

## Spontaneous fission of rutherfordium isotopes - total kinetic energies

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**Abstract.** The isotopes  $^{255,256,258}\text{Rf}$  were produced in the fusion-evaporation reactions  $^{50}\text{Ti} + ^{207,208}\text{Pb}$  and  $^{50}\text{Ti} + ^{209}\text{Bi}$  at GSI Darmstadt, using the velocity filter SHIP. Total kinetic energies of fragments from spontaneous fission for these isotopes were evaluated with a correction to pulse-height defect.

### 1 Introduction

Spontaneous fission (SF) is a decay mode with large impact on the stability of nuclei in the transfermium region. These isotopes are stabilized against SF mainly by microscopic shell effects [1]. Systematic studies of SF properties allow us to understand these effects and determine the production possibilities for the heaviest nuclei. Up to now, only a few results (often based on limited statistics of SF events) with measured total kinetic energy (TKE) were obtained for rutherfordium ( $Z = 104$ ) isotopes [2–5].

### 2 Experiment

The experiment was performed at GSI Darmstadt in Germany. The heavy-ion  $^{50}\text{Ti}$  beam with typical energies from 225 to 243 MeV was delivered by the UNILAC accelerator. The fusion-evaporation reactions  $^{50}\text{Ti} + ^{207,208}\text{Pb}$  were used for production of the studied isotopes  $^{255,256}\text{Rf}$  via  $xn$  evaporation channels from the compound nuclei  $^{257}\text{Rf}$  and  $^{258}\text{Rf}$ . The reaction  $^{50}\text{Ti} + ^{209}\text{Bi}$  was used for indirect production of  $^{258}\text{Rf}$  via EC decay of  $^{258}\text{Db}$ .

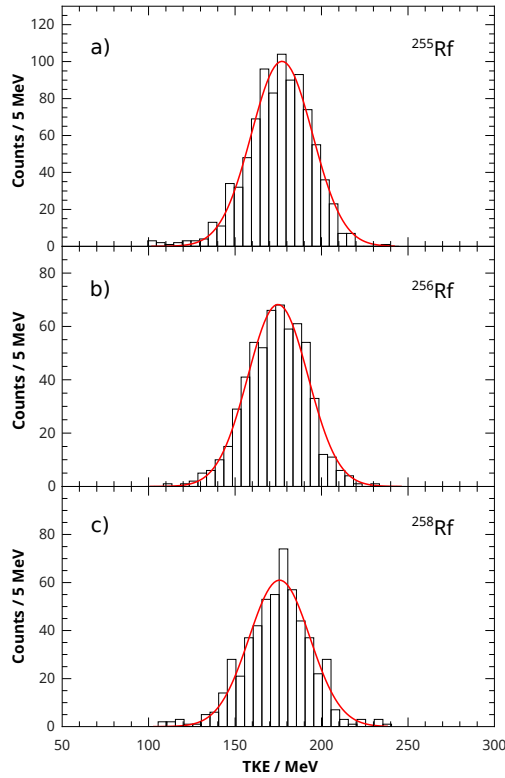
Evaporation residues (ER) were separated from the beam and other undesired background using the velocity filter SHIP [6] and delivered to a detection setup. After passing through the time-of-flight system they were implanted (6.5 – 6.8)  $\mu\text{m}$  deep into the 16-strip position-sensitive STOP detector. The same-type detectors arranged in the "BOX" geometry were placed in front of the

STOP detector to detect escaping particles. A clover detector with four Ge crystals was placed behind the STOP detector for  $\gamma$ - and X-ray detection.

