

INTERNATIONAL NUCLEAR DATA COMMITTEE

PROGRESS

IN

FISSION PRODUCT NUCLEAR DATA

Information about activities in the field of measurements and compilation/evaluations of fission product nuclear data (FPND)

collected

by

G. Lammer

Nuclear Data Section International Atomic Energy Agency Vienna, Austria

No. 5 June 1979

IAEA NUCLEAR DATA SECTION, KÄRNTNER RING 11, A-1010 VIENNA

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NOT FOR PUBLICATION

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FOREWORD

This is the fifth issue of a report series on Fission Product Nuclear Data (FPND) which is published by the Nuclear Data Section (NDS) of the International Atomic Energy Agency (IAEA). The purpose of this series is to inform scientists working on FPND, or using such data, about all activities in this field which are planned, ongoing, or have recently been completed.

The main part of this report consists of unaltered original contributions which the authors have sent to IAEA/NDS. Therefore, the IAEA cannot be held responsible for the information contained nor for any consequences resulting from the use of this information. The present issue contains also a section with some recent references relative to fission product nuclear data, which were not covered by the contributions submitted.

The types of activities being included in this report are measurements, compilations and evaluations of:

Fission product yields (neutron induced and spontaneous fission); Neutron reaction cross sections of fission products; Data related to the radioactive decay of fission products; Delayed neutron data of fission products; and Lumped fission product data (decay heat, absorption etc.)

The fourth issue of this series has been published in July 1978 as INDC(NDS)-95/G+P. The present issue includes contributions which were received by NDS between 1 August 1978 and 15 May 1979.

The next issue of this report series is envisaged to be published in June 1980.

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IV. Meetings on FPND

SUBMITTING CONTRIBUTIONS

The next issue is expected to be published in June 1980. All scientists who are presently working — or have recently completed work — in the field of FPND and who want to contribute to the 6th issue of this series, are kindly asked to send contributions to me between now and the end of April 1980, so that they reach NDS before 30 April 1980.

Those scientists or groups who have already contributed to the present issue and who want to leave their contribution(s) unchanged or who wish to suggest only slight changes, should write an appropriate note to me before the above deadline.

Format:

The size of one contribution should preferably not exceed one page. Of course, the number of contributions per working group or laboratory is not restricted. Similar experiments (or calculations, evaluations, etc.) performed by one person or group should preferably be combined to one contribution, if this is possible without loss of clarity.

The headings suggested for the 3 types of contributions are, for

Measurements:	Compilations:	Evaluations:
Laboratory and address: Names: Facilities:	Laboratory and address: Names:	Laboratory and address: Names:
Experiment: Method: Accuracy: Completion date: Discrepancies to other reported data: Publications:	Compilation: purpose: major sources of information: deadline of literature coverage: cooperation: other relevant details: computer file: completion date: Publications:	Evaluation: purpose: method: major sources of information: deadline of literature coverage: status: cooperation: other relevant details: computer file of compiled data: computer file of evaluated data: discrepancies encountered: completion date: Publications:

For the sake of consistency it is requested that the suggested headings be used as far as appropriate.

<u>Comments or suggestions</u> concerning the format, content and layout of this report series are most welcome and should be directed to me in time before the next issue.

I would like to thank the contributors for their cooperation.

G. Lammer

1. MEASUREMENTS

1.1. Fission yields

Fission. nuclide	incident neutron energy	page	Fission. nuclide	incident neutron energy	page
Th -2 32 U -2 32	pile 0.1 - 8 MeV thermal	(14) <u>28</u> 36 <u>16</u>	U-238	fast fast, 15 MeV 0.1 - 8 MeV spontaneous	32 38 5 36 41
U-2 33	thermal	<u>15</u>	Np-237	fast	38
	fast fast, 15 MeV	(28) (54) 5 38 5 5	Pu-239	thermal	15 <u>17</u> 21 27 (31)
V-2 35	thermal	$(8) \\ 10 \\ (11) \\ (12) \\ 15 \\ 17 \\ 21 \\ 21 \\ (8) \\ 10 \\ 10 \\ 10 \\ 10 \\ 10 \\ 10 \\ 10 \\ 1$		fast fast, 15 MeV	$ \begin{array}{c} 40 \\ 47 \\ (54) \\ 29 \\ (31) \\ (32) \\ 37 \\ 5 \end{array} $
		(28) (31) 39 53 (54)	Pu240	fast	(32) 38 45
	thermal (Se-83 only) thermal, fast, 3 MeV	1	Pa-241	thermal fast	17 <u>48</u> (32) 38
	fast	30	Pu-242	fast	38
	fast, 15 MeV	(32) 38 5	Am-241	fast 14 MeV	38 45 46
U-23 8	fast	(14)	Am-243	fast	38
		29	C f2 52	spontaneous	<u>18</u>

a) with respect to the earlier issues, underlined page numbers refer to new work, page numbers in brackets refer to unchanged contributions, and others refer to revised contributions.

FP isotope	reactions	incident neutron energy	page
Br-79 Br-81	} respars.	}3 - 1190 e v	20
Kr-85	transm. (res.)	1 – 1500 eV	(7)
Kr -86	1	1	<u>50</u>
Sr-86 Sr-87 Sr-88 Y-89 Zr-90 Zr-91	(n, y), res.	2.6 - 500 keV	50
Z r-91	(n,γ) , tot., res.	150 eV - 20 keV	(2)
Z r-92 Z r- 94	$\left.\right\}$ (n, γ), res.	2.6 - 500 keV	(50)
Z r- 96	(n,γ) , tot., res.	150 eV – 130 ke	V (2)
Nd-93	$\begin{pmatrix} n,\gamma\\ n,\gamma \end{pmatrix}$, res.	3 - 80 keV 2.6 - 500 keV	<u>22</u> (50)
Mo-92 Mo-94 Mo-95 Mo-96 Mo-97 Mo-98 Mo-100	(n,γ), res.	2.6 - 500 keV	50
Tc-99	transm. (res.)	1 – 1500 eV	7
Ru-100 Ru-101 Ru-102 Ru-104 Rh-103	(n,γ), res.	2.6 - 500 keV	50
Pd-104	(n,γ) , tot., scat.	below 50 keV	(4)
Pd-105	(n,γ) , res. (n,γ) , tot., scat.	2.6 - 500 keV below 50 keV	50 4
Pd-106	(n,γ) , res. (n,γ) , tot., scat.	2.6 - 500 keV below 50 keV	50 4
P d1 08	(n,γ) , res. (n,γ) , tot., scat.	2.6 - 500 keV below 50 keV	50 4
P d-11 0	(n,γ) , res. (n,γ) , tot., scat.	2.6 - 500 keV below 50 keV	50 (4)
Ag-107	tot., el., inel.	0.25 - 4.5 MeV	<u>37</u>
Cd-106 Cd-108 Cd-110	$\left.\right\}$ (n, γ), res.	2.6 - 500 keV	(50)

1.2. Neutron reaction cross sections

FP isotope	reactions	incident neutron energy	page
Cd-111 Cd-112 Cd-113 Cd-114 Cd-116 Te-122 Te-123 Te-124 Te-125 Te-126 Te-128 Te-130	$\left(n,\gamma\right)$, res.	}2.6 - 500 keV	(50)
I -12 7	(n, y)	3 - 80 keV	<u>22</u>
Cs-133	(n,γ) (n,γ) , res. transm. (res.)	3 - 80 keV 2.6 - 500 keV 1 - 1500 eV	(<u>50</u>) (7)
Cs-135 Cs-137	transm. (res.)	1 - 1500 eV	(7)
Ba-134 Ba-135 Ba-136 Ba-137 Ba-138 La-139 Ce-140 Pr-141 Nd-142) (n,γ), res.	2.6 - 500 keV	50
Nd-143	(n,γ)	5 - 300 keV	<u>19</u>
	(n,γ) , res.	2.6 - 500 keV	(50)
	integral (n,γ)	EBR-II	42
Nd-144	(n,γ), res.	2.6 - 500 keV	(50)
	integral (n,γ)	EBR-II	42
Nd-145	(n,γ)	5 - 300 keV	<u>19</u>
	(n,γ) , res.	2.6 - 500 keV	(50)
	integral (n,γ)	EBR-II	42
Nd-146	(n,γ)	5 - 300 keV	<u>19</u>
	(n,γ) , res.	2.6 - 500 keV	(50)
	integral (n,γ)	EBR-II	42
Nd-148	(n,γ)	5 - 300 keV	<u>19</u>
	(n,γ), res.	2.6 - 500 keV	(50)
	integral (n,γ)	EBR-II	42
Nd-150	integral (n, y)	EBR-11	42
Sm-147	(n,γ)	1 eV - 300 keV	<u>19</u>
	integral (n,γ)	EBR-II	42
Sm-149	(n,γ)	1 eV - 300 keV	<u>19</u>
	integral (n,γ)	EBR-II	42

1.2. Neutron reaction cross sections (continued)

FP isotope	reactions	incident neutron energy	page
S - 151	transm. (res.)	1 – 1500 eV	I
Eu Eu~151	{ (n,γ)	} 3 - 100 keV	<u>19</u>
Eu-151	integral (n,y)	ÉBR-II	42
	transm. (res.)	1 - 1500 eV	(7)
Eu-152	integral (n, γ)	EBR-II	42
	transm. (res.)	1 - 1500 eV	(7)
Eu-153	(n,γ)	3 - 100 keV	<u>19</u>
	integral (n,γ)	EBR-II	42
	transm. (res.)	1 - 1500 eV	(7)
Eu -15 4	integral (n,γ)	EBR-II	42
	transm. (res.)	1 - 1500 eV	(7)
Eu-155	transm. (res.)	1 - 1500 eV	(7)
T b1 59	(n,γ), res.	2.6 - 500 keV	50
	respars.	3 - 1190 eV	20
H 0-1 65 Tm-169	$\left.\right\}$ (n, γ), res.	2.6 - 500 keV	50

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1.2. Neutron reaction cross sections (continued)

1.3. Decay data

	-	page	FP isotope	data type	page
Zn-75 Zn-76 Zn-77 Zn-77 Zn-76 Ga-76 Ga-76 Ga-77 Ga-78 Ga-78 Ga-79 Ga-78 Ga-79 Ga-81 Ge-79 Ge-80 As-80 As-81 As-85 Br-85 Br-86 Br-87 Br-88 G	$\frac{2}{3}$ $\frac{3}{3}$, γ -spectra $\frac{2}{3}$ HL; γ , conv. el. $\frac{2}{3}$ $\frac{3}{3}$, γ -spectra	25 <u>26</u> 25 26 25 26	Br-89 Rb-91 Rb-93 Rb-94 Rb-95 Rb-96 Rb-97 Tc-106 Tc-107 Tc-108 In-120	β-, γ-spectra Q_{β} β strength funct. β strength funct. β-, γ-spectra β strength funct. β-, γ-spectra	$ \begin{array}{r} \frac{26}{25} \\ (8) \\ (8) \\ \frac{26}{(8)} \\ \frac{26}{(8)} \\ \frac{26}{(8)} \\ \frac{26}{(8)} \\ (24) \\ (24) \\ \frac{26}{25} \end{array} $

FP isotope Qβ In-121 Qβ In-122 HL; Qβ In-123 Qβ In-124 HL; Qβ In-125 Q8	data type γ, conv. el. γ, conv. el.	page 25 26 25 25 25 26 25	FP isotope Sb-130 Sb-131 Sb-132 Sb-134 Sb-135	data type $\left\{ \begin{array}{l} Q_{\beta} \\ \beta-, \gamma-\text{spectra} \end{array} \right\}$
In-121 Q _β In-122 HL; γ Q _β In-123 Q _β In-124 HL; γ Q _β In-125 Q _β	γ, conv. el. γ, conv. el.	25 26 25 25 25	Sb130 Sb131 Sb132 Sb134 Sb135	$\left\{ \begin{array}{l} Q_{\beta} \\ \beta \end{array} \right\}$ β -, γ -spectra
$ \begin{array}{ccc} In-126 & HL;, \\ Q_{\beta} \\ In-127 & Q_{\beta} \\ In-128 & HL; \\ Q_{\beta} \\ In-129 & Q_{\beta} \\ A = 133 \\ Sn-127 \\ Sn-132 \\ Sb-128 \end{array} \Big Q_{\beta} \\ \end{array} $	γ, conv. el. γ, conv. el.	25 26 25 25 26 25 25 (13) 25	Te-134 Te-135 I-137 I-138 I-139 Xe-133m Ba-139 Ba-140 La-140 Ce-144 Pr-144 Pr-145 Pr-146	$\begin{cases} Q_{\beta} \\ HL; \gamma, conv. \\ \beta-, \gamma-spectra \\ \beta-, \gamma-spectra \\ conv. coeff. \\ HL, absol. \gamma- \\ absol. \gamma-int. \end{cases}$

1.3. Decay data (continued)

25 <u>26</u> y-spectra 25 26 y, conv. el. 26 y-spectra γ -spectra **2**6 • coeff. (K,L) (33) absol. y-int. 44 l. γ-int. <u>35</u> l. y-int. 44

page

1.4. Delayed neutron data

FP isotope or fissile	data type	page	FP	'isotope or fissile	data t ype	page
Ga-79 Ga-80 Ga-81 Ga-82 Ga-83 Br-87 Br-87 Br-87 Br-88 Br-88 Br-89 Br-89 Br-90 Br-91 Br-92 Rb-91 Rb-92	$ \left. \right\} P_{n} \\ \left. \right\} E-spec. \\ P_{n} \\ \right\}$	25 25 25 25 25 (9) 25 (9) 25 (9) (9) (9) (9) (9) (9) (9) (9) (9) 25		Rb-93 Rb-94 Rb-95 Rb-96 Rb-97 In-127 In-128 In-129 In-130 In-131 In-132 Sb-134	E-spec. Pn E-spec. Pn E-spec. Pn E-spec. Pn Pn E-spec.	(8) 25 (8) 25 (8) 25 (8) 25 25 25 25 25 25 25 25 25 25 25 25 25

TT	isotope fissile	or	data t y pe	page
	Sb-135 Sb-135 I-137 I-137 I-138 I-138 I-138 I-139 I-139 I-140 I-141 Cs-141 Cs-141 Cs-142 Cs-143 Cs-144 Cs-145		P _n	(9) 25) 2(92) 2(92) 2(92) 2(92) 255 255 25 25 25 25 25 25 25 25 25 25 2
	0-237		£quii. spectra	24

1.4. Delayed neutron data (continued)

1.5. Decay heat

Fissionable isotope	neutron energy	radiation	page
Th-232 U nat. U-235	fast fast thermal fast fast	 β, γ; total β, γ; total β, γ; total β, γ; total β 	} (23) 49 (23) 29
U-23 8	fast	β, γ; total	(23)
Pu-239	thermal fast fast	$\begin{array}{c} \beta, \gamma; \text{ total} \\ \beta, \gamma; \text{ total} \\ \beta \end{array}$	49 (23) 29
Pu-241	thermal	β, γ; total	49

•

2. COMPILATIONS AND EVALUATIONS

Data category	page	
Fission yields	58 (66) (67) 74 <u>75</u> <u>76</u> (82)	(theor. prediction) (compil., Crouch) (Crouch) (ENDF/B-V) " " (charge dist.)
Cross sections	60 62 63 <u>72</u> 76 (78) 79	(CNEN-CEA) (Nd-isotopes) (RCN-2) (BNL-325) (ENDF/B-V)
Decay data	26 57 (61) 69 73 <u>76</u> 79	(avg β-, γ-energies) (for 14 F.P.) (Compil., JNDC) (UK) (ENDF/B-V)
Delayed neutrons	(61) <u>68</u> 70	(compil., JNDC) (HL, P _n) (equilibrium spectra)
Decay heat	(61) 79	(JNDC) (ENDF/B-V)

I. MEASUREMENTS

(revisions with respect to the last issue are marked by a vertical bar in the margin)

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		BELGIUM
Laboratory	:	Centre d'Etude de l'Energie Nucléaire CEN/SCK B-2400 MOL (Belgium)
Names	:	P. del Marmol, P. Fettweis
Facilities	:	BR1 Reactor
Experiment	:	The independent isomeric yield ratio of ^{83m} Se/ ^{83g} Se is being determined in thermal neutron fission of ²³⁵ U.
Method	:	Fast radiochemical separation of Se followed by γ -counting of the resulting activities.
Completion	date :	The counting statistics of the measurements expected to be completed in 1978 were too low : The experiment is being repeated.

