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MSRs: potential for modularity and sizing

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Interregional Workshop on Advances in Design of Generation-IV
Small and Medium Sized or Modular Reactors (SMRs)

Safety consideration:

Basic safety features

Source term mobility

Criticality safety

Fuel cycle considerations

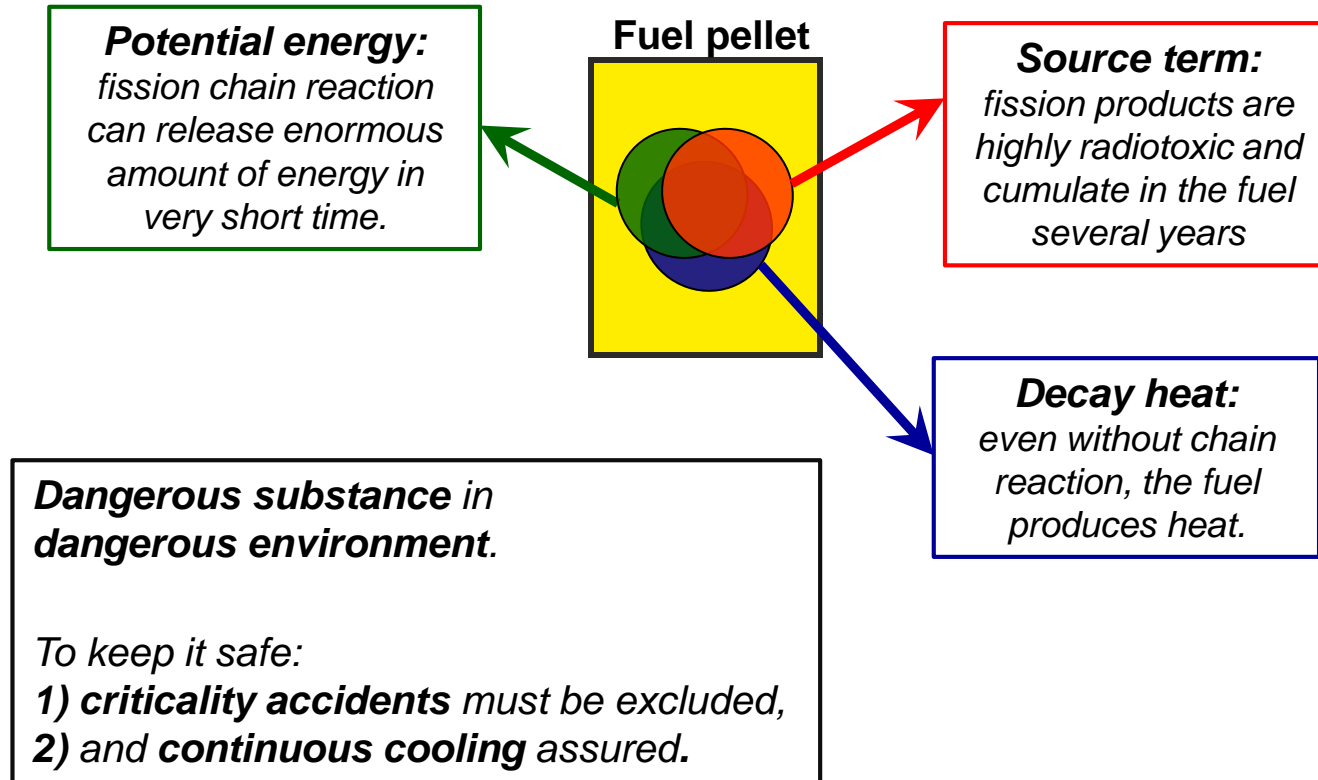
MSR breeding capability assessment

Core size and transition to Th-U cycle

Breed and burn cycle

Safety considerations: source term and driving forces

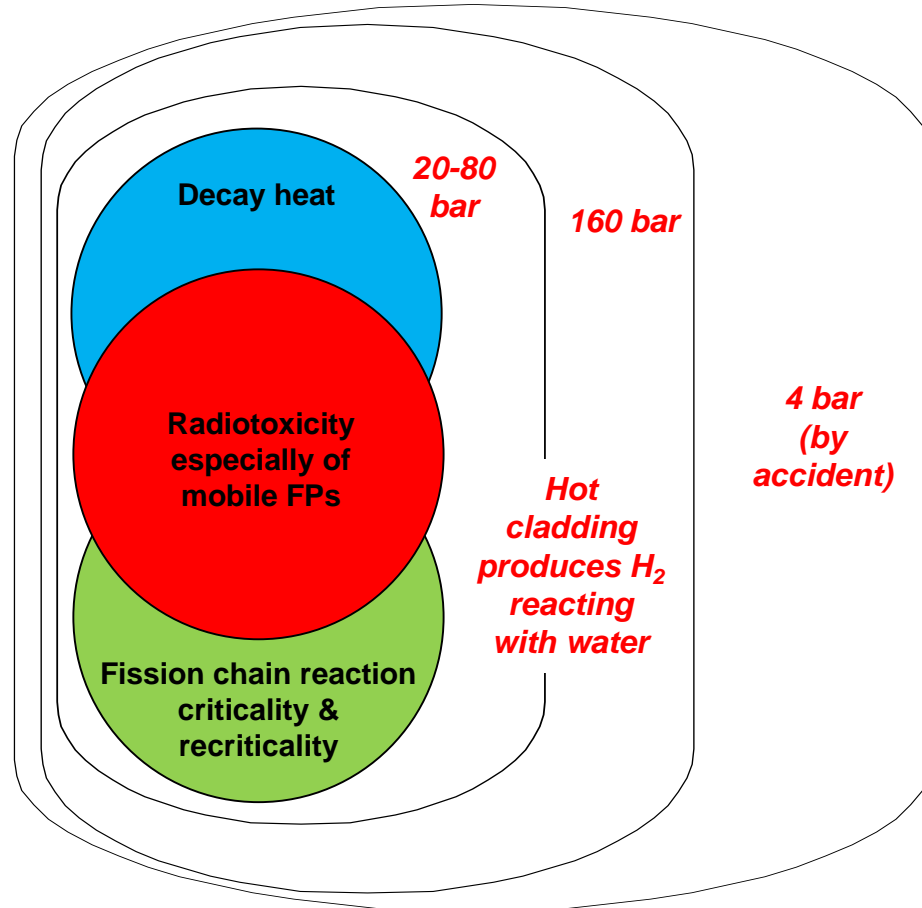
Why is nuclear safety so particular?



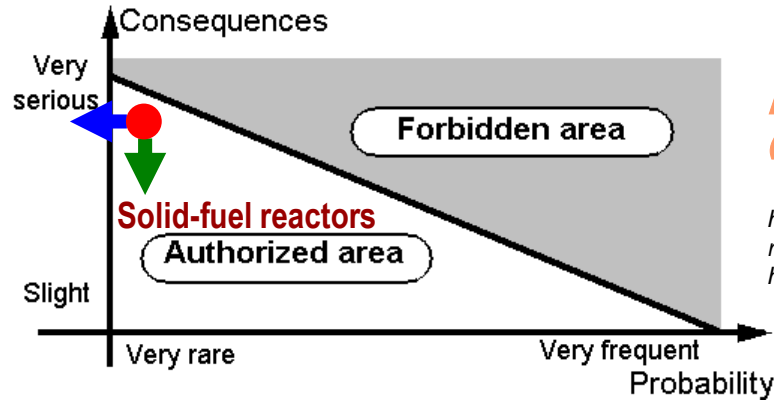
PWR overview: confined hazard

PWR

- *Low criticality & recriticality risks.*
- *Pressurized pin.*
- *Pressurized coolant.*
- *Reactive cladding.*
- *Containment should withstand the steam pressure (& non-condensables).*



Risk = probability X consequences (Farmers curve)



*Farmer curve illustration
(log-log scale)*

<http://www.iaea.org/ns/tutorials/regcontrol/assess/assess3212.htm>

2 basic directions to improve the position of solid-fuel reactors:



Reducing consequences = reducing source term:

*Decreasing installed power or fuel burn-up
(small modular reactors, shorter fuel cycle)*



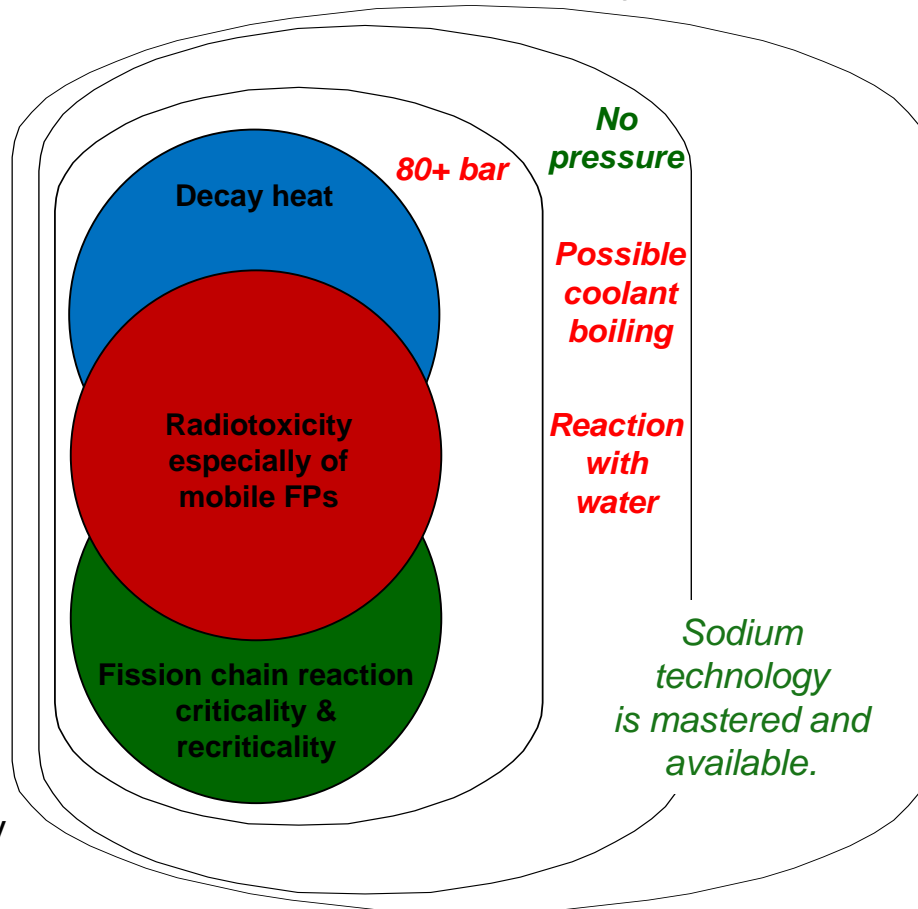
Adding barriers or decreasing the frequency of their failure:

*Adding second or third containment...?
Passive systems, redundancy, diversity, underground reactors, etc.
Filtered venting of containment = clever chemical barrier.*

Sodium cooled fast reactor safety

SFR

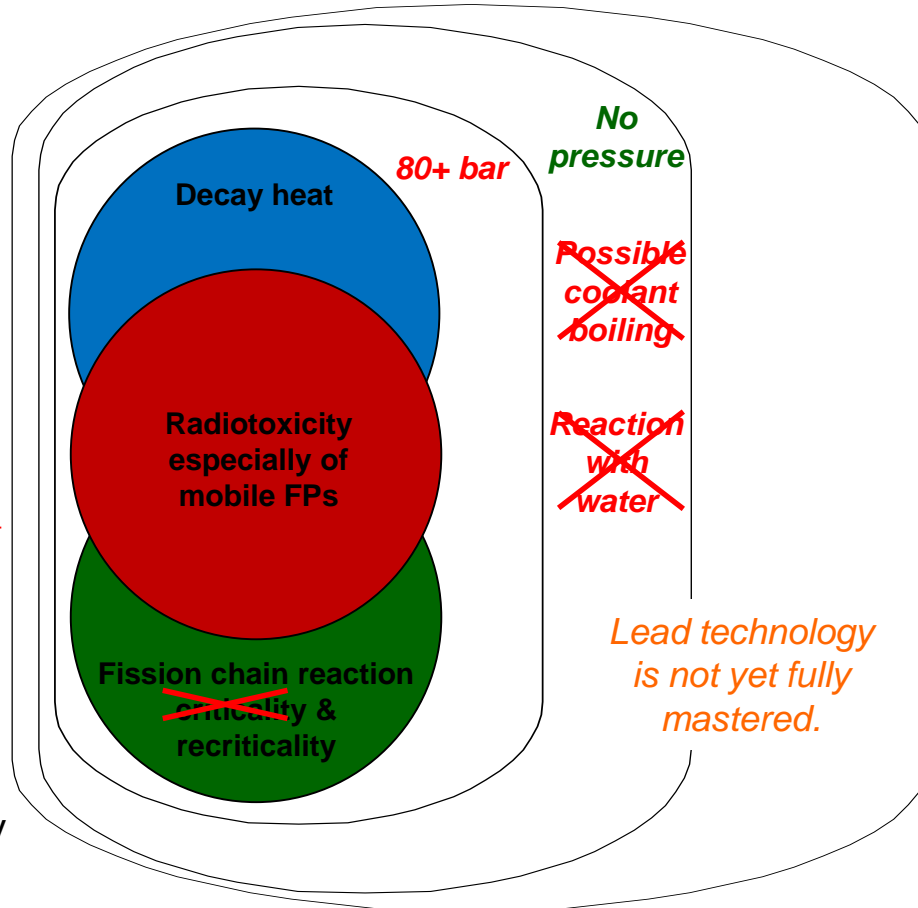
- **Higher burn-up =**
 - 1) higher source term of radiotoxicity
 - 2) higher decay heat
 - 3) higher pin pressure
- **High criticality & recriticality risks.** (enriched fuel, positive void effect)
- Coolant not pressurized but reacting with water.
- **Pool type design** may prevent LOCA accidents.



Lead cooled reactor safety (comparison to SFR)

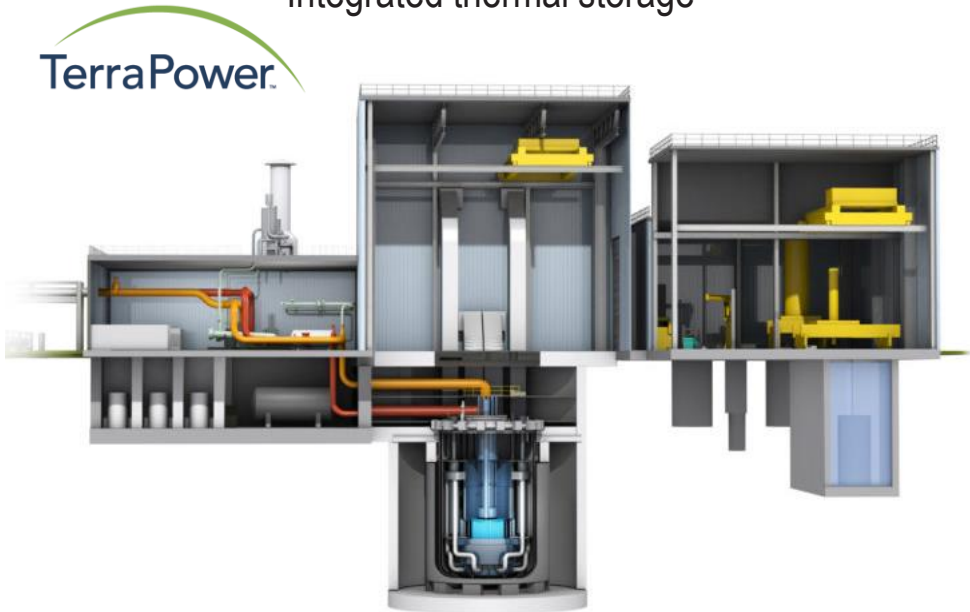
LFR

- **Higher burn-up =**
 - 1) higher source term of radiotoxicity
 - 2) higher decay heat
 - 3) higher pin pressure
- **High criticality & recriticality risks.**
(enriched fuel, ~~positive void effect~~)
- Coolant not pressurized ~~but reacting with water.~~
- **Pool type design** may prevent LOCA accidents.

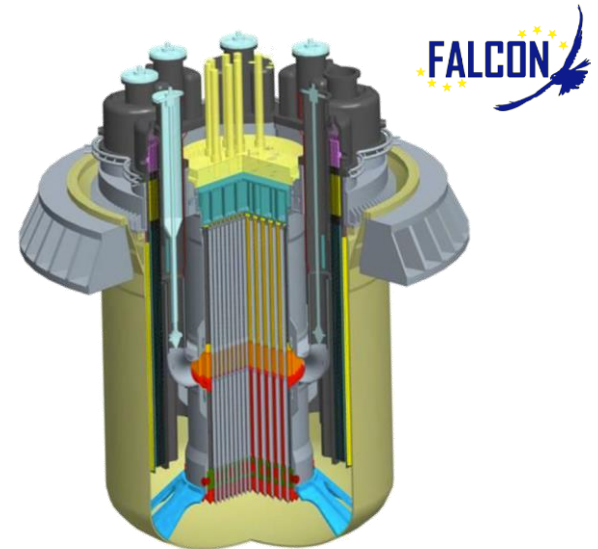


Two examples of LFR and SFR SMRs

Natrium (US)
Small modular SFR
345 MWe
Integrated thermal storage

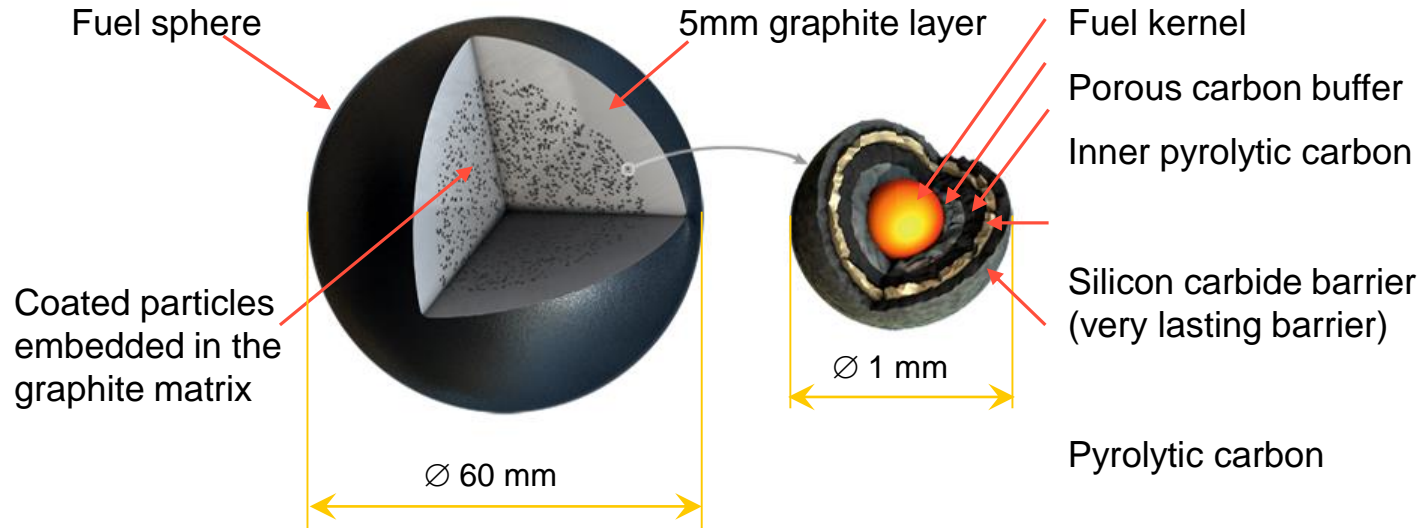


ALFRED (EU)
Small modular LFR
125 MWe
Oxygen control to limit corrosion



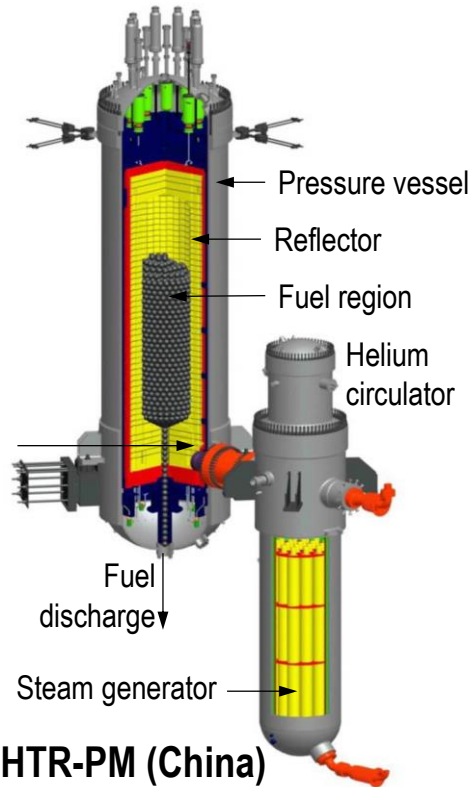
HTR with TRISO particles – different confinement

- **TRISO** (*TR*istructural-*ISO*tropic) **coated particles**



- **Tight up to 1400-1600 °C.**
- *Even above, no wild changes, melting, or degradation, just FPs release.*
- *If cooled by salt with low pressure, no driving force for FPs release.*
- *Very complicated reprocessing, but suitable for deep burnup.*

Two examples of HTR



MSR with liquid fuel

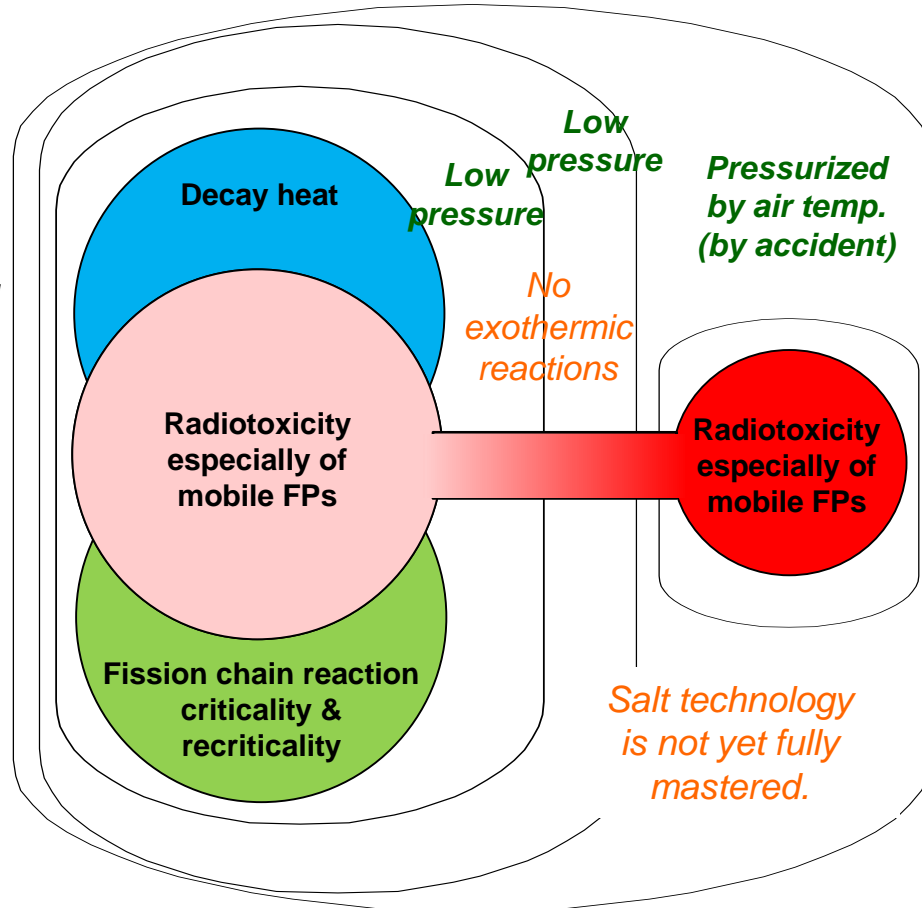
- **Liquid fuel**, similarly as the solid fuel, retains: uranium, TRU, and soluble FPs.
- On the other hand, it provides **zero confinement for mobile FPs!**
- Not only that the **mobile FPs** are not confined, they **must be removed** from the fuel.
- **Decoupling of mobile FPs from major heat sources** may strongly change **safety philosophy**.
- Such a decoupling is possible only in systems with **liquid fuel**.
- Or in reactor with vented assemblies (engineering challenge).



