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Estimation of Tritium Recovery from a Molten Salt Blanket of FFHR

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1. Background study :

Characteristics of Flibe as a self-cooled blanket

- Advantages

- (1) liquid blanket, low reactivity with air or water
- (2) low MHD effect
- (3) TBR >1 with addition of Be and low tritium solubility
→ FFHR or HYLIFE-II

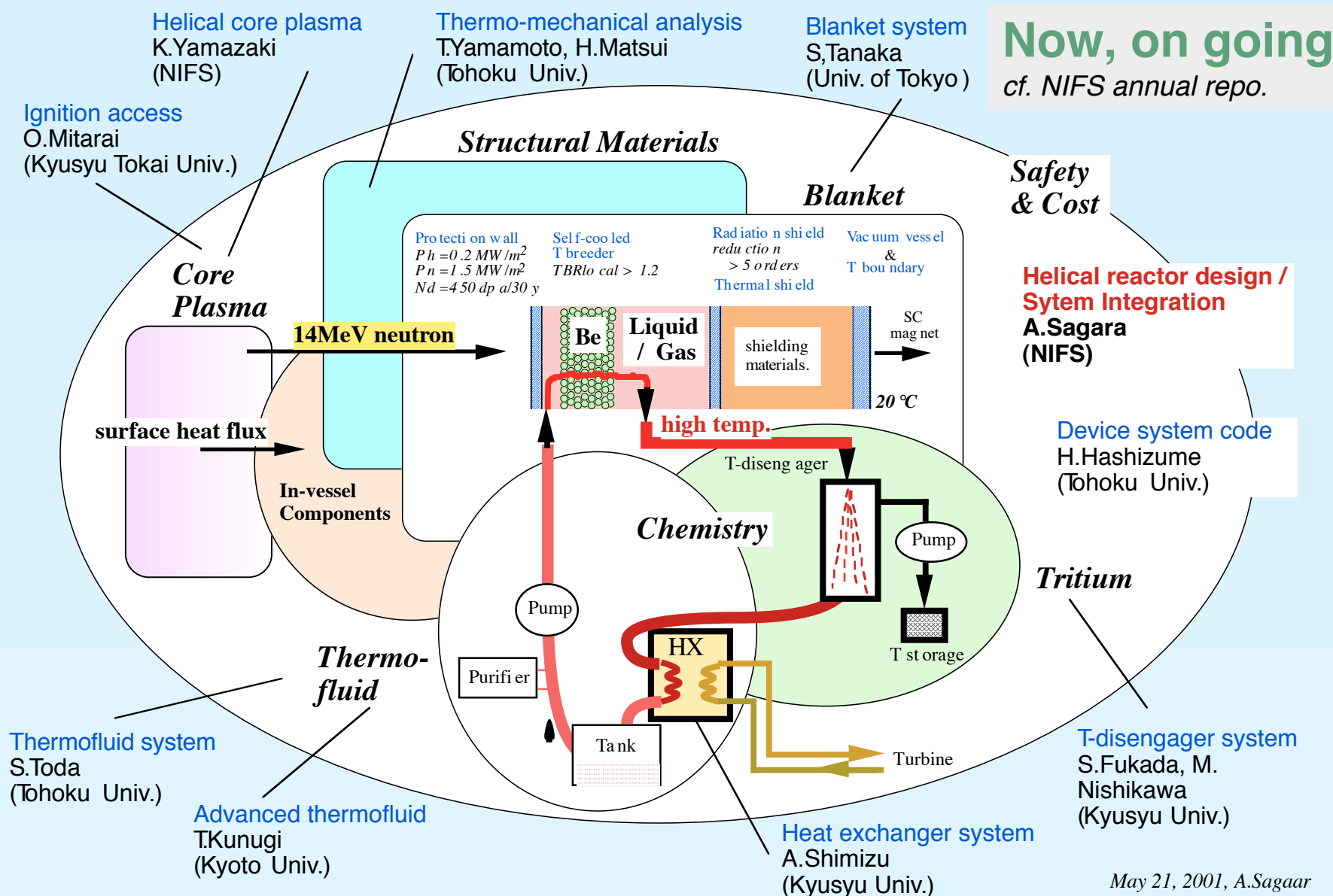
- Disadvantages

- (1) less Flibe-facing structural materials under neutron irradiation
- (2) unknown tritium behavior or recovery in Flibe is not studied.
- (3) possibly high tritium leak to the environment
- (4) issues of Be safety → JUPITER-II

____ Relating to this study



Design activities in NIFS Collaboration



May 21, 2001, A.Sagaar

1. Background study :

Tritium recovery from Flibe relating to the things

(1) Physical Properties (Underlined properties are not available or not reliable)

- Tritium solubility (TF or T_2)
- Tritium diffusivity, permeation coefficient or mass-transfer coefficient
- Viscosity, thermal diffusivity or heat-transfer coefficient
- Phase diagram, vapor pressure, free energy change of fluoride formation

Estimated from H_2 , D_2 or other data

(2) Affecting factors

- Chemical forms of tritium in Flibe , T_2 , TF or T_2O , or Flibe purification
- REDOX control condition, change from TF to T_2
- Corrosive HF or TF in Flibe
- Isotopic effect and coexistent components

to use purified Flibe and REDOX control by Be

1. Background study : Necessary conditions imposed on apparatus of tritium recovery from Flibe

- All tritium generated in a blanket is recovered
- Very low tritium leak to the environment
- Simple apparatus and easy to operate
- Operation over melting temperature
- High operation safety and material compatible with HF