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E.E.C. Belgium

(same as in INDC(NDS)-95)

Laboratory and address:	JRC, CBNM, Geel, Belgium ⁺ CNEN, Bologna, Italy
Names:	A. Brusegan, C. Coceva ⁺ , F. Corvi, P. Giacobbe ⁺ , M. Magnani ⁺ , G. Rohr.
Facilities:	Neutron time-of-flight spectrometers at the 60 MeV Linac (pulse width: 23 nsec).
Experiment:	Resonance parameters for 91 Zr and 96 Zr.
	Separated isotopes: 89% enriched ⁹¹ Zr and 57% ⁹⁶ Zr.
	1) Capture γ -ray measurements (⁹¹ Zr only).
	Energy range: 150 eV - 3200 eV.
	2) Capture measurement.
	Energy range ⁹¹ Zr: 150 eV - 20 keV 96Zr: 150 eV - 100 keV.
	3) Total cross section measurements.
	Energy range $\begin{array}{rrr} 91\\ 96\\ Zr: & 150 \text{ eV} - 14.8 \text{ keV}\\ 36\\ Zr: & 150 \text{ eV} - 130 \text{ keV}. \end{array}$
Method:	l) High and low energy γ -ray spectra measurements.
	Detector: GeLi gamma ray spectrometer. Flightpath: 13 m. Sample thickness: 14.4 [.] 10 ⁻³ at/barn.
	2) Capture cross section measurements.
	Detector: C_6F_6 detectors using Maier-Leibnitz method. Flightpath: 60 m. Sample material: ZrO Sample thicknesses: $91Zr$: $7.310 \cdot 10^{-3}$ at/barn $96Zr$: $4.780 \cdot 10^{-3}$ at/barn Neutron flux: boron slab with C_6F_6 detectors. Normalization: Ag using black resonance technique.
	3) Total cross section measurements.
	Detectors: NaI(T1) crystals with a boron slab. Flightpath: 100 m. Sample material: ZrO ₂ enriched to 57% 96 and 89% 2r. Sample thicknesses: 91 Zr 0.8 · 10 ⁻³ at/barn 2.4 · 10 ⁻³ at/barn 14.4 · 10 ⁻³ at/barn

E.E.C. Belgium

(cont'd)

		⁹⁶ Zr	$0.8 \cdot 10^{-3} \\ 3.5 \cdot 10^{-3} \\ 4.3 \cdot 10^{-3}$	at/barn at/barn at/barn
Accuracy:	Expected on final resona	ince par	ameters	
	91 Zr g Γ_n , $\Gamma_{\dot{\gamma}}$ between	5% and	20%	
	96 Zr g Γ_n , Γ_γ between	10% and	20%	
,	depending on the energy of the resonances.	range a	nd the stre	ngth
Expected completion date:	⁹¹ Zr : analysis finish	ied		
	⁹⁶ Zr : end of 1978			
Publications :	A. Brusegan, F. Corvi P. Giacobbe and M. Ma Neutron Resonance Par on Neutron Physics an Harwell, September 19	, G. Ro gnani, ameters d Nucle 78, p.	hr, C. Coc of ⁹¹ Zr, ar Data fo 706	eva, Proc. Int. Conf. r Reactors,

.

E.E.C., Belgium

Laboratory and address :	JRC, CBNM, Geel, Belgium S.C.K./C.E.N., Mol, Belgium
Names :	P. Staveloz, ⁺ E. Cornelis, ⁺ L. Mewissen, ⁺ F. Poortmans G. Rohr, R. Shelley, J. Winter, T. van der Veen
Facilities :	Neutron time-of-flight spectrometers at the 150 MeV Linac (pulse width : 4 nsec)
Experiments :	Resonance parameters for ^{104,105,106,108,110} Pd below 50 keV
	 Capture cross section measurements : ¹⁰⁵Pd and ¹⁰⁸Pd : measurements and analysis completed.
	2) Total cross section measurements : ¹⁰⁵ Pd and ¹⁰⁸ Pd : measurements and analysis completed. ¹⁰⁶ Pd : measurements completed, analysis in progress. ¹¹⁰ Pd : measurements nearly completed.
1	 Scattering cross section measurements : ¹⁰⁵Pd and ¹⁰⁸Pd : measurements and analysis completed.
Method :	1) Capture cross section measurements.
	Detector : $C_{6}D_{6}$ - detectors using Maier Leibnitz method. flight path length : 60 m.
	2) Total cross section measurements.
	Detector : ³ He gaseous scintillators. flight path length : 30 m and 60 m. cooled samples at liquid nitrogen temperature.
	3) Scattering cross section measurements.
	Detector : ³ He gaseous scintillators. flight path length : 30 m.
Accuracy :	Mostly between 5% and 20% on resonance parameters $g\Gamma_n$ and Γ_γ depending on the energy range and on the strength of the resonance.
Expected completion date:	Pd and Pd completed
	analysis of data for other isotopes : June 1980.
Publications :	Neutron Resonance Parameters of ¹⁰⁵ Pd and ¹⁰⁸ Pd, P. Staveloz, E. Cornelis, L. Mewissen, F. Poortmans, G. Rohr, R. Shelley, T. van der Veen, Proc. Int. Conf. on Neutron Physics and Nuclear Data for Reactors, Harwell, September, 1978, p. 701.

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FRANCE

- Laboratory : Service RADIOCHIMIE ET PHENOMENOLOGIE Centre d'Etudes de BRUYERES-LE-CHATEL B.P. n° 561 - 92542 MONTROUGE CEDEX - FRANCE.
- Names : J. LAUREC A. ADAM.
- <u>Facilities</u>: PROSPERO Critical assembly and LANCELOT 14 Mev neutrons generator (S.E.C.R./C.E. VALDUC) Radiochimical Laboratory Calibrated Ge-Li spectrometers.
- $\begin{array}{l} \underline{Experiments}: & \text{Determination of cumulative yields of some fission products} \\ & (\begin{array}{c} {}^{95}\text{Zr}, \begin{array}{c} {}^{97}\text{Zr}, \end{array} \begin{array}{c} {}^{99}\text{Mo}, \end{array} \begin{array}{c} {}^{103}\text{Ru}, \begin{array}{c} {}^{105}\text{Rh}, \end{array} \begin{array}{c} {}^{127}\text{Sb}, \end{array} \begin{array}{c} {}^{131}\text{I}, \end{array} \begin{array}{c} {}^{132}\text{Te}, \end{array} \begin{array}{c} {}^{140}\text{Ba}, \end{array} \\ & {}^{141}\text{Ce}, \end{array} \begin{array}{c} {}^{143}\text{Ce}, \end{array} \begin{array}{c} {}^{144}\text{Ce}, \end{array} \begin{array}{c} {}^{147}\text{Nd} \end{array} \begin{array}{c} \text{for} \end{array} \begin{array}{c} {}^{233}\text{U}, \end{array} \begin{array}{c} {}^{235}\text{U}, \end{array} \begin{array}{c} {}^{238}\text{U} \end{array} \begin{array}{c} \text{and} \end{array} \begin{array}{c} {}^{239}\text{Pu}, \end{array} \\ & \text{with fission spectrum and 14,7 Mev neutrons.} \end{array} \end{array}$
- <u>Method</u>: The fission number is measured by a fission chamber. The fission products activities of fissile target nuclides are determined by gamma direct spectrometry measurements with calibrated Ge-Li spectrometers. The targets and chamber deposits masses are determined by alpha and mass spectrometries.
- <u>Accuracy</u>: 3 to 5 %; the branching ratio error is not included; this last error is variable from one isotope to the other (1 % to 5 %).

Completion : end of 1979.

Publication : internal reports ; C.E.A. report to be published.

FRANCE

Laboratory and address :	Département de Recherche Fondamentale Laboratoire de Chimie Physique Nucléaire Centre d'Etudes Nucléaires de Grenoble 85 X - 38041 GRENOBLE CEDEX - France.
Names :	J. BLACHOT, J. CRANÇON, Ch. HAMELIN, G. LHOSPICE et A. MOUSSA.
Facilities :	Melusine reactor (thermal neutron and caramel system for fast neutrons) 3 MeV neutrons generator and high flux reactor of I.L.L.
Experiment :	The fractionnal independent yield of ${}^{86}\textsc{Br}$, ${}^{89}\textsc{Rb}$, ${}^{134}\textsc{I}$, ${}^{135}\textsc{Xe}$, ${}^{136}\textsc{m+g}\textsc{I}$, ${}^{136}\textsc{Cs}$, ${}^{137}\textsc{Xe}$, ${}^{144}\textsc{La}$ isotopes and the fractionnal cumulative yields of the parents have been or would be measured for $\frac{1}{2}$ ${}^{235}\textsc{U(n_{f},f)}$, ${}^{235}\textsc{(n_{3MeV},f)}$.
<u>Method</u> :	Radiochemical separations for Cs136 and Ge/Li gamma spectrometry. For the others, direct growth and decay activities are measured with a Ge/Li detector and recorded in a multispectrum mode by a 4K multichannel analyser.
Accuracy :	The average relative uncertainty of our measurements is between 5 and 10%.
Completion date :	December 1980.
Publication :	International Symposium on Physics and Chemistry of Fission - 14/18 May 1979 - Jülich.

.

Laboratory and address:

Institut für Reine und Angewandte Kernphysik der Universität Kiel (IKK), D-2054 Geesthacht, Reaktorstation

Names:

P. FISCHER, U. HARZ, P. PODEWILS, H.G. PRIESMEYER

Facility:

Fast Chopper Neutron Time-of-Flight Spectrometer, 42 m flight path in front of beam hole of 5 MW FRG-1 reactor. 15 ns/m nominal resolution, with special equipment for transmission measurements on highly radioactive samples; 11 Li-6 glass-scintillation detectors; max. rotor speed 15 000 upm; min. burst width 0.64 μ s; min. time channel width 100 ns.

Experiments:

Neutron resonance investigation by transmission measurements between 1 eV and 1.5 keV on separated stable or radioactive isotopes of special interest to reactor physics (especially fission products), gross fission products. Possibility of extending energy range to thermal region with crystal spectrometer, which can be made available.

Completed:	Tc 99 transmission with different sample thickness to clear up discrepancies in Γ of first resonance; Data published in Proceedings of "Neutron Physics and Nuclear Data for Reactors and Other Applied Purposes" Harwell, September 1978, page 718.
Ongoing:	Cs 133/135/137 FP mixture for isotope identification of 42.8 eV and 880 eV resonances; Pu 240 precision measurement for resonance parameter determination of 1.056 eV resonance.
Planned:	EuO isotope mixture from fast reactor control rod (transmission and resonance identification); Kr transmission (stable isotopes), in order to prepare Kr 85 measurement, Sm 151 transmission.

Method:

Sample-in-beam, sample-out-of-beam transmission measurement; black resonance or boron filter background determination technique.

Accuracy:

For resonance parameters: about 5 % or better, depending on statistical accuracy desired.

(same as in INDC(NDS)-95)

Laboratory and address:	Institut für Kernchemie Universität Mainz Postfach 3980 D-6500 Mainz, Germany
Names [*] :	 H. Ohm, A. Schröder, KL. Kratz (Kernchemie Mainz), K.D. Wünsch, G. Jung (Univ. Giessen/ILL Grenoble), C. Ristori, J. Crancon (CEN Grenoble) * This work is a collaboration with: Institut Laue-Langevin - BP 156 - 38042 Grenoble. II. Physikal.Institut, Univ. Giessen - D-6300 Giessen. Laboratoire de Chimie Physique Nucléaire - DRF - CENG - 38041 Grenoble.
Facilities:	Alkali isotope separator OSTIS installed at the Grenoble high-flux reactor.
Experiment:	Energy spectra of β -delayed neutrons have been measured in coincidence with γ -rays depopulating excited states in the respective neutron final nucleus.
	With these data and the information from neutron singles and γ -ray spectra β -strength functions which extend to near Q _p of six Rb isotopes have been constructed (A = 89,91, 93-95,97), their shapes exhibiting - in contrast to the present theoretical concepts - pronounced resonances and a systematic behaviour as a function of mass number an Q _p as expected from general nuclear strucutre considerations. The particular importance of these investigations lies in the fact that the shape of the β -strength function is decisive not only in predictions of β -decay half- lives and β -delayed neutron emission probabilities, but also for radioactive decay heat analyses, espe- cially for "loss-of-coolant accidents" (LOCA). Work is progressing.
Publications:	<pre>KL. Kratz et al., Proc. Int. Workshop, Hirschegg, Austria, AED-Conf. 77-017-001 ff (1977) 208. KL. Kratz et al., NEANDC(E)-192, vol. 5, p. 72 (1978).</pre>

(same as in INDC(NDS)-95)

Laboratory and address:	Institut für Kernchemie Universität Mainz Postfach 3980 D-6500 Mainz, Germany
Names:	W. Rudolph, KL. Kratz, G. Herrmann
Facilities:	TRIGA Mark II reactor
Experiment:	The fractional independent fission yields and β^- -delayed neutron abundances of the following nuclides were determined in thermal- neutron induced fission of ^{235}U : $^{87-92}Br$, 134 , ^{135}Sb , $^{137-141}I$.
	With these data the neutron emission probabilities of the above precursors were obtained.
Method:	After pulsed irradiation (30 msec FWHM), fast chemical separations by hydride volatilization for Sb, and by formation of volatile methyl- compounds through recoil reactions for Br and I. γ -spectroscopic measurements via daughter nuclides, and neutron counting
Accurary:	Error margins between ± 10 % and ± 60 % of the resulting value.
Publications:	W. Rudolph et al., J. Inorg. Nucl. Chem. <u>39</u> (1977) 753. KL. Kratz, Radiochim. Acta <u>25</u> (1978), 1.

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Laboratory:	Institut für Kernchemie			
	Universität Mainz			
	Postfach 3980			
	D-6500 Mainz, Germany			
Names:	M. Weis, H.O. Denschlag			
Facilities:	TRIGA Mark II Reactor			
Experiment:	The fractional independent or fractional cumulative			
	(FC) yields of the following nuclides were determined			
	in the fission of ²³⁵ U by thermal neutrons:			
	Nb-96, Nb-97(m+g), Nb-98m, Y-99 (FC), Zr-99,			
	Nb-99m, Nb-99g, Zr-101 (FC), Nb-101, Mo-101,			
	Zr-102 (FC), Nb-102(m+g), Nb-104(FC), Nb-105 (FC).			
Method:	Fast radiochemical separation of Nb or Zr after pulsed irradiation.			
Accuracy:	Generally $\pm 5\%$ (relative to value).			
Completion date:	1979			
Publications:	Jahresberichte 1975, 1976 and 1977			
	Institut für Kernchemie			
	Universität Mainz			
ļ	M. Weis, Dissertation, Mainz 1979			

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Laboratory:	Institut für Kernchemie Universität Mainz
	Postfach 3980
	D-6500 Mainz, Germany
Names:	G. Fischbach, H.O. Denschlag
Facilities:	TRIGA Mark II Reactor
Experiment:	The fractional cumulative ([*] independent) yields of the following isotopes were measured in ²³⁵ U(n _{th} ,f): Ba-141 [*] , Ba-143 to Ba-146,
	La-146, Ce-146, Ce-148 and Ce-149
Method:	Fast radiochemical separations and direct γ -ray
	<pre>spectroscopic measurement and/or indirect measurements</pre>
	via daughter nuclides.
Accuracy:	Varying
Completion date:	1979
Publications:	Jahresberichte 1975, 1976 and 1977
	Institut für Kernchemie
	Un ^m iversität Mainz

- 12 -GERMANY, FED. REP.

(same as in INDC(NDS)-95)

Laborat ory:	Institut für Kernchemie Universität Mainz
	Institut Lave Langevin
	29 Granoblo Englevin
	So drenoble, france
Names:	H.O. Denschlag, Z. Alfassi (U. Beersheva, Israel), J. Blachot (CENG, Grenoble), H. Braun, W. Faubel, T. Izak-Biran (SOREQ, Israel), H. Meixler, G. Paffrath,
	W. Pörsch, H. Schrader, G. Siegert, T. Tamai (KURRI, Japan),
	A.C. Wahl (U.St.Louis, Mo,USA), K. Wolfsberg (LASL, USA)
Facilities:	LOHENGRIN Mass-separator for unslowed fission products at ILL, Grenoble
Experiment:	The charge distribution in the heavy-mass-peak fission products from 235 U(n _{th} ,f) is being measured at various well defined kinetic energies (excitation energies) of the fission fragments.
Method:	Fission fragments separated according to mass (Resolution $\frac{M}{\Delta M}$ = 400) and kinetic energy (resolution ~2 MeV) are
	intercepted on a moving transport tape, transported continuously or discontinuously in front of a Ge(Li) γ -ray detector, and counted via the γ -rays emitted in their β -decay.
Accuracy:	Varying
Completion:	1979
Publication:	Progress reports are appearing since 1975 in Jahres- berichte, Institut für Kernchemie, Universität Mainz and Annex to the Annual Reports, Institut Laue Langevin, Grenoble.

(same as in INDC(NDS)-95)

Laboratory:	Institut für Kernchemie Universität Mainz Postfach 3980 D-6500 Mainz, Germany
Names:	H. Braun, H.O. Denschlag
Facilities:	TRIGA Mark II Reactor
Experiment:	Yields and decay properties of the fission product chain with mass number $A = 133$ are being redetermined.
Method:	Radiochemical and by mass-spectrometrv
Completion date:	1979
Publications:	Jahresbericht 1977 Institut für Kernchemie Universität Mainz

Laboratory	Physikalisch-Technische Bundesanstalt
and address:	D-3300 Braunschweig, Bundesallee 100
Name: Facilities:	K. Debertin ²⁵² Cf-source;
Experiment:	Determination of 238 U-fission yields in the fast neutron spectrum of a 252 Cf- source.
Method:	The 252 Cf-source is mounted 15 m above ground in the open air. 238 U-samples are irradiated in a 1 cm distance. Fission product activities are determined. by measuring the γ -ray spectrum with a calibrated Ge(Li)-spectrometer.
Accuracy:	<u>+</u> 2 % to <u>+</u> 5 % (1 σ uncertainty)
Completion date:	1978
Publication:	Proceedings of the International Conference on Neutron Physics and Nuclear Data for Reactors and Other Applied Purposes, Harwell, 1978, page 229.

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Laboratory and	Address: Badiochemistry Division Bhabha Atomic Research Centre Trombay, Bombay 400085, India
Names: B	lamaswami A, Naterajan V., Sampath Kumar R., and R.H. Iyer.
Facilities : H	Ligh Resolution Ge(Li) detector multichannel analyser, Solid State track detector and Optical Microscope.
Experiment : M	Leasurement of the absolute fission yields in the neutron induced fission of actinide isotopes.
Method : A r i a s t	Absolute yields - Short-lived fission products (half lives ranging from few minutes to few days) in the thermal neutron induced fission of ²³⁵ U, ²³⁹ Fu and ²³³ U and in the fast fission ²³² Th are being determined using high resolution gamma ray expectrometry. The total number of fissions in the sample was obtained by track etch technique using mica as the erack detector.
Accuracy : 1 Completion: W Data a	to 5% for the asymmetric fission products. Nork on ²³⁵ U and ²³⁹ Pu were completed. Work on other nuclides are in progress.
Remarks : I e e f	In this technique, errors arising out of inarruracies in the stimation of fission cross section due to variation of neutron energy spectrum, flux determination and in the estimation of the fissile atoms are eliminated.
Publications :T References :1	 be published. Lyer, R.H., Sampath Kumar R., and Choudhuri N.K., Nucl. Inst. Methods, <u>115</u>, 23(1974)
3	 Ramaswami A, Natarajan V, Sampath Kumar R., and Iyer R.H. Presented in "Nuclear Physics and Solid State Physics Symposium" December 1978, held in Bombay, India. 'Absolute fission yield measurements by track-etchcum-gamma Spectrometry'. Ramaswami A, Natarajan V, Srivastava B.K., Sampathkumar R., Choudhuri N.K., and Iyer R.H., JINC (in press).