Change of safety philosophy: decoupling from heat sources

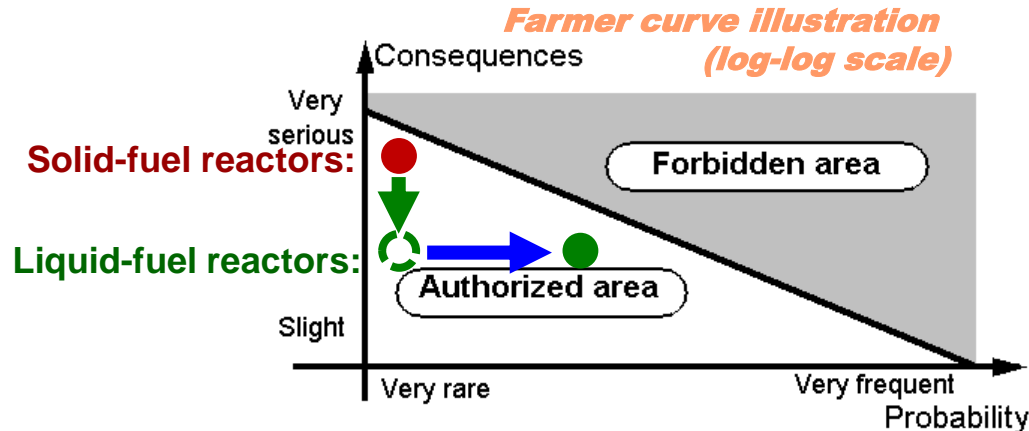
MSR

- *If the mobile FPs would be decoupled from the major heat sources, the risk would be strongly reduced.*
- *Deterministic reduction of serious hazard and consequences.*
- *Regulation for reprocessing plants may be applicable to the removed FPs.*



Risk = probability X consequences (MSR case)

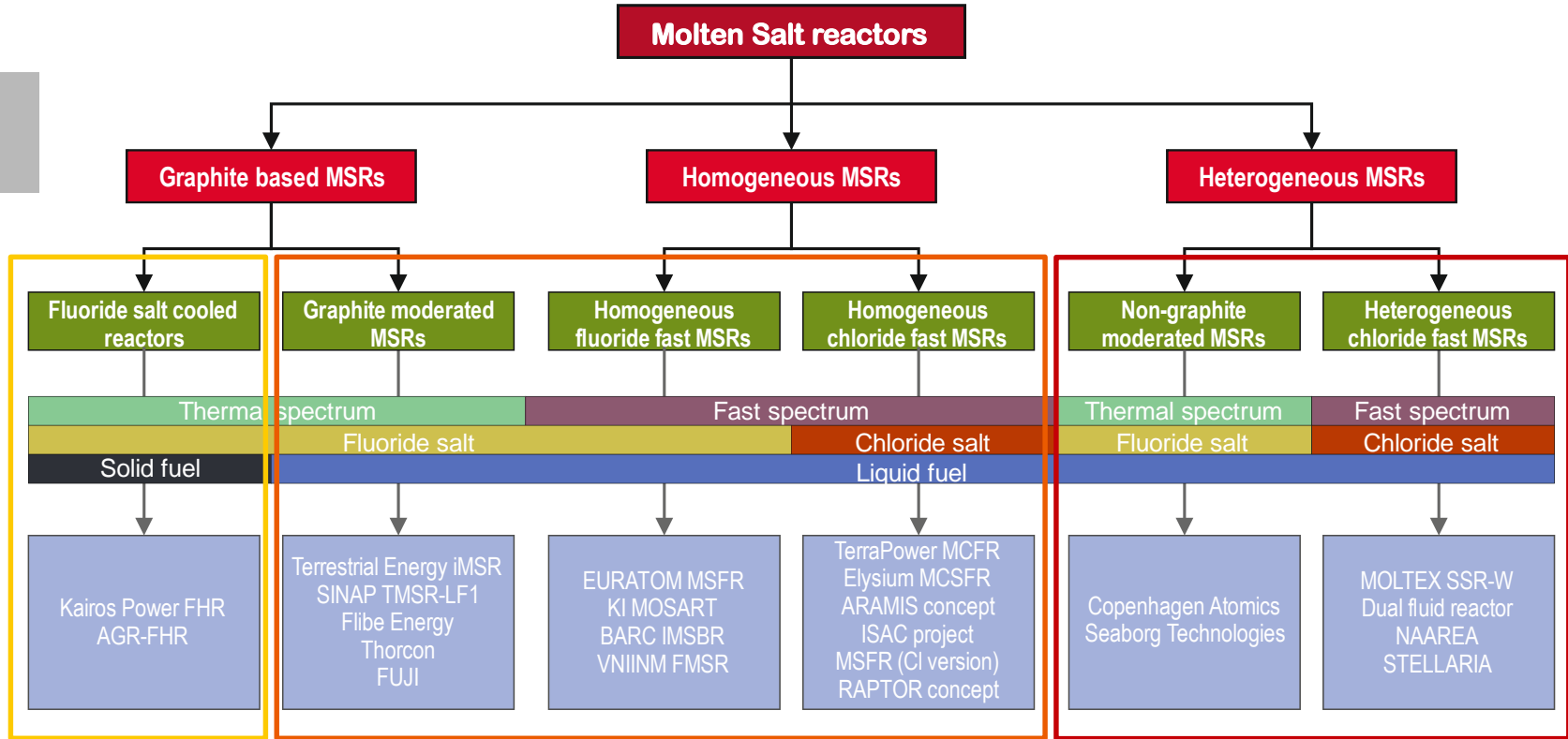
- **Technological requirement** of volatile FPs **continuous removal** may, together with the negative temperature feedbacks (criticality safety) and possible passive fuel drainage, provide “**deterministic**” safety.



<http://www.iaea.org/ns/tutorials/regcontrol/assess/assess3212.htm>

- ▼ **Online removal of highly mobile gaseous and volatile FP**
 (Ac and remaining FP are embedded in the salt)
- ➡ **Daily removal = possible reduction of classical barriers**
 (safety standards for reactor X reprocessing plant)

MSR as a reactor category rather than a single system



**HTR-like safety
(salt as a coolant)**

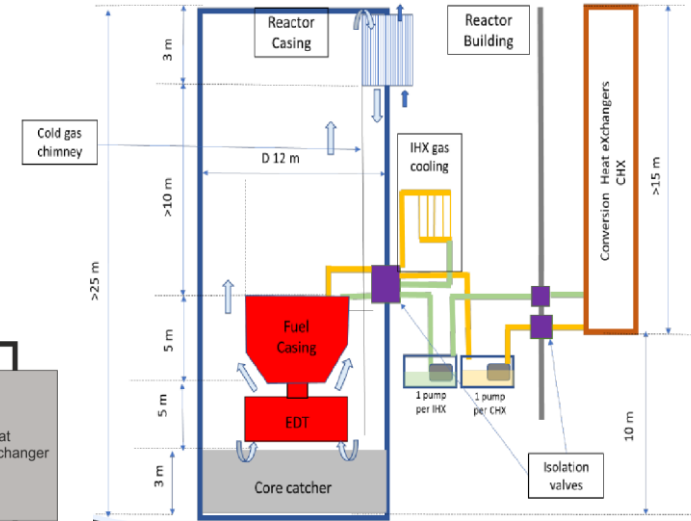
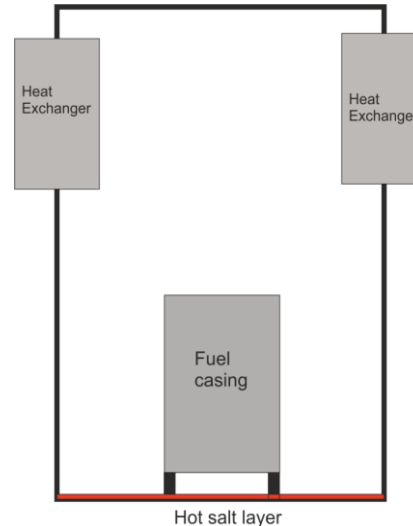
**Circulating fuel possibly with online
cleanup / offline reprocessing**

**Circulating or solid like fuel
(possibly sealed in pins)**

Safety considerations: source term mobility

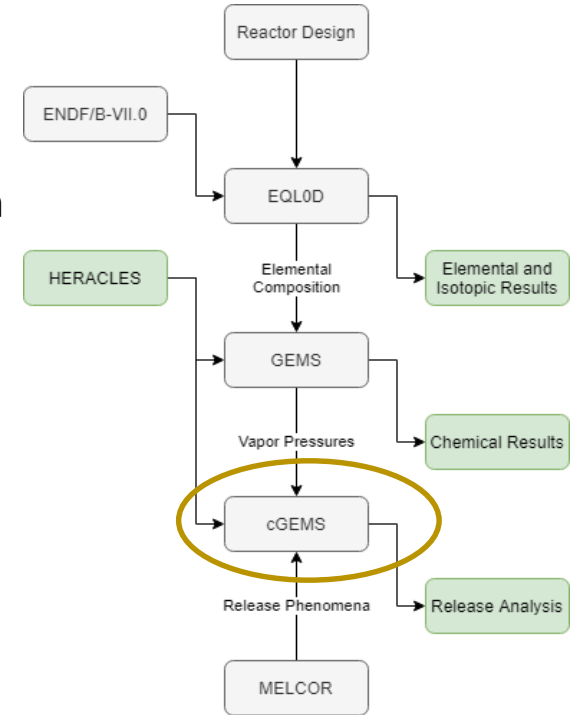
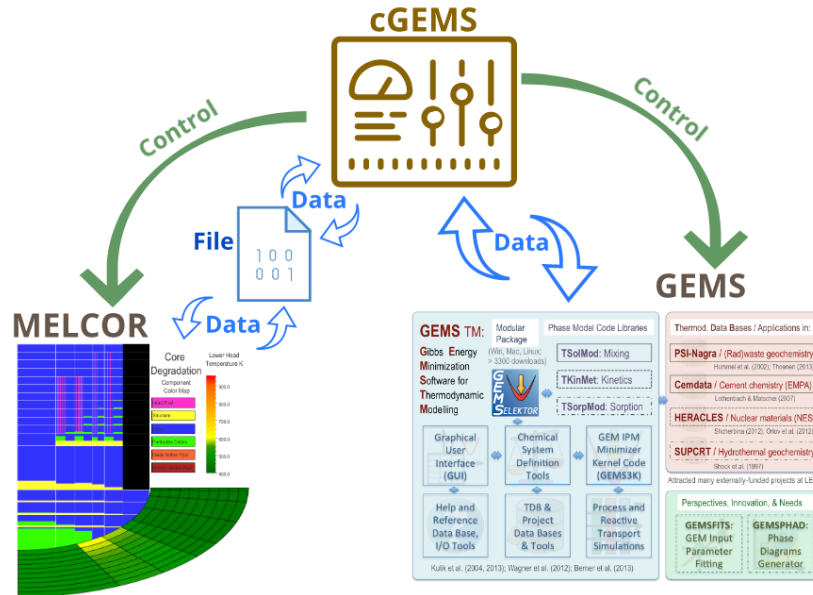
Simulation of severe accident in MSFR

- In the frame of SAMOSAFER project the deliverable **Aerosols formation and filtration in accidental conditions** was resealed. It is not public, but the results have been also published e.g. in Journal of Nuclear Materials:
J. Kalilainen, S. Nichenko, J. Krepel:
“Evaporation of materials from the molten salt reactor fuel under elevated temperatures”
<https://doi.org/10.1016/j.jnucmat.2020.152134>
- The simulated scenario assumes **salt spill at the bottom of the containment.**



From the SAMOFAR Final meeting, E. Merle et al.

- cGEMS code characteristics:
 - Composition of the salt from the EQL0D simulations
 - Uses the updated HERACLES database in GEMS software
 - Li, F, U, Th, Cs, Ba, Pu, Sr, La, Zr, Ce, Np, Nd extended system
 - The data exchange between MELCOR and GEMS: cGEMS

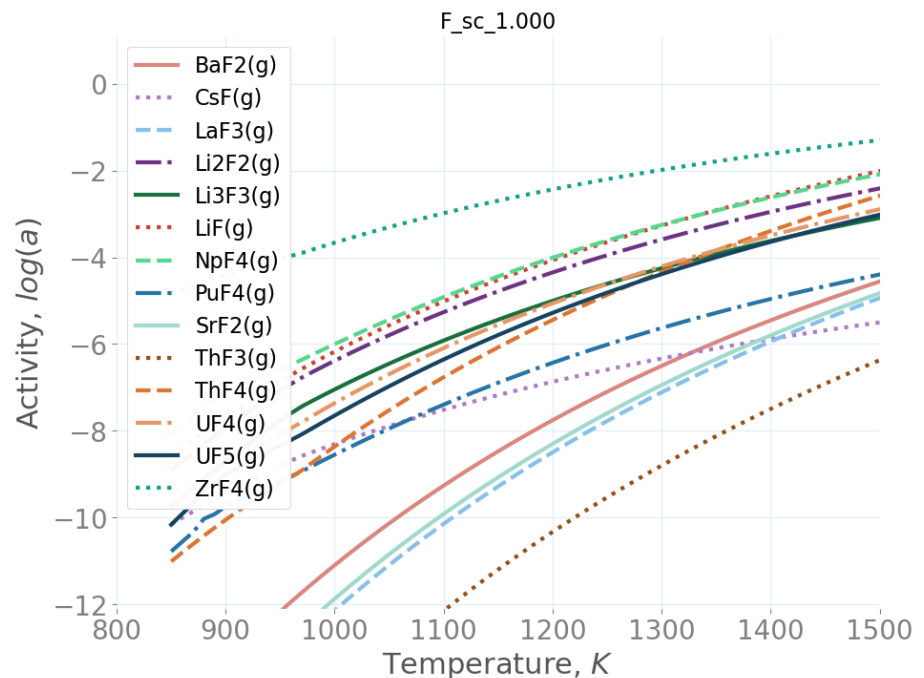


- For the simulation Heracles database of the GEMS code was extended.

Species	Changes Made
ThCl ₄	Imported as is from literature
Np	Imported as is from literature
PuCl ₃	Adjusted previously existing data entry to conform with literature melting point
UCl ₃	Missing liquid phase data manually matched based on literature values
NpF ₃	Missing liquid phase constructed from melting-/boiling points and similarity to UF ₃
AmF ₃	Solid adjusted and liquid designed from assumed similarity to UF ₃
ZrF ₄	Imported as is from literature
NdCl ₃	Imported as is from literature
PrCl ₃	Imported as is from literature
PrF ₃	Imported as is from literature
Na ₂ ThCl ₆	Created in GEMS function ReacDC
Pr	Imported as is from literature

Additional Changes were made to:
NpF₄, NdF₃, SrF₂, LaF₃, CeF₃, BaF₂, CsF

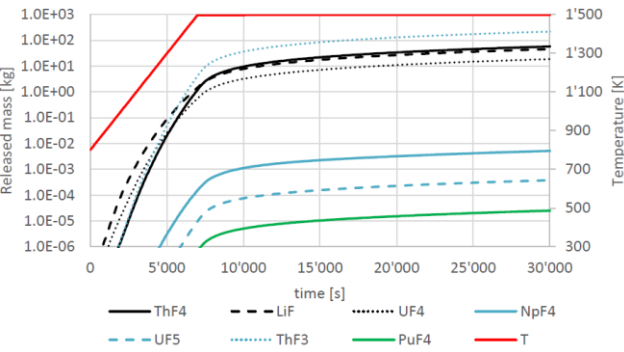
- GEMS code was applied to obtain the vapor pressures.



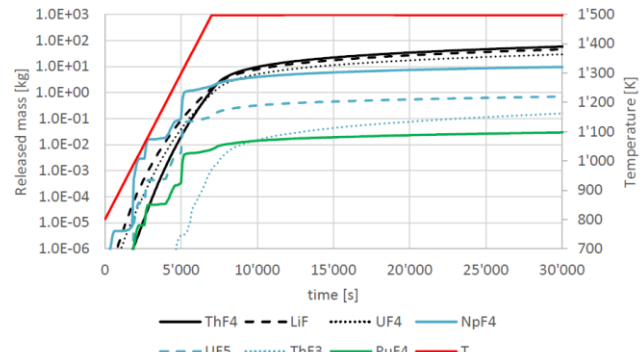
Compounds activity (proportional to vapor pressure) as a function of temperature

Released mass sensitivity to redox

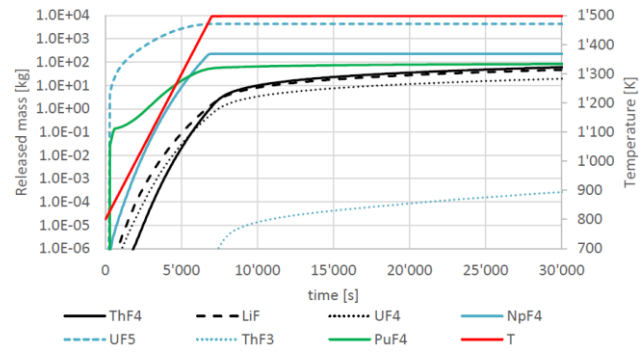
Fluorine: -1% mol.



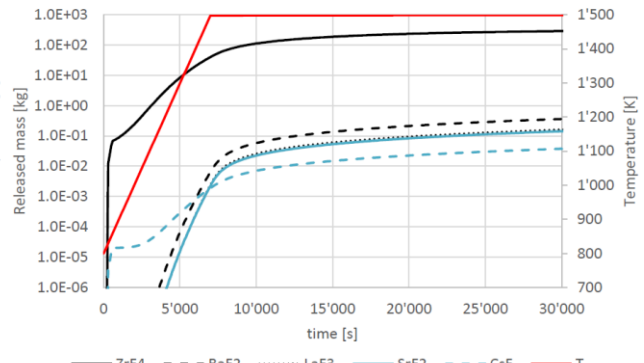
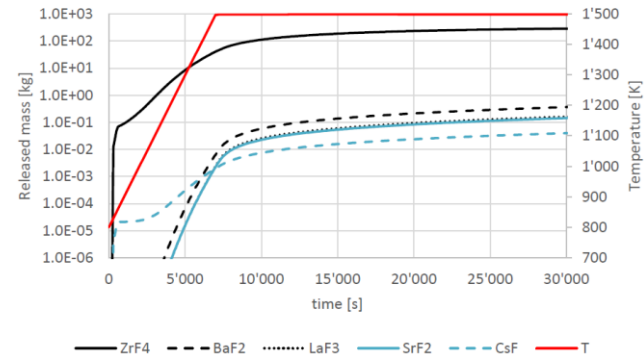
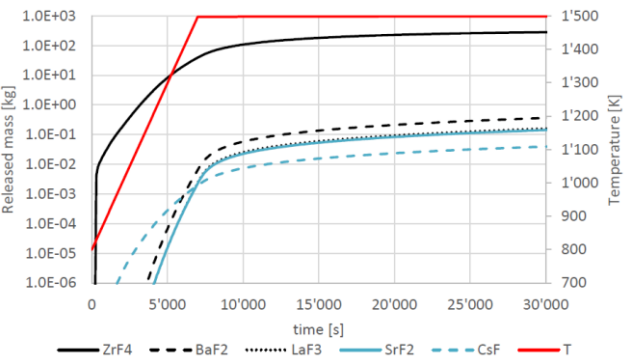
Stoichiometric



+1% mol.



