

Production of ^{265}Sg for chemical studies using the gas-jet transport system coupled to the RIKEN gas-filled recoil ion separator

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1. Introduction

RIKEN GARIS as a pre-separator for SHE chemistry

Breakthroughs in SHE chemistry

- Chemical and physical experiments under low background condition
- Stable and high gas-jet transport efficiency
- New chemical systems that were not accessible before

Development of a gas-jet transport system coupled to GARIS

[JNRS **8**, 55 (2007).; EPJD **45**, 81 (2007).; JNRS **9**, 27 (2008).]

→ Production of $^{261}\text{Rf}^{a,b}$ in the $^{248}\text{Cm}(^{18}\text{O},5n)^{261}\text{Rf}^{a,b}$ reaction

[Chem. Lett. **38**, 426 (2009).; PRC **83**, 034602 (2011).]

This work

Aqueous chemistry of Sg at RIKEN

– based on successful collaboration on aqueous chemistry of Rf and Db at JAEA –

- Production of ^{265}Sg and its homologues ^{173}W and ^{90}Mo



- Decay properties of $^{265}\text{Sg}^{a,b}$

Previous decay studies of ^{265}Sg

Düllmann and Türler: PRC **77**, 064320 (2008).

$^{248}\text{Cm}(^{22}\text{Ne},5;4n)^{265,266}\text{Sg}$: reanalysis \rightarrow 36 events on $^{265}\text{Sg}^{a,b}$

$^{208}\text{Pb}(^{70}\text{Zn},n)^{277}\text{Cn}$: SHIP/GARIS + FPD \rightarrow 4 events

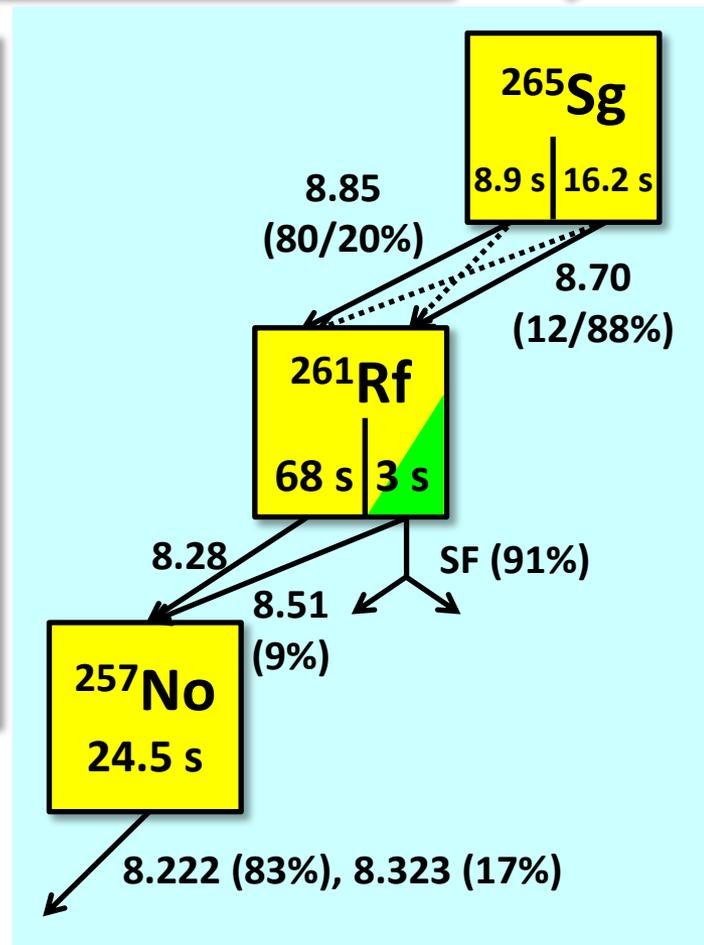
$^{248}\text{Cm}(^{26}\text{Mg},5n)^{269}\text{Hs}$: Gas-jet + COLD/CALLISTO/COMPACT \rightarrow 20 events



Refs. $^{248}\text{Cm}(^{22}\text{Ne},5n)^{265}\text{Sg}$	Method	E_{beam}	No of. events	σ [pb]
Lazarev <i>et al.</i> (1994)	DGFRS	116	4 ^{a)}	80
		121	6 ^{a)}	320
Gregorich <i>et al.</i> (1996)	MG	121	3	
Türler <i>et al.</i> (1998,1999)	OLGA	121/123	19	206
Türler <i>et al.</i> (1998)	PSI Tape	119	1 ^{b)}	78
Dressler <i>et al.</i> (2000)	PSI Tape	116	1 ^{b)}	73
Hübener <i>et al.</i> (2001)	HITGAS	119	2 ^{b)}	92

a) No $T_{1/2}$ data

b) Only sensitive to α -SF chains



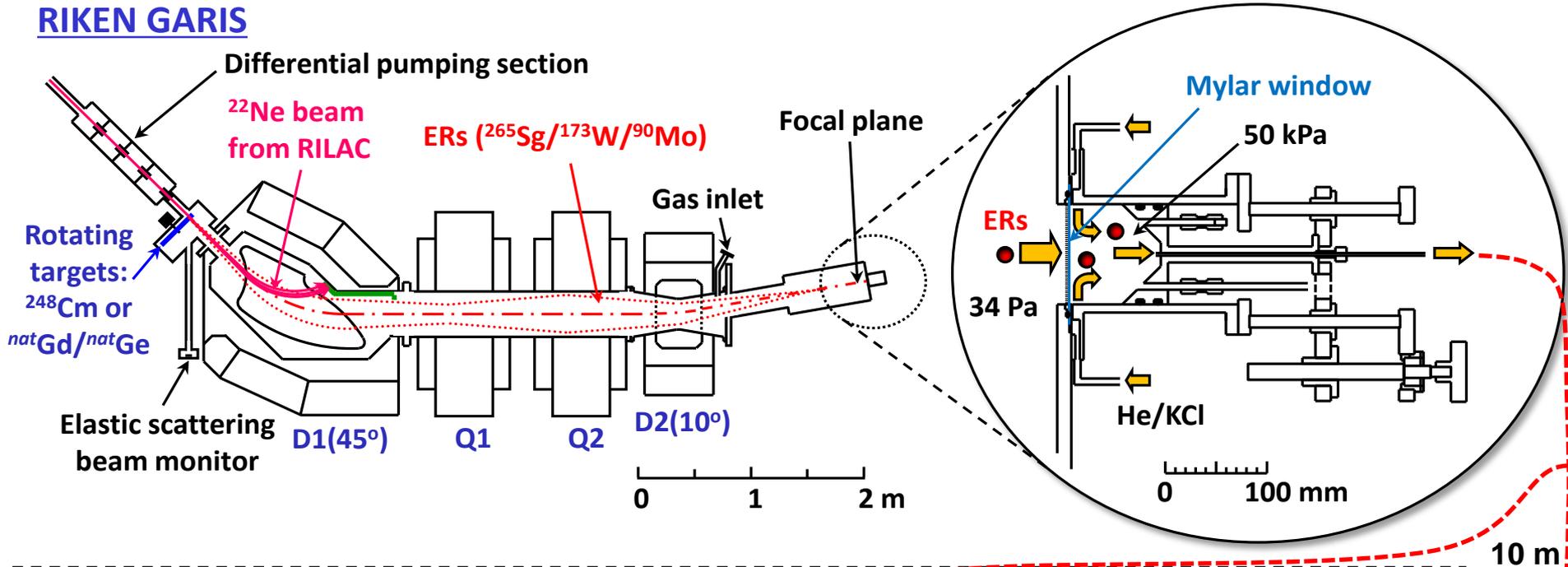
For future chemical studies with ^{265}Sg

