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S. Mialle, Alexandre Quemet, A. Ponvienne, A. Gourgiotis, M. Aubert, et al.. The use of total evaporation method using Channeltron electron multipliers by thermal ionization mass spectrometry for europium isotope ratio measurements on picogram sample amounts. *International Journal of Mass Spectrometry*, 2012, 309, pp.141-147. 10.1016/j.ijms.2011.09.014 . cea-03470721

HAL Id: cea-03470721

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The use of total evaporation method using Channeltron electron multipliers by thermal ionization mass spectrometry for europium isotope ratio measurements on picogram sample amounts

S. Mialle^a, A. Quémet^a, A. Ponvienne^a, A. Gourgiotis^a, M. Aubert^a, H. Isnard^{a,*}, F. Chartier^b

^a Commissariat à l'Energie Atomique, Saclay, DEN/DPC/SECR/LANIE, 91191 Gif-sur-Yvette Cedex, France

^b Commissariat à l'Energie Atomique, Saclay, DEN/DPC, France

ARTICLE INFO

Article history:

Received 19 April 2011

Received in revised form

15 September 2011

Accepted 15 September 2011

Available online 23 September 2011

Keywords:

Europium

TIMS

Channeltrons

Total evaporation

Isotope ratio

ABSTRACT

Thermal ionization mass spectrometry (TIMS) is a well established instrumental technique for providing accurate and precise isotope ratio measurements for elements with low first ionization potential. In the nuclear domain, particularly for the study of spent nuclear fuel sample solutions, the reduction of quantities is an important issue in order to decrease the analyst exposition to the radioactive samples. This paper presents experimental results obtained for picogram quantities of samples using electron multipliers (Channeltrons) implanted on the new generation of TIMS associated with the total evaporation (TE) acquisition method. TE method presents some advantages for the measurement of low quantities of samples in relation to the whole sample consumption and the integration of the whole signal. For this study, a Thermal Ionization Mass Spectrometer from IsotopX (Isoprobe-T) equipped with a multi-collector system including Faraday cups and four movable Channeltrons is used. This work is focused on lanthanide elements, and particularly europium, in relation to the importance of their characterization in the spent fuel to validate neutronic calculation codes. In a first approach, europium isotopic ratios by total evaporation method using Faraday cups are obtained for quantities down to 250 pg. The results obtained by TE (Faraday cups only) on such low quantities confirm the potentialities of the method. A reproducibility of 2.5‰ ($k=2$), was obtained on the $^{153}\text{Eu}/^{151}\text{Eu}$ ratio ($n=11$), for natural samples with an accuracy around 1‰.

In a second approach the same procedure is tested on Channeltron detectors by reducing the amounts of samples (from 50 pg to 1 pg). In order to obtain accurate isotope results, intrinsic parameters such as applied voltages, dead time and noise were determined. A Channeltron specific intercalibration gain method was also developed, as this parameter was found to be the major source of uncertainties using this type of detectors. The reproducibility on the $^{153}\text{Eu}/^{151}\text{Eu}$ ratio was respectively found around 3% ($k=2$). These results show the potentialities of isotopic measurements of sub-nanograms quantities and the different sources of uncertainties using total evaporation and Channeltrons on the Isoprobe-T.

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1. Introduction

Thermal ionization mass spectrometry (TIMS) is a recognized technique for the determination of lanthanides on a broad range of samples [1–7]. For several years, many advances have been made for isotopic determination of lanthanides to improve analytical precision for very low quantities of element (few nanograms and lower) [8–10]. In nuclear industry the reduction of quantities is an important issue in order to minimize the analyst's exposure to the radiation of the samples. Concerning TIMS, many studies have focused on the enhancement of the ionization efficiency. These

methods aim to reduce the oxide formation by understanding the associated mechanisms [11], improve beam focus and ion transmission by reducing the spatial extent of the sample load, or enhance the work function of the ionization filament (see for a review [12]). Instead of the measurement of lanthanide metallic ion, several authors use lanthanide single oxide ion measurements using O_2 or different oxide activators on filaments [8,9,13–15] increasing therefore the ionization efficiency. Another major development for low quantity measurements is the use of photomultipliers, hybrid detector like Daly electrode [16], or electron multipliers (EM). The latter can be divided in two categories: discrete dynodes EM and continuous dynode EM (channel electron multipliers or Channeltrons). Since the original idea suggested by Farnsworth [17] to create this type of detector and their conception [18,19], several channel electron multiplier characteristics have been improved.

* Corresponding author. Tel.: +33 1 69 08 80 79.

E-mail address: helene.isnard@cea.fr (H. Isnard).

Channeltrons, which were generally implanted in axial position, are now movable due to their miniaturized size on the new generation of TIMS [20–23]. In addition, data acquisition methods have been significantly improved. In order to achieve a suitable precision on isotope ratios [6], multistatic measurements using Faraday cup detectors associated with an internal normalization are traditionally used. Another method, called total evaporation (TE) [24], extensively used in the nuclear domain, was originally developed as an instrumental mass fractionation correction technique. The assumption made is that mass fractionation is minimized using total evaporation of the sample [4,10,25–29]. Another significant advantage of this technique is that very low amounts of samples are required [10].