Laboratory & Address	Radiochemistry Division Bhabha Atomic Research Centre Trombay, Bembay 400085	
Names	: S.B. Manehar, P.P. Venkatesan, S.M. Dest Satya Prakash and M.V. Ramaniah	nmukh
Facilities	High Resolution Ga(Li) gamma spactrometric system, beta proportional counters and lo back ground counters	ic ow
Experiment	Determination of fission yield in the neutron induced fission of ²³² U.	
Methed	232 Fission yields in neutron fission of were determined employing direct counting of catchere on Ge(Li) as well as using radiochemical techniques. Yields were determined by comparison method using 235U and 99Mo as internal standard.	J
Accuracy	: About 4-8% for asymmetric masses 12-15% for symmetric masses	
Completion date	: Already completed	
Publications	: To appear shortly in Phy. Rev. C.	

Laboratory & Address	:	Radiochemistry Division Bhabha Atomic Research Centre Trombay, Bombay 400 085.
Names	:	S.A. Chitambar, S.N. Acharya H.C. Jain, C.K. Mathews and M.V. Ramaniah.
Facilities	t	CH-5 Mass spectrometer with thermoionic source assembly.
Experiment	:	Determination of fission yield in thermal neutron induced fission of 233U, 235U, 239pu and 241pu.
Method	t	Fission yields in thermal neutron induced fission of $233_{\rm U}$, $235_{\rm U}$, $239_{\rm Pu}$ and $241_{\rm Pu}$ have been determined for about 20 mass nos.in each of the fissioning system by employing mass spectrometric techniques for the determination of relative yields.
Accuracy	:	About 2-3 percent for asymmetric masses.
Completion date	:	March 1980.

Laboratory & Address	: Radiochemistry Division Bhabha Atomic Research Centre Trombay, Bombay 400085
Names	: S.8. Manchar, Tarun Datta, S.S. Rattan, Satya Prakash and M.V. Ramaniah
Facilities	: High resolution Ge(Li) gamma spectrometric system.
Experiment	: Charge Distribution in nuclear fission Determination of fractional cumulative yields of 252 and I in the spontaneous fission of Cf
Method	: Direct catcher foil counting on Ge(Li)
Accuracy	: <u>+</u> 5%/
Completion date	: Already completed
Publication	Physical Review, C, <u>17(</u> i) 188 (1978)

JAPAN

Laboratory and a	ddress: Nuclear Phy Japan Atom Tokai-Mura	vsics II Laboratory, ic Energy Research Institute, , Naka-Gun, Ibaraki-Ken, Japan.
Names:	A. Asami, Y. Nakajin Y. Kawarasaki, Y. Fu T. Yamamoto, M. Sug Y. Kanda, T. Kawano	na, M. Mizumoto, M. Ohkubo, uruta (JAERI) imoto (Tohoku Univ.) (Kyushu Univ.)
Facilities:	120 MeV linac neutro	on TOF spectrometer.
1. Experiment:	Neutron capture cros keV region.	ss section measurements in the
Method:	3500 l liquid scint a resolution of l.9	illator tank at 52 m flight path with to 0.5 nsec/m.
Analysis:	Multiple scattering Monte Carlo) in the Self shielding corre	correction and self shielding (Schmitt, sample and neutron detector. ection (Dresner, Macklin).
(1) Samples:	1 ⁵¹ Eu, ¹⁵³ Eu, Eu. (samples are enriched from ORNL.	Chemical form Eu ₂ O ₃ , Separated isotope d to over 96 % for each isotope, loaned
Accurac	y:	6 to 10 %.
Energy	region:	3 to 100 keV.
Expecte	d completion date:	^{151,153} Eu May 1979. Eu Sep. 1979.
Publicatio	n:	
	A. Asami et al., Ne ments of ^{151,153} Eu a Neutron capture cros	eutron capture cross section measure- and Eu, Topical Conf. of Technique on as section measurements, ORNL, 1978.
	M. Mizumoto et al., of ¹⁵¹ Eu and ¹⁵³ Eu 1	Average neutron capture cross sections From 3 to 100 keV, to be published.
(2) Samples:	^{143,145,146,148} Nd, e Nd ₂ O ₃ in chemical fo	enriched to over 91 % for each isotope, orm, loaned from ORNL.
Energy	region:	5 to 300 keV.
Accurac	y:	8 to 30 %.
Expected	d completion date:	Dec. 1979.
Publication	n:	
	V Nakaiima ot al	Noutron cantuma cross contion

Y. Nakajima et al., Neutron capture cross section measurements of Nd-143, Nd-145, Nd-146 and Nd-148, Int. Conf. on Neutron Physic and Nuclear Data, Harwell., 1978, page 438.

JAPAN

(cont'd)

(3) Samples: 147,149 Sm, enriched to over 97 % for each isotope, Sm₂O₃ in chemical form, loaned from ORNL.

Energy region: 1 eV to 300 keV.

Status: Measurements in progress.

2. Experiment: Neutron resonance parameters.

Method: A 6 Li-glass neutron detector and a Moxon-Rae detector at 47 m flight path.

Analysis: Atta-Harvey code and Monte Carlo code MCRTOF.

(1) Sample: Tb.

Results:	Resonance parameters for 209 levels
	including 50 newly discovered ones
	in the region 3 to 1190 ev.

Feb. 1978.

Completion date:

Publications:

M. Ohkubo, Y. Kawarasaki, Slow neutron resonances in Tb-159, JAERI-M 7545 (1978), also to be published in J. Nucl. Sci. Tech.

(2) Samples: ^{79,81}Br, enriched to over 97 % for each sample, NaBr in chemical form, loaned from ORNL.

Status:

Measurements in progress.

<u>JAPAN</u>

Laboratory and address:	Institute of Atomic Energy, Kyoto University, Uji, Kyoto 611, Japan
Names:	Tomota Nishi, Ichiro Fujiwara and Nobutsugu Imanishi
Facilities:	5 MW research reactor [Research Reactor Institute, Kyoto University]
Experiment:	Cumulative and independent fission-yields of some fission products in the thermal-neutron induced fission of 233 U, 235 U and 239 Pu.
Method:	Radiochemical for fission yields; Instrumental with GeLi detectors.
Accuracy:	Errors range from 7% to 20% with different combinations of the fission products and the fissile isotopes.
[Expected] comple	etion date:)
Publication:	

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Nuclides		Completion date	Publication
128,130,132 _{Sn} , ¹³³ S	b [Cum.]		N.Imanishi, I.Fujiwara and
^{128,130,132} Sb ^{m,g} ,			T.Nishi, Nucl. Phys. <u>A263</u> ,
¹³¹ Sb, ^{131,133} Te ^{m,g}	[Ind.]	Sep. 1975	141(1976)
135 _I	[Cum.]		T.Nishi, I.Fujiwara and
131,133 _I ,			N.Imanishi, Inter. Conf. on Nucl
132,134,136 m,g I	[Ind.]	Dec. 1976	Structure,Tokyo, Sep. 1977
133,135 _{Xe} m,g	[Ind.]	Dec. 1976	ibid.
¹³⁸ Cs ^m ,g	[Ind.]	May. 1978	
90 _{Rb} m,g	[Ind.]	End of 1979	
148 _{Pm} m,g	[Ind.]	End of 1979	

JAPAN

Laboratory and address:	Research Laboratory for Nuclear Reactors, Tokyo Institute of Technology 2-12-1, O-okayama, Meguro-ku, Tokyo
Names:	N. Yamamuro, K. Saito, T. Wada (TIT) Y. Fujita, K. Kobayashi (Research Reactor Institute, Kyoto University)
Facilities:	46-MeV Electron Linear Accelerator (Research Reactor Institute, Kyoto University)
Experiment:	Capture Cross Section Measurements of 93 Nb, 127 I, and 133 Cs from 3 to 80-KeV using time-of-flight method
Method:	Gamma-rays from the neutron capture processes were detected by a C_6F_6 or a C_6D_6 liquid scintillation detector. Neutron flux impinging on the sample was measured by ${}^{10}B$ (93%) disk placed at the sample position. The absolute values of cross section were determined by normalizing to the 24-KeV cross sections measured with Fe-filtered method. Corrections for self-shielding and multiple scattering were performed using average cross sections. These data are currently examined for the correction for resonance self-shielding.
Accuracy:	Error of absolute cross section at 24-KeV is about 5% Statistical error of measured cross sections is 2 to 4%
(Expected) Completion Date:	June, 1979 for 93 Nb and 127 I Sept., 1979 for 133 Cs
Publications:	 N. Yamamuro et al., J. Nucl. Sci. and Technol. <u>15</u> 637 (1978) N. Yamamuro et al., Proc. Inter. Conf. Neutron Physics and Nuclear Data for Reactors and other Applied Purposes AERE Harwell Sept., 1978, page 432.

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JAPAN

(same as in INDC(NDS)-95)

Laboratory and address:	Nuclear Engineering Research Laboratory Faculty of Engineering University of Tokyo
	2-22 Shirane Shirakata, Tokai-mura Naka-gun Ibaraki, Japan

Names: M. Akiyama and S. An

Facilities: Fast Neutron Source Reactor "YAYOI"

Experiment: Fission Product Decay Heat for Fast-Neutron Fission of 235 U, 238 U, Nat.U. and 232 Th for Cooling Times of 10^3 to 10^6 secs.

Method: Samples have been irradiated for 5 to 10 min. with fast neutrons. Gamma-ray energy spectra have been measured using NaI detector and beta-ray energy spectra have been obtained using plastic scintillation detector combined with $\Delta E/\Delta x$ type proportional counter to eliminate gamma-ray effects. Counting times have been chosen to provide good statistics within the time range of interest. Total energy release rates for beta and gamma-rays have been obtained to integrated beta and gamma-energy spectra respectively and summed to obtain the fission product decay heat.

The same experimental program has been planed for the more short cooling times using the fast pneumatic-tube irradiation facility and other fissile materials (239 Pu and 233 U).

Accuracy:	<10% (expected)		
(Expected) Completion Date:	December 1978 for the more short cooling times August 1979 for ²³⁹ Pu and ²³³ U.		
Discrepancies to other Reported Data:	Preliminary Data are in good agreement with other recent experiments.		
Publications:	M. Akiyama et al., "Kyodo Riyo Seika Hokoku-sho at Nuclear Engineering Research Laboratory U. of Tokyo, 1976, 1977.		

SWEDEN

Laboratories:	Department of Nuclear Chemistry Chalmers University of Technology S-412 96 GÖTEBORG Sweden
	Department of Nuclear Chemistry University of Oslo Oslo 3 Norway
	Institut für Kernchemie Johannes Gutenberg Universität Postfach 3980 D-6500 MAINZ Germany
Names:	The SISAK Collaboration:
	G. Skarnemark and K. Brodén (Göteborg)
	D. Eriksen and I. Haldorsen (Oslo)
	N. Kaffrell, J. Stachel and N. Trautmann (Mainz)
Facilities:	Two SISAK systems for studies of radionuclides with half-lives >0.5 s.
Experiments:	$T_{1/2}^{-determinations, \gamma-singles, time-\gamma-\gamma}$ coincidence, $\gamma-\gamma$ angular correlation and β -radiation measurements, at present on $^{106-108}$ Tc.
Method:	Fast chemical on-line separations. The γ -measurements
	are carried out on flow cells or ion exchange columns. Ge(Li)-detectors are used.
Discrepancies to other data:	There are very few data available in this region.

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SWEDEN

	Laboratory and address:	The Studsvik Science Research Laboratory S-611 82 Nyköping, Sweden
	Facility:	The OSIRIS on-line separator has been used to extract selected nuclei from thermally fissioned 235U.
1.	Names:	K Aleklett, O Glomset, P Hoff, E Lund and G Rudstam
	Experiment:	Delayed neutron emission probabilities, P_n -values have been deduced for the precursors: $79-83_{Ga}$, $87-89_{Br}$, $92-96_{Rb}$, $127-132_{In}$, $134,135_{Sb}$, $137-139_{I}$ and $141-145_{Cs}$.
	Method:	Neutron and beta activities are measured simul- taneously in MCA mode. The neutron counter con- sists of 2.9 ³ He counters imbedded in parafine and the beta particles are detected with a 2 mm plastic detector. A Ge(Li) detector monitors the source purity. The calibration of the detectors have been performed most carefully and the P_n - values have been determined with an accuracy of typically 5 %.
	Completion date:	Completed and ready for publication.
2.	Names:	K Aleklett, E Lund and G Rudstam
	Experiment:	Total beta decay energies and atomic masses have been deduced for the following nuclides: $75-78_{Zn}$, $76-79_{Ga}$, $79,80_{Ge}$, $80,81,83_{As}$, $85-89_{Br}$, $120-129_{In}$, $127,132_{Sn}$, $128,130-132,134_{Sb}$ and $134,135_{Te}$.
	Method:	Beta particles were recorded in coincidence with gamma rays depopulating known levels in the daughter nucleus. The end-point energies of the beta- spectra were determined, and by adding the level energy the total beta-decay energies were obtained. The beta-particles were recorded in a Si(Li) de- tector and the gamma-rays in a Ge Li) detector or two NaI (T1) crystals. In the future the experi- mental arrangement will be improved by using two Ge (Li) detectors for the gamma registration.
	Publications:	K Aleklett, E Lund, G Rudstam, Nucl Phys A <u>281</u> , 213 (1977), K Aleklett, E Lund, G Nyman, G Rudstam, Nucl Phys <u>A285</u> , 1 (1977) E Lund, K Aleklett, G Rudstam, Nucl Phys <u>A286</u> , 403 (1977) K Aleklett, E Lund and G Rudstam, Phys Rev <u>C18</u> , 462 (1978) K Aleklett, E Lund and G Rudstam, Total β-decay energies and masses of ⁸⁵⁻⁸⁹ Br, to be published in Z Physik A.
	Completion date:	Indefinite

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SWEDEN

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3. Name: P Hoff Nuclear spectroscopic studies of the beta-decays of 88Br and 138I for determination of Experiment: decays of 05Br and 1351 for determination of delayed neutron emission to excited states in 87 Kr and 137 Xe respectively, are completed. Si-milar studies of the decays of 81 Ga, 89 Br, $^{93-96}$ Rb, 135 Sb and 139 I, complemented with theoretical calculations of the delayed neutron branching ratios, are under way. Absolute γ -ray intensities have been deduced for the decays. Publications: P Hoff, Levels of ¹³⁷Xe and ¹³⁸Xe populated in the decay of ¹³⁸I, J Inorg Nucl Chem (in press). P Hoff, The decay of ⁸⁸Br, Physica Scripta (in press). B Fogelberg, H Tovedal 4. Names: Nuclear spectroscopic studies of short lived Experiment fission product nuclei by means of y-ray and conversion electron spectroscopy and measurements of level half-lives. Studies of the decays of even mass isotopes of In with A = 120 - 128 were completed during 1978. A project consisting of comprehensive studies of the decays of ¹³⁷I and ⁸⁷Br is expected to be completed during 1979. Completion Indefinite for the nuclear spectroscopy project date: as such. B Fogelberg and P Carlé, Levels and transition probabilities in 120,122,124,126,128Sn studied Publication: in the decay of In isotopes, Nucl Phys (in press). G Rudstam and K Aleklett 5.Names: Experimental determinations of the average ra-Experiment: diation energies of individual fission products. The measurements of average beta energies are completed and the experimental determinations of average gamma energies are in progress. The average beta and gamma energies in the decay Evaluation: of 382 known fission products have been determined. As far as possible the values are based on experi-mental data direct determinations, published decay schemes, and a study of beta strength functions. In cases for which no experimental data exist the average energies have been derived using extrapolated beta strength function. G Rudstam and K Aleklett, The energy distri-**Publications:** bution of antineutrinos originating from the decay of fission products in a nuclear reactor, The Studsvik Science Research Laboratory Report NFL-2 (1978). G Rudstam and K Aleklett, Average beta and gamma energies of fission products, The Studsvik Science Research Laboratory Report NFL-7 (1979). G Rudstam, Detailed comparison between decay heat data calculated by the summation method and integral measurements, The Studsvik Science Research Laboratory Report NFL-9 (1979).

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SWEDEN

Laboratory and address:	Neutron Physics Laboratory AB Atomenergi Fack S-611 01 NYKÖPING Sweden
Names:	P-I Johansson and G Nilsson
Facilities:	6 MeV VdG accelerator PDP-15 Computer 24 k memory (on line) NaI(T1) and Ge(Li) spectrometers, β-spectrometer CDC-CYBER 73 omputer (off line)
Experiment:	The objective of the experiment is to improve on the accuracy of currently available fission pro- duct decay heat data by means of radiometric study of small uranium specimens at cooling times longer than 3 seconds after irradiation with thermal neutrons.
	The residual power of gamma and beta radiation from thermal fission of 2^{35} U has been obtained with an accuracy of \pm 7 % in the time interval 10 sec to 25 min after fission. Measurements are in progress for studying also the residual power due to β -emission from 2^{39} Pu.
Method:	A facility for thermal neutron irradiation of fissile specimens using a VdG accelerator has been built. Specimens are transported between the neutron source and a spectrometer by means of a pneumatic system.
	The absolute number of fissions in the sample is determined by two independent methods: a) by uti- lizing an absolutely calibrated fission chamber with an active volume of about the same size as the samples, b) by counting the number of gamma quanta emitted from fission products with known yields and decay properties.
	The beta radiation is measured with Si(Li) crys- tal. Sample transportation, irradiation and counting times are handled by a PDP-15 computer. Spectra are automatically stored on magnetic tape for off-line data analysis, i.e. the trans- formation from measured pulse height spectra to energy spectra.
Accuracy:	Better accuracy than \pm 10 % is expected for the total energy released as β or γ -radiation from the fission products at any time between a few seconds and 30 minutes after fission.
Completion date:	Measurements on gamma-radiation from 235 U were completed in December 1976 and the study on beta emission were completed in March this year. The measurement on 239 Pn will be completed in Octo- ber 1979.