Decay properties (E_{α} and $T_{1/2}$) and cross section ?

\rightarrow Systematic studies on $^{248}\text{Cm}(^{22}\text{Ne},5n)^{265}\text{Sg}^{a,b}$

2. Experimental

Experimental setup



Chemistry laboratory

Direct catch: $^{173}\text{W}/^{90}\text{Mo}$

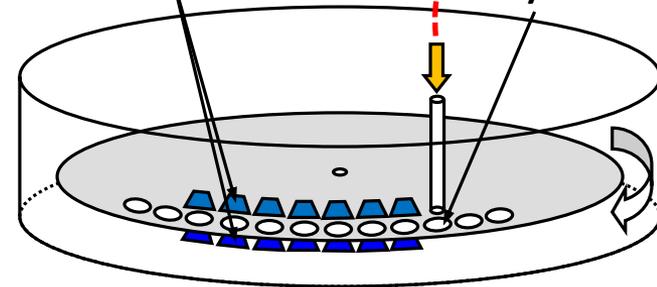
Glass filter
GB-100R

→ Y-spectrometry
with Ge detector

MANON for α spectrometry: ^{265}Sg

Si PIN photodiodes

Mylar foil



Experimental conditions

Nuclide	$^{265}\text{Sg}^{a,b}$ (Z=106)	^{173}W (Z=74)	^{90}Mo (Z=42)
Half-life	8.9, 16.2 s [*]	7.6 min	5.67 h
Reaction	$^{248}\text{Cm}(^{22}\text{Ne},5n)$	$^{nat}\text{Gd}(^{22}\text{Ne},xn)$	$^{nat}\text{Ge}(^{22}\text{Ne},xn)$
Beam energy (MeV)	118	←	←
Beam intensity (pμA)	4	←	0.25
Target (μg/cm ²) on 2-μm Ti	280/230 (Cm ₂ O ₃)	340 (Gd ₂ O ₃)	290 (Ge)
Recoil energy (MeV)	9.4	14.1	25.8
Magnetic rigidity (Tm)	1.73–2.16	1.50–1.93	0.985
GARIS He (Pa)	34	←	←
RTC Mylar window (μm)	0.65	←	←
Honeycomb grid (%)	84	←	←
Gas-jet He (kPa)	50	←	68
Chamber depth (mm)	40	100	100
He flow rate (L/min)	2.0/2.5	2.0	4.0
KCl generator (°C)	600/605	620	620
Aerosol collection (s)	20	120	300
α- or γ-spectrometry	MANON	Ge detector	Ge detector

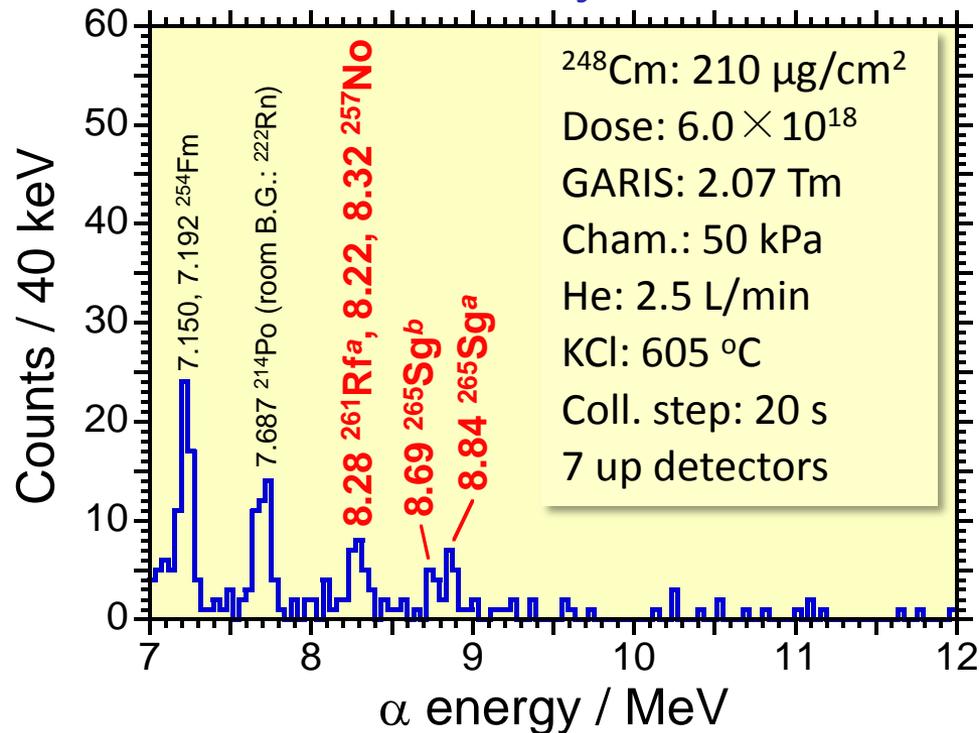
* Düllmann and Türler: PRC **77**, 064320 (2008).