Actinides and salt species

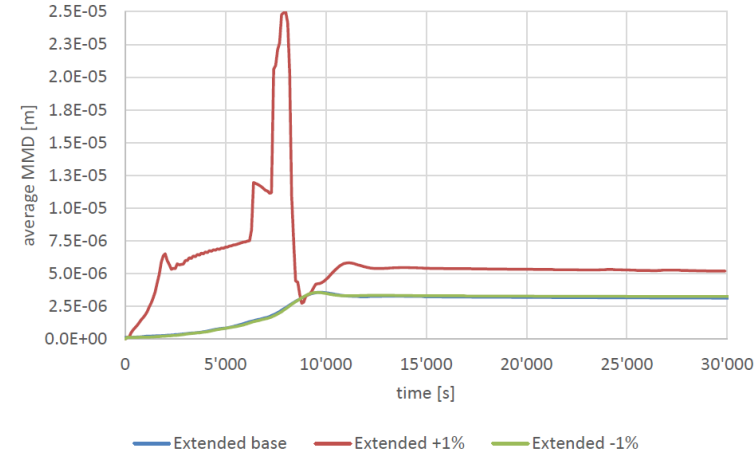
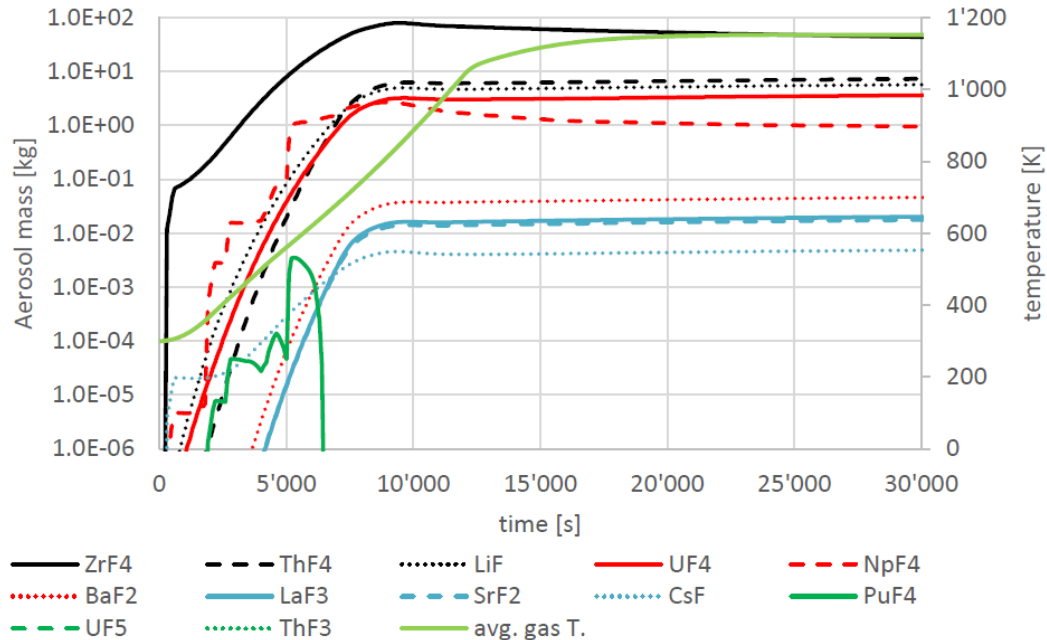


Fission products species

Total released mass during the accident (salt heat up from 800°C to 1500°C)

Released aerosols mass (cGEMS)

- Characterization of released aerosols

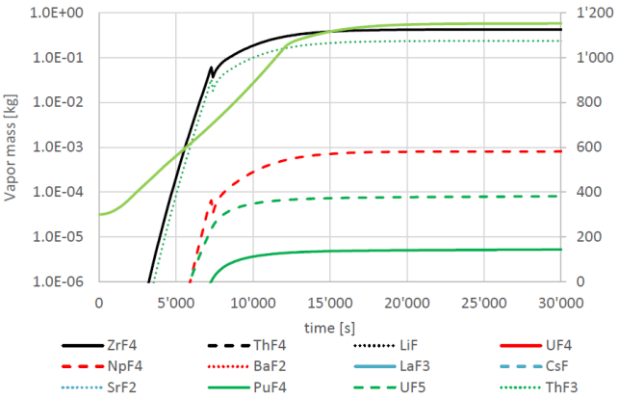


Total released mass of aerosols (left) and their mass median diameter (MMD) during the accident (salt heat up from 800°C to 1500°C)

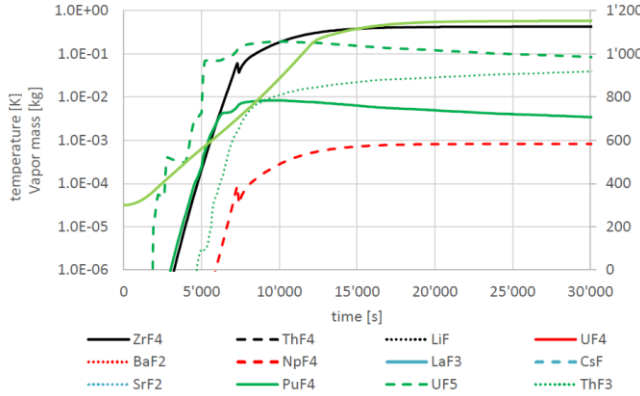
Released vapors mass (cGEMS)

- Characterization of released vapors

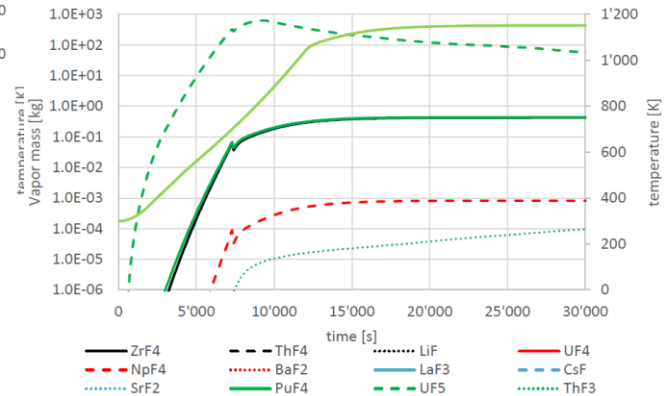
Fluorine: -1% mol.



Stoichiometric



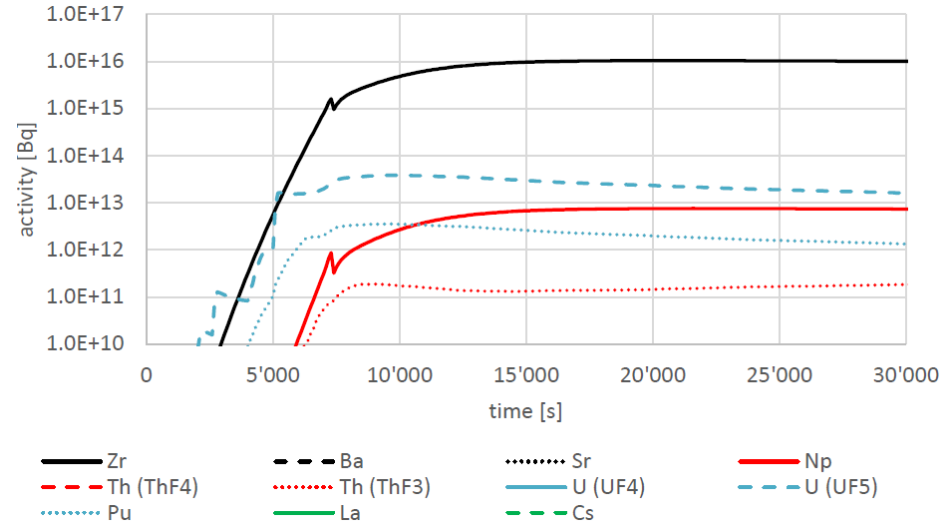
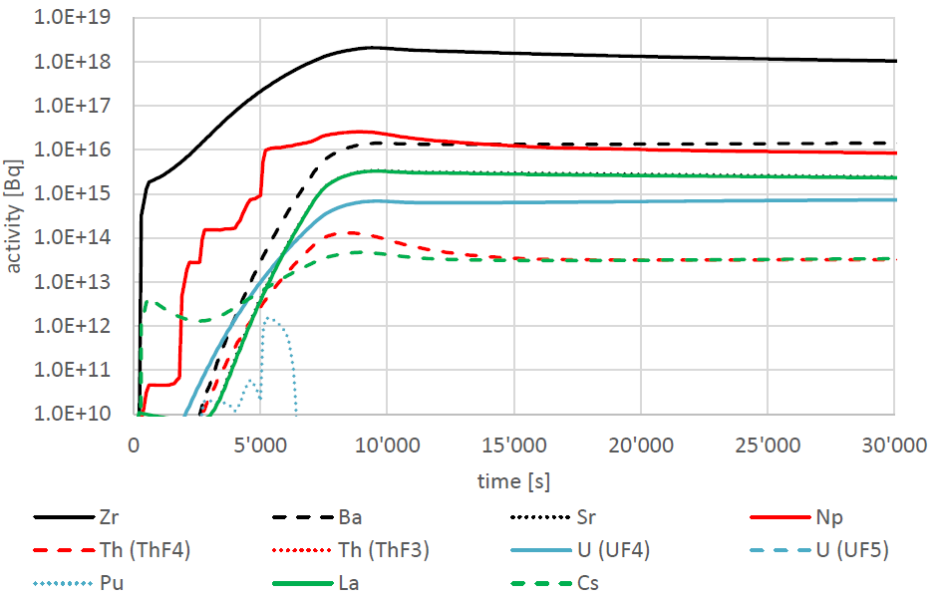
+1% mol.



— avg. gas T.

Total released mass of vapors during the accident (salt heat up from 800°C to 1500°C)

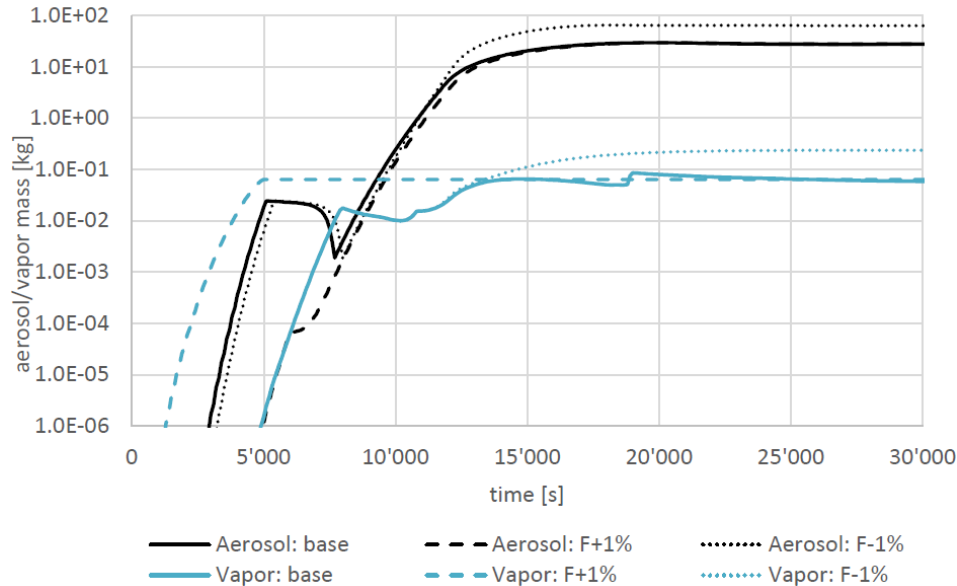
- Characterization of released activity in form of aerosols and vapors



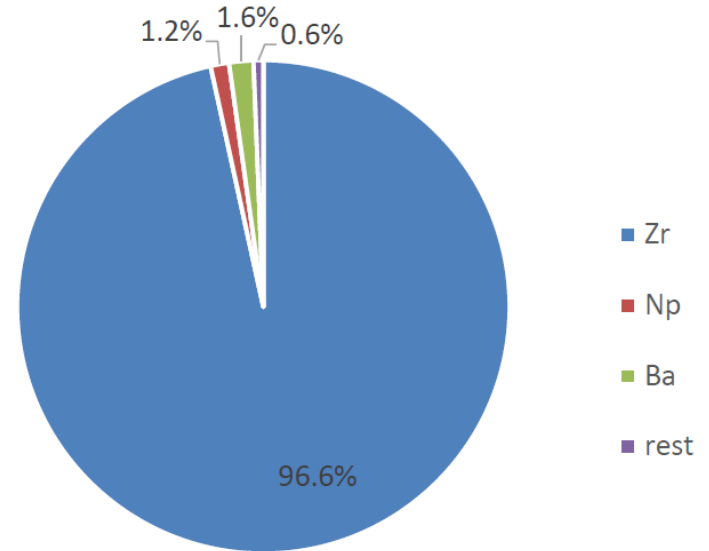
Total released activity in form of aerosols (left) and vapors (right) during the accident (salt heat up from 800°C to 1500°C)

ZrF₄ as the major component?

- Based on the applied reprocessing scheme, ZrF₄ in form of aerosols seems to be the major activity carrier during the postulated accident.



Total released activity in form of aerosols and vapors during the accident (salt heat up from 800°C to 1500°C)



Activity break-down at the end of simulation (t=30'000s) of the accident (salt heat up from 800°C to 1500°C)

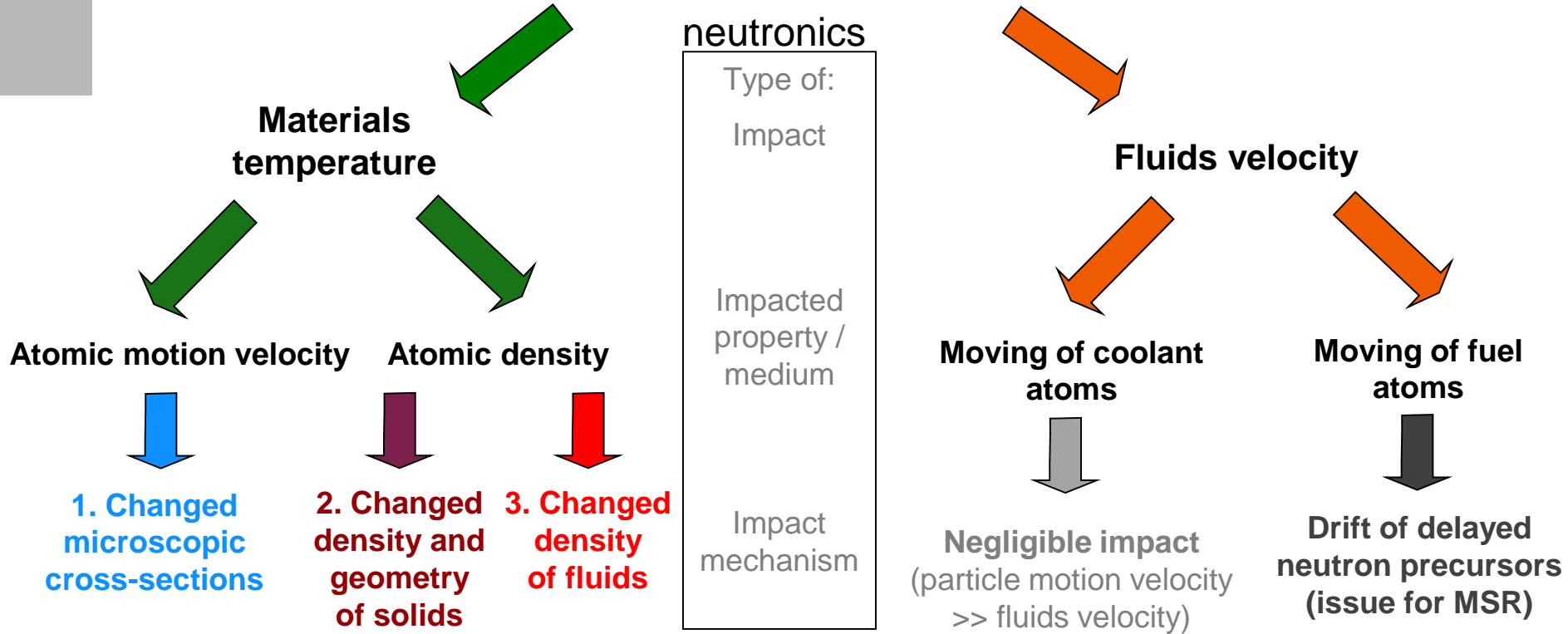
Summary – source term

- Localization and mobility of source term in MSR strongly depends on the in-situ salt treatment.
- Off-gas system may represent its major component.
- Source term released from the salt after vessel damage is determined by the salt treatment during nominal operation.
- In presented study ZrF_4 was not removed from the fuel salt during nominal operation, hence it dominates in the source term.
- The distributed source term might increase both capital and operation costs and bigger units might be more profitable.

Safety considerations: criticality safety

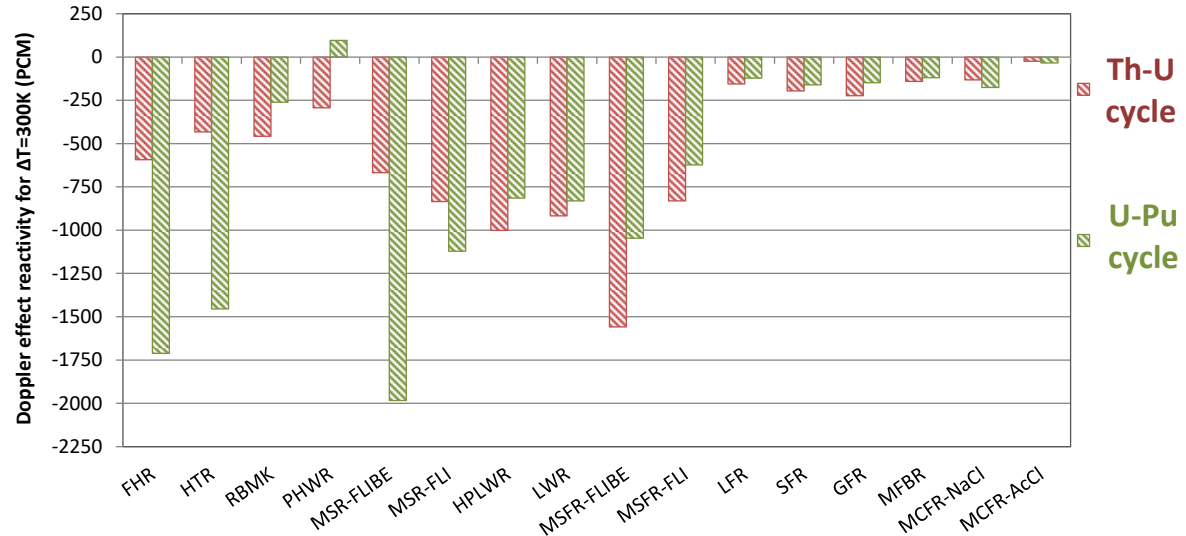
Criticality safety: thermal feedback coefficients

Thermal-hydraulics
interaction with
neutronics



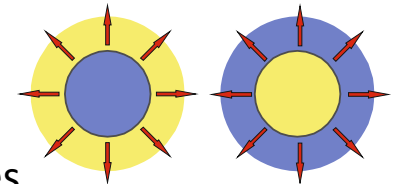
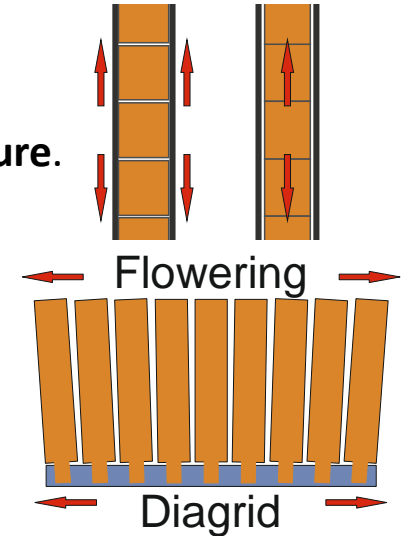
1. Changed microscopic Cross-sections (XS)

- **Doppler broadening of XS** result in **negative reactivity** practically in all reactor with substantial share of heavy nuclei with capture XS resonances (^{238}U and ^{232}Th).
- As an example reactivity introduced by **300K** fuel temperature increase is shown.
- *The figure shows results for iso-breeding fuel composition. It is not a value for standard fuel composition.*
- *No time to discuss PHWR. Have a look on the refered paper.*



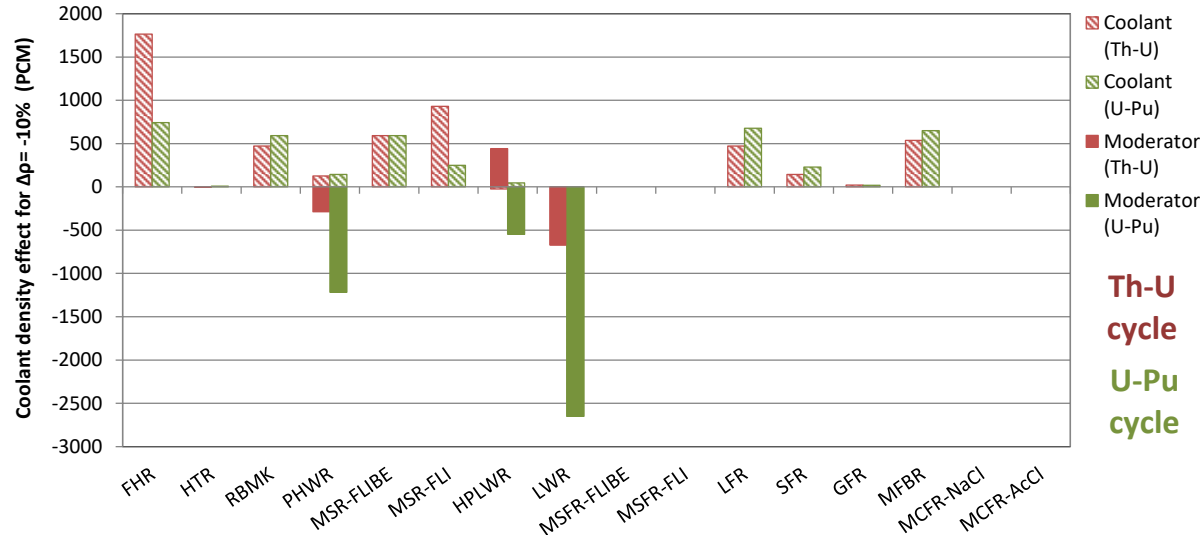
2. Changed density and geometry of solids

- **Temperature increase** of solid materials results in their **thermal expansion**.
- **Active core** is delimited by **fuel presence**.
- Active core **axial** expansion is driven by **fuel or cladding temperature**.
(cladding temperature drives fuel position in case of closed gap between them)
- Active core **radial** expansion is driven by **diagrid expansion** and assemblies **flowering effect**.
- **Non-fuel solid materials** can expand:
 - **outwards** from the active core, e.g., **cladding** with open gap
 - **inwards** to the active core, e.g., control rods and **control rods drivers expansion**
- Expansion of **solid materials can expel fluids** from the core.
 - e. g., cladding radial expansion expels sodium from the core.
 - in rare cases it can be the opposite, e. g., fluid in expanding tubes.