- Operated under low water vapor condition



→ Permeation window, He-Flibe counter-current column,
Vacuum disengager

2. Design study :

Design of tritium-recovery apparatus for FFHR

Assumptions

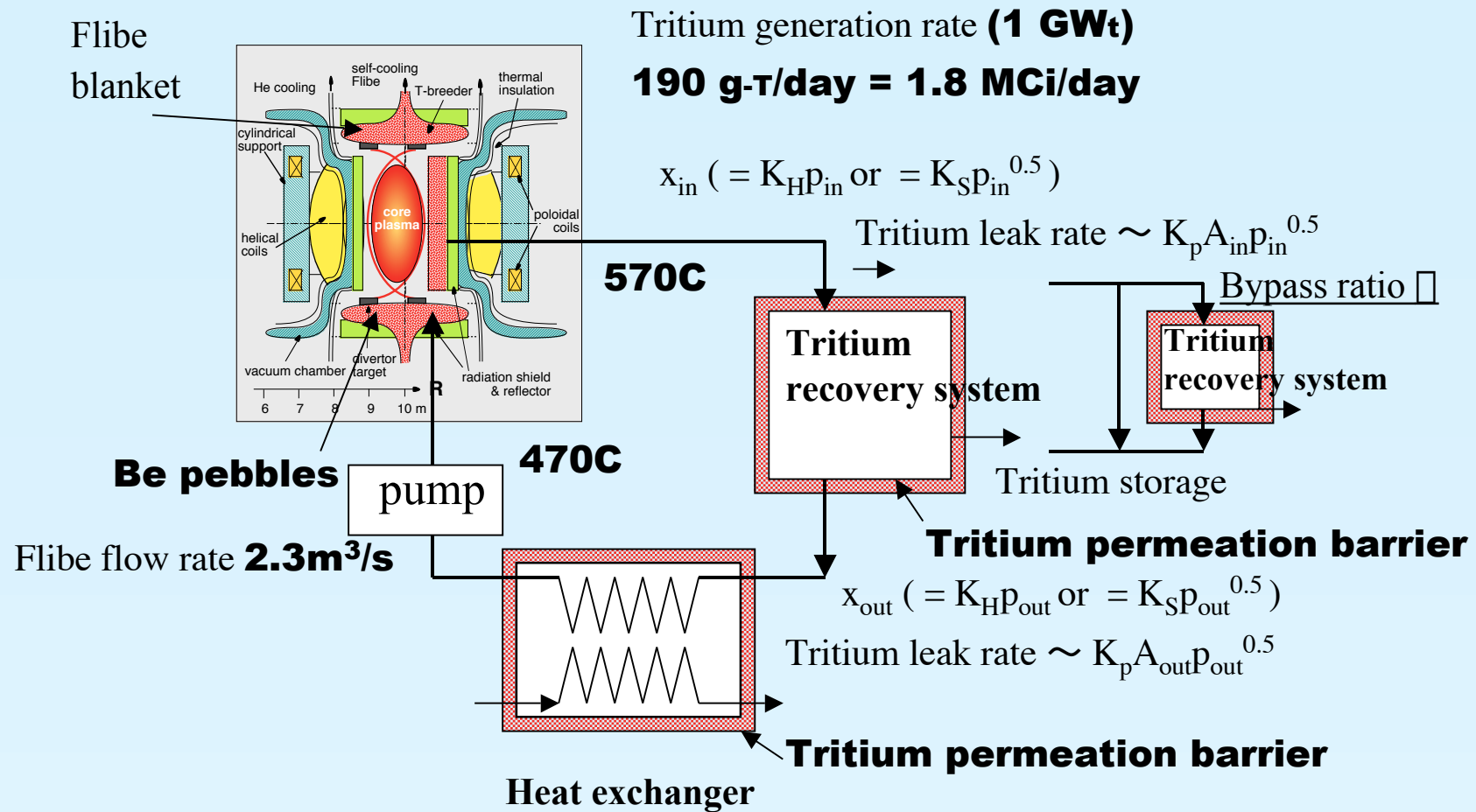
- Based on FFHR design
- Steady-state operation
- Tritium generated in a blanket is transferred by Flibe and is recovered by a tritium recovery apparatus.
- No isotope effect in properties
- No decomposition of Flibe by $V \times B$
- We focused on whether or not tritium can be recovered by a realistic scale of apparatus.

Table 1 FFHR parameters on tritium recovery

Fusion power	1GWt
Tritium generation rate	190g-T/day
Coolant	LiF(64%)-BeF ₂ (36%)
Coolant flow rate in blanket	2.3m ³ /s
Colant inlet temperature	470°C
Coolant outlet temperature	570°C
Tritium leak rate	10 Ci/day
Major radius	10 m
Neutron wall loading	1.7 MW/m ²
TBR	>1.2

What is the rate-determining step of tritium recovery

2. Design study : Flow of Flibe and tritium



Tritium leak rate to the environment $\sim$ **10 Ci/day**

$$K_p A_{in} p_{in}^{0.5} + K_p A_{out} p_{out}^{0.5} + \text{others}$$

2. Design study : Evaluation of tritium leak

- If there is **no barrier for tritium leak**,

$$Q_{\text{leak}} = 2 \times 10^{-3} \text{ mol-T/s} = 5 \text{ MCi/day} \rightarrow \text{huge tritium leak}$$

$$(A=10^2 \text{ m}^2, p_{\text{T2}}=10^3 \text{ Pa}, K_p=3.15 \times 10^{-11} \text{ mol/msPa}^{0.5} \text{ (ferrite)}, t=5 \text{ mm})$$

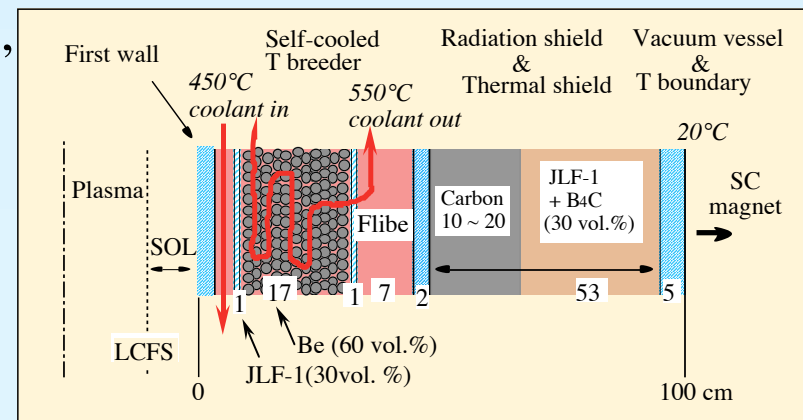
impossible to lower the tritium leak rate, $< 10^{-8}$ Pa

