



Summary of RED-IMPACT results on the Impact of P&T on the High Level Waste Management

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on behalf of the RED-IMPACT collaboration*

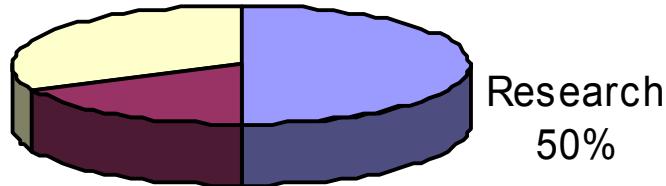
**10th International Exchange Meeting on Partitioning and
Transmutation, IEMPT10
Mito (Japan) 6-10 October 2008**

RED-IMPACT

Impact of Partitioning, Transmutation and Waste Reduction Technologies on the Final Nuclear Waste Disposal

23 partners + 2 subcontractors

Nuclear
Industry &
Utilities



KTH-Sweden: Coordinator
FZJ-Germany: Co-Coordinator
Belgium: BN; SCK-CEN
Czech Republic: NRI; RAWRA
EC: ITU-Karlsruhe
France: Areva ANP, CEA; COGEMA
Germany: FANP; GRS; IER; KKP
Netherlands: NRG
Romania: CITON
Slovakia: DECOM, VUJE
Spain: CIEMAT; EA; ENRESA
UK: NexiaSolutions; NIREX; UC

EC CONTRACT NO. FI6W-CT-2004-002408
Duration: March 2004 – September 2007



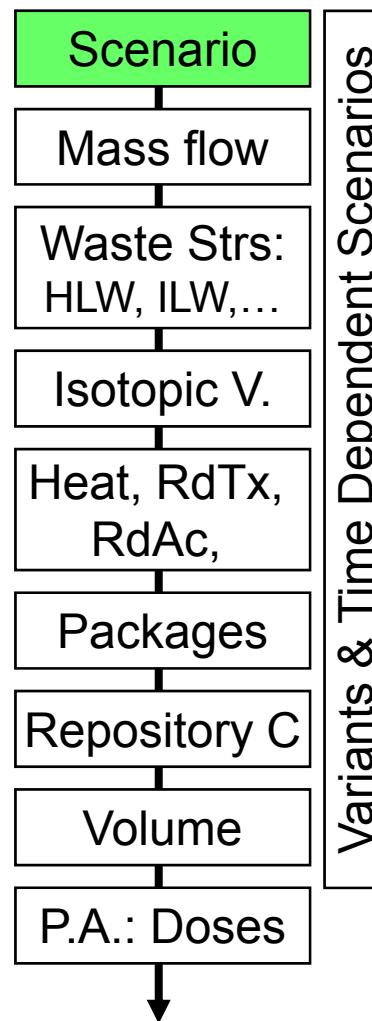
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After a comprehensive inventory of existing and foreseen nuclear fuel cycle facilities in Europe, including a review of worldwide ongoing R&D programs on P&T, and assuming the potential evolution of P&T technologies and advanced fuel cycles (including Gen III & Gen IV reactors and ADS), the project has estimated the effect of P&T on waste management when using existing deep geological repository designs.

- **WP1:** Review of waste management and transmutation strategies, selection of fuel cycles scenarios
- **WP2:** Feasibility of the industrial deployment of selected scenarios and their impact on waste management
- **WP3:** Assessment of waste streams, waste features, leach resistance, heat generation, reprocessing capability etc. for selected fuel cycles.
- **WP4:** Assessment of the benefits of P&T/C in advanced fuel cycles for waste management and geological disposal.
- **WP5:** Economic, environmental and societal assessment of fuel cycle strategies
- **WP6:** Synthesis and dissemination of results

Scenarios for Fuel Cycles



- **Industrial scenarios @ equilibrium**

- **Scenario A1: (reference)** : once through open cycle with Gen-II / III reactors
- **Scenario A2** : mono-recycling of plutonium in Gen-III reactors (+ variants)
- **Scenario A3** : Multirecycling of Pu (only) in Sodium Fast Reactors (EFR)

- **Innovative scenarios @ equilibrium**

- **Scenario B1** : Multi-recycling of Pu & MA in Sodium Fast Reactors (EFR) **plus advanced PUREX**
- **Scenario B2** : mono-recycling of plutonium in Gen-III reactors and burning of Minor Actinides in ADS **plus advanced PUREX & PYRO**
- **Scenario B3** : mono-recycling of plutonium in Gen-III reactors + burning of plutonium in Gen-IV fast reactors + burning of Minor Actinides in ADS including advanced PUREX and PYRO (*reduced efforts*)

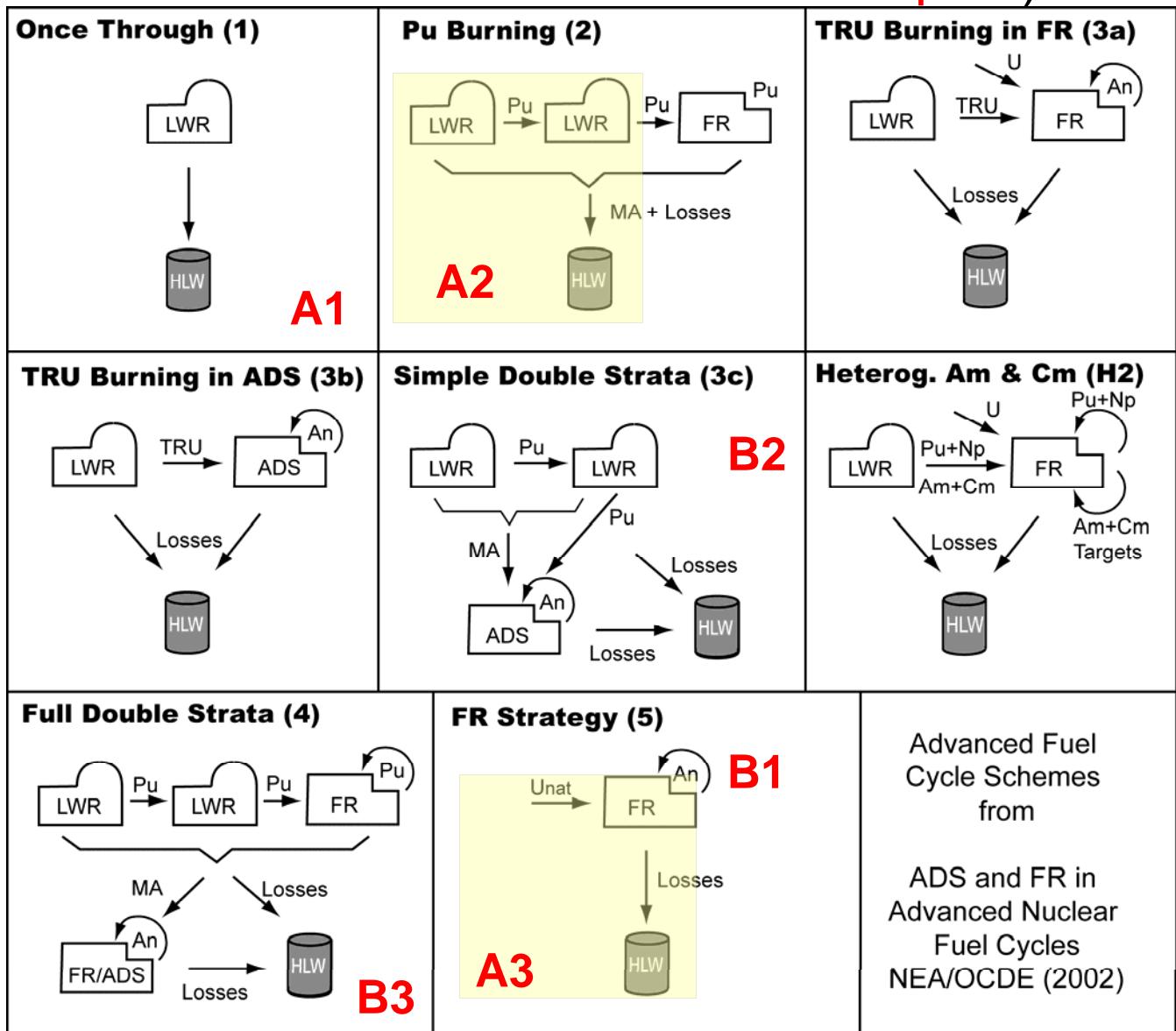
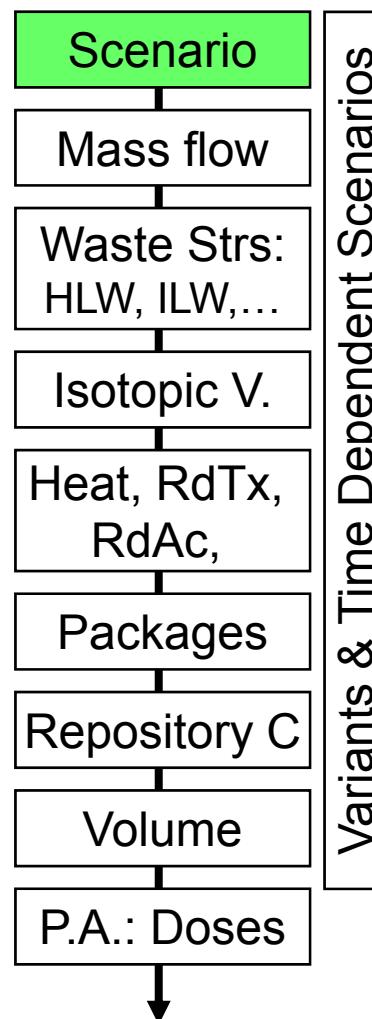
- **Transition Scenarios**



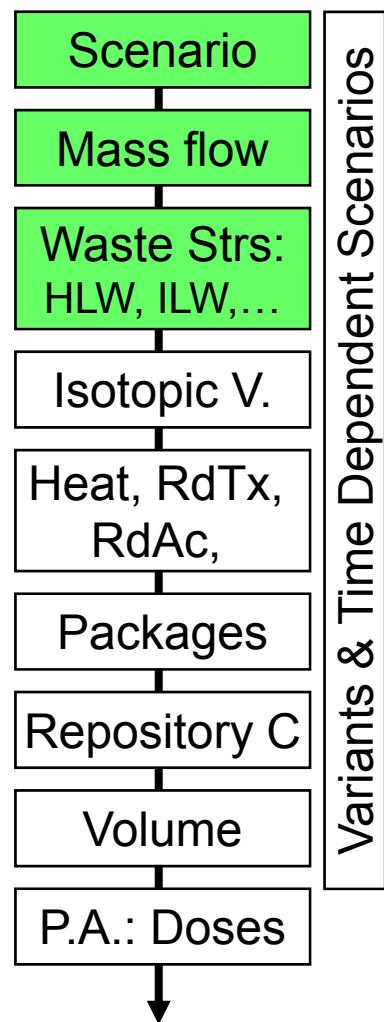
Scenarios for Fuel Cycles

(NEA /

Red-Impact)



Scenarios for Fuel Cycles



Pu reprocessing in LWR (single)

The reference cooling time is 50 years
 UO₂ Initial enrichment in ²³⁵U = 4.20% /
 MOX 8.5% Pu

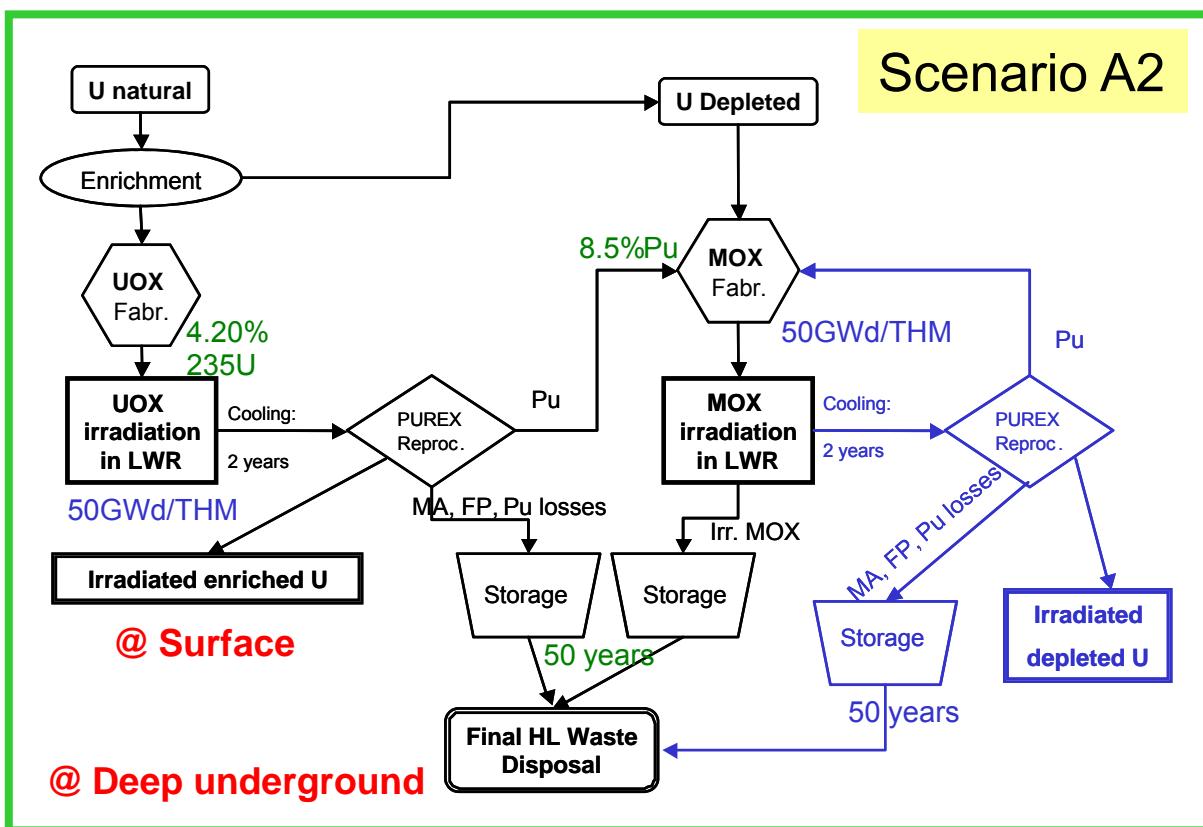
Variants:

- Multiple recycling of Pu in LWR
- Different reactors and fuels for Pu recycling
- Different fuel burnup for Pu recycling

Only presently deployed actinide (Pu) recycle

HLW Waste Streams:

- Irradiated U (enriched)
- MOX Spent fuel
- Purex HLW:
FF, 100% MA
U & Pu losses



