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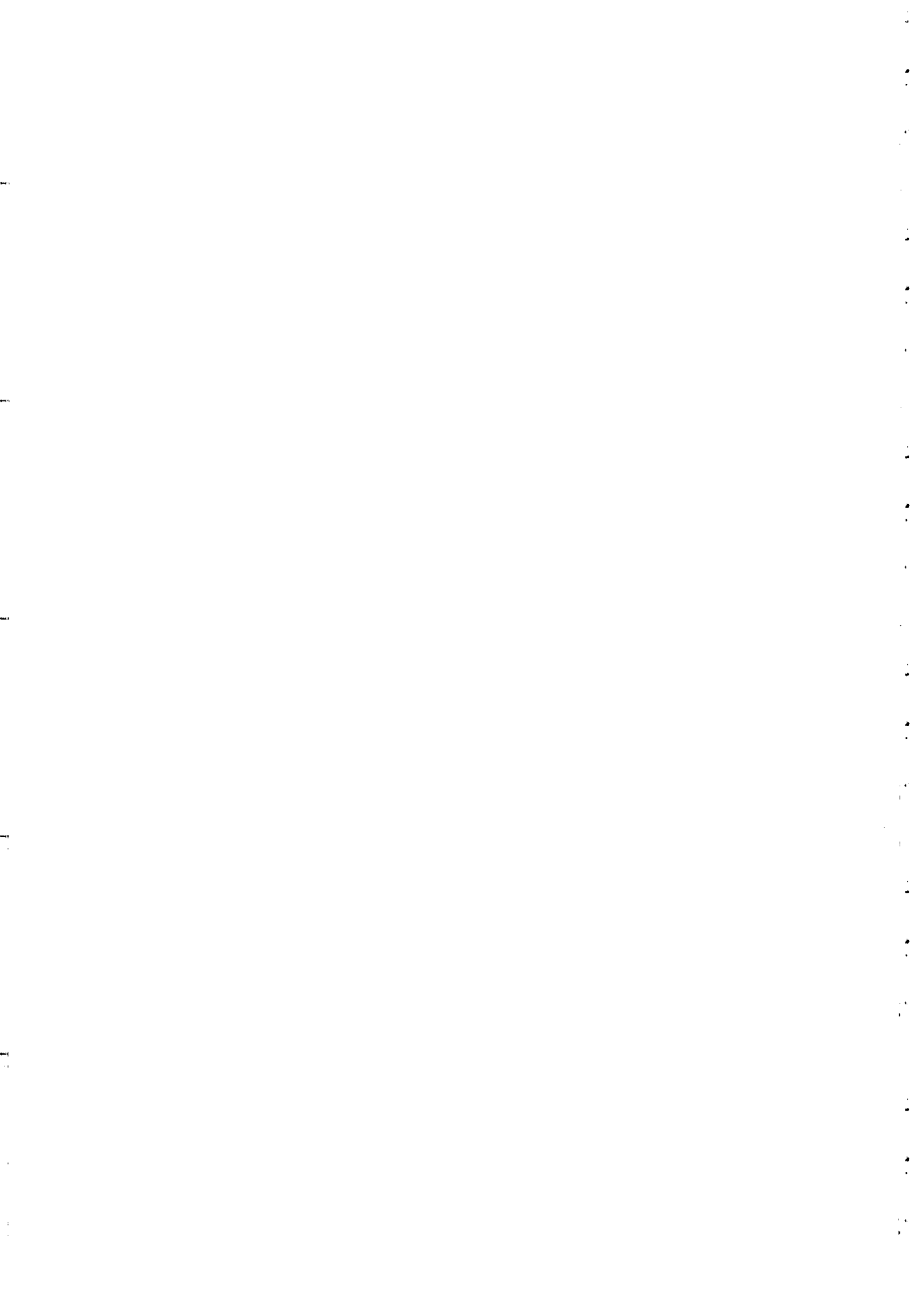
Workshop on
**Nuclear Reaction Data and Nuclear Reactors:
Physics, Design and Safety**

13 March - 14 April 2000

Miramare - Trieste, Italy

Long Term Radiotoxicity

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**WORKSHOP ON NUCLEAR REACTION DATA
AND NUCLEAR REACTORS:**

Physics, Design, Safety

Trieste, ITALY

Prof. Igor SLESSAREV
CEA, FRANCE

LONG TERM RADIOTOXICITY

April 12 2000

**LONG -TERM RADIOACTIVITY AND
RADIOTOXICITY OF NUCLEI
as the result
of NUCLEAR ENERGY PRODUCTION**

GROUPS of LONG-TERM RADIOACTIVE WASTES:

1.ACTINIDES

2.Long-Lived FISSION PRODUCTS (LLFP)

3.LANTANIDES (LA)

(in connection with discharge fuel reprocessing needs)

Quantitative measure of RADIOTOXICITY

Factor of dose "par ingestion" or "par inhalation"

$$R(\text{Sv}) = F_d (\text{Sv/Bq}) \times A(\text{Bq})$$

Table. Periods and factor of dose "par ingestion"

Nuclides	Life period (y)	Emission	F_d
Tc-99	2 10⁵	β-	0.64 10⁻⁹
I-129	1.6 10⁷	β-	0.11 10⁻⁶
Pu-238	88	α	0.23 10⁻⁶
<u>Pu-239</u>	2.4 10⁴	α	<u>0.25 10⁻⁶</u>
Am-241	432	α	0.20 10⁻⁶
Am-243	7.4 10³	α	0.20 10⁻⁶
Cm-244	18	α	0.16 10⁻⁶

TOXICITY

TABLEAU

On donne les logarithmes décimaux des facteurs de danger ingestif (en Sv/g initial du noyau considéré) à diverses échéances. Les familles de décroissance pour les actinides sont les suivantes:

- 1 décroît sur Pb206
- 2 " " Pb207
- 3 " " Pb208
- 4 " " Bi209

Th²³²

-2.28 -2.28 -2.28 -2.29 -2.28 -2.28 -2.28

Noyau	Famille	10 ² ans	10 ³ ans	10 ⁴ ans	10 ⁵ ans	10 ⁶ ans	10 ⁷ ans	10 ⁸ ans
U232	3	5.22	1.35	-3.74	-	-	-	-
U233	4	1.47	1.78	2.40	2.43	0.72	-16.43	-
U234	1	1.22	1.24	1.65	2.35	1.57	-9.49	-
U235	2	-2.19	-1.78	-0.96	-0.31	-0.26	-0.26	-0.30
U236	3	-0.79	-0.79	-0.79	-0.80	-0.81	-0.92	-1.88
U238	1	-3.08	-3.08	-3.06	-2.53	-1.66	-1.62	-1.62
Np237	4	1.44	1.44	1.45	1.56	1.63	0.37	-12.29
Pu236	3	5.23	1.36	-3.73	-	-	-	-
Pu238	1	5.46	2.41	1.64	2.34	1.56	-9.49	-
Pu239	2	3.43	3.41	3.30	2.17	-0.27	-0.27	-0.31
Pu240	3	3.99	3.94	3.53	-0.40	-0.81	-0.93	-1.89
Pu241	4	5.13	4.50	1.44	1.55	1.62	0.36	-12.30
Pu242	1	2.20	2.20	2.20	2.12	1.40	-1.63	-1.63
Pu244	3	-0.14	-0.10	0.08	0.17	0.16	0.13	-0.18
Am241	4	5.11	4.49	1.44	1.55	1.62	0.36	-12.30
Am242m	1	5.61	3.97	1.79	2.30	1.53	-2.39	-2.39
Am243	2	3.94	3.92	3.68	2.33	-0.27	-0.28	-0.32
Cm242	1	5.45	2.40	1.63	2.33	1.55	-9.50	-
Cm243	2	5.13	3.41	3.30	2.17	-0.27	-0.28	-0.32
Cm244	3	4.68	3.94	3.52	-0.40	-0.82	-0.93	-1.90
Cm245	4	3.93	4.11	3.85	1.59	1.62	0.35	-12.31
Cm246	1	4.13	4.07	3.51	2.12	1.40	-1.63	-1.64
Cm247	2	0.59	0.62	0.82	1.06	1.06	0.89	-0.21
Cm248	3	2.84	2.84	2.83	2.75	1.96	0.13	-0.19

Pu-231

4.06 4.06 3.98 3.16 -5.1 -27.7

Noyau	10 ² ans	10 ³ ans	10 ⁴ ans	10 ⁵ ans	10 ⁶ ans	10 ⁷ ans	10 ⁸ ans
Se79	0.77	0.77	0.73	0.31	-3.85	-	-
Zr93	-1.41	-1.41	-1.41	-1.43	-1.60	-3.38	-
Nb94	0.99	0.97	0.84	-0.50	-13.84	-	-
Tc99	-0.67	-0.67	-0.69	-0.81	-2.08	-14.80	-
Pd107	-3.15	-3.15	-3.15	-3.16	-3.20	-3.62	-7.78
Sn126	0.76	0.76	0.73	0.46	-2.25	-	-
I129	-0.32	-0.32	-0.32	-0.32	-0.33	-0.51	-2.23
Cs135	-1.09	-1.09	-1.09	-1.10	-1.22	-2.40	-14.18

ACTIVITY

TABIEAU

On donne les logarithmes décimaux des activités (en Bq/kg initial du noyau considéré) à diverses échéances. Les familles de décroissance pour les actinides sont les suivantes:

- | | | |
|---|-------------|-------|
| 1 | décroit sur | Pb206 |
| 2 | " | Pb207 |
| 3 | " | Pb208 |
| 4 | " | Bi209 |