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SWITZERLAND

Laboratory and address: Eidg. Institut für Reaktorforschung, CH-5303 Würenlingen Institut für anorganische, analytische und physikalische Chemie, Universität Bern, CH-3012 Bern, Switzerland H.R. von Gunten Name: Facility: Swimming-pool type reactor (SAPHIR) Determination of independent and cumula-Experiments: tive yields in the fission of 232 Th, 233 U, 235 U, 239 Pu, and other nuclides. Absolute yields in reactor neutron fission of 232 Th. Independent yields of 148m Pm and 148g Pm (thermal neutron fission of 233 U and 239 Pu). Radiochemical and instrumental (GeLi). Method: 5 - 10 % Accuracy: Measurements completed: $\begin{vmatrix} 148 \\ \text{Pm} \\ 232 \\ \text{Th} \\ 148 \\ \text{Pm} \\ \text{completed}, \text{ report in preparation} \\ \text{completion date uncertain} \end{vmatrix}$ - T. Kaiser and H.R. von Gunten, Indepen-dent yields of ⁹⁶Nb and 51 min ⁹⁸Nb in Publications: the thermal neutron-induced fission of $23_{3_{\rm U}}$, $23_{\rm U}$ and 23_{9} Pu, J. Inorg. Nucl. Chem. <u>40</u>, 377 (1978). - T. Kaiser and H.R. von Gunten, Cumulative mass yields in the neutron-induced fission of 239 Pu at the resonance energy of 0.3 eV. Phys. Rev. Cl7, 1510 (1978). - H. Gäggeler and H.R. von Gunten, Charge distribution in the photofission of $238_{\rm U}$ with Bremsstrahlung of E 20-50 MeV. J. Inorg. Nucl. Chem. 40^{max} , 1859 (1978).

Laboratory and Address: AEE Winfrith

UKAEA Atomic Energy Establishment Winfrith Dorchester, Dorset DT2 8DH

Names:

M. F. Murphy, W. H. Taylor

Experiment: Measurement of gross beta-decay power from products of ²³⁹Pu and ²³⁵U fission in a fast reactor. Irradiation period 10⁵ seconds, detection continued up to 3.10⁷ seconds after shutdown. Experiment completed.

Method: Thin deposits of ²³⁹Pu and ²³⁵U irradiated with catcher foils at centre of Zebra core with neutron energy spectrum close to that of fast power reactor. Fissions monitored by absolute (Alpha-calibrated) counters. Catcher foils transferred rapidly to scintillation detector, current output from photomultiplier being measure of beta power. Calibrated using standard Sr-90/Y-90 source. Results of various subsidiary experiments have confirmed the validity of the experimental method and determined the necessary corrections.

Accuracy: Target accuracy is \pm 7% (standard error) on absolute beta power as function of time from 30 seconds to 1 year after irradiation.

Completion Date:

Complete.

Publication:

An unclassified Winfrith report (AEEW-R1212) has been published.

Laboratory and Address	AERE Harwell	UKAEA, AERE, Harwell, Oxfordshire OX11 ORA U.K.
Names:	J. G. Cuninghame, H. H. Wi	llis
Facilities:	ZEBRA - BIZET	
Experiment:	To measure the effect of c neutron spectrum on fissic	hange of reactor on yields.
Method:	Four irradiations, each of ²³⁸ U and two ²³⁹ Pu metal b 100mg weight have been mad inner core and two in the samples of each of the fis counted directly on a cali detector while the other used to prepare purified s fission products of very 1 As, Ag, Cd, Sn, Sb and Rar More irradiations will hav out and we are still some able to determine the numb absolutely but preliminary promising.	two ²³⁵ U, two peads of approx. The; two were in the outer. One of the sile materials was brated Ge(Li) was dissolved and samples of certain ow yield, viz. The Earths. The to be carried way from being pers of fissions results are
Accuracy:	Expected <u>+</u> 10%	
Completion date:	Expected end - 1979	

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Laboratory and Address:	AERE Harwell	UKAEA AERE, Harwell Oxfordshire, OX11 ORA
Names:	I. C. McKean and E. A. C. C.	couch
Experiment:	³ H yield in thermal and fast U and Pu isotopes	t fission spectra for
Facilities:	GLEEP and 'ZEBRA' Reactors	
Method:	The tritium produced in fiss tritiated water, separated f products and measured by lic counting. A preliminary exp completed in which solutions irradiated in a thermal flux irradiated in GLEEP (235 U + Irradiations in ZEBRA-BIZET are planned for mid - 1979.	sion is converted to from other fission quid scintillation periment has been s of 235 U were s. Samples have been 239 Pu in solution). (235 U + 239 Pu metal)

Accuracy:

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<u>+</u> 10%

(same as in INDC(NDS)-95)

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Laboratory and Address:	DNPDE	Dounreay Nuclear Power Development Establishment, UKAEA, Northern Division, Thurso, Caithness, Scotland KW14 7TZ
Names:	W. Davies, V.M. Sinclair	
Facilities:	PFR	
Experiment:	The measurement of the absolute yields of 90Sr, 137Cs, 144Ce, 143,145,146,148,150Nd and perhaps other fission products, from the fission of 235U, 238U, 239Pu, 240Pu and 241Pu	
	In progress	
Method:	Twelve sealed stainless s irradiated. Of these,	teel capsules are to be
	3 capsules contain ²³⁵ U a dioxide, 3 capsules contain ²³⁹ Pu plutonium dioxide, 2 capsules contain ²³⁸ U a with an isotopic analysis 1 capsule contains ²⁴⁰ Pu of plutonium with an isot 1 capsule contains ²⁴¹ Pu of plutonium with an isot 2 capsules contain no add The ²³⁵ U and ²³⁹ Pu capsul powder mixed with the fis heat transfer reasons. It is expected that the ² receive irradiation corre up of the fissile materia 0.7% burn-up, the ²⁴⁰ Pu c and the ²⁴¹ Pu capsule to	s highly enriched uranium as low ²⁴⁰ Pu content s depleted uranium dioxide of 99.7% ²³⁰ U, as a dried aqueous solution opic analysis of 99% ²⁴⁰ Pu, as a dried aqueous solution opic analysis of 93% ²⁴¹ Pu, and ed fissile material. es contain stainless-steel sile material dioxide for ^{35U} and ²³⁹ Pu capsules will sponding to about 16% burn- 1, the ²³⁰ U capsule to about apsule to about 4% burn-up about 23% burn-up.
	A set of capsules identic except for irradiation in and analysed alongside th being to improve the reli	al to the irradiated set the reactor will be dissolved is irradiated set, the objective ability of the analyses.
	The aim is to correlate 1 irradiation with the amou formed, for each capsule, absolute measurements of obtained.	oss of fissile material during nts of fission products (except ²³⁸ U) to enable fission yields to be
Accuracy:	⁺ 2% for 2350 and 23 ⁺ 6% for 2380, 240Pu	99Pu fission yields a and ²⁴¹ Pu fission yields
Expected completion date:	1980	

(same as in INDC(NDS)-95)

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Laboratory and Address	N.P.L. Teddington	Division of Radiation Science and Acoustics National Physical Laboratory Teddington Middlesex TW11 OLW	
Names:	P. Christmas, P. Cross	;	
Facilities:	π√ 2 Beta-ray spectrome	$\pi\sqrt{2}$ Beta-ray spectrometer, isotope separator	
Experiment:	Determination of K & L coefficients of ^{133m} Xe	internal conversion	
Method:	Peak-to-Beta-Spectrum (PBS). The study of instrumental distortions of beta-spectra in the $\pi\sqrt{2}$ spectrometer has been completed and submitted for publication, together with a paper describing the application of the PBS method to ¹³⁷ Cs. The results quoted in the latter include precise values for the internal conversion coefficients and gamma-ray intensity per disintegration for the 662 KeV transition.		
Accuracy:	Target is <u>+</u> 1 per cent	•	
Completion Date:	Uncertain, due to lack	of effort.	

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Laboratory and address:	Birmingham Radiation Centre	University of Birmingham P.O. Box 363 Birmingham B15 2TT United Kingdom
Names:	J.G. Owen, J. Walker, D.R. Weaver	
Facilities:	3MV Dynamitron accelerator	
Experiment:	Delayed neutron spectrum measurements following mono- energetic fast neutron induced fission in 235 U	
Method:	³ He spectrometer; cyclic irradiation and counting to give near equilibrium contributions from all delayed neutron groups	
Accuracy:	Uncertainties on response function and efficiency measurements of the ³ He spectrometer are estimated to give rise to + 5% error on the final spectra. When combined with statistical errors in the delayed neutron counts the overall error is estimated as + 7% in the range 0.1-1MeV. Above 1MeV an uncertainty of 15-20% is expected. Below 0.1MeV corrections due to scattering are less certain and work is in hand to assess their influence on the accuracy	
Completion date:	Measurements at 0.94, 1.44, 1.76 a complete. Further measurements a with other fissioning nuclides are	and 6.0MeV are at other energies and e in progress
Discrepancies with other reported data:	This work shows significant number above 1MeV as does the measurement This is in marked disagreement with with proton recoil spectrometers Woodruff) which show very little H At delayed neutron energies less shows considerable emission as doe and Woodruff; the spectral shapes however.	rs of delayed neutrons t of Evans and Krick. th measurement made (e.g. Eccleston and high energy contribution. than 0.1MeV this work es that of Eccleston s are very different,
Publications:	Owen J.G., Walker J., Weaver D.R. 'Energy spectra of delayed neutron Proceedings of the International (Data and Nuclear Data for Reactors 1978. OECD Nuclear Energy Agency	ns from uranium fission'. Conference on Neutron s, Harwell, September y (1978) 606-609.
	Walker J., Weaver D.R., Owen J.G. 'Delayed neutron measurements with accelerator'. Proceedings of the the Application of Small Accelerat 1978.	n a 3MV Dynamitron e Fifth Conference on tors, Texas, November

United Kingdom

Laboratory and address	:	University of London Reactor Centre, Silwood Park, Sunninghill, Ascot, Berkshire, SL5 7PY.	
Names	:	B.J. Olomo, T.D. Mac Mahon.	
Facilities	:	$4\pi\beta-\gamma$ coincidence system, Ge(Li) detectors.	
Experiment	:	Measurement of absolute gamma-ray emission probabilities in fission products.	
Method	:	Thin sources prepared from solutions of the radioisotope. Total disintegration rate measured in $4\pi\beta$ - γ coincidence system. Gamma-ray emission rates determined using calibrated Ge(Li) detectors. Combination of results gives number of gamma rays emitted per disintegration.	
Accuracy	:	Target accuracy is 1% (1 σ), accuracies achieved within range 0.8 - 1.2%, limited mainly by efficiency calibration of Ge(Li) detectors.	
Completion date :		¹⁴⁴ Ce/ ¹⁴⁴ Pr : completed early 1979.	
		¹⁴⁰ Ba/ ¹⁴⁰ La : Aug/Sept 1979.	
Results	:	$E_{\gamma}(keV)$ $I_{\gamma}(\%)$ $I_{\gamma}(\%)$ Ref 1	
		¹⁴⁴ Ce 134 10.69 ± 0.12 11.09 ± 0.16	
		¹⁴⁴ Pr 696 1.484 \pm 0.012 1.342 \pm 0.013	
		¹⁴⁴ Pr 1489 0.276 ± 0.002 0.279 ± 0.003	
		¹⁴⁴ Pr 2186 0.786 \pm 0.009 0.700 \pm 0.010	
		The above results have not yet been published.	
Reference 1	:	K. Debertin et al. Ann. Nucl. Ener. 2, 37-43 (1974).	

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Laboratory and address:	Argonne National Laboratory 9700 South Cass Avenue Argonne, Illinois 60439 USA
Names:	L. E. Glendenin, J. E. Gindler, J. W. Meadows
Facilities:	Fast-neutron generator facility (FNGF)
Experiment:	Determination of fission yields for mono- energetic neutron-induced fission as a function of incident neutron energy over the range 0.1 to 8 MeV.
Method:	Yields determined (1) radiochemically with either β - or γ -counting and (2) by γ -counting irradiated foils of fissionable material. Neutrons produced by Li-p or D-d reaction. Flux monitored with fission chamber utilizing as the fission source the same material as that being irradiated. Absolute yields determined from flux measurements and/or 200% normalization of mass-yield distribution.
Accuracy:	Yields > 1% determined by γ -counting: 3-5% Yields < 1% determined by γ -counting: 5-20% Yields determined radiochemically with β -counting: 10-20%
Completion date:	Measurements of ²³⁸ U(n,f) for 1.5, 2.0, 3.9, 5.5, 6.9, and 7.7 MeV neutrons (published Jan. 1978). ²³² Th(n,f) in progress; work to be completed by mid-1979. Continuing program for other fissile and fertile nuclides.
Publications:	"Mass distributions in monoenergetic-neutron- induced fission of ²³⁸ U", S. Nagy, K. F. Flynn, J. E. Gindler, J. W. Meadows, and L. E. Glendenin, Phys. Rev. <u>C17</u> , 163 (1978).
	"Fission yields for fast-neutron fission of uranium-238", J. E. Gindler, L. E. Glendenin, J. W. Meadows, and K. F. Flynn, Nucl. Sci. Eng. <u>70</u> , 101 (1979).

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Laboratory and Address

Argonne National Laboratory 9700 South Cass Avenue Argonne, Illinois USA

Names

A. Smith, P. Guenther, G. Winkler* and J. Whalen

Facilities

Monoenergetic pulsed-beam accelerator

Experiment

Fast-Neutron Total and Scattering Cross Sections of ¹⁰⁷Ag in the MeV Region

Neutron total cross sections are measured from 0.25 to 4.5 MeV at intervals of $\sqrt{10}$ keV. Neutron differential elastic- and inelastic-scattering cross sections are measured from 1.5 to 4.0 MeV at intervals of ≤ 0.2 MeV. Cross sections for scattering into more than 20 energy groups are determined. Cross sections calculated from an optical-statistical model are in quantitative agreement with measured neutron inelastic-scattering cross sections. In the context of this model, significant dependence of the inelastic-scattering process on parity and/or deformation is not in evidence. The interpretation of the observed neutron inelastic-neutron-scattering results is consistent with previous reported J^{T} assignments and the systematics of nuclear-level-densities.

Publication

Argonne National Laboratory Report, ANL/NDM-46 (1979)

*Permanent address: Institut fuer Radiumforschung und Kernphysik, A-1090 Vienna, Boltzmanngasse 3, Austria.

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FPND NEWSLETTER CONTRIBUTION - I

Laboratory:Idaho National Engineering LaboratoryAddress:Allied Chemical Corporation
550 Second Street
Idaho Falls, Idaho 83401
United StatesName:William J. Maeck

Experiment: Fast Reactor Fission Yields and Determination of Burnup
for Fast Reactor Fuels

A program is in progress at the Idaho Chemical Processing Plant (ICPP) laboratories to accurately measure absolute fast reactor fission yields for ²³³U, ²³⁵U, ²³⁸U, ²³⁷Np, ²³⁹Pu, ²⁴⁰Pu, ²⁴¹Pu, ²⁴²Pu, ²⁴¹Am, and ²⁴³Am. The irradiations were conducted in EBR-II, Row-8. The analytical measurements have been completed for all samples except ²⁴³Am. Reports have been issued giving fast reactor yield data for ²³³U, ²³⁵U, ²³⁸U, ²³⁷Np, ²³⁹Pu, ²⁴⁰Pu, ²⁴¹Pu, ²³⁵U, ²³⁸U, ²³⁷Np, ²³⁹Pu, ²⁴⁰Pu, ²⁴¹Pu, and ²⁴²Pu.

<u>Method</u>: The principal measurement technique is isotope dilution mass spectrometry for the isotopes of Kr, Rb, Sr, Zr, Mo, Ru, Xe, Ce, Ba, La, Ce, Nd, and Sm. The number of fissions is established by the summation of the total atoms in the heavy mass peak.

<u>Accuracy</u>: In general, the uncertainties associated with 233 U, 235 U, 237 Np, 239 Pu, 241 Pu, and 242 Pu yields range from 1.0-1.5% relative, and for 238 U and 240 Pu yields, the uncertainties range from 1.5-3.0% relative.

<u>Future Work:</u> Data reduction, statistical analysis and calculation of the fast fission yields for 241 Am is in progress.

To augment the fast yield data obtained from the irradiation in Row-8 of EBR-II, samples of 233 U, 235 U, 239 Pu, and 241 Pu which were irradiated in Row-2 of EBR-II are being analyzed. These data will provide more

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information relative to the effect of neutron energy on fission yields. The analysis of the 233 U and 235 U samples has been completed and analysis of the 239 Pu and 241 Pu samples is in progress.

The fabrication of samples for irradiation in the Fast Flux Test Facility (FFTF) at Hanford, Washington, is in progress. These samples will be irradiated in late 1979 and will be used to provide FFTF benchmark fast yield data for ²³³U, ²³⁵U, ²³⁹Pu, and ²⁴¹Pu.

<u>Special Comments</u>: All yield data reported from this work are associated with a measured or calculated neutron spectrum. The study to correlate yields with neutron energy is continuing.

<u>Publications</u>: The results of this measurement program are published in a series of Allied Chemical Corporation - Idaho Chemical Programs reports designated ICP-1050-I, II, III, etc. The following reports are available from the National Technical Information Service, U.S. Dept. of Commerce, 5285 Port Royal Road, Springfield, Virginia, 22161, USA.