Scenarios for Fuel Cycles



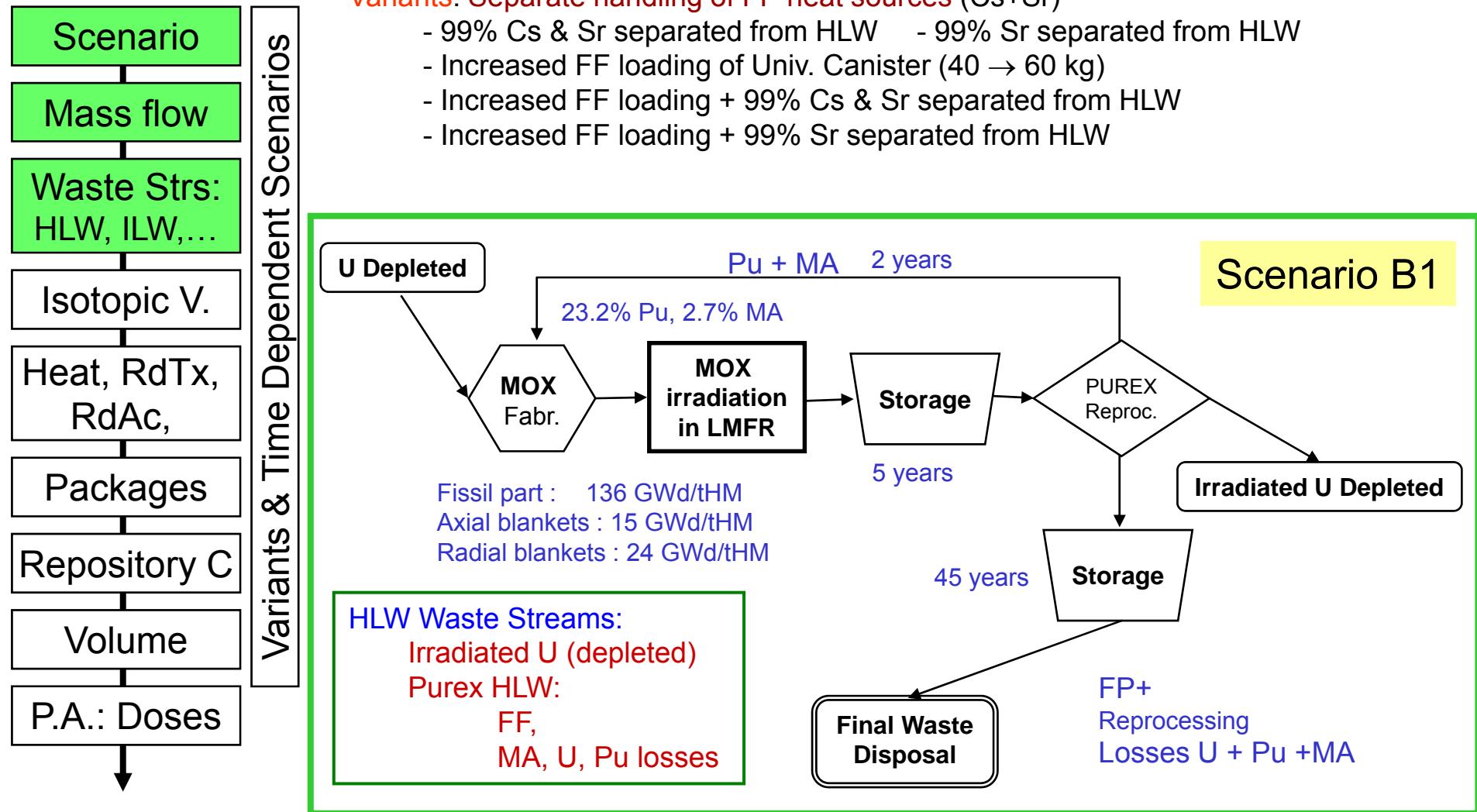
Generation IV fast reactor

EFR CD9/91, Core, Axial and Radial Blankets

23.2% Pu and 2.7% MA in "MOX"

Variants: Separate handling of FP heat sources (Cs+Sr)

- 99% Cs & Sr separated from HLW - 99% Sr separated from HLW
- Increased FF loading of Univ. Canister (40 → 60 kg)
- Increased FF loading + 99% Cs & Sr separated from HLW
- Increased FF loading + 99% Sr separated from HLW



Scenarios for Fuel Cycles

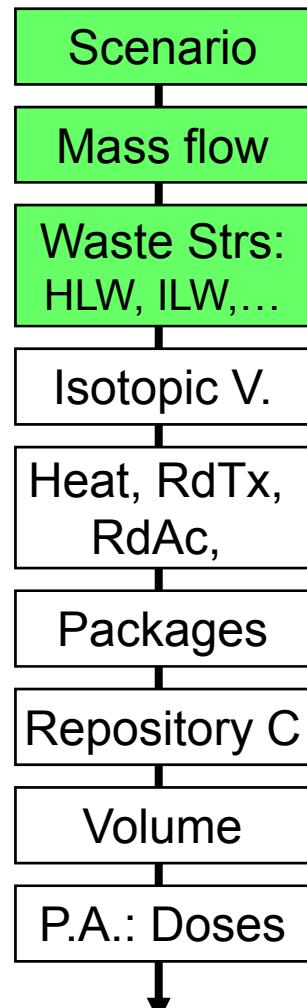


Simplified double strata: with LWR UOX, Mono-recycling of PU in LWR and multi-recycling of Pu+MA in fast spectrum ADS (**no Fast reactor**)

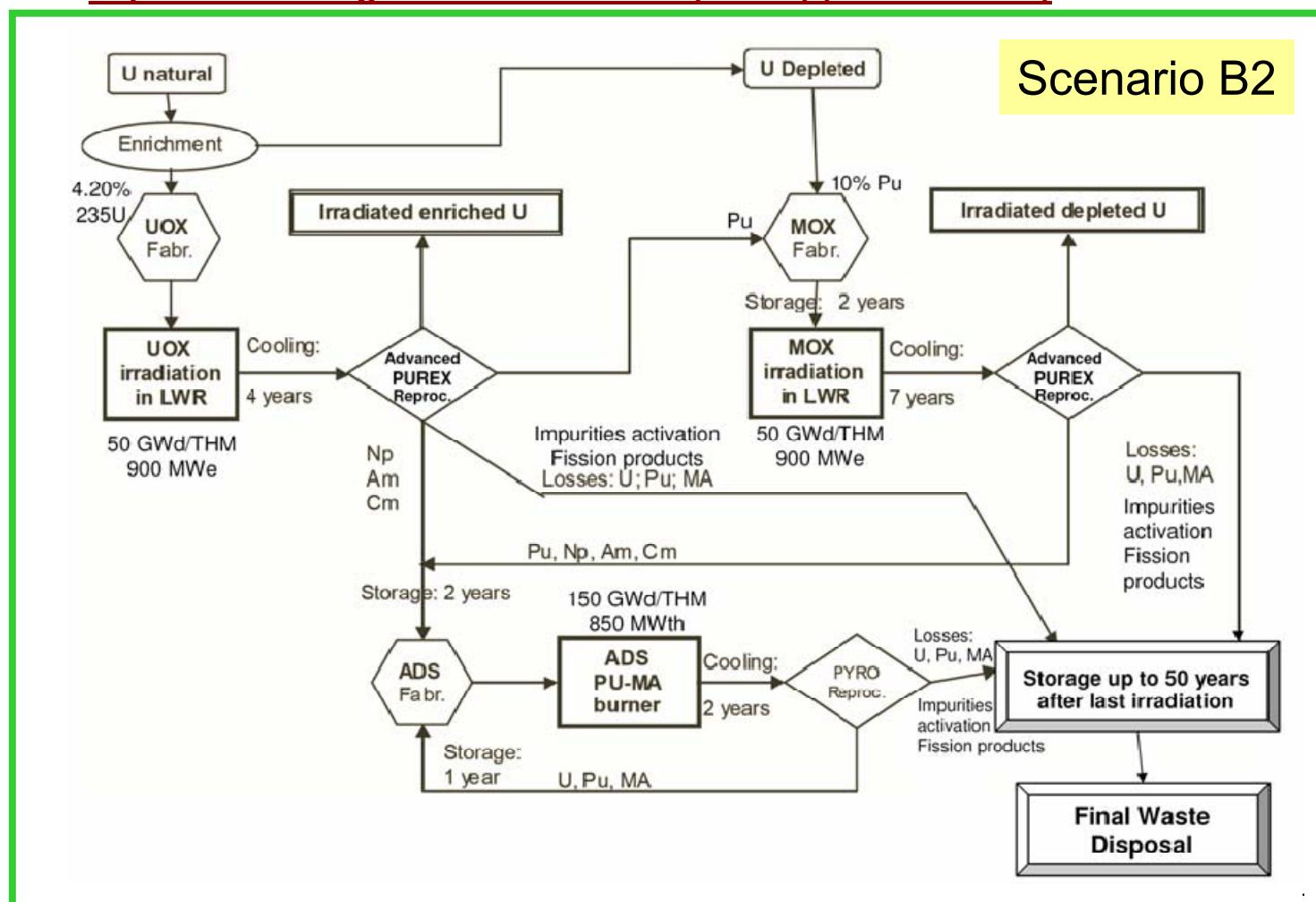
Variants: ADS with TRU Oxide fuel in ZrO_2 inert matrix

(+ 8.5% Pu content in MOX and 5 years LWR UOX spent fuel cooling time)

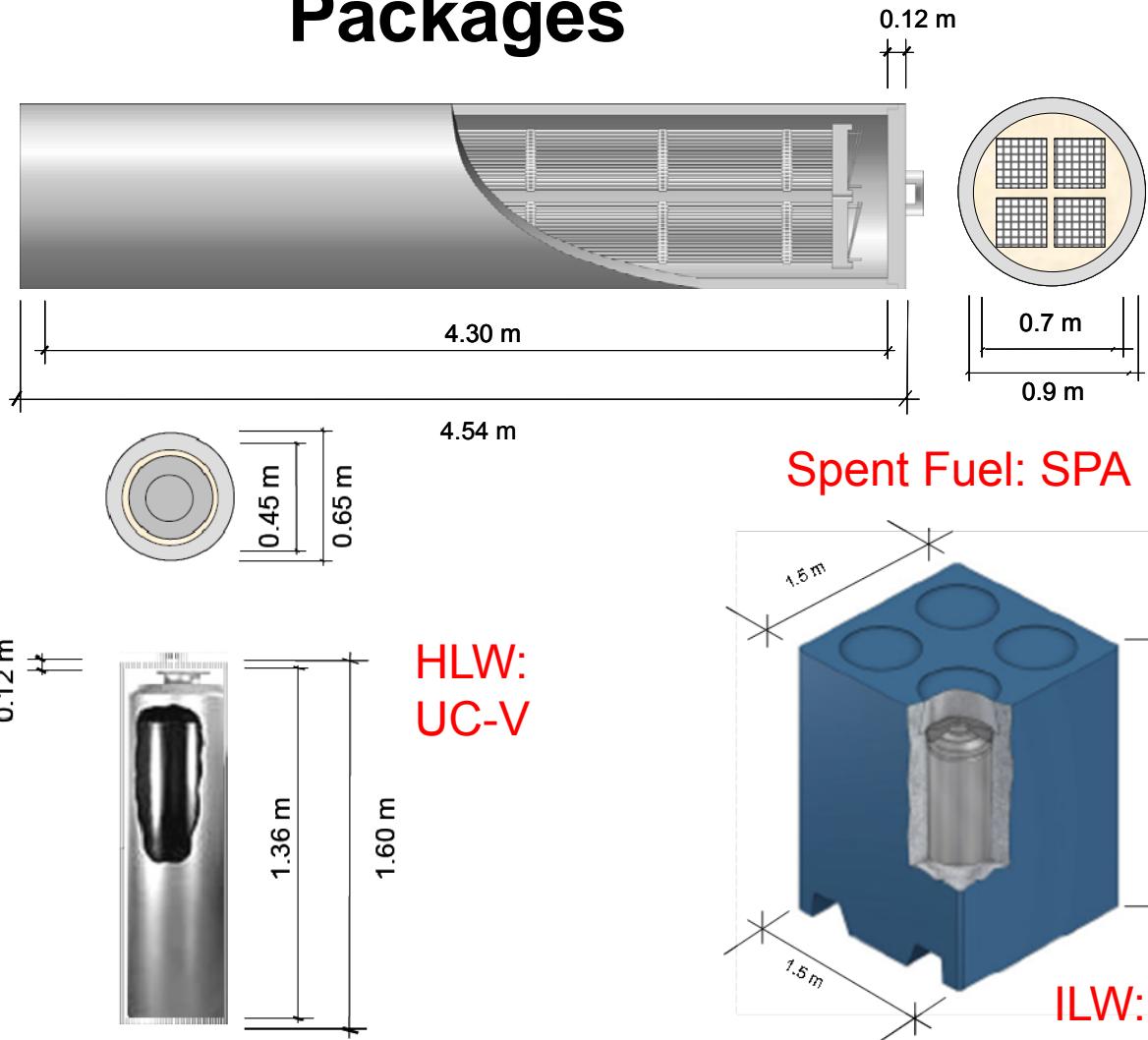
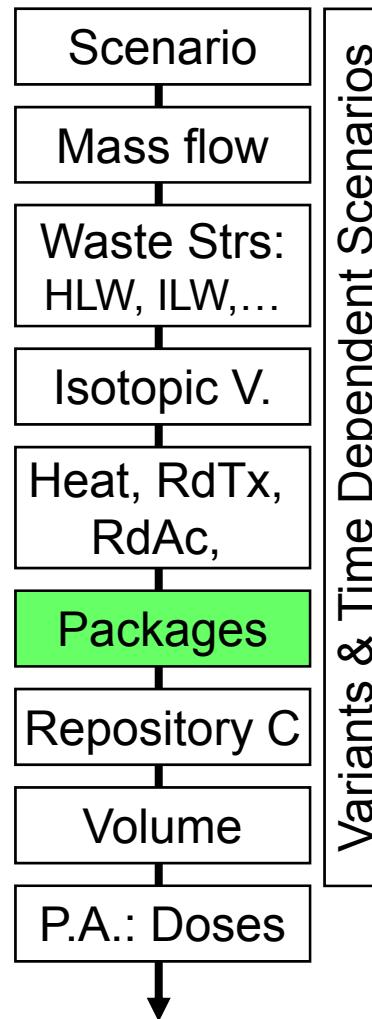
Separate handling of FP heat sources (Cs+Sr) (studied in B1)



Variants & Time Dependent Scenarios

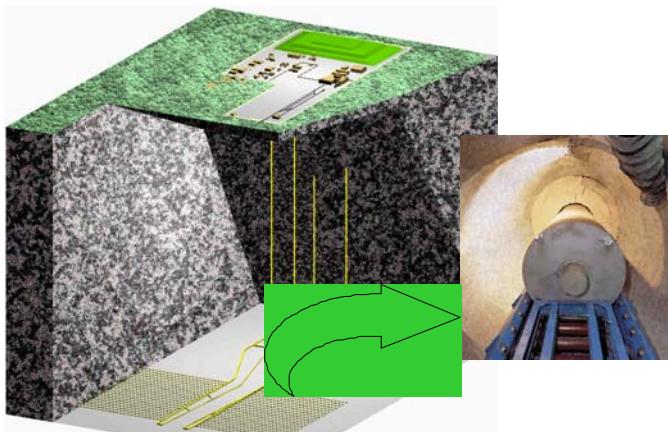
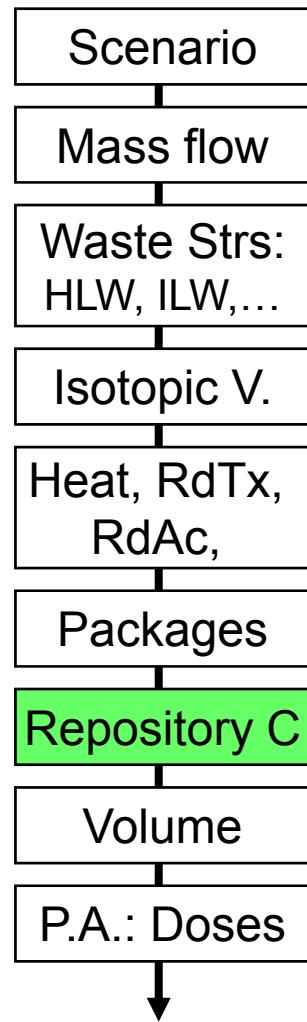


Packages

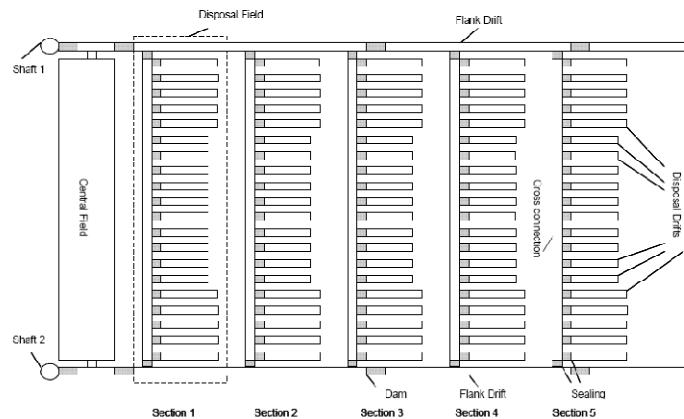


Number of packages, volume and gallery length, heat and radiation levels (n, γ)

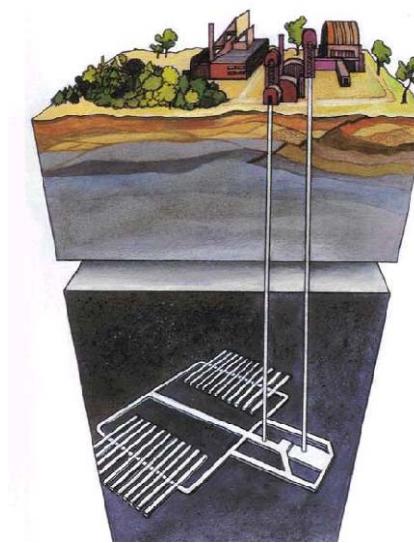
Deep Geological Repositories



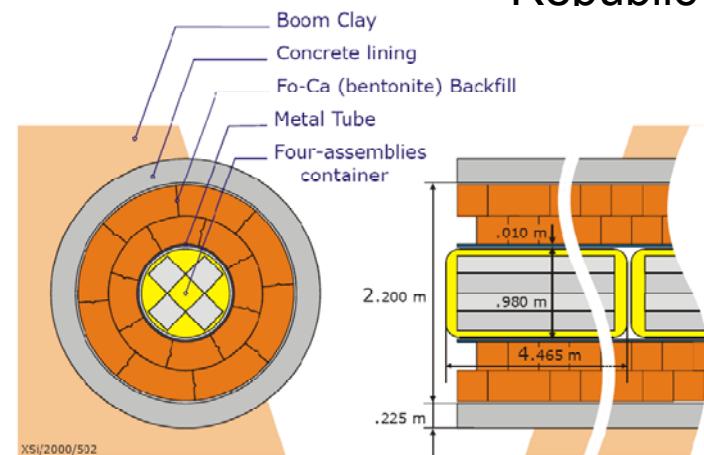
Granite Spain



Salt Germany



Granite
Czech
Republic



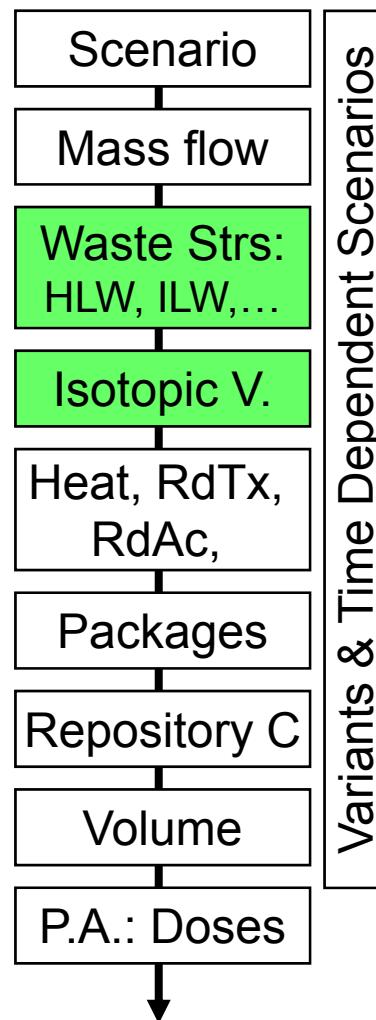
Boom Clay Belgium



HLW: Waste Streams and Isotopic v.