3. Results and discussion

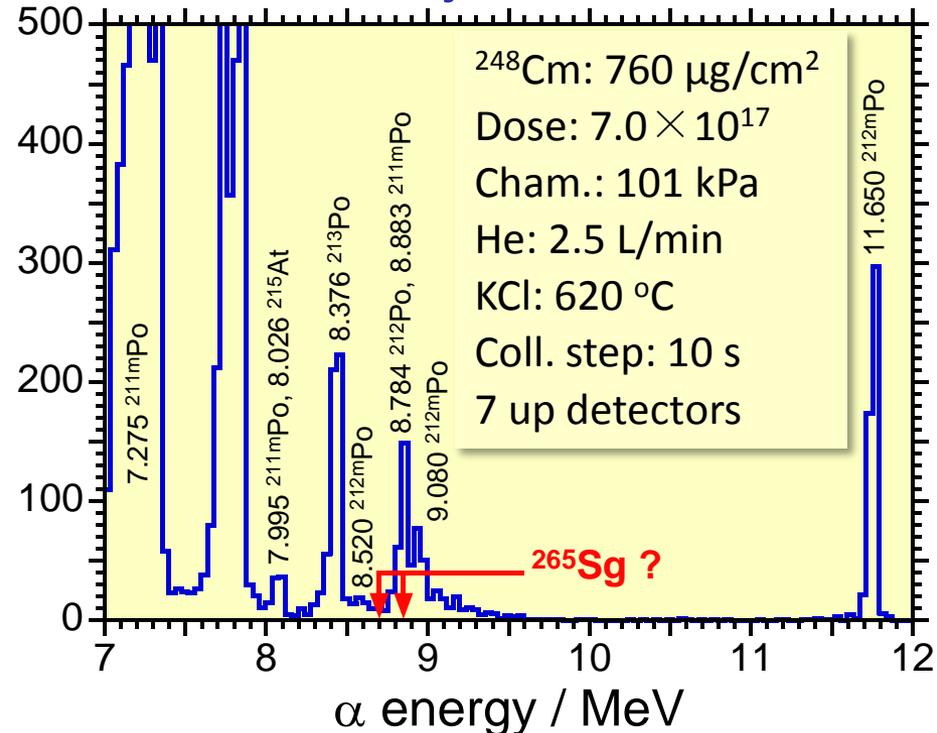
3.1. $^{248}\text{Cm}(^{22}\text{Ne},5n)^{265}\text{Sg}$

Beam time	Magnetic rigidity (Tm)	Beam dose ($\times 10^{18}$)
Sep. 30–Oct. 6, 2008	1.73	2.07
	1.94	1.91
	2.16	1.57
	2.04	0.639
Sep. 19–23, 2009; July 16–20, 2010	2.07	11.2

RILAC + GARIS + Gas-jet + MANON



AVF + Gas-jet + MANON



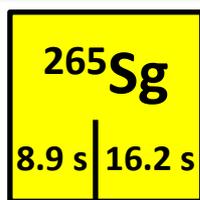
Search for α - α / α -SF correlations

$E_\alpha = 8.0\text{--}9.0$ MeV; $E_{\text{SF}} \geq 30$ MeV

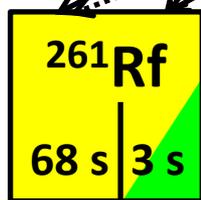
$\Delta T = 226$ s

	Observed	Random
α - α	98	< 5.3
α - α - α	18	< 0.1
α -SF	18	< 2.0

Düllmann and Türlér (2008)



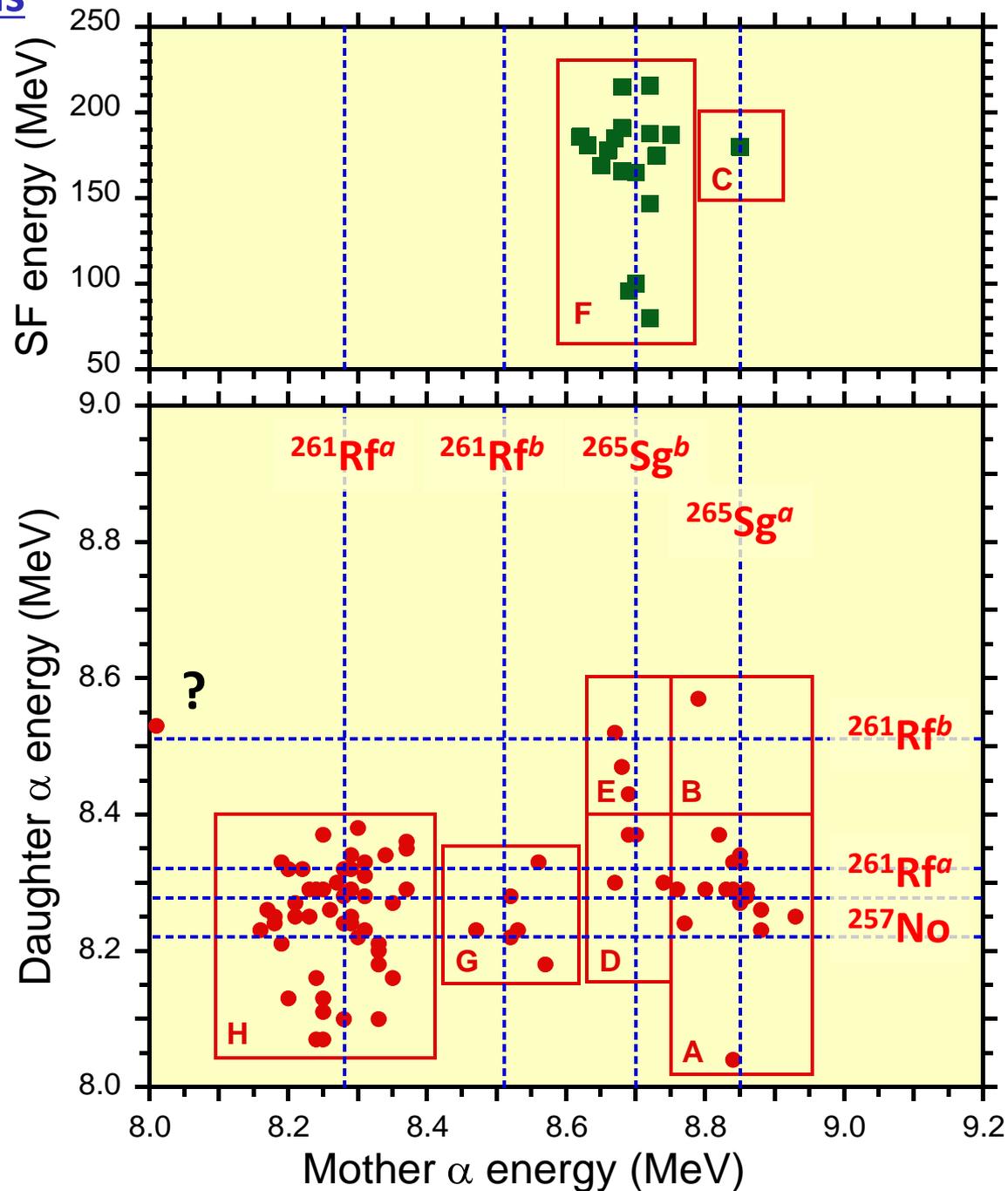
8.85 (80/20%)
8.70 (12/88%)



8.28
8.51 (9%)
SF (91%)



