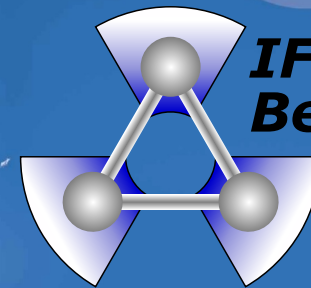


The Dual Fluid Reactor

An environmental-friendly nuclear concept for
cost-efficient electricity and fuel
with no need for geological waste storage



**IFK
Berlin**

*Institute for
Solid-State Nuclear Physics
gGmbH*

dual-fluid-reactor.org



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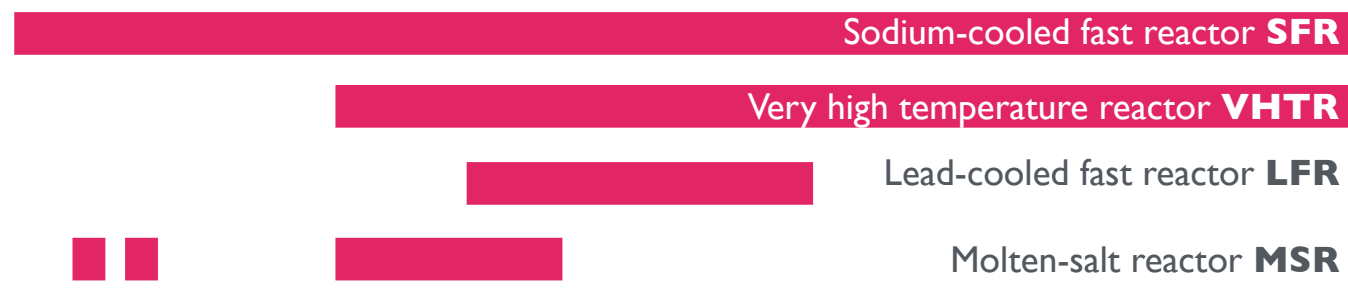
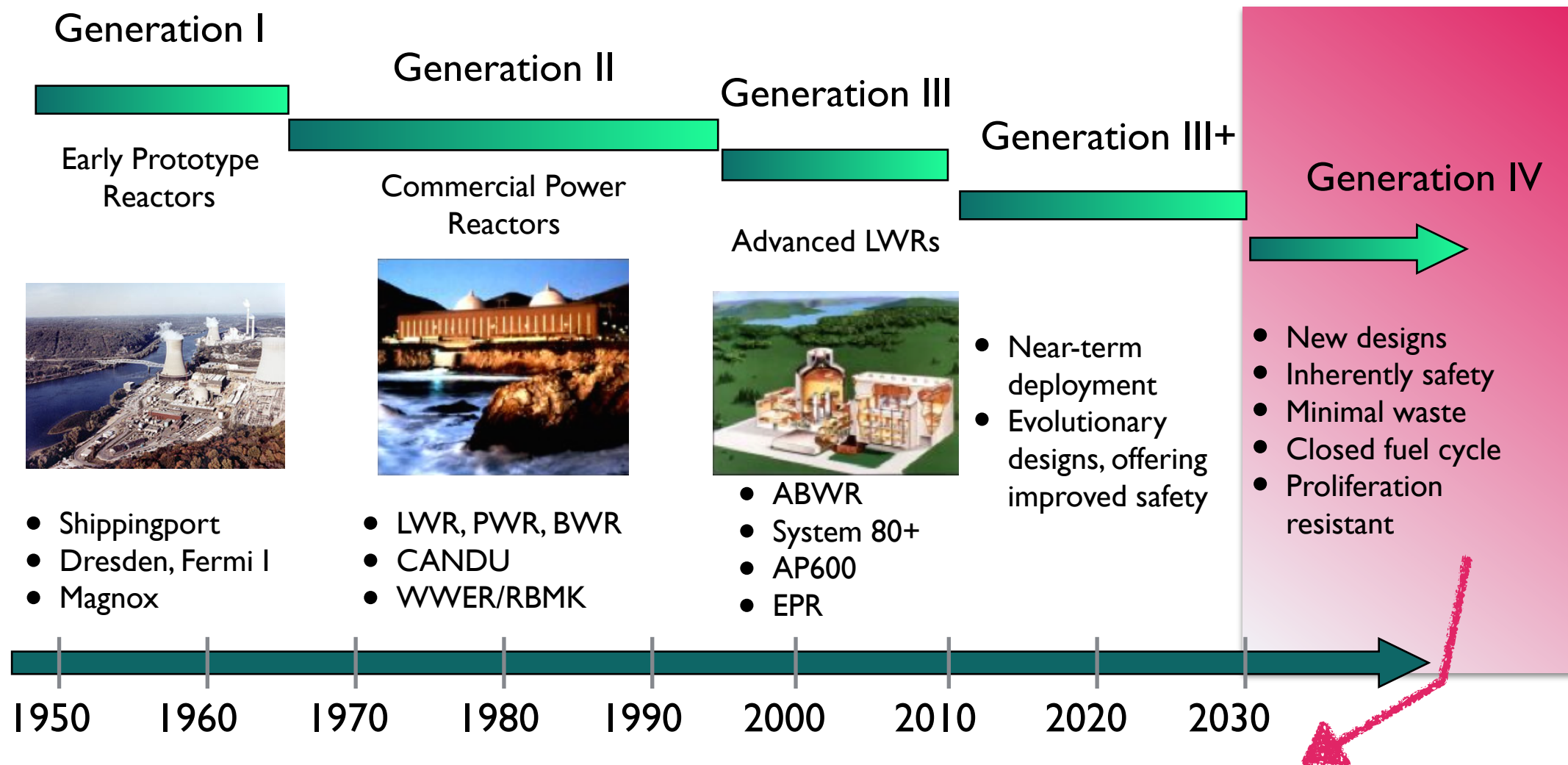


Dr. rer. nat. Götz Ruprecht



Technische Universität München
Department for Nuclear Technology
Prof. Dr. Rafael Macián-Juan

Reactor development - where we are



Generation IV

Not like fusion research. They all have been running in the past.

The main problem with Generation IV is the economy: They still use **solid fuel rods** and therefore still require the **expensive fuel cycle**. Only exception: The molten salt reactor **MSR**.

The Dual Fluid Reactor

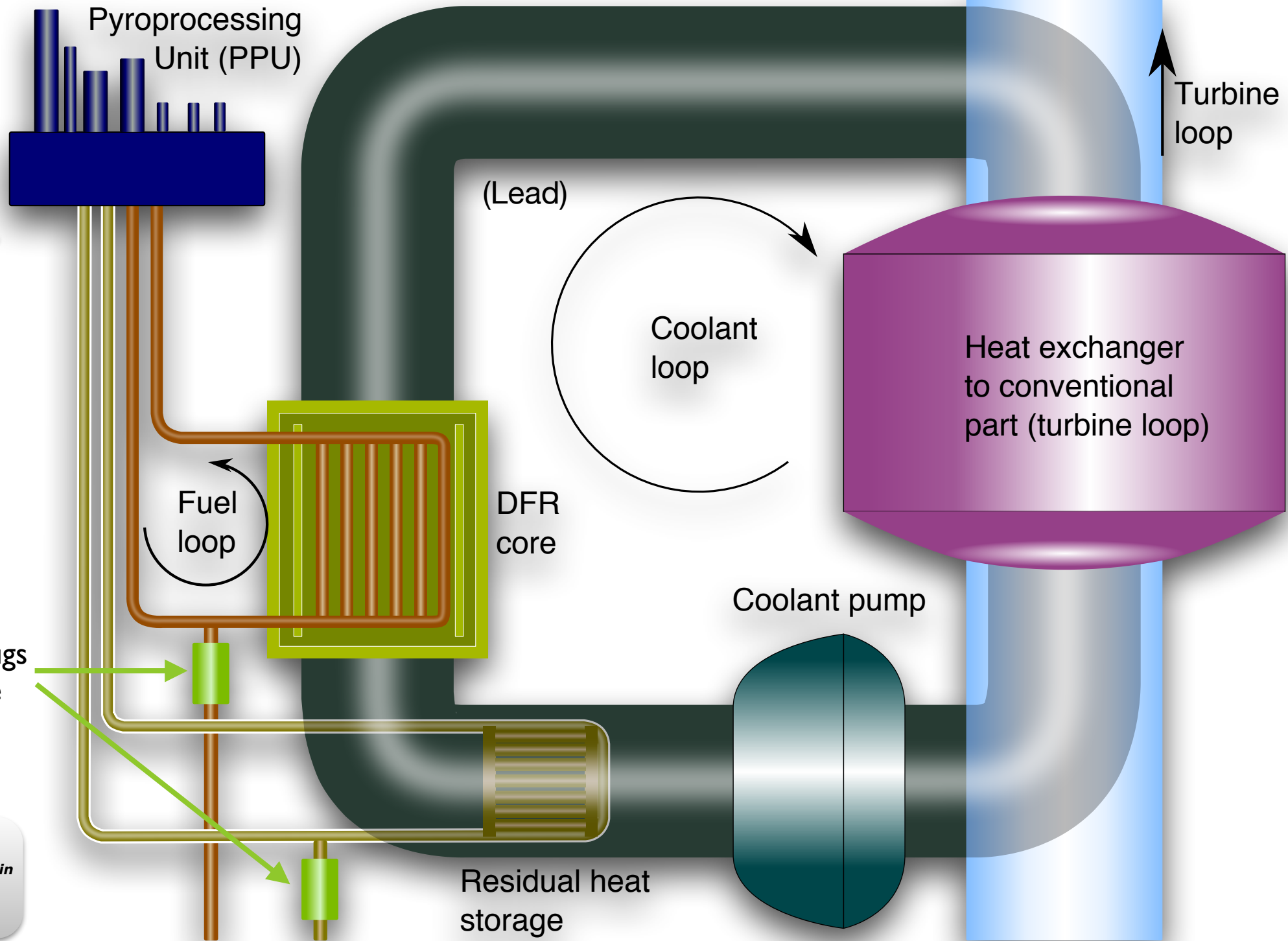
A concept beyond Generation IV

- Natural Uranium
- Depleted Uranium
- Thorium
- Used fuel elements



- Fission products
- Med. radioisotopes
- Fissile material

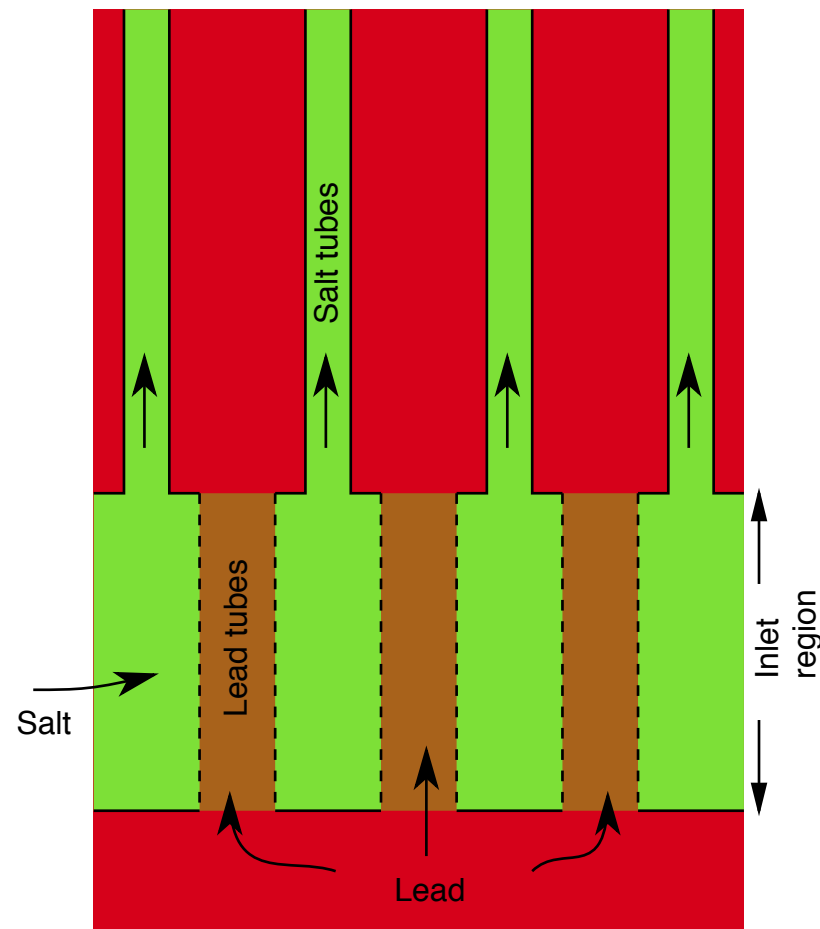
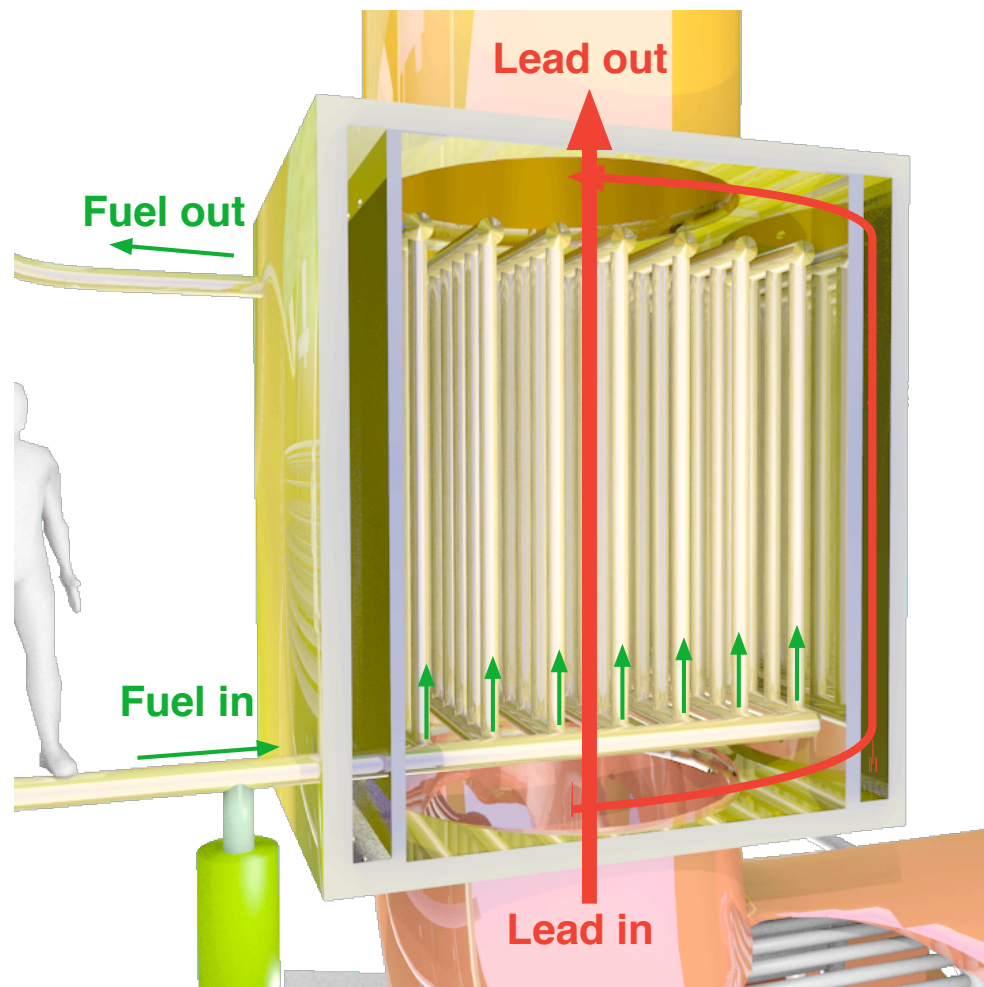
Melting fuse plugs = run-away safe



International patent protection for the Dual Fluid principle since Sep. 2011



The Dual Fluid Principle



International patent application PCT/DE2012/000957

- Nuclear reactor including a primary duct for **continuous** insertion and discharge of liquid fuel into and out of a core vessel wherein the fuel duct is lead through the core vessel, characterized by
- a **secondary duct** for a liquid coolant wherein the coolant enters the aforementioned core vessel via an inlet, **passing and bathing the primary duct** and leaving the core vessel via an outlet.