The objective of this work is to study the potentialities and limitations of the TE method associated with Channeltron electron multipliers for low quantities europium isotopic ratio measurements. Europium was chosen in relation to its natural isotopic ratio close to 1 ($^{153}\text{Eu}/^{151}\text{Eu} = 1.0916 \pm 0.0019$) [30] and because of its interest for nuclear applications [31,32]. The first part of this paper presents the optimization of the Channeltron parameters. A detailed description of the settings for this type of detector and a method for dead time determination will be presented. Then, investigations on low sample amounts will be presented in two steps. First, a data set was collected using total evaporation method on Faraday cups in order to obtain reference values (sample amounts down to 250 pg). In a second step, the method was transposed on Channeltrons in static multicollection mode for picogram quantities of sample. Reproducibility, accuracy and major sources of uncertainties on such low amounts will be discussed.

2. Experimental

2.1. Reagents

For the optimization of Channeltron parameters, uranium NIST 500 standard solutions from the National Institute of Standard and Technology (NIST) starting for U_3O_8 powder dissolved in HNO_3 14 M are used. These solutions are re-concentrated at $2 \mu\text{g}/\mu\text{L}$ using sub-boiling HNO_3 0.5 M. This acid was obtained by dilution of sub-boiling 14 M nitric acid, using an EVAPOCLEAN system (Analab, France) and deionized water (resistivity $18.2 \text{ M}\Omega\text{cm}$) obtained from a Milli Q system (Millipore, Milford, MA, USA).

Europium solutions are prepared from a NIST SRM 3117a sealed borosilicate glass ampoule diluted in HNO_3 sub-boiling 0.5 M in order to obtain 1, 0.5 and $0.25 \text{ ng}/\mu\text{L}$ solutions for Faraday cup measurements and 50, 25, 10, 5 and $1 \text{ pg}/\mu\text{L}$ solutions for Channeltron measurements. The concentrations of these solutions were checked using an X series Inductively Coupled Plasma Quadrupole Mass Spectrometer (ICP-QMS) (Thermo Electron, Winsford, UK). For inter-Channeltron gain measurements, europium solution ($1 \mu\text{g}/\mu\text{L}$) in 2% HNO_3 from certified natural solutions (SPEX CertiPrep) is used.

2.2. Instrumentation

All the measurements were carried out on an Isoprobe-T, developed by IsotopX (Manchester, UK). This spectrometer is equipped with a glove box. A triple Re-filament configuration is used to control independently the evaporation and the ionization temperatures. This triple configuration is constituted of three filaments positioned in central, inside and outside positions. These filaments (thickness $30 \mu\text{m}$, width 0.7 mm), provided by ATES (Castelnaud Le Lez, France), are obtained from a 99.995% pure Re alloy. All filaments are beforehand degassed under a 10^{-7} mbar vacuum by warming up at different currents (respectively 1–3 A during 30 min

then 4 and 5 A during 1 h each). $1 \mu\text{L}$ of diluted solutions (described in Section 2.1) are deposited on the outside Re filament. During the deposit, the filament is heated up to 0.8 A. Then the current is progressively increased to 2 A during 10 s.

The analyzer is a low resolution magnetic field (resolution around 400). The multicollector is equipped with 9 movable Faraday cups, a fixed Daly electrode and 4 movable Channeltrons. Faraday cups are equipped with high-ohmic resistors ($10^{11} \Omega$). The amplifiers are maintained at a low pressure (10^{-2} mbar) and at a constant temperature (16°C) in a box, called Decabin, in order to reduce the Johnson noise. A WARP Filter (Wide Aperture Retarding Potential) is located before the electrode to improve abundance sensitivity [33] down to the ppb level. Four independently movable Channeltrons are located in low positions relative to the axial Faraday cup but only two of these (L4 and L5) were used in this study. The saturation level of these detectors is around 200,000 cps (counts per second). As these detectors are not equipped with any filters, abundance sensitivity was evaluated around 2 ppm as proportion of ^{238}U ion ion measured at mass 237, in accordance to literature data [34].

2.3. Channeltron optimization

2.3.1. Applied voltages

On this instrument three parameters have to be optimized for each Channeltron: multiplier voltage (MV), discriminator voltage (DV), and entrance voltage (EV). MV is the voltage applied between the input and the output of the detector; DV reduces the dark noise (when no signal is applied) and has also an influence on the measured intensities. EV corrects for asymmetrical peak shape due to the incidence angle of the ion beam with the Channeltron. There is a last common parameter: CT Suppressor (CT Sup), which is used to prevent both electrons escaping from Channeltrons and stray electrons entrance, has to be determined. As no significant influence on signal intensities was observed for different CT Sup values, according to manufacturer recommendations, CT Sup was fixed at -90 V for this study.