3. Changed density of fluids

- **Decreased density** result typically in **positive reactivity**.
Only for moderating fluid it is negative (reduced scattering XS dominate).
- As an example **10% density reduction** is shown. (*infinite lattice simulation = 0 for hom. reactors*)
- *The figure shows results for iso-breeding fuel composition. It is not a value for standard fuel composition.*
- *In MSR-FLI and MSR-FLIBE two effects take a part: reduced salt capture and changed fuel-to-moderator ratio, Doppler effect is not included.*



4. Fuel circulation

- In many MSR concepts fuel circulate through the reactor and primary circuit.
- Obviously, actinides and fission product circulate too.
- As a consequence, decay heat and delayed neutrons release is distributed.
- Using simple 1D approximation, the equation for precursors writes as:

$$\frac{\partial}{\partial t} C(z, t) + \frac{\partial}{\partial z} (vC(z, t)) = Q(z, t) - \lambda C(z, t)$$

~~$$\frac{dC(t)}{dt} = Q(t) - \lambda C(t)$$~~

- Assuming constant velocity v and characteristic line $z=vt$ we can integrate it:

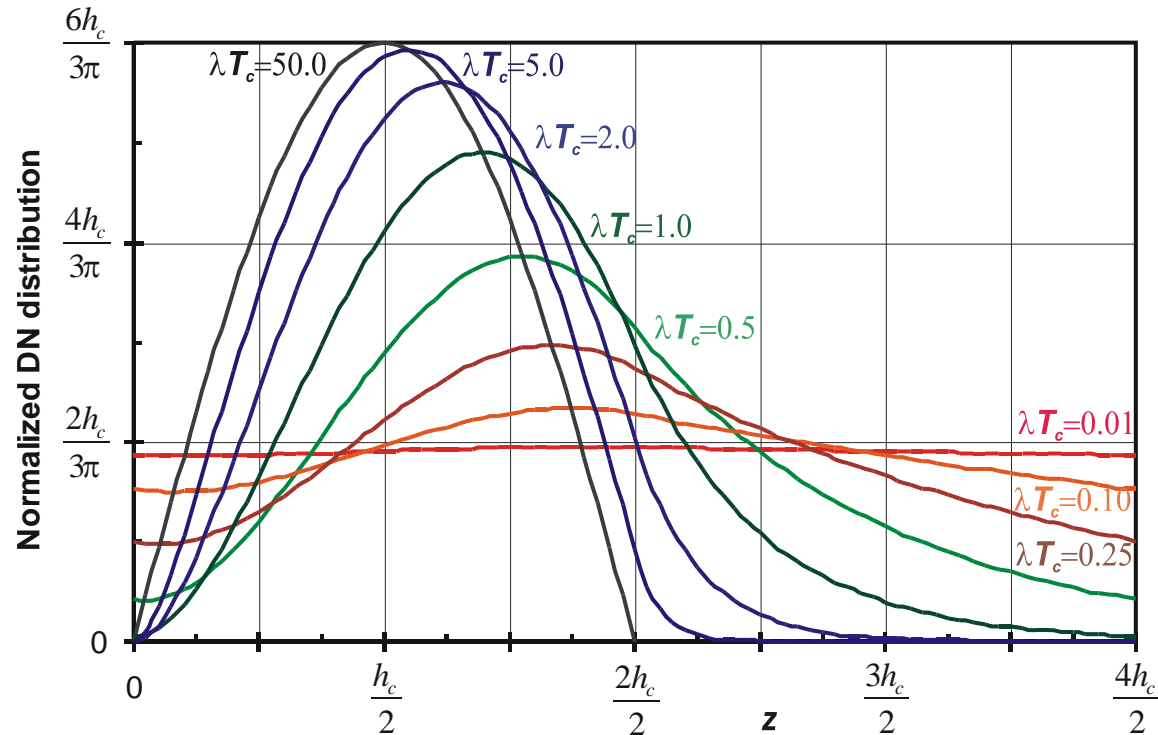
$$C(z, z/v) = e^{-\frac{\lambda}{v}(z-z_0)} \left(C(z_0, z_0/v) + \int_{z_0}^z \frac{Q(z', z'/v)}{v} e^{\frac{\lambda}{v}(z'-z_0)} dz' \right)$$

- Lets assume also sinus shape of the neutron flux:

$$Q(z) = \frac{\pi}{2h_c} \sin(z\pi/h_c) \text{ for } z = (0, h_c)$$

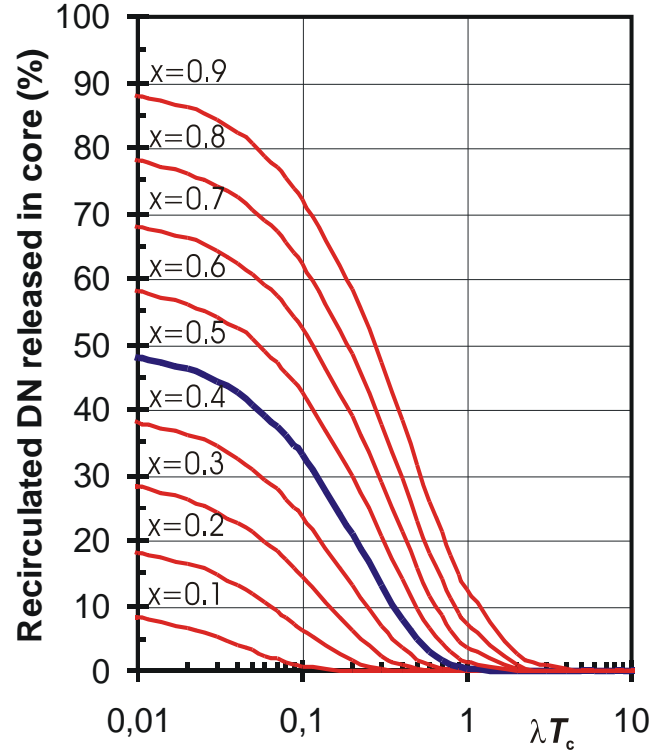
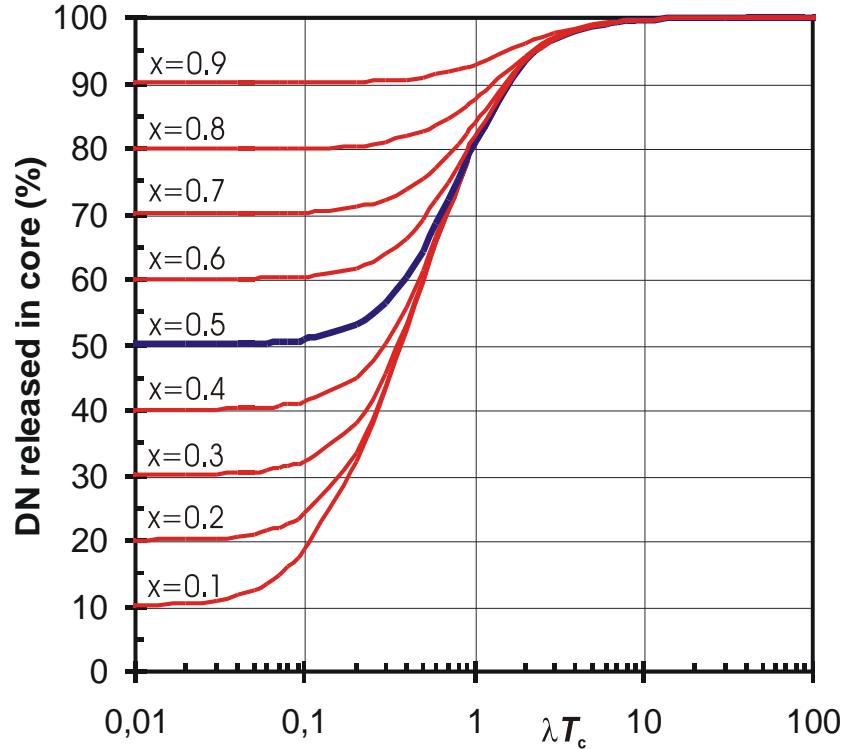
4. Fuel circulation

- Distribution of delayed neutrons.



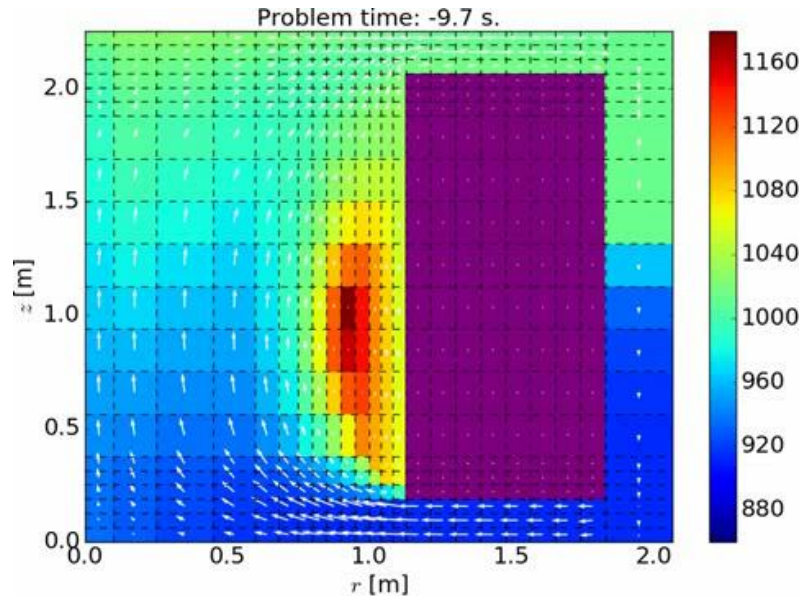
4. Fuel circulation

- Share of delayed neutrons released in the core.



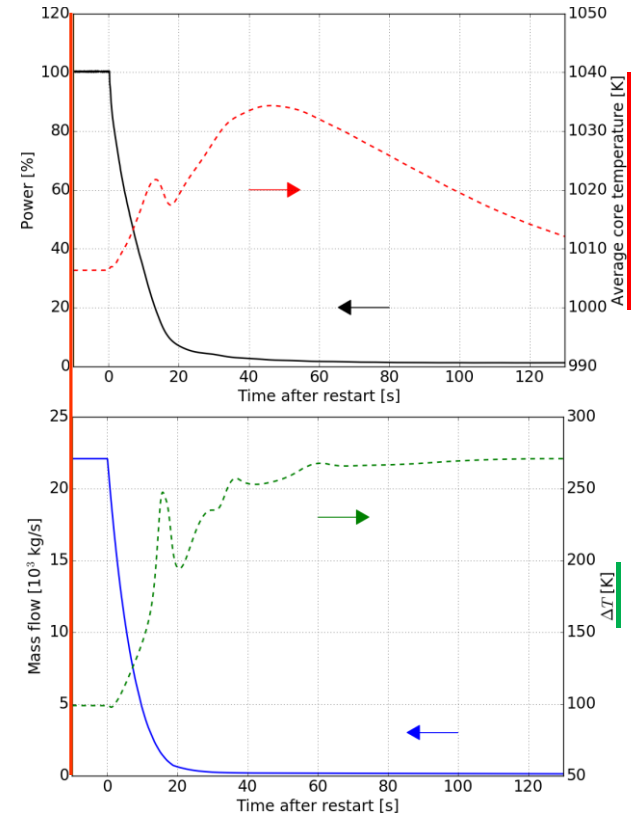
TRACE application to MSFR

- **TRACE-PARCS** system code application to **MSFR**.
- Applying TRACE vessel component.
- Applicable for 3D transients (2D symmetric transients).
- Capability of system analysis with acceptable accuracy and CPU time.



ULOF- unprotected loss of flow

E. Pettersen, LRS, MSc thesis 2016



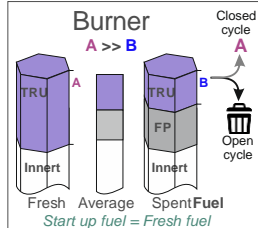
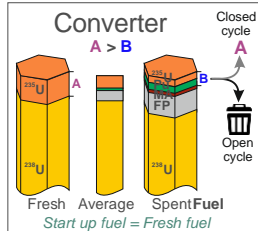
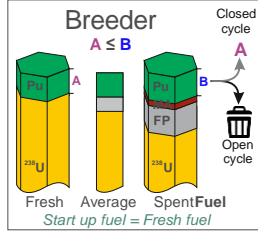
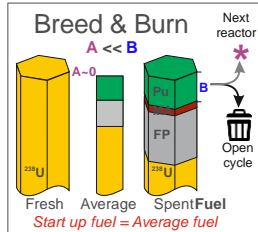
Summary – criticality safety

- MSR can be designed with negative Doppler effect.
- In case of waste burner fueled by minor actinides, the weaker or even positive, Doppler effect can be compensated by negative fuel salt expansion coefficient.
- MSR can be designed as a breeder in Th-U or U-Pu cycle with all thermal feedback coefficients being negative.
- Small size of the core or low power is not necessary for that.
- Fuel circulation redistributes delay neutron and decreases their weight.
- The only sizing consideration related to safety is decay heat removal.

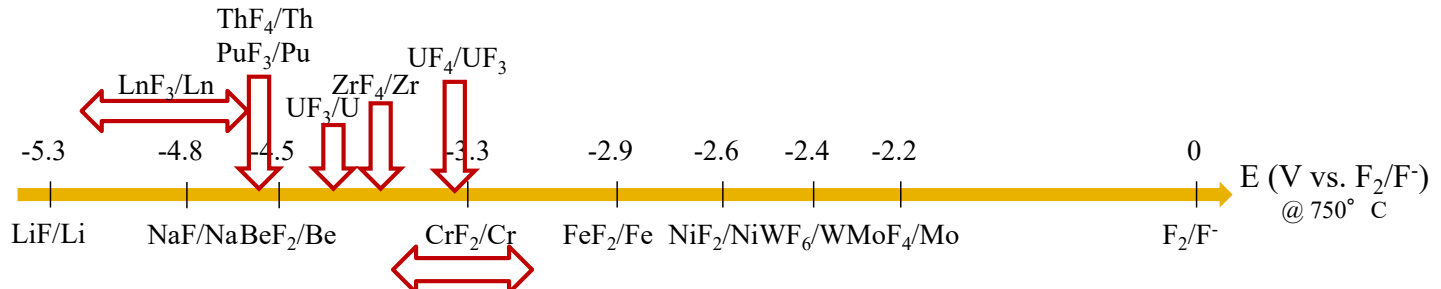
Fuel cycle considerations

MSR fuel cycle choices and options

Neutron economy



- The carrier salt (liquid fuel solvent) is an ionic liquid and can be irradiated without a limit.
- No matter the fuel cycle, it is always good to keep actinides as long as possible in the reactor (high burnup).
- At best only Fission Products (FPs) should leave the core.
- Unfortunately, FPs (Lanthanides) tends to leave the fuel salt as last.

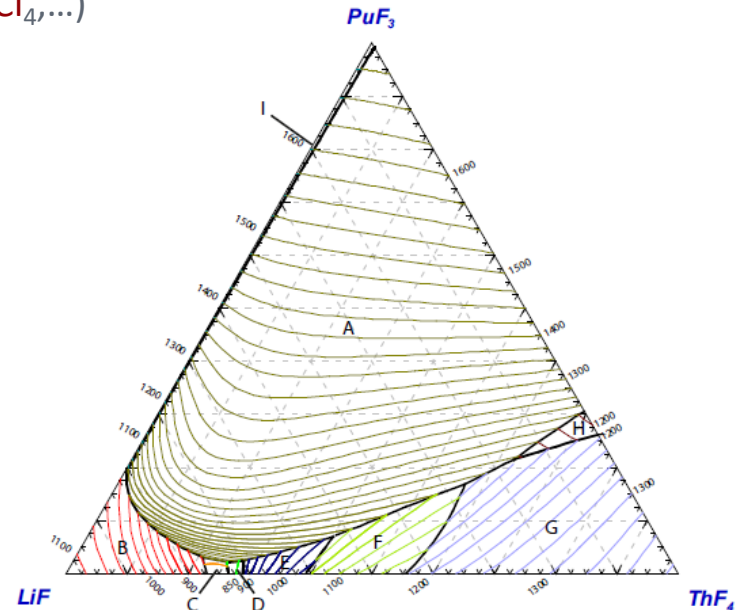


Relative stability of redox couples, Gibbs energy : $\Delta E \text{ (V)} = -\Delta_r G \text{ (J/mol)} / nF$, $E^\circ(\text{F}_2/\text{F}^-) = 0 \text{ V}$

Adopted from: Laurent Cassayre, LGC.CNRS.FR, Corrosion in molten salt reactors, EVOL Winter School, 4-6 November 2013, IPNO, France

Reasonable temperature window for operation: interval between salt melting and structural materials failure

- Typically eutectic mixture of carrier salts (LiF, BeF₂, NaF, LiCl, NaCl,...) and actinides salts (ThF₄, UF₄, PuF₃, PuCl₃, UCl₃, ThCl₄,...)
- MSRE salt, **T_{melt.} = 432 °C**
65% LiF - 29.1% BeF₂ - 5% ZrF₄ - 0.9% UF₄
- MSBR, Th-U equilibrium cycle, **T_{melt.} = 500 °C**
71.7% LiF - 16% BeF₂ - 12% ThF₄ - 0.3% UF₄
- MSFR, Th-U equilibrium cycle, **T_{melt.} = 560 °C**
78% LiF - 17.6% ThF₄ - 4% UF₄ - 0.2% PuF₃
- MSFR, Pu started Th-U cycle, **T_{melt.} = 625 °C**
78% LiF - 16% ThF₄ - 6% PuF₃
- MCFR, Pu started U-Pu cycle, **T_{melt.} = 565 °C**
60% NaCl - 35% UCl₃ - 5% PuCl₃
- MCFR, Pu started Th-U cycle, **T_{melt.} = 425 °C**
55% NaCl - 39% ThCl₄ - 6% PuCl₃
- Generally **solubility limits** (e.g. PuF₃) and **actinides density** compete with **melting temperature**.



LiF-ThF₄-PuF₃ ternary phase diagram
w/ fixed 1% mol UF₄ concentration

E. CAPELLI et al., "Thermodynamic Assessment of the
LiF-ThF₄-PuF₃-UF₄ System," *J. Nucl. Mater.*, **462**, 43 (2015).

Reprocessing as “à la carte “ choice

Fuel salts components:

1. Carrier salt (LiF, NaCl,...)
2. Fertile actinides (^{232}Th and ^{238}U).
3. Fissile actinides (^{233}U and ^{239}Pu).
4. Minor actinides (MA).
5. FPs.

Reprocessing strategies:

Salt treatment / reprocessing techniques:

- Gaseous and volatile FPs removal (off-gas system).
- Metallic FPs removal (sponge filter or by off-gas sys.).
- Molten salt / liquid metal reductive extraction.
- Electro-separation processes.
- Compound evaporation or possibly precipitation.
- Fluoride volatilization techniques, fluorination of the molten salt mixture.

Salt removal from the core	Removed salt share	Fissile fuel recycling	Fissile fuel return after reprocessing	Carrier salt cleaning	Carrier salt return after reprocessing	Reprocessing waste immobilization
Continuous or Batch-wise	From 0.1% to whole salt volume	In-situ or Ex-situ	ASAP or with months or years of delay	In-situ or Ex-situ	ASAP or with months or years of delay	In-situ or Ex-situ

Fuel cycle assessment from neutronics perspective

5 major fuel cycle types (*reactor physics perspective*)

- I. Enriched uranium burning** (^{235}U burning) **Resources utilization: <1% of nat. U**
- Enrichment level can range from 0.7% to 20%.
 - The cycle is generally open and “waste” intensive.
 - However, irradiated U and generated Pu can be recycled as MOX fuel.
- II. Closed Th-U cycle** (^{232}Th burning in closed cycle) **<95% of ^{232}Th**
- Actinides recycling in a breeder reactor fueled by ^{232}Th .
- III. Closed U-Pu cycle** (^{238}U burning in closed cycle) **<95% of nat. U**
- Actinides recycling in a breeder reactor fueled by ^{238}U .
- IV. Breed-and-burn U-Pu cycle** (^{238}U burning in open cycle) **cca 20-30% nat. U**
- Open cycle, Ac. are not recycled, but the reactor acts as a breeder.
 - “Waste” intensive cycle, however fuel can be reused by another reactor.
- V. Synthetic actinides burning** **not relevant**
- Cycle dedicated to minimization of existing synthetic actinides.
- **Combination or transition between above cycles**
 - e.g. actinides from I. or V. can acts as an initial or add on fuel for II. - IV. or vice versa.

5 fuel cycle performance parameters

I. Breeding capability

- How many neutrons can be captured by ^{232}Th or ^{238}U so that the reactor is still critical.
- BTW: Uranium enrichment reduces ^{238}U capture, hence also the breeding capability.
- It is about neutron economy.

II. Achievable burnup

- Is limited by Fission Products (FPs) neutron capture and by fuel irradiation stability.
- Depends on initial reserve of fissile material and its renewal (breeding capability).

III. Initial fissile mass

- It is determined by neutron economy and spectrum type of the reactor.
- Higher burnup may impose higher initial fissile mass reserve.

IV. Means of criticality maintenance

- Ac. irradiation and FPs creation results in reactivity oscillations / swing.
- Compensation option for reactivity swing differ between reactor types.