DFR power plant

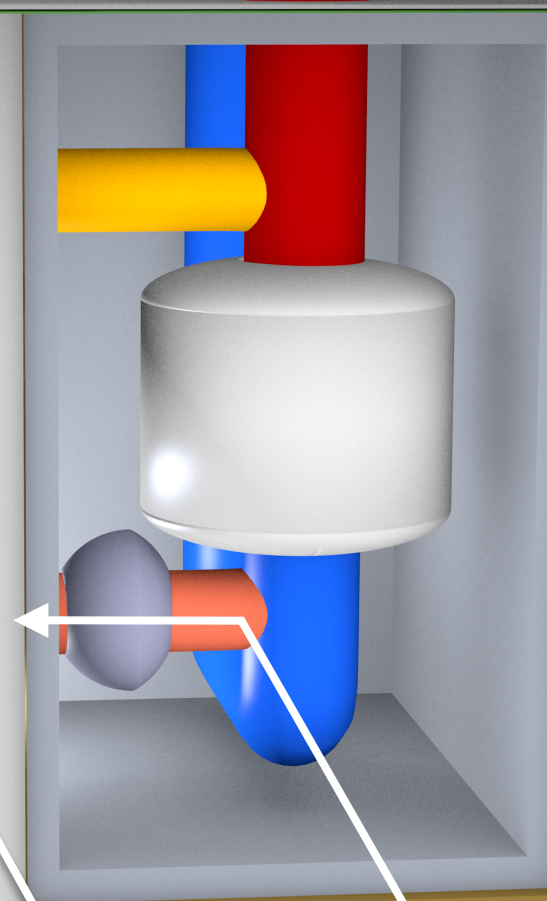
<http://festkoerper-kernphysik.de>



Pyrochemical
Processing
Unit (PPU)

On-site storage. Pre-assorted
rare metals available after
latest 300 years.

Sub-critical fuel tanks



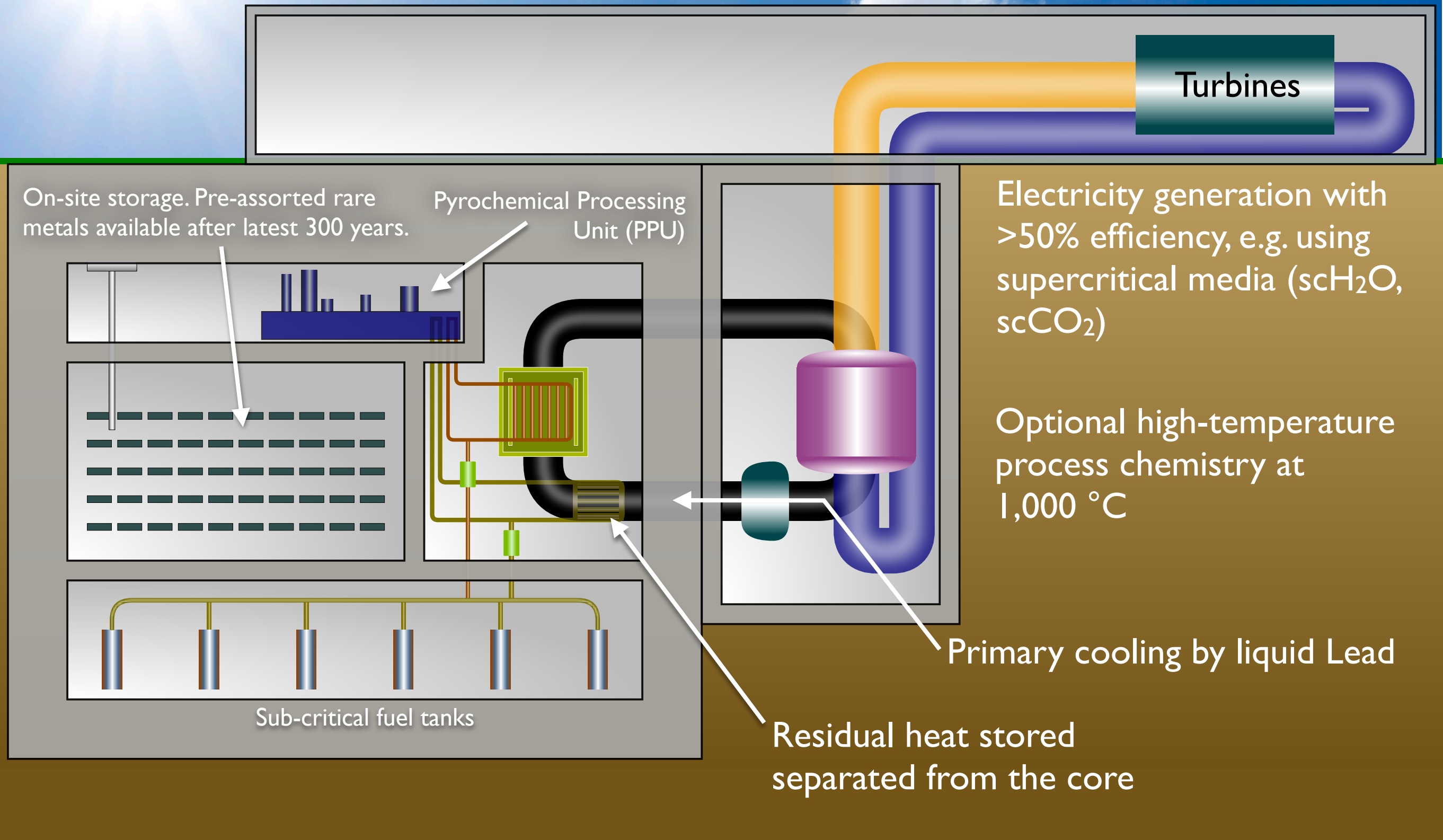
Electricity generation with
>50% efficiency, e.g. using
supercritical media (scH_2O ,
 scCO_2)

Optional high-temperature
process chemistry at
 $1,000\text{ }^\circ\text{C}$

Primary cooling by liquid Lead

Residual heat stored
separated from the core

DFR power plant



DFR Control

- Highly negative temperature coefficient due to thermal expansion of the liquid fuel
 - Temperature rises → Fission rate and heat production drop
 - Temperature drops → Fission rate and heat production rise
- Therefore, the temperature is held homeostatic at 1000 °C
 - no material stress on power change
- Therefore, power is fully regulated by heat extraction
 - Load-following operation in the grid
- Therefore, also qualified for rapidly changing power demand in chemical plants (process heat)
- No mechanical regulation equipment needed
- The reactor can be on „stand by“ in a critical state at zero power output → Safe operation mode

DFR Safety

Safety advantages

- Highly negative temperature coefficient → Self regulation
- Residual heat disposal by natural convection
- Overheat protection by passive melting fuse plugs
- No static overpressure

Problem

Risk of leakages

More volatile activity due to high operating temperature and liquid fuel

Raised proliferation risk due to online fuel processing

Solution

No overpressure, survey and containment without much effort

Will be extracted/absorbed in the online fuel processing, containment

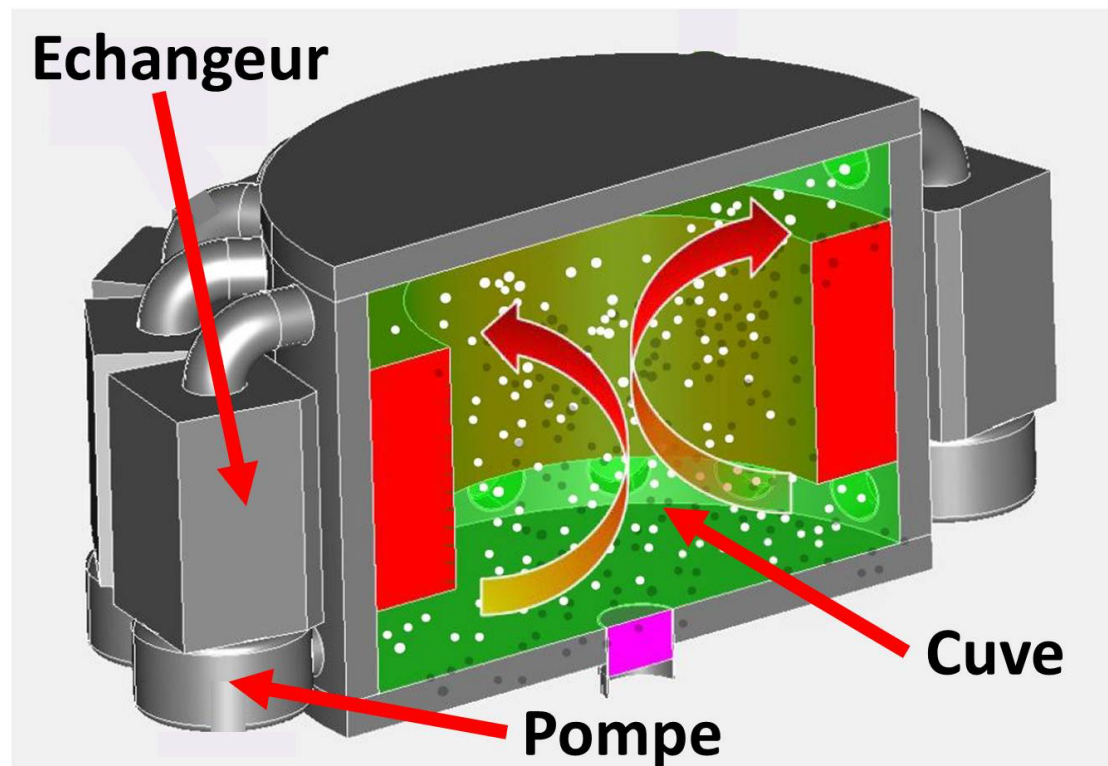
Fixed piping, encapsulation and monitoring is easier to accomplish due to compact size

Why is the DFR not an MSR?

Molten Salt Reactor (MSR)

Single fluid

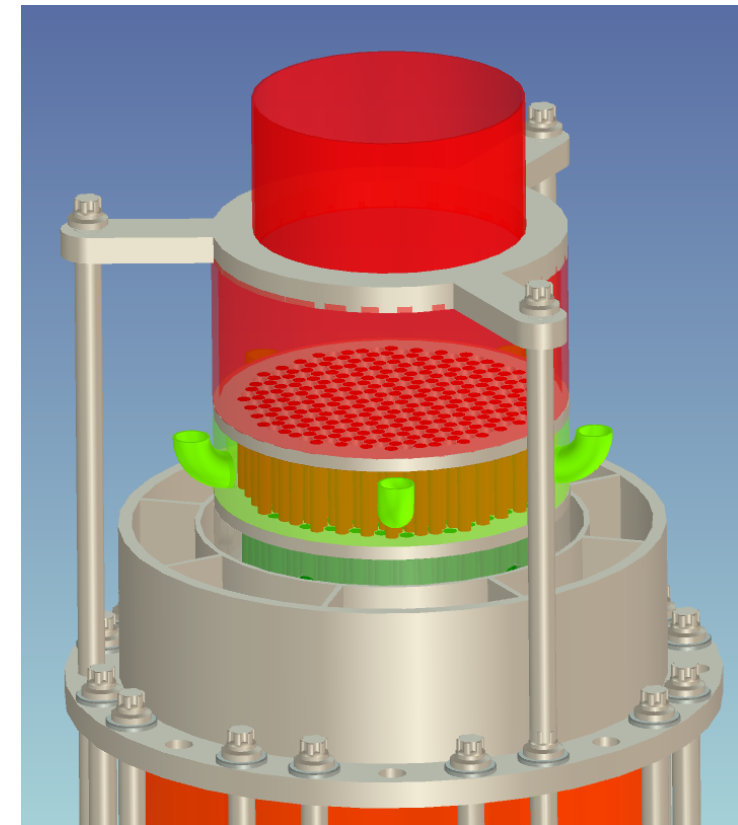
- Homogeneous core
- Heat removal by salt
- Fuel limited to salt



Dual Fluid Reactor (DFR)

Two fluids

- Heterogeneous core
- Heat removal by second fluid
- Fuel liquid less constrained



The double function of fuel providing and heat removal in the MSR limits its power density. **This limitation is not present at the DFR.**