Noyau	Famille	10 ² ans	10 ³ ans	10 ⁴ ans	10 ⁵ ans	10 ⁶ ans	10 ⁷ ans	10 ⁸ ans
U232	3	15.39	11.52	-2.72	-	-	-	-
U233	4	11.59	11.79	12.31	12.34	10.62	-6.52	-
U234	1	11.36	11.37	11.58	12.12	11.33	0.27	-
U235	2	8.21	8.24	8.47	8.90	8.95	8.94	8.90
U236	3	9.38	9.38	9.38	9.38	9.37	9.25	8.20
U238	1	7.57	7.57	7.57	7.70	8.21	8.24	8.23
Np237	4	10.71	10.72	10.75	11.10	11.34	10.08	-2.58
Pu236	3	15.40	11.53	-2.72	-	-	-	-
Pu238	1	14.46	11.67	11.57	12.12	11.32	0.26	-
Pu239	2	12.36	12.35	12.24	11.11	8.94	8.94	8.90
Pu240	3	12.92	12.88	12.47	9.41	9.36	9.25	8.20
Pu241	4	14.16	13.42	10.74	11.09	11.33	10.07	-2.59
Pu242	1	11.16	11.16	11.15	11.08	10.36	8.23	8.22
Pu244	3	9.30	9.31	9.38	9.42	9.42	9.41	9.18
Am241	4	14.03	13.41	10.74	11.09	11.33	10.07	-2.59
Am242m	1	14.75	13.05	11.51	12.03	12.41	7.47	7.46
Am243	2	13.17	13.13	12.84	11.26	8.93	8.93	8.89
Cm242	1	14.45	11.67	11.56	12.11	11.31	0.26	-
Cm243	2	14.23	12.34	12.23	11.10	8.93	8.93	8.89
Cm244	3	13.85	12.87	12.46	9.40	9.35	9.24	8.19
Cm245	4	13.12	13.22	12.94	11.08	11.33	10.07	-2.59
Cm246	1	13.05	12.99	12.44	11.08	10.36	8.22	8.22
Cm247	2	9.84	9.87	10.06	10.23	10.22	10.05	8.99
Cm248	3	11.20	11.20	11.19	11.11	10.35	9.40	9.17

PHYSICS of TRANSMUTATION: POTENTIAL, principal CONSTRAINTS

Neutronics plays a most important role in transmutation potential

Fig. "Transmutation sieve"

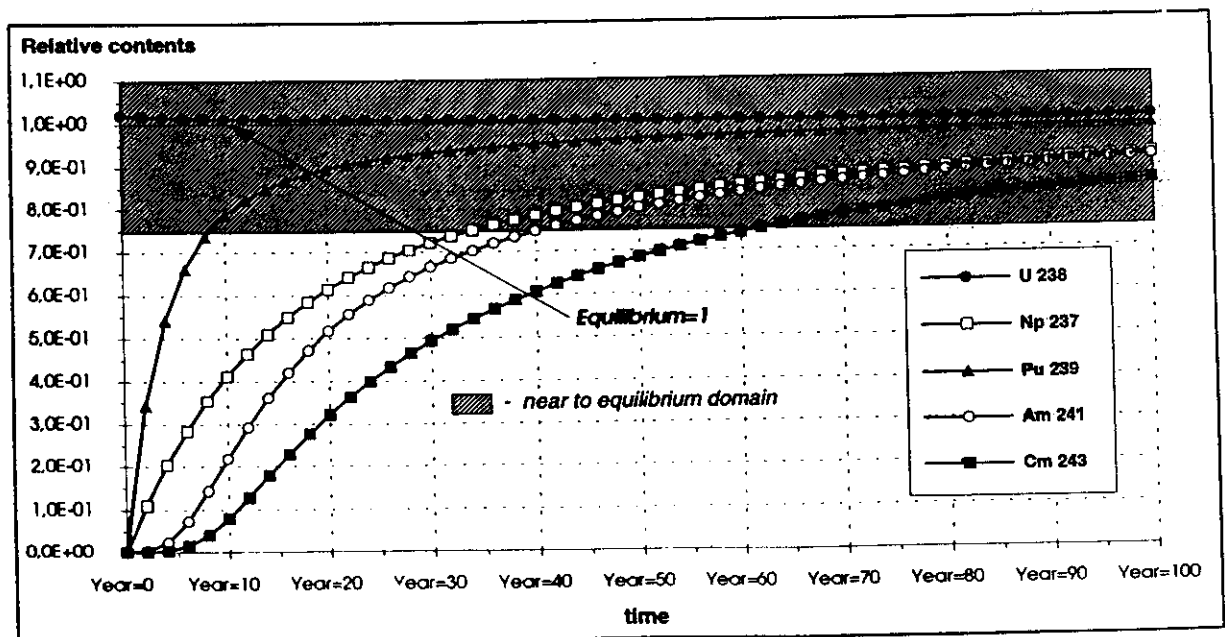
Overall neutron balance for nuclide family library (at equilibrium)

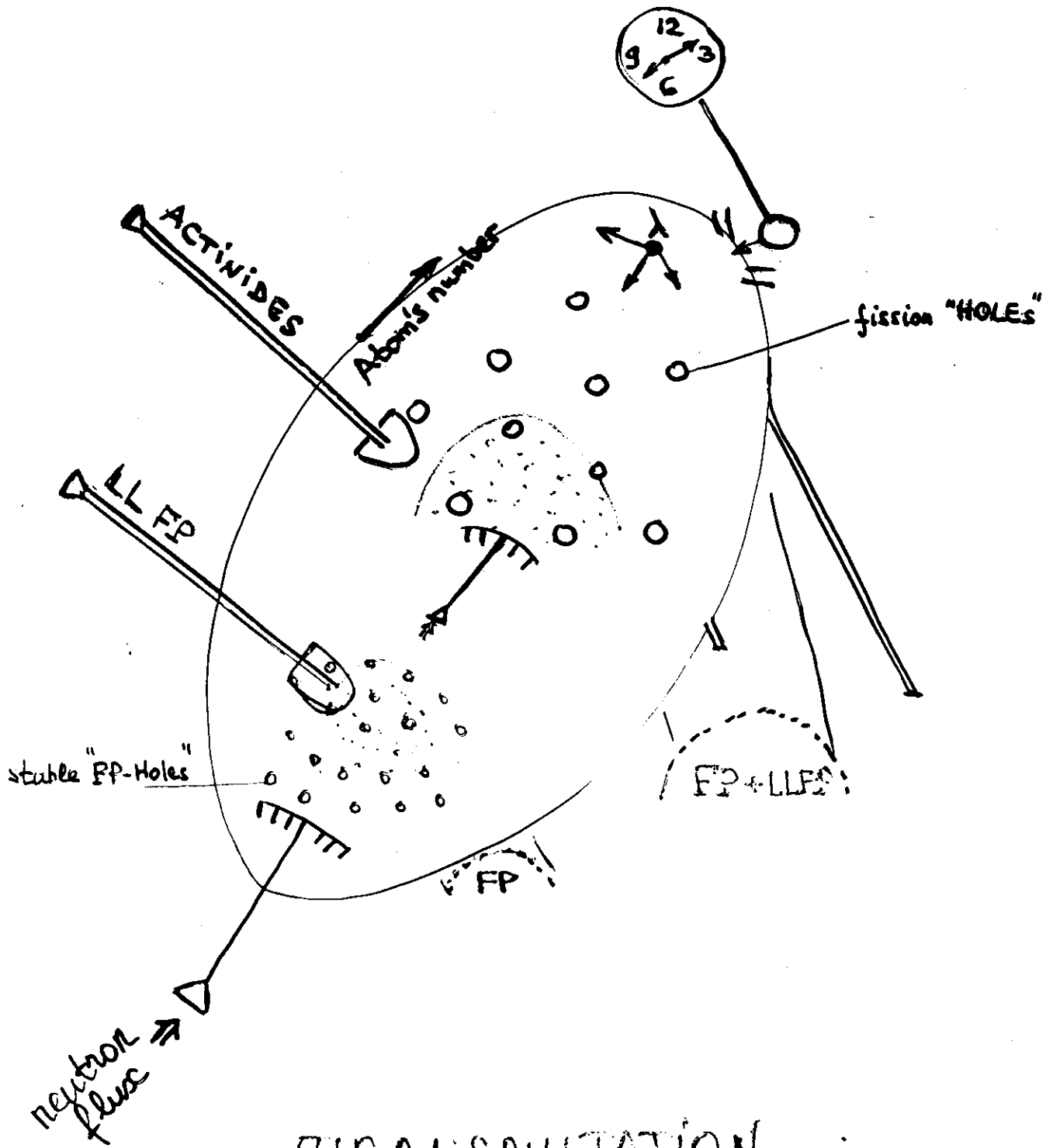
Equilibrium:

"Family" is a set of nuclei (including the initial nucleus) which have been produced both by transmutation in a neutron flux Φ of the nucleus as well as of its subsequent products, and by decay of nuclei.

The equilibrium state of a family in a given nuclear system is the relative concentration distribution N of this family members for the asymptotic ($t \rightarrow \infty$) condition when a given system is continuously fed by the initial nucleus only

The analysis of a transient time shows that in the case of the main families (such as ^{238}U , ^{235}U) the rate of convergence to equilibrium even for standard neutron fluxes level is high enough.





TRANSMUTATION
"SIEVE"

Equilibrium concentrations have been obtained as solution of the following equations (BATEMAN):

$$dN_J/dt = (M\Phi - \Lambda - L) \cdot N_J - S_J = 0$$

where :

M -- the operator of a nucleus transmutation by means of the corresponding neutronic processes (capture, fission, (n,2n), (n,3n) reactions, etc.),

Φ -- a neutron flux,

Λ -- the operator of decay : α , β^+ , β^- , etc.,

L -- the operator of nuclei "*losses*"
(more generally, the operator of nuclei extraction rate) from fuel cycle
(e.g. losses during reprocessing),

S_J -- a source of the J-family (reactor feed rate).

Family Overall Neutron Consumption

Decay, irradiation and regular discharge (or losses) of every nucleus J (family "father") in a neutron flux produce at equilibrium (see equation (1)) the J-nucleus family.

Then one can calculate the total number of neutrons D_J which have been consumed by this family to be burnout transmuted to either stable or short lived nucleus and then extracted from this system).