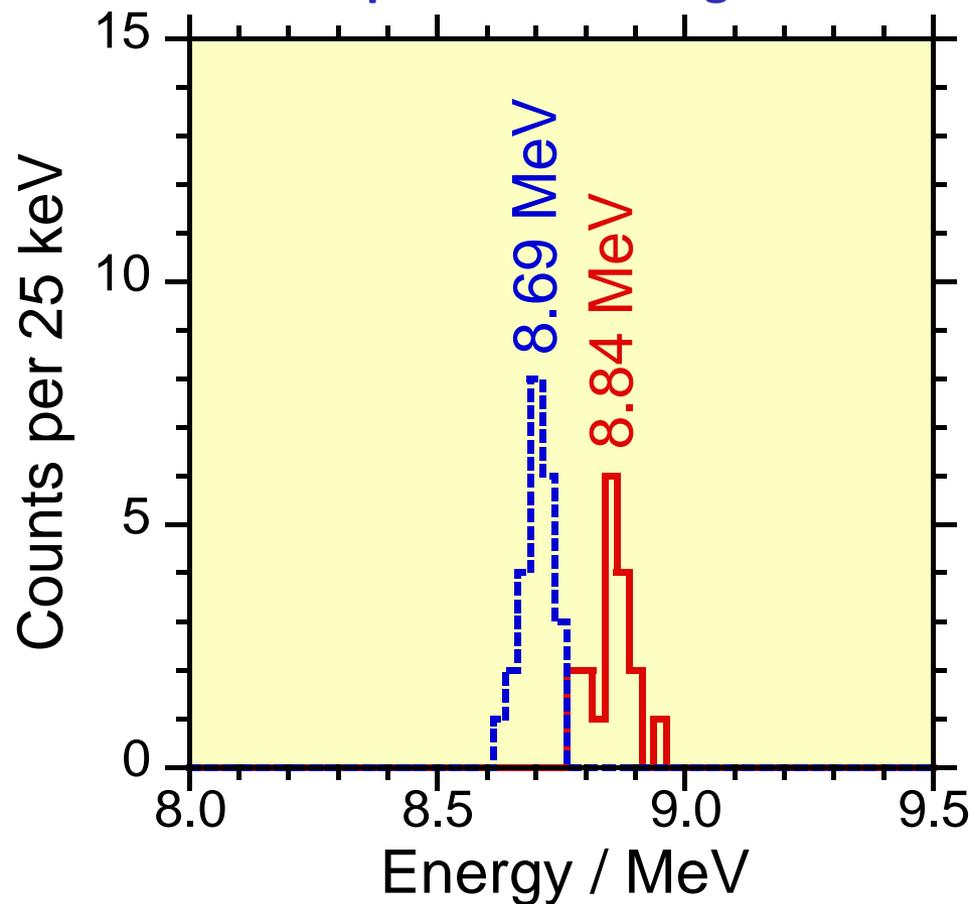
8.222 (83%), 8.323 (17%)



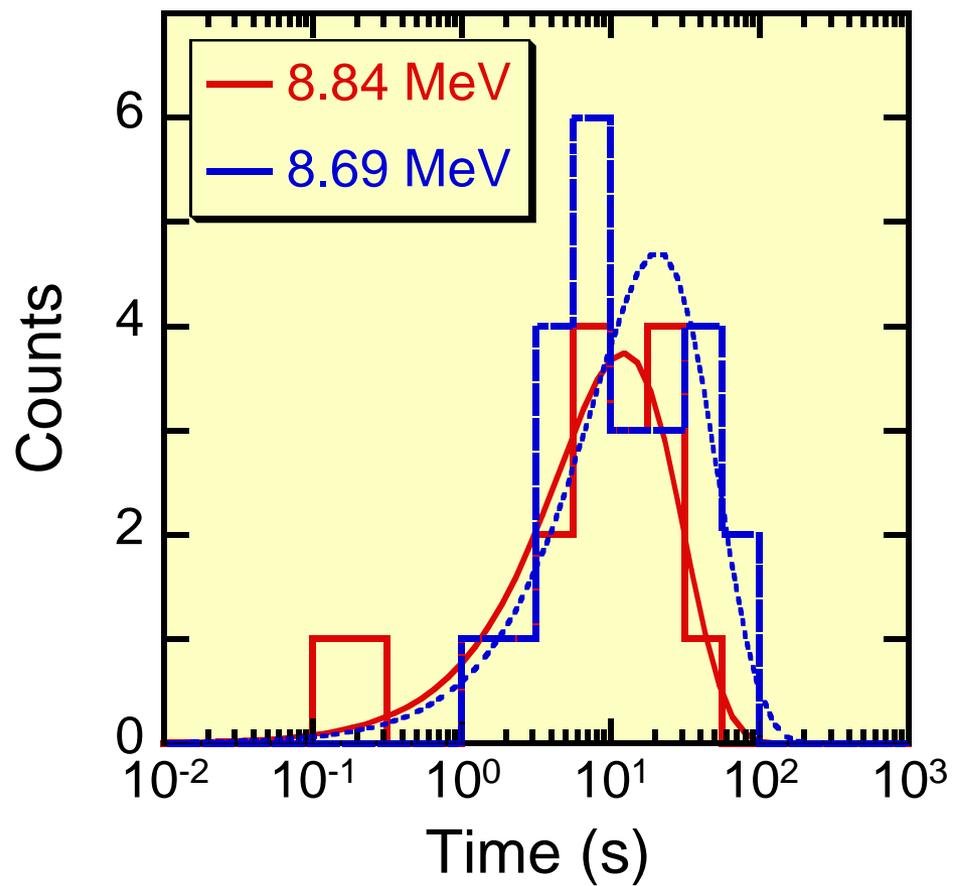
α energy and half-life of $^{265}\text{Sg}^{a,b}$

This work					Düllmann and Türler (2008)		
	n	E_α [MeV]	$T_{1/2}$ [s]	b_{SF} [%]	n	E_α [MeV]	$T_{1/2}$ [s]
$^{265}\text{Sg}^a$	18	8.84 ± 0.05	$8.5^{+2.6}_{-1.6}$	< 50	20	8.85	$8.9^{+2.7}_{-1.3}$
$^{265}\text{Sg}^b$	24	8.69 ± 0.05	$14.4^{+3.7}_{-2.5}$	< 51	24	8.70	$16.2^{+4.7}_{-1.9}$