Difference to MS(F)R concepts

- The DFR concept does not rely on molten-salt. There are 2 development threads

DFR/m with **molten-metal** fuel

DFR/s with **molten-salt** fuel

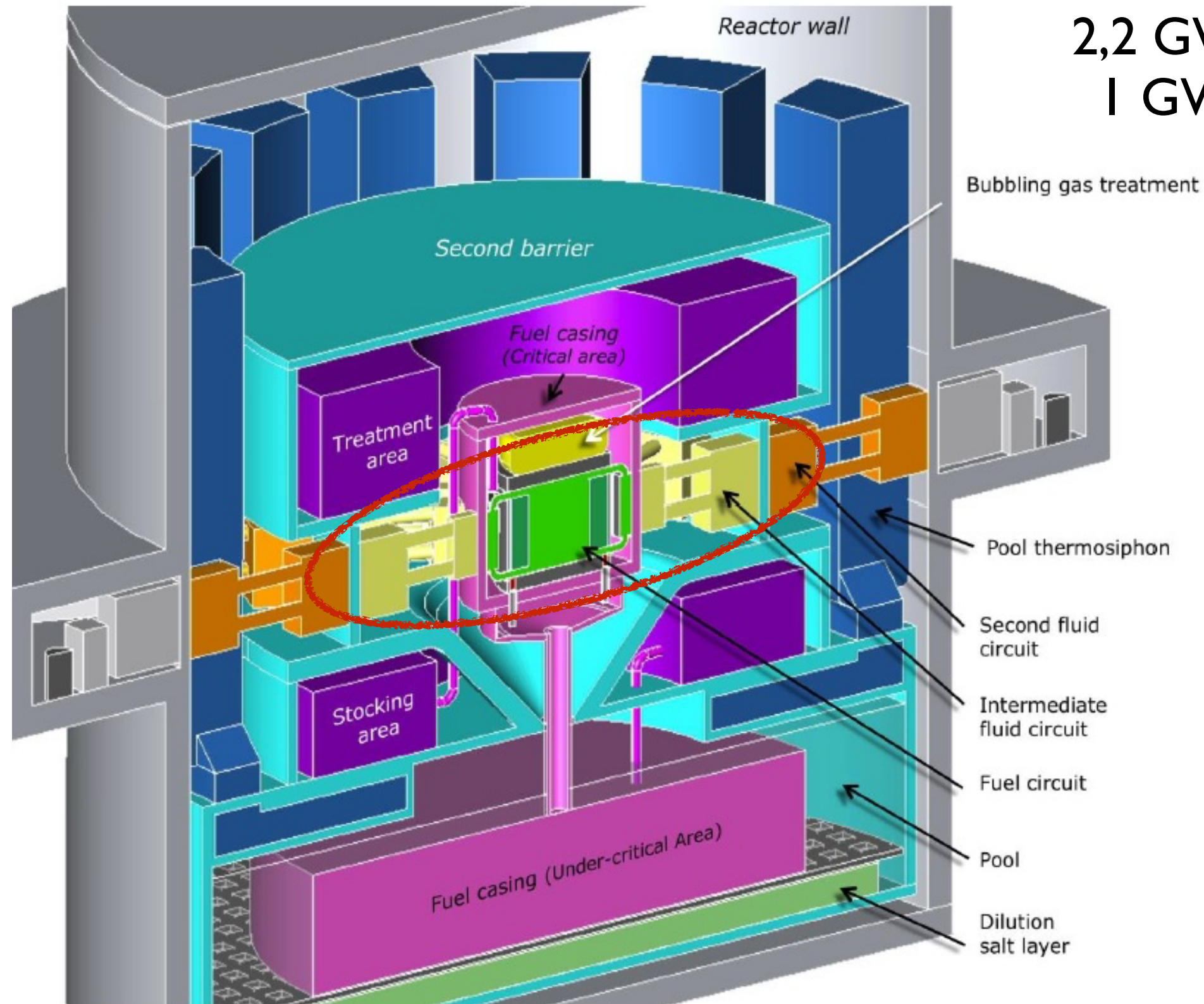
- DFR/s is quite different from MSR
 - The salts are **undiluted** (No Li, Be, or other carrier salts)
 - The salts are **chlorides** (UCl_3 , ThCl_3 , PuCl_3 ...)
 - The reprocessing is based on **distillation/rectification**, a very simple physical separation processes.
- There is no need for liquid/liquid extraction as for fluorbased MSR concepts, or wet-chemical processes.

Power Density: DFR vs MSR

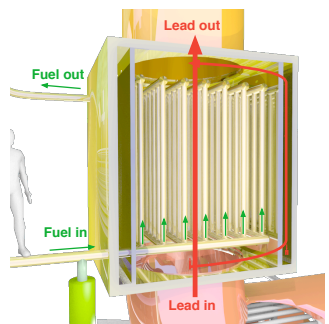
SAMOFAR

2,2 GW_{th}

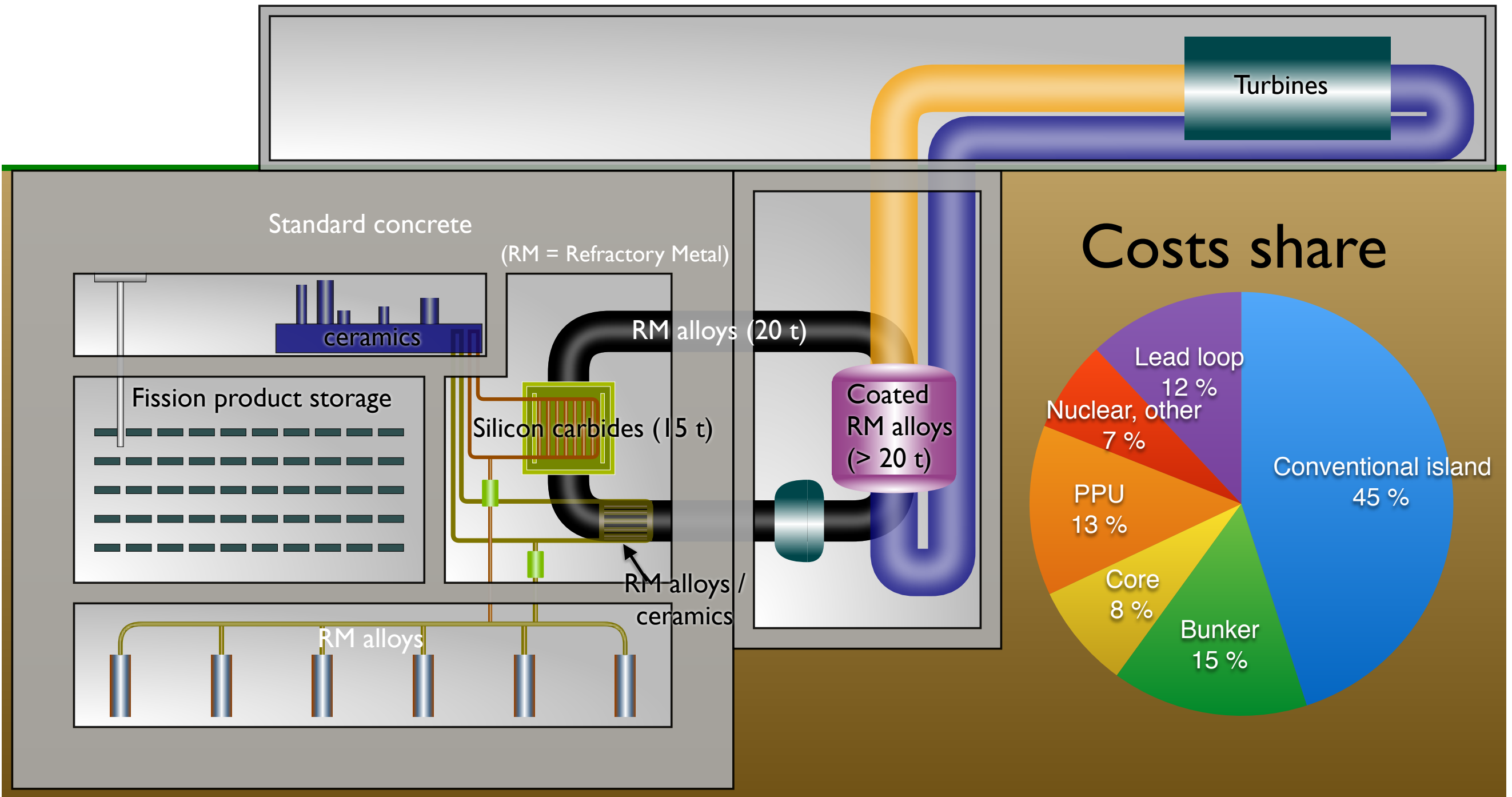
1 GW_e



DFR
3 GW_{th}
1,5 GW_e

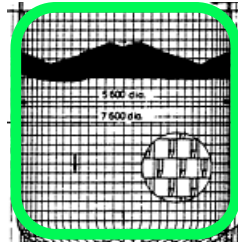


Materials in the DFR Power Plant



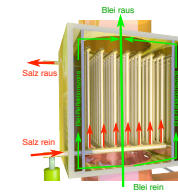
THTR vs DFR

THTR: 300 MW_e



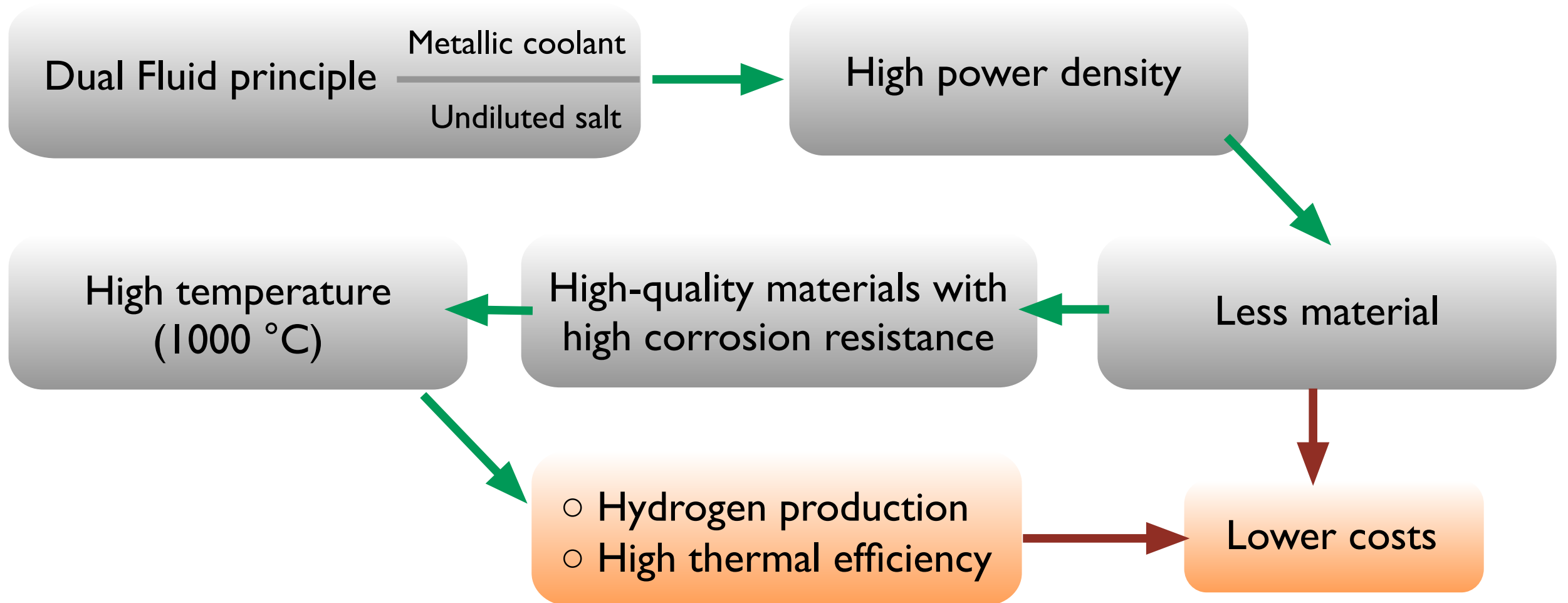
- Infrastructure: Enrichment, fuel element production
- Not thermal and no fuel processing
→ needs final geological deposit
- The GenIVVHTR concept provides even a second intermediary He loop!
- Scalability: max. 300 MW, otherwise risk of core meltdown
- 3 €/Watt, 5 ct/kWh

DFR: 1.500 MW_e



- No further infrastructure
- Fast reactor + online reprocessing
→ no need for final geological deposit
- 1/8 of the volume and 5-fold power
→ 40-fold power density
→ 1/40 of material expenses
- Much higher scalability
- 1 €/Watt, 0,65 ct/kWh

Why is the DFR so efficient?



Salts and Lead at 1000 °C

How is this possible?

- *Outside* the nuclear industry suitable materials are known since a long time
- Focus of nuclear industry so far was on finding *cheap* materials (usually steel alloys) that are corrosion resistant.
- The DFR can afford *expensive* materials due to the low material consumption

Possible Materials

Silicon Carbide (SiC)

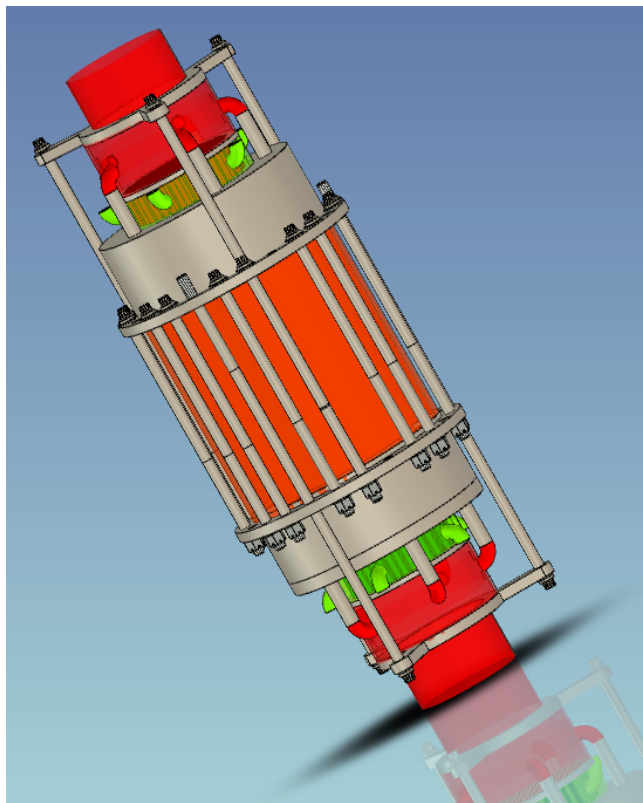
Refractory metal alloys and ceramics

DFR bei 1000 °C

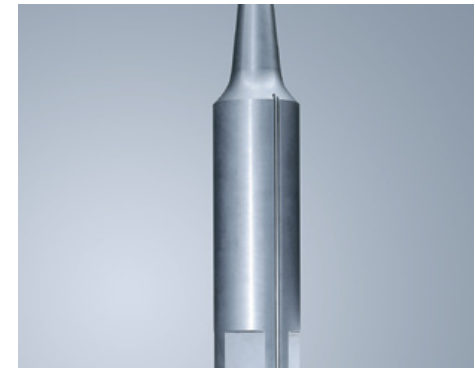
Umspült von flüssigem Blei und Metall-/Salzbrennstoff in einem Neutronenbad
Gibt es dafür Materialien?