If a given system is fed by a set of J-nuclei then the potential neutron surplus G will be defined by a sum of all D_J (neutron consumption), the neutron losses of this system and supplementary neutrons (μ) (in ADS) :

$$G (\text{neutrons / fission}) = -\sum_J X_J \cdot D_J - 0.3 + \mu = \sum_J X_J \cdot G_J + \mu = G_0 + \mu .$$

where X_J is a fraction of J-nucleus in fuel feed.

SPECIFIC FEATURES OF TRANSMUTATION. NEUTRONICS

Actinide Toxicity after Fuel Irradiation

Individual nuclear family toxicities depends on

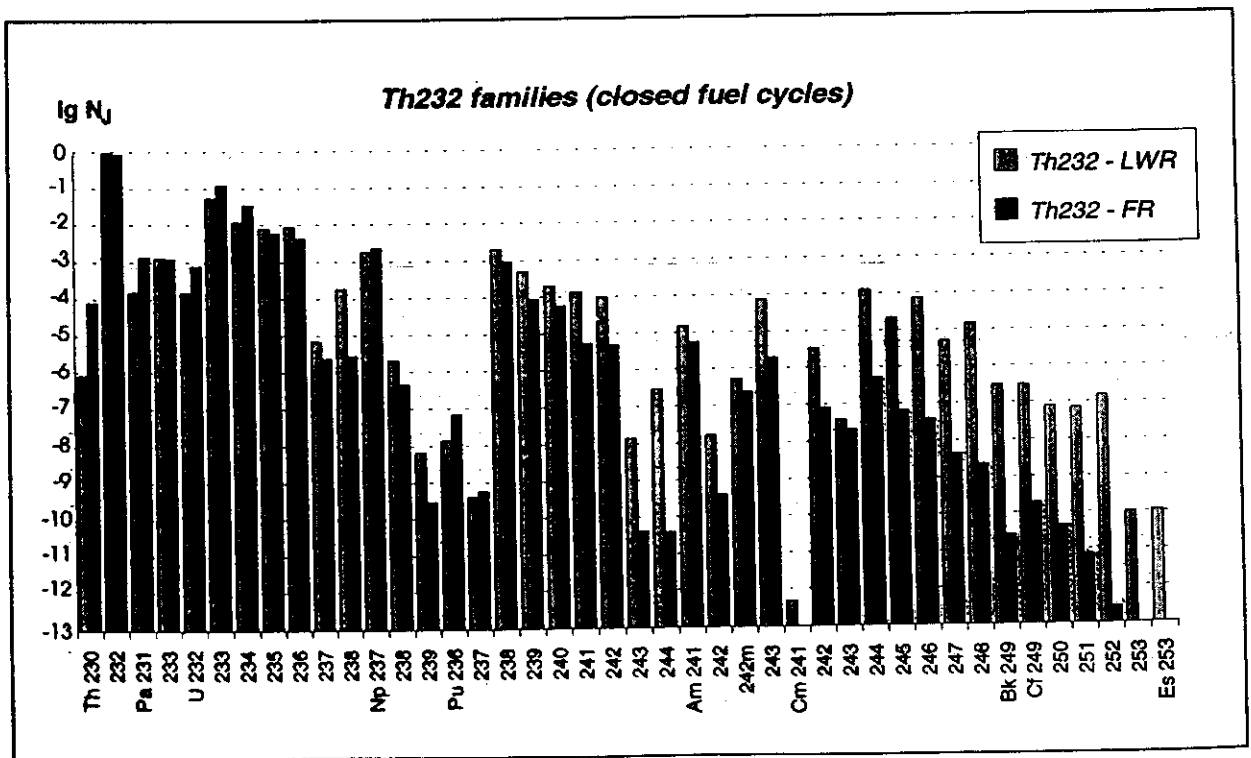
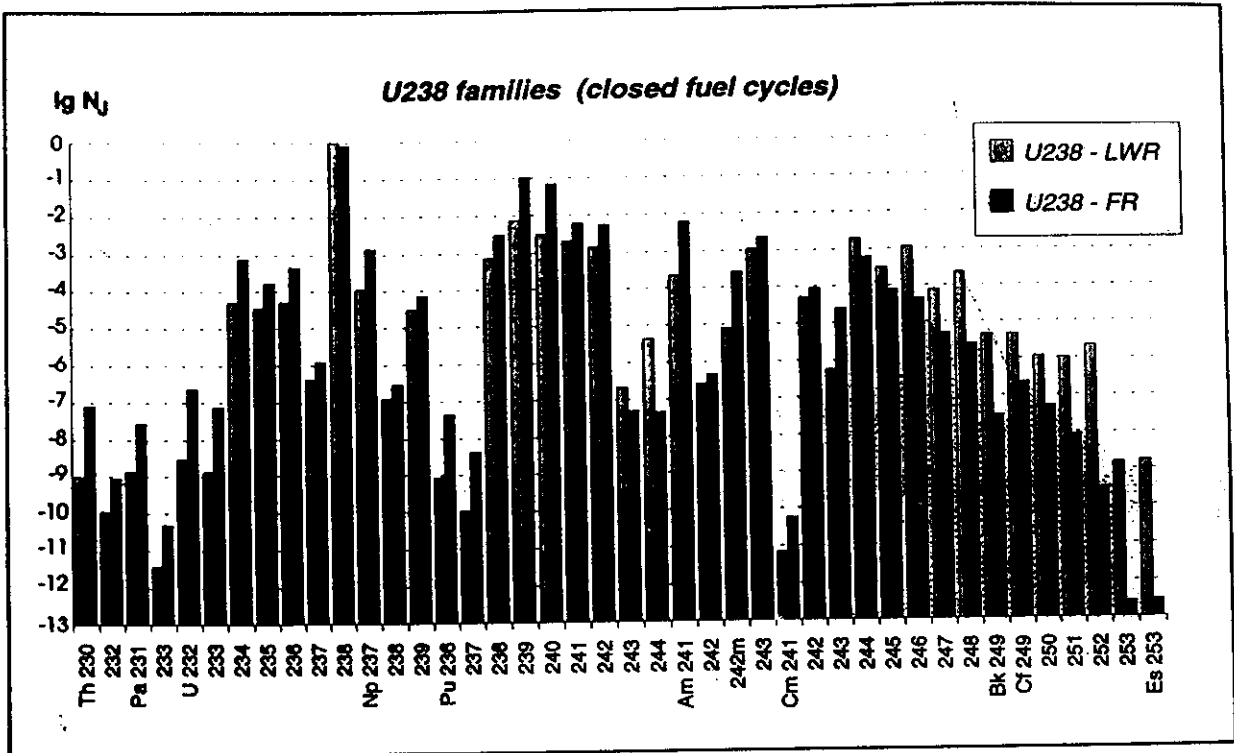
- *initial nuclide ("father")*
- *neutron flux level,*
- *neutron spectrum*
- *fuel cycle parameters*
(burnup via discharge period, losses, etc.)

TABLE II

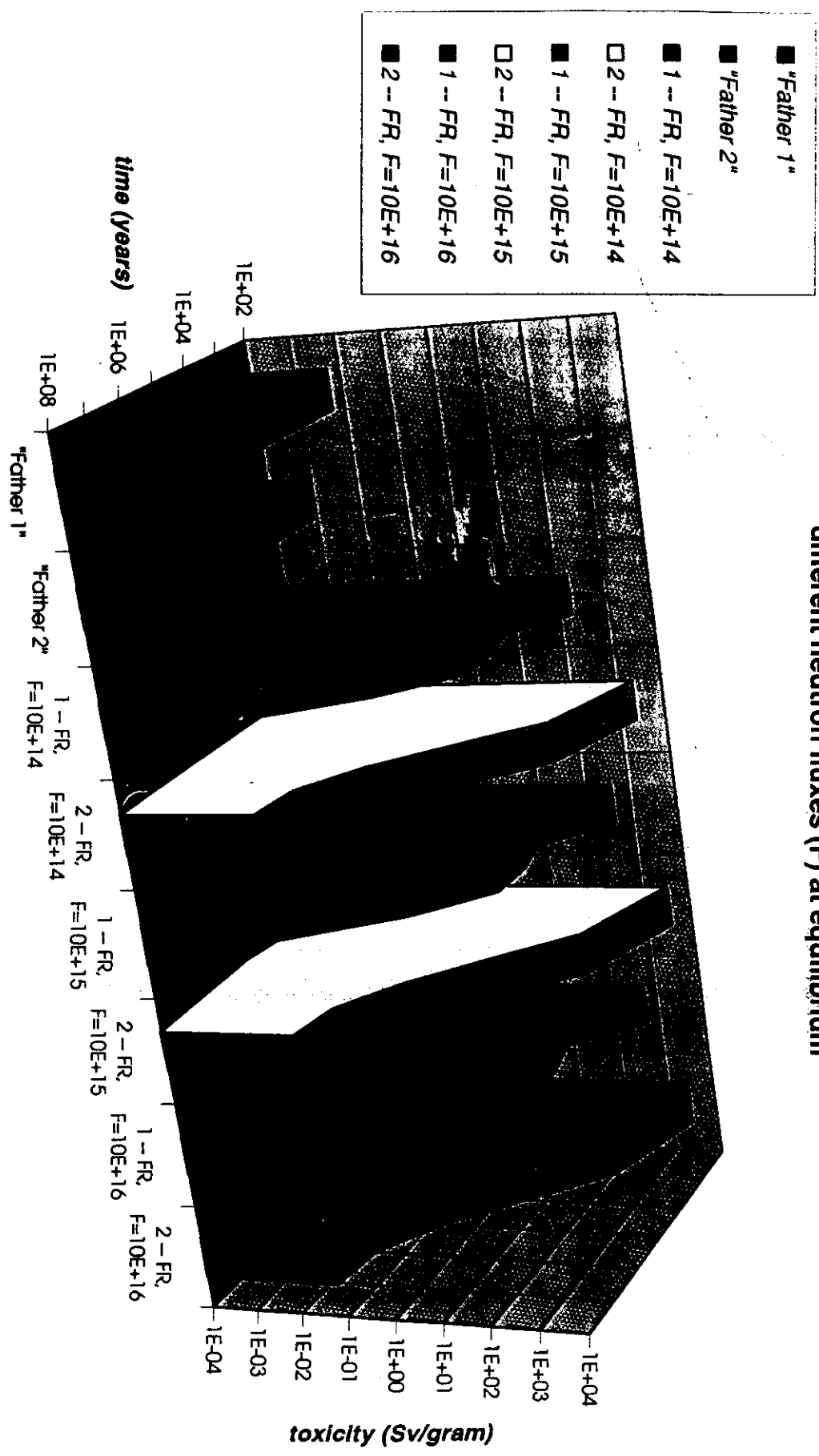
Normalised toxicities of J-families T_J at equilibrium (Sv/gram); T_J have been averaged on Short ($10^2 \div 10^4$ years) and Long ($10^2 \div 10^6$ years) time intervals (negligible losses rates)