α spectrum of $^{265}\text{Sg}^{a,b}$

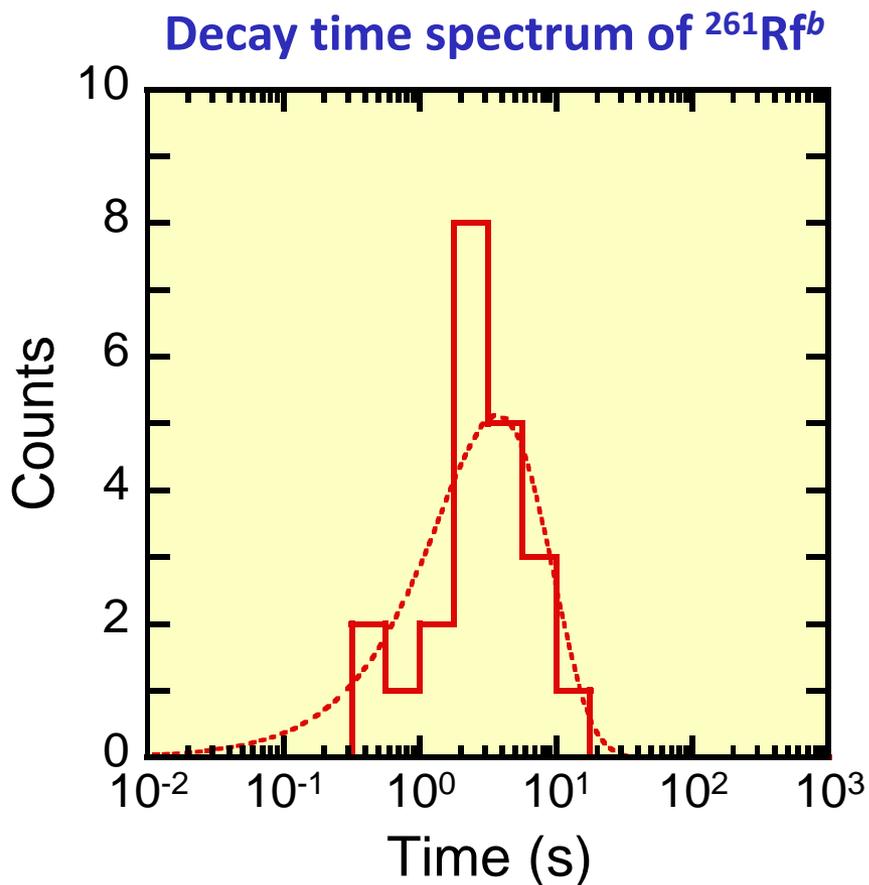


Decay time spectrum of $^{265}\text{Sg}^{a,b}$

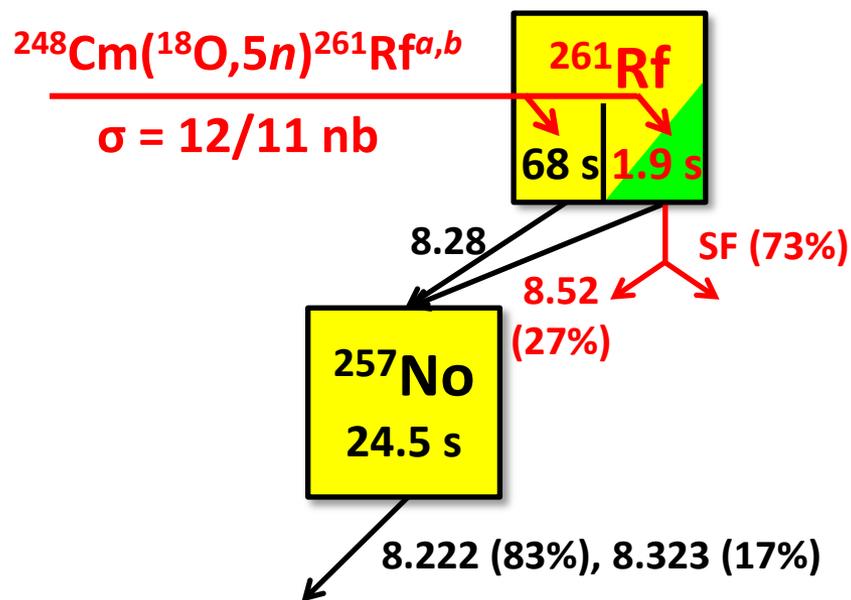


α energy and half-life of $^{261}\text{Rf}^{a,b}$ and ^{257}No

	This work				References			
	n	E_α [MeV]	$T_{1/2}$ [s]	b_{SF} [%]	n	E_α [MeV]	$T_{1/2}$ [s]	b_{SF} [%]
$^{261}\text{Rf}^a$	48	8.27 ± 0.06	(33^{+12}_{-7})		-	8.28 ± 0.02^a	68 ± 3^b	$< 10^a$
$^{261}\text{Rf}^b$	25	8.51 ± 0.06	$2.6^{+0.7}_{-0.5}$	82 ± 9	88	8.52 ± 0.05^d	1.9 ± 0.4^d	73 ± 6^d
^{257}No	54	$8.07 - 8.38$	23^{+4}_{-3}		-	$8.222, 8.323^c$	24.5 ± 0.5^c	



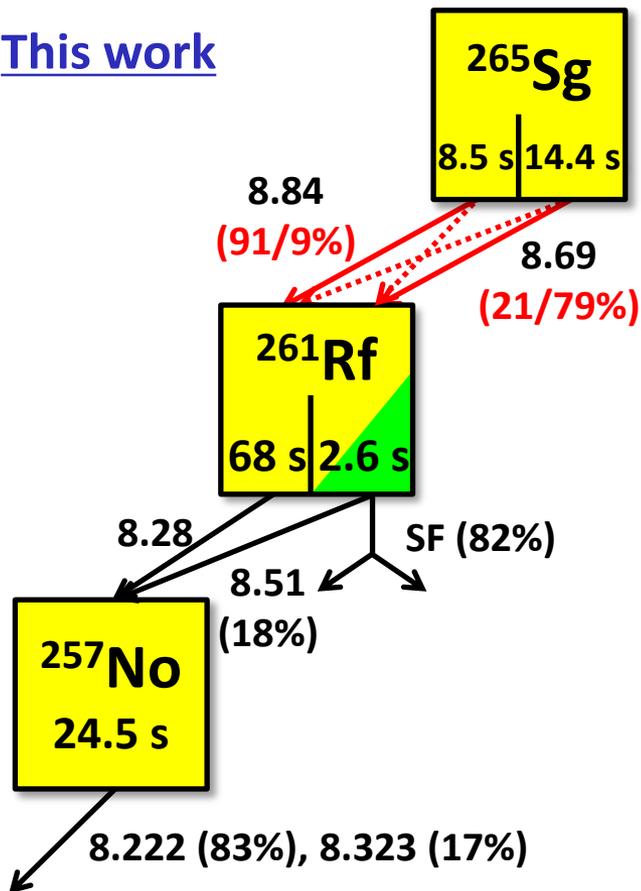
- a) Table of Isotopes, 8th ed.
- b) Asai, private communication.
- c) Asai *et al.*, PRL **95**, 102502 (2005).
- d) Haba *et al.*, PRC **83**, 034602 (2011).