- W. J. Maeck, Editor, "Fast Reactor Fission Yields for ²³³U, ²³⁵U, ²³⁸U, ²³⁹Pu and Recommendations for the Determination of Burnup on FBR Mixed-Oxide Fuels: An Interim Project Report," Allied Chemical Corporation -Idaho Chemical Programs Rept., ICP-1050-I (January 1975).
- W. J. Maeck, Editor, "Fast Reactor Fission Yields for ²³⁹Pu and ²⁴¹Pu," Allied Chemical Corporation - Idaho Chemical Programs Rept., ICP-1050-II (August 1977).
- W. J. Maeck, W. A. Emel, A. L. Erikson, J. E. Delmore, J. W. Meteer, "Fast Reactor Fission Yields for ²³⁷Np," Allied Chemical Corporation -Idaho Chemical Programs Rept., ICP-1050-III (September 1977).
- W. J. Maeck, R. L. Eggleston, A. L. Erikson, R. L. Tromp, "Fast Reactor Fission Yields for ²⁴⁰Pu and ²⁴²Pu," Allied Chemical Corporation -Idaho Chemical Programs Rept., ICP-1050-IV (February 1979).

FPND NEWSLETTER CONTRIBUTION - II

Laboratory:Idaho National Engineering LaboratoryAddress:Allied Chemical Corporation
550 Second Street
Idaho Falls, Idaho 83401
United StatesName:William J. Maeck
Experiment:Experiment:Thermal Fission Yields for 235U and 239Pu

An existing experiment has been extended to remeasure the major fraction of the mass yield curve for the thermal fission of ^{235}U and ^{239}Pu . The need for this program resulted from serious discrepancies in some measured relative isotopic ratios for certain isotopes in the thermal fission of ^{239}Pu . New yield values, based on the analysis of six ^{235}U samples (three at

] 1 a/oF and three at \sim 35 a/oF) have been published.

<u>Method</u>: The principal measurement technique is isotope dilution mass spectrometry for the isotopes of Kr, Rb, Sr, Zr, Mo, Ru, Xe, Cs, Ba, La, Ce, Nd, and Sm. For the ²³⁵U samples, the number of fissions was established by two methods: 1) the summation of the total atoms in the heavy mass peak, and 2) the heavy element difference technique.

<u>Accuracy</u>: The uncertainties associated with these new 235 U yields range from 0.5-1.5% relative. A comparison of the number of fission determined by the summation technique and the heavy element difference technique show an agreement of $\pm 0.5\%$ relative.

<u>Future Work</u>: Analysis of six 239 Pu samples (three at 1 a/oF and three at \sim 40 a/oF) has been completed and data reduction is in progress. A final report giving new 239 Pu thermal yield data will be issued late in 1979.

Data Discrepancies: See Publication 1.

Publications:

- W. J. Maeck, W. A. Emel, J. E. Delmore, F. A. Duce, L. L. Dickerson, J. H. Keller, R. L. Tromp, "Discrepancies and Comments Regarding ²³⁵U and ²³⁹Pu Thermal Fission Yields and the Use of ¹⁴⁸Nd as a Burnup Monitor," Allied Chemical Corporation - Idaho Chemical Programs Rept., ICP-1092 (December 1976).
- W. J. Maeck, W. A. Emel, F. A. Duce, R. L. Tromp, J. W. Meteer, "Absolute Thermal Fission Yields for ²³⁵U," Allied Chemical Corporation - Idaho Chemical Programs Rept., ICP-1142 (September 1978).

FPND NEWSLETTER CONTRIBUTION - III

Laboratory:	Idaho National Engineering Laboratory			
Address:	Allied Chemical Corporation 550 Second Street Idaho Falls, Idaho 83401 United States			
Name:	William J. Maeck			
Experiment:	Natural Fission Reactor Studies: ²³⁸ U Spontaneous Fission Yields			

In the process of analyzing approximately 25 rich uranium ore samples for fissiogenic ruthenium, a preliminary estimate of the 238 U spontaneous fission yields for 99 Ru, 101 Ru, 102 Ru, and 104 Ru has been obtained. The measurement technique was mass spectrometry.

After correction for the natural Ru component and the fissiogenic component resulting from 235 U induced fission, the best estimate for the isotopic composition of 238 U spontaneous fission Ru is:

99	0.236
101	0.285
102	0.314
04	0.165

Using a value of 6.0% for the 238 U spontaneous fission yield of 99 Mo, the preliminary 238 U spontaneous fission yields for the Ru isotopes are:

⁹⁹ Ru	6.0%	(relative	to	⁹⁹ Mo)
¹⁰¹ Ru	7.25			
^{l 02} Ru	8.0			
^{L04} Ru	4.2			

These values have been updated and supersede those reported in INDC (NDS)-86.

A new ore sample, believed to have a very high fraction of spontaneous fission is being hi-graded and will be analyzed to give improved ^{238}U spontaneous fission data for ruthenium. A new mass spectrometric measurement procedure for ruthenium has been developed which gives more reliable data on the 10-25 ng level.

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Laboratory and address:	daho Nati G&G Idaho . O. Box daho Fall .S.A.	ional Engineering Laboratory o, Inc. 1625 s, Idaho 83401
Names:	. D. Hark	ker, R. A. Anderl
Experiment:	ntegral c eactor ty	cross-section measurements in fast- vpe environments.
<u>Method</u> :	ultiple s eodymium, ⁴³ Nd, ¹⁴⁴ ⁺⁹ Sm, ¹⁵¹ iated in osimetry oth at th eflector. ere deter sing Ge(L ode was u hape info eaction r easured u esults ar roup cross echniques	samples of highly-enriched isotopes of samarium, and europium, specifically, Nd, ¹⁴⁵ Nd, ¹⁴⁶ Nd, ¹⁴⁸ Nd, ¹⁵⁰ Nd, ¹⁴⁷ Sm, Eu, ¹⁵² Eu, ¹⁵³ Eu, and ¹⁵⁴ Eu were irra- a row 8 position of EBR-II. Samples and sets of spectrum monitors were located he reactor midplane and in the axial Reaction rates for the spectrum monitors rmined by the gamma-spectrometric technique (i) spectrometers. A spectrum-unfolding used to obtain flux and neutron-spectral prmation from the monitor reaction rates. rates for the fission product samples were using mass-spectrometer techniques. The re used in the integral testing of multi- s sections by least squares adjustment
Accuracy:	% to 10%	
Measurements Completed:	ntegral c ⁺⁴ Nd, ¹⁴⁵ ⁵⁴ Eu are	apture reaction rates obtained for ¹⁴³ Nd Nd, ¹⁴⁷ Sm, ¹⁴⁹ Sm, ¹⁵¹ Eu, ¹⁵² Eu, ¹⁵³ Eu and summarized in Table 1.
<u>Completion Date</u> :	dditional ay to imp nese meas ne neutro ill be co	mass spectrometry measurements are under- prove the accuracy of the Eu reaction rates. surements, a detailed characterization of on environment and integral data testing ompleted by the end of 1979.
Publications:	. A. Ande sotopes c rans. Am.	erl <u>et al</u> , "EBR-II Irradiation of enriched of Neodymium, Samarium, and Europium," Nucl. Soc. <u>28</u> , 745 (1978).

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(cont'd)

TABLE I

INTEGRAL CAPTURE REACTION RATES MEASURED FOR ISOTOPES OF Nd, Sm AND Eu IRRADIATED IN A ROW 8 POSITION OF EBR-II.

Isotope	Axial ^a	Reaction Rate ^b	Reaction Rate (Fission Product)
	Position	(rps/atom) x 10 ¹⁰	Reaction Rate (²³⁵ U(n,f))
¹⁴³ Nd	+3.8	5.026(1.1)	.215(3.9)
	+38.8	8.746(.6)	.380(3.6)
144 Nd	+3.8	1.060(2.0)	.0459(3.8)
	+38.8	.993(1.1)	.0431(3.7)
¹⁴⁵ Nd	+3.8	7.645(1.0)	.327(3.8)
	+38.8	14.55(.6)	.632(3.6)
¹⁴⁷ Sm	-3.8	23.11(.6)	.988(3.7)
	+31.2	44.81(1.2)	1.88(3.8)
¹⁴⁹ Sm	-3.8	40.90(2.4)	1.75(4.4)
	+31.2	81.51(3.3)	3.43(4.8)
¹⁵¹ Eu	+3.8	54.0(6.0)	2.34(6.8
	+38.8	112.9(6.0)	4.90(6.9)
¹⁵² Eu	-3.8	51.6(10.0)	2.23(11.0)
	+31.2	74.6(10.0)	3.14(11.0)
¹⁵³ Eu	+3.8	29.6(5.0)	1.28(5.9)
	+38.8	61.9(4.0)	2.69(5.3)
¹⁵⁴ Eu	-3.8	37.5(8.1)	1.62(8.7)
	+31.2	62.0(8.2)	2.61(9.0)

^a Axial location is the axial distance from reactor midplane at 0 to the axial midpoint of the experiment capsule.

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^b Example, reaction rate for 143 Nd(n, γ) 144 Nd at axial position of +3.8 cm is (5.026 x 10⁻¹⁰ ± 1.1%)rps/atom.

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Idaho National Engineering Laboratory EG&G Idaho, Inc. P. O. Box 1625 Idaho Falls, Idaho 83401 USA
R. J. Gehrke, R. G. Helmer
l) $4\pi \beta$ - γ coincidence counting system 2) Calibrated Ge(Li) spectrometers
Determination of absolute Y-ray emission probabilities for important fission-product isotopes.
The decay rates are determined by the $4\pi \beta - \gamma$ coincidence counting system, which has two separate pulse-processing systems. One system is based on fixed pulse widths. The other is based on variable pulse widths and an overlap coincidence circuit. The dead time of the beta, gamma and coincidence channels is measured by counting the pulses from a 10 MH clock. The variable pulse width system is useful in measuring the γ -ray emission probabilities of short-lived (< 30 m) fission products, where high count rates are needed. The γ -ray emission rates are determined by Ge(Li) spectrometers whose efficiencies have been measured to an accuracy of $\pm 1\frac{1}{2}\%$ (1 σ) between 0.3 and 2 MeV.
\pm 1% to \pm 5% (1 uncertainty)
Emission probability of the prominent 165.9 -keV γ ray from 139 Ba decay has been determined with a precision of \sim 1.1% (1σ level) and the 139 Ba half-life has been remeasured with a precision of \sim 0.3% (1σ level). From this former datum, it is possible to derive more precise emission-probability values for the γ rays from the 139 Xe and 139 Cs parent-members of the A-139 decay chain
Measurement activity is an on-going effort. ¹³⁹ Ba measurement completed in March, 1979. ¹⁴⁵ Pr and ¹⁴⁶ Pr measure- ments to be completed by October, 1979
L. O. Johnson and R. J. Gehrke "A High Rate $4\pi \beta - \gamma$ Coincidence Counting System", IEEE Transactions on Nuclear Science, Vol. NS-26, No. 1, 476 (1979).

U.S.A.

LABORATORY	Lawrence Livermore Laboratory University of California P.O. Box 808 Livermore, CA 94550, U.S.A.	McClellan Central Laboratory 1155th Technical Operations Squadron McClellan AFB, CA 95652, U.S.A.
1. NAMES	D. R. Nethaway A. L. Prindle	W. A. Myers M. V. Kantelo R. L. Osborne
FACILITY	FLATTOP Critical Assembly (Pu), Lo Laboratory.	s Alamos Scientific
EXPERIMENT	Measure fission yields (both total independent yields) for fission of by fission-spectrum neutrons.	chain yields and Pu-240 induced
METHOD	Measurements were made both by doi and by direct counting with Ge(Li) accuracy of the measurements is ab yields are based on a comparison w yields and on a normalization of t	ng chemical separations detectors. The out ± 5%. Absolute ith U-235 fission he mass-yield curve.
COMPLETION DATE	The experiment is finished.	
PUBLICATION	Phys. Rev. C <u>18</u> , 1700 (1978), (Repo	rt UCRL-80020).
2. NAMES	D. R. Nethaway M A. L. Prindle R D. H. Sisson	. V. Kantelo . A. Sigg
EXPERIMENT	Measure fission yields for fission fission-spectrum neutrons.	of Am-241 induced by
METHOD	Measurements are being made by doin on the irradiated Am-241 samples an catcher-foil technique. Absolute y a comparison with U-235 fission yie tion of the mass-yield curve.	g chemical separations d by using the recoil ields will be based on lds and on a normaliza-

<u>COMPLETION DATE</u> The measurements are finished, and we plan to start writing a report for publication soon.

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LABORATORY	Lawrence Livermore Laboratory University of California P.O. Box 808	McClellan Central Laboratory 1155th Technical Operations Squadron
	Livermore, CA 94550, U.S.A.	McClellan AFB, CA 95652, U.S.A.
3. <u>NAMES</u>	D. R. Nethaway A. L. Prindle D. H. Sisson	M. V. Kantelo R. A. Sigg
FACILITY	Livermore ICT Facility (14-MeV neut	tron source)
EXPERIMENT	Measure fission yields for fission by 14.8-MeV neutrons.	of Am-241 induced

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METHOD Measurements were made by doing chemical separations on the irradiated Am-241 samples, and by using the recoil catcher-foil technique. The accuracy of the measurements is about ± 5%. Absolute yields are based on a normalization of the mass-yield curve.

COMPLETION DATE | The experiment is finished.

<u>PUBLICATION</u> A rough draft has been written which will be issued as a Lawrence Livermore Laboratory Report and then submitted to Phys. Rev. C.

U.S.A.

Laboratory and Address:		Oak Ridge National Laboratory P. O. Box X, Building 6010 Oak Ridge, Tennessee 37830, U.S.A.		
Names:		J. K. Dickens and J. W. McConnell		
Facilities:		Fast Rabbit Transport Station at the Oak Ridge Research Reactor (ORR)		
Experiment:		Absolute yields of forty-nine fission products, representing thirty-six mass chains created by thermal-neutron fission of ²³⁹ Pu have been determined.		
Method: A one microgram neutrons. Follo counting area. using a large-vo between 1550 and		a sample of ²³⁹ Pu was irradiated for 100 sec with thermal owing irradiation the sample was moved to a low-background Gross fission product gamma-ray spectra were obtained olume Ge(Li) detector. Counting intervals were initiated d 2.6 x 10 ⁶ sec following the end of the irradiation.		
Accuracy:		Absolute lo uncertainties range between 2.5 and 25%, made up of 2.0% uncertainty assigned to detector efficiency, 1.3% uncertainty in determining the number of fissions created in the sample, and un- certainties in peak extraction and in branching ratios and lifetimes given in the literature.		
Completion Date:		January 1979		
Discrepancies to Other Report Data:		Cumulative fission yields agree well with previous measurements and recommended evaluations except for mass chains between 101 and 107, for which the present results are larger than previously determined.		
Publications:		J. K. Dickens and J. W. McConnell, "Fission-product Yields for Thermal-neutron Fission of ²³⁹ Pu, (pre- print available May 1979).		

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U.S.A.

Laboratory and address:	Oak Ridge National Laboratory P. O. Box X, Building 6010 Oak Ridge, Tennessee 37830, U.S.A.		
Names:	J. K. Dickens		
Facilities:	Fast Rabbit Transport Station at Oak Ridge Research Reactor (ORR)		
Experiment:	Absolute yields of seventeen fission products, representing sixteen mass chains created by thermal-neutron fission of ²⁴¹ Pu have been determined.		

- Method: A one microgram sample of ²⁴¹Pu was irradiated for 50 sec with thermal neutrons. Following irradiation the sample was moved to a low-background counting area. Gross fission-product gamma-ray spectra were obtained using a large volume Ge(Li) detector. Six counting measurements were made between 17 and 210 hours after the irradiation.
- Accuracy: Between 4% and 15% (1 σ), made up of 2.5% uncertainty assigned to detector efficiency, 2.8% uncertainty in determining the number of fissions created in the sample, and the remainder due to uncertainties in peak extraction and in branching ratios and lifetimes given in the literature. Uncertainties assigned to nine of the measured yields are smaller than existing evaluated uncertainties for these yields.
- September 1978 Completion Date: Discrepancies to Other Report Data: Data agree with evaluation of Crouch (E. A. C. Crouch, Atomic Data and Nucl. Data Tables 19, 419 (1977)) for fifteen of the measured values. J. K. Dickens, "Fission Yields for Thermal-Neutron Fission of ²⁴¹Pu," Nucl. Sci. Eng. (in press). Publications:

- 49 -<u>U.S.A.</u>

Laboratory and address	Oak Ridge National Laboratory P. O. Box X, Building 6010 Oak Ridge, Tennessee 37830		
Names:	J. K. Dickens and R. W. Peelle		
Facilities:	Fast Rabbit Transport Station at Oak Ridge Research Reactor (ORR)		
Experiment:	Total Beta and Gamma Energy Release for Thermal- Neutron Fission of ²³⁵ U, ²³⁹ Pu, and ²⁴¹ Pu for Cooling Times of 2 to 14000 secs.		
Method: Microgram san with thermal gamma-ray ene tillation det time interval to differenti obtain total data have bee	es of 235 U, 239 Pu, and 241 Pu have been irradiated for short periods utrons, and returned pneumatically to a counting area. Beta- and cy spectra of moderate resolution have been obtained using scin- tors (NE110 for beta rays and NaI for gamma rays) for selected within the time range of interest. The spectra have been reduced production cross sections do/dE and have been integrated to tergy release rates for beta and gamma rays (separately). These summed to obtain the total energy release.		
Accuracy:	3% (10) for 235 U and 239 Pu, 4% for 239 Pu		
Completion Date:	May 1977 for ²³⁵ U, December 1977 for ²⁴¹ Pu, and August 1978 for ²⁴¹ Pu		
Discrepancies to Other Report Data:	Data are in reasonable agreement with other recent experiments and with results of summation calculations. However, present results are up to 15% smaller than recent data from Los Alamos; these discrepancies have not been resolved.		
Publications:	J. K. Dickens, J. F. Emery, T. A. Love, J. W. McConnell, K. J. Northcutt, R. W. Peelle, and H. Weaver, "Fission-Product Energy Release for Times Following Thermal-Neutron Fission of ²³⁵ U Between 2 and 14000 Seconds," ORNL/NUREG-14 (October 1977).		
	J. K. Dickens, J. F. Emery , T. A. Love, J. W. McConnell, K. J. Northcutt, R. W. Peelle, and H. Weaver, "Fission-Product Energy Release for Times Following Thermal-Neutron Fission of ²³⁹ Pu Between 2 and 14000 Seconds," ORNL/NUREG-34 (April 1978).		
	J. K. Dickens, T. A. Love, J. W. McConnell, J. F. Emery, K. J. Northcutt, R. W. Peelle, and H. Weaver, "Delayed Beta- and Gamma- Ray Production Due to Thermal-Neutron Fission of ²³⁵ U, Spectral Distributions for Times After Fission Between 2 and 14000 Sec: Tabular and Graphical Data," NUREG/CR-0162, ORNL/NUREG-39 (August 1978).		
	J. K. Dickens, J. F. Emery, T. A. Love, J. W. McConnell, K. J. Northcutt, R. W. Peelle, and H. Weaver, "Fission-Product Energy Release for Times Following Thermal-Neutron Fission of ²⁴¹ Pu Between 2 and 14000 Seconds," NUREG/CR-0171, ORNL/NUREG-47 (August 1978).		