Siliziumkarbid (SiC)

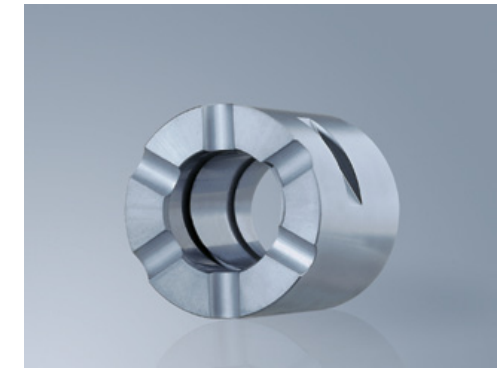
DFR/s-Kern



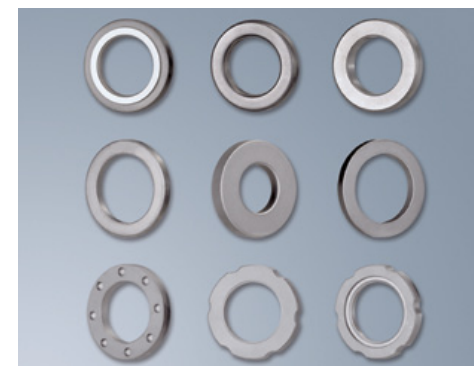
SiC-Maschinenteile →



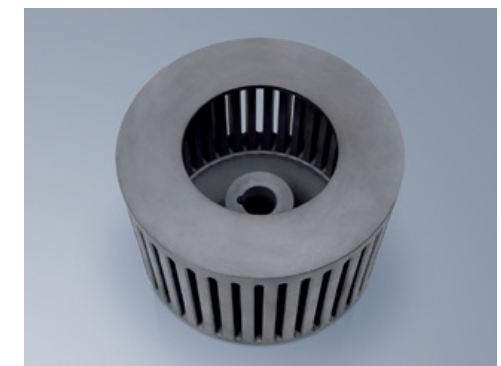
EKasic® Siliziumkarbid Pumpenwellen



Gleitlager aus EKasic® Siliziumkarbid, werden z.B. in hochwertigen Chemie- und Industriepumpen, sowie in Rührwerken für die chemische, pharmazeutische und Lebensmittelindustrie verwendet



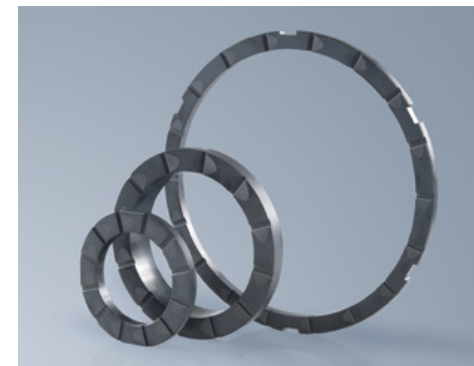
Gleitringdichtungen aus EKasic® Siliziumkarbid eignen sich besonders für Medien, die stark beansprucht sind, z.B. durch Verunreinigung, abrasion und/oder Korrosion



Sichterräder aus EKasic® Siliziumkarbid finden Verwendung in der chemischen, pharmazeutischen, Lebensmittel-, mineralien-, metall- und Recycling-Industrie zur Herstellung von Pulvern, Granulat und Schüttgut.



Laserstrukturierte Gleitringdichtungen aus EKasic® Siliziumkarbid (links: radiallager, rechts: axiallager), werden z.B. in hochbeanspruchten Chemiepumpen, in magnetkupplungen für hermetisch dichte Pumpen sowie in Rührwerken für chemische und pharmazeutische Verfahren verwendet



Gasdichtungsringe aus EKasic® Siliziumkarbid werden zur Abdichtung von Kompressoren und Rührwerken für die Erdöl- und Gasverarbeitungsindustrie eingesetzt

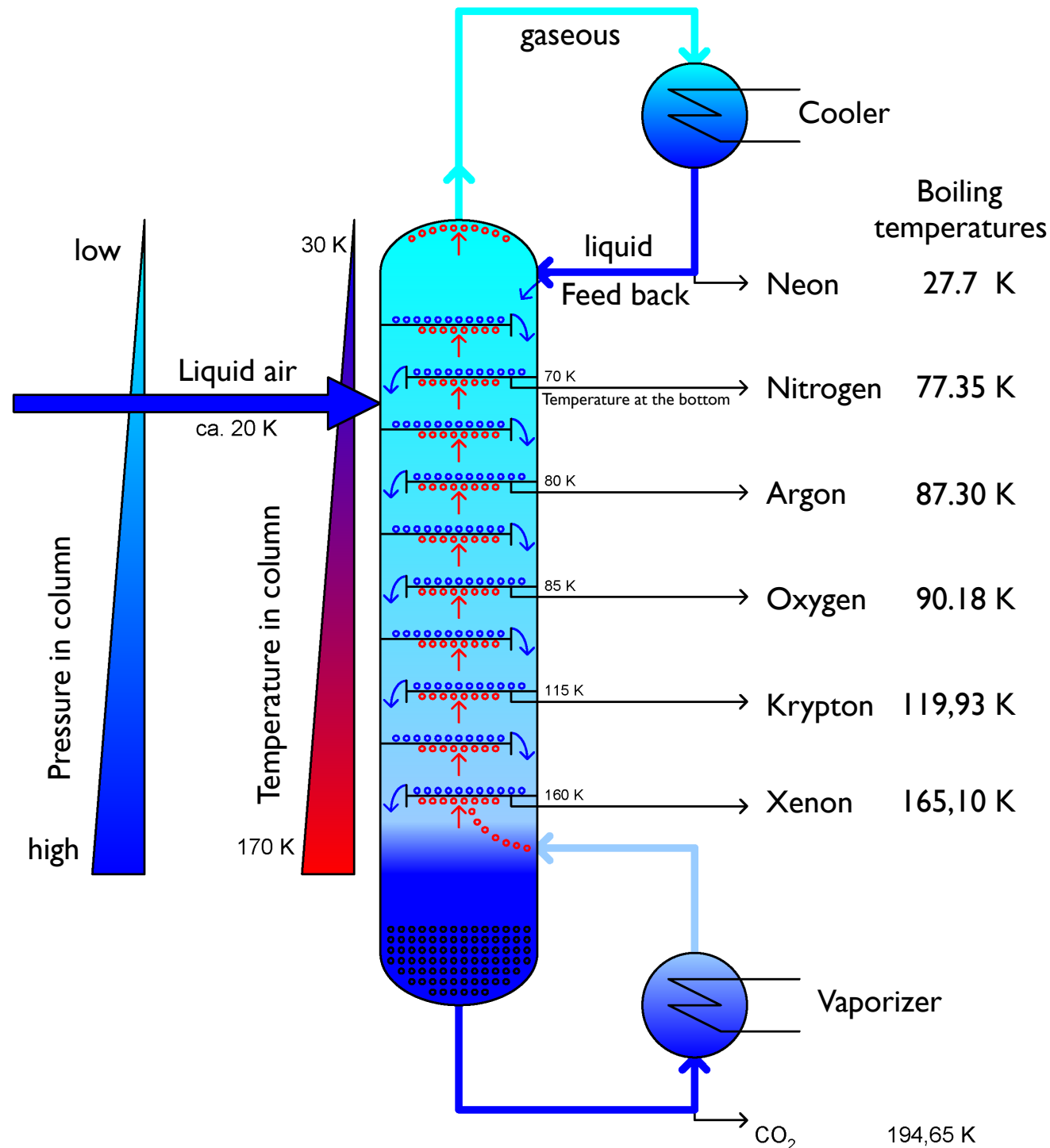
Salt Reprocessing with the PPU

(Pyrochemical Processing Unit)

Well-known techniques from the industrial chemistry:

Partitioning of the salt components by distillation.

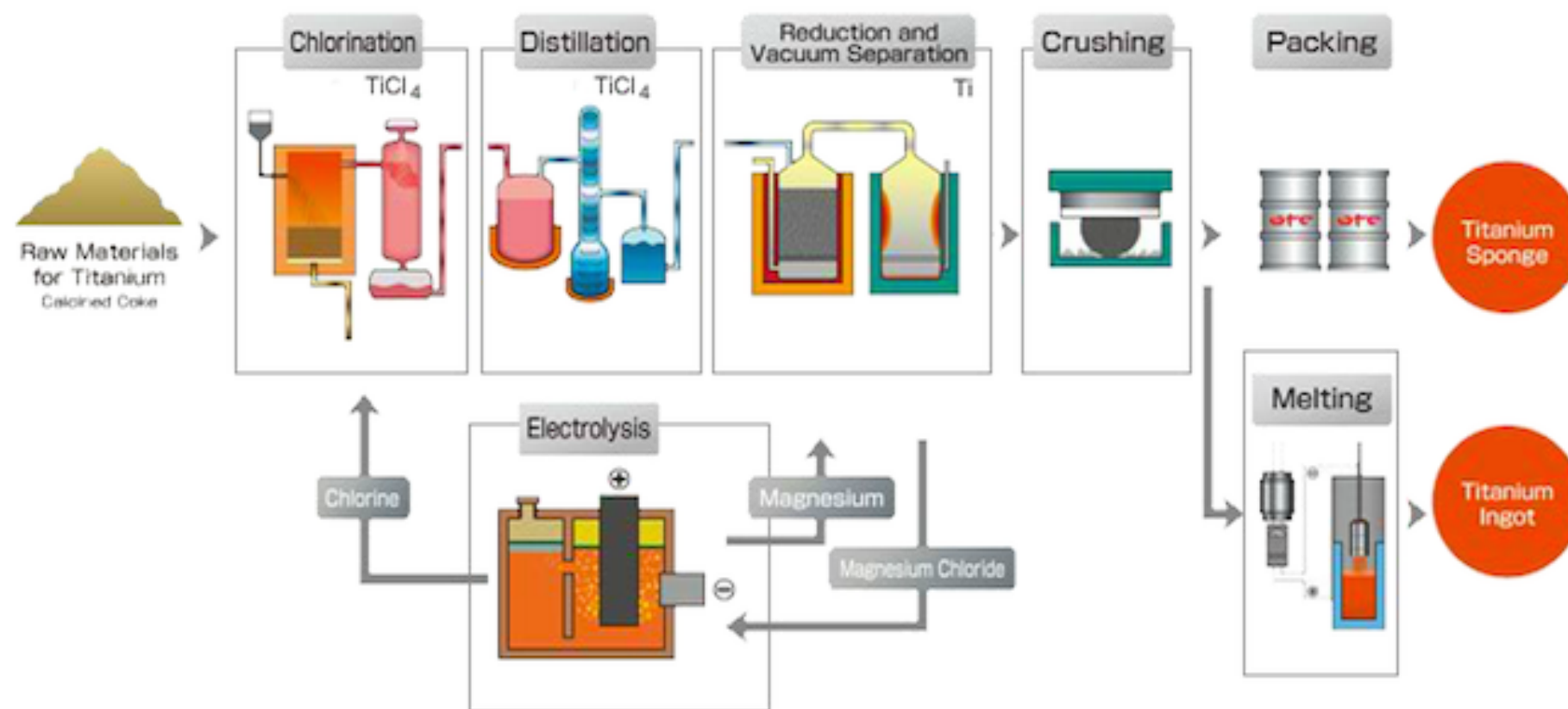
No wet-chemical techniques with large amounts of medium-active chemical waste



Example: Rectification of liquid air

The Kroll Process

- State-of-art since a long time
- Used for all metals of the Ti group (e.g. Zr, Hf)
- Developed in the 1930ies
- Titanium ore is reduced and chloridized
- Distillation to the single chlorides at clearly above 1000 °C to 1400 °C
- Then reduction of TiCl_4 with alkaline metals
- High purity Titanium is sold for 10 \$/kg

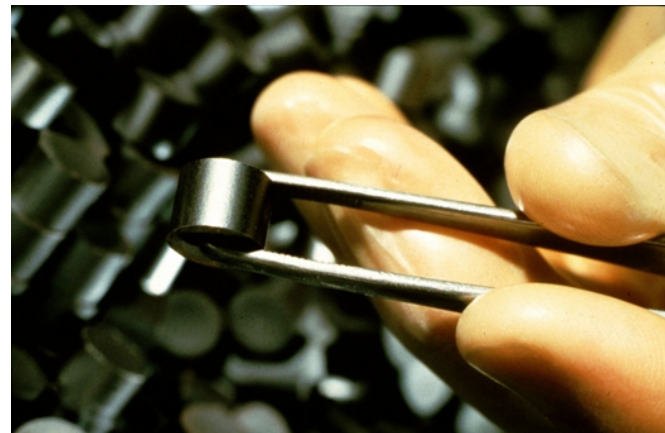


Ti-Produktion at Osaka Titanium Technologies
<http://www.osaka-ti.co.jp/e/>

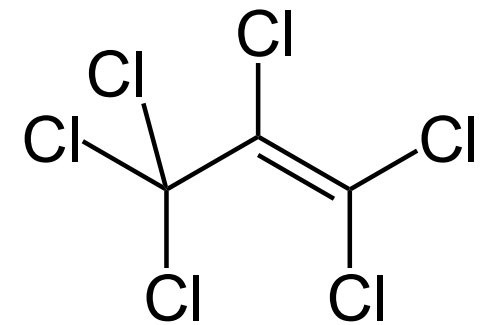
Waste Pre-conditioning

Treatment of fuel elements

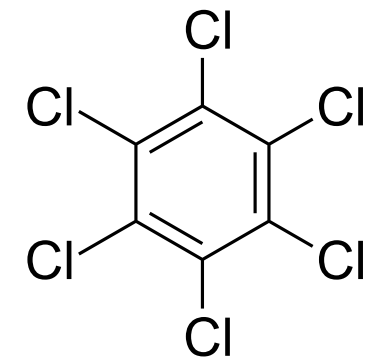
- Opening of the bundles and tubes
- Separation of pellets from tube
- Cleaning of the tube material
- Redox reaction with chlorocarbons
- Distillation / Infusion PPU



Hexachloropropene



Hexachlorobenzene



Uranium dioxide

Uran trichloride + carbone oxides

P&T strategy with the DFR

Step 1

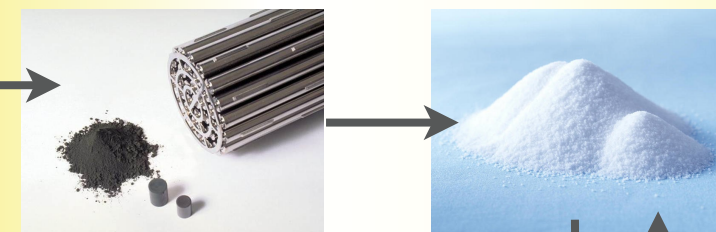
Pre-conditioning plant

All amounts are for the heavy-metal (HM) part. Tons are metric tons

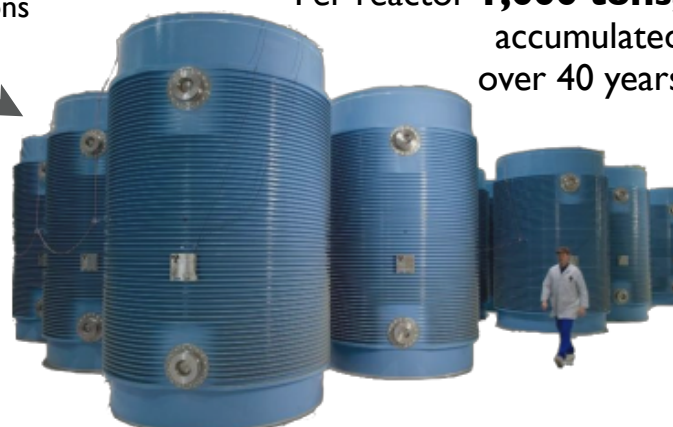
Per reactor **1,000 tons**, accumulated over 40 years

25 tons / year, in case the reactor is still in operation

- Dismantling of the fuel elements
- Chopping of the pellets
- Conversion of the oxides into chlorides



Today's pressurized water reactor (1400 MW_e)



Chloride salts

Chlorine (reprocessed)

Mass reduction:
Factor 33 within 2 years

Step 2 Partitioning plant

Fission product storage

30 tons
Thereof 3 tons long-lived
Max. 300 years of storage time

Energetically **not usable**

PPU

Max. throughput:
500 tons / year

Actinide storage

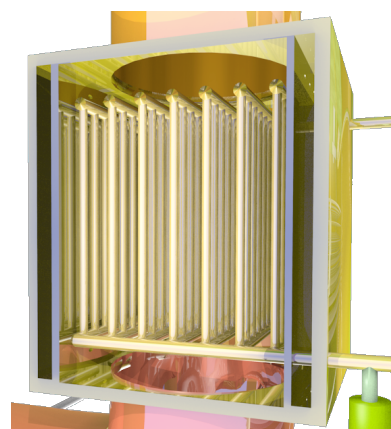
960 tons Uranium
9 tons Plutonium
1 tons minor Actinides