- If the heat exchanger or the tritium recovery system is **surrounded by a stagnant Flibe (or Flinak) layer (barrier)**,

$$Q_{\text{leak}} = 6 \times 10^{-10} \text{ mol-T/s} = 1.6 \text{ Ci/day} \rightarrow \text{allowable tritium leak}$$

$$(A=10^2 \text{ m}^2, p_{\text{T2}}=10^3 \text{ Pa}, D_{\text{T}}=5 \times 10^{-9} \text{ m}^2/\text{s}, K_{\text{H}}=3.2 \times 10^{-7} \text{ mol/m}^3 \text{ Pa}, t=50 \text{ cm})$$

- Other reliable tritium barrier with a factor of 10^6 is OK, if available (e.g. alumina)



2. Design study : Material balance of tritium in Flibe

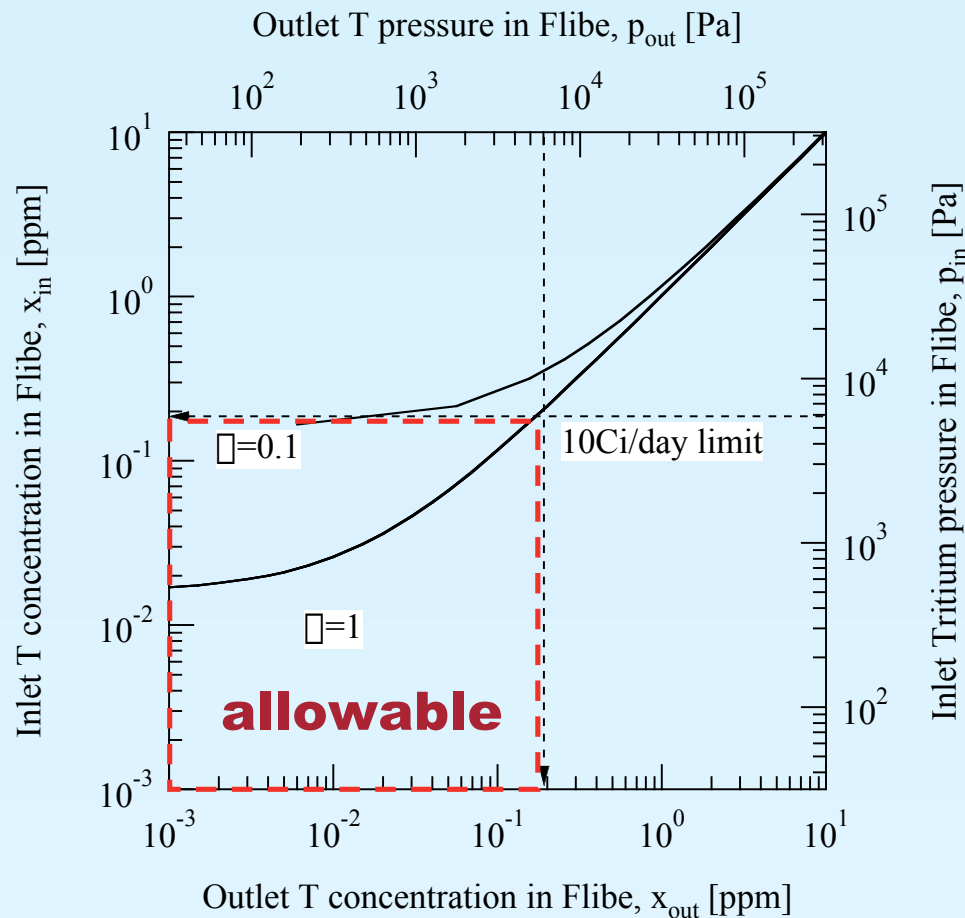


Fig. 2 Relation between c_{in} and c_{out}

$$Q_T = \dot{W}_{Flibe} C_{Flibe} (x_{T,in} - x_{T,out}^b)$$

$$x_{T,out} = \beta x_{T,out}^b + (1 - \beta) x_{T,in}$$

$$p_{T,in} = K_H x_{T,in}$$

$$p_{T,out} = K_H x_{T,out}$$

$$Q_{leak} = K_p A p_{T,in}^{0.5} + K_p A p_{T,out}^{0.5}$$

Necessary conditions

$$Q_T = 1.8 \text{ MCi/day}$$

$$Q_{leak} < 10 \text{ Ci/day}$$

Design of economic apparatus

$\beta=0.1$ is possible

if 50cm shield is present

Permeation window

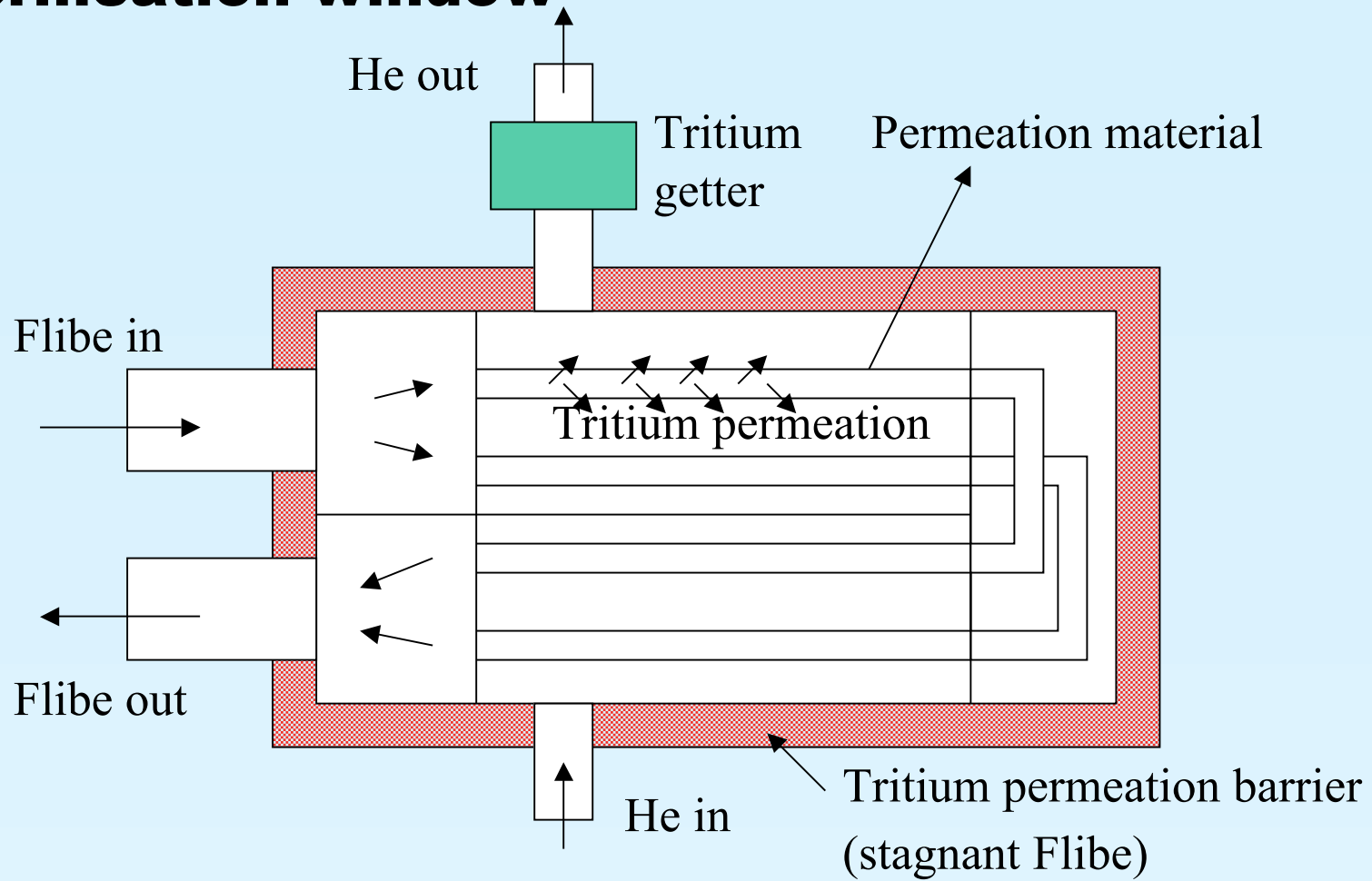
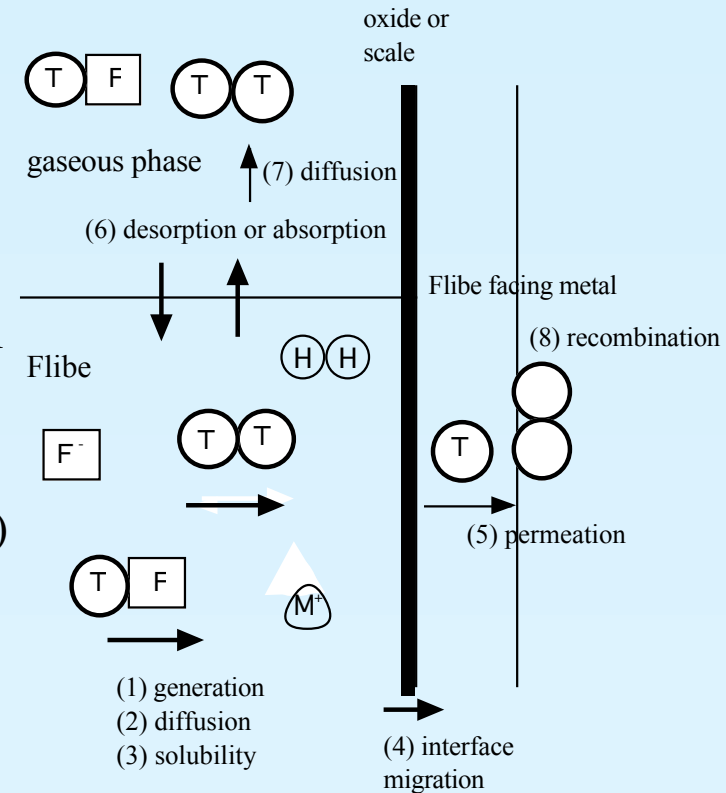