	Thermal Reactors ($\Phi = 10^{14}$ n/cm ² s)		Fast Reactors ($\Phi = 10^{15}$ n/cm ² s)		Super-thermal Reactors ($\Phi = 10^{16}$ n/cm ² s)		Natural Toxicity ($\Phi = 0$ n/cm ² s)	
	T_J Short	T_J Long	T_J Short	T_J Long	T_J Short	T_J Long	T_J Short	T_J Long
"fathers" of families								
²³²Th	21	4.6	38	11	41	3.8	0.005 2	0.0052
²³¹Pa	5364	47	7583	31	3633	36	10570	0.3
²³³U	221	53	117	62	168	23	111	46
²³⁵U	372	19	92	19	173	2.5	0.037	0.48
²³⁸U	97	1	841	5.7	275	3.5	0.000 8	0.0068
²³⁷Np	1436	71	318	65	1984	55	28	39
²³⁸Pu	2120	77	751	86	4411	63	164	90
²³⁹Pu	4851	46	4290	28	4610	58	2300	13
²⁴⁰Pu	5952	50	6025	22	4723	59	5730	0.41
²⁴¹Pu	5834	62	4506	63	4661	62	1430	38
²⁴²Pu	5139	58	3106	56	5883	60	160	63
²⁴¹Am	3869	72	3826	67	6019	63	1410	38
^{242m}Am	6222	60	5128	74	6125	62	1180	85
²⁴³Am	6277	55	6557	32	6120	61	6500	17
²⁴²Cm	2051	78	798	86	2626	86	161	88
²⁴⁴Cm	6135	53	6264	19	6058	61	6140	0.4
²⁴⁵Cm	5942	101	7939	71	5876	97	9640	52

Equilibrium concentration distributions

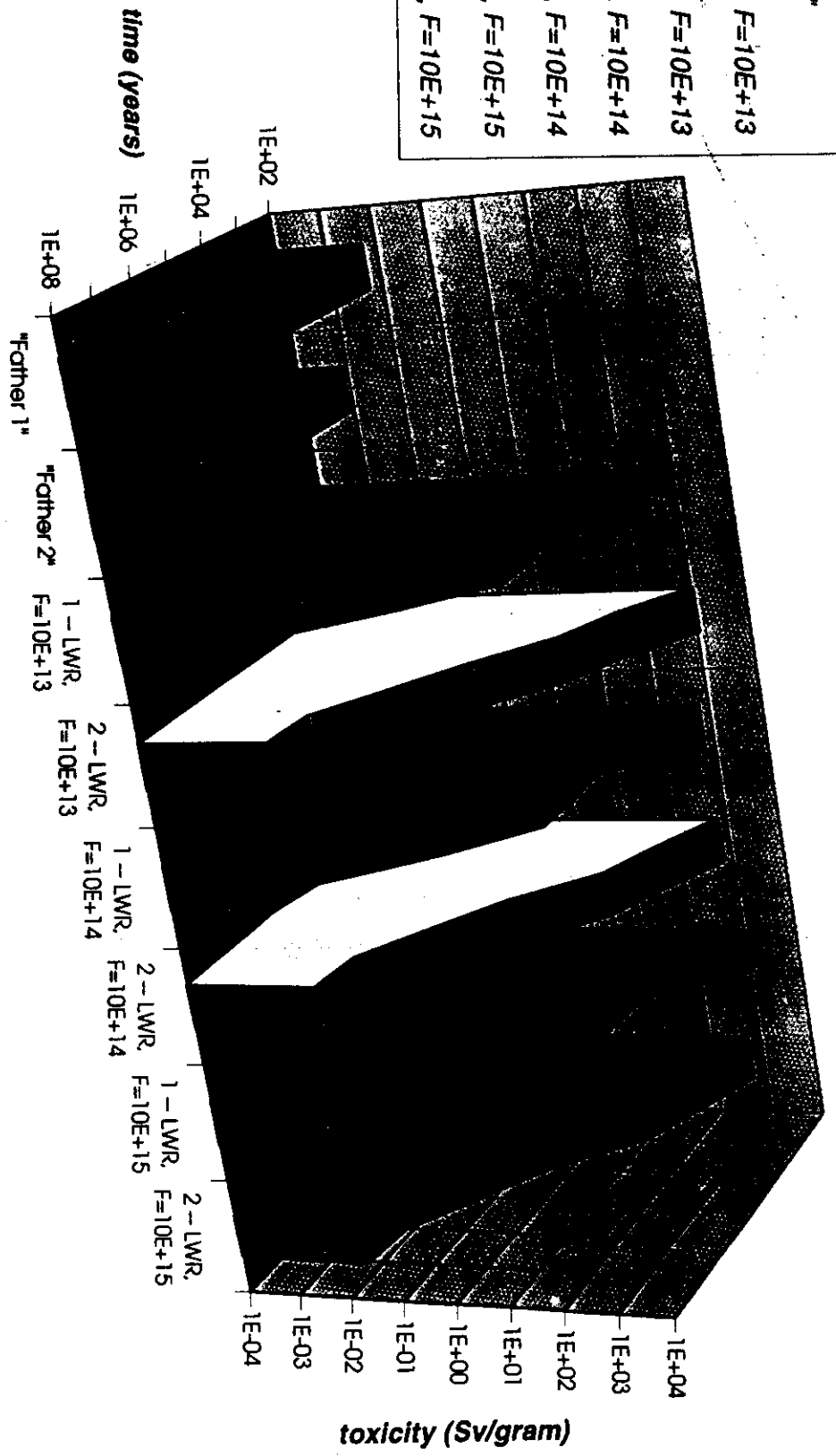


Toxicity of Th232 ("father 1"), U238 ("father 2") and their "families" in FR with different neutron fluxes (F) at equilibrium

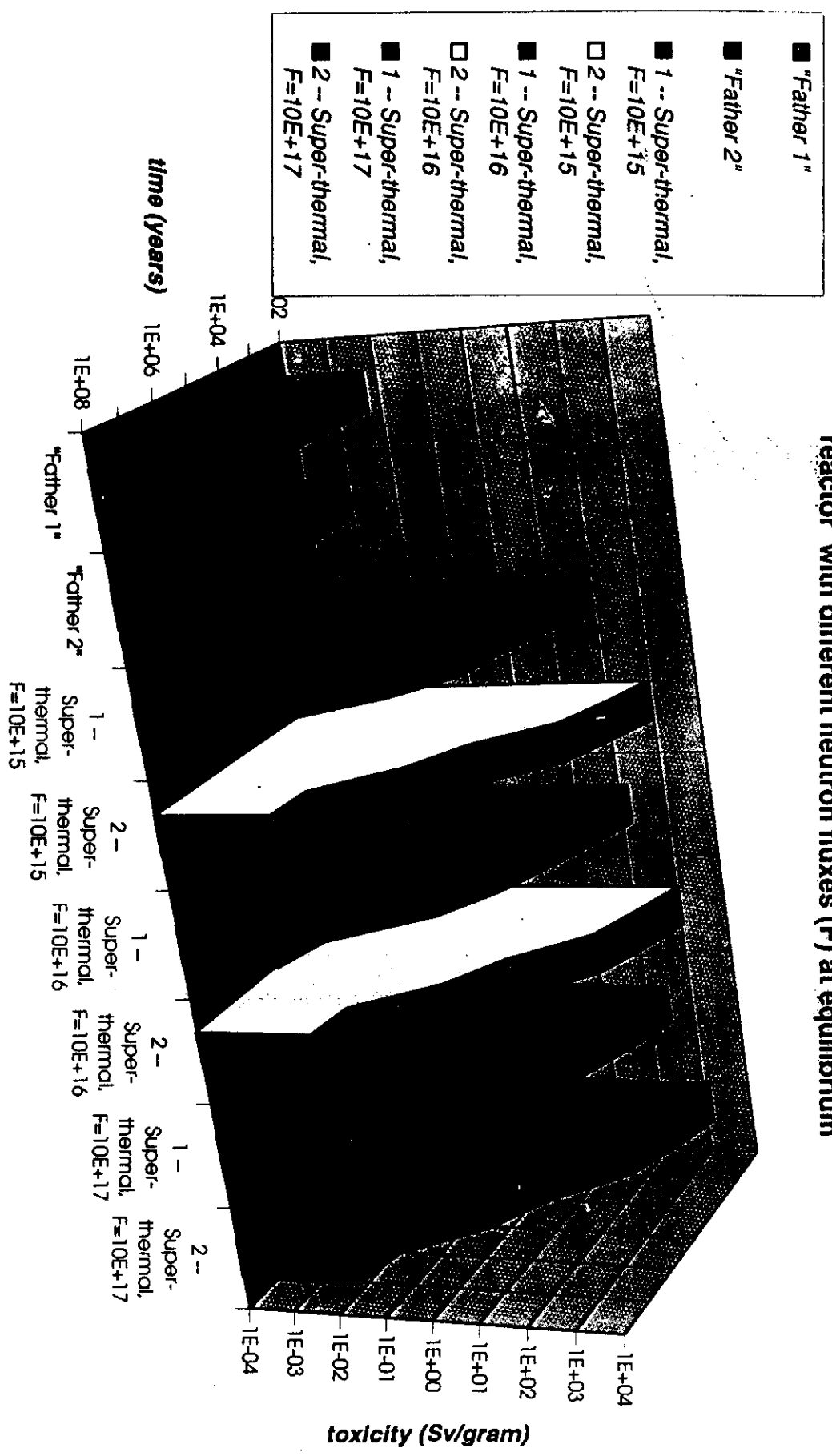


Toxicity of Th232 ("father 1"), U238 ("father 2") and their "families" in LWR with different neutron fluxes (F) at equilibrium

