics than achievable with exotic atoms. The geometrical conditions of the characterisation were equivalent to the pionic hydrogen experiment. The measurement could be understood in terms of a Monte-Carlo X-ray tracking routine allowing, however, for an additional Gaussian broadening. From this a prescription for extrapolating the result to different energies became possible.

In future it is planned to improve on the peak/background ratio by insertion of a proper slit of tapered material with a hole of 1 mm(vertical) x 30 mm (horizontal) near the plasma chamber in the direction of the crystal.

In addition the achievable pressure will be optimised in reducing the surface of the iron insertion pieces drastically. The use of M1 transitions from helium-like chlorine (supplied from the gaseous compound CHClF_2 , chlorodifluoromethane) and helium-like sulfur from SO_2 is envisaged for future ex-

periments as a countercheck of the assumptions and conclusions made here. The energies at 2.430 and 2.756 keV, respectively, are close to the 2p-1s and the 3p-1s transition energies in pionic hydrogen.

Additionally a thorough investigation of X-ray intensities as a function of its radial origin inside the plasma is feasible with the special set-up of the crystal spectrometer. The described set-up allows to determine the Doppler broadening from the ion velocity distribution directly with crystals with better resolution. Similarly the observation of such a broadening as a function of the power and frequency injected into the ECR source may be studied.

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Figure captions

Fig.1. Set up of the PSI ECRIT together with the Bragg crystal spectrometer

Fig. 2. The spectral region for argon X-rays from Ar^{14+} , Ar^{15+} and Ar^{16+} (M1 transition) as emitted from the PSI ECRIT and measured with a Si (111) crystal. The energy values are from [7]. One channel (pixel) corresponds to 79 meV.

Fig. 3. The spectrum of the Ar^{16+} 3104 eV M1 line measured with a quartz ($10\bar{1}$) crystal is shown together with a fit for a detector position of 12.5 mm out of focus in direction of the crystal.

Fig. 4. The spectrum of the Ar^{16+} 3104 eV M1 line measured with a quartz ($10\bar{1}$) crystal is shown for the focal position together with a fit. A circular aperture with a diameter of 60 mm had been applied.

Table 1

The FWHM in μrad of the theoretical response function is compared with the measurement for a Si (111) as well as a quartz ($10\bar{1}$) crystal for different openings (rectangular or circular) of the diffraction surface. The last column shows the FWHM for a Gaussian to be folded with the theoretical response function in order to fit the measurement best.

Crystal/aperture	Theor. resp. fct.	Measurement	Gaussian
Si/Rect. 60	116.0(0.5)	127.4(1.7)	38.0(4.0)
Si/Circ. 60	114.5(0.5)	121.5(2.2)	30.3(3.5)
Si/Rect. 40	108.5(0.6)	119.8(2.2)	34.6(3.2)
Si/Circ. 40	108.7(0.6)	117.5(2.1)	34.2(2.7)
quartz/Rect. 60	109.4(0.3)	121.5(1.8)	38.0 (4.1)
quartz/Circ. 60	106.2(0.3)	110.2(1.4)	21.5 (3.2)
quartz/Rect. 40	98.8(0.4)	104.2(1.7)	25.1 (2.9)
quartz/Circ. 40	98.4(0.4)	104.2(1.6)	25.6 (3.1)