Energetically **usable**

Step 3 Transmutation plant

DFR core
3 GW thermal

- Energetic usage:
- High temperature heat
 - Electricity generation



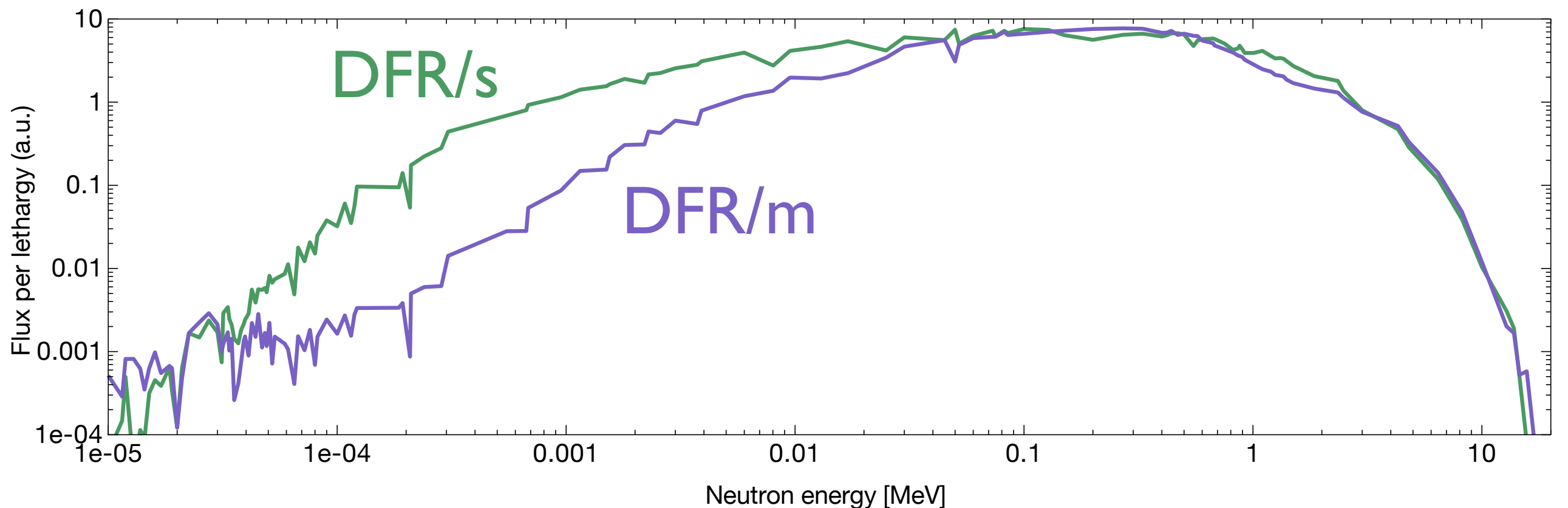
Incineration of actinides
in the DFR core

Consumption: 1.2 tons / year
(Range of the waste for many hundred years)

Optional additional transmutation of up to
0.3 tons / year long-lived fission products

DFR: Metal vs Salt Fuel

3 GW _{th} , 1.5 GW _{el}	DFR/s	DFR/m
Fuel	Undiluted Act.-Cl3 salt, density 3500 kg/m ³	Pure eutectic with >70% actinides, density 16500/9500 kg/m ³
Critical with	20.5/18.3/15 HM mass-% reactor-Pu/ ²³⁵ U/ ²³³ U	8.4/8.8/8.8 HM mass-% reactor-Pu/ ²³⁵ U/ ²³³ U
Blanket	cylindrical, thickness 1 m, height 5.5 m (100 m ³)	No blanket, thicker reflector (Pb coolant) 0.5 m
Structural Material	pure high-density SiC, 3210 kg/m ³	ZrC-20mass%TiC, 6100 kg/m ³



DFR neutronic results

Assumed fuel salt density: 3.5 g/cm³, no burnup, ²³⁸U fast fission ignored for CR calculation
 Composition: ³⁷Cl, 68.5 mole-%, actinides balance (material below and ²³⁸U/²³²Th balance)

Fissile Material	Type	Fuel Enrichment	ν	CR
U-235	Salt	~19.5%	2.47	< 0.9
Pu-239	Salt	~17%	2.90	< 1.2
Reactor-grade Pu ^a	Salt	~22%	2.92	1.2 (1.25)
U-233	Salt	15.5%	2.52	1.1
Reactor-grade Pu ^a	Metal	< 9%	2.92	> 1.6

$$CR = \frac{\text{Cap}(^{238}\text{U}/^{232}\text{Th}) + \text{Cap}(^{238}\text{Pu}) + \text{Cap}(^{240}\text{Pu}) + \text{Cap}(^{242}\text{Pu})}{\text{Total fission and capture } (^{233}\text{U} + ^{235}\text{U} + ^{239}\text{Pu} + ^{241}\text{Pu})}$$

a) 45 GWd/t Pu

DFR: Metal vs Salt Fuel

	DFR/s (hexag.)	DFR/m (hexag.)
fission zone DxH [m]	2.8 x 2.8	3 x 2.6
outer / inner tube diameter [mm]	18 / 15	24 / 20
Pitch-to-diameter ratio	1.25	1.25
mean linear power density [W/cm]	850	1250
mean temperatures fuel inlet / outlet [K]	1270 / 1540	1350 / 1650
temperatures coolant inlet / outlet [K]	1030 / 1300	1070 / 1370
conversion ratio U-Pu / Th-U cycle at start	> 1.2 / 1.1	1.7 / 1.1
²³⁴ U, ²⁴⁰ Pu, ²⁴² Pu burnable? (resp. CR)	no (1.2 / 1.1)	yes (2.1 / 1.3)
²³⁸ U fast fission / all fission	6%	20%
fiss. zone volume [m ³] (fuel fraction)	21 (32%)	23.6 (32%)
fuel / coolant velocity (m/s)	1.2 / 3.6	0 / 2.6

DFR/s Thermohydraulics

DFR Core

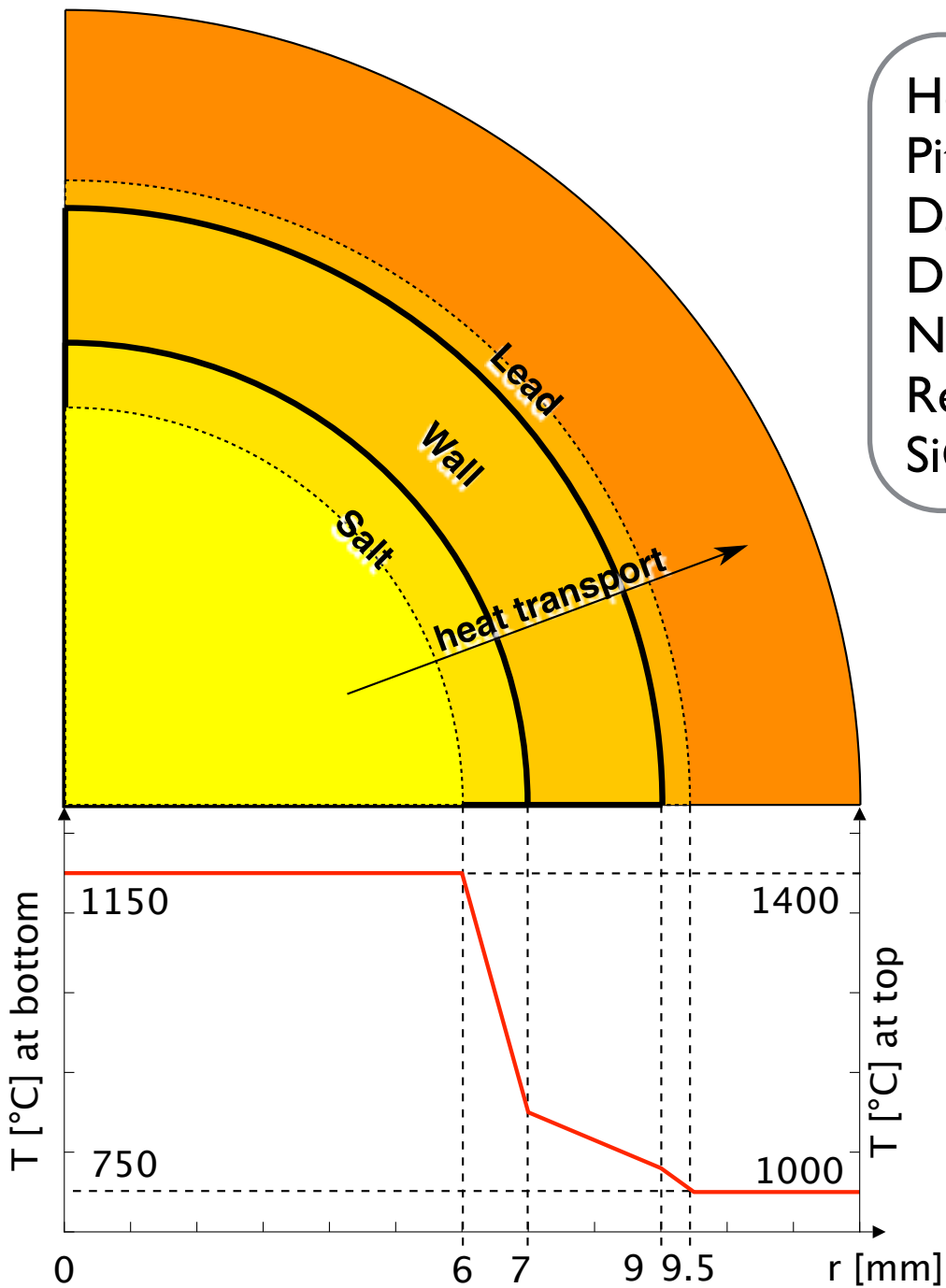
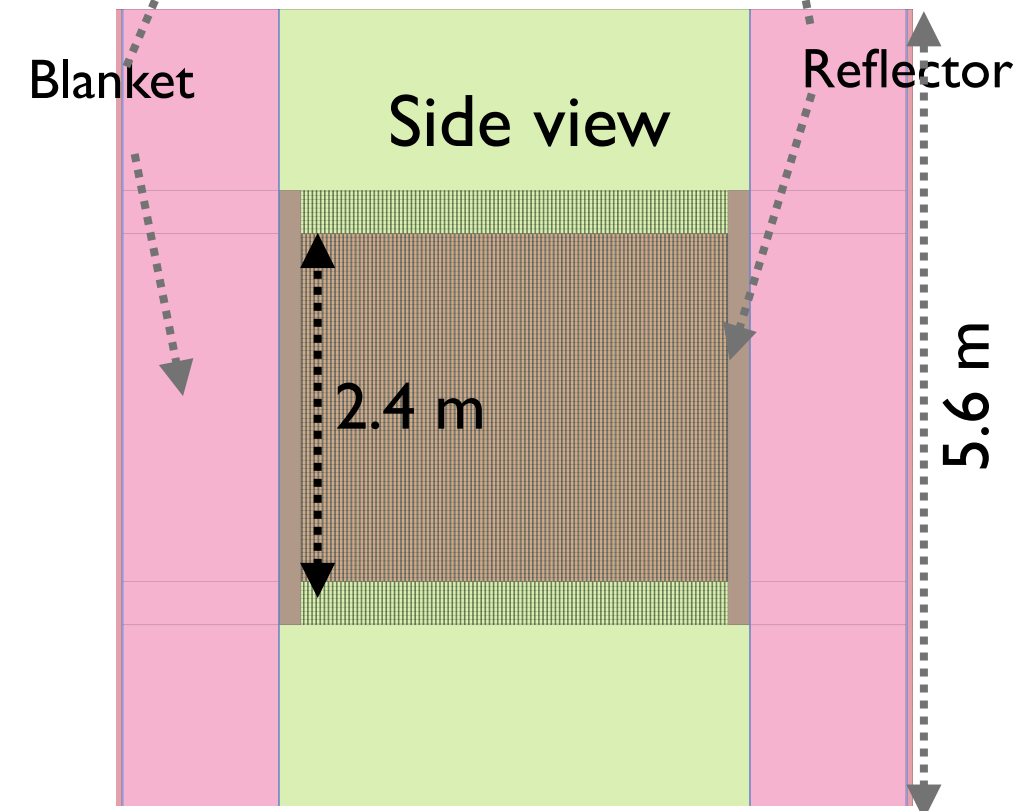
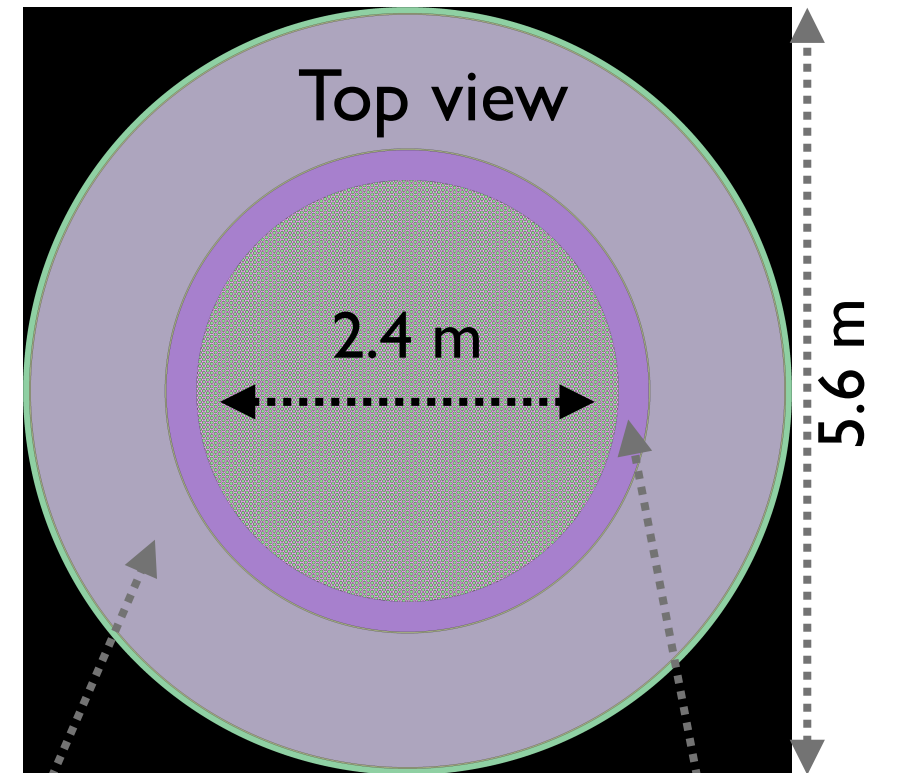
Hexagonal lattice
 Pitch-to-diameter (P/D) ~ 1.15
 $D_o = 19(18)\text{mm}$
 $D_i = 14(15)\text{mm}$
 Nominal power: 3 GW_{th}
 Reflector thickness: 0.2m
 SiC density 3.2 g/cm^3

UCI3 (theor. !):