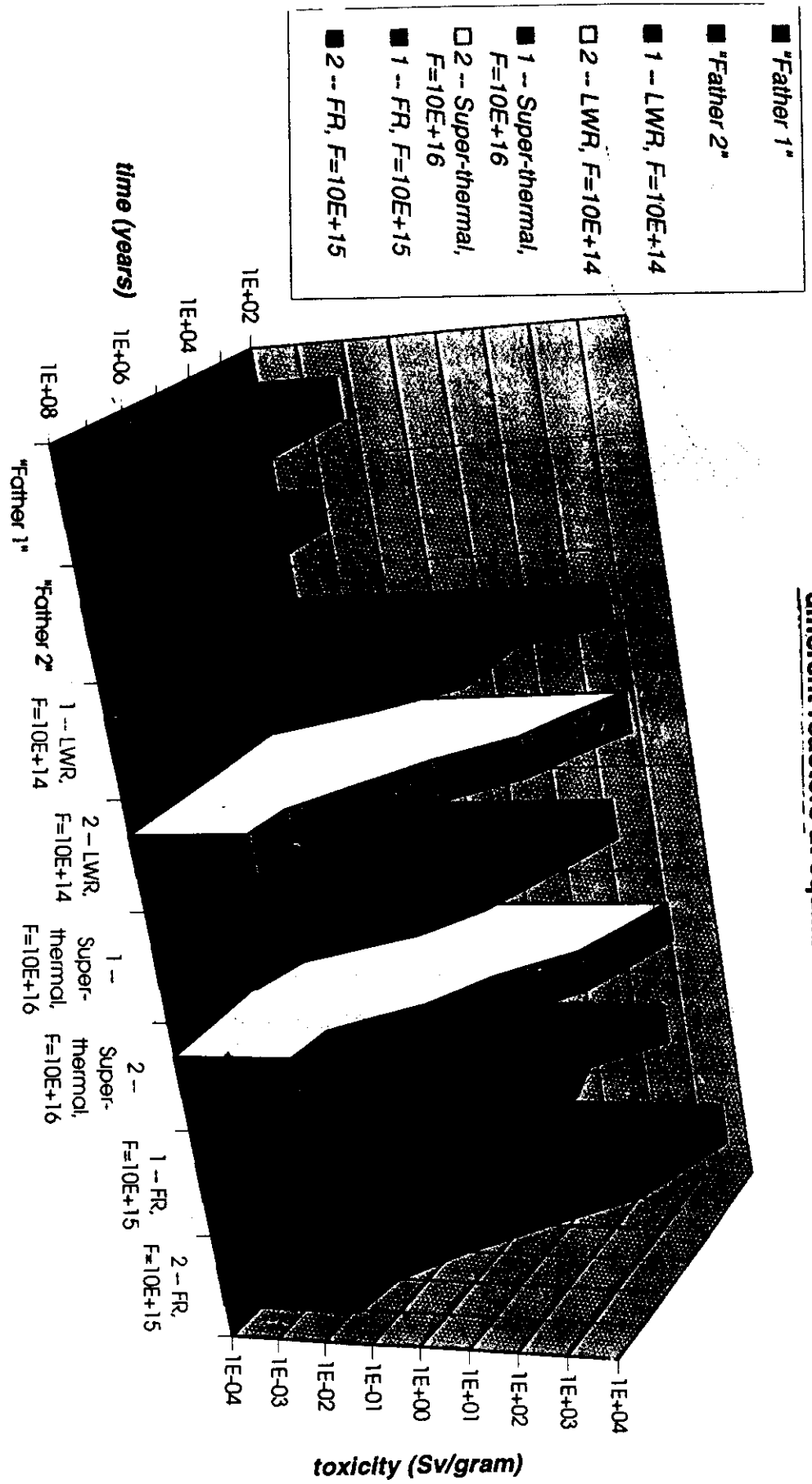
- "Father 1"
- "Father 2"
- 1 - LWR, F=10E+13
- 2 - LWR, F=10E+13
- 1 - LWR, F=10E+14
- 2 - LWR, F=10E+14
- 1 - LWR, F=10E+15
- 2 - LWR, F=10E+15



Toxicity of Th232 ("father 1"), U238 ("father 2") and their "families" in Super-thermal reactor with different neutron fluxes (F) at equilibrium

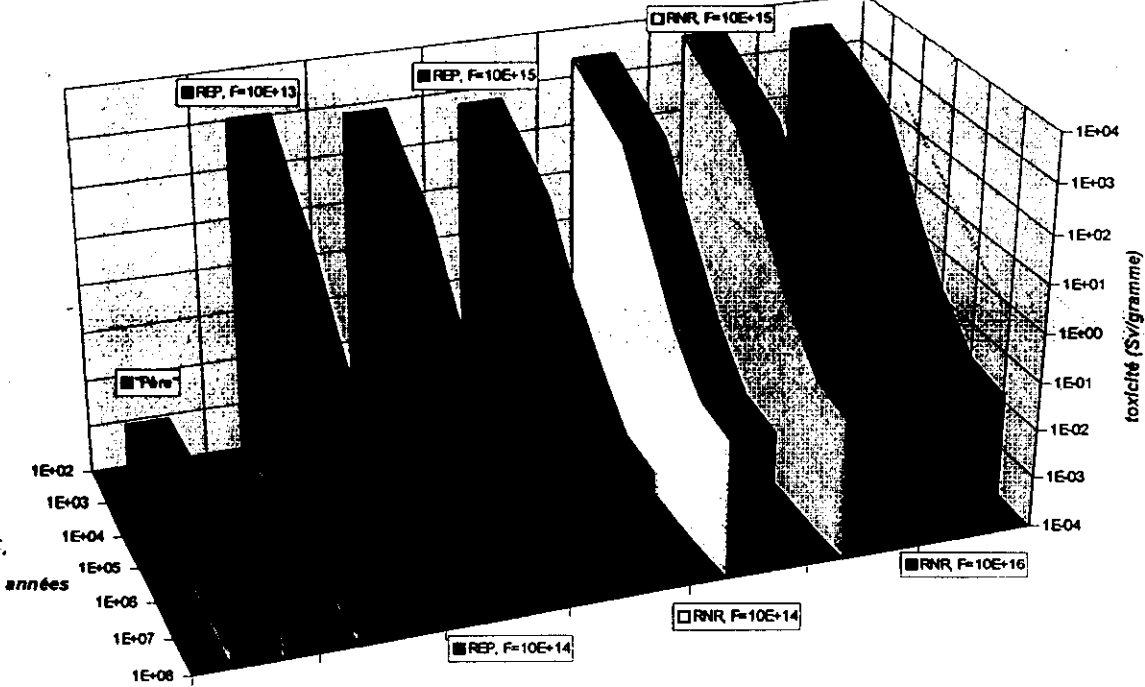


Toxicity of Th232 ("father 1"), U238 ("father 2") and their "families" in different reactors at equilibrium