First, the multiplier voltage is optimized with DV set up to default values defined by the constructor (around 0.9 V). A rhenium filament is heated up in order to have an ^{187}Re signal around 100,000 cps on the Daly electrode. This ion beam is then measured on Channeltron and MV is increased gradually in order to reach the optimum signal intensity. Both dark noise and Channeltron response are measured for each MV value. Fig. 1a shows the Channeltron response with respect to MV, whereas no significant dark noise variation was measured. An intensity plateau is observed between 2.6 and 2.9 kV. In order to optimize the detector life MV has to be set above the “knee” [35] (Fig. 1a). The optimized voltage is 2.6 kV for L4 and 2.7 kV for L5. This voltage has to be readjusted with respect to the detector behaviour.

The optimum DV is then set up when maximal intensity and minimal noise are obtained. The noise is measured when the isolation valve is closed. Fig. 1b shows that the optimal signal to noise ratio is obtained for a DV value close to 0.7 V for the L5 detector. The same approach lead to an optimum DV value of 1.2 V for the L4 detector.

EV is therefore optimized by using an automatic *peak flat test* implemented on the instrument software. EV is optimized by comparison of signal intensities at the peak centre (PKC) and on the both sides of the peak centre. EV is then fixed when all intensities are in the range $0.995 < I/I_{\text{PKC}} < 1.005$. Following this procedure EV voltage is -17 V for L4 and -33 V for L5. As Channeltron peak shapes are dependent on the ion velocity and detector position, this set up has to be readjusted when a new element is analyzed or a new detector configuration is used.

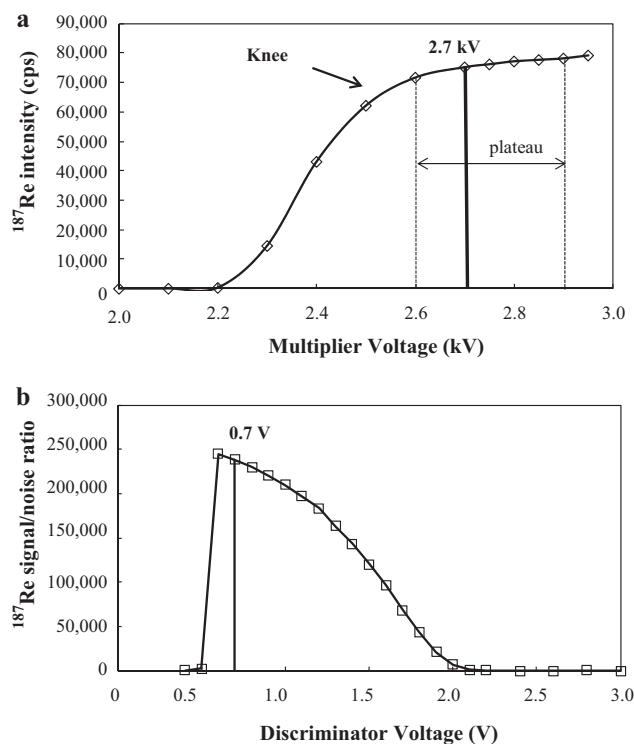


Fig. 1. Optimization of the multiplier voltage and discriminator voltage for the Channeltron L5. (a) ^{187}Re signal intensity from the filament expressed in count per second (cps) versus multiplier voltage (kV). The optimum value is obtained on the plateau (just above the “knee”) and (b) ^{187}Re signal to noise ratio versus discriminator voltage (V). The optimized value is determined when the maximum ratio signal/noise is obtained.

2.3.2. Dead time optimization

Dead time, the minimum time interval required to detect independently two ions arriving at an electron multiplier, can impact the observed ratio to a varying degree depending on the count rate [36–39]. The following formula is traditionally used to apply dead time correction [40,41]:

$$I_{\text{corrected}} = \frac{I_{\text{measured}}}{1 - \tau I_{\text{measured}}} \quad (1)$$

where τ is the dead time, $I_{\text{corrected}}$ the dead time corrected intensity and I_{measured} the measured signal intensity. An accurate and precise determination of the dead time prevents any signal losses and some non linearity effects [37,42].

In this study the dead time calculation is performed following the method described by Vanhaecke et al. [43]. Experimentally, uranium certified standard NIST 500 was chosen because it has two certified uranium isotope ratios. $^{235}\text{U}/^{238}\text{U}$ is close to unity ($^{235}\text{U}/^{238}\text{U} = 0.9997 \pm 0.0014$) and can be used for accurate mass-fractionation correction whereas $^{234}\text{U}/^{238}\text{U}$ ratio = 0.01042 ± 0.00002 can be used for dead-time evaluation. A specific acquisition method was used for this purpose: (1) Channeltron dead time is set initially at 0 ns; (2) ^{234}U is measured on the studied Channeltron and (3) ^{238}U and ^{235}U are measured on Faraday cups. Then, the ratio $^{234}\text{U}/^{238}\text{U}$ is determined at increasing ^{234}U signal intensities: 1×10^4 , 5×10^4 , 1×10^5 , 1.5×10^5 and 1.75×10^5 cps. Then, for each intensity rate on the Channeltron, relative isotope ratios were plotted as a function of the values used for the dead time correction leading to curves with a point of intersection as shown in Fig. 2a. The intersection of lines gives the experimentally determined dead time value. In our case, dead time was estimated at 150 ns for L4 and 290 ns for L5. Comparing to SEM or Daly electrode classical dead time values ($\tau = 10\text{--}50$ ns)