Fig. 3 Permeation window system

2. Design study : Tritium transfer process in Flibe-facing material

Processes include

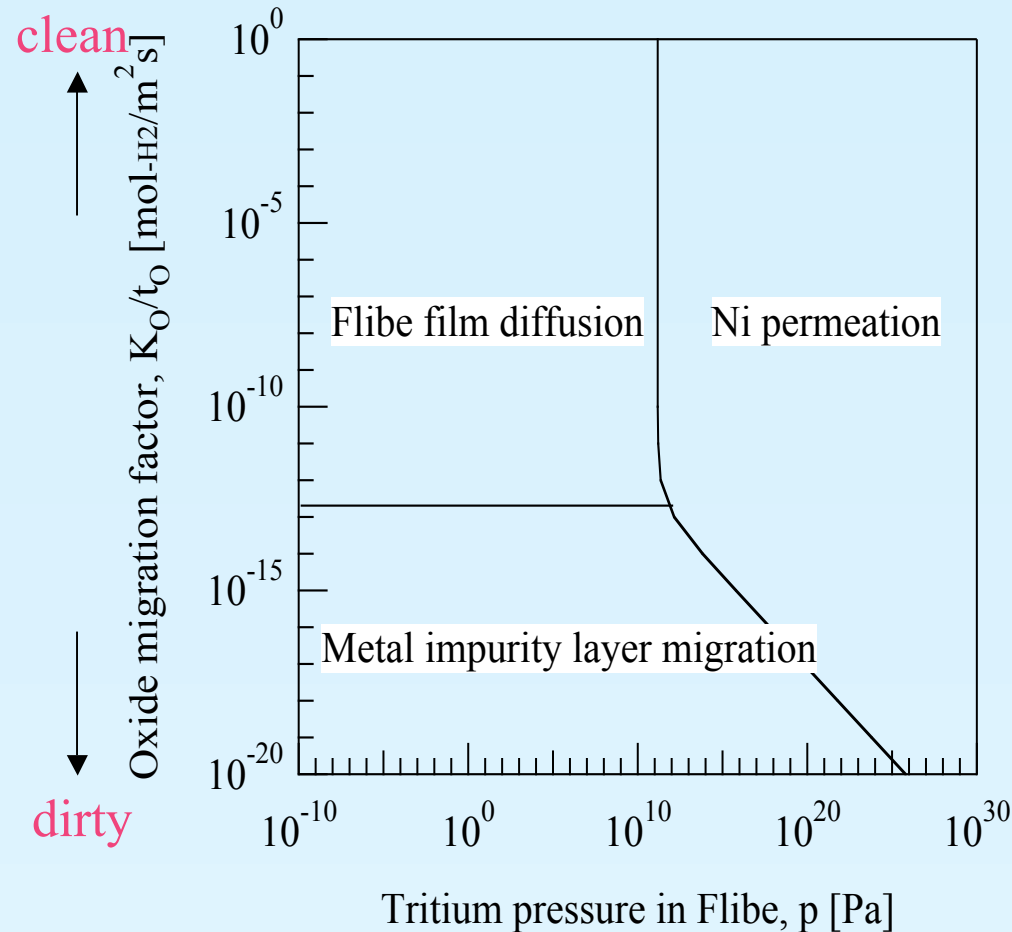
- (1) Tritium (TF or T₂) generation rate in Flibe
 - (2) Tritium (TF or T₂) **diffusion** in Flibe
 - (3) Tritium (TF or T₂) **solution** in Flibe
 - (4) Tritium **migration** from Flibe surface to metal surface (impurity layer)
 - (5) Tritium (T) **permeation** through metal
 - (6) **Desorption** or absorption of tritium (TF or T₂) from Flibe
 - (7) Tritium (TF or T₂) **diffusion** in gas
 - (8) Tritium **recombination** on metal surface
- () unique to Flibe



If H₂ or H₂O coexists in Flibe, HT or HTO is also present.

Fig. 4 Mass transfer processes from Flibe to gas phase

2. Design study : Rate-determining step of permeation window



$$\begin{aligned}
 j &= 2k_M(c_{F,B} - c_{F,1}) && \text{Fluid Flibe film} \\
 &= \frac{2K_O}{t_O}(p_1 - p_2) && \text{Impurity migration} \\
 &= \frac{K_p}{t_p}(\sqrt{p_2} - \sqrt{p_3}) && \text{Permeation} \\
 &= 2k_d c_{s,3}^2
 \end{aligned}$$

Fig. 5 Rate-determining step of the overall tritium permeation

2. Design study :

Design equations of permeation window

- Permeation material : Ni, or ferritic steel
- Upstream side : (purified) Flibe
- Downstream side : He purge
- Overall resistance of mass-transfer of tritium permeation

$$\left(\frac{1}{k_{L,0} H_{T_2}} \right)^{1/2} = \left(\frac{1}{k_L H_{T_2}} \right)^{1/2} + \left(\frac{t_s}{K_p} \frac{(P_{1,in}^{0.5} - P_2^{0.5}) - (P_{1,out}^{0.5} - P_2^{0.5})}{\ln \left(\frac{P_{1,in}^{0.5} - P_2^{0.5}}{P_{1,out}^{0.5} - P_2^{0.5}} \right)} \right)^{1/2}$$

Diffusion in Flibe Permeation through wall

$$\frac{k_L d}{D_L} = 0.023 \left(\frac{\rho_L d v}{\mu_L} \right)^{0.8} \left(\frac{\rho_L}{\mu_L D_T} \right)^{1/3} \left(\frac{\rho_{L,b}}{\rho_{L,w}} \right)^{0.14}$$

- Tritium generated in blanket, Q_T , is all recovered by permeation window.

$$Q_T = A k_{L,0} \frac{(c_{1,in} - H_{T_2} p_2) - (c_{1,out} - H_{T_2} p_2)}{\ln \left(\frac{c_{1,in} - H_{T_2} p_2}{c_{1,out} - H_{T_2} p_2} \right)}$$

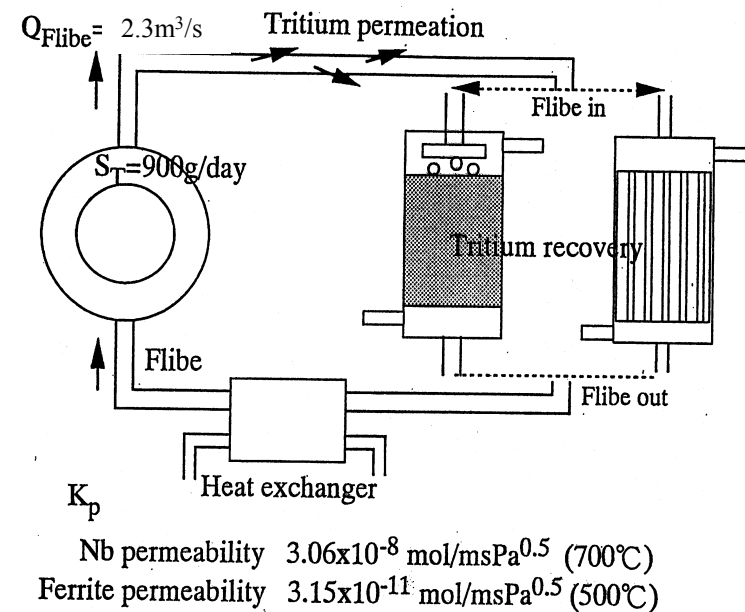


Fig. 5 Permeation window tritium recovery system

2. Design study :

Estimation of permeation window area

- Tritium recovery is easy at high tritium pressure (concentration)
⇒ high tritium leak
- Tritium recovery is difficult at low tritium pressure (concentration)
⇒ large apparatus
- Operation at around $x_T=0.2$ ppm satisfies 10Ci/day limit and 190g/day recovery
- Resistances of Flibe-film diffusion is controlling and permeation gives a small effect