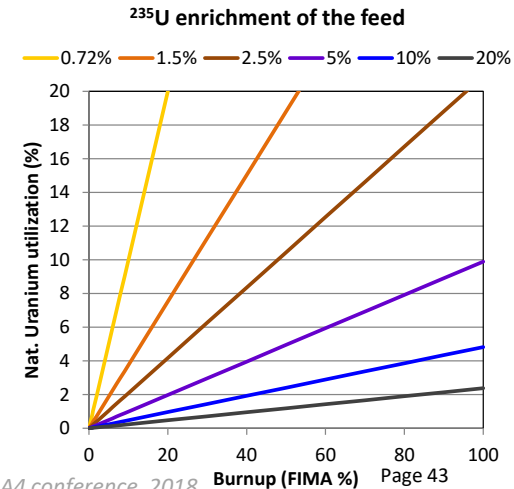
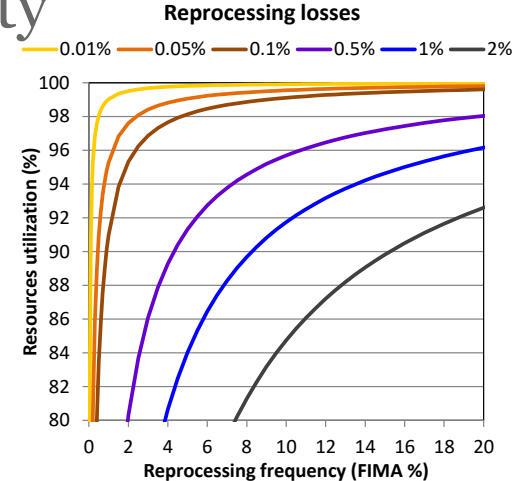
V. Transmutation capability

- “Neutron costs” and “speed” of synthetic actinides fission.
- Synthetic Ac. compatibility with the fuel and fabrication process.

Ac. recycling vs. breeding capability

- Fuel cycle can be closed for: **Burner, Convertor, and Breeder.**
- Recycling does **not make sense** for **B&B** reactor.
However, the fuel can be recycled and used in other reactor.
- **Recycling** is the ultimate **waste reducing** option.
- **Recycling in a breeder** => **highest resources utilization.**
It is limited by reprocessing losses.
- **Recycling in a convertor** => **medium resources utilization.**
It is limited by enrichment process and reprocessing losses.
- **Recycling in a burner** => mainly **waste minimization.**
- **Fuel recycling / long irradiation** leads to:

equilibrium fuel composition.



Equilibrium fuel composition

- When fuel cycle **parameters**: power, reprocessing scheme, feed composition, etc. are **fixed**, reactor will converge to equilibrium state / **fuel composition**.

- The composition depends on **feed** type ^{238}U or ^{232}Th

- and on the reactor **spectrum**.

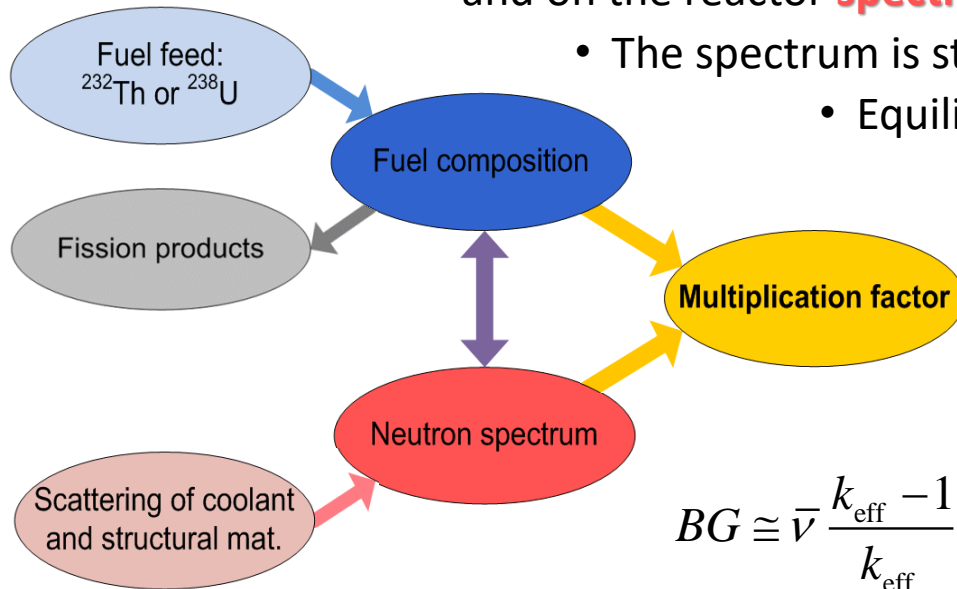
- The spectrum is strongly determined by **scattering materials**.

- Equilibrium composition and spectrum determine the **multiplication factor**.

- Equilibrium is **inherent core state**.
(Bateman's matrix eigenstate:
composition, spectrum, reactivity)

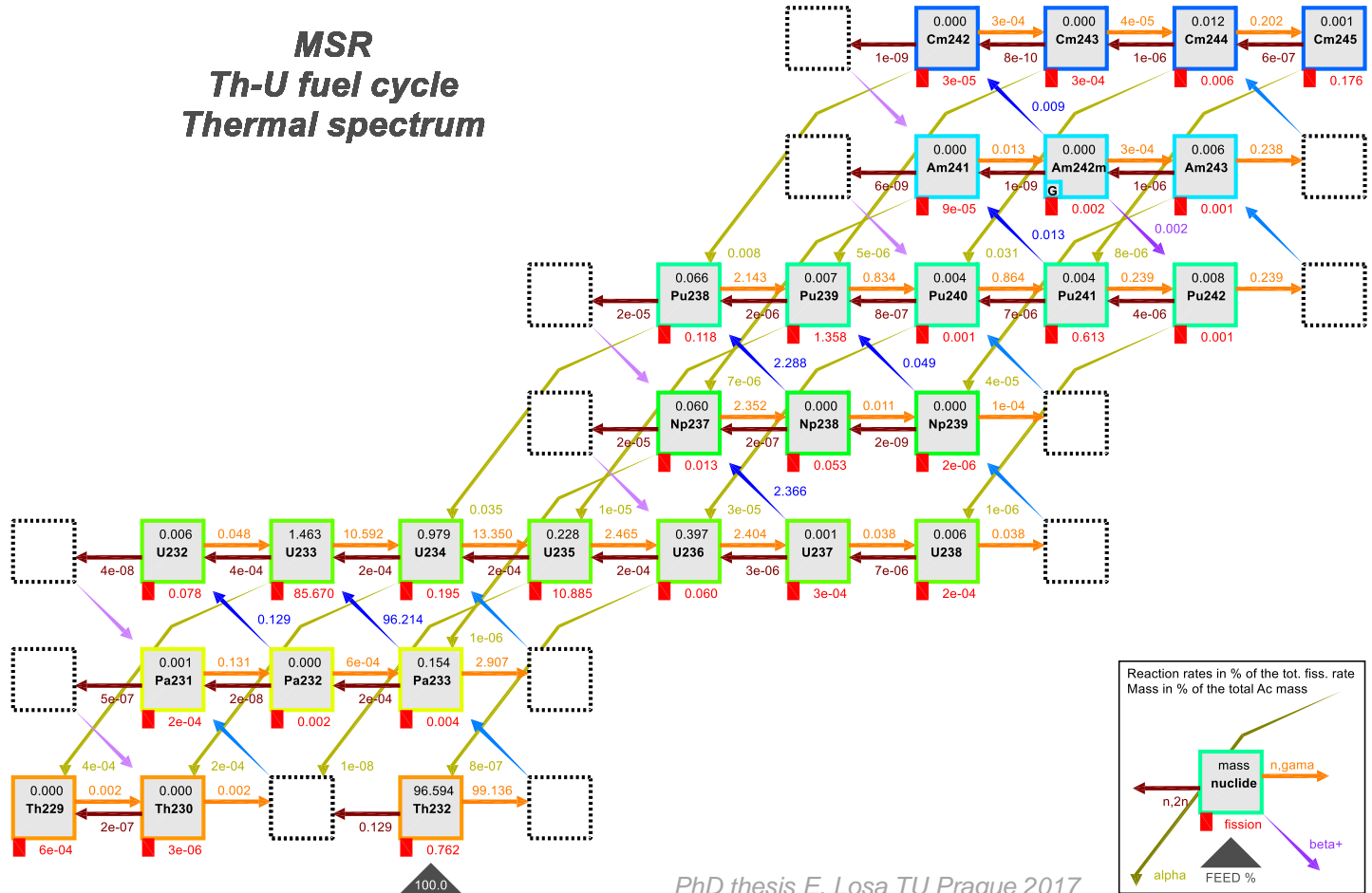
- Equilibrium **reactivity** indicates neutron **efficiency** of the core and its **capability for breeding**.

$$BG \cong \bar{\nu} \frac{k_{\text{eff}} - 1}{k_{\text{eff}}}$$



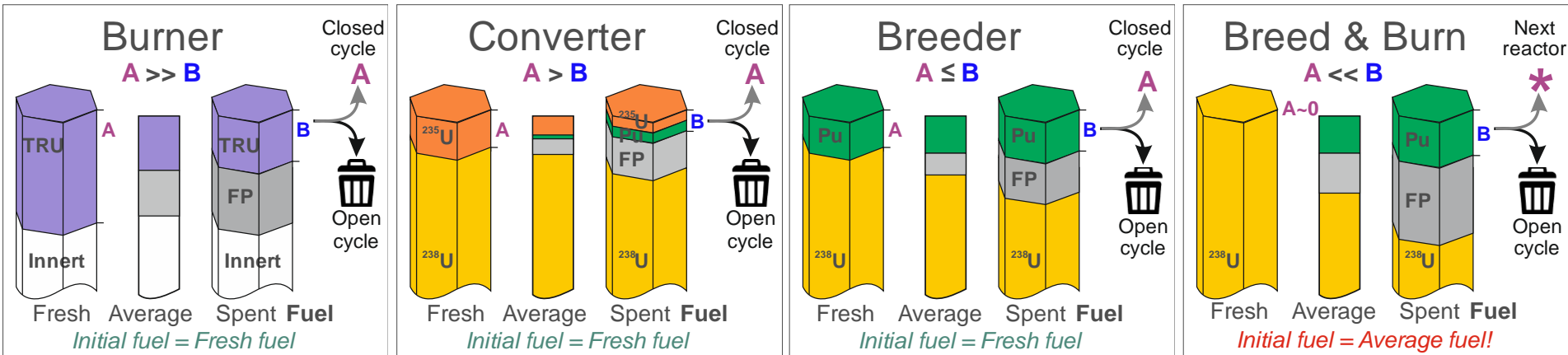
Equilibrium fuel composition (= irradiation chain)

MSR
Th-U fuel cycle
Thermal spectrum



Reactor classification by breeding capability

Neutron economy



- Burner is typically used for synthetic Ac and excludes fertile isotopes as ^{238}U or ^{232}Th .
- Converter, e. g. PWR or DMSR, is usually operates in open fuel cycle and burns ^{235}U .
- Breeder profit from neutronics advantages only in the closed cycle. For Iso-breeding (EU) or Break-even (US) reactor $\text{Pu} \Rightarrow A=B$.
- Extreme breeder can be operated in Breed & Burn mode. It can have high fuel utilization even without reprocessing.

Breed & Burn (B&B): special open cycle mode

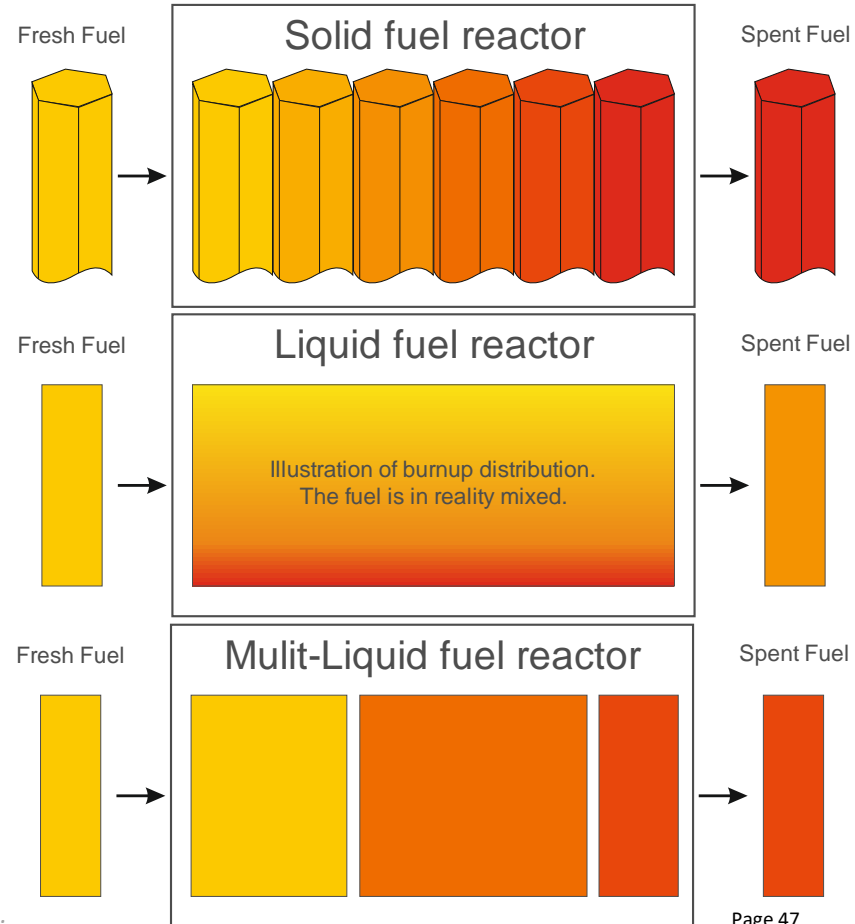
- Initially the fissile fuel will be **bred** and firstly later it will be **burned**.
- The B&B cycle in **liquid fuel** reactor substantially differs from **solid fuel**.

Discharged fuel:

Most burned in solid fuel case

Average burned in liquid fuel case.

- To increase the burnup and reduce the core size (**single-fluid** layout can be bulky), **multi-fluid** layout can be used.



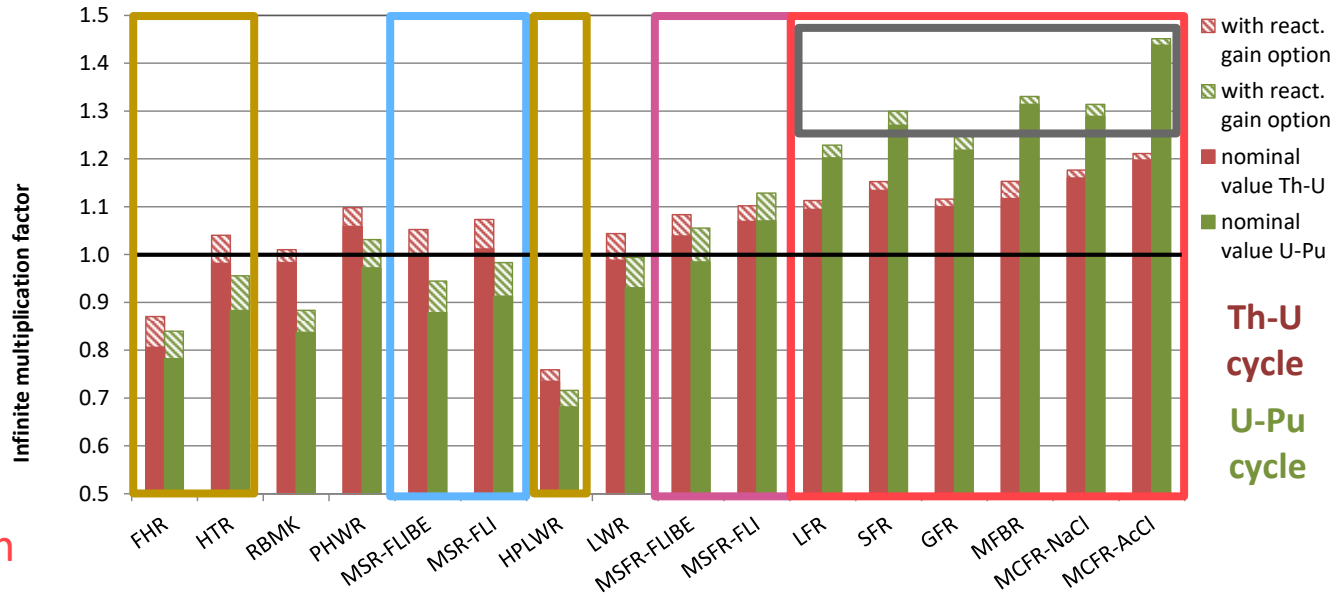
Equilibrium cycle comparison for 16 reactors

- **Assumptions:**
 - 1) infinite lattice.
 - 2) neglected fission products.
 - 3) design as is, no additional optimization.
 - 4) ENDF/B-VII.0
- 8 thermal and
- 8 fast reactors were compared in both U-Pu and Th-U cycles.

<i>Solid fuel thermal reactors</i>				<i>Solid fuel fast reactors</i>			
Reactor name (and label)	Lattice geometry	Name and short name	Lattice geometry	Reactor name (and label)	Lattice geometry	Name and short name	Lattice geometry
Fluoride high temperature reactor PB-AHTR (FHR)		Pressurized heavy water reactor (PHWR)		European lead system (LFR)		Gas cooled fast reactor (GFR)	
High temperature reactor (HTR)		High performance light water reactor (HPLWR)		European sodium fast reactor (SFR)		Metal fueled fast breeder reactor (MFBR)	
Reaktor balshoi moshnosti kanalnyj (RBMK)		Light water reactor VVER-1000 (LWR)		<i>Liquid fuel fast reactors</i>			
				Molten salt fast reactor fueled by LiF-BeF ₂ -AcF ₄ (MSFR-FLIBE)		Molten salt fast reactor fueled by NaCl-AcCl ₄ (MCFR-NaCl)	
<i>Liquid fuel thermal reactors</i>				Molten salt fast reactor fueled by LiF-AcF ₄ (MSFR-FLI)		Molten salt fast reactor fueled by AcCl ₄ (MCFR-AcCl)	
Thermal MSR fueled by LiF-BeF ₂ -AcF ₄ (MSR-FLIBE)		Thermal MSR fueled by LiF-AcF ₄ (MSR-FLI)					

Equilibrium multiplication factor (reactivity)

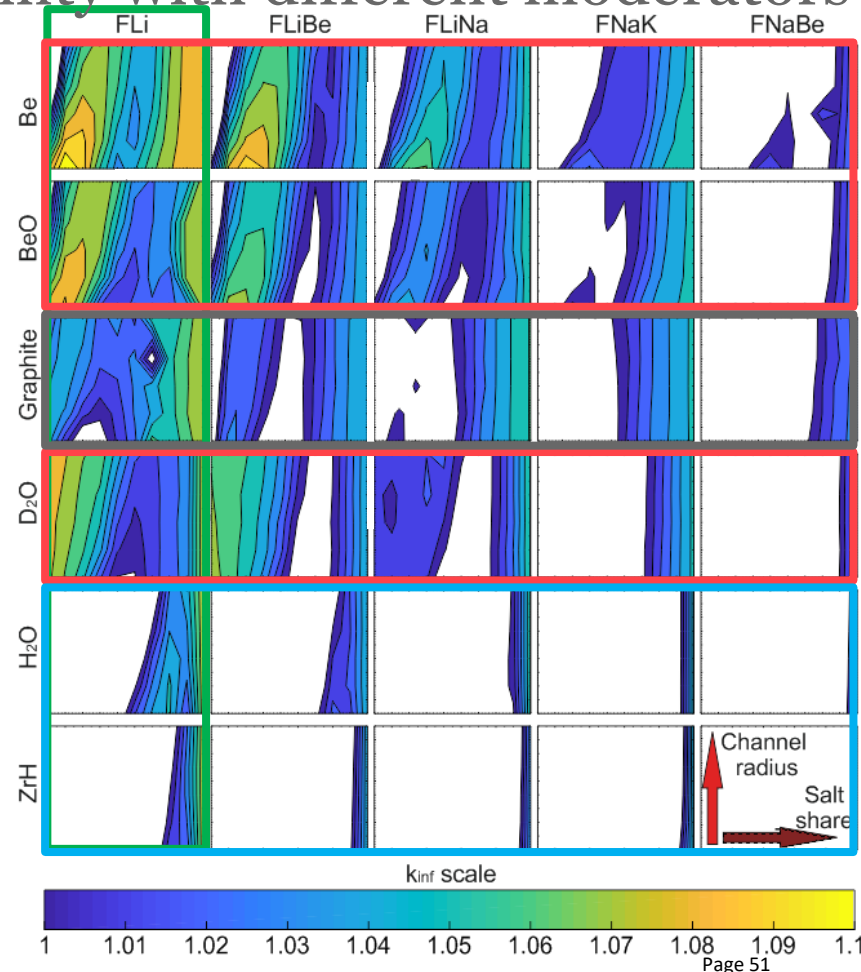
- Better performance: **Th-U in thermal and U-Pu in fast spectra.**
- Graphite moderated MSR only in Th-U.
- Fluorides fast MSFR possible in both cycles *(equal performance almost epithermal)*.
- GenIV fast reactors can all breed in closed cycle.
- B&B possible for U-Pu.
- SCWR, FHR, HTR???



Breeding capability of moderated MSRs

Th-U breeding capability with different moderators

- **5 fluoride salts** were analyzed with **6 selected moderators**.
- Equilibrium k_{inf} is presented as a function of **salt share** and **channel radius**.
- **FLi** salt is neutronically the best.
- Good results for **Be, BeO, and D₂O**; however, they are not compatible with the salt without cladding (SiC..?).
- Hydrogen based moderators **ZrH and H₂O** not applicable for closed cycle.
- **Graphite** is not the best moderator, but the only one directly compatible with salt.