3.5 g/cm^3
 2 W/(m K)
 0.5 mPa s
 400 J/(kg K)
 $v = 1\text{ m/s}$
 $Re = 140,000$
 $Pr = 0.09$

Lead coolant:

9.7 g/cm^3
 23 W/(m K)
 1 mPa s
 140 J/(kg K)
 $v = 4\text{ m/s}$
 $Re = 360,000$
 $Pr = 0.006$



Fuchs-Nordheim and simple reactivity investigations (1st pulse)

Conditions for application:

- $\rho \gg \beta \rightarrow$ Inserted reactivity much larger than delayed neutron fraction
- Adiabatic fuel heating (always given in fast reactors and high reactivity insertion rate)
- Reactivity coefficients constant over relevant temperature range

	LWR	DFR
Inserted reactivity ρ	0.01	0.01
β (reactor-grade Pu)	0.0035	0.0035
Reactivity coefficient α	-3 pcm/K	-50 pcm/K
Fuel heat capacity C_p	40 MJ/K	9 MJ/K
Prompt neutron lifetime	60 μ s	6 μ s (!)
Max. temp. change ΔT_{fuel}	400 K	26 K
Temperature after pulse ΔT_{fuel}	200 K	13 K
Pulse duration Δt_{peak}	40 ms	4 ms
Pulse energy E_{peak}	20 GJ	0.2 GJ
Pulse power Δp_{peak}	1 TW	0.1 TW

$$\Delta P_{\text{peak}} = C_p(\rho - \beta)^2 / (2\Lambda\alpha)$$

$$\Delta T_{\text{fuel}} = 2(\rho - \beta) / \alpha$$

$$\Delta T_{\text{fuel}} = (\rho - \beta) / \alpha$$

$$\Delta t_{\text{peak}} = \sim 4\Lambda / (\rho - \beta)$$

$$E_{\text{peak}} = 2C_p(\rho - \beta) / \alpha$$

- DFR reactivity coefficient (almost) independent on fuel composition (liquid fuel)
- LWR temperature changes far too high \rightarrow control rods needed

Dual Fluid Reactor

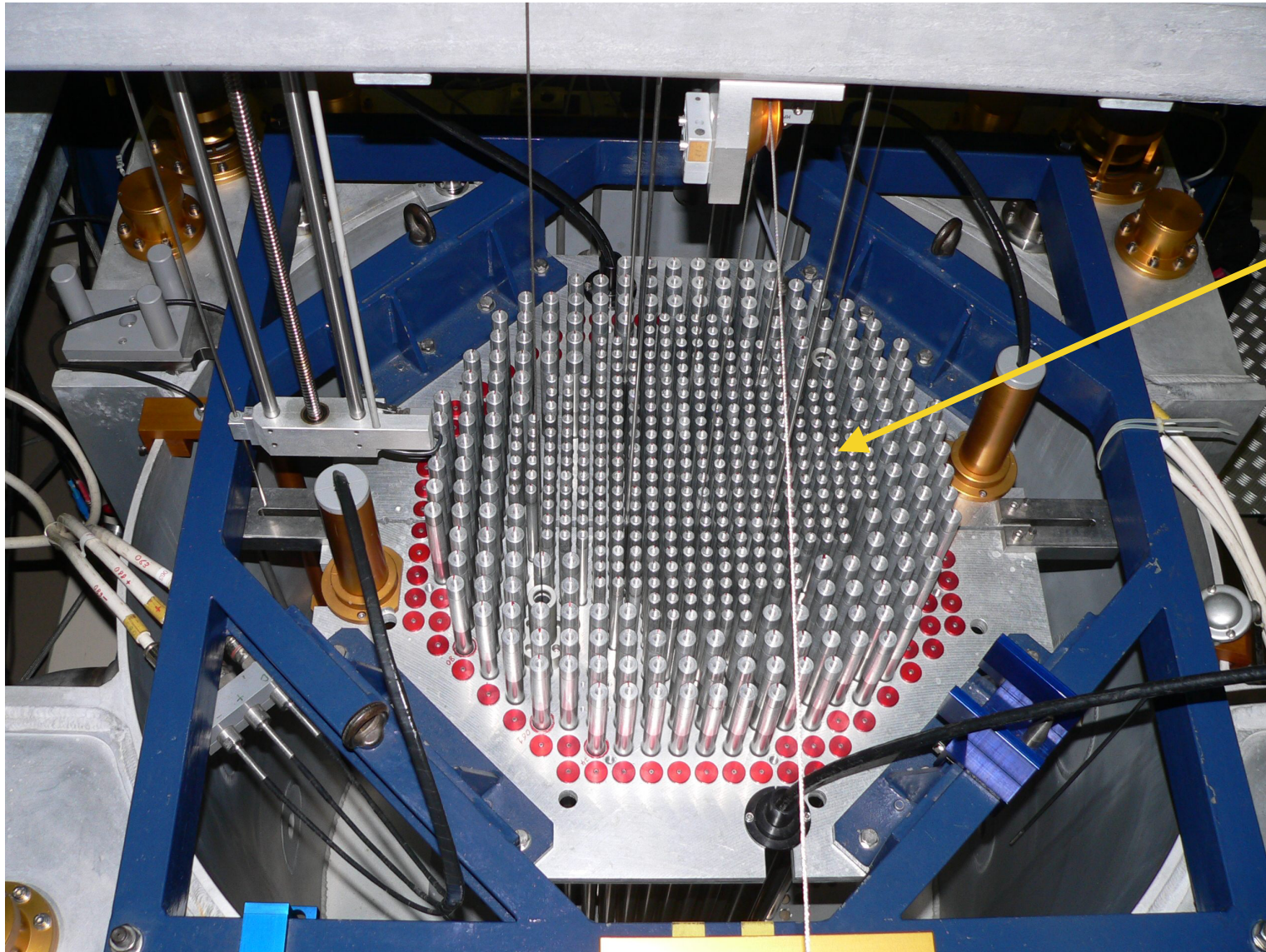
Applications, economic and financial aspects



Institut für Festkörper-Kernphysik Berlin

Institute for Solid-State Nuclear Physics Berlin

Today's reactor designs



Almost all are water-moderated and based on **solid fuel rods:**

- Expensive external fuel cycle
- Using **only 1%** of the mined Uranium
- **99% waste** that needs geological storage
- Low power density

They also work at **high pressure**

Today's nuclear reactors are more effective than other power generating systems, but **nuclear power can do much better!**

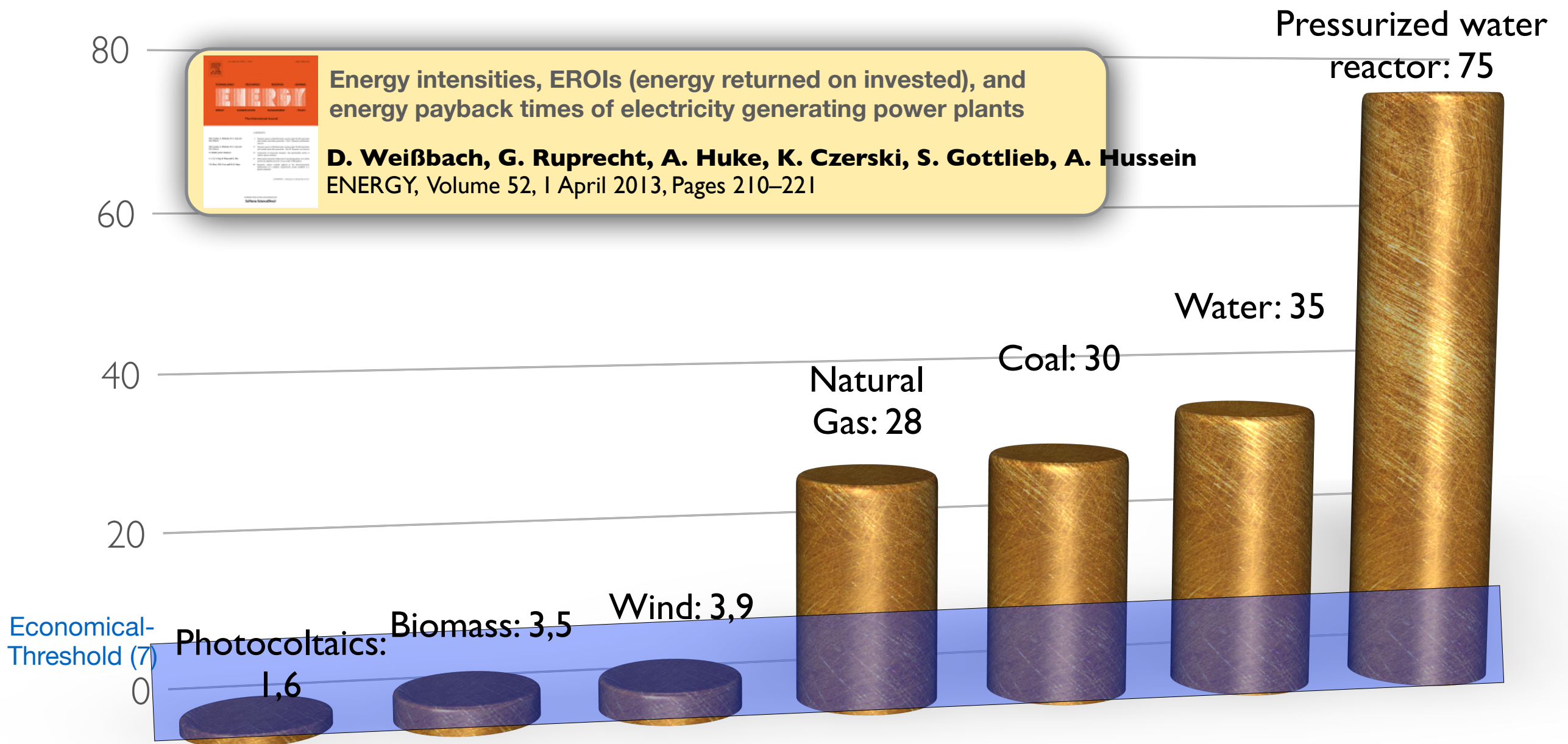
Energy Return on Invested - EROI

During the entire life-cycle of the plant:

Produced electricity E_{out}

divided by the expended energy (construction, operation, deconstruction) E_{in} .

$$EROI = \frac{E_{out}}{E_{in}}$$



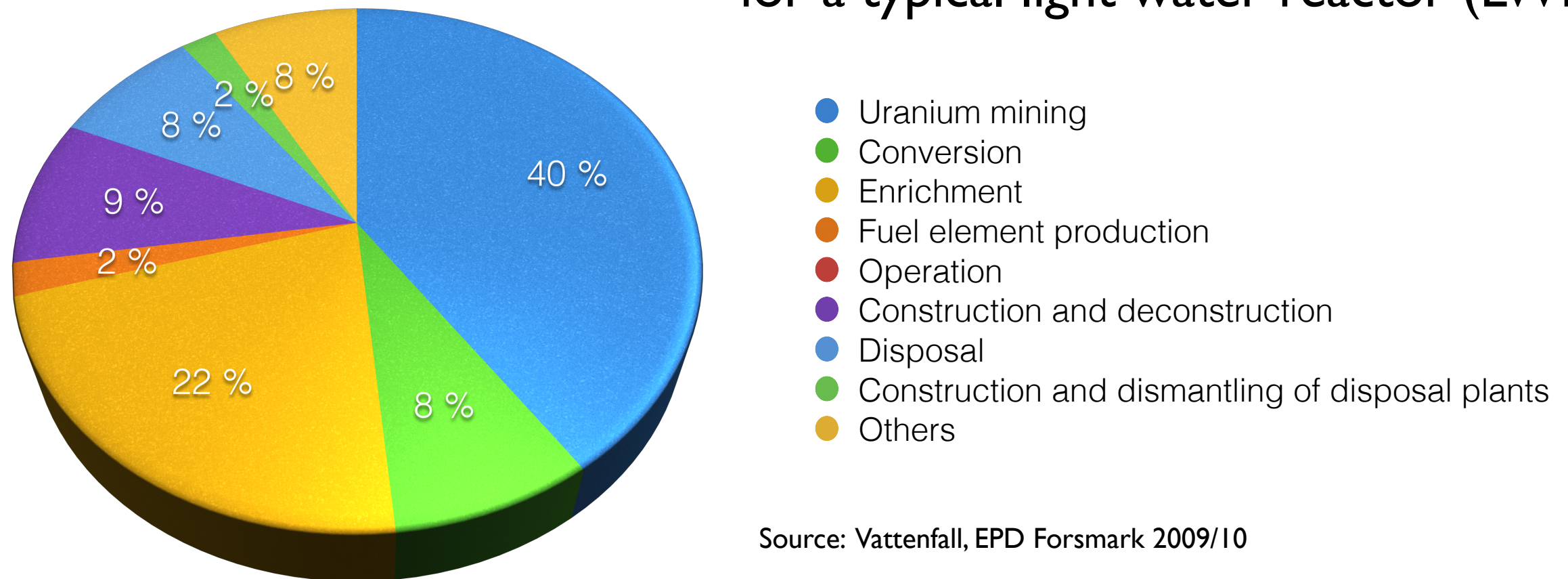
Is the EROI for PWRs large?

EROI	Energy release on combustion/fission
Pressurized water reactor: 75	Uranium nucleus: 200.000.000 eV
Factor 2,5	Factor 100 million
Coal power plant: 30	Hydrocarbon atom: 2 eV

What's going wrong here?

The expensive nuclear fuel cycle today

Contributions to the energy demand in the nuclear power production for a typical light water reactor (LWR)



EROI: 29 → 75 → 105 → 115

100% Diffusion → 17% Diffusion + Centrifuge (today) → 100% Centrifuge → 100% LASER

How efficient is the DFR

If these expenses are reduced to DFR level, EROI and costs change to...

	Enrichment		Uranium Supply			Dismantling						
		↑		↑		↑						
	LWR		Construction and Operation		Fuel Cycle		DFR/s	DFR/m				
EROI:	75	→	115	→	120	→	390	→	1000	→	2000	5000
Costs: cent/kWh overnight	2.7	→	2.3	→	1.5	→	1.1	→	0.8	→	0.65	?

From LWR to DFR.

Many steps are repealed or reduced, increasing the EROI and decreasing the costs.

DFR/m comes close to the theoretical limit of nuclear energy, dominated by the Uranium mining expense.