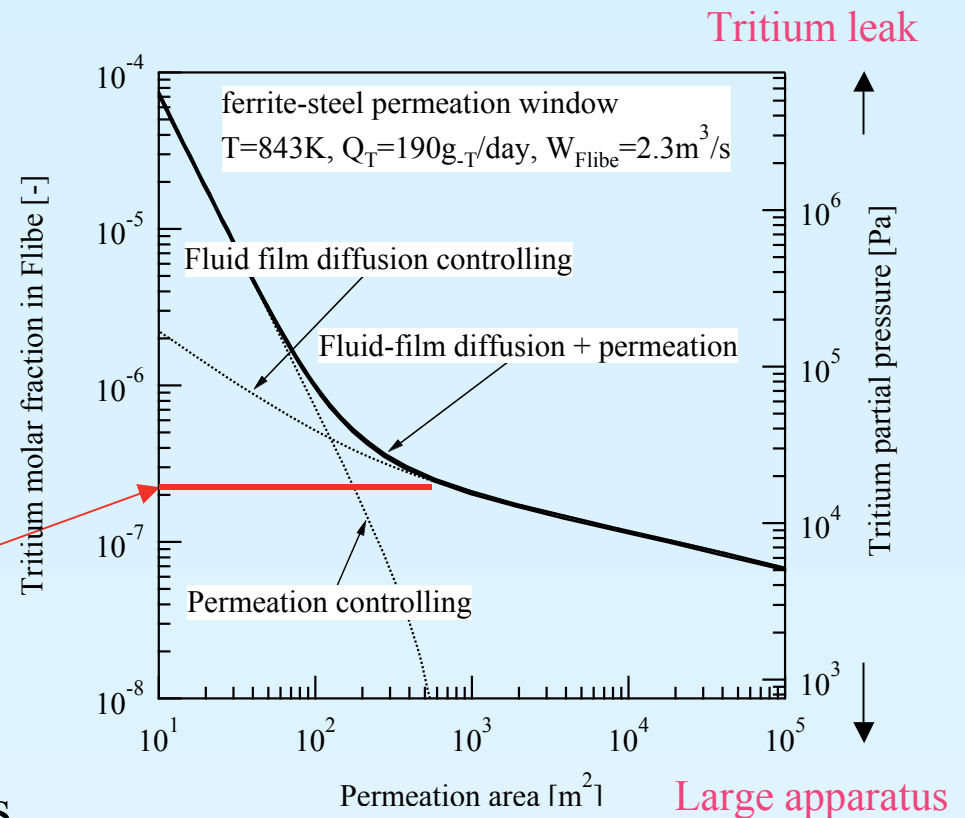


Fig. 6 Permeation window area

($\square=1$, $t=1\text{mm}$)