Red-Impact- HLW hypothesis



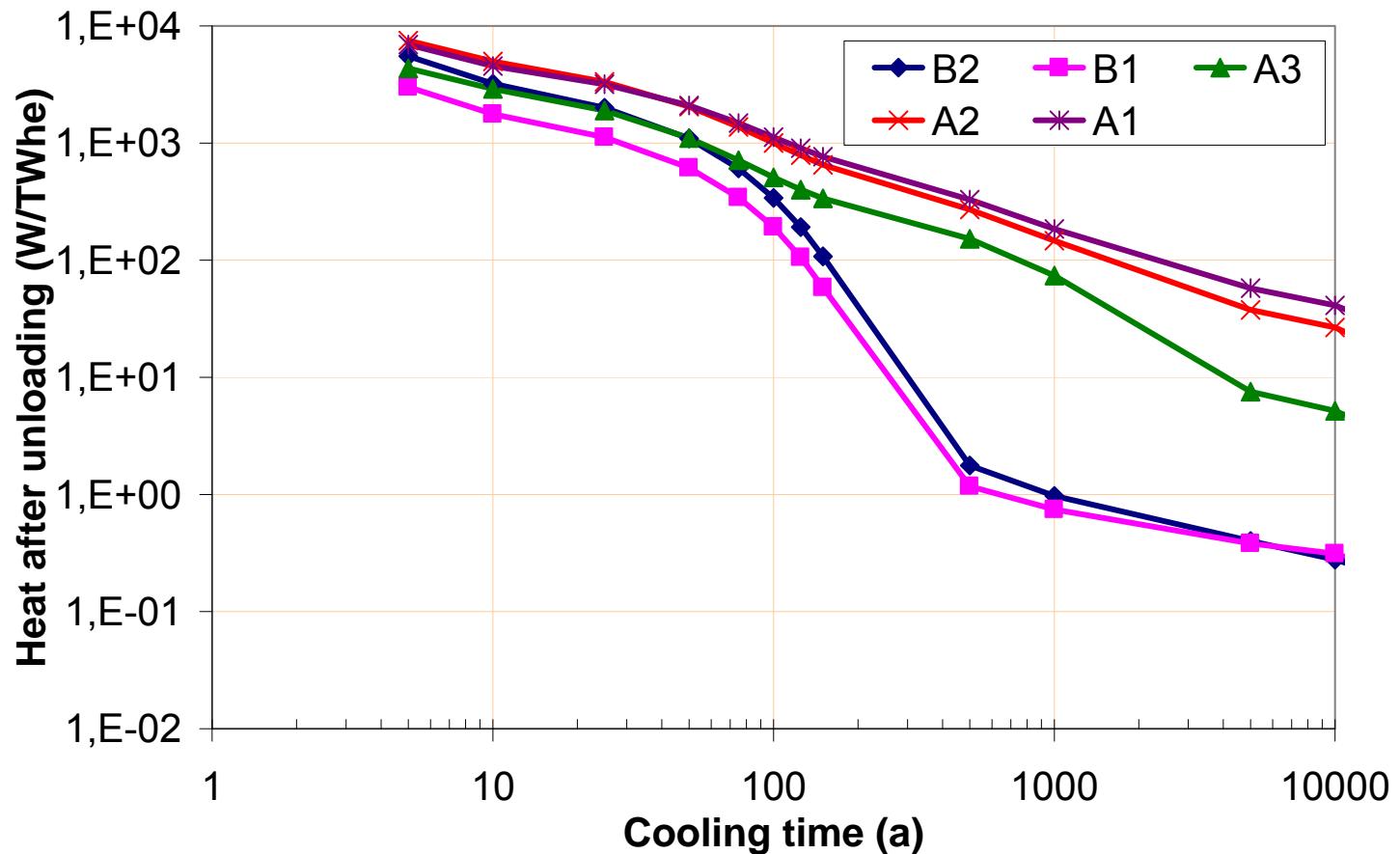
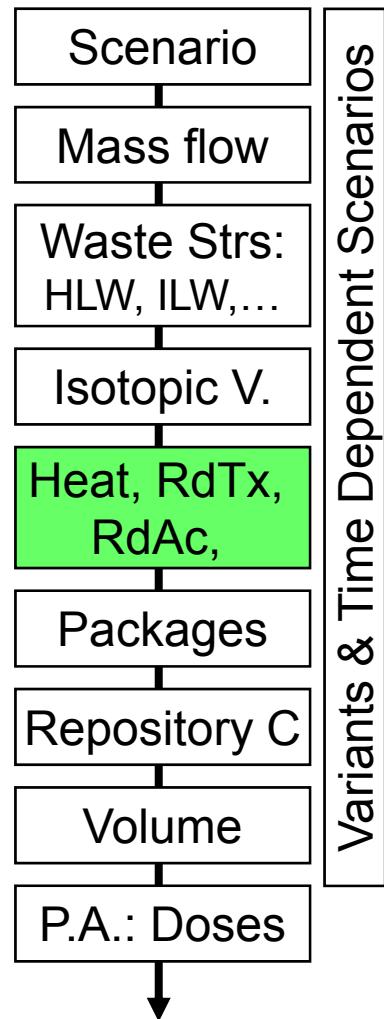
Fractions of main chemical elements incorporated into the vitrified HLW (%)

SCENARIO		A2	A3	B1	B2		
REPROCESSING TYPE		Standard PUREX	Standard PUREX	Extended PUREX	Extended PUREX	Extended PUREX	PYRO
REPROCESSED FUEL		UOX (PWR)	Blankets + Core (FR)	Blankets + Core (FR)	UOX (PWR)	MOX (PWR)	ZrN +TRUN (ADS)
ACTINIDES (An)	U and Pu	0.1	0.1	0.1	0.1	0.1	0.1
	MA	100	100	0.1	0.1	0.1	0.1
FISSION PRODUCTS (FP)	Noble Gases	0	0	0	0	0	0
	Iodine	1	1	1	1	1	1
	Noble Metals	100	100	100	100	100	0
	Others	100	100	100	100	100	100
LIGHT ELEMENTS (Fuel Impurities)	H	0	0	0	0	0	0
	Carbon	10	10	10	10	10	10
	Chlorine	1	1	1	1	1	1
	Others	100	100	100	100	100	100
Fuel Assembly Structural (particles)		0	0	0	0	0	0
Zr of the ADS fuel matrix (also for the Zr-fission product)		-	-	-	-	-	1

Derived magnitudes: Heat Load



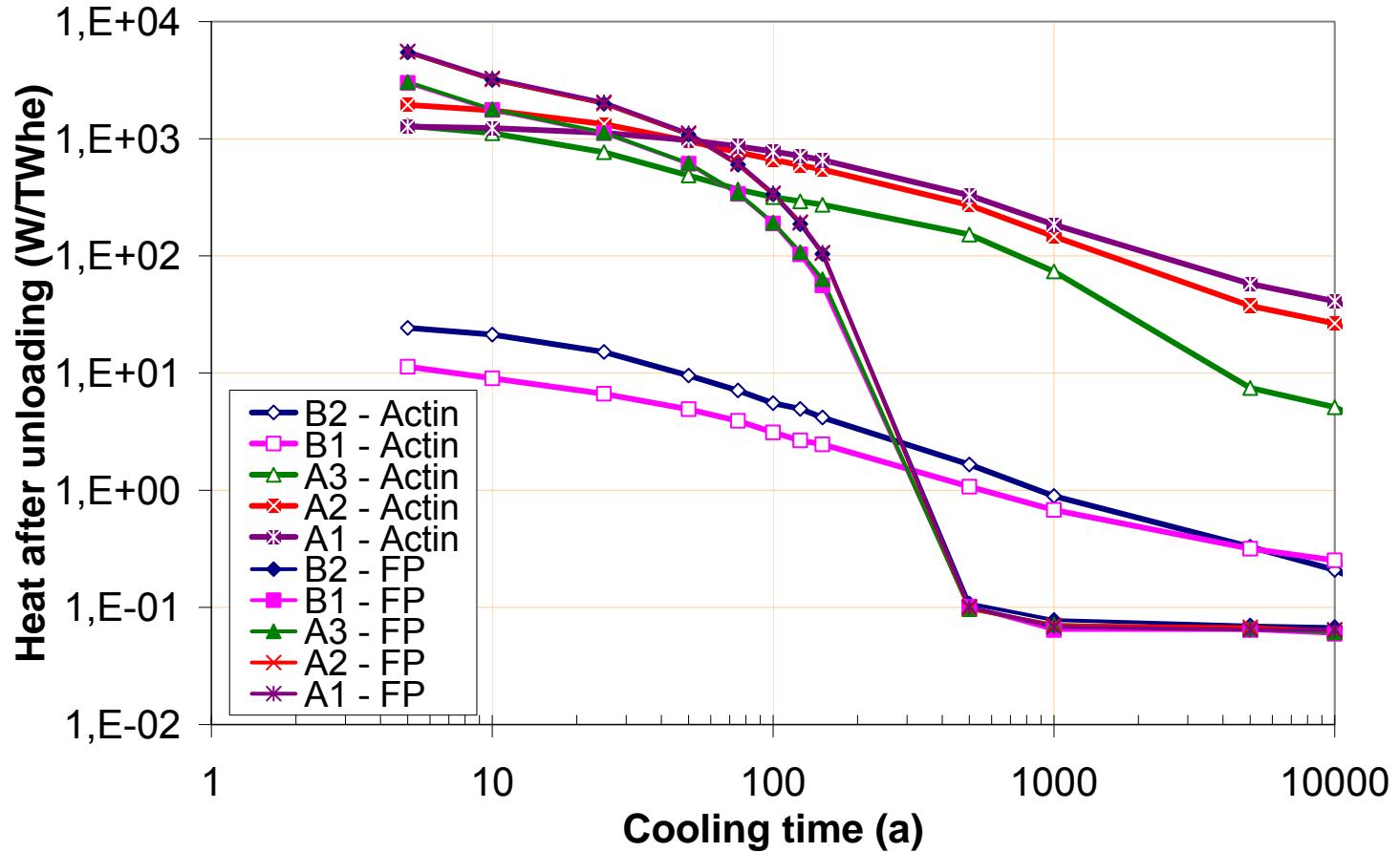
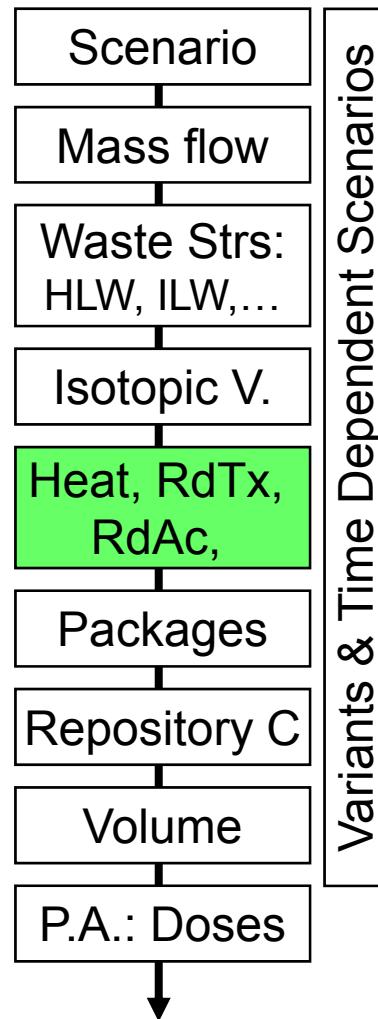
Red-Impact HLWs (total) thermal power evolution with time



Derived magnitudes: Heat Load



Red-Impact HLWs (total) thermal power evolution with time
(Actinides and Fission Fragments contributions)