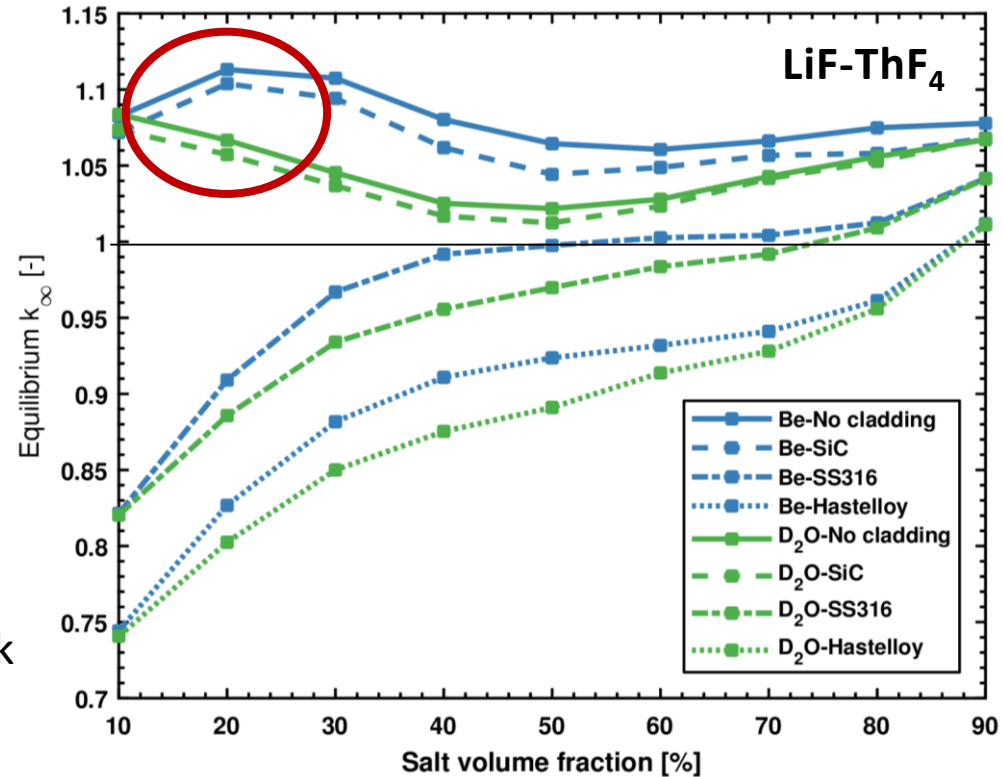


Neutronics impact of cladding (^7LiF salt example)

- **LiF salt** combined with **Be** and **D₂O** moderators was selected to analyze the impact of cladding:
Hastelloy,
SS316,
and SiC.
- Only **SiC** seems to have acceptable low parasitic neutron capture.
- Purely from neutronics perspective **Heavy Water Boiling MSR** would work

HWB-MSR 😊

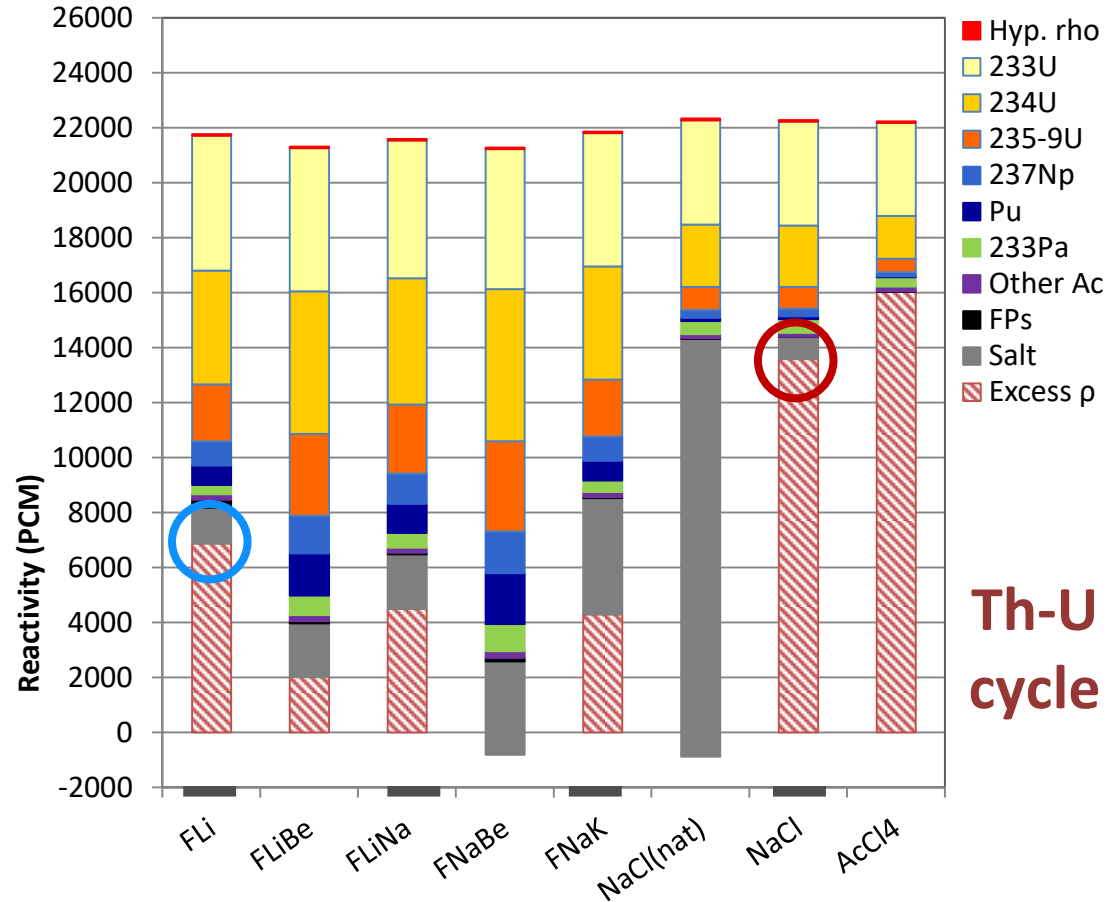
Thesis topic for super hero...?



Breeding capability of homogeneous fast MSR

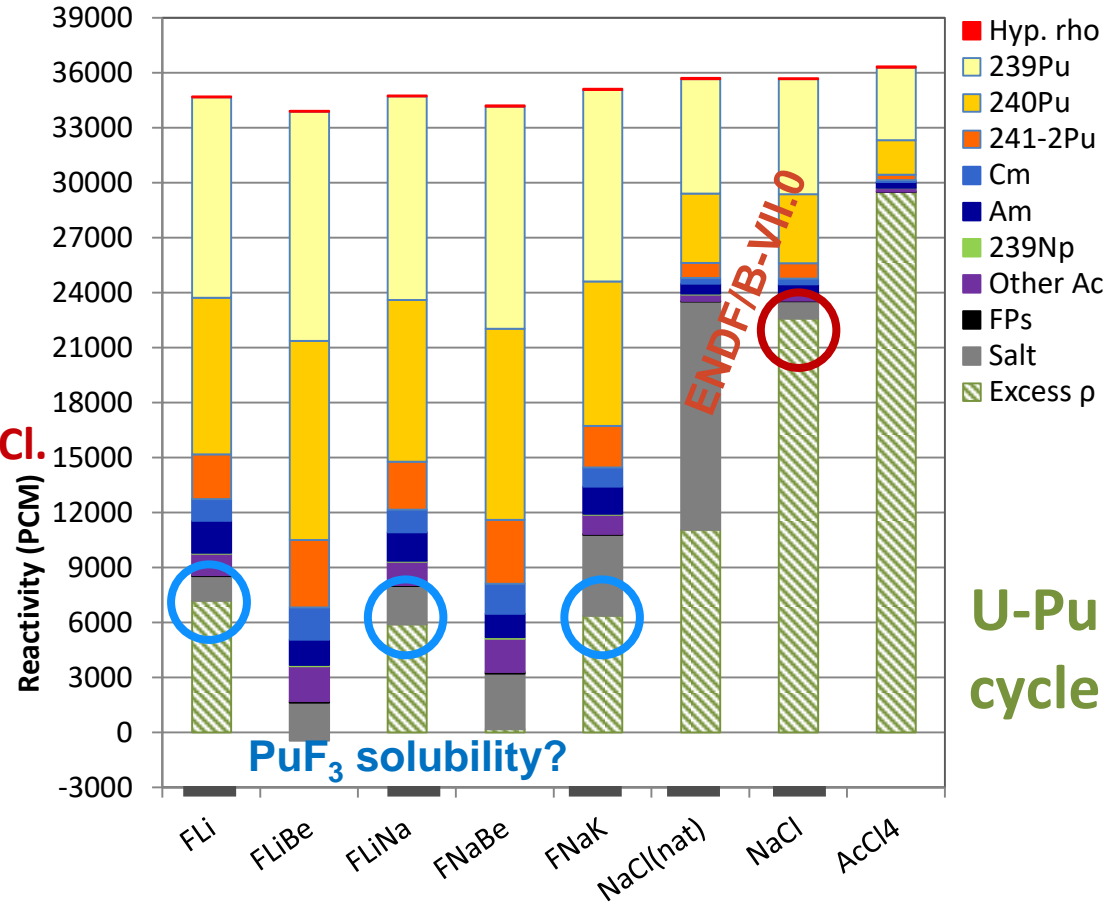
Th-U cycle performance without moderator

- **8 salts** were evaluated:
FLi, FLiBe, FLiNa, FNaBe, FNaK, NaCl (nat), Na³⁷Cl, Ac³⁷Cl.
32% AcCl₃ 32% AcCl₃ 100% AcCl₄
- **4 options in Th-U** (reasonable melting point and reactivity):
FLi, FLiNa, FNaK, Na³⁷Cl.
- **Na³⁷Cl** provides the highest excess in Th-U of 13000 pcm.
- **FLi** is best fluoride salt with 6000 pcm.



U-Pu cycle performance without moderator

- **8 salts** were evaluated:
FLi, FLiBe, FLiNa, FNaBe, FNaK, NaCl (nat), Na³⁷Cl, Ac³⁷Cl.
32% AcCl₃ 32% AcCl₃ 100% AcCl₃
- **5 options in U-Pu** (reasonable melting point and reactivity):
FLi, FLiNa, FNaK, NaCl (nat), Na³⁷Cl.
- **Na³⁷Cl** provides the highest overall excess of 22000 pcm.
- **FLi, FLiNa, FNaK** have similar performance of ~6000pcm, PuF₃ solubility is the major limiting issue.



Self-sustaining breeder core size estimate

Core radius estimate in Th-U cycle

- Bare core criticality line.

$$k_{inf} = 1 + M^2 B^2$$

Derived from Fermi theory of bare "thermal" reactor:

$$k_{eff} = k_{inf} P_1 P_2 = k_{inf} \frac{e^{-\tau B^2}}{1 + L^2 B^2} \cong k_{inf} \frac{1}{1 + (\tau + L^2) B^2} = k_{inf} \frac{1}{1 + M^2 B^2}$$

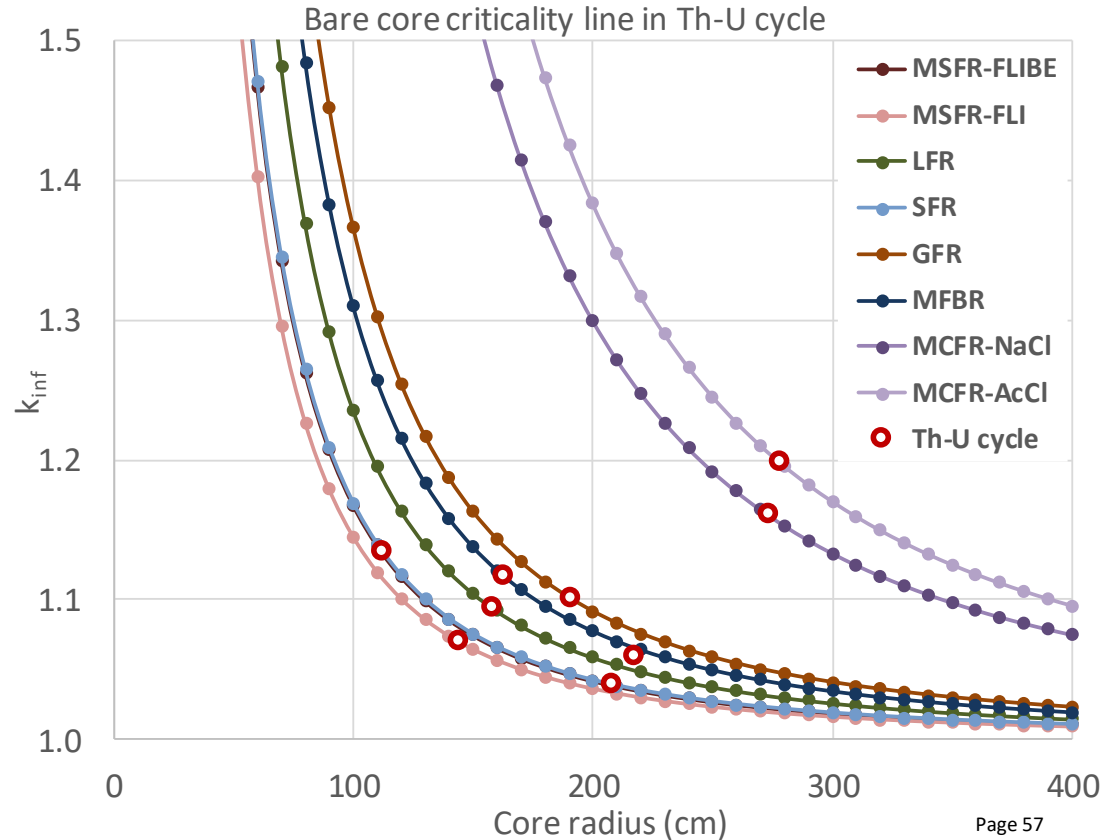
- Buckling for a cylinder:

$$B^2 = \left(\frac{\pi}{h} \right)^2 + \left(\frac{2.405}{r} \right)^2$$

- Minimal volume for given B^2 :

$$\frac{h}{r} = \frac{\sqrt{2}\pi}{2.405} \cong 1.85$$

- Core radius estimate in Th-U cycle =>**



Core radius estimate in U-Pu cycle

- Bare core criticality line.

$$k_{inf} = 1 + M^2 B^2$$

Derived from Fermi theory of bare "thermal" reactor:

$$k_{eff} = k_{inf} P_1 P_2 = k_{inf} \frac{e^{-\tau B^2}}{1 + L^2 B^2} \cong k_{inf} \frac{1}{1 + (\tau + L^2) B^2} = k_{inf} \frac{1}{1 + M^2 B^2}$$

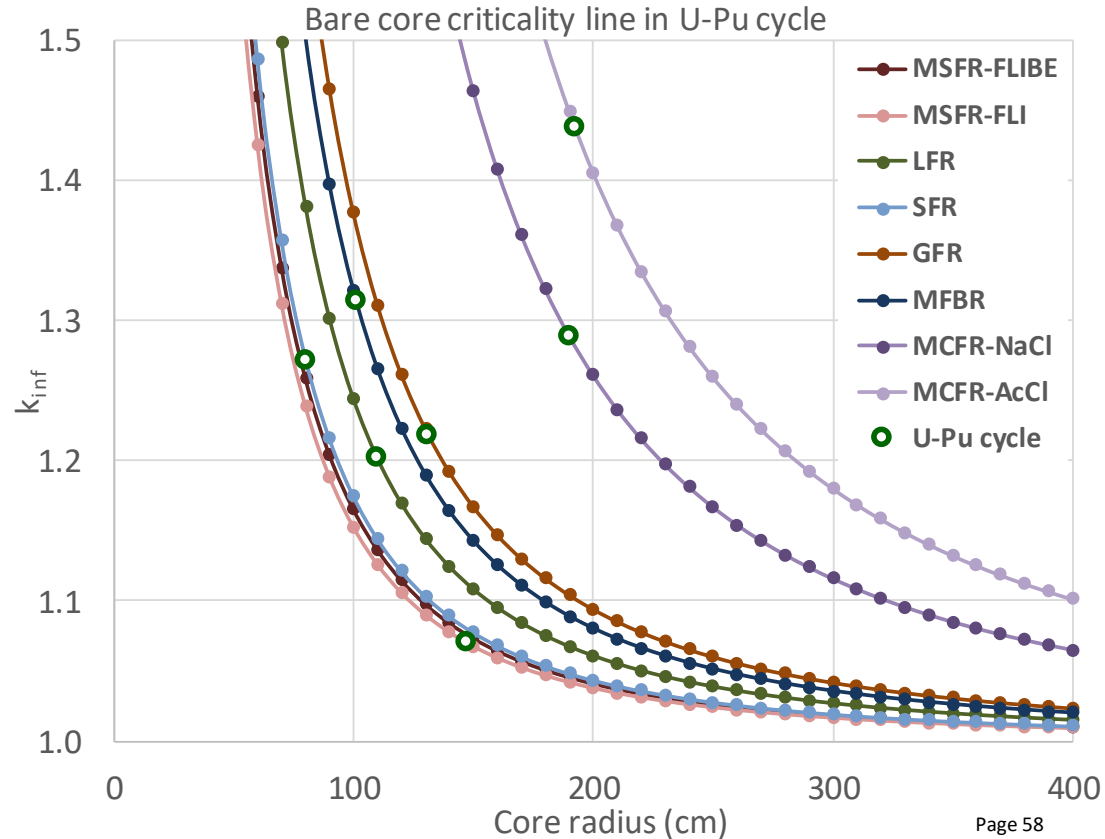
- Buckling for a cylinder:

$$B^2 = \left(\frac{\pi}{h} \right)^2 + \left(\frac{2.405}{r} \right)^2$$

- Minimal volume for given B^2 :

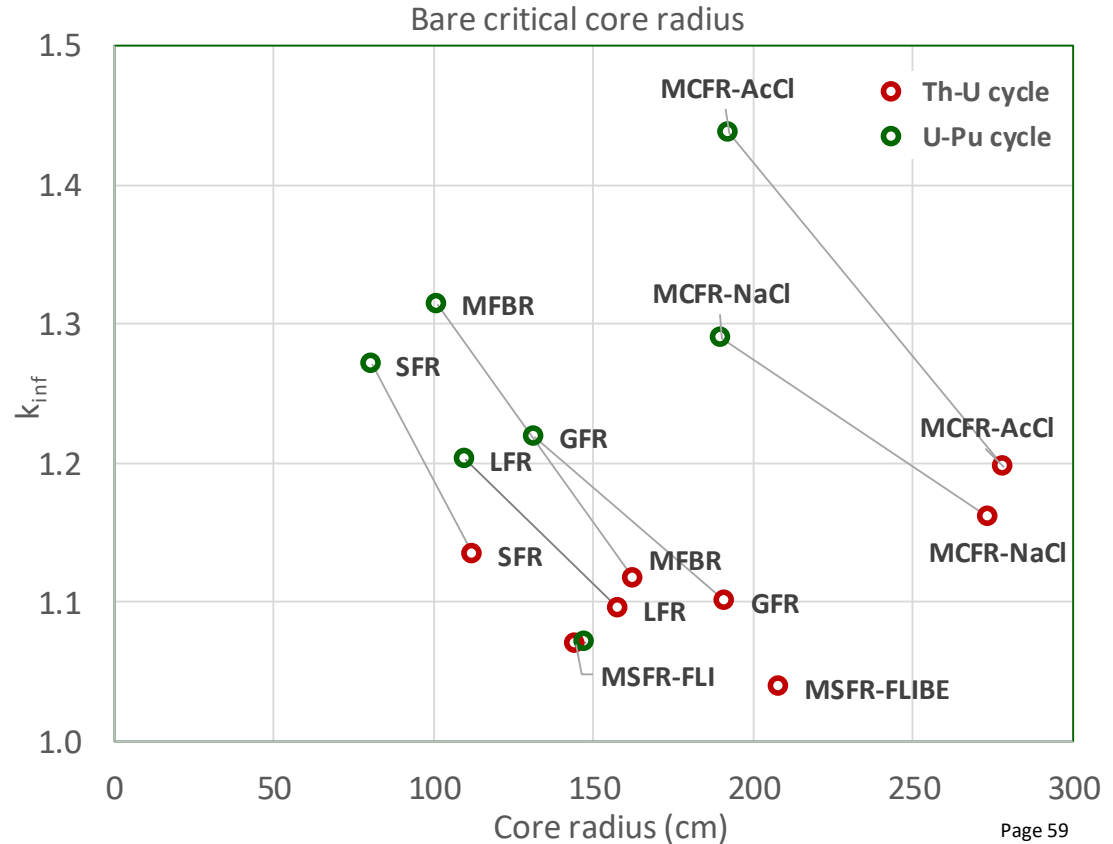
$$\frac{h}{r} = \frac{\sqrt{2}\pi}{2.405} \cong 1.85$$

- Core radius estimate in U-Pu cycle =>**



Core radius estimate: Th-U cycle X U-Pu cycle

- By all other fast reactors **U-Pu cycle provides smaller cores.**
- **SFR** is the **most compact** bare iso-breeding core in both cycles.
- **MCFR** is the **biggest** bare iso-breeding core in both cycles.
- **MSFR-FLI** is the **smallest MSR** core and it has the same core size for both cycles. (**very soft fast spectrum**)



Core radius estimate in Th-U cycle

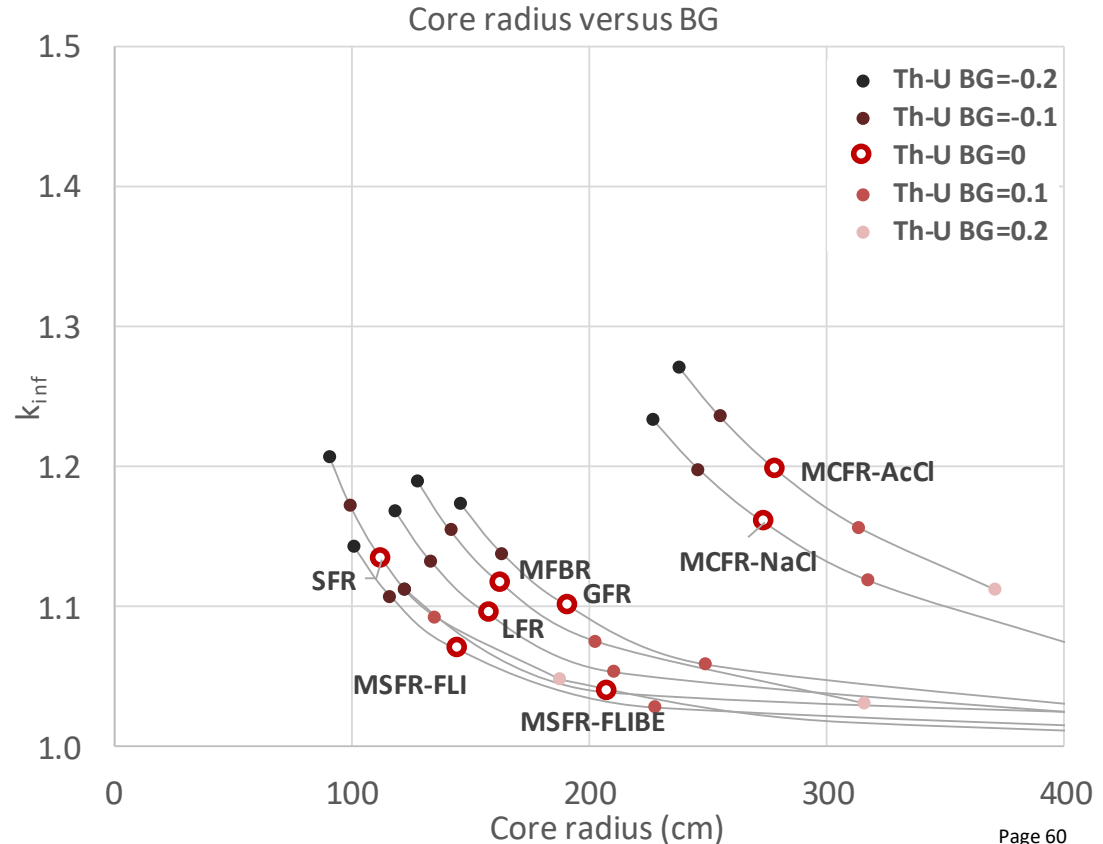
- Combining these two equations:

$$k_{eff} \cong k_{inf} \frac{1}{1 + M^2 B^2}$$

$$BG_{per} \cong \bar{v} \frac{k_{eff} - 1}{k_{eff}}$$

- Bare core size can be estimated for several BG values.