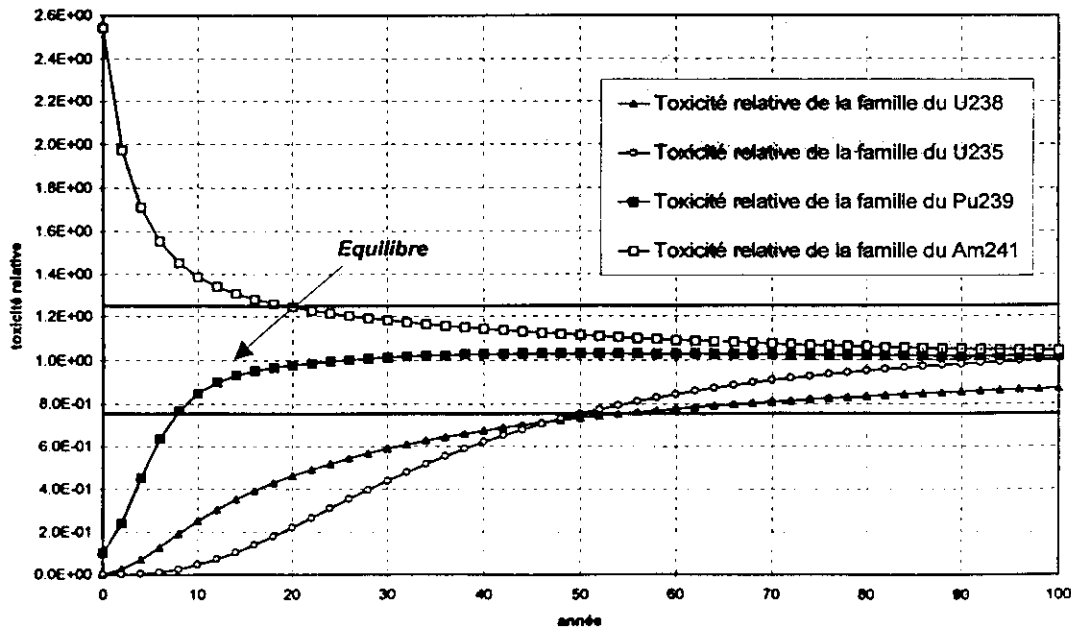


Toxicités à l'équilibre

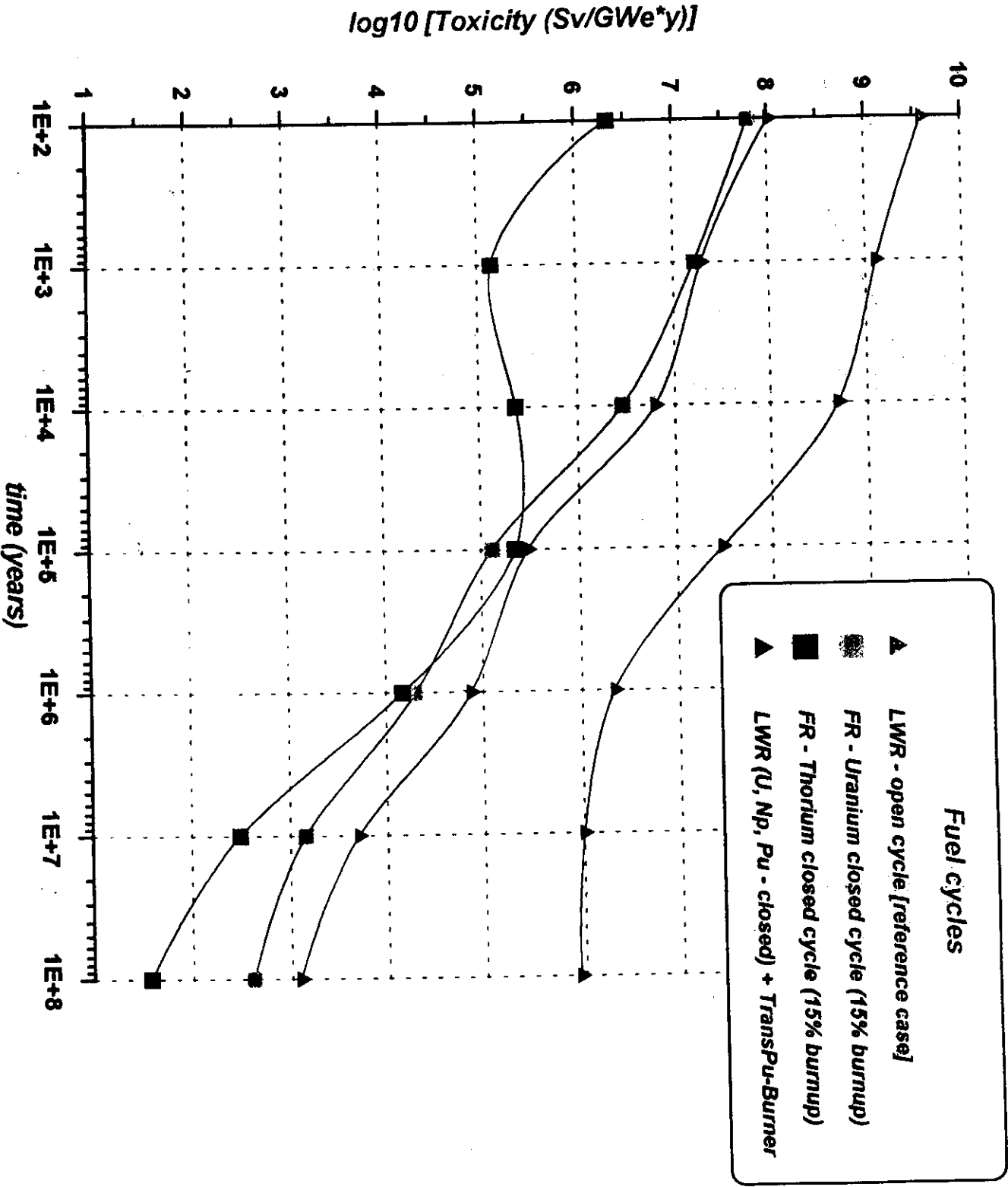
Toxicité du U238 ("père") et de sa "famille" dans les différents réacteurs et les flux neutroniques (F) en équilibre



Convergence relative de la toxicité (après 100 ans de refroidissement) pour les familles principales en REP standard



Fuel Waste Toxicity reduction potential



LLFP TRANSMUTATION

is IMPORTANT

because of

- non-negligible toxicity,
- essential MOBILITY

Time T_J^{transm} needed to incinerate half of initial mass which is a function of the cross-section $\sigma'_{n,\gamma}$ (barns) and of the neutron flux Φ (n/cm²s) is:

$$T_J^{transm} = \frac{\ln 2}{\sigma'_{n,\gamma} \Phi \times 3.16 \times 10^7} \text{ years}$$

Transmutation under neutron flux can be reasonable

if $T_{1/2} \gg T_J^{transm}$ and

if this time is technologically acceptable

TABLE I

Parameters of long lived nuclei to be eventually transmuted in a fast (E_n (neutron energy) = 0.2 MeV, JEF-2.2) and "thermal" ($E_n = 1$ eV, JEF-2.2) spectra with standard flux levels : $\Phi = 10^{15}$ (n/cm²s) $\Phi = 10^{14}$ (n/cm²s) correspondingly

Isotopes, J	$\sigma_{n,\gamma}^J$ (barns)		$T_{1/2}$ (years)	T_J^{transm} (years)		Radio-toxicity in (Sv/g) at $t = T_J^{transm}$	Recommendation to transmutation
	fast spectrum	thermal spectrum		fast spectrum	thermal spectrum		
⁷⁹ Se	0.03	0.1	6.5×10^4	7.3×10^2	2.2×10^3	6.0	questionable
⁹⁰ Sr	0.01	0.14	29	2.2×10^3	1.6×10^3	-	non-transmutable
⁹³ Zr	0.03	0.28	1.5×10^6	730	790	0.04	questionable
⁹⁴ Nb	0.04	2.2	2.0×10^4	5.5×10^2	1×10^2	9.0	questionable or transmutable
⁹⁹ Tc	0.2	4.3	2.1×10^5	110	51	0.2	transmutable
¹⁰⁷ Pd	0.5	0.3	6.5×10^6	44	730	0.0007	transmutable
¹²⁶ Sn	0.005	0.05	1×10^5	4.4×10^3	4.4×10^3	4.0	questionable
¹²⁹ I	0.14	4.3	1.6×10^7	160	51	0.5	transmutable
¹³⁵ Cs	0.07	1.3	2.3×10^6	310	170	0.08	transmutable
¹³⁷ Cs	0.01	0.02	30	2.2×10^3	1.1×10^4	-	non-transmutable
¹⁵¹ Sm	0.7	700	89	31	0.3	-	non-transmutable or questionable

AXIOMS

A1. For a given neutron spectrum:

**THE MAXIMUM SPECIFIC (per power unit) RATE OF
LLFP-TRANSMUTATION
(the number of transmutations per fission)
IS EQUAL TO THE NEUTRON SURPLUS PRODUCED BY
(FUEL + S).**

A2.

**THE REAL LLFP-TRANSMUTATION RATE IS DEFINED
BY THE DISTRIBUTION OF THE NEUTRON SURPLUS
BETWEEN COMPETITORS: such as LLFP,
CONSTRUCTION MATERIALS, NEUTRON LEAKAGE, etc**

If required transmutation rates are comparable in the value with real neutron
surpluses of fuels

(industrial transmutation scale!)

there is the single principal way to increase the LLFP
transmutation rate:

**the proper choice of both neutron
spectrum and fuel:**

TWO LLFP TRANSMUTATION PROBLEMS

- **TRANSMUTATION RATE MAXIMISATION**
(number of LLFP-transmutations per fission)
via
NEUTRON PRODUCTION POTENTIAL ENHANCEMENT
- **LLFP INVENTORY MINIMISATION**

OVERALL NEUTRON CONSUMPTION OF LLFP NEEDED FOR THEIR TRANSMUTATION

DEFINITION : J-family is a set of nuclei (including the initial nucleus J) which have been produced both by the transmutation in a neutron flux Φ of the J nucleus as well as of its subsequent products, and by decay of nuclei.

The overall neutron consumption D_j of the LLFP family (J-family) can be defined as the total number of neutrons which have to be spent to incinerate the whole family. Removal of stable and short lived fission products can be considered as a single sink of long-term radioactive fission products.

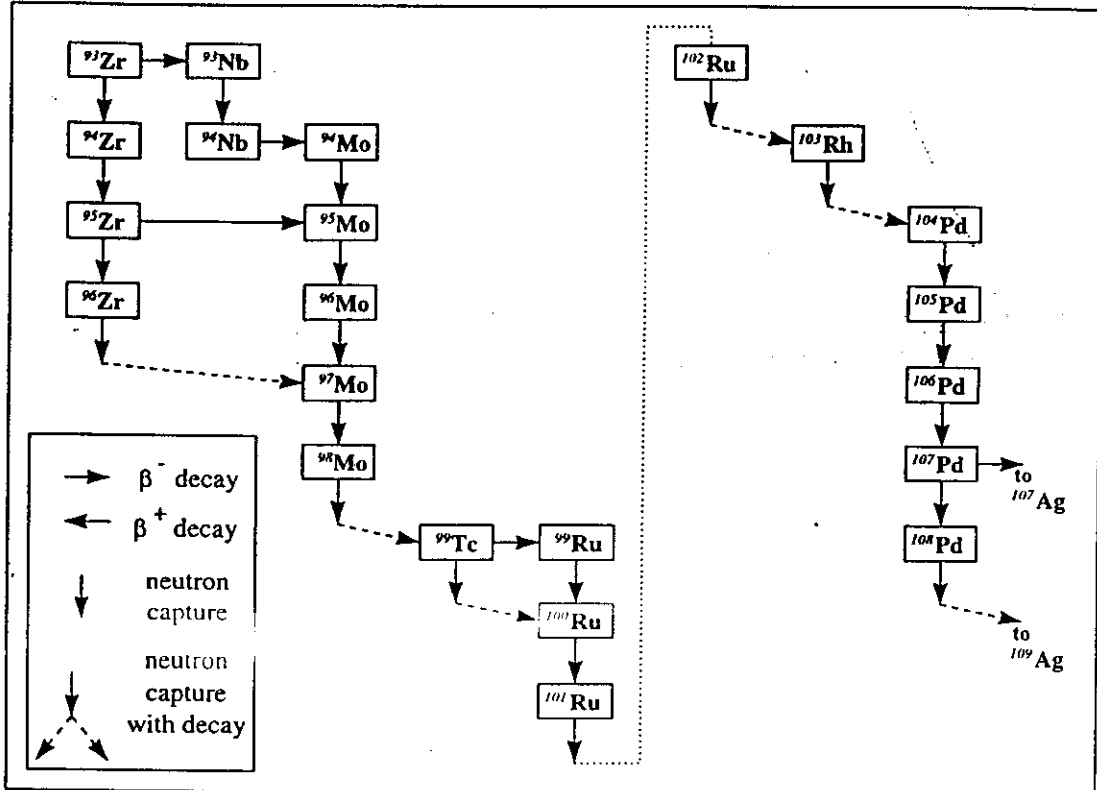


Fig. 1. LLFP transmutation scheme.