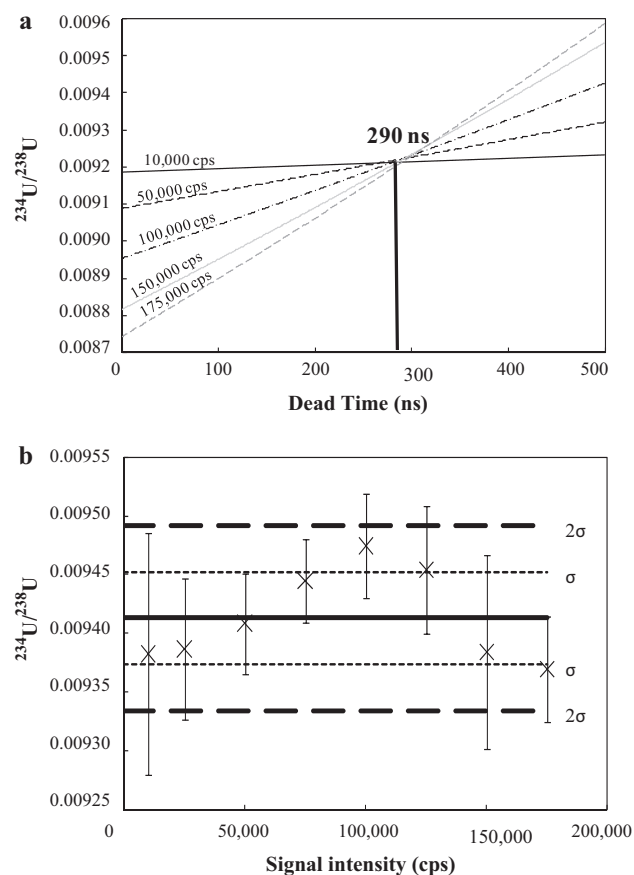


Fig. 2. Dead time and test of linearity for the Channeltron L5 performed on uranium NIST 500. (a) $^{234}\text{U}/^{238}\text{U}$ isotope ratio (given in atom ratio) versus dead time values for different signal intensities. The intersection of the plots gives the optimum dead time. (b) Example of linearity test performed on $^{234}\text{U}/^{238}\text{U}$ isotope atom ratio (no gain correction) at different intensities of the ion beam. The solid line represents the mean of these measurements and the thin and thick dashed lines represent respectively the mean $\pm \sigma$ and the mean $\pm 2\sigma$. It shows no discernible trend with the signal at the uncertainty of the measurements.

[35,44,45], the ones obtained for Channeltrons are much higher, which was already observed for this kind of detector [43,46].

Several linearity tests are performed for each Channeltron on uranium NIST 500 standards in order to validate the above determined dead time values [35]. The linearity of the detector is characterized by an invariance of the measured isotopic ratio whatever signal intensity collected. Experimentally, ^{238}U is measured on a Faraday cup and ^{234}U on a Channeltron. Measurements are performed, like for the dead time optimization, at different ion currents without any gain corrections. Fig. 2b shows an example of linearity test performed for a Channeltron. For both L4 and L5, the results demonstrate that the two detectors are linear from 5000 cps to 175,000 cps. For each linearity test, the average of eight measurements (from 5000 cps to 100,000 cps) is given with a reproducibility <1%.

2.3.3. Noise and efficiency

Noise measurements were performed, with closed isolation valve, on both detectors (L4 and L5). Each measurement takes around 20 min and the integration time is fixed at 5 s for a sequence. A noise level of about 0.2 cps (counts per second) for L4 and 0.3 cps for L5 was observed.

Finally, efficiencies of Channeltrons compared to were estimated at around 95% for europium standard and around 70% for Uranium. These measurements performed by comparing a similar ion beam on both detectors (close to 1×10^5 cps) show that the

efficiency of this kind of detectors decreases when the ratio m/z increases.

2.4. Total evaporation methods

The “total evaporation” method, described first by Romkowski et al. [24] is based on the collection and integration of the overall ion beams generated by the total evaporation of a small sample amount in a short time. This method was initially developed in order to circumvent mass fractionation induced by thermal ionization sources [4,10,25–27].

A total evaporation analysis can be divided in 3 parts: adjustment, acquisition and shutdown phases [4]. For each step, some parameters are changed following the sample amount or the kind of detector. In the next part, the acquisition parameters used in the case of multi-collection mode with Faraday cups and Channeltrons is described. The advantage of this collection mode is to take account the ion beam fluctuations.