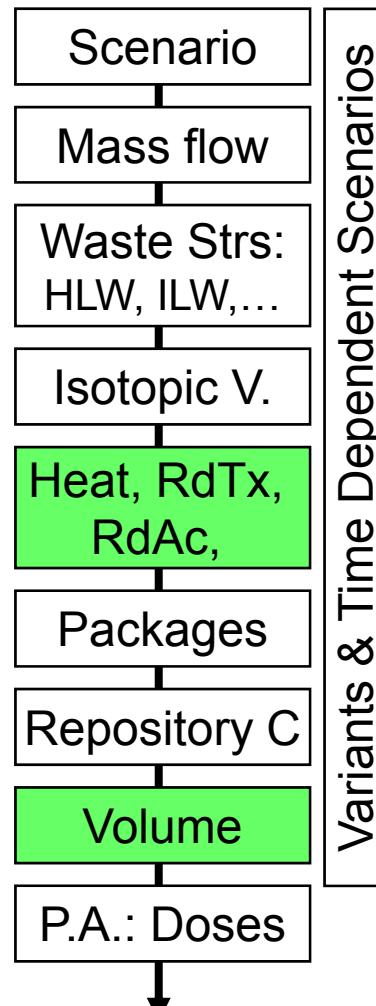




Derived magnitudes: Heat Load



**Red-Impact HLWs thermal power
(total per waste stream)**



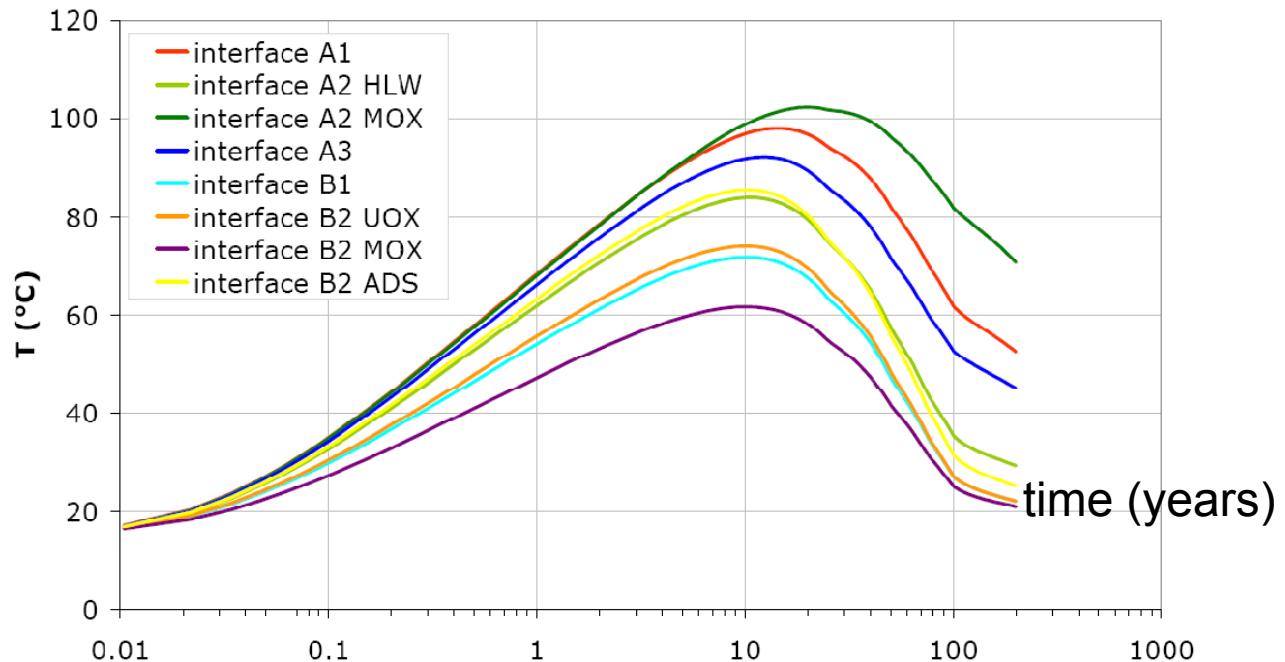
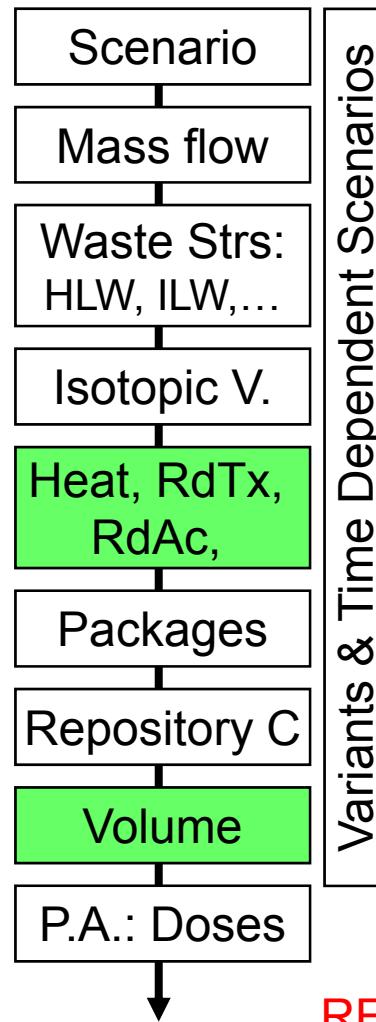
HLW total thermal power per TWheh in the scenarios

Scenario	A1	A2			A3	B1	B2			Total B2
Fuel Type	LWR-UOx	LWR-MOx	LWR-UOx (reproc.)	Total A2	FR-MOx (reproc.)	FR-MOx (reproc.)	LWR-UOx (reproc.)	LWR-MOx (reproc.)	ADS-TRU-Nitride	Total B2
Thermal Power (W) after 50 years (emplacement time)	2096	836	1205	2042	1270	713	869	86	145	1100
Thermal Power (W) after 150 years	766	457	275	732	384	73	83	9	17	108
Thermal Power (W) after 1000 years	185	117	27	144	82	2	0	0	1	1
Thermal Power (W) after 10000 years	43	24	2	26	7	2	0	0	0	1

HLW Heat Load -> Capacity



The main thermal limitation for the repository concept is that the maximum temperature at the gallery lining / Boom Clay interface has to remain below 100 °C. For granite repositories the temperature in the bentonite buffer has to remain lower than 100 °C



Red-Impact temperature at the interface between the gallery lining and the Boom Clay (Belgian repository concept for disposal in clay)

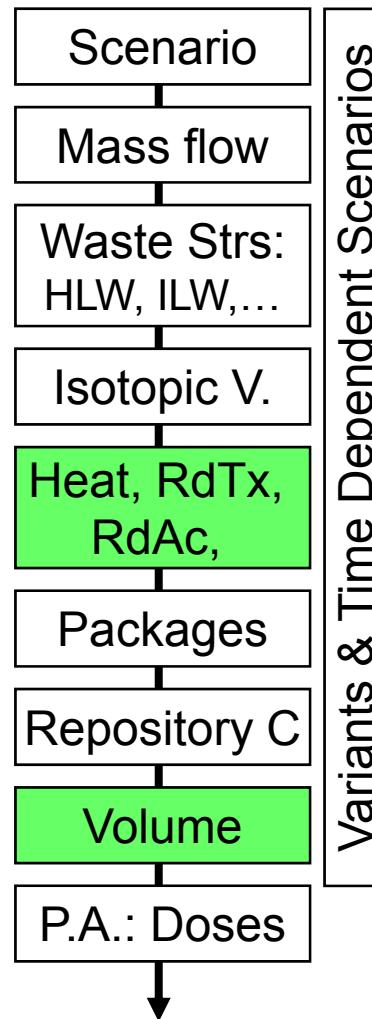
Length of the needed HLW disposal galleries B1/A1 reduced by 3

RED-Impact cases for different Granite/Clay repositories $1.6 < B1/A1 < 6$

HLW Heat Load -> Capacity



Cs & Sr separation from HLW



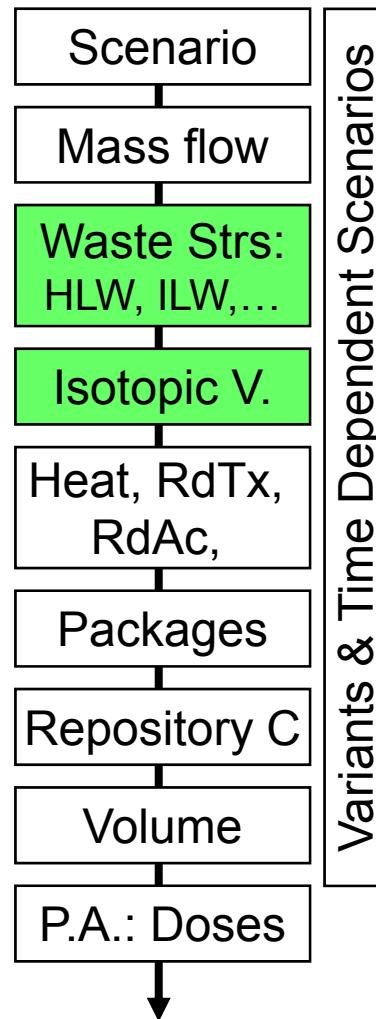
- Two variant scenarios of B1 in which it is assumed that **only Sr or both Cs and Sr are separated** from the HLW prior to vitrification studied.
- **Separating Sr only**, results in a further reduction of the needed gallery length with a factor 1.5, or about a **factor 5** in comparison with the "once through" reference fuel cycle A1.
- Separating Cs and Sr makes that the **thermal output of the vitrified HLW is so low that cooling times are not longer needed**, or that **other repository concepts** can be used.
- However, for the estimation of the needed gallery length one should not forget that the separated Cs **has to be disposed** of in a geological repository also, because it contains the **long-lived ¹³⁵Cs isotope** (half-life 2.3 million years).
- **Separating Cs and Sr**, if it is assumed that the conditioned Cs-waste is disposed after a cooling time of **100 years** (50 more than normal), the **needed gallery length can be reduced** with a factor 4 in comparison with scenario B1 and with a **factor 13** in comparison with scenario A1.
- Longer cooling times would increase the reduction factors.



ILW: Waste Streams and Isotopic v.



Large amounts of ILW can compromise the capacity improvements from the HLW heat reductions



Red-Impact – ILW hypothesis

Fractions of main chemical elements incorporated into the compacted ILW (%)

SCENARIO		A2	A3	B1	B2		
REPROCESSING TYPE		Standard PUREX	Standard PUREX	Extended PUREX	Extended PUREX	Extended PUREX	PYRO
REPROCESSED FUEL		UOX (PWR)	Blankets + Core (FR)	Blankets + Core (FR)	UOX (PWR)	MOX (PWR)	ZrN +TRUN (ADS)
ACTINIDES (An)	U and Pu	0.02	0.02	0.02	0.02	0.02	0.02
	MA	0.02	0.02	0.02	0.02	0.02	0.02
FISSION PRODUCTS (FP)	Noble Gases	0	0	0	0	0	0
	Iodine	1	1	1	1	1	1
	Noble Metals	0.2	0.2	0.2	0.2	0.2	100
	Others	0.2	0.2	0.2	0.2	0.2	0.2
LIGHT ELEMENTS (Fuel Impurities)	H	40	40	40	40	40	40
	Carbon	0	0	0	0	0	0
	Chlorine	0.2	0.2	0.2	0.2	0.2	0.2
	Others	0.2	0.2	0.2	0.2	0.2	0.2
Fuel Assembly Structural		100	100	100	100	100	100
Zr of the ADS fuel matrix (also for the Zr-fission product)		-	-	-	-	-	100



Number of Waste Packages



Red-Impact- HLW waste packages

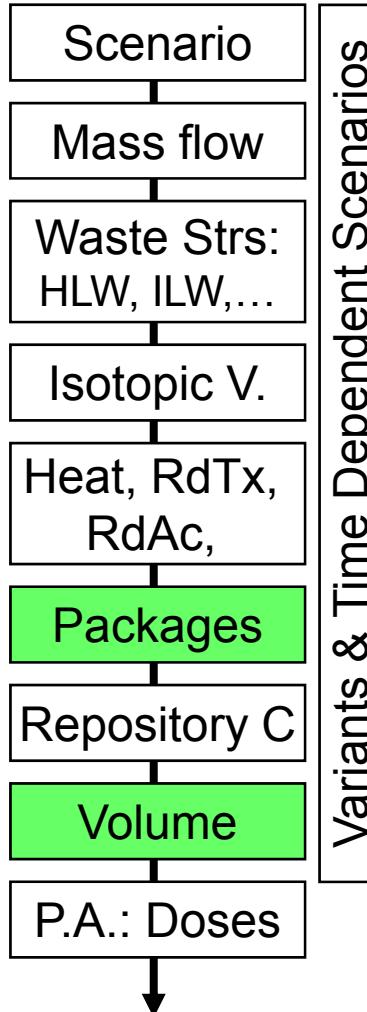
HLW forms and packages production per scenario

Scenario	A1	A2		A3	B1	B2		
Fuel Type	LWR-UOx	LWR-MOx	LWR-UOx (reproc.)	FR-MOx (reproc.)	FR-MOx (reproc.)	LWR-UOx (reproc.)	LWR-MOx (reproc.)	ADS TRU-Nitride (reproc.)
Waste Forms per tHM	2.18	2.21	1.12	2.09	1.98	1.08	1.07	2.61
tHM / TWeh	2.46	0.25	2.21	1.15	1.15	1.91	0.24	0.12
Waste Forms per TWeh/h	5.35 SFA	0.54 SFA	2.48 UC-V	2.40 UC-V	2.27 UC-V	2.07 UC-V	0.26 UC-V	0.33 UC-V
Waste Forms per HLW Package	4	1	1	1	1	1	1	1
Waste Packages per TWeh	1.34	0.54	2.48	2.40	2.27	2.07	0.26	0.33
Package Volume (m³)	2.89	1.51	0.53	0.53	0.53	0.53	0.53	0.53
HLWs Volume (m³/TWeh)	3.87	0.82	1.32	1.27	1.21	1.10	0.14	0.17
HLWs TOTAL Volume (m³/TWeh)	3.87	2.14		1.27	1.21	1.41		

Red-Impact- ILW waste packages

ILW forms and packages production per scenario

Scenario	A1	A2		A3	B1	B2			
Fuel Type	LWR-UOx	LWR-MOx	LWR-UOx (reproc.)	FR-MOx (reproc.)	FR-MOx (reproc.)	LWR-UOx (reproc.)	LWR-MOx (reproc.)	ADS-TRU Nitride (reproc.)	ADS (operation)
Waste Forms per tHM	-	-	1.00	4.10	4.10	1.00	1.01	6.22	0.27
tHM / TWeh	2.46	0.25	2.21	1.15	1.15	1.91	0.24	0.12	0.12
Waste Forms per TWeh/h	-	-	2.21	4.71	4.71	1.91	0.24	0.78	0.03
Waste Forms per ILW Package	-	-	4	4	4	4	4	4	4
Waste Packages per TWeh	-	-	0.55	1.18	1.18	0.48	0.06	0.19	0.01
Package Volume (m³)	-	-	4.5	4.5	4.5	4.5	4.5	4.5	4.5
ILWs Volume (m³/TWeh)	-	-	2.49	5.31	5.31	2.15	0.27	0.87	0.04
ILWs TOTAL Volume (m³/TWeh)	-	2.49		5.31	5.31	3.33			

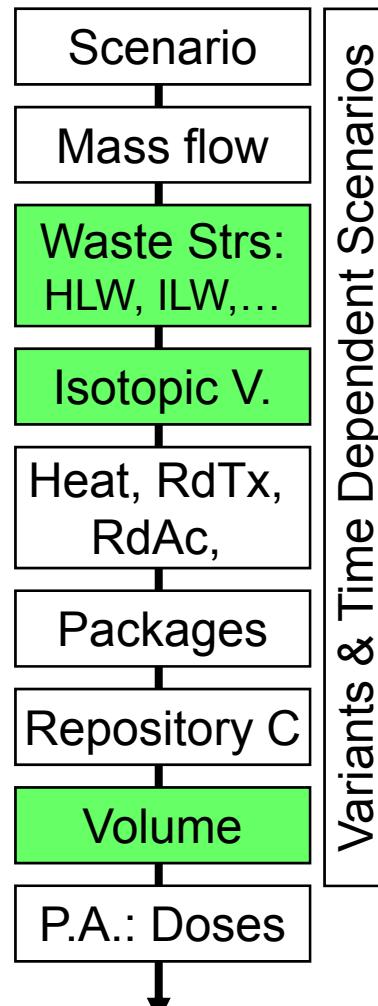




ILW: Waste Streams and Isotopic v.