For comparison:

Wind and PV: 1-4
 Fossil fuels: 30
 Hydro: 35

Nuclear:

Today's LWRs: 75
 Theoretical limit: 10,000

Energy Efficiency of Power Plants

Efficiency by **EROI** (Energy Return on Energy Invested)
see Weißbach et al., *Energy*, vol. 52 (2013), pp. 210–221

Dual Fluid Reactor

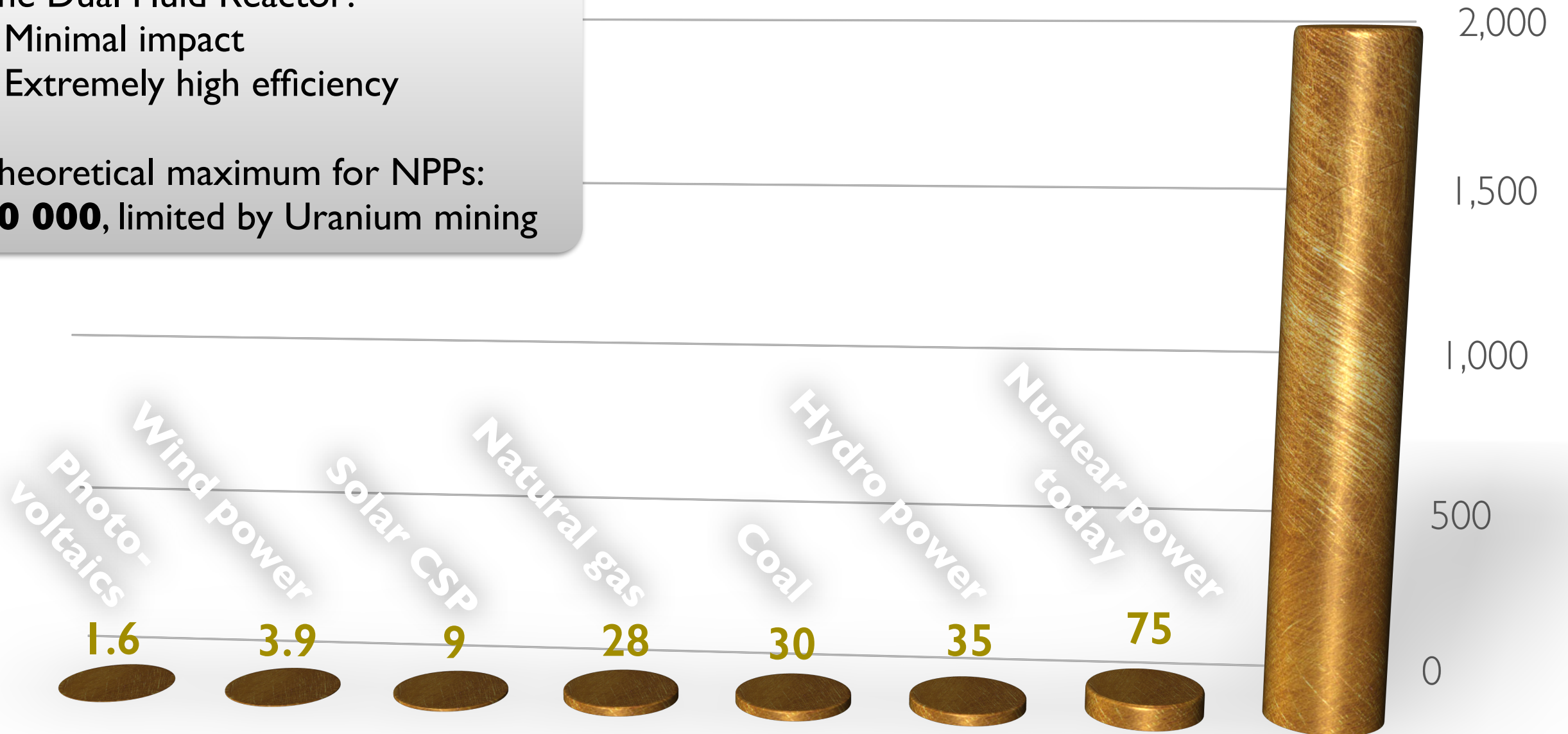
The Dual Fluid Reactor:

- Minimal impact
- Extremely high efficiency

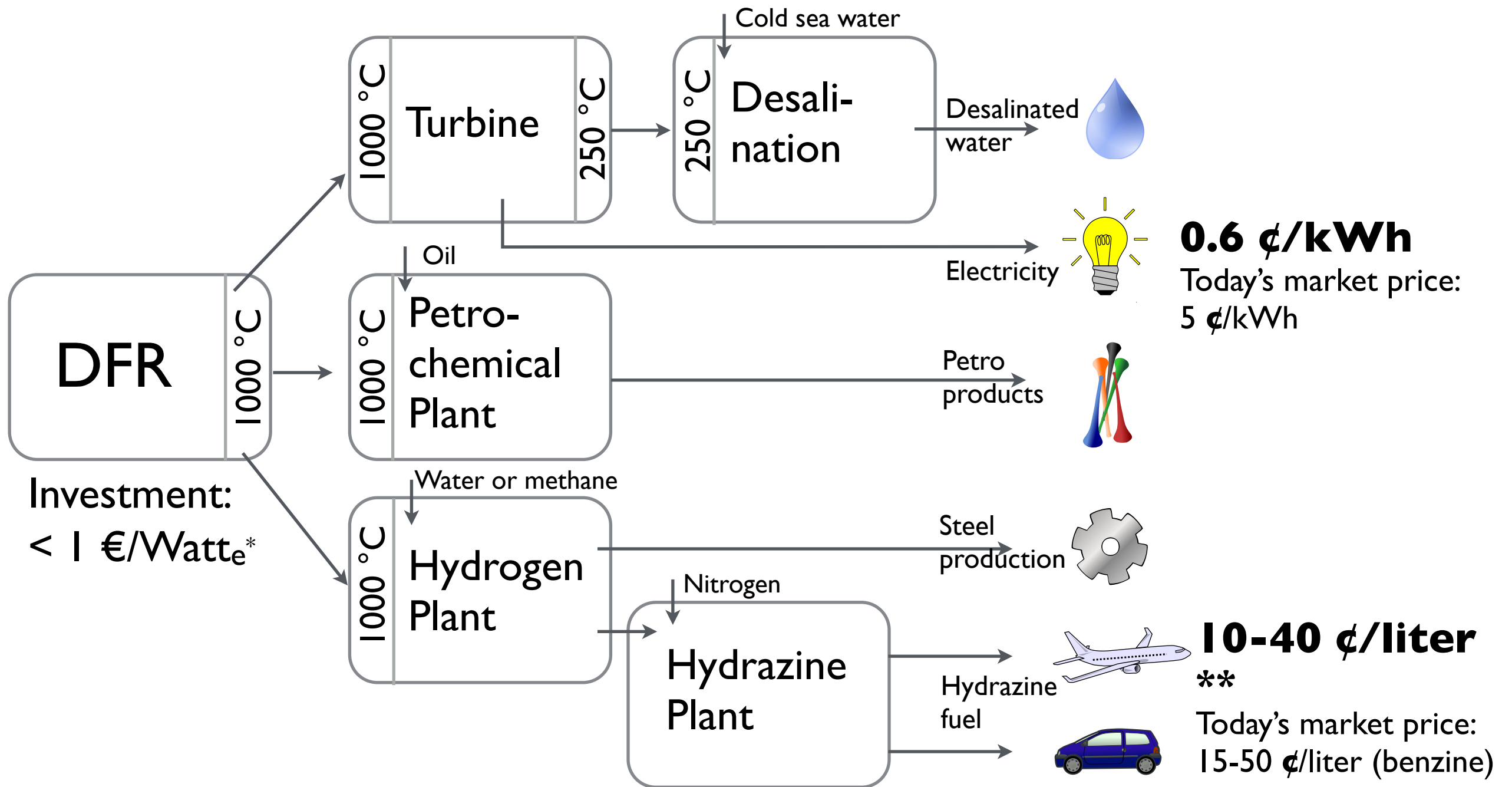
Theoretical maximum for NPPs:

10 000, limited by Uranium mining

2000



DFR applications



* Overnight costs
** Gasoline equivalent

Hydrazine-based fuel cell **electro mobility** also possible with 1.5 ¢/km* and ranges of more than 1,000 km

Hydrogen production

- Water dissociation by high temperature
- Hot-ELLY, KfA Jülich for THTR
- Sulfur Iodine process for VHTR (GenIV)

- Gasoline synthesis by coal hydration similar to crude oil reforming

- Lignite transport by ship to the NPP where they are anyway for cooling.
- Process heat by DFR from nuclear waste

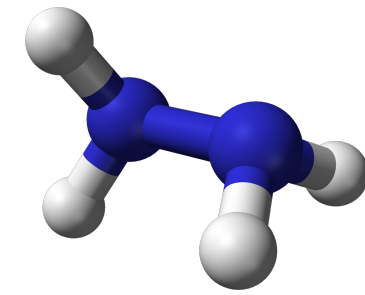
- Alternatively CO₂ usage from power plant exhaust



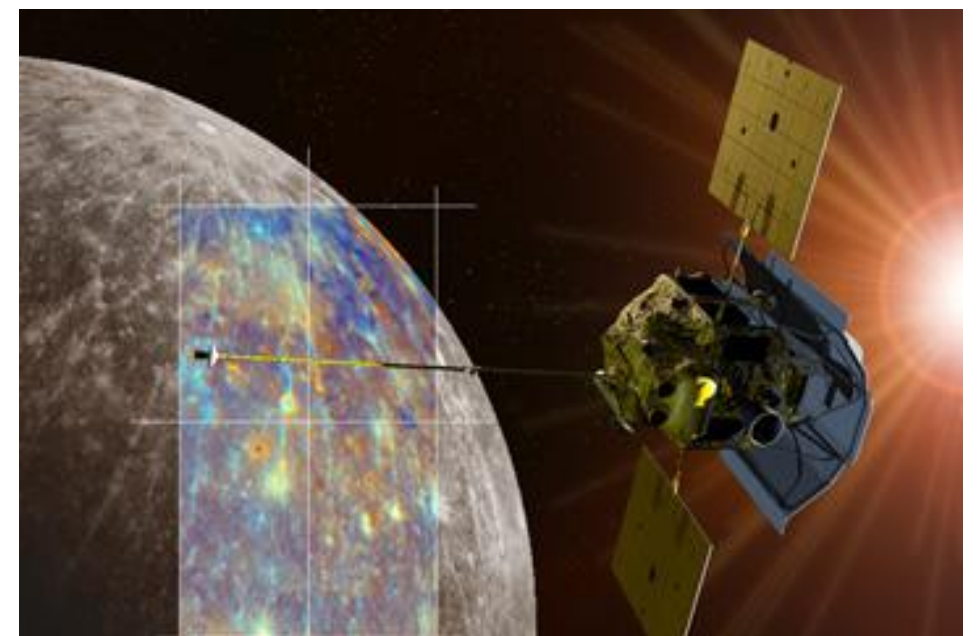
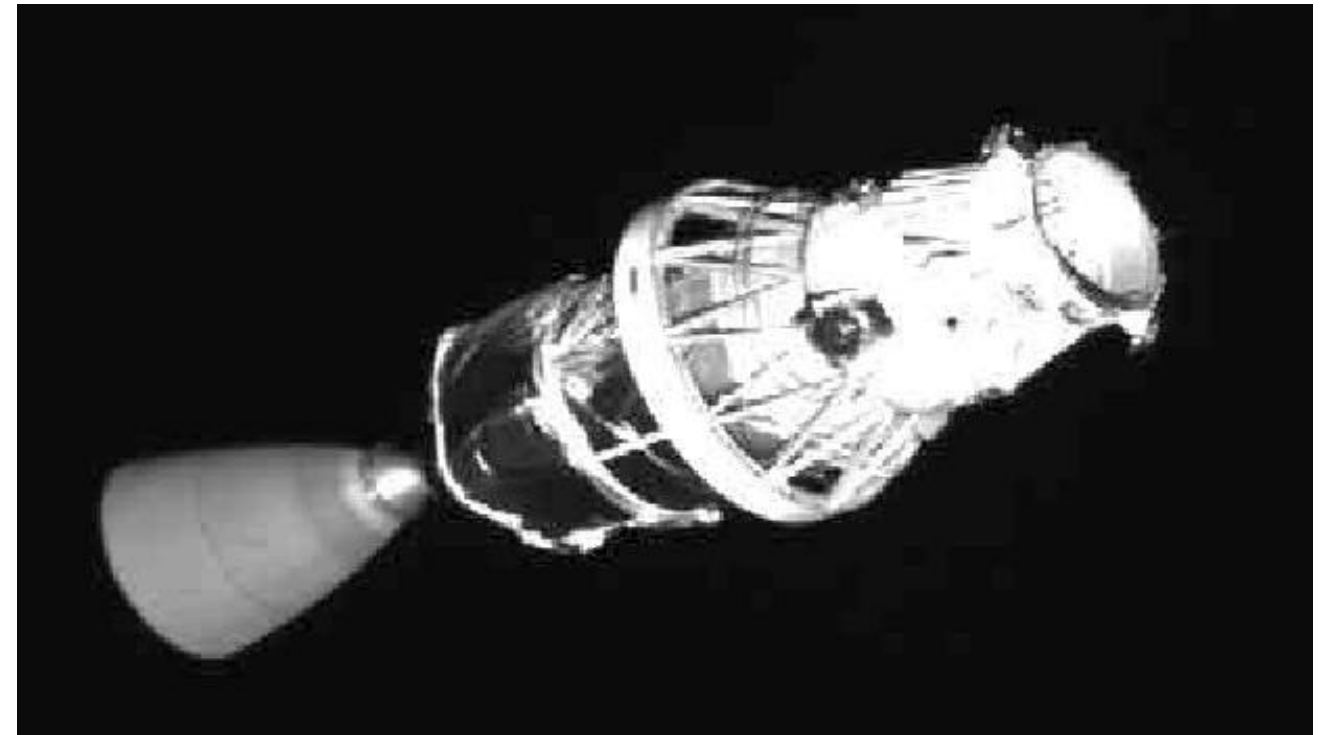
Bundesarchiv, B 145 Bild-FD50662-0023
Foto: Reinck | 29. August 1979



NtL: Hydrazine

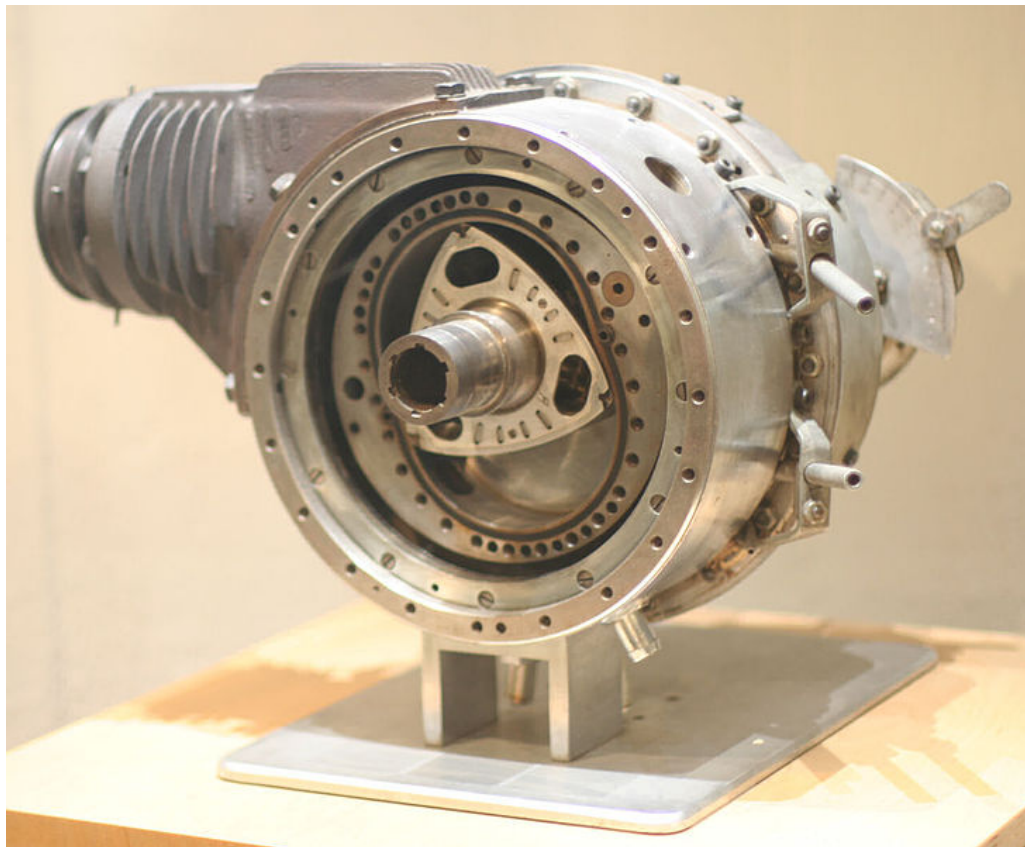
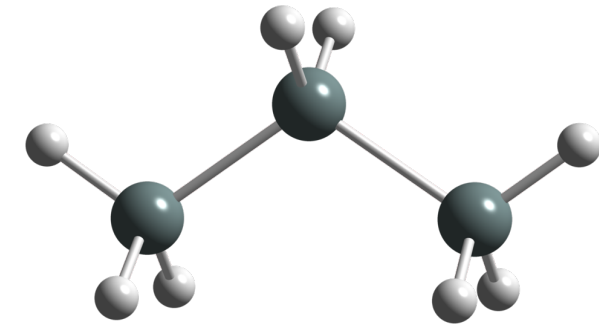