2.4.1. Multi-static Faraday cup measurements

First, Faraday cups are moved in order to collect the europium isotopes and intercalibration gains are measured daily by an automatic process. This is carried out using a highly stable current which is successively applied to each amplifier. Reproducibility of electronic gains obtained by this procedure was better than 0.001%. The adjustment phase is the same for all the measurements done on Faraday cups whatever the quantity of sample deposited. During this step, filaments are gradually pre-heated at 4.5 A for the centre and 0.7 A for the side in 10 min. The ionization filament temperature is then adjusted for a 80 mV signal on the ^{187}Re ion beam. At this stage, peak centre and lens focus are performed and the centre filament temperature is kept constant during the overall analysis. Then, the side temperature is increased to reach a 5 mV signal on the ^{153}Eu ion beam. This ion current is the minimum value required to perform a peak centre. No focus is performed on the ^{153}Eu to avoid further signal loss.

The acquisition phase begins with a peak centre measurement. Baselines are measured at ± 0.5 u with a 5 s integration time. Then several parameters are modified in relation to the quantity of sample deposited on the filament. The aiming signal intensity (maximal instantaneous collected ion intensity) is set by the operator. The selected values are 1 V on the major isotope for samples ranging from 50 ng to 100 ng and 0.050 V for 1 ng and lower sample amounts (called in the following “low quantity sample”). Each isotope is collected with an integration time of 10 s for 100–50 ng samples and 5 s for low quantity samples. Continuous monitoring of the ion current decay rate governs the side filament temperature steps. The maximum side filament current is set at 4 A whatever the sample analyzed. The termination signal condition, which is the ion current intensity for the major isotope below which the analysis is stopped, is set at 5 mV for all samples. Then, the shutdown phase, during which filament currents are set at 0 A. Gain and baseline corrections are performed for each integration. Then the isotopic ratio is calculated on the basis of the integrated ion current for each isotope.

2.4.2. Multi-static Channeltron measurements

The acquisition procedure described above has to be adapted for the multi-static Channeltron measurements. The modifications applied are described below.

2.4.2.1. Intercalibration gain method. Channeltrons are moved in order to collect europium ion beams but, in opposition to Faradays cups, no automatic procedure is available to evaluate the relative detector response. A procedure was then developed for Channeltron gain calibration. In a first step, the ionization filament

Table 1

Successive sequences used to measure intercalibration gain on Channeltrons.

Channeltron	L4	L5
Sequence 1	^{151}Eu	–
Sequence 2	^{153}Eu	^{151}Eu
Sequence 3	–	^{153}Eu

is gradually heated to 4.5 A. Successive peak centre, lens focus, and ionization filament temperature adjustments are performed in order to obtain a 80 mV signal on the ^{187}Re ion beam measured on Faraday cup. In a second step, the side filament temperature is adjusted to obtain a ^{153}Eu signal of 100,000 cps on Channeltron. Peak centre, lens focus and baselines (± 0.5 u) are performed before gain acquisition (5 s integration time). Data are acquired in multi-dynamic collection mode with three measurements sequences (see Table 1). The procedure includes 4 blocks of 10 cycles. Gains are calculated by sequential ^{151}Eu and ^{153}Eu signal measurements on Channeltron following Eq. (2):

$$G_{Lx/Ly}(i) = \frac{({}^i\text{Eu})_{Lx}}{({}^i\text{Eu})_{Ly}} \quad (2)$$

where i stands for the considered europium isotope, x and y stand for the considered Channeltrons. This procedure is applied before and after each analysis. Gain values and reproducibilities will be discussed in Section 3.

2.4.2.2. Total evaporation using Channeltrons. In order to avoid pre-emitting signal on Eu no current was applied on side filaments while the centre filament is warmed up at 4.5 A. The ^{187}Re ion beam is still optimized and focused at 80 mV on a Faraday cup. The side filaments are then heated to reach a ^{153}Eu ion beam of 5000 cps on Channeltron. Peak centre and baseline measurements are performed before the beginning of the acquisition phase. During this step ion beams are collected (5 s integration time) on both Channeltrons (^{151}Eu on L5 and ^{153}Eu on L4) with an aiming intensity sets at 100,000 cps. The maximum side filament current is set at 4 A. The termination intensity is fixed at 5000 cps. The shutdown phase is already described in Section 2.4.1.

3. Results and discussion

3.1. Multi-static Faraday cup measurements

Natural europium isotopic measurements have been performed by TE method during several years in our Laboratory (50–100 ng deposits, $n=34$). For 26 measurements the cumulated ^{153}Eu signals are included between 50 and 200 V. Four measurements have a cumulated ^{153}Eu signal <50 V and four values are above 200 V. Two values were rejected after performing a statistical test on the $^{153}\text{Eu}/^{151}\text{Eu}$ measured ratios. The average value obtained on these 32 measurements is 1.0927 ± 0.0011 ($k=2$) which is consistent with IUPAC reference value 1.0916 ± 0.0019 for natural europium [1,30] within uncertainties. During this work, ten measurements were performed with a 1 ng deposit with cumulated ^{153}Eu signals ranging from 0.3 to 20 V (1 rejection). Six measurements were performed with a 0.5 ng deposit and the collected ^{153}Eu signals ranging from 0.2 and 7 V (no rejection) for these analyses. Eleven measurements were performed with a 0.25 ng deposit with cumulated ^{153}Eu signals ranging from 0.2 to 1.5 V (1 rejection). The average $^{153}\text{Eu}/^{151}\text{Eu}$ values obtained by TE for lower quantities: 1 ng, 0.5 ng and 0.25 ng are respectively equal to 1.0928 ± 0.0011 ($k=2$, $n=9$), 1.0931 ± 0.0005 ($k=2$, $n=6$) and 1.0939 ± 0.0027 ($k=2$, $n=10$). The results obtained are all consistent with the laboratory reference value within uncertainty (Fig. 3a). As expected, a higher reproducibility is observed for the lowest amounts of sample (0.25 ng).