Counter-current extraction tower

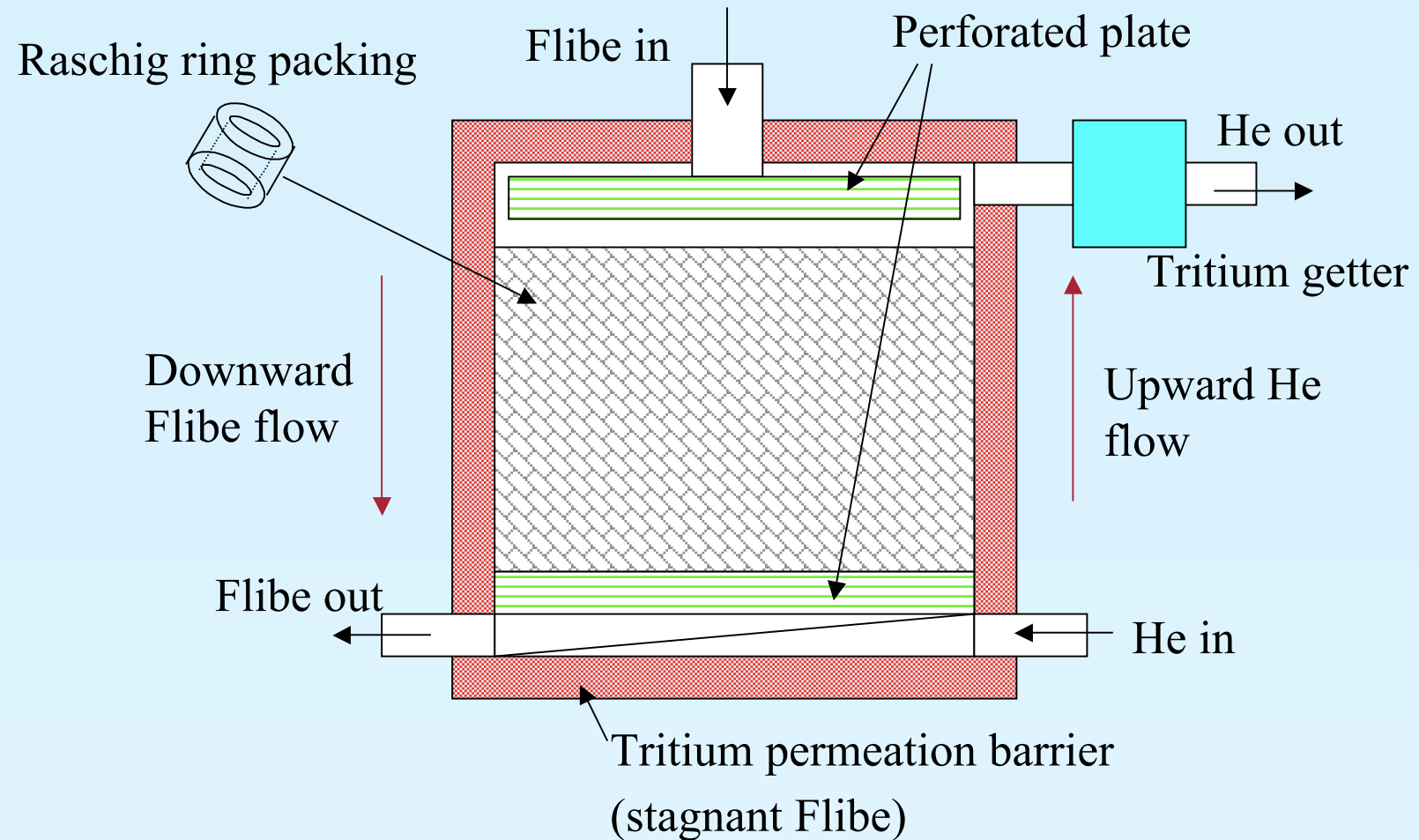


Fig. 8 Flibe-He counter-current extraction tower

2. Design study :

Design equations for He extraction tower

- Material balance equation

$$Ldy = Gdx$$

$$k_L a_v \Delta_s (y - y_i) dz = k_G a_v c_t (x - x_i) dz$$

- Gas-phase height transfer unit

$$H_G = \frac{G}{k_G a_v c_t} = 3.07 \frac{G^{0.32}}{L^{0.51}} \frac{\Delta_G^{\frac{2}{3}}}{D_G}$$

- Flibe-phase height transfer unit

$$H_L = \frac{L}{k_L a_v \Delta_s} = \frac{1}{430} \frac{L^{0.22}}{\Delta_L} \frac{\Delta_L^{0.5}}{D_L}$$

- Concentration profile (Solution)

$$\frac{y}{y_{in}} = \frac{\exp\left[1 - \frac{K_C L}{G H_{0,L}} \frac{z}{H_{0,L}}\right] - \frac{K_C L}{G}}{\exp\left[1 - \frac{K_C L}{G H_{0,L}} \frac{h}{H_{0,L}}\right] - \frac{K_C L}{G}}$$

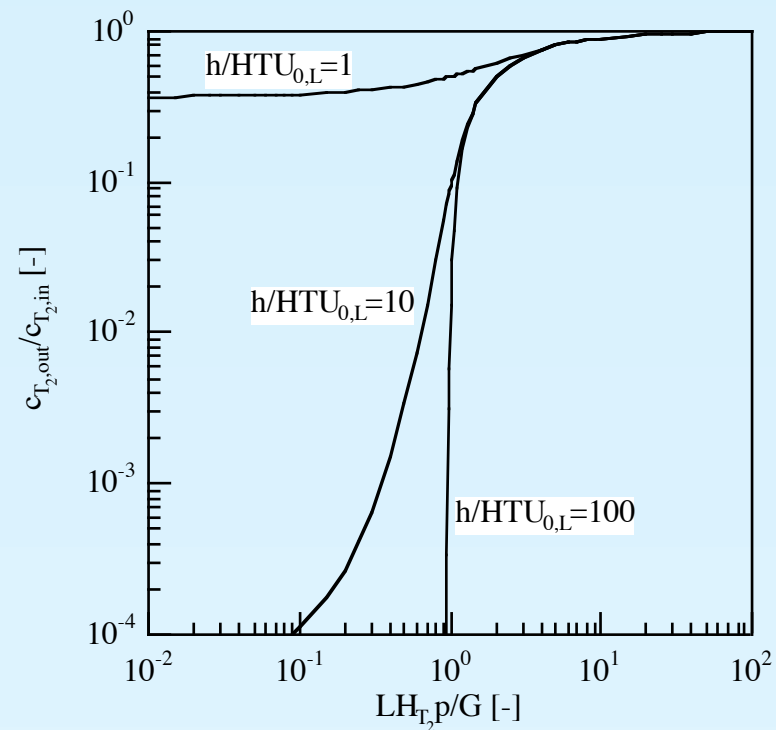


Fig. 9 Concentration change in extraction tower

Rate-determining step is liquid phase diffusion.

2. Design study :

Tritium concentration in He extraction tower

- Tritium concentration in He almost increases logarithmically.
- Tritium concentration in Flibe almost decreases logarithmically.
- 10^5 of DF for Flibe-tritium system is possible by 5 m extraction tower.
- 190 g/day tritium can be recovered by one extraction tower.
- Values of H_L are unsure because they are not determined by Flibe experiment but by water-hydrogen system.
- $(HTU)_L \gg (HTU)_G$

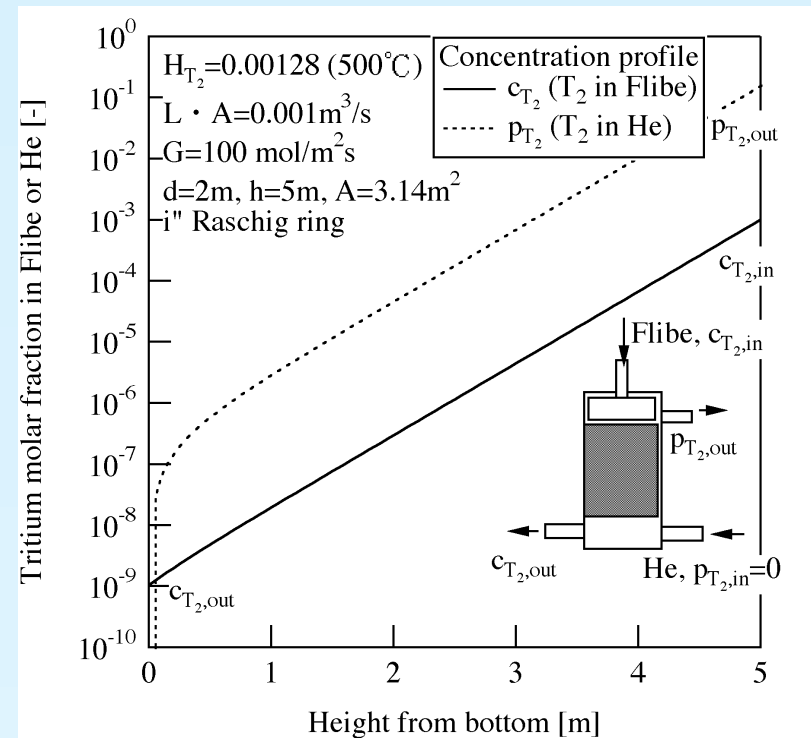


Fig. 10 Tritium concentration in He extraction tower

($\square = 0.1$)