Red-Impact relevant fission and activation products in the solid wastes (HLWs + ILWs)



Nuclides	Mass of main Fission and Activation Products in Solid Wastes (g/TWeh)									
	A1		A2		A3		B1		B2	
	HLW	ILW	HLW	ILW	HLW	ILW	HLW	ILW	HLW	ILW
Fission Prod.	125810	-	106110	195	89723	189	89936	189	103868	191622(*)
C-14	0.3	-	0.03	0.01	0.2	2.3	0.2	2.3	3.1	0.1
Cl-36	6.0	-	0.3	0.5	0.002	0.0004	0.002	0.0004	0.04	0.6
Se-79	17	-	17	0.03	16	0.03	16	0.03	17	0.1
Sr-90	571	-	542	1.0	249	0.5	250	0.5	513	1.2
Nb-94	18	-	2.5	15	0	79	0	78.6	0.7	21
Tc-99	2931	-	2920	6.5	2298	209	2317	208	2969	10.9
Pd-107	837	-	937	1.5	1207	2.4	1217	2.4	828	296
Sn-126	80	-	85	0.1	108	0.2	109	0.2	93	0.3
I-129	634	-	82	5.7	6.7	6.6	6.7	6.7	6.9	9.3
Cs-135	1633	-	1750	2.9	4270	8.5	4270	8.5	2336	6.5
Cs-137	1405	-	1405	2.5	1137	2.3	1133	2.3	1455	3.8

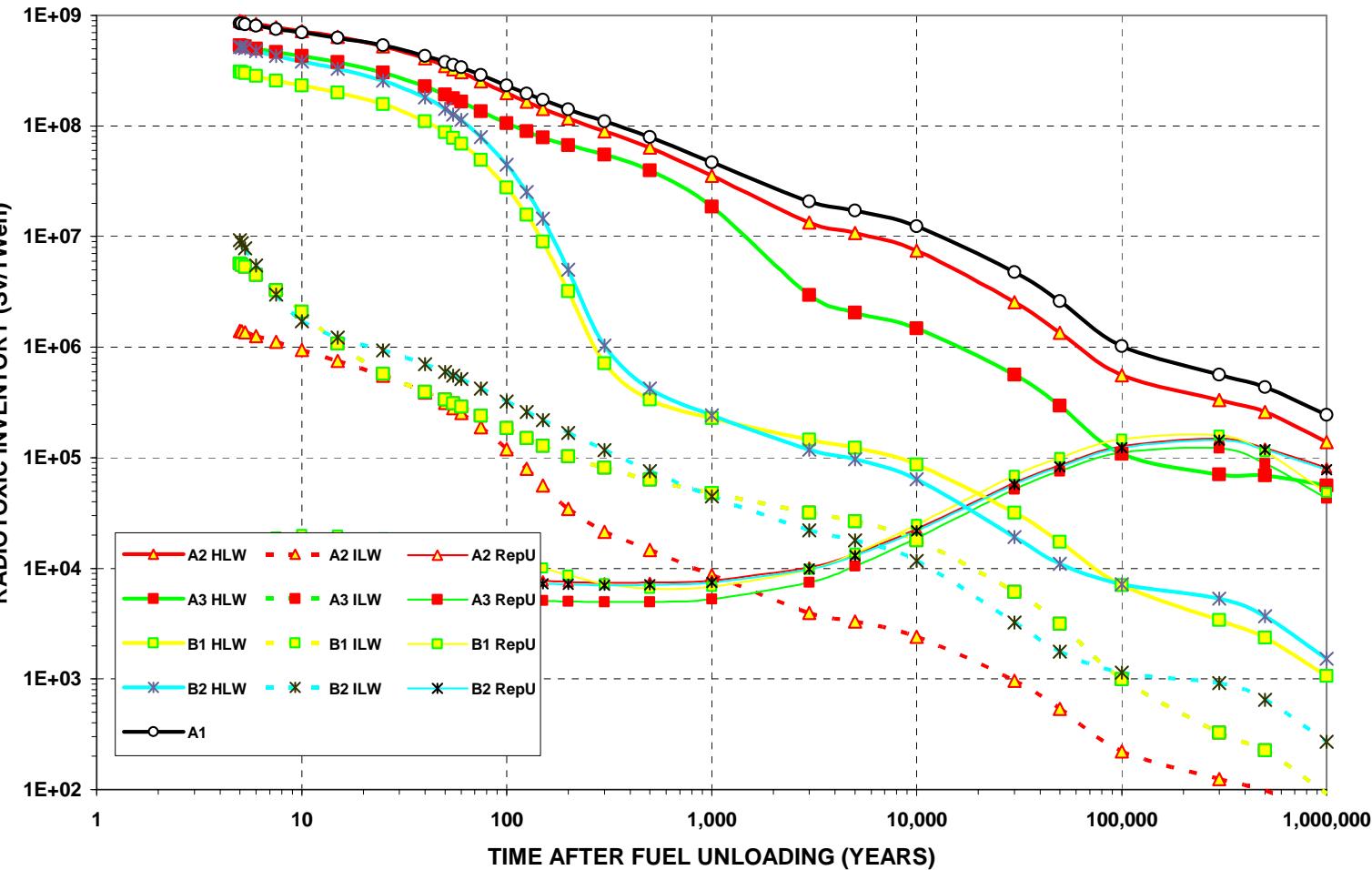
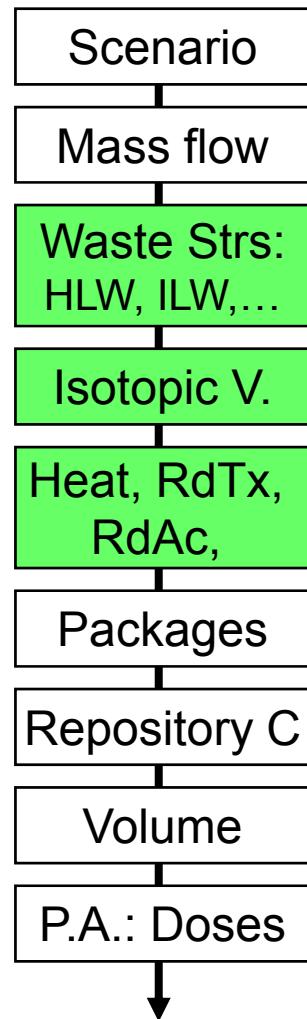
(*) Zr of the ADS fuel matrix included.

AI = Activation of impurities in fuel or structural materials

HLW + ILW: Radiotoxicity

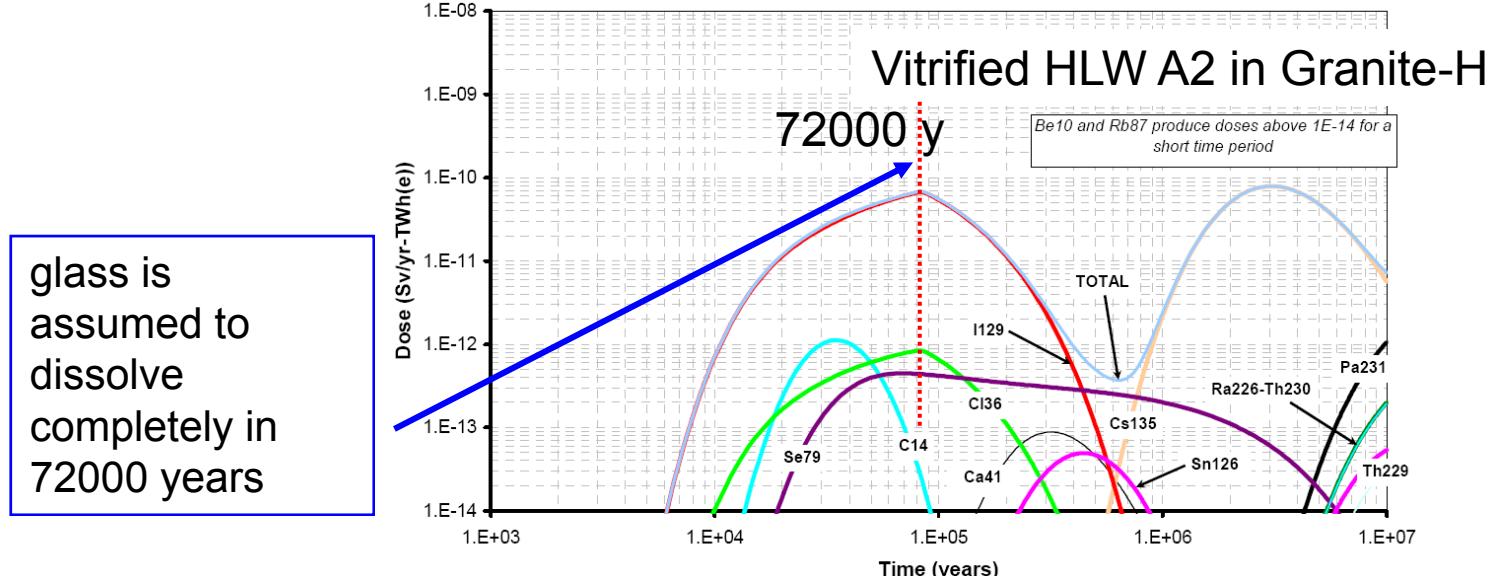
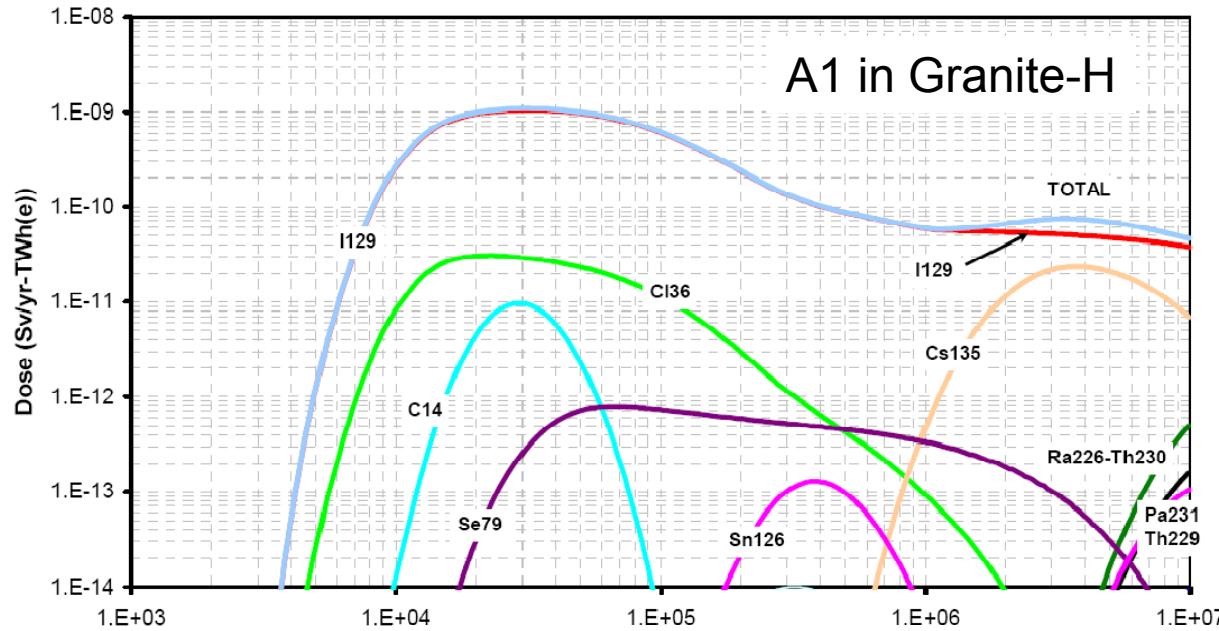
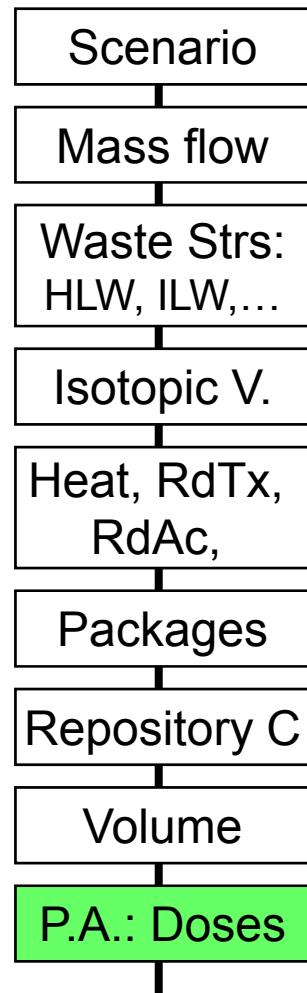


Red-Impact HLWs and ILWs radiotoxicity evolution with time



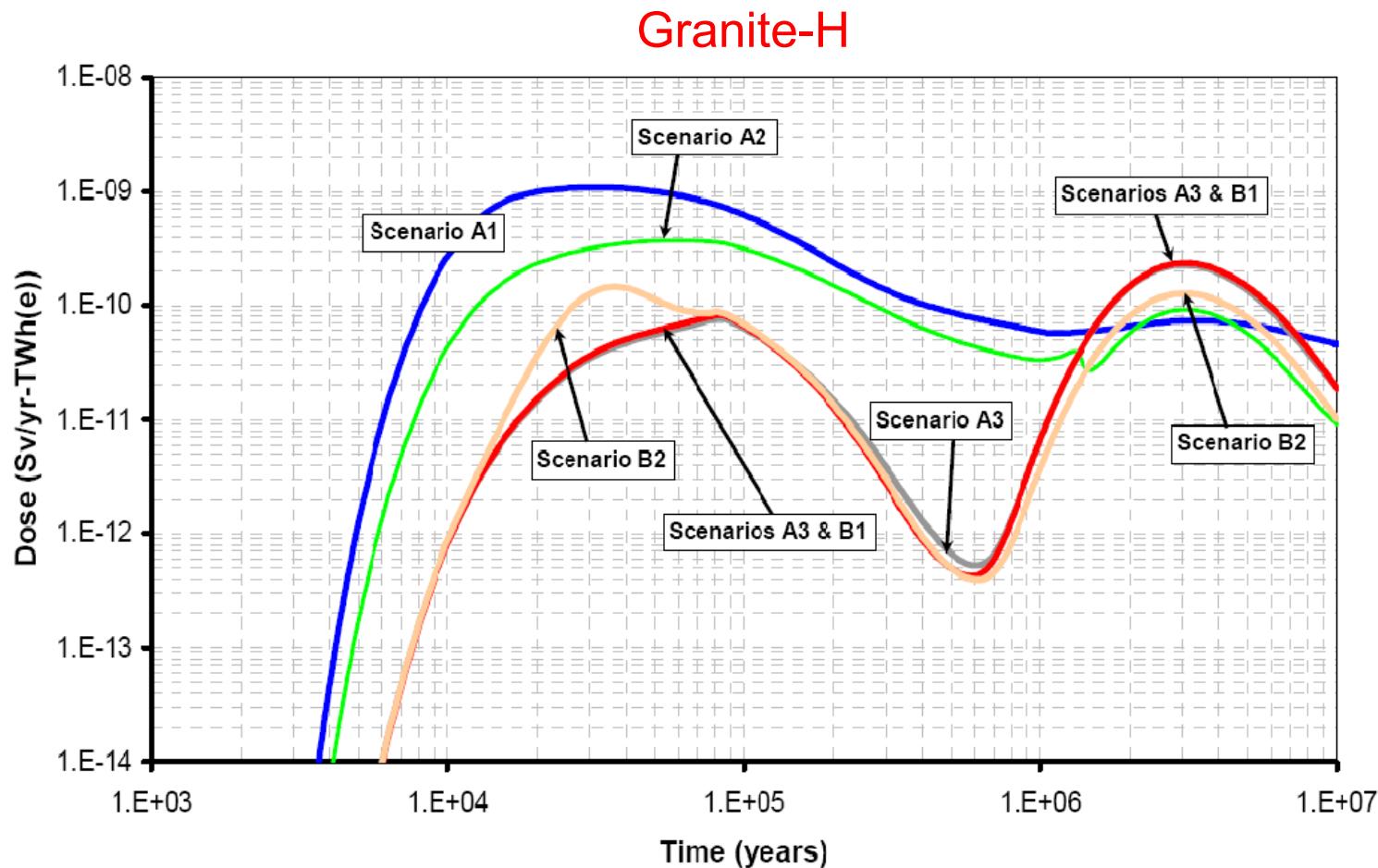
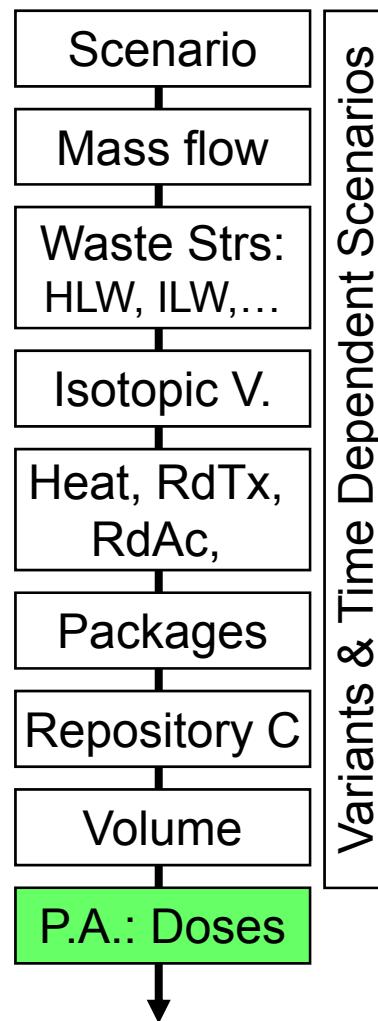