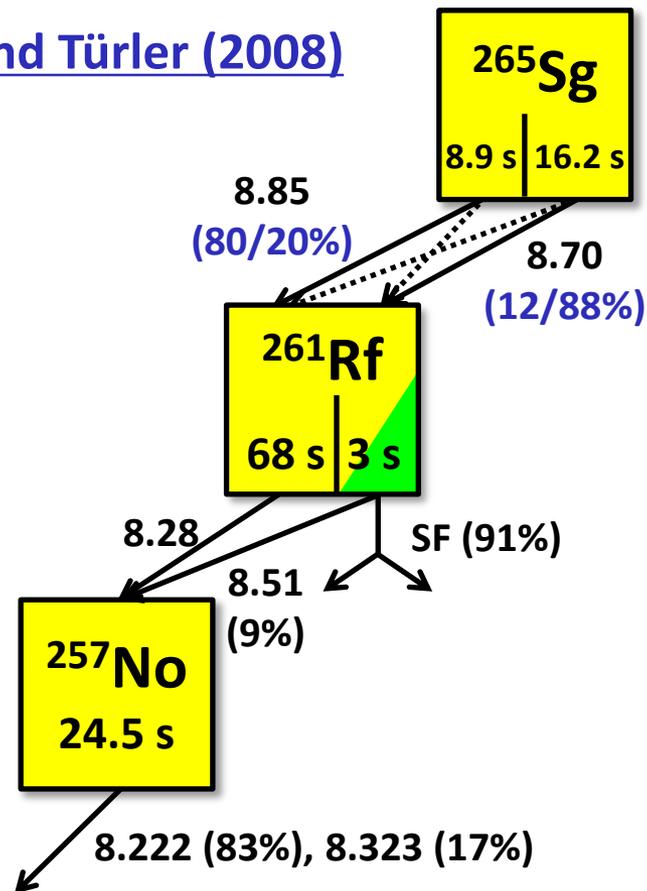
Decay patterns observed in the chain $^{265}\text{Sg}^{a,b} \rightarrow ^{261}\text{Rf}^{a,b} \rightarrow ^{257}\text{No}$

^{265}Sg state	→	^{261}Rf state	No. of events		Branching ratio [%]	
			(obs.)	(corr.)	This work	Düllmann and Türler (2008)
<i>a</i>	→	<i>a</i>	16	19.9	91	80
	→	<i>b</i>	2	2.0	9	20
<i>b</i>	→	<i>a</i>	4	5.1	21	12
	→	<i>b</i>	19	19.0	79	88

This work



Düllmann and Türler (2008)



Cross section

Cross section at 118 MeV [pb]

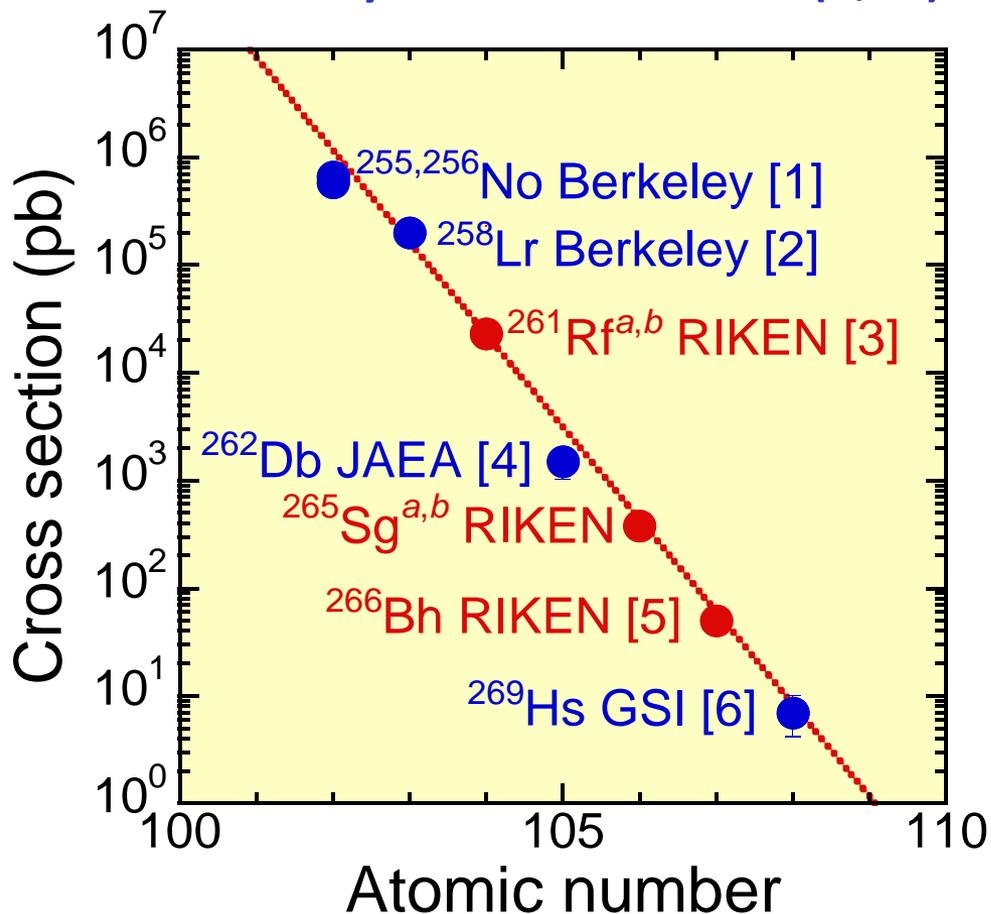
$^{265}\text{Sg}^a$	180^{+80}_{-60}
$^{265}\text{Sg}^b$	200^{+60}_{-50}



$$\sigma(^{265}\text{Sg}^{a+b}) = 380^{+90}_{-70} \text{ pb}$$
$$\sigma(^{265}\text{Sg}^a)/\sigma(^{265}\text{Sg}^b) = 1.3 \pm 0.6$$