	<u>Laboratory</u> : Oak Ridge National Laboratory, Bldg. 6010, P. O. Box X, Oak Ridge, TN 37830			
Names: R. L. Macklin				
<u>Facilities</u> : Oak Ridge Electron Linear Accelerator (ORELA) Flight Path 7				ar Accelerator (ORELA)
	Experiment:	xperiment: Fast Neutron (n,γ) Cross Sections E _n = 2.6 - ∿500 keV Target-isotopes: see table below		
	Method: Tota Nucl 1270	al Pi 1. In 0-12	rompt Photon Energy nstr. and Meth. <u>91</u> , 79 (1975).	Detectors, Neutron Time-of-Flight. 565-571 (1971), Phys. Rev. <u>C11</u> ,
	Accuracy: 2-	5% i:	n cross section, 20 .	2% resolution (FWHM)
	[Expected] co	omplo	etion date: See tab	le below
	<u>Publications</u>	:	μ	
	<u>Isotopes</u>		Completion Date	Publications
	86 _{Kr}		indefinite	Data taking planned for 1979
	86,87,88 _{Sr}		1979, plan to remeasure ⁸⁷ Sr w. D. Drake (LASL)	"Valence Neutron Capture in ⁸⁸ Sr", J. W. Boldeman, B. J. Allen, A. R. de L. Musgrove, Nucl. Phys. <u>A269</u> , 397 (1976)
l	89y		1976	J. Boldeman et al, Nucl. Sci. Engr. <u>64</u> , 744 (1977)
1	90,91,92,94Zr	r	1976	"The 292.4 eV Neutron Resonance Parameters of 91Zr", R. L. Macklin, et al, Nucl. Sci. Engr. <u>62</u> , 174 (1977) "keV Neutron Capture in Zirconium- 91", J. W. Boldeman et al, AAEC/E 367 "The Neutron Total and Capture Cross Sections of 92,94Zr", J. W. Boldeman et al, Nucl. Phys. <u>A263</u> , 389 (1976) "High Resolution Neutron Transmission and Capture for 917"
				et al, Austral. J. Phys. <u>30</u> , 391 (1977)
	93 _{Nb} ·		1976	R. L. Macklin, "Neutron Capture Cross Section of Niobium from 2.6 to 700 keV", Nucl. Sci. Engr. <u>59</u> , 12 (1976)

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Isotopes	Comple	tion Date	Publications
92,94,95,96,97,9	98,100 _{ΜΟ}	1976	"Neutron Resonances in 100Mo and Valence Neutron Capture", H. Weigmann et al., Phys. Rev. <u>C</u> (in press) "Resonance Parameters, Capture Gamma- Rays and Reaction Mechanisms in 98,100Mo + n", H. Weigmann et al., Proc. Spec. Mtg. on Structural Materials for Fast Reactors (Geel, Belgium Dec. 1977). "Average Neutron Resonance Parameters and Radiative Capture Cross Sections for the Isotopes of Molybdenum", A. R. de L. Musgrove et al., Nucl. Phys. <u>A270</u> , 108 (1976)
100,101,102,104 _R	u 1979		analysis proceeding
103 _{Rh}	1979		analysis proceeding
104,105,106,108,	110 _{Pd}	1979	"104,105,106,108,110Pd(n,γ) Cross Sections above 2.6 keV", R. L. Macklin et al., Nucl. Sci. Engr. (in press 1979)
106,108,110,111, 113,114,116 _{Cd}	112,	1978	"Neutron Capture Resonance Parameters and Cross Sections for the Even-A Isotopes of Cd", A. R. Musgrove et at., J. Phys. G, <u>4</u> , 771 (1978)
122,123,124,125, 126,128,130 _{Te}		1977	R. R. Winters (Denison University, ORNL Consultant) analyzing the data.
133 _{Cs}	indefi	nite	data taken
134,135,136,137,	138 _{Ba}	1976	"keV Neutron Resonance Capture in Barium-137", A. R. de L. Musgrove et al., Aust. J. Phys. <u>29</u> , 157 (1976) "keV Neutron Capture Cross Sections of 134Ba and 136Ba", Nucl. Phys. <u>A256</u> , 173 (1976). See below w. 140Ce
139 _{La}	1 9 78		"Resonant Neutron Capture in ¹³⁹ La", A. R. Musgrove et al., Austral. J. Phys. <u>31</u> , 47 (1978)

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	Isotopes	<u>Completion Date</u>	<u>Publications</u>
	140 _{Ce}	1978	"Resonance Neutron Capture in ¹³⁸ Ba and ¹⁴⁰ Ce and the Prompt Neutron Correction to Gamma-Ray Detectors", A. R. Musgrove et al., Austral. J. Phys. (in press) "Non-Statistical Neutron Capture in 140Ce", A. R. Musgrove et al., Austral. J. Phys.
	141pr	1977	Data taken, analysis at Lucas Heights
	142,143,144,145,	146,148 _{Nd}	"Non-Statistical Effects in the Radiative Capture Cross Sections of the Neodymium Isotopes", A. R. Musgrove et al., AAEC/E 401
	159 _{Tb}	1978	"Neutron Capture Cross Section of 159Tb From 2.6 to 700 keV", M. Mizumoto et al., Phys. Rev. <u>C17</u> , 522 (1978)
•	165 _{Ho}	1976	"The 165 Ho(n, γ) Standard Cross Sections from 3 to 450 keV", R. L. Macklin, Nucl. Sci. Engr. <u>59</u> , 231-236 (1976)
	169 _{Tm}	1979	Data taken, analysis by D. Drake, J. Malanify (LASL) and R. Macklin (ORNL), "Fast Neutron Capture Cross Sections of 169Tm, 191Ir, 193Ir and 175Lu for $3 \leq E_n \leq 2000$ keV", R. L. Macklin et al., Los Alamos Report LA-7479-MS (1978)

For most of these measurements, see also: Proceedings of an International Conference on Neutron Physics and Nuclear Data for Reactors and Other Applied Purposes, Harwell, 1978, page 449.

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CONTRIBUTION TO "PROGRESS IN FISSION PRODUCT NUCLEAR DATA"

<u>U. S. A.</u>

Laboratory and Address:

Pacific Northwest Laboratory P. O. Box 999 Richland, WA 99352 USA

Names: P. L. Reeder and R. A. Warner

Facilities: SOLAR - Spectrometer for On-Line Analysis of Radionuclides. This is an on-line mass spectrometer which incorporates a 235U target in a surface ionization source located in the thermal column of a 1 MW TRIGA reactor at Washington State University, Pullman, WA.

Experiment: Isomer yield ratios for $^{235}U + n_{+h}$.

- Method: Ratios of independent yields of fission product isomers are being measured for thermal neutron fission of 235U by use of an on-line mass spectrometric technique. A short burst of neutrons from the TRIGA reactor is used to produce various isomers of Br, Rb, In, I and Cs fission products within the surface ionization source. Selective ionization performs the rapid chemical separations and magnetic analysis performs the mass separation to give the desired nuclides as a beam of ions. Ions are collected on a moving tape collector system for a short time interval during and after the neutron pulse. The radioactive decay of the two isomers is followed by beta and gamma counting to determine the relative yield of each isomer.
- Accuracy: The final accuracy will probably depend more on how well the decay schemes are known for particular cases than on statistical uncertainties.

Completion Date: It is hoped to complete this work by spring of 1980.

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Laboratory and address: University of Illinois Nuclear Radiation Laboratory Nuclear Engineering Program Urbana, Illinois 61801 U.S.A.

Names: Bernard W. Wehring

- Facilities: Illinois Advanced TRIGA 1.5-MW Nuclear Reactor, HIAWATHA Fission-Fragment Mass Spectrometer.
- Experiment: Direct Physical Measurement of the Primary Postneutron-Emission Nuclide Yields in Thermal-Neutron Fission of U-235, Pu-239, and U-233
- Method: The fission-fragment recoil mass spectrometer HIAWATHA, consisting of a cylindrical focusing electrostatic analyzer and time-offlight system, is used to determine fragment masses while fragment energy loss is used to identify fragment atomic numbers in a multiparameter experiment. All fragment velocities and charge states are measured.
- Accuracy: 0.5-amu mass resolution, achieved, about 1-Z atomic-number resolution, achieved, 1% standard error (relative error) in largest mass yield, achieved, 0.02-0.1% standard error (absolute error) in nuclide yields, achieved.

Completion date:

Publications:

- Gino Dilorio, "Direct Physical Measurement of Mass Yields in Thermal Fission of Uranium 235," Ph.D. Thesis, University of Illinois at Urbana-Champaign, 1976.
- Gino DiIorio and B. W. Wehring, "Direct Physical Measurement of Mass Yields for 235 U(n_{th},f)," Trans. Am. Nucl. Soc. <u>24</u>, 459 (1976).
- Gino Dilorio and B. W. Wehring, "HIAWATHA, A Fission-Fragment Recoil Mass Spectrometer," Nucl. Instr. Methods <u>147</u>, 487 (1977).
- R. B. Strittmatter and B. W. Wehring, "Direct Physical Measurement of Nuclide Yields for ²³⁵U(n_{th},f)," Trans. Am. Nucl. Soc. <u>27</u>, 862 (1977).
- R. B. Strittmatter, "Nuclide Yields for Thermal Fission of Uranium 235," Ph.D. Thesis, University of Illinois at Urbana-Champaign, 1978.
- R. B. Strittmatter and B. W. Wehring, "Direct Measurement of Nuclide Yields in Thermal-Neutron Fission Using HIAWATHA," Proceedings of the International Conference on Neutron Physics and Nuclear Data for Reactor and other Applied Purposes, Harwell, September 25-29, 1978.

II. COMPILATIONS AND EVALUATIONS

(revisions with respect to the last issue are marked by a vertical bar in the margin)

FFANCE

- LABORATORY AND ADDRESS : LABORATOIRE DE METROLOGIE DES RAYONNEMENTS IONISANTS CEN SACLAY BP 2 91190 GIF SUR YVETTE
- NAMES : J. LEGRAND, F. LAGOUTINE

EVALUATION : Radionuclide decay data

- PURPOSE : To provide recommended values and uncertainties for the nuclear data, including half-lives, Q-values, branching fractions for the various decay modes, energies and intensities of all emitted radiations (e.g. β , C.E, γ , X-ray) ; K and total ICC.
- METHOD : Selection of sources materials. Data evalued on the basis of the rules established by the European group.
- STATUS : Evaluation of the following F.P. ${}^{86}_{\text{Rb}}$, ${}^{89}_{\text{Sr}}$, ${}^{90}_{\text{Sr}}$, ${}^{91}_{\text{Y}}$, ${}^{95}_{\text{Zr}}$, ${}^{106}_{\text{Ru}}$, ${}^{106}_{\text{Rh}}$, ${}^{127m}_{\text{Te}}$, ${}^{131}_{\text{I}}$, ${}^{131m}_{\text{Xe}}$, ${}^{131m}_{\text{Te}}$, ${}^{137}_{\text{Cs}}$, ${}^{140}_{\text{Ba}}$, ${}^{141}_{\text{Ce}}$.

COMPLETION DATE : 1977 - 1979

PUBLICATION : Table de radionucléides vol. 1 - 2 - edition CEA-LMRI ISBN-2-7272-0004-8. Some 80 nuclides are available, including the F.P.

PHONED ACTIVITIES : continuous evaluation

INDIA

Laboratory	and	address:	Health Physics Division
			Bhabha Atomic Research Centre
			Bombay 400 085, India.
Names:			P.P.Chakraborty, D.N.Sharma, M.R.Iyer
			and A.K.Ganguly.

THEORETICAL COMPILATION

- (1) <u>Type of data</u>: Mass yield and Elemental yield prediction of fragments for spontaneous fission of various fissioning nuclides.
- (2) <u>Purpose</u>: To predict mass and elemental yield of fragments from various fissioning nuclei using a model which is independent of any input fission data.
- (3) <u>Major source of information</u>: The stable neutron numbers as a function of Z of nuclei obtained from the Nuclear Mass Tables is the only input required for the calculations. Using this input data the Order Disorder Model is applied to the fissioning nucleus to predict yield values.
- (4) <u>Results and discrepancies</u>: The various characteritics of the mass yield distribution, viz., peak-to-valley ratios, bunching of higher mass peaks etc. agree well with those of experimental data. The absolute values have some discrepancies with experimental values, though an improvement in the right direction has been noticed by using the later data on nuclear stability. This leads to the conclusion that the accuracy of the predictions depend on the accuracy of the only input data used in the calculations viz., the stable

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<u>INDIA</u> (cont'd)

neutron number values. The yield distribution predicted for trans-californium nuclei show the onset of symmetric division for fission of higher mass number nuclei. (5) <u>Completion date</u>: The first stage of the computation is completed by February 1977 and further improvement of the predictions is under progress using the improved stability data of nuclei.

(6) Publications: (i) Chakraborty P.P.

Asymmetry of Mass and Charge Distribution in Low Energy Fission and Fissile Material Identification Techniques

Ph D Thesis, Gujarat University, India 1977

(ii) Chakraborty P.P., Sharma D.N., Iyer M.R. and Ganguly A.K. Asymmetry of Mass and Charge Division

To be published.

ITALY

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Laboratory and address:	CNEN, Centro di Calcolo Via Mazzini, 2 - 40138 Bologna, Italy				
Names:	F. Fabbri, T. Martinelli, E. Menapace, M. Motta G.C. Panini, G. Reffo, M. Vaccari, A. Ventura				
Evaluation:	The revision of the previous evaluations concerning Pd-105, Nd-143, Sm-149, Sm-151 on the basis of recent experimental microscopic and integral data				
Purpose:	Estimate of long term reactivity changes and FP. accumulation in fast reactor				
Method:	Calculations by BW-single and -multilevel for- malism (resonance region) and by statistical and optical models				
Major sources of information:	NEUDADA, CINDA, Nuclear Data Sheets				
Deadline of literature coverage: December 1978					
Status:	The full evaluation and file compilation in the range 0-15 MeV of the before mentioned isotopes is in progress and the results are under discussion with the cooperating parts (see below)				
Cooperation:	CEA - Cadarache and Saclay and ECN Petten				
Other relevant details:	25 group cross sections at infinite dilution and 0°K temperature have to be generated for each evaluated isotope				
Computer file of evaluated dat	Computer file of evaluated data: ENDF/B format.				
Publications:	Tables of Neutron Resonance Data and Parameters for 63 Fission Products Evaluated and Compiled at the Bologna Centre (1977). A. Montaguti, M. Vaccari, CNEN-RT/FI(78)23.				
	Multigroup Cross-Sections of 63 Fission Product Nuclei from Different Nuclear Data Files.				

A. Montaguti, G.C Panini, M. Vaccari, CNEN-RT/FI(78)16.

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JAPAN
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(same as in INDC(NDS)-95)

Japanese Nuclear Data Committee / Decay Heat Nuclear Data Working Group R. Nakasima (Hosei University) group leader M. Yamada (Waseda University), T. Tamai (Kyoto University) 1. Otake and A. Zukeran (Power Reactor and Nuclear Fuel Development Corp.) S. Iijima, T. Murata, T. Yoshida and M. Iida (Nippon Atomic Industry Group Co.) T. Hojuyama (Mitsubishi Atomic Power Industry) K. Umezawa, T. Tasaka, Z. Matumoto and T. Tamura (Japan Atomic Energy Research Institute) M. Akiyama (University of Tokyo) T. Yamamoto (Tohoku University) 1. Compilation: Decay data and delayed neutron data Purpose: For summation calculation of decay heat Major Sources of Information: Journals, Nuclear Data Sheets and ENSDF Deadline of Literature Coverage: None Cooperation: None Computer File: Nuclear structure data file NDFILE and retrieval program ABEG. ORNL Evaluated Nuclear Structure Data File, ENSDF. Expected Completion Date: Continuous compilation Publication: in plan 2. Evaluation: Estimation of decay and delayed neutron data Sensitivity study for decay heat Purpose: Making more reliable estimation of released beta and gamma energies for short-lived fission products Estimation of delayed neutron emission probability Estimation of the effects of experimental errors in decay data Method: Application of gross theory of beta decay Search and use of systematics Major Sources of Information: Own compiled data Deadline of Literature Coverage: None Status: Estimation of released beta and gamma energies completed, but minor correction required Estimation of delayed neutron emission probability almost completed. but quite preliminary Computer program for sensitivity studies in preparation Cooperation: None Computer File of Evaluated Data: in plan Discrepancy Encountered: Gamma ray assignments in the case of beta decays of both the ground and isomeric states, for example Rb-90 Intensity of the ground state beta Publication: T. Yoshida, Nucl. Sci. Eng. 63 (1977) 376

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JAPAN

Laboratory and address : Japanese Nuclear Data Committee/FPND W.G., Japan Atomic Energy Research Institute, Tokai-mura, Naka-gun, Ibaraki, Japan.				
Names : S. Iijima, M. Kawai, T. Murata, T. Yoshida (Nippon Atomic Industry Group Co., Ltd.) S. Igarasi, T. Nakagawa, Y. Kikuchi, Z. Matumoto, H. Nishimura(JAERI) H. Matsunobu (Sumitomo Atomic Energy Industries, Ltd.) H. Sasaki (Mitsubishi Atomic Industries, Inc., now at PNC) T. Aoki (Fuji Electric Co.) K. Maki, A. Zukeran (Hitachi Ltd.) T. Watanabe (Kawasaki Heavy Industries) I. Otake (PNC) R. Nakasima (Hosei Univ.)				
Evaluation : Neutron cross sections of Nd isotopes.				
Method : Calculation with spherical optical model and statistical theory, Single and multi-level BW formula in thermal and resonance regions. Optical model parameters are determined by SPRT method. Level density parameters are re-evaluated. Source : NEUDADA, CINDA, and the recent capture data for Nd isotopes at JAERI and Lebedev institute. Integral data from STEK and CFRMF.				
Deadline of literatur coverage : Spring, 1979				
Status : Progress is very slow. Difficulties are encountered in the consistent determination of level density parameters <u>a</u> , T, etc.				
Other relevant details :				
The evaluation of 68 nuclides was completed in Aug., 1977, and the file is available from NEA Data Bank. Integral test calculation using STEK reactivity data and CFRMF acti- vation data was completed recently. Results are being examined.				