Individual Dose from repository: HLW



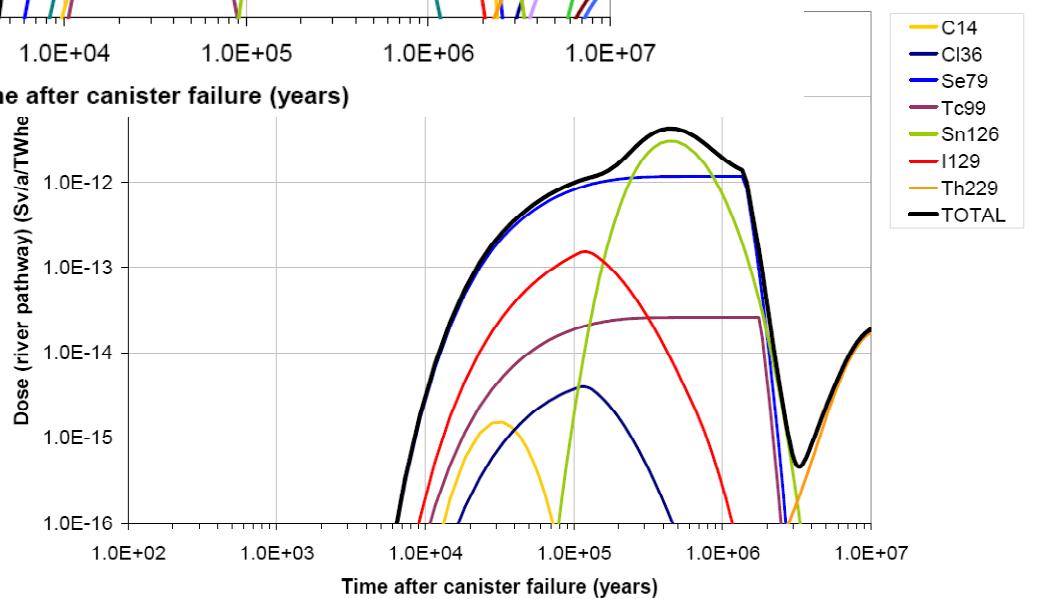
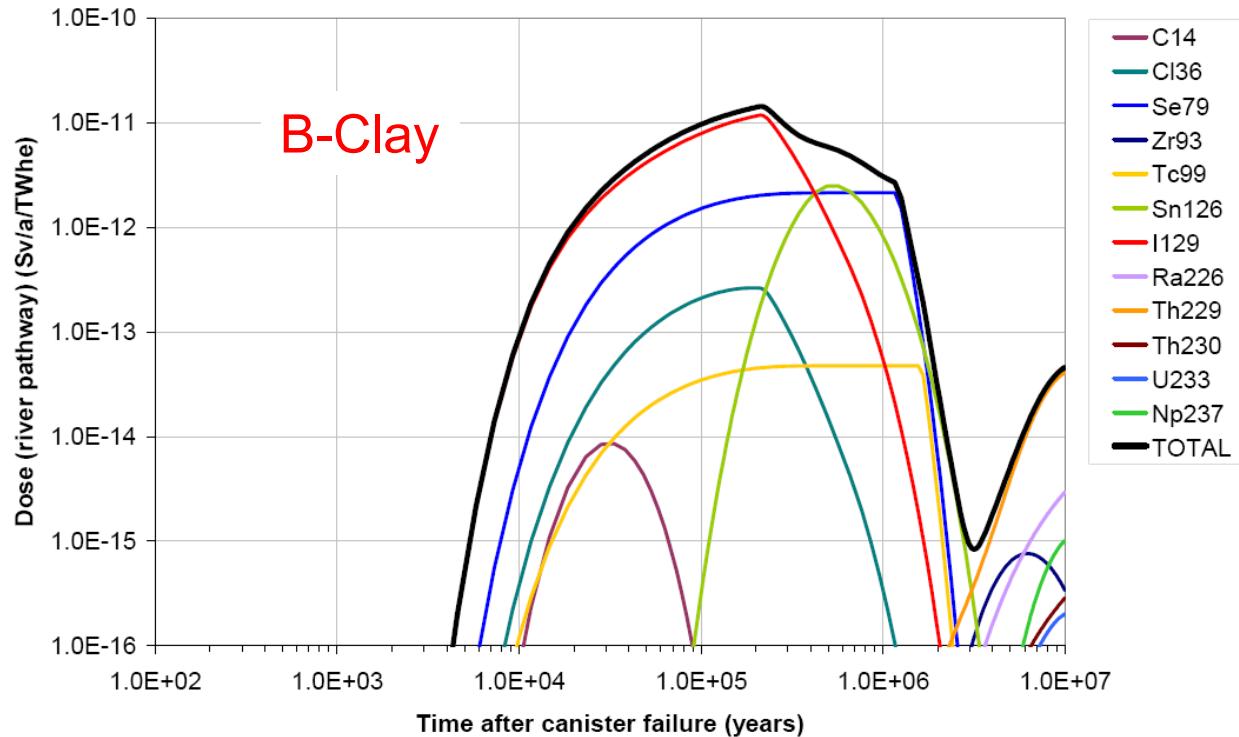
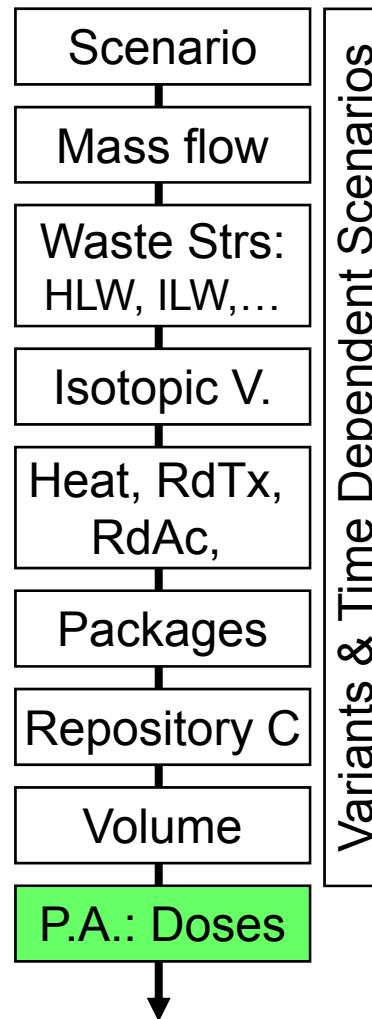
glass is assumed to dissolve completely in 72000 years

Individual Dose from repository: HLW

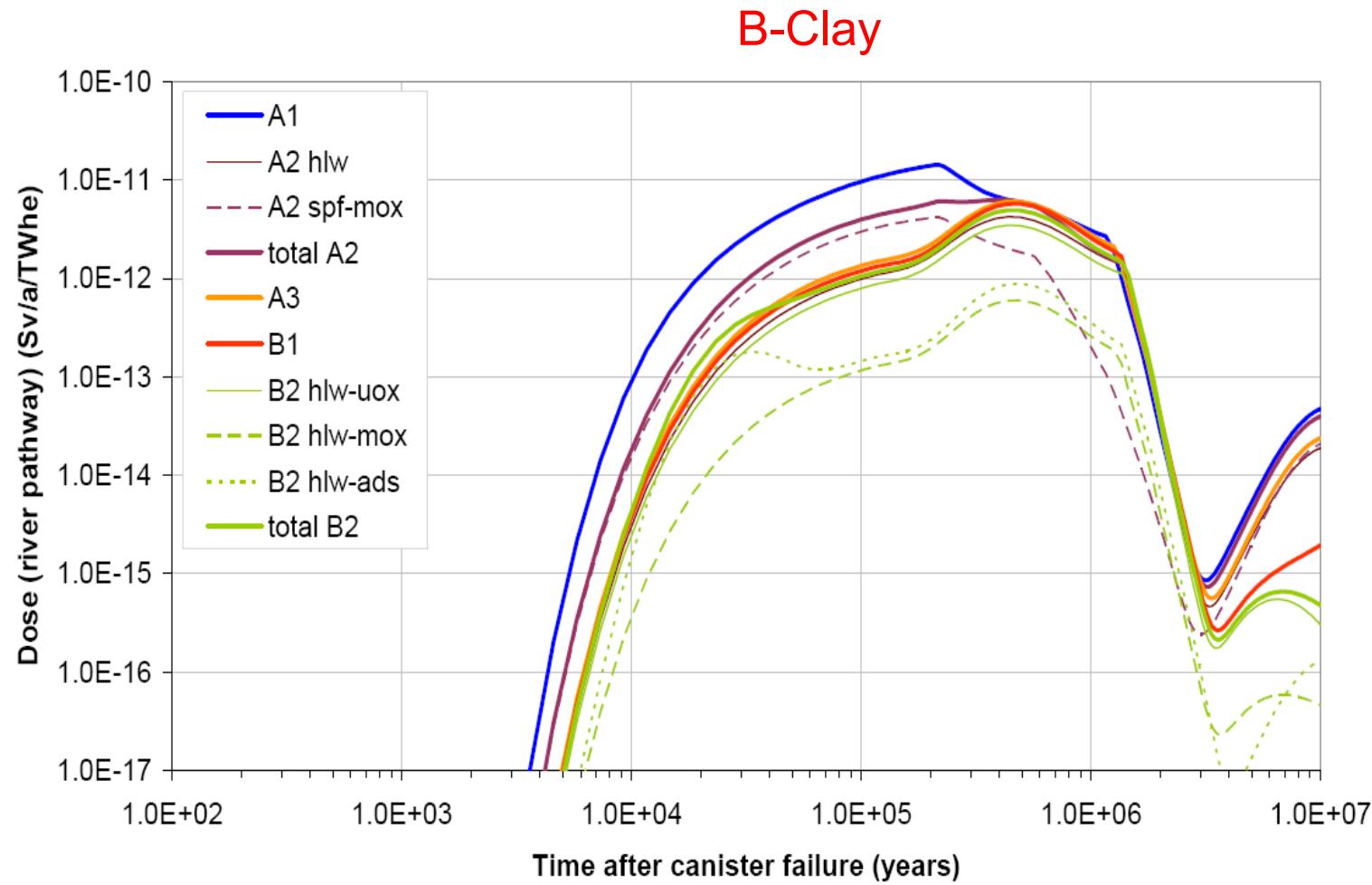
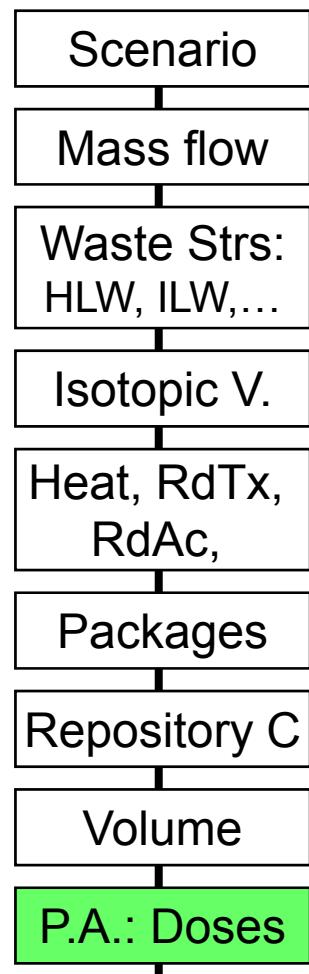


Note that the matrix lifetime has a strong effect on doses due to the mobile fission and activation products such as ^{129}I , ^{14}C and ^{36}Cl