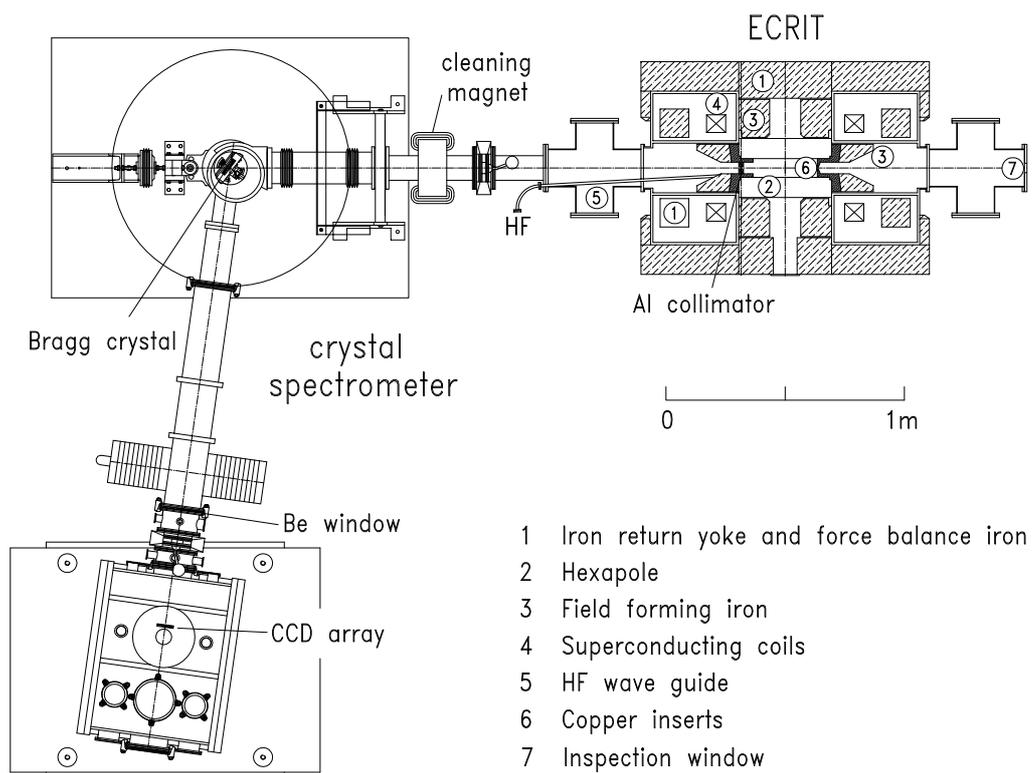


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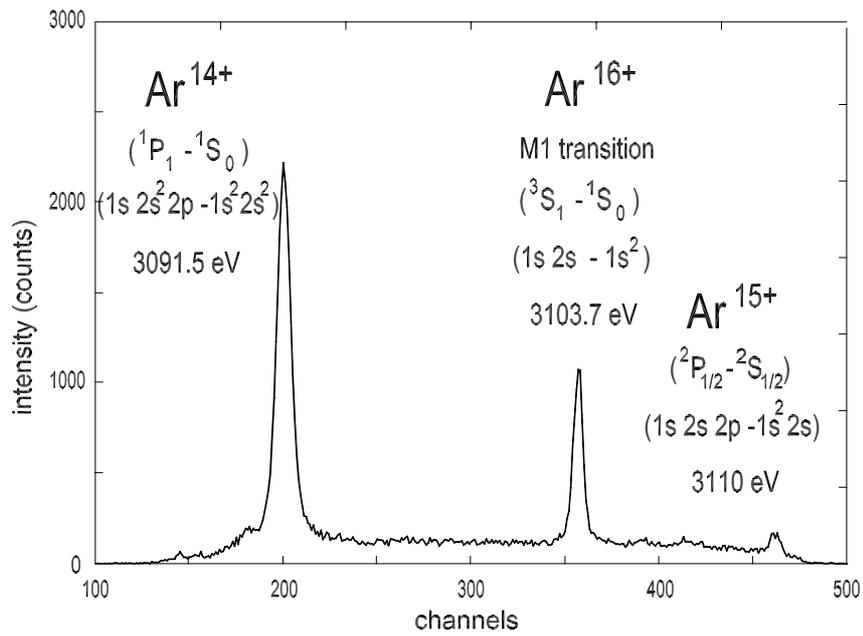


Fig. 2. The spectral region for argon X-rays from Ar¹⁴⁺, Ar¹⁵⁺ and Ar¹⁶⁺ (M1 transition) as emitted from the PSI ECRIT and measured with a Si (111) crystal. The energy values are from [7]. One channel (pixel) corresponds to 79 meV.