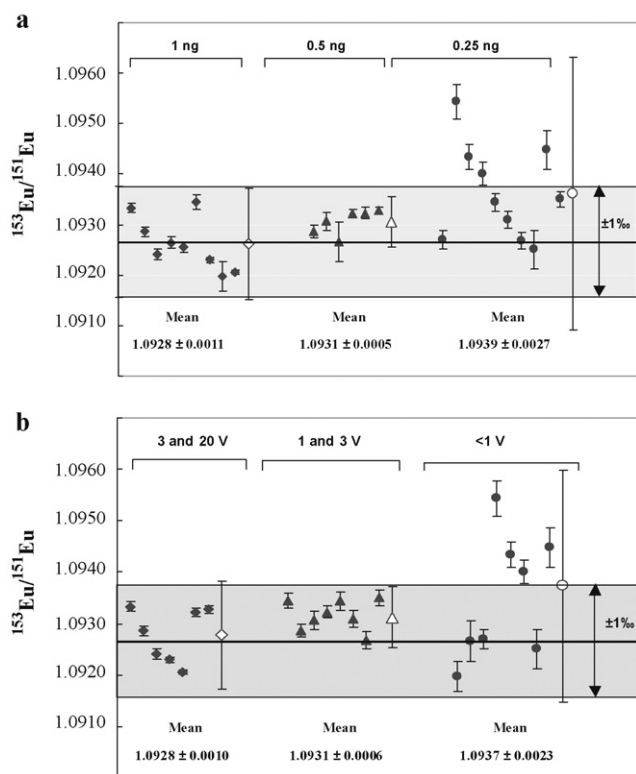


Fig. 3. Results of TE method for $^{151}\text{Eu}/^{153}\text{Eu}$ ratio (given in atom ratio) for samples of 250 pg to 50 ng of europium NIST standard solution using static multicollection mode with Faraday cup detectors. Each data point is represented with an uncertainty applying the counting statistic $\sigma_N = \sqrt{N}/N$, where N is the number of ions. Dash area corresponds to the reference value obtained for quantity of 50 ng ($^{151}\text{Eu}/^{153}\text{Eu} = 1.0927 \pm 0.0011$). (a) The data are represented based on the mass of deposit on the filament. The mean values of each data series are indicated with their uncertainties expressed as the standard deviation at a confident level of $k=2$. (b) The data are represented based on total signal collected during the analysis. The mean values of each data series are indicated with their uncertainty expressed as the standard deviation at a confident level of $k=2$.

Fig. 3b shows the $^{153}\text{Eu}/^{151}\text{Eu}$ ratio versus cumulated ^{153}Eu signal. We observed a significant increase of the standard deviation when cumulated ^{153}Eu signal decreases. As observed in Fig. 3b, a cumulated ^{153}Eu signal close to 1 V is required in order to obtain a reproducibility ($k=2$) of 1%.

3.2. Multi-static Channeltron measurements

3.2.1. Intercalibration gains and uncertainty

As described in Section 2.4.2, Channeltron–Channeltron intercalibration gain must be carefully determined in order to obtain accurate isotope ratio. Several tests were conducted to evaluate the intercalibration gain dependence with respect to the atomic mass of the studied element. Several elements (K, Mo, Eu, Re, U) were tested for this purpose. A clear decrease of the L4/L5 gain is observed with increasing m/z . This phenomenon is ascribed to detection efficiency dependence with respect to ion mass, incidence angle, ion charge and velocity [47]. However, no significant variation linked to this phenomenon was observed for the two europium masses (151 u and 153 u, $\Delta M=2$ u) and the average of gains calculated on each isotope is determined.

As indicated in Section 2.4.2, average gain (i.e. average gain value before and after analysis) is used for measurement corrections in order to account for time variations (following Eq. (3)):

$$G_{Lx/Ly} = \frac{(G_{Lx/Ly})_{t1} + (G_{Lx/Ly})_{t2}}{2} \quad (3)$$

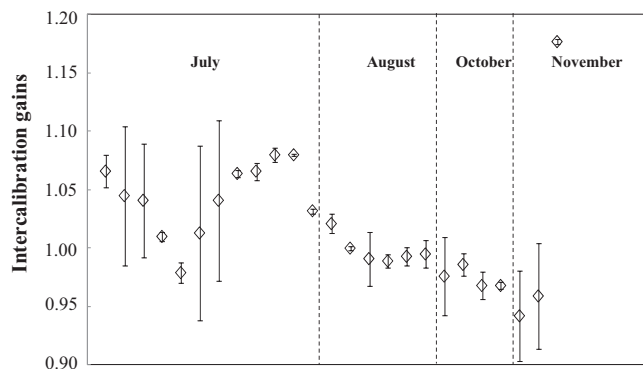


Fig. 4. Evolution of the gain values during the examined time period, July through November. Each interval included between the two vertical lines represents one day. Each gain is indicated with its uncertainty expressed as standard deviation at a confident level of $k=1$. The uncertainties are evaluated by the relative difference of the gain value before and after measurements of the sample.

where $G_{Lx/Ly}$ stands for average gain, $(G_{Lx/Ly})_{t1}$ is the gain before the analysis and $(G_{Lx/Ly})_{t2}$ is the gain after analyses.