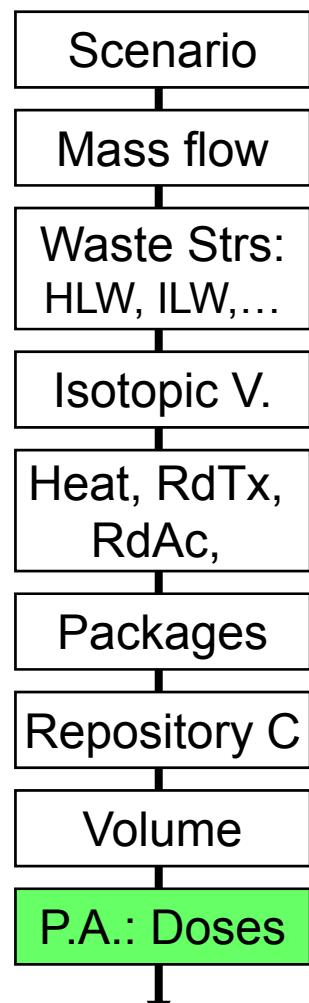
Individual Dose from repository: HLW



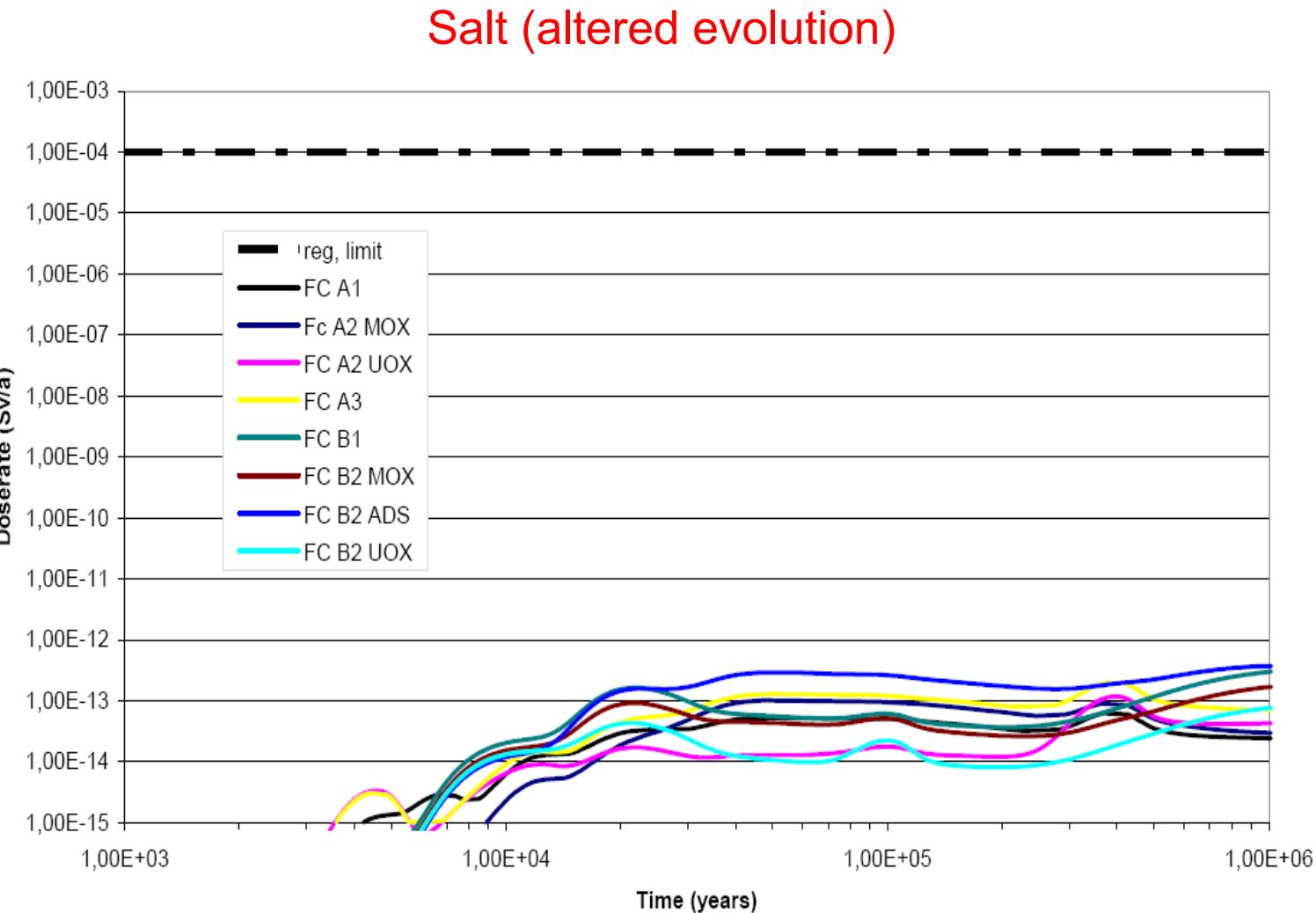
Individual Dose from repository: HLW



Individual Dose from repository: HLW

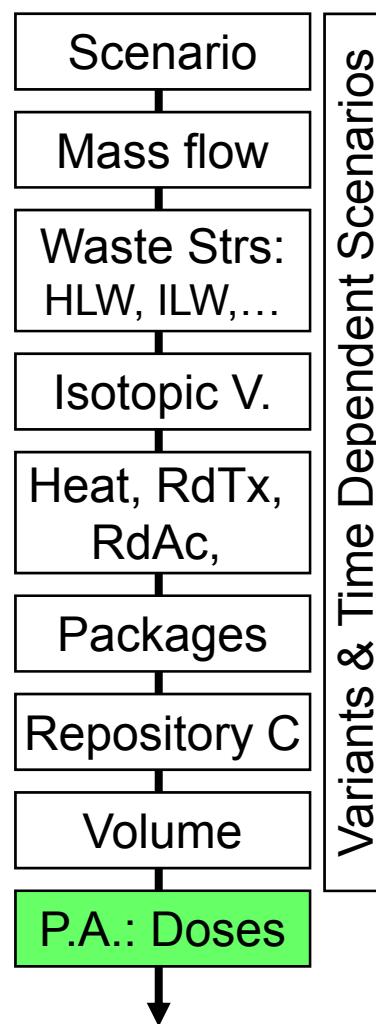


Variants & Time Dependent Scenarios





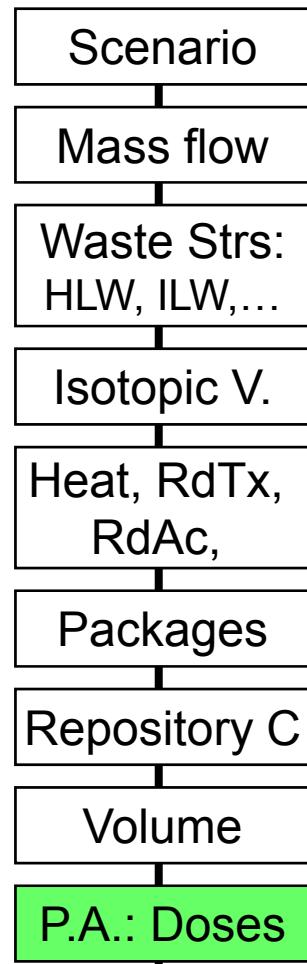
Individual Dose from repository: HLW



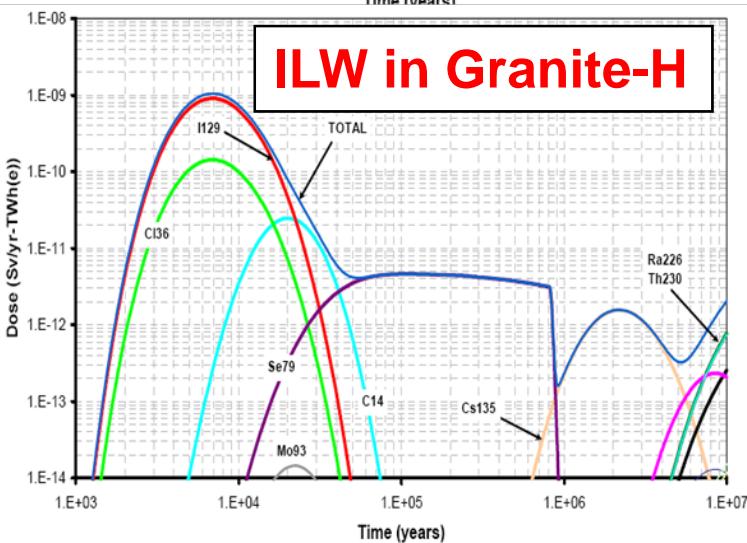
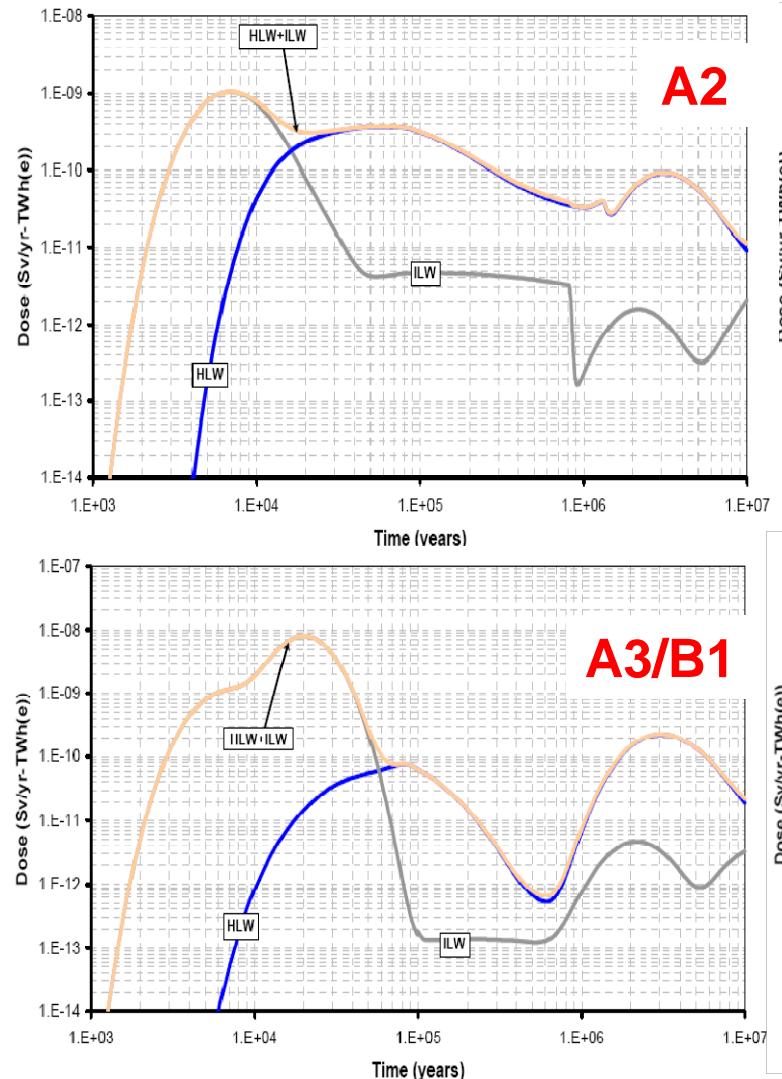
- For all considered host formations, the impact of P&T on the maximum dose is limited, because the maximum dose is essentially due to long-lived fission products.
- One of the most important contributors to the total dose is **129I**. The amount of **129I** going into the repository as HLW very strongly depends on the fraction of spent fuel that is reprocessed.
- For those long-lived fission products that pass to the HLW during reprocessing (**79Se**, **126Sn** and **135Cs**) doses arising from the different scenarios are quite similar.
- For **129I** and **14C** and **36Cl** (activation products) the inventories in the HLW are much smaller than in the original fuel, because **large fractions of those elements have been released as effluents**.
- The decrease in doses due to **129I**, **14C** and **36Cl** in the scenarios with reprocessing is a consequence of the reduced inventory in the HLW.
- The **transmutation of most actinides** in fast reactors or accelerator-driven systems in case of advanced fuel cycle scenarios **has little impact on the resulting doses**, because the low solubility of the actinides in reducing conditions and the strong sorption on minerals present in the buffer and host formation .
- When the waste disposal configuration is adapted to the thermal output of the disposed waste, the higher disposal density in case of advanced fuel cycles can result in a decrease of the release rate of solubility limited radionuclides (**79Se**, **99Tc** and **126Sn**,) in case of disposal.



Individual Dose from repository: ILW



Variants & Time Dependent Scenarios

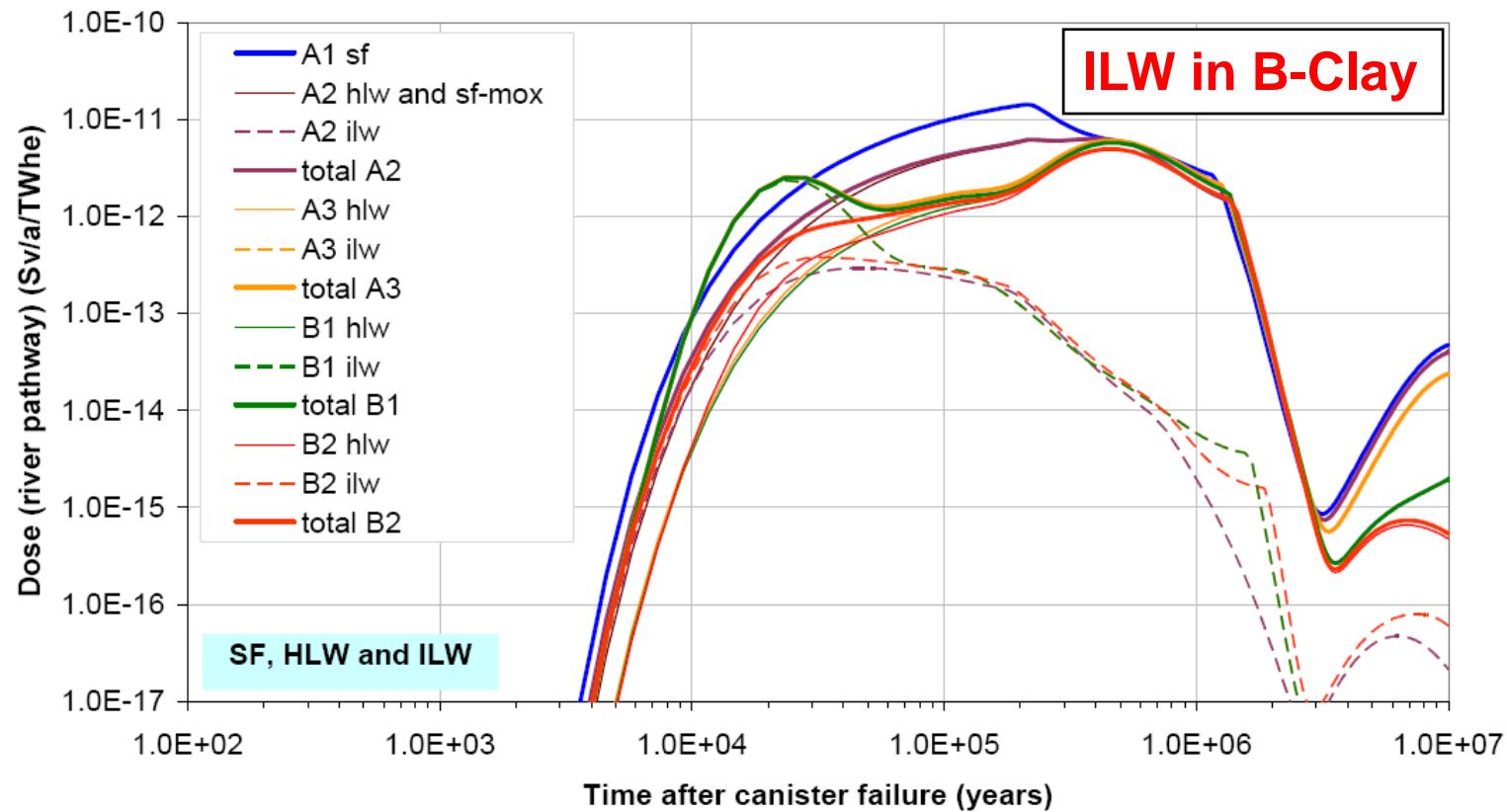
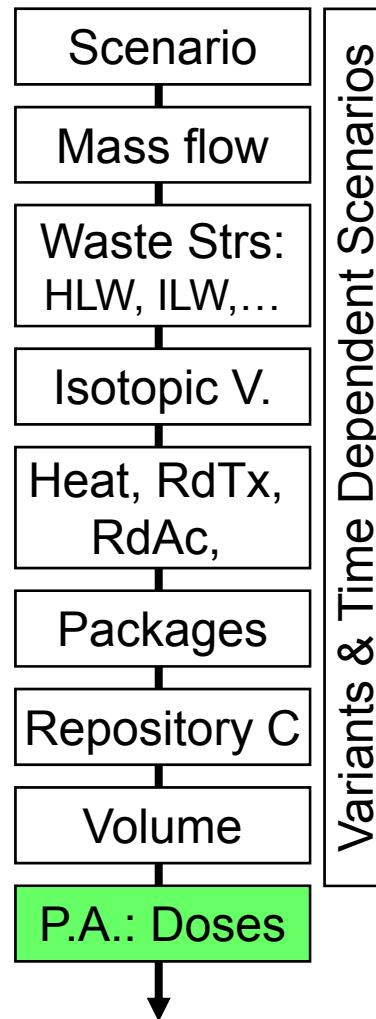


Dose from ILW dominated by I129, Cl36, C14, Cs135.
It was assumed that the waste matrix does not provide any isolation

Individual Dose from repository: ILW

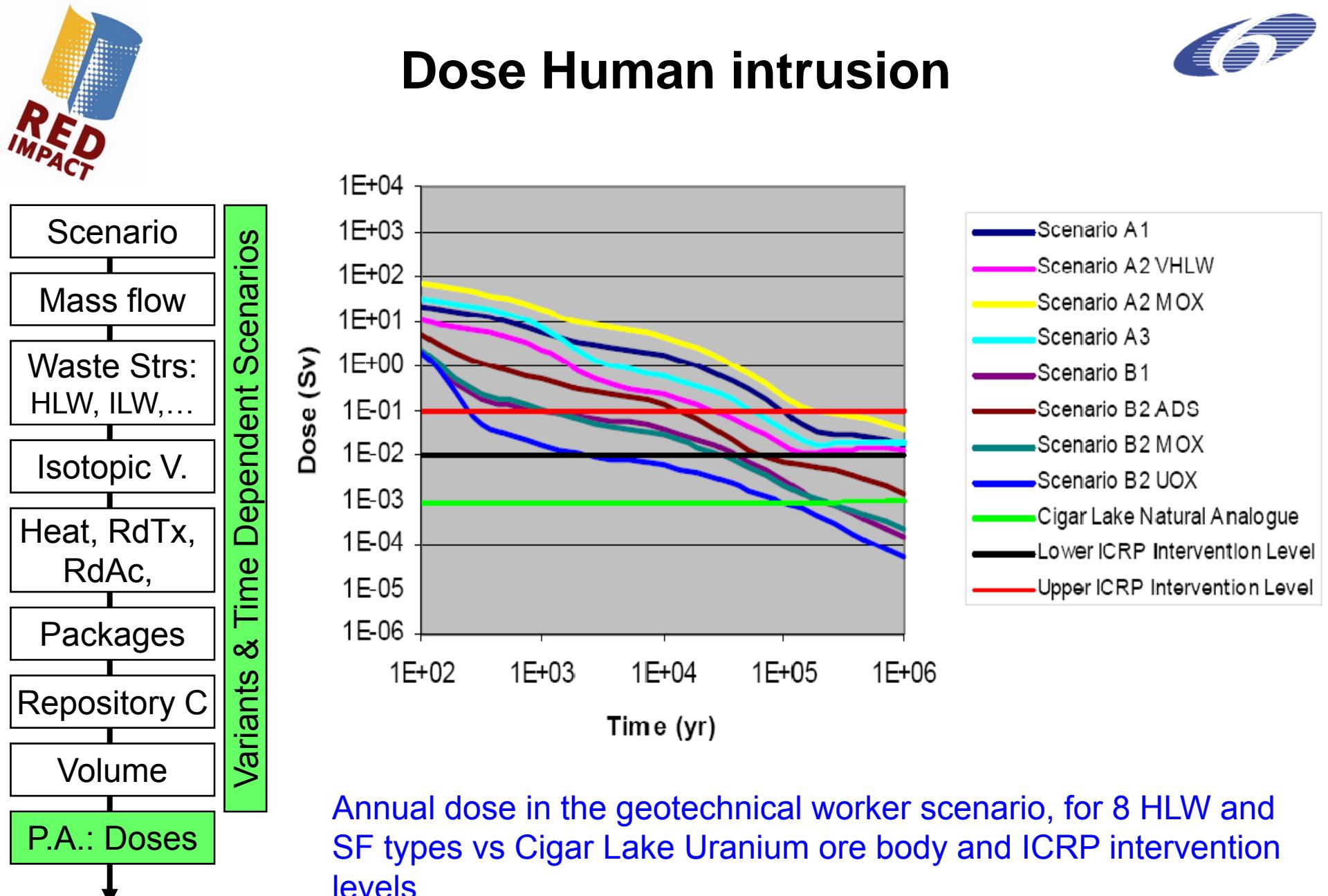


In B-Clay, the about 50-m thick clay barrier spreads the release of mobile fission and activation products from the host clay formation into the surrounding aquifer layers over several tens of thousands of years, minimizing their radiotoxicity at release time.



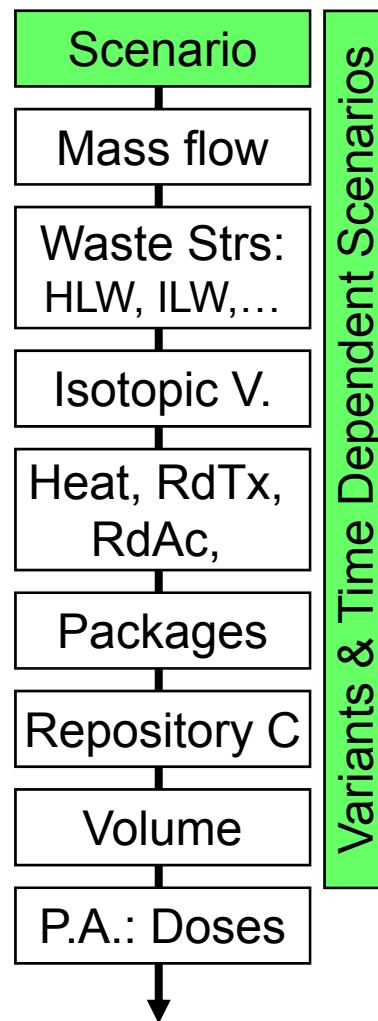
The main contributors to the total dose are ¹²⁹I, ³⁶Cl and ⁷⁹Se.
The matrix lifetime was taken equal to 1000 years.