Th-U cycle =>



Core radius estimate in U-Pu cycle

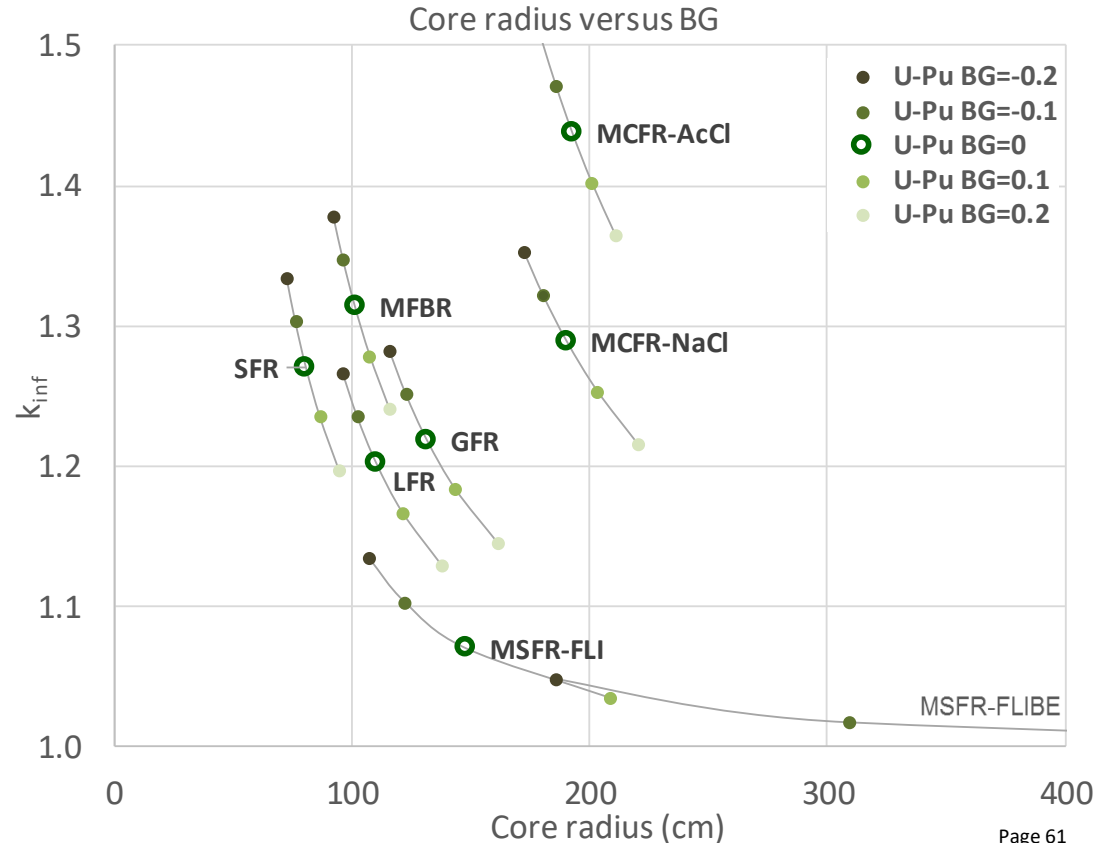
- Combining these two equations:

$$k_{eff} \cong k_{inf} \frac{1}{1 + M^2 B^2}$$

$$BG_{per} \cong \bar{v} \frac{k_{eff} - 1}{k_{eff}}$$

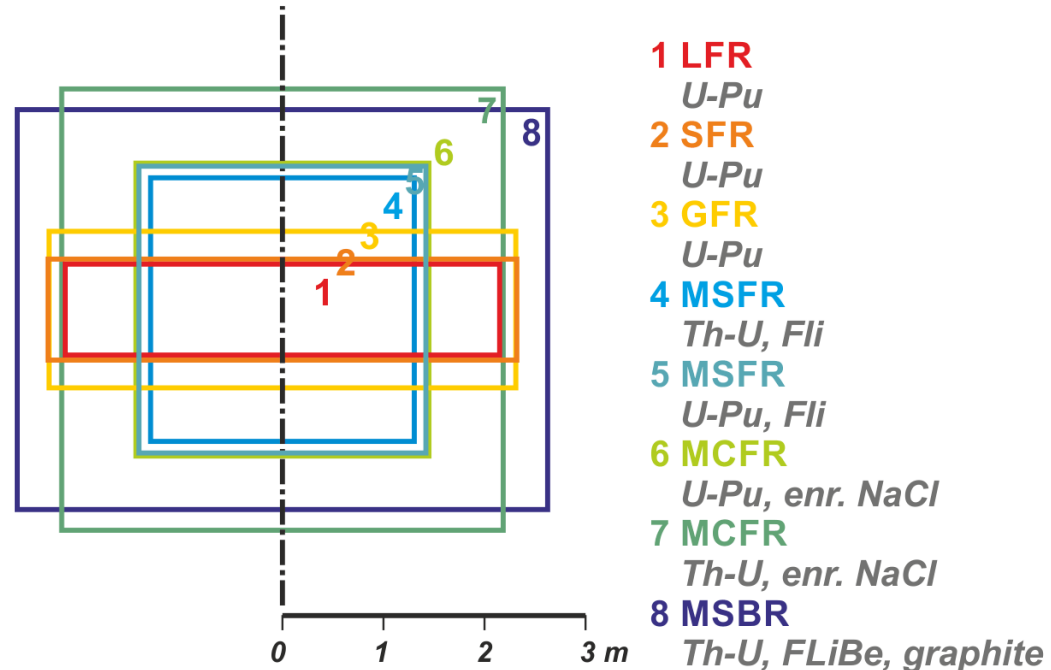
- Bare core size can be estimated for several BG values.

U-Pu cycle =>



Self-sustaining breeder in closed cycle

- Using 1m **hastelloy reflector** core size was estimated for **single-fluid** designs.
- It was compared with classical fast reactors.
- MSFR (**Fli**) in Th-U (4) is compact.
- MSFR (**Fli**) in U-Pu (5) is bigger.
- MCFR (**Na³⁷Cl**) in U-Pu (6) is comparable to MSFR in U-Pu (5).
- MCFR (**Na³⁷Cl**) in Th-U (7) is big.
- MSBR (ORNL design, 13% salt).



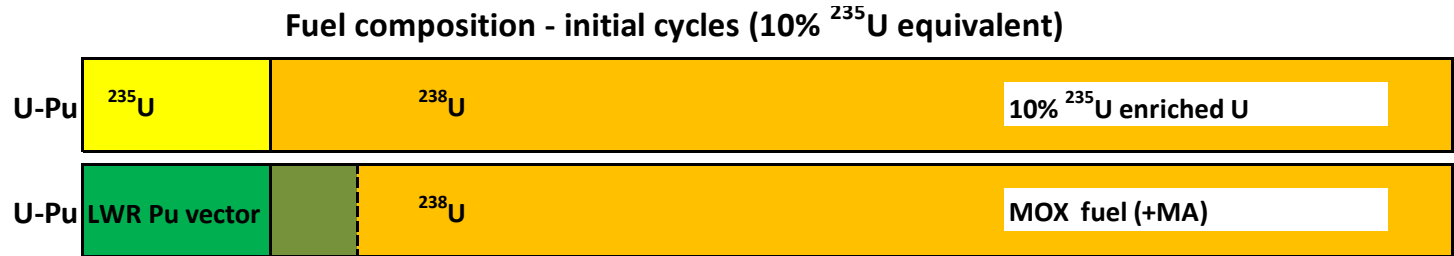
Transition to closed Th-U and U-Pu cycle

5 major fissile materials to start the Th-U cycle

Material	RG_Pu	LEU	HEU	U233	WG_Pu
Fissile isotope(s)	$^{239}\text{Pu}, ^{241}\text{Pu}$	^{235}U	^{235}U	^{233}U	^{239}Pu
Fissile isotope share	~60%	1-20%	21-95%	60-100%	>93%
“Availability”	medium	high	high	low	medium
Proliferation risk	medium	medium	high	high	high

RG_Pu and LEU as initial fuel load

- Both **RG_Pu** and **LEU** are very natural option to start the **U-Pu** cycle.



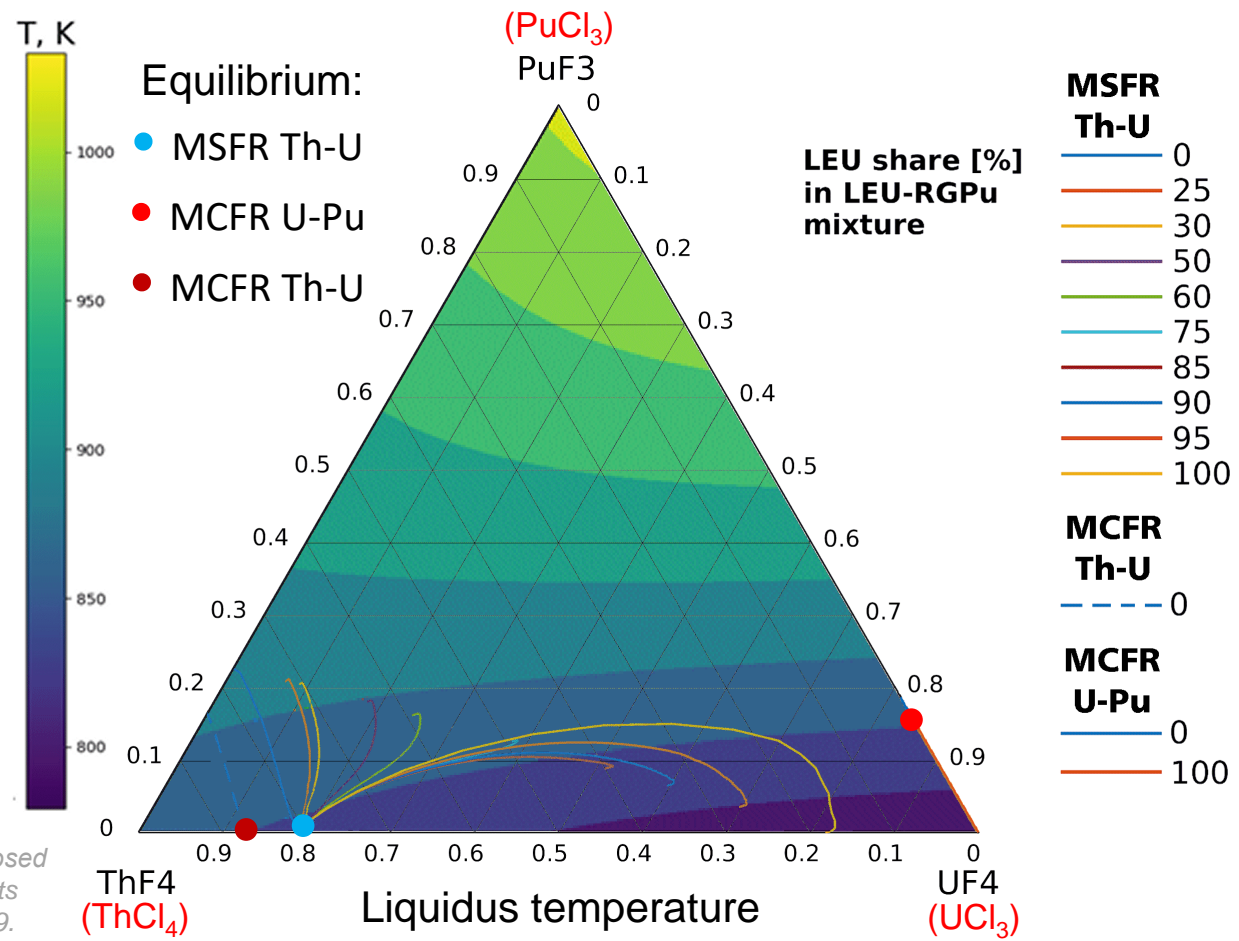
- Starting **Th-U** cycle with **LEU** induces ^{238}U presence in the core.



- Starting **Th-U** cycle with **RG_Pu**, **LEU** or their mixture introduces strong **perturbation**.
- Pu and ^{235}U & ^{238}U** are not presented in the salt at equilibrium Th-U cycle.



Transition to Th-U cycle in MSFR (and U-Pu MCFR)

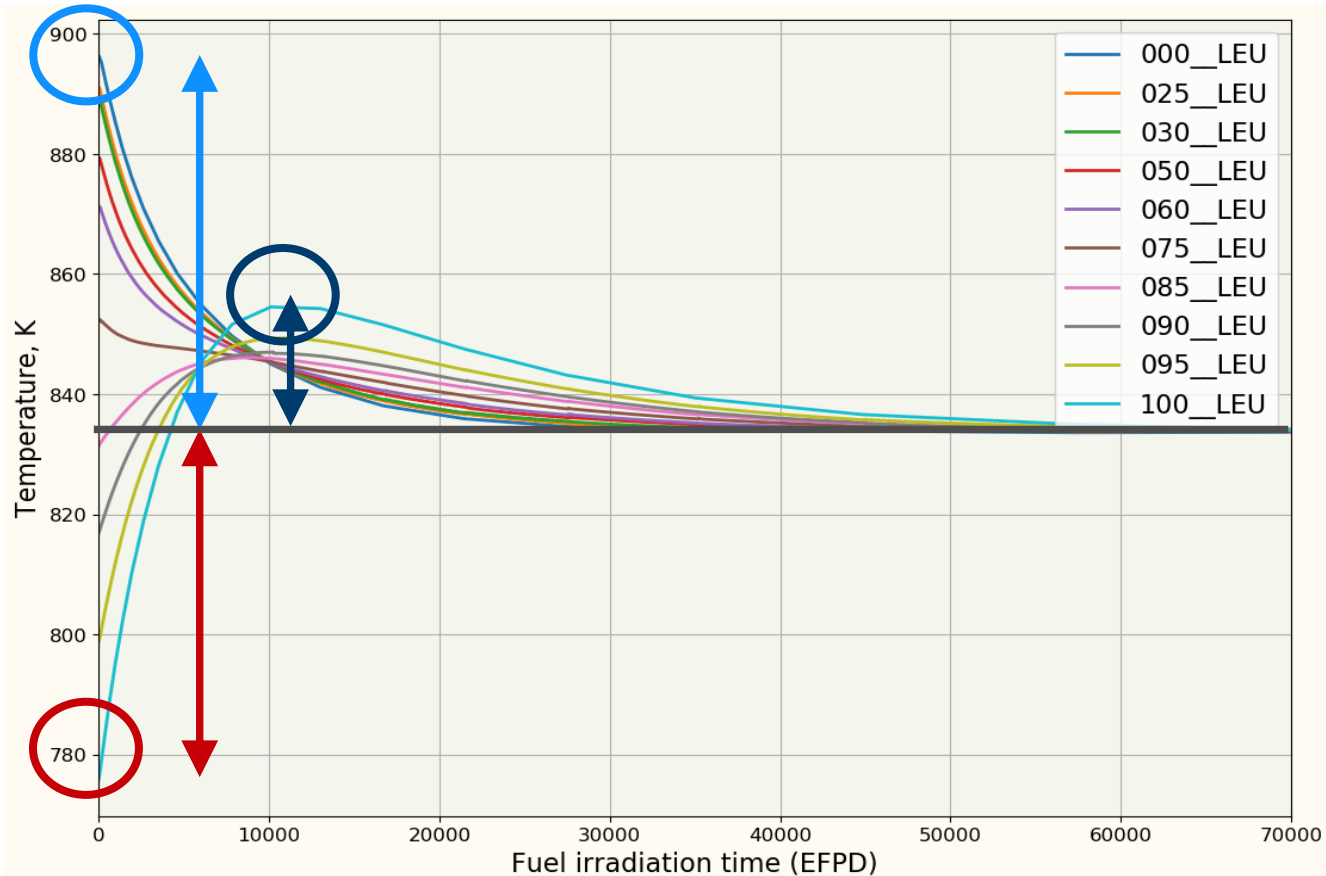


Křepel, J. et al., Transition to closed Th-U fuel cycle in fluoride salts based fast MSR. ICAPP 2019.

Evolution of the liquidus temperature

- **PuF₃** presence increases the liquidus temperature.
- **UF₄** presence decreases the liquidus temperature.
- The temperature for equilibrium Th-U cycle is cca **835K**.
- The difference is: up to **-55K** and **+60K** or **+20K**.

LiF-ThF₄-UF₄-PuF₃
(at **76.6%** LiF).



Self-sustaining breeding in open cycle (B&B)

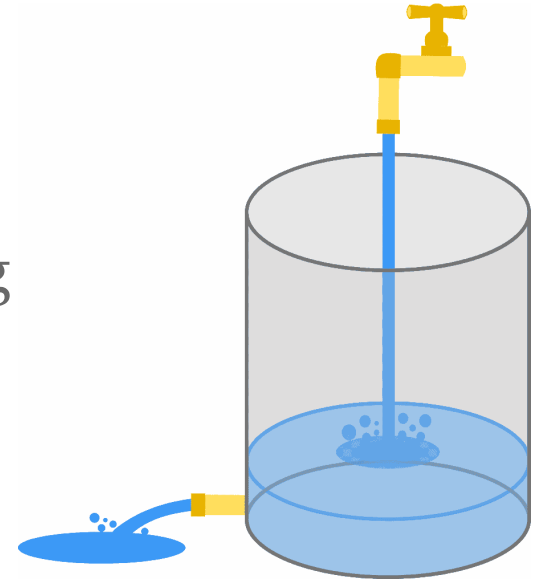


Illustration of tap-like reactor

<https://www.subpng.com/>

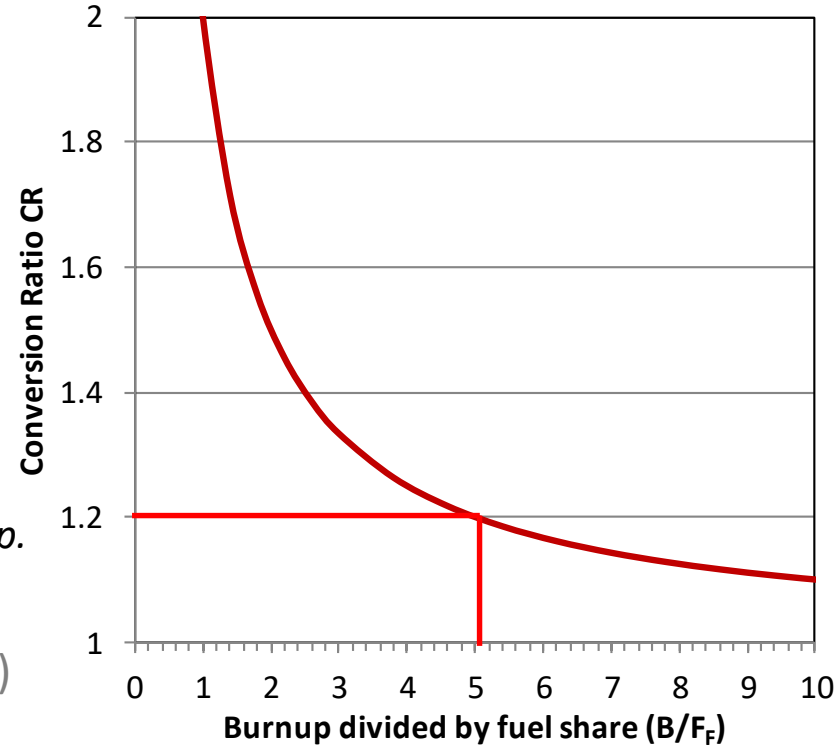
Trivial criteria for breed-and-burn cycle operation

- In B&B cycle conditions:
 - fresh fuel is only fertile material
 - spent fuel is not recycled.
- B&B trivial criterion (tautology): $I = II$
 - Fissile Fuel F_F share in the discharged fuel.
 - New fissile fuel bred in the discharged fuel.

$$F_F = B(CR - 1) \Rightarrow \frac{1}{CR - 1} = \frac{B}{F_F}$$

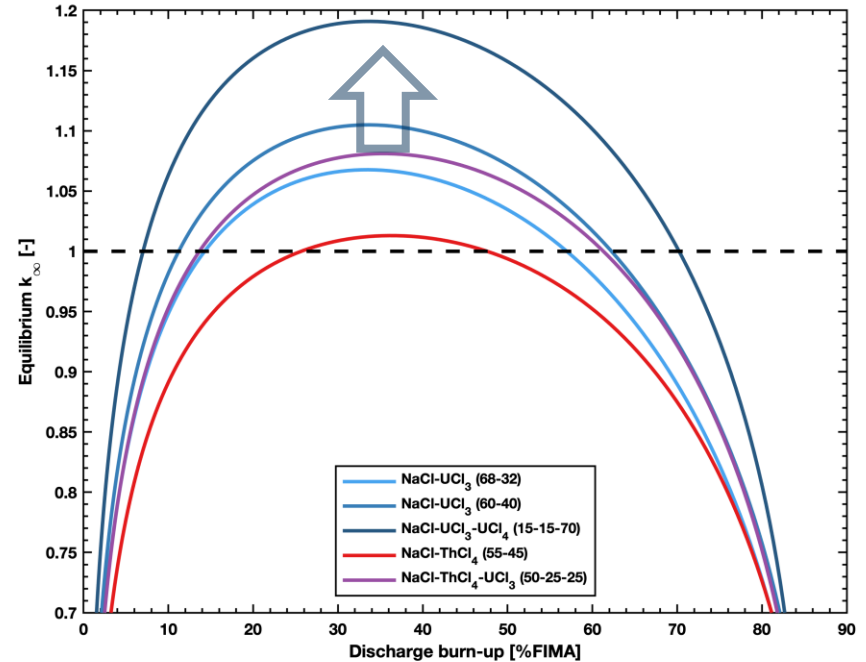
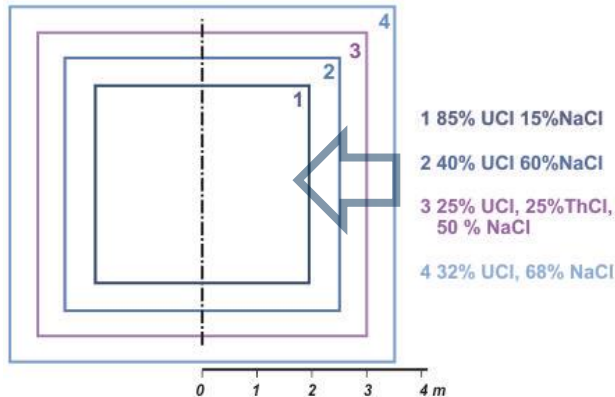
where CR is conversion ratio and B is the fuel burnup.