Computer file of evaluated data : JENDL (ENDF/B-4 format)

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JAPAN

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Expected completion date : The work is largely behind schedule.

Discrepancies encountered :

Summarized in No. 4 of publication list.

Recent publications :

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- H. Matsunobu and T. Watanabe, compilation of measured capture cross sections for JENDL FPND file, JAERI-M 7568 (1978).
- 2. Z. Matsumoto, T. Murata and R. Nakasima, Level scheme for some fission product nuclides. Comparison of level scheme used by JAERI and Petten, JAERI-M 7734 (1978).
- 3. S. Iijima et al., J. Nucl. Sci. Technol. 14 161 (1977).
- 4. S. Iijima, IAEA-213, Petten 1977, Review Paper No. 9.
- 5. H. Nishimura et al., Integral test of JENDL-FP data file, JAERI-M report (to be published shortly)

NETHERLANDS

	Laboratory and address	Netherlands Energy Research Foundation (ECN) Postbus 1, 1755 ZG Petten, The Netherlands. Telephone: (02246) - 6262, telex: 57211 reacp nl
	Names	J.W.M. Dekker, H. Gruppelaar, R.J. Heijboer and A.J. Jansser
1	<u>Evaluation</u>	 RCN-2 evaluation of neutron cross sections (σ_t, σ_e, σ_{nγ}, σ_{nn}, -matrix, σ_{n2n}) for about 60 fission products in the energy range of 10⁻³ eV to 15 MeV, in KEDAK type format, for the following elements, Zr, Nb, Mo, Tc, Ru, Rh, Pd, Ag, Cd, Te, I, Xe, Cs, La, Ce, Pr, Nd, Pm, Sm, Eu, Gd, Tb. Generation of group cross sections for fast reactor calculations based on RCN-2 evaluation in the 26-group ABBN scheme with a fast reactor flux weighting spectrum and error files for capture group constants, including 26×26 covariance matrices. Adjustment of capture group constants based on (2) and integral STEK+CFRMF measurements. Generation of an adjusted point cross sections library based on STEK+CFRMF integral data. Calculation of pseudo-fission-product cross sections.
	Purpose	Fast breeder power-reactor data needs.
	Method	Calculation with multilevel Breit-Wigner formula, optical model and revised statistical model, taking into account all available experimental information.
	Major sources of information	BNL-325, NEUDADA, CINDA, Nuclear Data Sheets, recent liter- ature, integral data mainly from STEK and CFRMF.
	Status	 (1-3) Completed and reported for 40 isotopes; planned for 1979: new evaluations for Eu-isotopes, revisions for important fission products; planned for 1980: 25 other isotopes. (4) In progress, completed 1979-1980. (5) Updating in 1980.
	Computer file	 KEDECN, KEDAK type format (co-operation with Dr. B. Goel, KfK, Karlsruhe), has been sent to NEA Data Bank at Saclay.
	Completion date	1979/1980.
1	Recent publi- cations	 H. Gruppelaar, Tables of RCN-2 fission-product cross section evaluation, part <u>1</u>, ECN-13 (1976), part <u>2</u>, ECN-33 (1977), and part <u>3</u> (in press). J.W.M. Dekker, Tables and figures of adjusted and un- adjusted capture group cross sections based on the RCN-2 evaluation and integral measurements in STEK, part <u>1</u>, ECN-14 (1977), part <u>2</u>, ECN-30 (1977), and part <u>3</u>, ECN-54 (1979).
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NETHERLANDS

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- (2,3) J.W.M. Dekker and H.Ch. Rieffe, Adjusted capture cross sections of fission-product nuclides from STEK reactivity worths and CFRMF activation data, ECN-28 (1977) and part 2, ECN-55 (1979).
 (5) R.J. Heijboer, Pseudo fission-product cross sections for a 1200 NUC fission and product cross sections
- (5) R.J. Heijboer, Pseudo fission-product cross sections for a 1300 MWe fast breeder reactor (status July 1978), ECN-52 (1978).

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UNITED KINGDOM

(same as INDC(NDS)-95)

Laboratory and Address:	AERE Harwell	UKAEA AERE, Harwell, Oxfordshire, OX11 ORA
Name:	E.A.C. Crouch	
<u>Compilation</u> :	Chain, Cumulative and Ind yields for all neutron in with neutrons of energy up spontaneous fission. On	ependent fission product duced fission reactions p to 14 MeV, including going compilation.
Purpose:	Basic data for fission yi	eld evaluation.
Sources:	Journals, Proceedings of Sother open literature, Pr work is complete but unli	Learned Societies, or oject reports if the kely to be published.
Deadline:	No results prior to 1950	are collected.
Cooperation:	We are prepared to exchan	ge files with other groups.
Computer File:	Information held in stand	ard forms on Computer Files.
Completion Date:	Continuous compilation.	
Publications:	AERE R6642 'A library of product yields maintained computer methods'. 'Part I: The establishme E.A.C. Crouch, December 1	neutron induced fission and interrogated by nt of the library'. 970.
	AERE R7207 'A library of product yields maintained computer methods'. 'Part II: The interrogat E.A.C. Crouch, August 197	neutron induced fission and interrogated by ion of the library'. 2.
·	Fission Product Yields fr E.A.C. Crouch. Atomic Data and Nuclear D May 1977. (Contains experimental va after fitting to conserve	om Neutron-Induced Fission - ata Tables, Vol. 19, 5, lues and adjusted values
	Laboratory and Address: Name: Compilation: Purpose: Sources: Deadline: Cooperation: Computer File: Completion Date: Publications:	Laboratory and Address: AERE Harwell Name: E.A.C. Crouch <u>Compilation</u> : Chain, Cumulative and Ind yields for all neutron in with neutrons of energy u spontaneous fission. On Purpose: Basic data for fission yi Sowrces: Journals, Proceedings of other open literature, Pr work is complete but unli Deadline: No results prior to 1950 Cooperation: We are prepared to exchan Computer File: Information held in stand Completion Date: Continuous compilation. Fublications: AERE R6642 'A library of product yields maintained computer methods'. 'Part I: The establishme E.A.C. Crouch, December 1 AERE R7207 'A library of product yields maintained computer methods'. 'Part II: The interrogat E.A.C. Crouch, August 197 Fission Product Yields fr E.A.C. Crouch. Atomic Data and Nuclear D May 1977. (Contains experimental va after fitting to conserve

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UNITED KINGDOM

Laboratory and Address:	AERE	Harwell	UKAEA AERE Harwell Oxfordabiro OX11 ORA
Name:	E.A.	C. Crouch	OXIOIUSIIITE OXII OKA
Evaluation:	(1)	Neutron induced fission fissile nuclides at ne MeV; chain yields and	n product yields for all utron energies up to 15 independent yields.
	(2)	Adjustments of the cha lated independent yield the conservation laws set'.	in yields and the calcu- ds to force agreement with i.e. to form a 'consistent
Purpose:	UKND	File to be used in Read	ctor design and operation.
Method:	(1)	The individual yields : chain and independent) and the means calculate	for a given reaction (both , are examined, weighted ed together with the errors.
	(2)	The evaluated yields at ation to fill missing independent yields by meters estimated from 1 are fitted by least squ conditions to give adju and independent yields	re augmented by interpol- values or in the case of calculation based on para- known values. The results uares to the conservation ustments for chain yields
		Almost complete - the s and the equality of yie elements. The set will to produce an estimate Fission nearer to expe vious sets.	fitting of conservation laws elds of complementary l be tested for its ability of after heat from ²³⁹ Pu erimental values than pre-
Sources:	Сотр	ilation mentioned above	
Deadline:	No r beli resu	esults prior to 1950 are eved to be complete up a lts included.	e collected. Compilations to end 1975, some 1976
Status:	Eval 1977	uation and Consistent so . Further development o	et complete at January continuing.
Co-operation:	We a	re prepared to exchange	files with other groups.
Computer Files of Compiled Data:	Comp	ilation as above.	· · ·
Computer File of Evaluated data:	Magn set :	etic tape or punched can in ENDF/BIV format.	rds of the consistent
Discrepancies found:	File disc:	s are compared with thos repancies found are reso	se of B.F. Rider and olved.
Publication:	Fiss E. A Atom: May	ion Product Yields from . C. Crouch. ic Data and Nuclear Data 1977.	Neutron-Induced Fission.

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UNITED KINGDOM

Laboratory and Address	AERE Harwell	UKAEA, AERE, Harwell, Oxfordshire, OX11 ORA U.K.
Names:	E. A. C. Crouch	
<u>Evaluation</u> :	Compilation and evaluation of delayed neutron emitter pemission probabilities of the emitters. Hence, using the consistent sets, calculation neutron yields.	of the half lives precursors and he delayed neutron fission product n of the delayed
Purpose:	UK Nuclear Data File for use design and operation calcula	e in Reactor ations.
Sources:	The open literature.	
Deadline:	Continuous compilation.	
Status:	Compilation of delayed neut	ron data proceeding.
Co-operation:	We are prepared to exchange other groups.	information with
Computer files:	Not yet implemented.	

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	Laboratory and Address		CEGB Berkeley	Berkeley Nuclear Laboratories Berkeley Gloucestershire GL13 9PB U.K.
	Working Group:	Mrs.	B.S.J. Davies A. Tobias J.R. Parkinson M.F. James A.L. Nichols D.G. Vallis K.M. Glover G. Evangelides	CEGB, BNL. CEGB, BNL. BNFL, Windscale. AEE, Winfrith. AEE, Winfrith. AWRE, Aldermaston. AERE, Harwell. Imperial College, London.
	1. <u>Compilation and</u> <u>Evaluation</u> :			
	Radionuclide	decay	data	
	purpose:		to provide a compre radioactive decay d branching ratios, me with associated unc	mensive, updated data file of ata, including half lives, Q-values an α , β and λ energies and intensities ertainties.
	progress:		the FPND set which A. Tobias with the become the recommen- in the U.K. It is	was formed by merging the data of JS ENDF/B IV decay data file has ded set for decay-heat calculations now known as UKFPDD-1.
			Data for about 80 m has been provisiona new set which will	uclides has been re-evaluated and lly merged with UKFPDD-1 forming a be tested in decay heat calculations.
2. Decay scheme calculations - the CASCADE Programme				
	purpose:		to compare experiment from a more basic date produce mutually con- for different radia	ntal data with decay data calculated ata set (e.g. US ENSDF file) and to nsistent catalogues of emission data tion types.
	progress:		the CASCADE code de CASCADE and its sui in all of the possi	velopment is nearing completion. Te of programmes are being tested ole options to eliminate any bugs.
			The major $\lambda - \lambda$ and can now be evaluated	λ - x and other coincidence data sets data automatically.
			The results can be a or V format. Decay of included in ENDF/B is be included in an El inclusion of $\lambda - \lambda$ of now being tested.	reproduced in standard ENDF/B IV and/ lata information not previously IV or V but evaluated by CASCADE can IDF/B IV or V <u>type</u> format (e.g. coincidence data tables). This is

completion date: expected complete by end of 1979.

Publications: FISP-5 An extended and improved version of the fission product code FISP.RD/B/N4303. TOBIAS, A.

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(cont'd)

TOBIAS, A.	A brief description of the ENDF/B-V format adopted for use in the U.K. Decay and Fission Product Data files. RD/B/N4423.
TOBIAS, A.	Extensions to COGEND for ENDF/BV output of spontaneous fission decay data. RD/B/N4309.
DAVIES, B.S.J. et al	The U.K. Chemical Nuclear Data Files: An evaluated set of radioactive decay data for reactor calculations. Presented at the International Conference on Neutron Physics and Nuclear Data for Reactors and other applied purposes, Harwell 25-28 Sept. 1978.
WOOLLAM, P.B.	An assessment of the data for decommissioning calculations on Ag-108 metastable RD/B/N4373

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UNITED KINGDOM

Laboratory and address:	Birmingham Radiation Centre	University of Birmingham P.O. Box 363 Birmingham B15 2TT United Kingdom	
Name:	D.R. Weaver		
Evaluation:	Equilibrium and near-equilibrium delayed neutron spectra		
Purpose:	For reactor physics calculations and analysis of delayed neutron yield measurements. The evaluation was recommended by the March 1979 Vienna Consultants' Meeting on Delayed Neutron Properties		
Method:	To be decided		
Deadline of literature coverage:	None		
Status:	April 1979: Collection of numerical data and uncertainty information begun		

U.S.A.

Laboratory and address:		National Nuclear Data Center Building 197D Brookhaven National Laboratory Upton, New York 11973		
Names	:	S. Mughabghab N. Holden		
Evaluation	:	Revised edition of BNL-325 Vol. 1		
Information Source	:	All experimental measurements of thermal cross sections, resonance parameters, and average resonance parameters.		
<u>Details</u>	:	BNL-325, Vol. 1, Resonance Parameters will appear in two parts: the first part covering elements and isotopes for Z=1-60; the second part, 61-99. The format will be the same as that of the third edition. Additional quantities such as incoherent scattering cross sections, 30 keV Maxwellian average capture cross sections, γ -ray strength functions will be included. The introduction will be expanded to incorporate among other things a table of resonance energies used as standards, calcu- lations of thermal cross sections according to the Lane-Lynn theory.		
Computer File	:	The resonance parameter data will be on a computer file and can be sent to users on request.		
Completion Date	:	It is hoped that Part 1 will be completed sometime during the summer of 1979; Part II by the year end.		
References:	:	S. F. Mughabghab and D. I. Garber BNL-325, Neutron Cross Sections, Vol. 1, Resonance Parameters (third edition) 1973.		

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Laboratory and address:	Idaho National Engineering Laboratory EG&G Idaho, Inc. P. O. Box 1625 Idaho Falls, Idaho 83401 USA
Names:	C. W. Reich, R. L. Bunting, R. G. Helmer, M. A. Lee
<u>Compilation</u> ;	Decay data for fission products. Quantities treated include: T_2 ; QB; branching fractions for the various decay modes; energies and intensities of all emitted radiations (e.g., B, γ , C.e., x-ray); K-, L- and total ICC; delayed-neutron energy spectra for individual precursors; uncertainties in all measured values.
Purpose:	Decay data file for ENDF/B.
Major sources of information:	Nuclear Data Sheets, journals and preprints of recent work.
Deadline of literature coverage:	Ongoing. For Version V of ENDF/B, cut-off date is approximately September, 1978
<u>Computer File</u> :	Decay data are included in ENDF/B Fission Product File. Tapes available through normal ENDF/B procedures. Evaluated decay data sets for 317 fission- product nuclides (and isomeric states) have been prepared for inclusion in the ENDF/B-V Fission-Product File.
Publications:	C. W. Reich, "Applications of Fission- Product Decay Data", in Proceedings of the Isotope Separator On-Line Workshop, U. S. DOE Report BNL 50847 (July, 1978) pp. 109-148.

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<u>U.S.A</u>.

Address:	General Electric Company Vallecitos Nuclear Center P. O. Box 460 Pleasanton, California 94566, USA				
Name:	Dr. Benjamin F. Rider				
Compilation:	Fission Product Yields for 30 Fissioning Systems U235T Pu239T U233F Pu241F U234F Pu238F				
	U235F Pu239F U233HE Pu242F U237F Am241F U235HE Pu241T U236F Th232H Pu240H Am243F U238F U233T Pu239H Np237F U234H Np238F U238HE Th232F Pu240F Ef252s U236H Cm242F				
Purpose:	For Burnup, fission rate, shielding, fission product source terms, inventory, and decay heat calculations. Basis for ENDF/B-V yields of USA National Nuclear Data Center.				
Sources:	CINDA, INIS Atomindex, Journals, Correspondence, Reports				
Deadline:	Ongoing				
Cooperation:	Brookhaven National Laboratory, Cross Section Working Evaluation Group (CSEWG) Evaluated Nuclear Data File (ENDF/B-V)				
Details:	Approximately 18,000 entries from 1200 references, 100% proof read against original publications in 1979 for reliability; intercompared in cooperation with E.A.C. Crouch compilation at Harwell for completeness and accuracy. Independent and cumulative yields from weighted averages of measured yields, unmeasured yields computed from best available models. Isomer ratios of independent yields to metastable and ground states from measurement and calculated from nuclear spins for unmeasured <i>isomers</i> of known spin states. Yields for and cumulative yields are before delayed neutron emission and cumulative yields are after delayed neutron emission. Charge balance adjusted to conserve protons, including ternary fission. Resulting yields give fission product decay heat power for U235T in good agreement with experiment.				