2. Design study : Analysis of vacuum disengager

Assumptions

- Flibe drops free fall from the top and dissolved tritium is desorbed by a vacuum pump.
- $d_p=2\text{mm}$, $h=10\text{m}$, $\square=0.1$
- Surface recombination resistance is ignored.
- Drop formation energy $\sim 1.4\text{kW}$
- Pumping power for perforation $\sim 160\text{kW}$

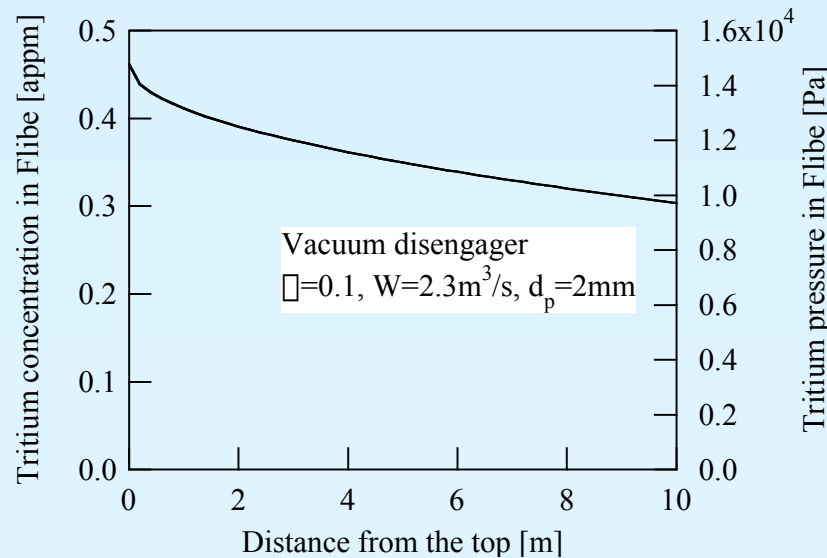
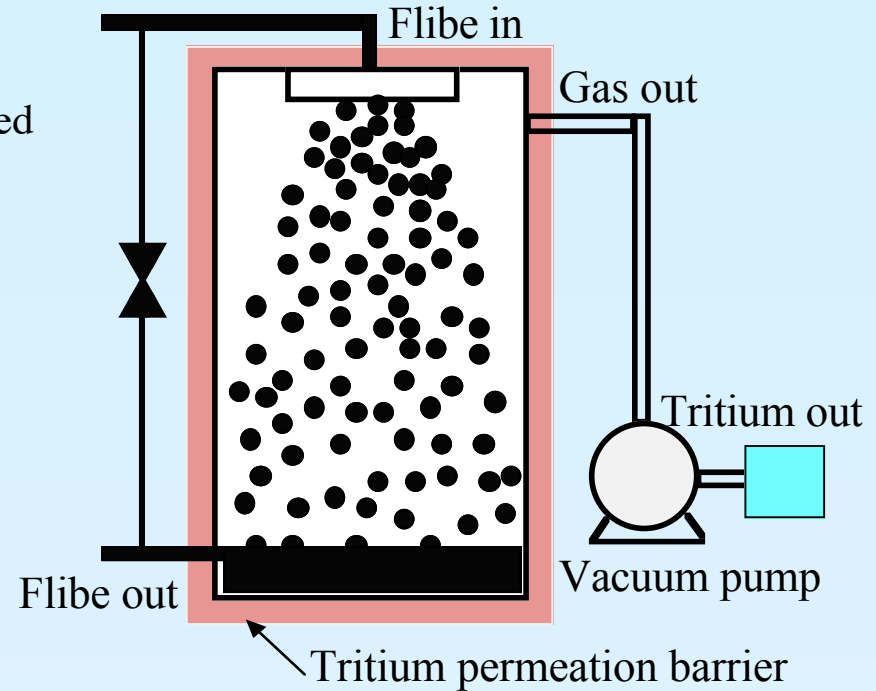


Fig.11 Concentration profile in disengager



$$\frac{x_{T,out}}{x_{T,in}} = \frac{6}{\square^2} \sum_{n=1}^{\infty} \frac{1}{n^2} \exp\left(-\square \frac{4Dn^2 \square^2 t}{d^2}\right)$$

$$t = \sqrt{\frac{2h}{g}}$$

$$Q_T = \square W_{Flibe} c_{Flibe} (x_{T,in} - x_{T,out})$$

2. Design study : Conclusions from design study

- **Need a tritium barrier** to control tritium leak below 10Ci/day
- As long as the tritium leak rate is lower than the permissible level (e.g. by **stagnant Flibe**), arranging the tritium recovery system in a by-pass line is effective.
- In that case, the recovery systems of the **permeation window, extraction column and vacuum disengager** are a realistic-scale apparatus.
- **Diffusion in fluid Flibe is the rate-determining step** in the permeation window and He-Flibe counter-current column.
- In order to raise the accuracy of the design, the mass-transfer coefficient from Flibe to extraction gas should be clearly determined.

2. Design study :

On **Flinabe** as a blanket material (LiF-NaF-BeF_2)

advantages

- Low melting point (=240C?), low operation temperature
- $\text{TBR} > 1$ may be possible with addition of Be.
- Other physical properties may be similar to Flibe

disadvantages

- Activation of Na (formation of ^{22}Na or ^{24}Na)
- There are no data available on tritium mass-transport



More data are necessary for the design of the tritium recovery system