Gain repeatability ranges between 0.03 and 0.2%. As shown in Fig. 4, where average gain variations obtained for 25 analyses performed during 5 months are reported, long term gain reproducibility ($n=25$) was found to be close to 5% ($k=1$). To take into account this reproducibility, we evaluated the relative gain difference before and after measurement following Eq. (4):

$$\Delta G = \frac{(G_{Lx/Ly})_{t1} - (G_{Lx/Ly})_{t2}}{G_{Lx/Ly}} \quad (4)$$

Final uncertainty on intercalibration gain was evaluated by combination of this relative gain difference and repeatability.

3.2.2. Total evaporation measurements

Several measurements using TE method with europium quantities varying from 100 pg to 1 pg and Channeltron detection were performed. 100 pg measurements were stopped due to a long analysis time (>1 h). By consequence only measurements ranging from 50 pg to 1 pg were really studied. The same acquisition method was used whatever the quantity deposited on the side filament.

The profiles observed by TE using Channeltron detector are comparable to those obtained using Faraday cup with the exception of very low quantity (1–10 pg) profiles which are very short transient signals. A pre-emitting ion beam (i.e. a short transient europium signal) in the beginning of peak integration was also observed for the large majority of data although side filaments are not heated up. This could be explained by the fact that the centre filament alone produces enough energy to partially evaporate and ionise europium on the side filament.

Table 2 reports the results obtained for each deposited quantity. The data are ordered in relation to the cumulated ^{153}Eu count rate ($>10^7$ cps, between 1 and 10^7 cps and $<10^6$ cps). Intercalibration gain and associated uncertainty are also reported together with raw and corrected $^{153}\text{Eu}/^{151}\text{Eu}$ ratio. Uncertainty on raw ratios is evaluated by applying the counting statistic \sqrt{N}/N , where N is the number of ions [10]. The corrected $^{153}\text{Eu}/^{151}\text{Eu}$ ratio uncertainty is expressed taking into account the combination of raw ratios and gain uncertainty (see Section 3.2.1). The $^{153}\text{Eu}/^{151}\text{Eu}$ ratio average value and relative standard deviation (RSD) (varying from 0.85% to 1.85%) were estimated for each cumulated count rate range (Fig. 5). Uncertainty calculation shows that the gain uncertainty (% range) budget is the major source of uncertainty for static collection mode measurements using Channeltrons. The average value obtained for 25 measurements on the $^{153}\text{Eu}/^{151}\text{Eu}$ ratio = 1.0920 (3.3%, $k=2$). This value is in good agreement (accuracy ranging from 0.5% to

Table 2
Results obtained with total evaporation method for different deposited amounts by 2 channel electron multipliers and classified by collected ^{153}Eu signal (respectively $>10^7$ cps, 10^6 – 10^7 cps, $<10^6$ cps). For each measurement the intercalibration gain is given with its associated uncertainty at a confident level of $k=1$. Raw measurements are given with their uncertainties at $k=1$ and corrected ratios with their uncertainties at a confident level of $k=1$ which are a combination of gain uncertainties and raw data uncertainties. Their means are given with the standard deviation at a confident level of $k=2$.