Dose Human intrusion



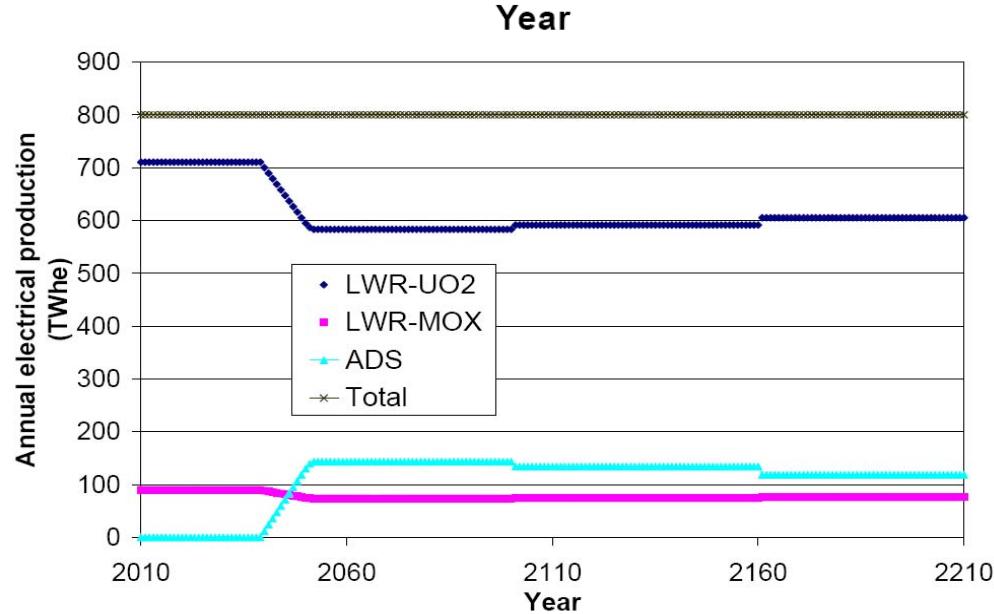
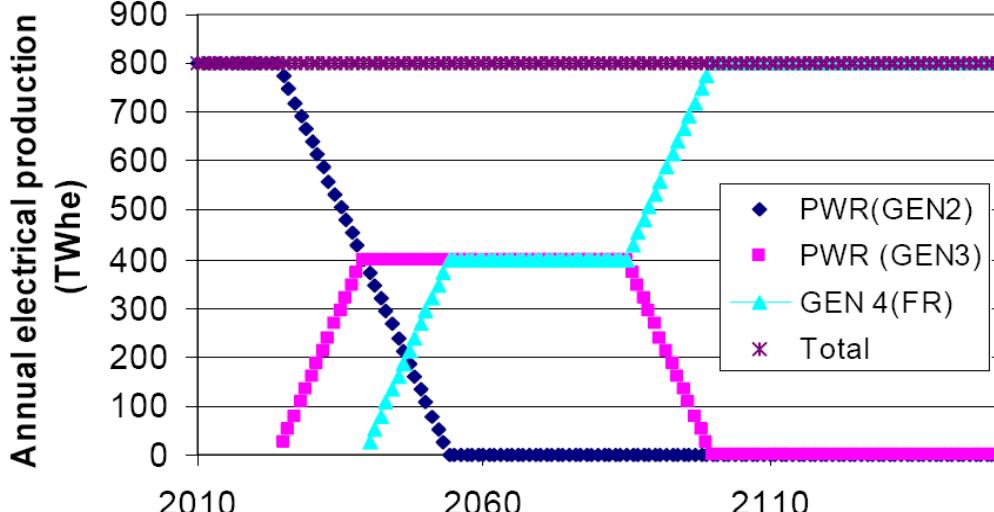
Annual dose in the geotechnical worker scenario, for 8 HLW and SF types vs Cigar Lake Uranium ore body and ICRP intervention levels

Time dependent Scenarios



Transition
Scenarios:
A1 to A3 or B1

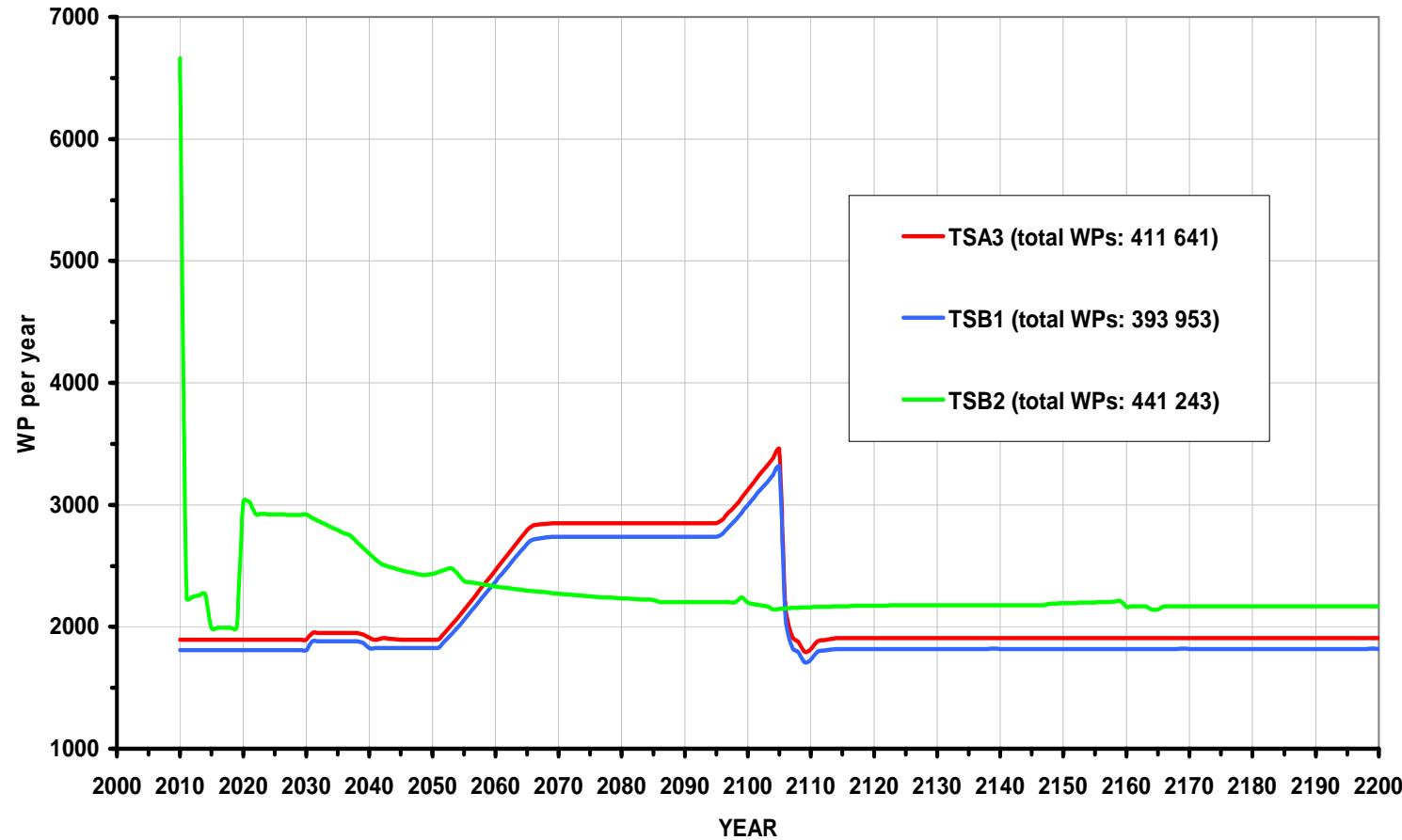
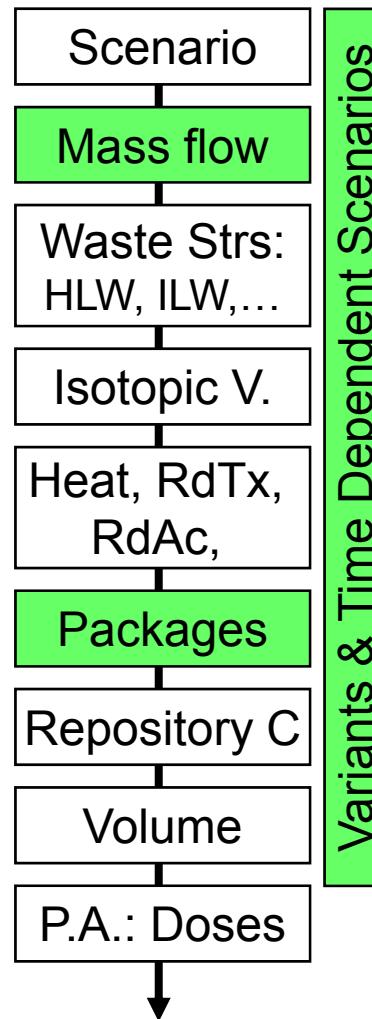
Transition
Scenarios:
A1 to B2



Time dependent Scenarios

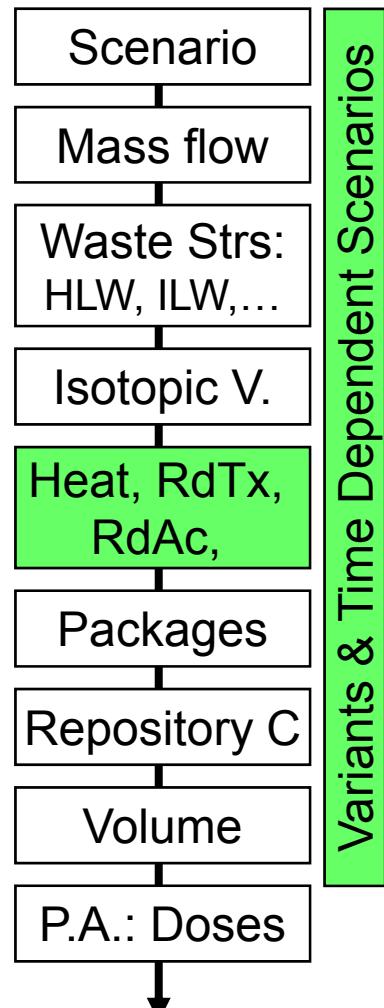


Annual production of HLW packages in Transition Scenarios

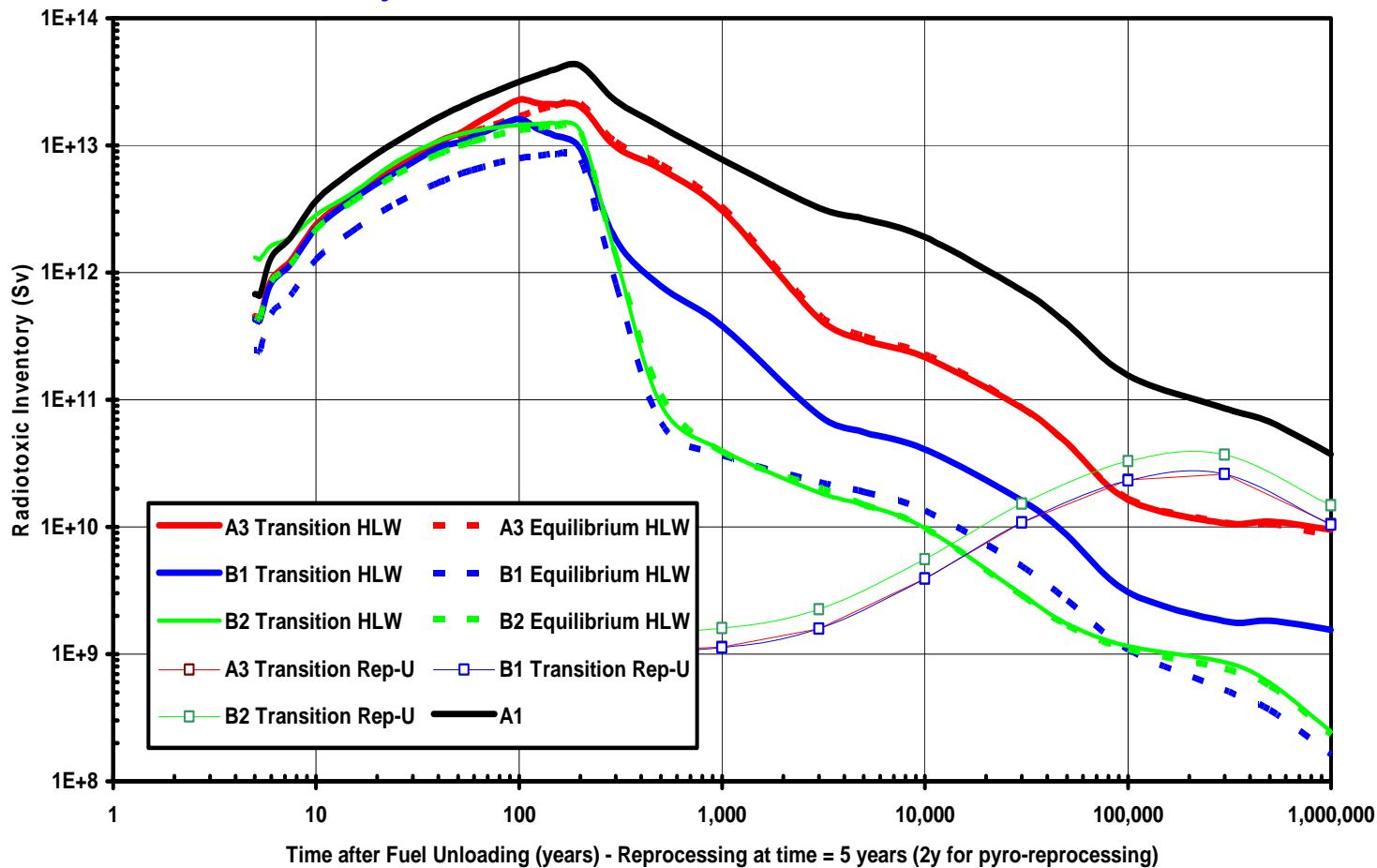


Detailed and well anticipated planning is needed to achieve equilibrated and optimized facilities and resources utilization.

Time dependent Scenarios



Radiotoxicity inventories for transition scenarios



Optimized program of reprocessing as a function of technology maturity to achieve maximum improvements on radiotoxicity inventory and capacity of the repository.

Red-Impact main conclusions

FP6 RED-IMPACT project has extended the 2002 & 2006 NEA/OCDE studies on P&T adding improvements on:

- A more complete evaluation of wastes:
 - intermediate level wastes, ILW,
 - more detailed structural materials and fuel impurities activation
 - the activation of all the specific components of the ADS (spallation target structural materials and coolant activation, spallation products activation,...).
- More details in the Cs & Sr separate handling scenarios,
- The performance assessment, including in this case the handling of the ILW, has been improved,
- A large effort to evaluate the transition process from the present European situation to hypothetical future scenarios.
- A first methodology to include indicators for all aspects related to waste management (technical, economical, social, environmental,...) in a single evaluation mechanism



Red-Impact main conclusions



- P&T reduces the final waste Radiotoxicity and its Thermal Load from the wastes to the repository, and strongly accelerates the Thermal load decay after 100 years.
- This will allow to reduce the gallery length required for HLW in granite and clay repositories, increasing their capacity (factor 1.6-6). The effect is enhanced by delaying disposal to >100y from discharge, separating Sr &Cs handling or both.
- There is small or no advantage from P&T on the dose to the average member of the critical group from the normal evolution scenarios. But significant advantage for improvable human intrusion scenarios.
- ILW could compromise the advantages if prevention is not taken:
 - Large volume of ILW (where to store then, compacted packages?, ...)
 - Dose to average individual larger than from HLW (how to isolate then?)
- Transition scenarios show the importance of long term planning of resources and infrastructure availability for the feasibility and performance of P&T.
- Transition scenarios indicates that using early P&T technology before is fully functional (MA handling) might limit the final performance for scenarios mainly oriented to reduction of wastes (like reduction of nuclear park).
- Independently of the minimization technologies used, there will be always need of highly isolated final repository for the residual (losses,...) HLW.