TABLE 2

Overall neutron consumption (D_j) of "transmutable" and "questionable" isotopes together with their yields (Y_j) per 1 fission in LWR (UOX) after 5 years of cooling time. Time interval between reprocessing steps - 3 years (removable nuclides : all fission products except all isotopes of *Zr, Tc, Pd, I, Cs, Sn, Nb, Se*)

Transmutable and questionable isotopes (J)	D_j , neutron/transmutation	Y_j , nuclei/fission in NP
^{93}Zr	2.01	0.050
^{99}Tc	1.01	0.055
^{107}Pd	2.04	0.015
^{129}I	1.008	0.009
^{135}Cs	1.002	0.017
^{126}Sn	~2	0.0012
^{94}Nb	0.985	6.3×10^{-7}
^{79}Se	~2	0.0004

TABLE 3

Overall neutron consumption (D_j^*) of "transmutable" nuclides (including all isotopes) together with their yields (Y_j) per 1 fission in LWR (UOX) after 5 years of cooling time. Time interval between fission products reprocessing - 3 years (removable nuclides : all fission products excluding all isotopes of Zr, Tc, Pd, I, Cs)

Transmutable nuclides, J	D_j^* (neutron/transmutation)	Y_j (nuclei/fission in NP)
all Zr	2.03	0.26
all Tc	1.01	0.055
all Pd	3.22	0.095
all I	1.01	0.011
all Cs	0.585	0.13

TABLE 4

Comparison of overall neutron consumption D (neutron/fission in NP)

and

LLFP transmutation effect (reduction of radio-toxicity) for groups of nuclei

$$D = D^* \times Y$$

Transmutable nuclides	D (neutron/fission in NP)	Radio-toxicity reduction factor for $t = 1000$ years (Sv/GW _e ×year)
⁹⁹ Tc	0.055	5500
¹²⁹ I + ⁹⁹ Tc	0.064	8160
¹²⁹ I + ⁹⁹ Tc + ¹³⁵ Cs	0.081	9060
all isotopes of Tc, I, Cs	0.15	9060
all isotopes of Tc, I, Cs + Zr	0.68 <too many!>	9930
all isotopes of Tc, I, Cs +Zr + Pd	0.98 <too many!>	9935

TABLE 5

RADIOTOXICITY of U being eliminated by NP (LWR's)

years	10^2	10^3	10^4	10^5	10^6	10^7	10^8
radio-toxicity	5×10^4	6×10^4	2×10^5	1×10^6	4×10^5	4×10^5	4×10^5

TABLE 6

Radio-toxicity of long-lived LLFP in the repository

after elimination of Tc, I, Cs (Sv/GW_e×year)

years	10^2	10^3	10^4	10^5	10^6	10^7	10^8
radio-toxicity	4.9×10^3	4.9×10^3	4.6×10^3	2.3×10^3	$\sim 0.5 \times 10^3$	~ 0	~ 0

AS THE RESULT:

$$\text{TOX}_{\text{LLFP}} < (\text{TOX}_{\text{U}})/10$$

(THERE IS A MARGIN TO COMPENSATE A MOBILITY OF THE REST LLFP's!)

IT CORRESPONDS TO THE GENERAL "CLEAN ENERGY PRINCIPLE":

**TOTAL WASTE TOXICITY PRODUCED BY
NUCLEAR POWER HAS NOT TO EXCEED THE
TOTAL TOXICITY ELIMINATED BY NUCLEAR
POWER**

Neutron SURPLUS PRODUCTION Potential (FUEL + EXTERNAL SOURCE)

$$G = -\sum_j f_j \times D_j - (CM + L) + S$$

$$G_{TRU} = 1.11 - 0.3 + 0.15 \Big|_{K_{eff} = 0.95} = 0.96 \text{ neutron/fission}$$

$$G_{TRPu} = 0.72 - 0.3 + 0.15 = 0.57 \text{ neutron/fission}$$

REMARK - PARK of LLFP-Burners

If the fast spectrum TRU transmuter ($G = 0.96$ neutron/fission) is used both for the TRU transmutation and Tc, I, Cs incineration, then the fraction ε :

$$\varepsilon \geq \frac{0.15 \text{ neutron / fission}}{G = 0.96 \text{ neutron / fission}} \approx 0.16$$

CONCLUSION

The industrial scale of the LLFP incineration requires an essential number of neutrons.

Hence, fast neutron spectrum and external neutron source (e.g. ADS) are favourable for this application.

RECOMMENDATIONS towards RADIOLOGICAL CLEAN NP

- to put "*transmutable*" (*Tc, I* and *Cs*) fission products in fast reactors and/or ADS,
- to put "*non-transmutable*" nuclides such as *Sr, Sm* to an interim storage for decaying (during some hundreds years) and eventually directly - to a geological repository,
- to put all "*questionable*" nuclides (*Se, Nb, Sn*) together with "*transmutable*" *Zr* and *Pd* to a geological repository (an interim storage can be used for the first hundreds years, similar to "*non-transmutable*" nuclides),
- to put all short lived nuclides into an interim storage and then to a geological repository.

To optimise LLFP transmutation, one needs

- *To use a spectrum with a highest neutron production surplus potential ⇒*

HIGHEST LLFP TRANSMUTATION RATE

(FAST neutron spectrum)

- *To minimise LLFP INVENTORY*

by

**neutron leakage minimisation,*

**minimisation of "parasitic" neutron consumption...),*

via

neutron spectrum thermalisation including LLFP location area,

LLFP resonance capture effects application (TARC),

.....

CONSUMPTION OF NEUTRONS BY LANTHANIDES

(D^* of the LANTHANIDES)

REASON: problems of separation LA and TRU

1. YIELDS

Example: fission's in fast spectrum

TOTAL YIELD: ~ 25 % of FP

Isotopes compositions:

La	0.146
Ce	0.309
Pr	0.166
Nd	0.231
Pm	0.0440
Sm	0.0740
Eu	0.0230
Gd	0.00686
Tb	0.000283
Dy	0.000174
Ho	0.0000066
Er	0.0000032
Tm	0.00000015
Yb	0.000000029
Lu	0.0000000007

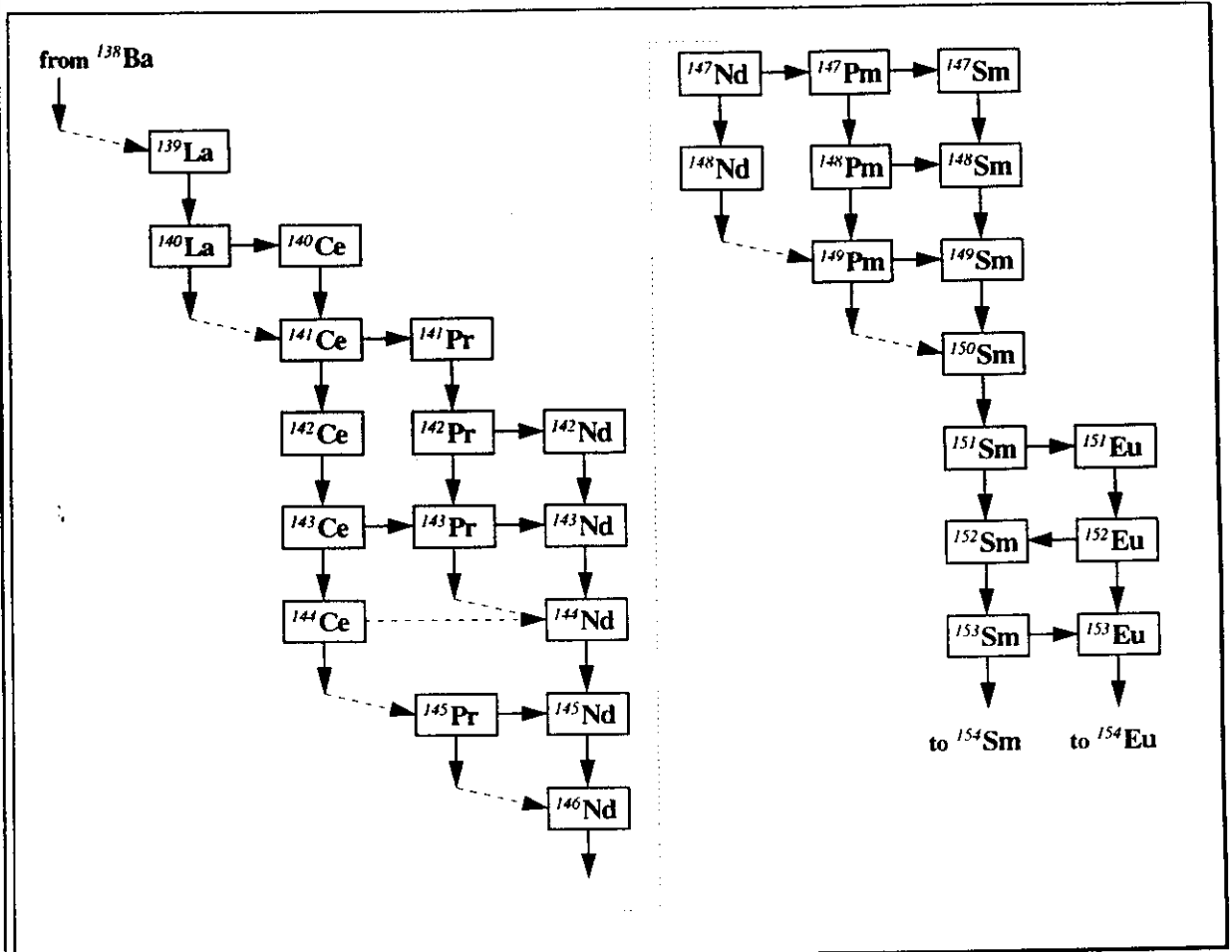
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Table. D* (neutrons per incineration) for Lantanides