Assumptions: GARIS eff. = 13%; gas-jet eff. = 50%; gas-jet transport time = 3 s

Cross section systematics for $^{248}\text{Cm}(X,5n)$

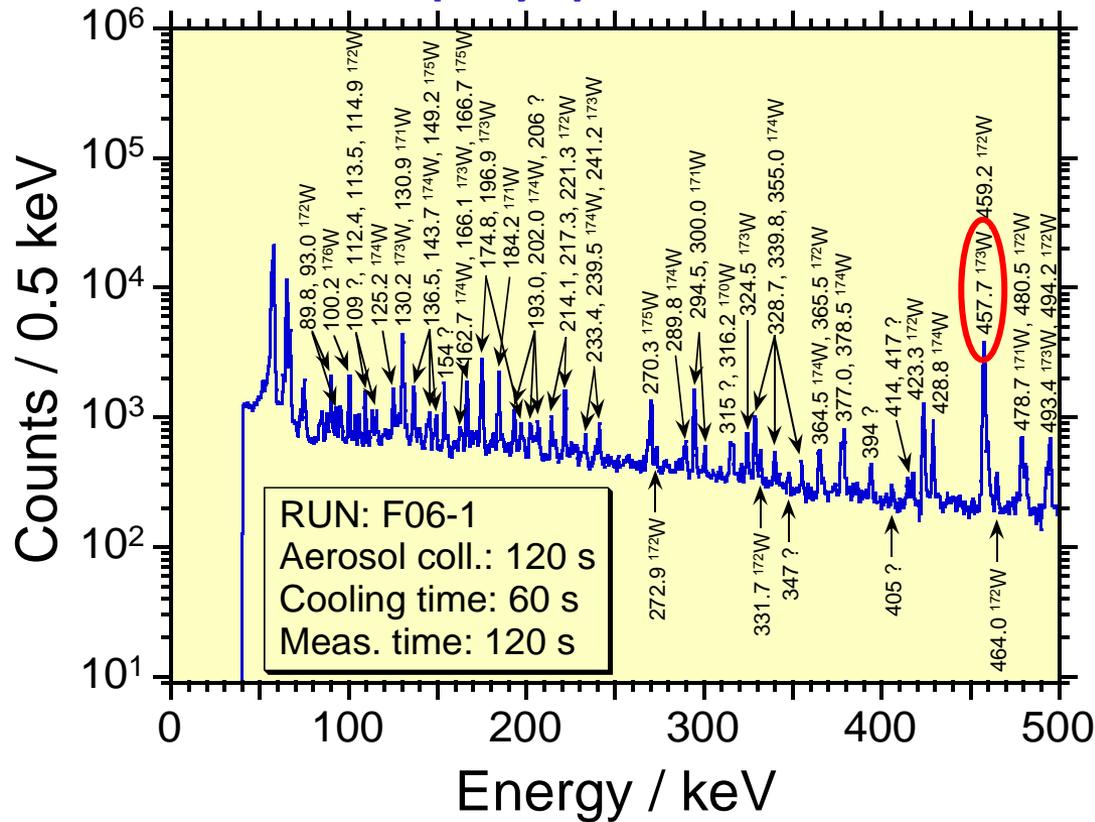


References

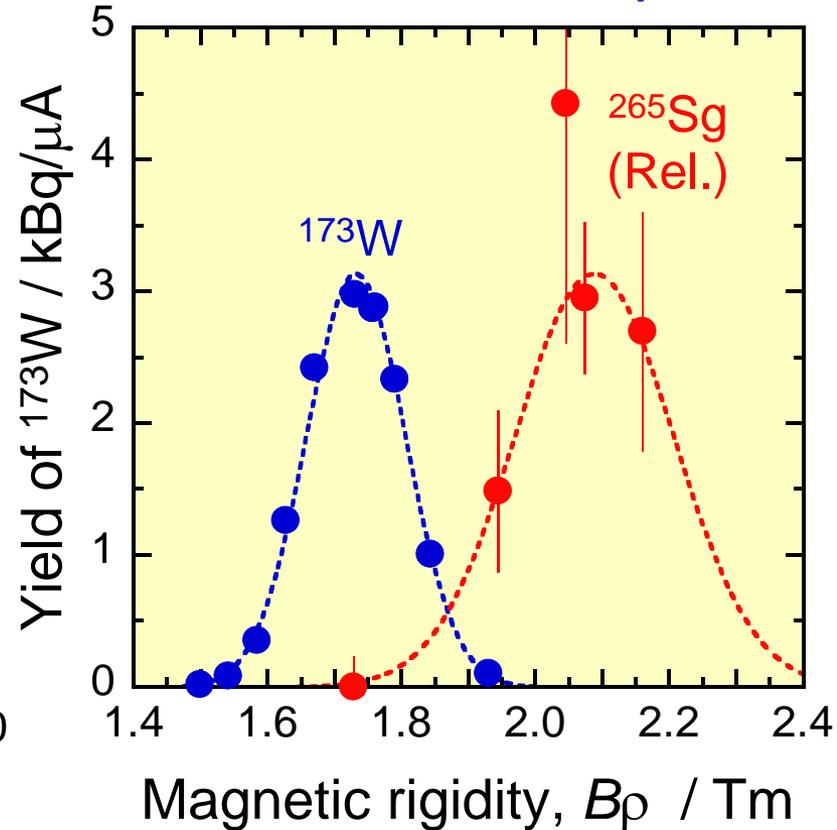
- [1] Sikkeland *et al.*, PL **172**, 1232 (1968).
- [2] Eskola *et al.*, PRC **4**, 632 (1971).
- [3] Haba *et al.*, PRC **83**, 034602 (2011).
- [4] Nagame *et al.*, JNRS **3**, 85 (2002).
- [5] Morita *et al.*, JPSJ **78**, 064201 (2009).
- [6] Dvorak *et al.*, PRL **100**, 132503 (2008).