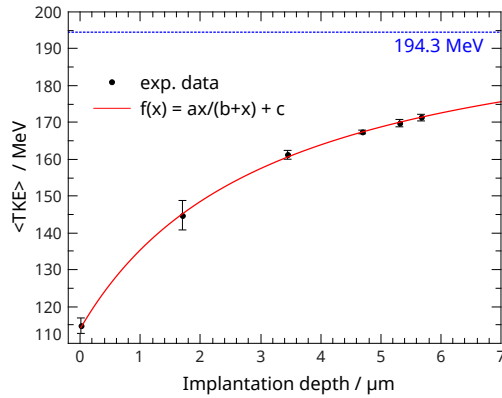
### 3 Results

In order to detect SF events from specific isotope, we used a time and position correlation search technique. In the case of  $^{255}\text{Rf}$ ,  $^{256}\text{Rf}$  we searched for correlations between an ER implantation signal and a high-energy signal corresponding to SF. For  $^{258}\text{Rf}$  instead of the ER implantation signal we used the low-energy signal corresponding to electrons originating from internal conversion process from de-excitation of levels populated after the EC decay  $^{258}\text{Db} \xrightarrow{EC} ^{258}\text{Rf}$ . The time conditions were set to  $\approx 5 \times$  half-life between signals for each isotope (8500 ms for  $^{255}\text{Rf}$ , 35 ms for  $^{256}\text{Rf}$  and 60 ms for  $^{258}\text{Rf}$ ) and position window of 1 mm in the detector for  $^{255}\text{Rf}$  and  $^{256}\text{Rf}$ . As position condition for  $^{258}\text{Rf}$ , we required only the same strip number since for many low-energy signals from electrons the position information was not registered. We collected several hundreds of SF events for  $^{255}\text{Rf}$ ,  $^{256}\text{Rf}$  and  $^{258}\text{Rf}$ . The TKE distributions of fission fragments for each isotope are shown in Fig. 1. The histograms contain all registered SF events - the cases when both fragments stayed in the STOP detector ( $\approx 60\%$ ) as well as events when one fragment escaped the STOP detector and was either detected in the BOX detector ( $\approx 30\%$ ) or escaped the detection setup completely ( $\approx 10\%$ ).

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**Figure 1.** Total kinetic energies for  $^{255}\text{Rf}$ ,  $^{256}\text{Rf}$  and  $^{258}\text{Rf}$  from all observed SF events for each isotope. Due to the pulse-height defect, the positions of gaussians are shifted to lower values.



**Figure 2.**  $\overline{TKE}$  from the SF of  $^{252}\text{No}$  vs. implantation depth of ER in the detector. Blue dashed line:  $\overline{TKE} = 194.3$  MeV of  $^{252}\text{No}$  from [3]. Red solid line: saturation-growth fit.

The crucial task for the evaluation of  $\overline{TKE}$  using Si detectors is the correction of deficit in measured energies. There are two main effects influencing the TKE measurements, discussed in previous studies [7–9] performed at SHIP. First, the energy calibration based on  $\alpha$ -decay energies is not valid for fission fragments due to the pulse-height defect (see e.g. [10]), resulting in a significant energy deficit. The second effect is a dependence of the detected TKE on the implantation depth of ER into the detector. In order to find the energy correction, we performed TKE measurements for  $^{252}\text{No}$  with known  $\overline{TKE} =$

194.3 MeV [3] at six different implantation depths [8]. Our observations (shown in Fig. 2) proved the strong non-linear saturation-like dependence of detected TKE on implantation depth (see [11] for more details). The correction to the energy deficit at given implantation depth was determined as a difference of the reference  $\overline{TKE}$  of 194.3 MeV and the value  $\overline{TKE}_{fit}$  from saturation-growth model fit of the six points (blue dashed and red solid lines in Fig. 2) as  $\Delta E = (194.3 - \overline{TKE}_{fit})$  MeV.

**Table 1.** In the table columns, from left to right, the isotope,  $\overline{TKE}$  evaluated in this work, reference value of  $\overline{TKE}$  and corresponding references are stated.

Isotope	$\overline{TKE}_{exp}$ [MeV]	$\overline{TKE}_{ref}$ [MeV]	Ref.
$^{255}\text{Rf}$	$199.5 \pm 2.7$	$199 \pm 3$	[4]
$^{256}\text{Rf}$	$198.7 \pm 2.8$	$198.9 \pm 4.4$	[2]
$^{258}\text{Rf}$	$198.2 \pm 3.0$	$197.6 \pm 1.1$	[2]

Considering the facts, that  $^{252}\text{No}$  is close in  $Z$ ,  $A$  and  $\overline{TKE}$  to studied Rf isotopes, we applied this correction to  $\overline{TKE}$  of  $^{255}\text{Rf}$ ,  $^{256}\text{Rf}$  and  $^{258}\text{Rf}$ . Corrected  $\overline{TKE}$  values for each isotope are summarized in Table 1 and compared to values from previous studies. The results are in a good agreement, which supports the validity of our correction.

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