Deposit (pg)	Cumulated count rate ^{153}Eu (cps)	Gain	Uncertainty % ($k=1$)	$^{153}\text{Eu}/^{151}\text{Eu}$ raw	Uncertainty % ($k=1$)	$^{153}\text{Eu}/^{151}\text{Eu}$ corr.	Uncertainty % ($k=1$)
50	1.3E+07	1.07	1.3	1.1595	0.025	1.0881	1.3
50	1.0E+07	0.94	4.1	1.0067	0.029	1.0692	4.1
50	9.8E+06	0.96	0.5	1.0408	0.029	1.0853	0.5
25	2.2E+07	1.04	4.7	1.1384	0.020	1.0937	4.7
25	6.3E+07	1.04	6.6	1.1537	0.011	1.1087	6.6
10	3.7E+07	1.01	7.4	1.0937	0.015	1.0795	7.4
	Mean			1.0988		1.0874	
	2SD			0.1269		0.0267	
	2RSD %			12		2.5	
	Accuracy (%)					-0.5	
50	2.0E+06	1.08	0.6	1.1896	0.064	1.1019	0.6
50	3.9E+06	1.08	0.1	1.1968	0.046	1.1085	0.2
25	4.2E+06	1.05	5.7	1.1555	0.044	1.1060	5.7
25	6.6E+06	1.06	0.3	1.1804	0.035	1.1091	0.3
25	4.0E+06	1.07	0.7	1.1789	0.045	1.1061	0.7
10	4.9E+06	0.99	0.6	1.0934	0.041	1.1060	0.6
5	2.0E+06	0.99	2.3	1.0982	0.065	1.1086	2.3
5	2.7E+06	0.98	3.4	1.0627	0.056	1.0894	3.4
1	5.7E+06	1.03	0.2	1.1530	0.037	1.1175	0.2
	Mean			1.1278		1.1059	
	2SD			0.0992		0.0184	
	2RSD %			8.8		1.7	
	Accuracy (%)					1.2	
10	6.9E+05	1.01	0.4	1.0634	0.11	1.0532	0.4
10	9.9E+05	0.99	0.8	1.0681	0.09	1.0753	0.8
10	5.9E+03	1.18	0.2	1.2905	1.18	1.0960	1.2
5	9.3E+05	1.00	0.2	1.1092	0.09	1.1093	0.2
5	8.9E+05	1.00	1.2	1.0850	0.10	1.0905	1.2
5	5.2E+05	1.02	0.8	1.0850	0.13	1.0629	0.8
1	2.4E+05	0.98	0.9	1.0718	0.19	1.0943	1.0
1	7.3E+04	0.99	1.0	1.0378	0.35	1.0525	1.1
1	1.1E+05	0.97	1.2	1.0622	0.27	1.0973	1.2
1	2.6E+04	0.97	0.3	1.0568	0.57	1.0919	0.6
	Mean			1.0930		1.0823	
	2SD			0.1440		0.0400	
	2RSD %			13.2		3.7	
	Accuracy (%)					-1.0	
	Mean, $n=25$					1.0920	
	2SD					0.0359	
	2RSD %					3.3	
	Accuracy (%)					-0.06	

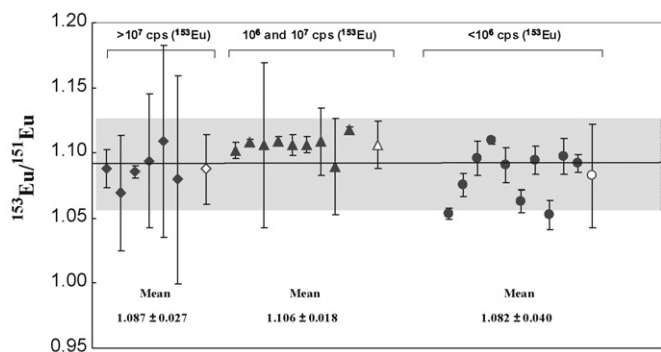


Fig. 5. Results of TE method and multistatic Channeltron detection for the $^{153}\text{Eu}/^{151}\text{Eu}$ ratio (given in atom ratio) with masses ranging from 1 pg to 50 pg. The results are classified in three categories according to the collected signal on the ^{153}Eu isotope. Each value is given with its uncertainty which is a combination of the gain uncertainty and counting uncertainty based on Poisson statistic. Each white point represents the mean for each category with its uncertainty at a confident level of $k=2$. The solid line represents the reference value of the laboratory at 50 ng and the gray area its uncertainty level.

1.2% depending of the collected signal) with TE Faraday cup measurements within uncertainty.

So, Channeltrons allow the measurements of picogram size samples dividing by 1000 and more the traditional quantity (50 ng) deposited on the filament in comparison to measurements using Faraday cups. A potential application of this type of detector is to measure the Eu and lanthanides isotopic composition in irradiated fuel samples of limited sample size. For example in the case of lanthanide transmutation target [48] irradiated in French fast neutron reactor, the quantity of products formed are very low (ng to pg quantities). The TE method on channeltron electron multipliers may hold potential for applications in this research field. Nevertheless this type of detector involves an uncertainty increase due to (1) higher uncertainty of the ion counting statistic and (2) intercalibration gain reproducibility which is not better than % level. However in several applications, Channeltrons could be used on peak jumping mode. For comparison at 100,000 cps (which are performed on L_4 detector) repeatability for europium measurements in peak jumping mode are around 0.2% (50 cycles, $k=1$) and reproducibility around 1.4% ($n=5$, $k=2$).

4. Conclusions

This study presents the optimization of Channeltron detectors implanted on an Isoprobe-T (IsotopX, UK) Thermal Ionization Mass Spectrometer. All Channeltron parameters are studied and for each detector noise, efficiency and dead time are evaluated. This type of detector is used to obtain europium isotopic ratio measurements in static multicollector mode using TE. The average value on the $^{153}\text{Eu}/^{151}\text{Eu}$ ratio for 25 measurements with quantity deposit ranging from 1 to 50 pg is equal at 1.092 ± 0.036 (SD at $k=2$) in agreement with reference value obtained on Faraday/Faraday static measurements ($^{153}\text{Eu}/^{151}\text{Eu} = 1.0927 \pm 0.0011$). The major source of uncertainty for this type of measurements is related to the intercalibration gain uncertainty which vary between 0.1% and 7% during all the analytical session.

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