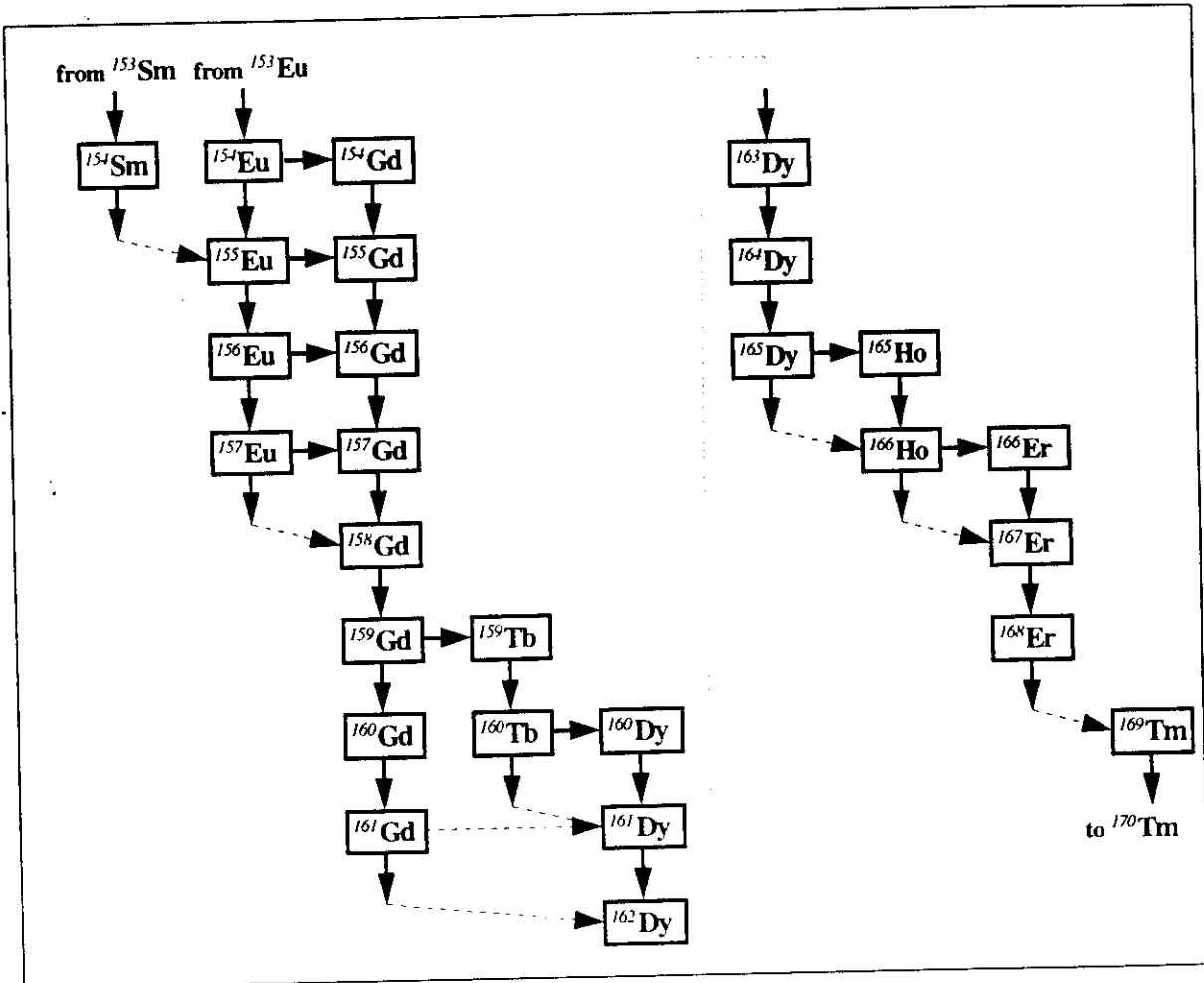
Spectrum/Flux	10^{13}	10^{14}	10^{15}	10^{16}
FAST	33.5			
THERMAL	33.9			

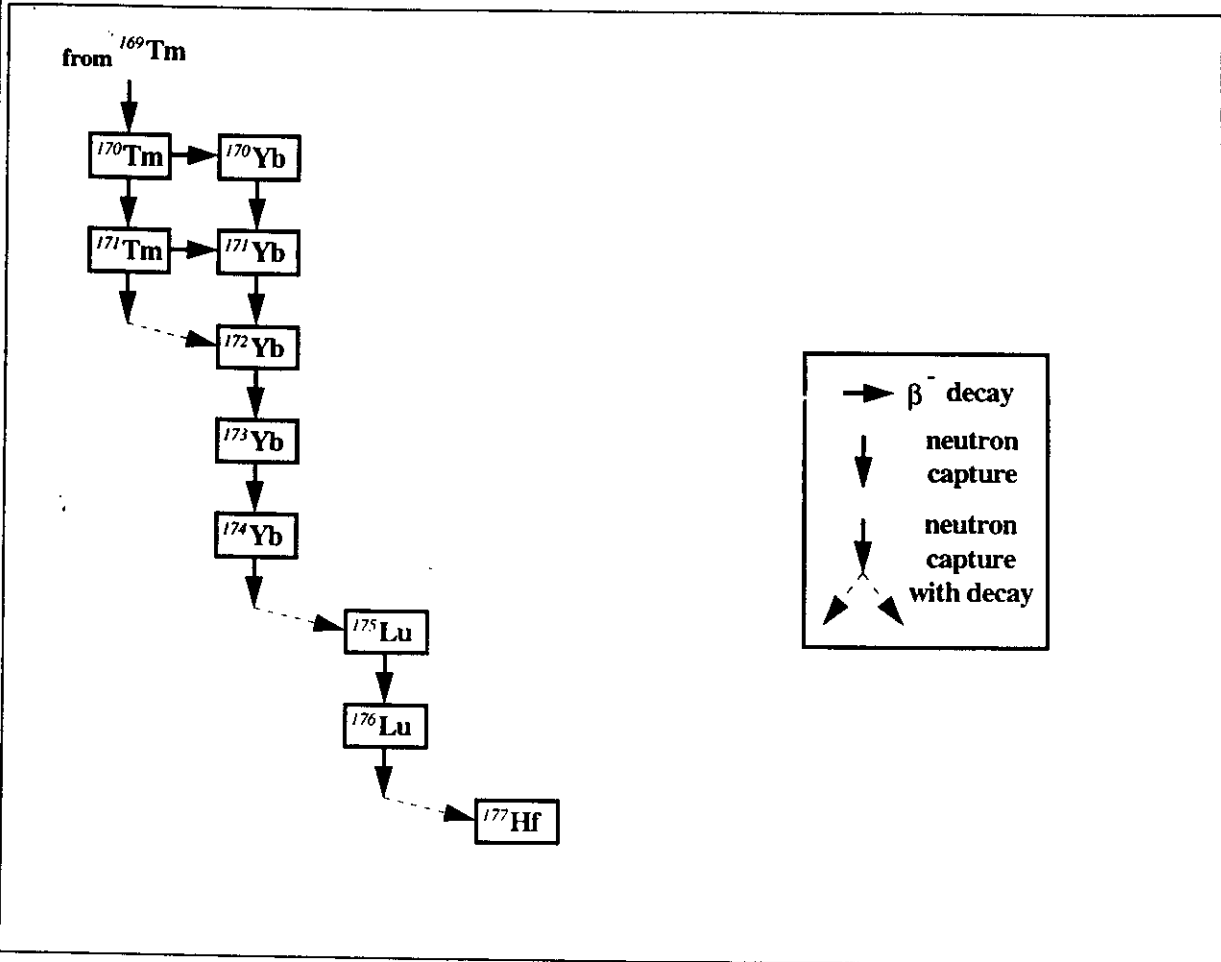
Table. D* (neutrons per incineration) for Lantanides family (Fast spectrum, Flux: 10^{15} n cm⁻²s⁻¹)

Isotope	D*
La	0.52
Ce	0.47
Pr	0.42
Nd	2.36
Pm	0.23
Sm	3.71
Eu	1.87
Gd	2.98
Tb	1.21
Dy	3.26
Ho	0.71
Er	2.55



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CONCLUSION:

- * The potential of LLFP (LA) incineration depends on the FUEL overall neutron production potential.

Table 1. The overall neutron production (-D) for TRU discharge of LWR being used in LLFP BURNERS as the fuel.

Spectrum type	$\Phi = 10^{14}$ n cm ⁻² s ⁻¹	$\Phi = 10^{15}$ n cm ⁻² s ⁻¹	$\Phi = 10^{16}$ n cm ⁻² s ⁻¹	$\Phi = 10^{17}$ n cm ⁻² s ⁻¹
	-D neutron/fission			
Fast (SPX)	0.72	1.15	1.31	1.34
Thermal (LWR-MOX)	0.03	0.30	0.47	0.50
Thermalised (CANDU)	0.21	0.43	0.53	0.60
Resonant (C.Rubbia)	0.40	0.46	0.55	0.69
Well-thermalised (Ch.Bowman)	0.25	0.43	0.52	0.60