3.2. $^{nat}\text{Gd}(^{22}\text{Ne}, xn)^{173}\text{W}$

γ -ray spectrum



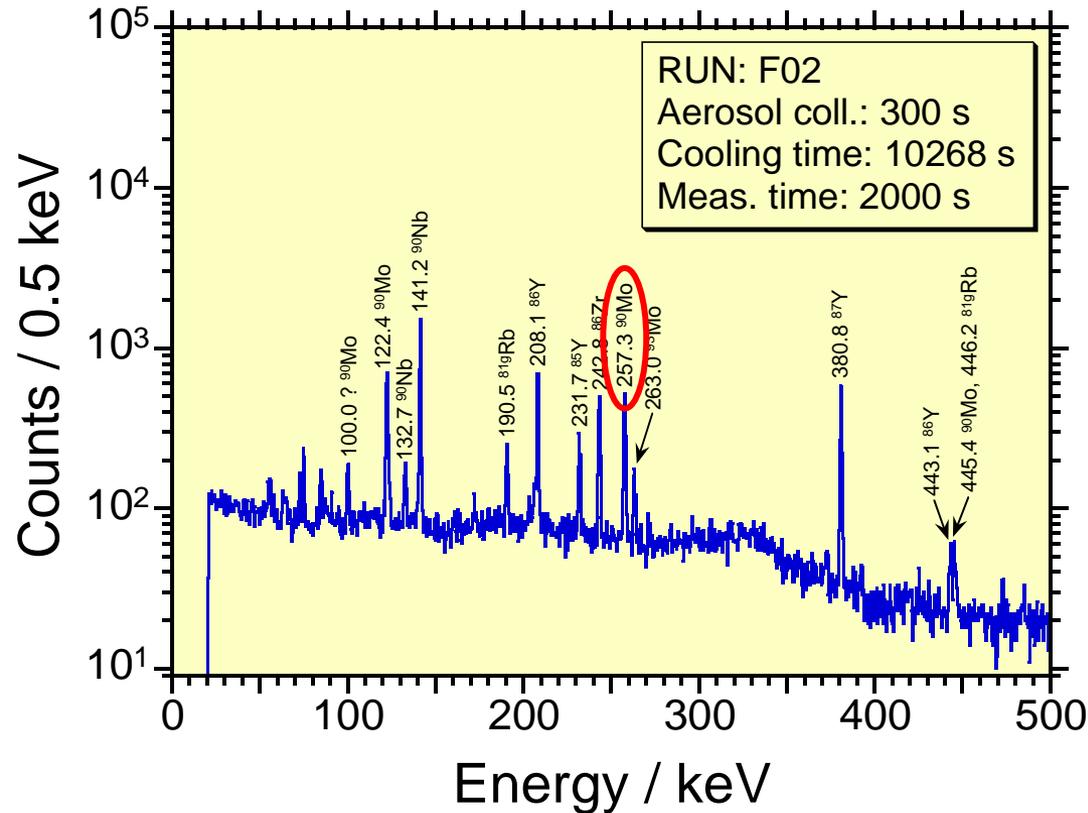
Yield of ^{173}W vs. B_p



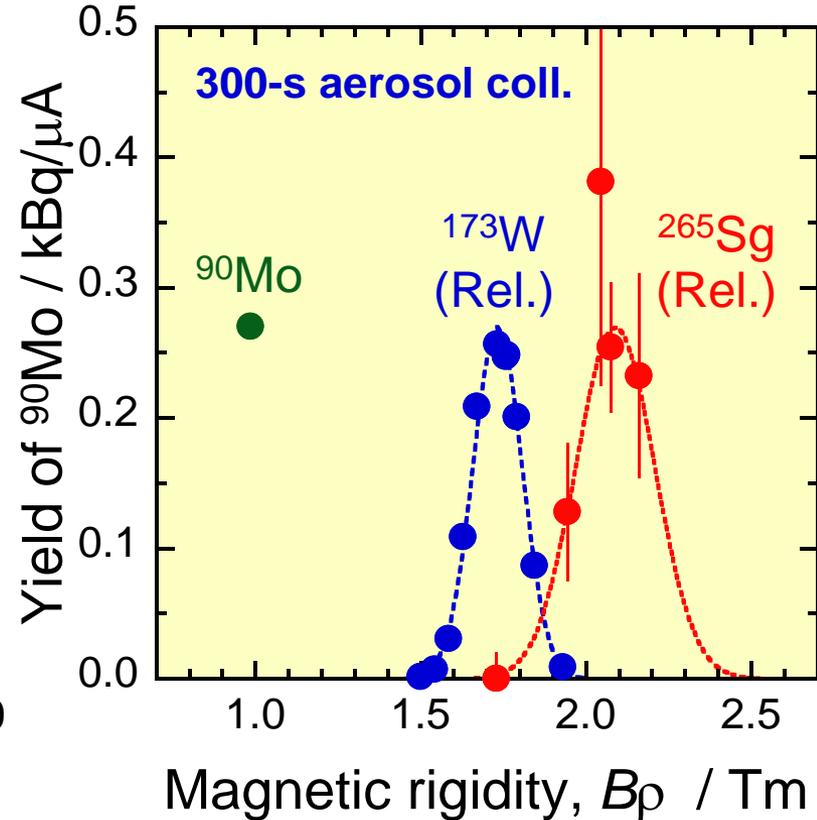
- ^{171}W ($T_{1/2} = 2.38$ min), ^{172}W (6.6 min), **^{173}W (7.6 min)**, ^{174}W (31 min), ^{175}W (35.2 min), and ^{176}W (2.5 h)
- **$B_p = 1.73 \pm 0.01$ Tm** and $\Delta B_p / B_p = 10.1 \pm 0.3\%$
- Yield (Chem. Lab.): **3.1 kBq/ μA** after 120-s aerosol collection
- Gas-jet eff.: **$65 \pm 1\%$**

3.3. $^{nat}\text{Ge}(^{22}\text{Ne},xn)^{90}\text{Mo}$

γ -ray spectrum



Yield of ^{90}Mo vs. $B\rho$



- ^{90}Mo ($T_{1/2} = 5.67$ h), ^{93}Mo , $^{90\text{m}}\text{Mo}$, $^{90\text{g}}\text{Mo}$, $^{89\text{a}}\text{Mo}$, $^{89\text{b}}\text{Mo}$, $^{88\text{g}}\text{Mo}$, $^{87\text{m}}\text{Mo}$, $^{87\text{g}}\text{Mo}$, $^{86\text{g}}\text{Mo}$, $^{89\text{m}}\text{Nb}$, $^{89\text{g}}\text{Nb}$, $^{87\text{m}}\text{Nb}$, $^{86\text{g}}\text{Nb}$, $^{85\text{g}}\text{Nb}$, $^{89\text{m}}\text{Zr}$, $^{89\text{g}}\text{Zr}$, $^{87\text{m}}\text{Zr}$, $^{86\text{g}}\text{Zr}$, $^{85\text{g}}\text{Zr}$, $^{84\text{m}}\text{Zr}$, $^{83\text{g}}\text{Zr}$, $^{81\text{g}}\text{Rb}$, and $^{81\text{m}}\text{Kr}$

- Yield (Chem. Lab.): **0.27 kBq/ μA** after 300-s aerosol collection (0.985 Tm)
(The optimum $B\rho$ and the gas-jet efficiency will be measured in the future.)

4. Summary

- $^{265}\text{Sg}^{a,b}$ for chemical studies of Sg were produced in the $^{248}\text{Cm}(^{22}\text{Ne},5n)$ reaction, and their decay properties were investigated using MANON under low background conditions attained by the GARIS gas-jet system.
- $^{265}\text{Sg}^b$ with $T_{1/2} = 14.4$ s and $\sigma = 200$ pb is available for aqueous chemistry of Sg.
- The homologues of Sg, ^{173}W and ^{90}Mo , were produced in the $^{nat}\text{Gd}(^{22}\text{Ne},xn)$ and $^{nat}\text{Ge}(^{22}\text{Ne},xn)$ reactions, respectively.

→ *We are ready for chemistry experiments of Sg together with its homologues!!*