- Reactor must be critical for CR , F_F , and B ,
e.g. for $CR=1.2$: $F_F=10\% \Leftrightarrow B=50\%$ ($1\% \Leftrightarrow 5\%$)
- Fuel utilization in B&B cycle?
It is equal to the burnup.



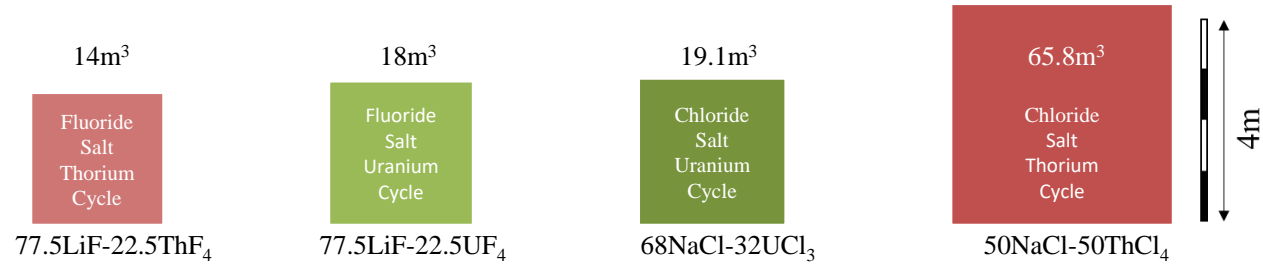
Self-sustaining breeder in open cycle (B&B)

- B&B is practically not possible in Th-U cycle.
- It is only possible in mixed U-Pu & Th-U cycle.
- B&B cores are bulky (chlorides = hard spectrum, but also high Migration area).
- The performance increases with growing actinides share in the core.



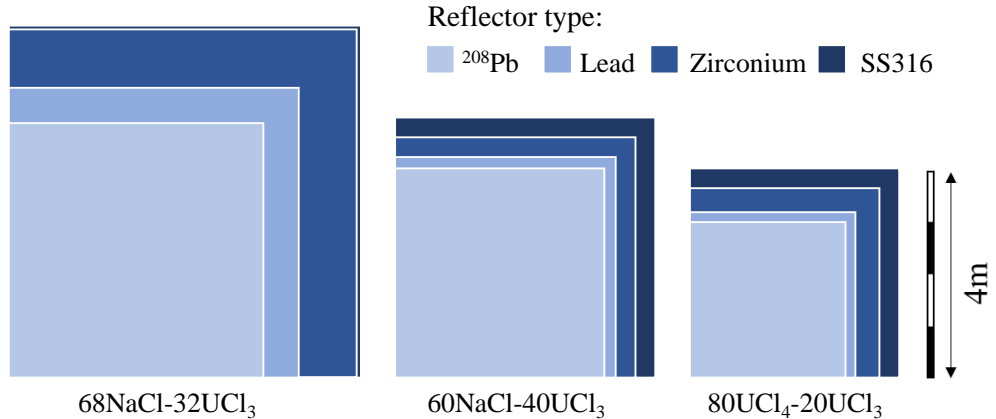
Core size comparison for closed and open B&B cycle

- Closed cycle



- Breed and burn open cycle

Critical core sizes

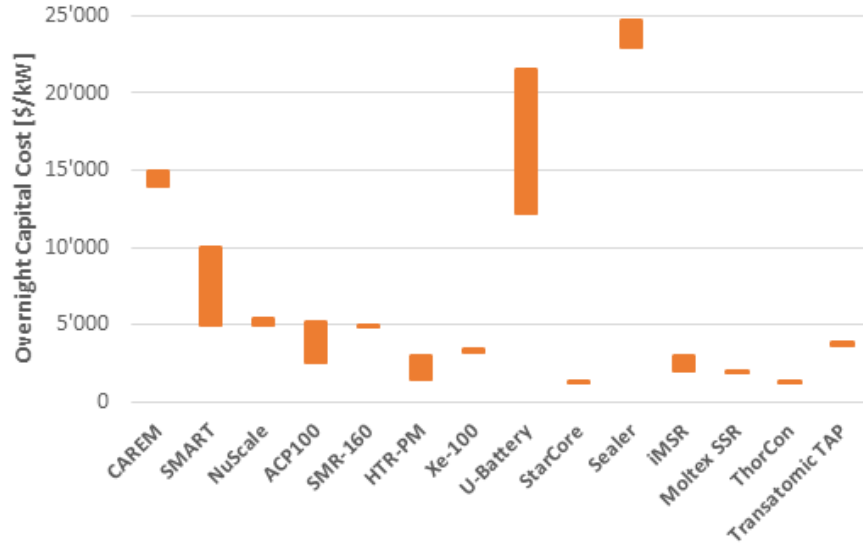


Summary – fuel cycle

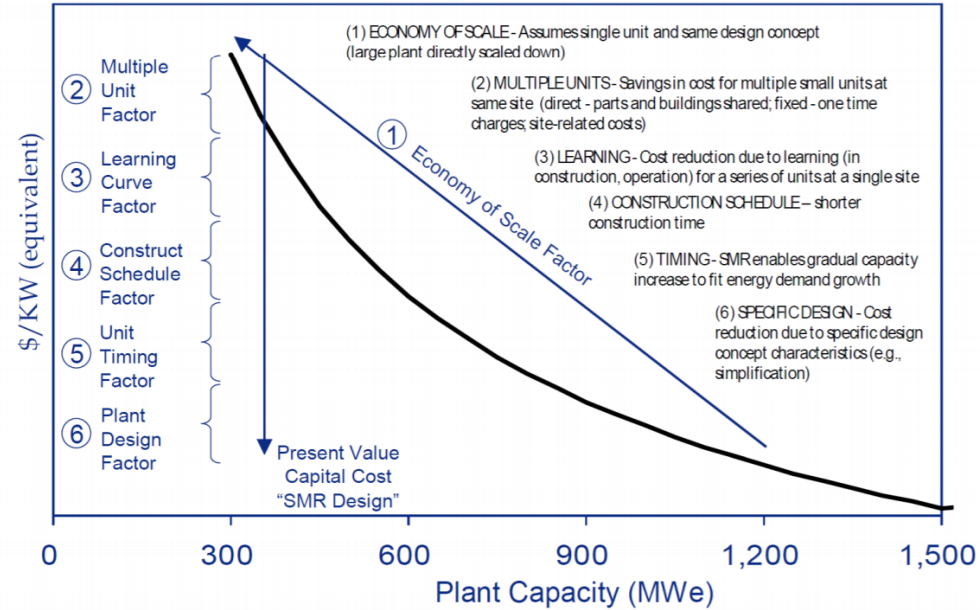
- MSR can be designed as very compact burner fueled by minor actinides or enriched uranium.
- MSR can be designed as a Th-U or U-Pu breeder.
- Fluoride salts are more convenient for Th-U cycle (thermal or fast).
- Chloride salts are more convenient for U-Th cycle (fast).
- Chloride salts are applicable for U-Pu breed-and-burn cycle (fast).
- The core size may be relatively large.
- The low-pressure vessel can be still manufactured in factory as a module.

Some general comments on size and modularity

Thesis of P. Dobrzynski, Estimating the Cost of Small Modular Reactors, 2017.

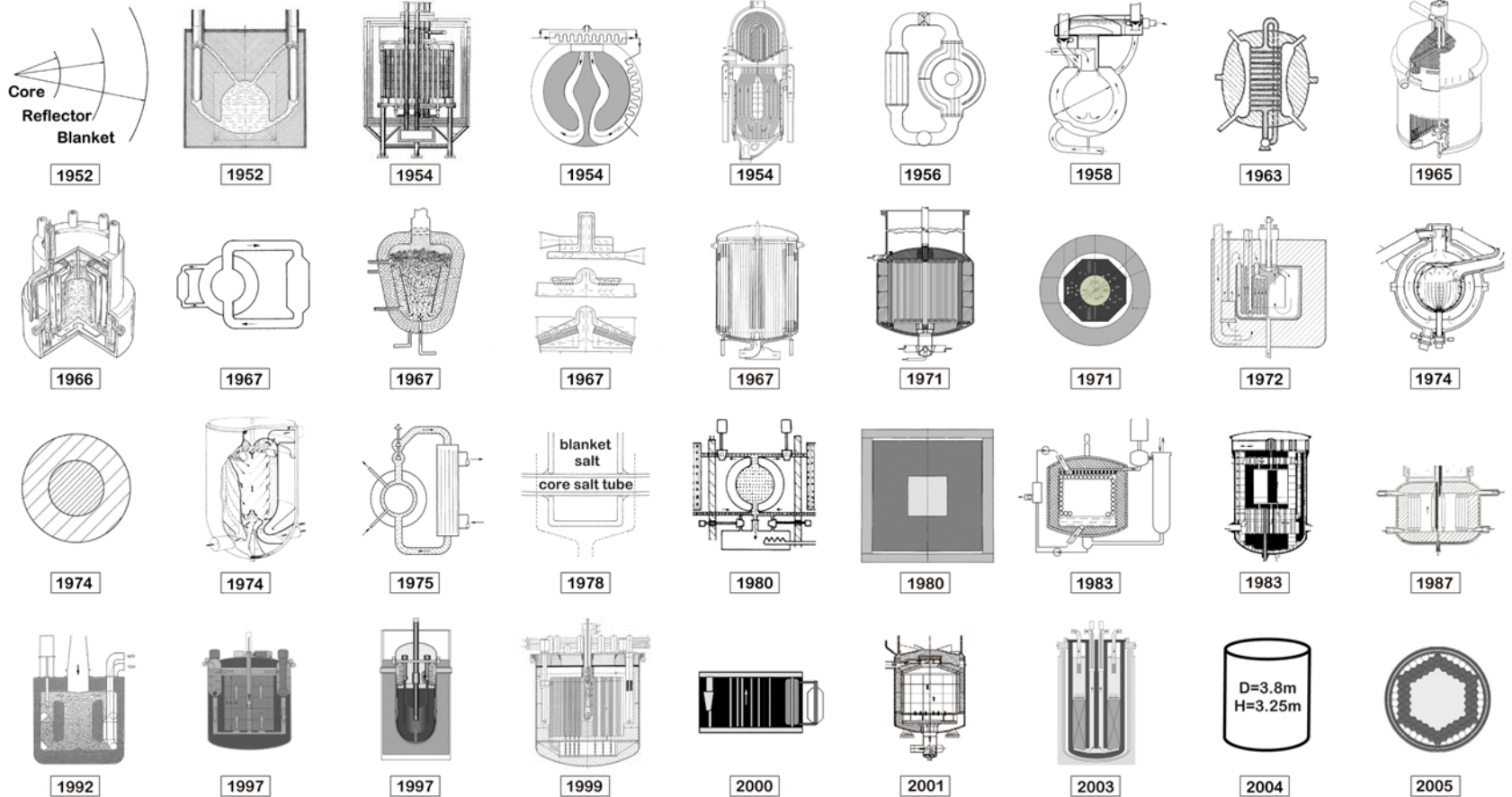
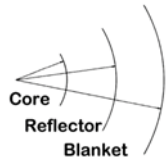


Available vendor estimates of the selected SMRs



Influence of SMR's features on the unit investment cost (Carelli, et al., 2008)

Recall of the variety



1952

1952

1954

1954

1954

1956

1958

1963

1965

1966

1967

1967

1967

1967

1971

1971

1972

1974

1974

1974

1975

1978

1980

1980

1983

1983

1987

1992

1997

1997

1999

2000

2001

2003

2004

2005

blanket salt
core salt tube

D=3.8m
H=3.25m

General conclusion

- MSR is not a reactor, it is entire category of reactors.
- MSR has potential to be safe, sustainable and economic at once.
- MSR is not yet safe, sustainable and economic...
- We should firstly utilize the potential.
- Safety evaluation would be premature for many concepts.
- Economy was also not evaluated, but prototypes would be expensive.
- Sustainability, or actually breeding capability, was evaluated.
- MSR can breed in both open and closed cycle.
- MSR might be designed as modular reactor.
- Preferably Big Modular Reactor (BMR).

**Thank you.
Questions?**



List of accomplished MSc and PhD theses

- Listed are also several theses accomplished in **cooperation** with other institutes or by visiting students from **POLIMI** and **CTU Prague**.

Name	Title	Thesis type / Status	Year
B. Hombourger	Parametric Lattice Study for Conception of a Molten Salt Reactor in Closed Thorium Fuel Cycle	Swiss nuclear MSc course / Defended	2013
C. Fiorina	The Molten Salt Fast Reactor as a Fast-Spectrum Candidate for Thorium Implementation	PhD thesis in cooperation with PSI / Defended at POLIMI	2013
V. Ariu	Heat exchanger analysis for innovative molten salt fast reactor	Swiss MSc course / Defended	2014
M. Auffero	Development of advanced simulation tools for circulating-fuel nuclear reactors	PhD thesis in cooperation with PSI / Defended at POLIMI	2014
H. Kim	Static and transient analysis of Molten Salt Reactor Experiment using SERPENT-2 / TRACE / PARCS codes	Swiss nuclear MSc course / Defended	2015
J. Choe	Empirical Decay Heat Correlations and Fission Products Behavior in MSRs	Swiss nuclear MSc course / Defended	2015
J. Bao	Development of the model for the multiphysics analysis of Molten Salt Reactor Experiment using GeN-Foam code	Swiss nuclear MSc course / Defended	2016
D. Pyron	Safety Analysis for the Licensing of Molten Salt Reactors	Swiss nuclear MSc course / Defended	2016
E. Pettersen	Coupled multi-physics simulations of the MSFR using TRACE-PARCS	MSc thesis in cooperation with PSI / Defended at Univ. Paris-Saclay	2016
N. Vozarova	Behaviour of fission products in the molten salt reactor fuel	Swiss MSc course in cooperation with JRC Karlsruhe / Defended at ETHZ	2016
M. Zanneti	Development of new tools for the analysis and simulation of circulating-fuel reactor power plants	PhD thesis in cooperation with PSI / Defended at POLIMI	2016
E. Losa	U-Pu and Th-U Fuel Cycle Closure	PhD thesis in cooperation with PSI / Defended at TU Prague	2017
B. Hombourger	Small modular Molten Salt Fast Reactor design for closed fuel cycle	PhD thesis in frame of SNSF project at PSI Defended at EPFL	2018
M. Di Filippo	Development of dedicated MSR burn-up tool	Swiss nuclear MSc course / Defended	2018
V. Sisl	Thorium Utilization in the Fuel Cycle of Advanced Nuclear Reactors	MSc thesis in cooperation with PSI / Defended at TU Prague	2018
T. Koivisto	Assessment of waste burning in open cycle of two fluids chloride MSR	MSc thesis in cooperation with PSI / defended at Aalto University in Finland	2019
V. Raffuzzi	Modelling of batch-wise operation of European Sodium Fast Reactor and Breed&Burn Molten Salt Reactor	Swiss nuclear MSc course / Defended	2019
J. Dietz	Chemical-Thermodynamic Modelling of the MSR-Related Systems Under Normal and Accident Conditions	Swiss nuclear MSc course / Defended	2020
R. Gonzales	Improved methodology for analysis and design of Molten Salt Reactors	PhD thesis in frame of H2020 project SAMOFAR / Defended at EPFL	2021
J. Santora	Two-fluid Breed & Burn MCFR parametric study	Swiss nuclear MSc course / Defended	2022
F. Borys	Identification and evaluation of transmutation criteria for selected reactors	Swiss nuclear MSc course	2023
M. Krstovic	Extended Point Kinetics Solver For MSR	Swiss nuclear MSc course	2023

Burnup definition

Burnup definition

- For solid fuel burnup is defined as:

$$B_{Gwd/tHM}(t) = \frac{\int_0^t P(t) dt}{M_{Ac}(0)}$$

$$B_{FIMA\%}(t) = \frac{\int_0^t F(t) dt}{N_{Ac}(0)}$$

$$FPS_{share} = \frac{M_{FPs}(t)}{M_{Ac}(0)} = \frac{M_{FPs}(t)}{M_{Ac}(t) + M_{FPs}(t)} = B_{FIMA\%}(t)$$

- For liquid fuel two definitions are possible:
- Differential

$$B_{Gwd/tHM}(t) = \frac{P(t)}{M_{Ac,in}^{\&}(t)}$$

$$B_{FIMA\%}(t) = \frac{F(t)}{N_{Ac,in}^{\&}(t)} \cong \frac{N_{Ac,in}^{\&}(t) - N_{Ac,out}^{\&}(t)}{N_{Ac,in}^{\&}(t)} \cong \frac{N_{FPs,off-gas}^{\&}(t) + N_{FPs,out}^{\&}(t)}{N_{Ac,in}^{\&}(t)}$$

- Integral

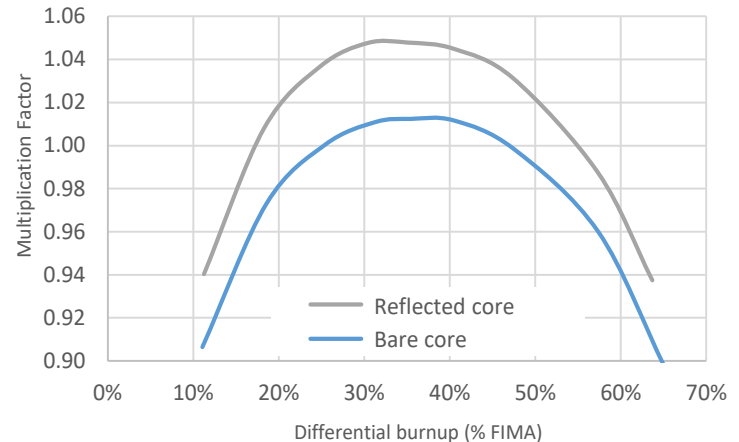
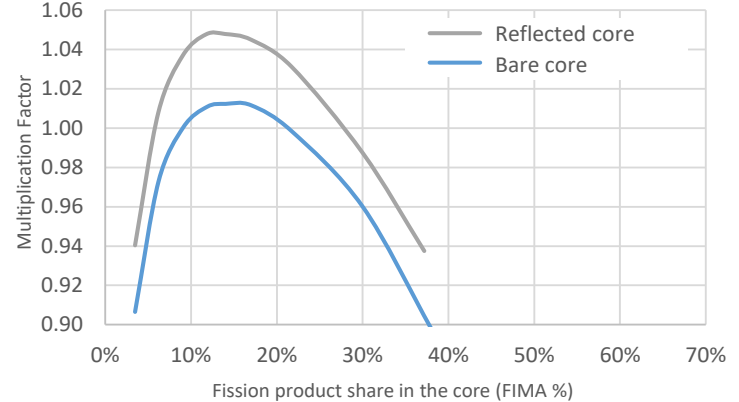
$$B_{Gwd/tHM}(t) = \frac{\int_0^t P(t) dt}{M_{Ac,core}(0) + \int_0^t M_{Ac,in}^{\&}(t) dt}$$

$$B_{FIMA\%}(t) = \frac{\int_0^t F(t) dt}{N_{Ac,core}(0) + \int_0^t N_{Ac,in}^{\&}(t) dt} = \frac{\int_0^t F(t) dt}{N_{Ac,core}(t) + \int_0^t N_{Ac,out}^{\&}(t) dt + \int_0^t F(t) dt}$$

Burnup definition

- Due to the continuous fission products removal by off-gas system, the burnup and the fission products share in the core differs.

$$FPs_{share,core} = \frac{M_{FPs,core}(t)}{M_{Ac,core}(t) + M_{FPs,core}(t)} \neq B_{FIMA\%}(t)$$



Burnup definition

- The differential and integral definition provide different values.
- The integral definition includes the initial core loading.
- For stabilized and long enough operation, they can be equal.

$$B_{GWd/iHM}(t) = \frac{P(t)}{M_{Ac,in}(t)}$$

 \Leftrightarrow

$$B_{GWd/iHM}(t) = \frac{\int_0^t P(t) dt}{M_{Ac,core}(0) + \int_0^t M_{Ac,in}(t) dt}$$