- Synthesis of ammonia from atmospheric nitrogen and water. (Haber-Bosch, SSAS)
Hydrazine synthesis (Pechiney-Ugine-Kuhlmann)
- Gasoline equivalent costs:
 - Ammonia: 20 c/l
 - Hydrazin: 40 c/l
- Fuel cell driven by nuclear-produced hydrazine is the only way of electro mobility with costs advantage against combustion engines:
 - Construction costs: Similar to Diesel engine
 - Fuel costs: 50% of today's gasoline

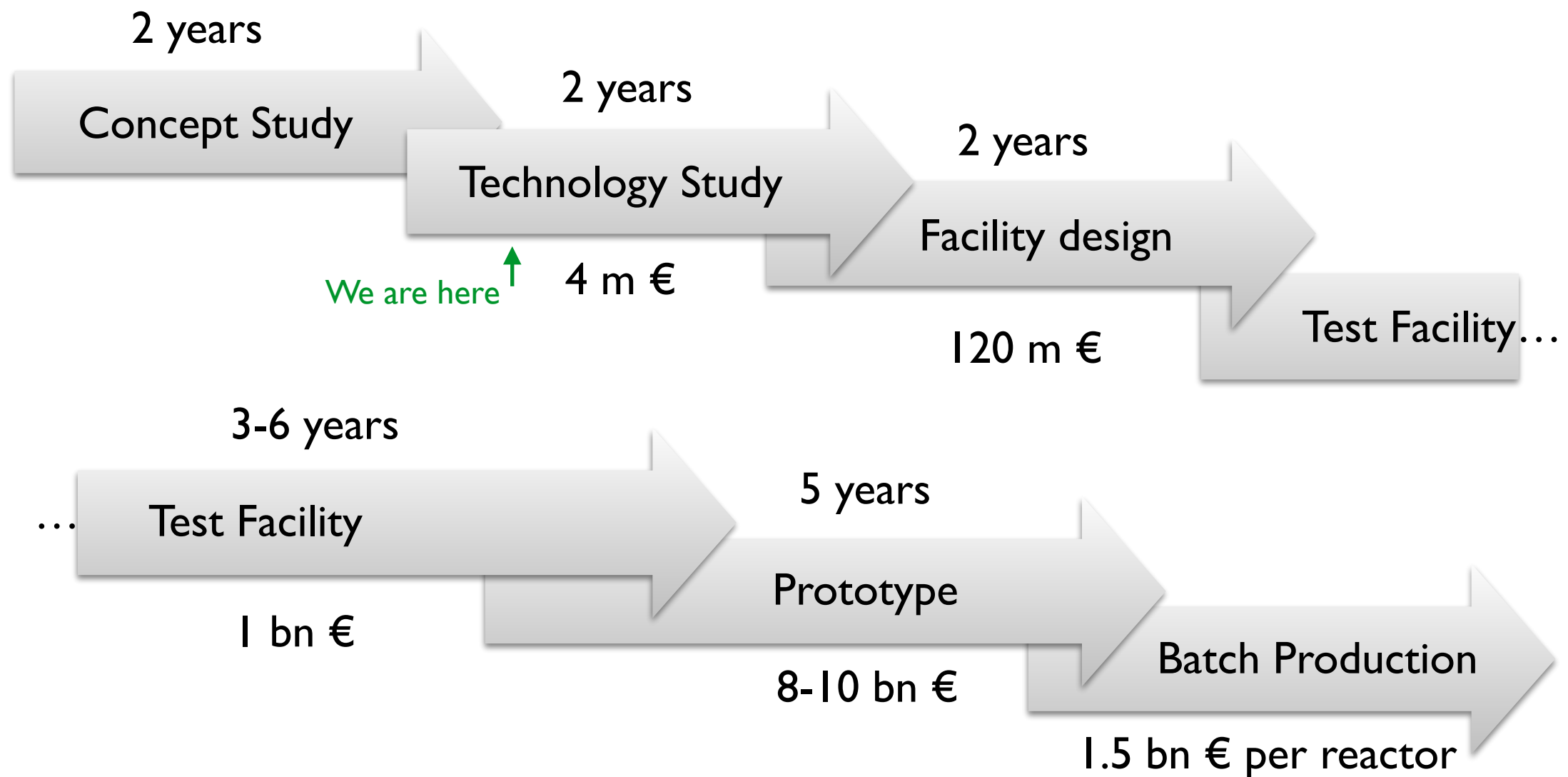


StL: Silane

- Müller-Rochow synthesis from quartz and water
- High power density
- Combusting at 1400 °C with air nitrogen
- Ideal as fuel for hypersonic aircrafts (SCRAM-jet), also for rockets
- Usage in Wankel engines for cars possible



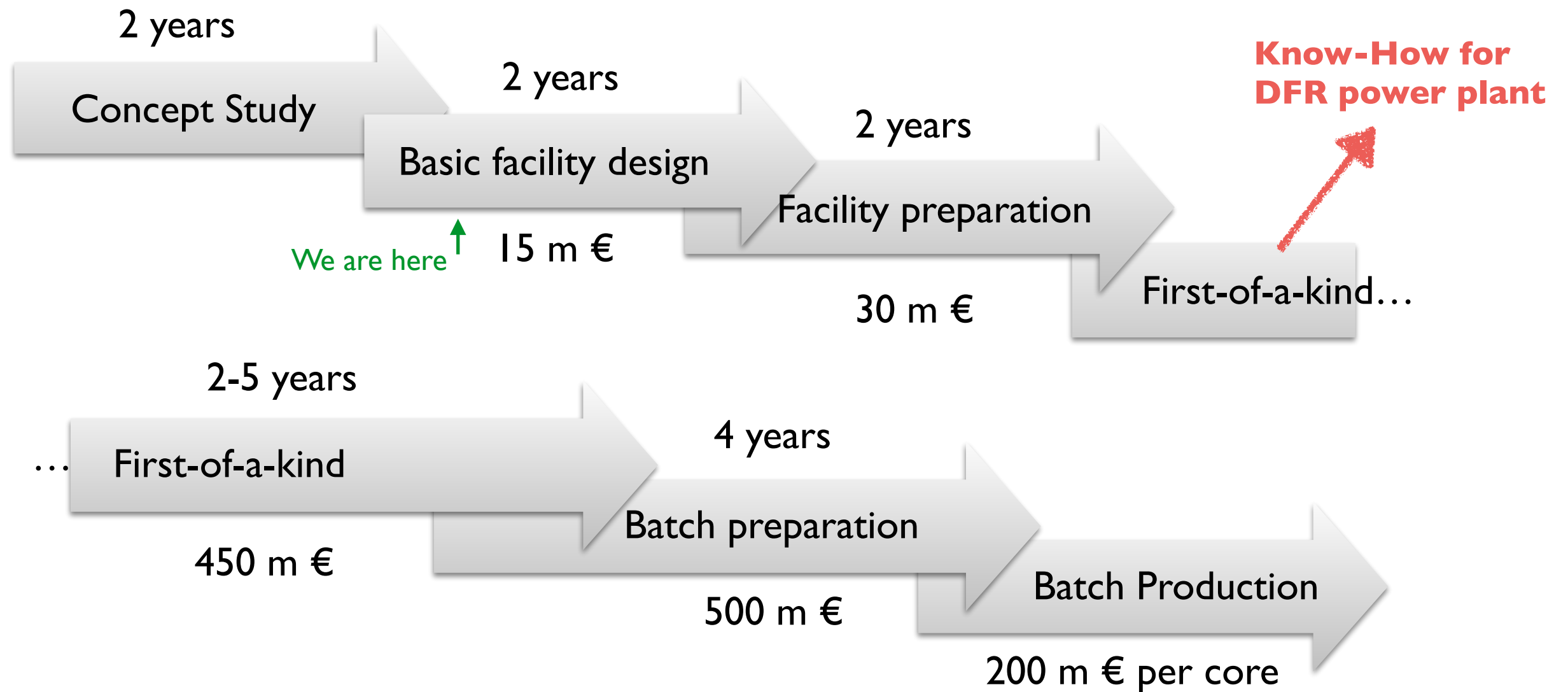
DFR development: Schedule and costs



Development of the DFR prototype: 10 years, 10 bn €
Serial type: 1.5 bn €

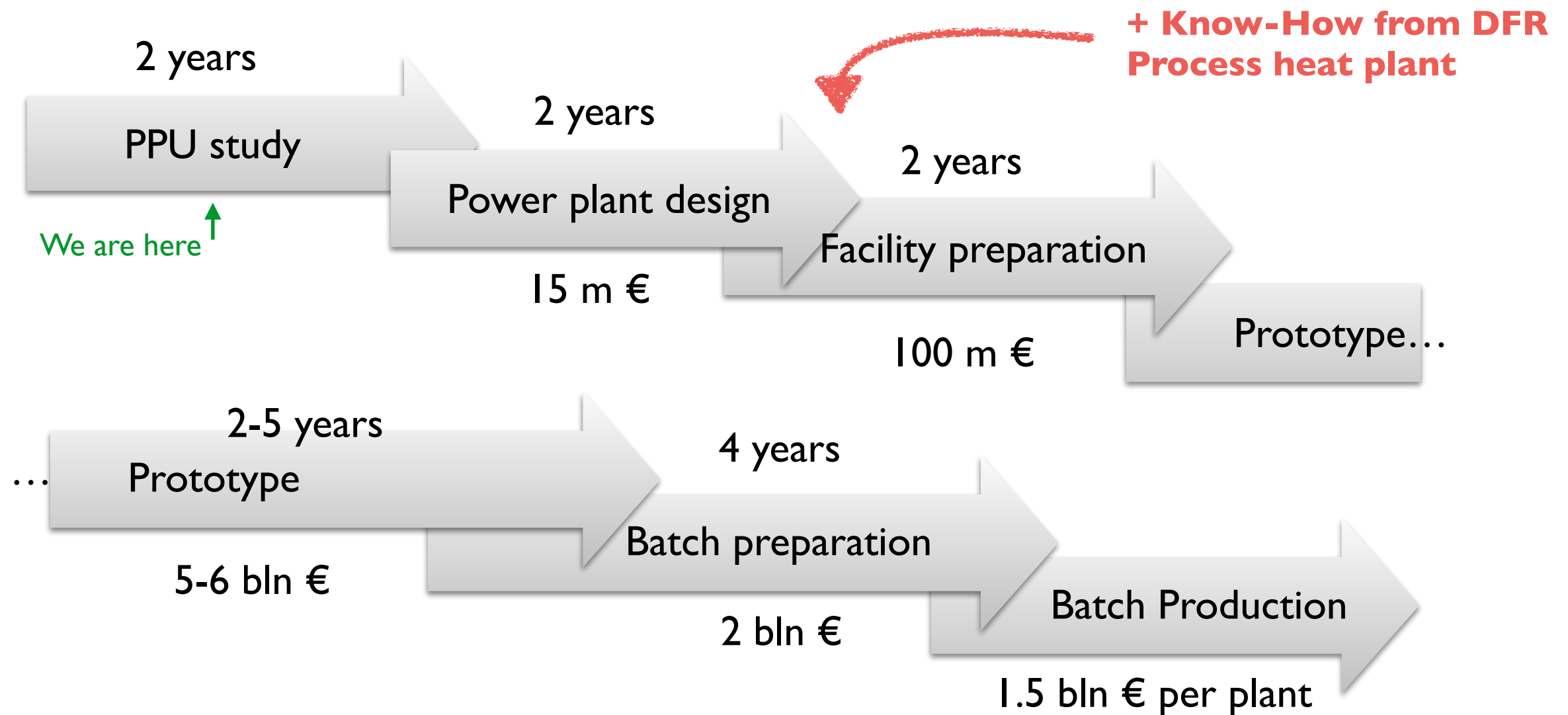
For comparison: Germany's Renewable Energy Law (EEG): **25 bn € per year**

DFR process heat plant development: Schedule and costs



Development of the DFR process heat plant ($300 \text{ MW}_{\text{th}}$):
 8 years, 1 bln €
 Serial type: 200 m €
 (scH₂O @ 1000 °C)

DFR power plant development: Schedule and costs



Development of the DFR power plant (1.5 GW_e):
8 years, 8 bln €
Serial type: 1.5 bln €

World Market

	Increase and Replacement*** until 2050 (PWh)	Plants*
Electricity	36	2740
Heat	40	3040
Transportation	42	1600
Process Heat	72	2740
Sum	190	14400
Investment Costs Process**		9110 Mrd. €
Investment Costs Electricity**		17340 Mrd. €
Investment Costs Total**		26500 Mrd. €

* 3 GW thermal or 1,5 GW electrical

** 2 € per Watt electrical, 0,7 € per Watt thermal

*** For transportation and heat full transition to synthetical fuels (with fuel cells) or electricity, for electricity and process heat 80% replacement of old plants

Key properties of the DFR

- Adiabatic power plant: No external fuel cycle needed
- Investment costs: 1 €/Watt^a → Comparable with coal power plants
- Energy efficiency (EROI) 20 times as high as for pressurized water reactors
- Electricity production costs: 0.6 ¢/kWh^a. Per serial DFR annual profit of 300 mio. € possible
- Oil-equivalent fuels can be produced for 50 US\$/barrel^{a,b} (0.3 €/liter^b)
- Electromobility based on hydrazine fuel cells possible with 1.5 ¢/km^a and ranges of more than 1,000 km

Alle costs are based on today's energy mix.

When the entire economy changes to DFR technology the costs further drop to the ratio of the EROIs.

a) Overnight costs

b) Energy equivalent

Supporters always welcome!



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