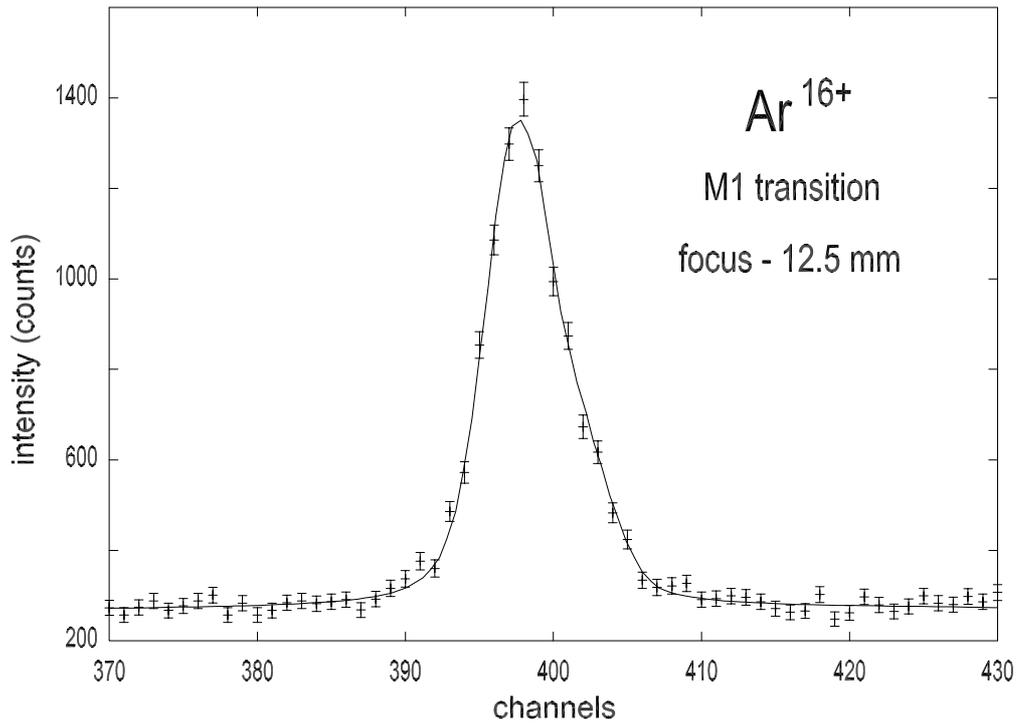


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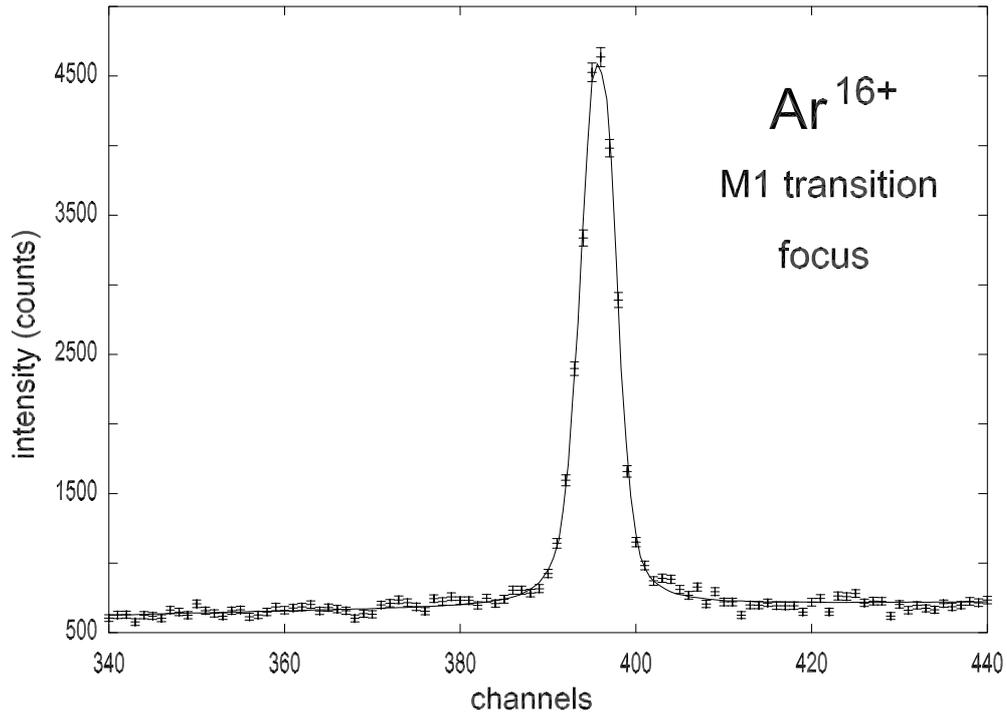


Fig. 4. The spectrum of the Ar^{16+} 3104 eV M1 line measured with a quartz ($10\bar{1}$) crystal is shown for the focal position together with a fit. A circular aperture with a diameter of 60 mm had been applied.