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Completion Date: March 17, 1979, for current version (revised annually).

Publications: "Compilation of Fission Product Yields", NEDO-12154-3A (1979), available on microfiche only from General Electric Co., Vallecitos Nuclear Center, P. O. Box 460, Pleasanton, California 94566, USA - ATTN: Dr. B. F. Rider. U.S.A.

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Laboratory and address:

Hanford Engineering Development Laboratory P.O. Box 1970 Richland, WA 99352

Names:

RE Schenter, FM Mann, DL Johnson, and F Schmittroth

Evaluation:

ENDF/B-V Fission Product Data File and Fission Yield Files

- A. Coordinate generation and testing of complete ENDF/B-V file which will contain cross sections, decay data and fission yields for 877 fission product nuclei and 11 fissionable nuclei. Coordination is part of the responsibility as Chairman of CSEWG (Cross Section Evaluation Working Group) Fission Product and Actinide Data Subcommittee. Two fission product working groups to this subcommittee are chaired by TR England (LASL) and CW Reich (INEL) and cover the areas of fission yields and experimental decay data, respectively. Evaluations to these files will be contributed by essentially all CSEWG member laboratories.
- B. Evaluate important FP cross sections for fast and thermal reactor application. These will mainly involve updating about 180 cross section evaluations from ENDF/B-IV with emphasis on capture. Use will be made of combining recent integral and differential data results from CFRMF, STEK, RPI and ORNL.
- C. Evaluate decay data parameters \bar{E}_{β} , \bar{E}_{γ} for "Theoretical" ("no line data") FP nuclides using extrapolated "fits" to known data and integral testing of recent decay heat measurements.

purpose:

Update ENDF/B Fission Product Data Files

completion dates:

ENDF/B-V FP file will be issued July 1979. ENDF/B-V fission yield files issued April - May 1979.

references:

R. E. Schenter, "Fission Product and Actinide Data Evaluations for ENDF/B-V," <u>Trans. Am. Nuc. Soc. 28</u> (1978) 738.

Additional references related to this work may be obtained from RE Schenter.

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Laboratory and Address:

University of California Los Alamos Scientific Laboratory P. O. Box 1663 Los Alamos, New Mexico 87545 (U.S.A.)

Names:

T. R. England (LASL)
R. E. Schenter (HEDL)
B. F. Rider (G.E.)
J. Liaw (U. of Oklahoma)

Compilation:

Library of evaluated fission product yields for Version V of the Evaluated Nuclear Data Files (ENDF/B-V).

Deadline of Literature Coverage:

Mid-1978, including recent unpublished data.

Cooperation:

Subcommittee consisting of members from major U.S.A. commercial and government laboratories.

Other Relevant Details:

Twenty yield sets for 11 fissionable nuclides $(^{233}, 235, 236, 238_{U}, 239, 240, 241, 242_{Pu}, 237_{Np}, 232_{Th}, and 252_{Cf})$. Each set contains ~ 1100 yields and uncertainties; independent yields before delayed neutron emission and cumulative yields (by A and Z) after delayed neutron emission are given. Yield distributions account for isobaric states, Z and N pairing effects, ternary fission and delayed neutron branching.

Completion Date:

August 1978 for compilation. Phase I testing completed. Phase II testing in progress.

Publications:

Report on Phase I testing, and other relevant details in progress.

Computer File:

Distributed by the National Nuclear Data Center at the Brookhaven National Laboratory.

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LABORATORY AND ADDRESS:

University of California Los Alamos Scientific Laboratory P O Box 1663 Los Alamos, New Mexico 87545 (USA)

NAMES :

T. R. England R. J. LaBauve W. B. Wilson

COMPILATION:

Library of processed 154-group ENDF/B-IV fission_product reaction cross sections.

PURPOSE:

Data file of multigroup values (10^{-5} eV-20 MeV) for use in collapsing to few-group values.

MAIN SOURCE OF INFORMATION:

ENDF/B-IV Fission-Product Data File

OTHER RELEVANT DETAILS:

Cross sections were processed into the Power Reactor Studies (PRS) 154-group structure described in Ref. 1 and 2, using the PRS Neutron Flux Weighting Function described in Ref. 1. Cross sections were processed at 900 or 1000 K at infinite dilution. A total of 181 nuclides are described with total, elastic, total inelastic, and radiative capture multigroup values. Additional neutron absorption reaction cross-section tabulations are given for 36 of the nuclides. A total of 960 multigroup cross-section tabulations are included in the data file, which is issued with a companion collapsing code TOAFEW.

COMPUTER FILE:

The data file and collapsing code are available from the Radiation Shielding Information Center, Oak Ridge National LAboratory, P O Box X, Oak Ridge, Tennessee 37830 (USA).

REFERENCES:

- W. B. Wilson, T. R. England, and R. J. Labauve, "Multigroup and Few-Group Cross Sections for ENDF/B-IV Fission Products; the TOAFEW Collapsing Code and Data File of 154-Group Fission-Product Cross Sections," Los Alamos Scientific LAboratory report LA-7174-MS (March 1978).
- 2. R. J. LaBauve and W. B. Wilson, "Proposal to Extend CSEWG Neutron and Photon Multigroup Structures for Wider Applications," Los Alamos Scientific Laboratory report LA-6240-P (February 1976).

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Laboratory and Address:

University of California Los Alamos Scientific Laboratory P. O. Box 1663 Los Alamos, New Mexico 87545

Names:

T. R. England R. J. LaBauve D. G. Madland W. B. Wilson

Cooperation:

R. E. Schenter, chairman of the ENDF/B actinide and fission product subcommittee, and F. Schmittroth of the Hanford Engineering Development Laboratory, P. O. Box 1970, Richland, Washington 93352.

Compilations:

A) Nuclide Parameter Evaluated Compilations

1) β and γ decay energies, branching fractions [decay and (n,γ)], half-lives, Q-values and cross sections for 824 fission products are tabulated in Ref. 1. This is a basic data set that includes the major types of parameters, with corrections, from ENDF/B-IV, except for yields and the energy dependence of cross sections.

2) Multigroup β and γ spectra are tabulated in Ref. 2 for 180 fission product nuclides. The β spectra are given in 75 groups and the γ spectra in 150 groups. These data are based on ENDF/B-IV.

3) Multigroup cross sections are compiled in Ref. 11, processed from ENDF/B-IV.

4) Few group fitted spectral functions available in Ref. 9.

5) Comparisons with experiment and a new decay heat standard are presented in Ref. 14. Reference 15 is a code incorporating the pulse function data of the new ANS Decay Heat Standard.

B) Evaluations

1) Yield distribution (pairing effects) and branching to isomeric states are evaluated and modeled in Refs. 3 and 4. Estimated values from the modeling are also included in these references.

2) Ternary fission is evaluated and compiled in Ref. 5.

3) β and γ spectra, decay heating and absorption buildup are evaluated by comparison with experiment in Refs. 6-13.

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Purpose:

Research by the Los Alamos nuclear data group (T-2) is directed at improvement in the national data file ENDF/B and at the use of these data in, e.g., determining a new decay heat standard, absorption buildup, etc.

References

- T. R. England and R. E. Schenter, "ENDF/B-IV Fission Product Files: Summary of Major Nuclide Parameters," Los Alamos Scientific Laboratory report LA-6116-MS [ENDF-223] (October 1975).
- T. R. England and M. G. Stamatelatos, "Multigroup Beta and Gamma Spectra of Individual ENDF/B-IV Fission-Product Nuclides," Los Alamos Scientific Laboratory report LA-NUREG-6622-MS (Decamber 1976).
- 3. D. G. Madland and T. R. England, "The Influence of Pairing on the Distribution of Independent Yield Strengths in Neutron-Induced Fission," Los Alamos Scientific Laboratory report LA-6430-MS [ENDF-240] (July 1976).
- D. G. Madland and T. R. England, "Distribution of Independent Fission-Product Yields to Isomeric States," Los Alamos Scientific Laboratory report LA-6596-MS [ENDF-241] (November 1976).
- 5. D. G. Madland and Leona Stewart, "Light Ternary Fission Products: Probabilities and Charge Distributions," Los Alamos Scientific Laboratory report LA-6783-MS [ENDF-247] (April 1977).
- D. G. Foster, Jr. and T. R. England, "Time-Dependent Spectra of Photons and Spontaneous-Fission Neutrons for Applied Problems," <u>Invited Paper</u>, Trans. Am. Nucl. Soc. <u>23</u>, 551 (1976).
- 7. T. R. England and M. G. Stamatelatos, "Beta and Gamma Spectra and Total Decay Energies from Fission Products," Trans. Am. Nucl. Soc. <u>23</u>, 493 (1976).
- M. G. Stamatelatos and T. R. England, "Fission-Product Gamma-Ray and Photoneutron Spectra and Energy-Integrated Sources," NUREG-0155 [LA-NUREG-6345-MS] (Issued December 1976) (See also Addendum 1, March 1977).
- R. J. LaBauve, et al., "The Application of a Library of Processed ENDF/B-IV Fission-Product Aggregate Decay Data in the Calculation of Decay-Spectra," LA-7483-MS (September 1978).
- M. G. Stamatelatos and T. R. England, "Short Irradiation Fission-Product Beta Spectra and Total Energy: Calculations Versus Experiment," (ANS Summary accepted for ANS Annual Meeting June 12-17, 1977).

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(cont'd)

- 11. T. R. England, W. B. Wilson, and M. G. Stamatelatos, "Fission-Product Data for Thermal Reactors
 Part 1
 A Data Set for EPRI-CINDER Using ENDF/B-IV
 Part 2
 Users Manual for EPRI-CINDER Code and Data," Los Alamos Scientific Laboratory reports LA-6745-MS and LA-6746-MS (December 1975) [To be issued by EPRI ~ March 1977].
 12. W. B. Wilson and T. R. England, "Status of Fission-Product Data for Absorption
- Calculations," LA-UR-78-1452, (May 1978).
- 13. E. T. Jurney, P. J. Bendt, and T. R. England, "Fission Product Gamma Spectra," LA-7620-MS (January 1979).
- 14. T. R. England, R. E. Schenter, and F. Schmittroth," Integral Decay-Heat Measurements and Comparisons to ENDF/B-IV and V," NUREG/CR-0305 [LA-7422-MS] (August 1978).
- 15. W. B. Wilson, T. R. England, and R. J. LaBauve," DKPOWR: A Code for Calculating Fission-Product Decay Power (report in preparation).

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(same as in INDC(NDS)-95)

Laboratories Washington University, Dept. of Chemistry, St. Louis, MO., USA Los Alamos Scientific Laboratory, Group CNC-11, Los Alamos, NM USA

Names A. C. Wahl and K. Wolfsberg

Compilation and Independent yields and other data related to nuclear charge distribution in fission are being compiled and evaluated for low-energy fission processes (excitation energies up to ~ 20 MeV).

Purpose Development of systematics that will allow reliable estimates to be made for unmeasured independent yields and that will increase understanding of the fission mechanism.

Sources Journals, reports, preprints, and personal communications

- Method Data from various types of measurements are compared for evaluation of the reliability of the newer methods.
- Cooperation We are prepared to exchange files with other groups.
- Computer File Information is held in standard forms on computer files.
- Completions Continuous compilation
- Publications A. C. Wahl, A. E. Norris, R. A. Rouse, and J. C. Williams, in Proceedings of the Second International Atomic Energy Symposium on Physics and Chemistry of Fission, Vienna, Austria, 1969 (I.A.E.A.), p. 813.

K. Wolfsberg, Los Alamos Scientific Laboratory Report No. LA-5553-MS (1974).

A. C. Wahl, "Nuclear-Charge Distribution in Fission - Investigation of Systematics and Methods of Estimation of Independent Yields," Contribution to IAEA Petten Panel on Fission Product Nuclear Data - Sept., 1977. Published in: INDC(NDS)-87 (1978), 215.

111. RECENT PAPERS RELATED TO FPND

The papers listed below refer to activities relative to FPND which are not covered by the contributions contained in this issue. They are sorted according to

- 1. Fission yields
- 2. Neutron reaction cross sections
- 3. Decay data
- 4. Delayed neutron data
- 5. FP decay heat
- 6. FP data reviews.

Completeness of this Section has not yet been attempted.

1. Fission yields

Yields of Sn and Sb isotopes in the vicinity of the closed shell N=82 from reactor neutron induced fission of 232Th.

T. Izak-Biran, S. Amiel: J. Inorg. Nucl. Chem. <u>40</u> (1978) 937 [ind. yields of 131,132_{Sn.}, 131-133_{Sb}]

Fission product yields from 6-9 MeV neutron-induced fission of 235U and 238U.

T.C. Chapman, G.A. Anzelon, G.C. Spitale, D.R. Nethaway: Phys. Rev. C <u>17</u> (1978) 1089

 $[^{235}\text{U},\,^{238}\text{U}$ (n,f), $\text{E}_{n}\text{=}$ 6.0, 7.1, 8.1, 9.1 MeV; measured yields of 28 mass chains, table]

Absolute yields of 99 Mo and 111 Ag in the reactor neutron induced fission of 238 U.

S.G. Marathe, V.K. Rao, V.K. Bhargava, S.M. Sahakundu, R.H. Iyer:

J. Inorg. Nucl. Chem. <u>40</u> (1978) 1981

The fine structure of the fission yields of heavy nuclei.

K.A. Petrzhak, E.V. Platygina, V.F. Teplykh: Soviet Atomic Energy <u>42</u> (1977) 337

 $[^{232}\text{Th}, ^{235}\text{U}, ^{239}\text{Pu} (n,f), E_n=15 \text{ MeV}; relative yields of 131-135Xe isotopes}]$

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A general correlation for independent fission product
yield uncertainties.
B.I. Spinrad, C.H. Wu:
Nucl. Scie. and Engg. <u>66</u> (1978)421
[correlation between experiment/theoretical yield ratios
and the distance of the FP from Z<sub>p</sub>]
Prediction of fission product mass yields and related
parameters.
T. Yamamoto, K. Sugiyama:
Ann. Nucl. Energy <u>5</u> (1978) 613
[<sup>232</sup>U, <sup>238</sup>Pu thermal fission, <sup>234,236</sup>U, <sup>240,242</sup>Pu fission
with E<sub>n</sub>=2MeV; A=71-163 chain yields predicted by semi-
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with $E_n=2MeV$; A=71--163 chain yields predicted by semiempirical method]

Analytic formula for isobaric charge distribution of 235 U fission products by thermal neutrons, taking parity effects into account.

A.A. Byalko, V.M. Zhivun, V.V. Kovalenko, A.B. Koldobskij: Soviet Atomic Energy <u>43</u> (1978) 657

Light-charged-particle emission in keV neutron-induced fission of 239Pu.

B. Krishnarajulu, S. Sen, G.K. Mehta J. Phys. G <u>5</u> (1979) 319

2. Neutron reaction cross sections

Neutron capture and transmission measurements on fission product Pd-107.

U.N. Singh, R.C. Block, Y. Nakagome Nucl. Sci. and Engg. <u>67</u> (1978) 54 $[E_n \leq 700 \text{ eV}; \text{ table of resonance parameters}]$ Neutron total cross section measurement on 140_{Ce}. H.S. Camarda Phys. Rev. C <u>18</u> (1978) 1254

[E_n=20-240 keV; table of resonance-parameters]

Capture cross section measurements of ¹⁴¹Pr.

P.S. Feigenbaum, U.N. Singh, R.C. Block

 $[E_n = 20 \text{ eV} - 60 \text{ keV}; \text{ table of newly observed resonances}]$

Uncertainties on a pseudo fission product for fast power reactors.

G. Langlet, P. Copps, J.P. Doat: NEACRP-L-190 (1977)

A review of multigroup nuclear cross section processing. Proceedings of a Seminar Workshop, Oak Ridge, Tenn., March 14-16, 1978.

Compiled by D.K. Turbey, H.R. Hendrickson: ORNL/RSIC-41 (Oct. 1978)

Graphs of neutron capture cross sections of fission product isotopes from FPLIB-65/ENDF/B-IV.

M. Mattes: ZAED-10-1 (1978)

[FPLIB-65 is a 65-group cross section library generated from ENDF/B-IV]

3. Decay data

Decay energies of gaseous fission products and their daughters for A=88 to 93 and A=138 to 192.

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[Calculation of envelopes for d.n. spectra of ⁸⁵As, ^{88,89}Br, 135Sb, ¹³⁶Te, ^{138,139}I, ¹⁴³Cs; compared to experiments]

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[Contents of Session I: Sensitivities and Uncertainty Analysis for Fast and Thermal Reactors:

- "The Use of Cross-Section Sensitivities in the Analysis of Fast Reactor Integral Parameters," P.J. Collins and N.J. Lineberry (Argonne National Laboratory, Idaho).
- 2. "Advances in Fast Reactor Sensitivity and Uncertainty Analysis," J.H. Marable and C.R. Weisbin (Oak Ridge National Laboratory).
- 3. "Controlled Cross-Section Adjustment by Integral Data," U. Salmi, J.J. Wagschal, A. Ya'ari, and Y. Yeivin (Racah Institute of Physics, The Hebrew University, Israel)
- 4. "Sensitivity Analysis Applied to the Calculation of Detector-Response Kernels," W.H. Scott, Jr. (Science Applications, Inc., La Jolla), and P.J. McDaniel, J.H. Renken, and S.A. Wright (Sandia Laboratories, New Mexico)

- "Sensitivity of Water Reactor Fuel Cycle Parameters and Costs to Nuclear Data," M. Becker, D.R. Harris, B. Quan, and J.M. Ryskamp (Rensselaer Polytechnic Institute).
- 6. "Nonlinear Model Fitting of Thermal Neutron Data and its Application in Resonance Parameter Uncertainty Analysis," J.K. Thompson (Battelle Pacific Northwest Laboratories).
- 7. "Fission Product Decay Power and Uncertainties After Realistic Reactor Operating Histories," T.J. Trapp (Battelle Pacific Northwest Laboratory), and B.I. Spinrad (Oregon State University).]

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