Fluorex Reprocessing System for the Thermal Reactors Cycle

AND

Future Thermal/Fast Reactors (Coexistence) Cycle

Tokyo Electric Power Company,

Hitachi, Ltd.,

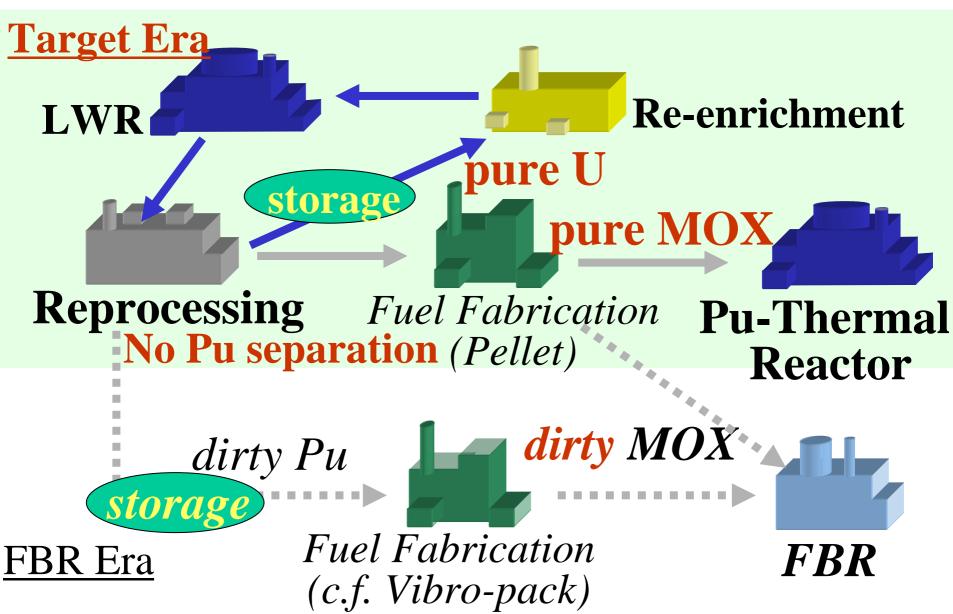
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OUTLINE OF THE PRESENTATION

A new reprocessing system, **FLUOREX**, is a hybrid system that combines fluoride volatilization and solvent extraction methods.

Technical feasibility of the fluorination and interface process is confirmed.

Future Nuclear Fuel Cycle Flow in Japan



Solution for the requirement: FLUOREX process

Key Technology for FLUOREX Process

(1)Selective Vaporization of U

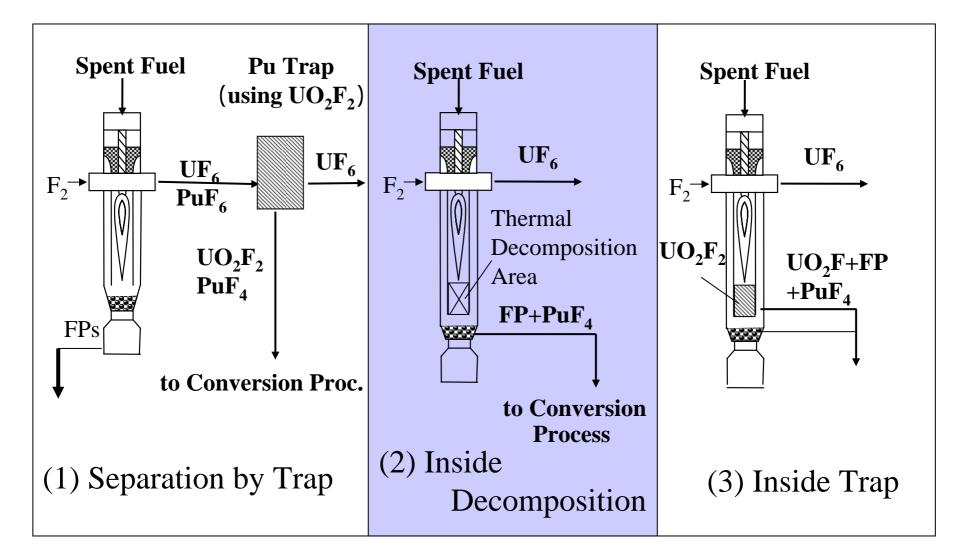
Problem

Vaporization of PuF_6 in the high temperature flame

- $UO_2 + 3 F_2$ $UF_6(gas)$ (Exothermic)
- $PuO_2 + 3F_2$ $PuF_6(gas)$ significant at high temperature

Suppression of Pu Volatilization at Fluorination

Three approaches to Suppress Pu Volatilization

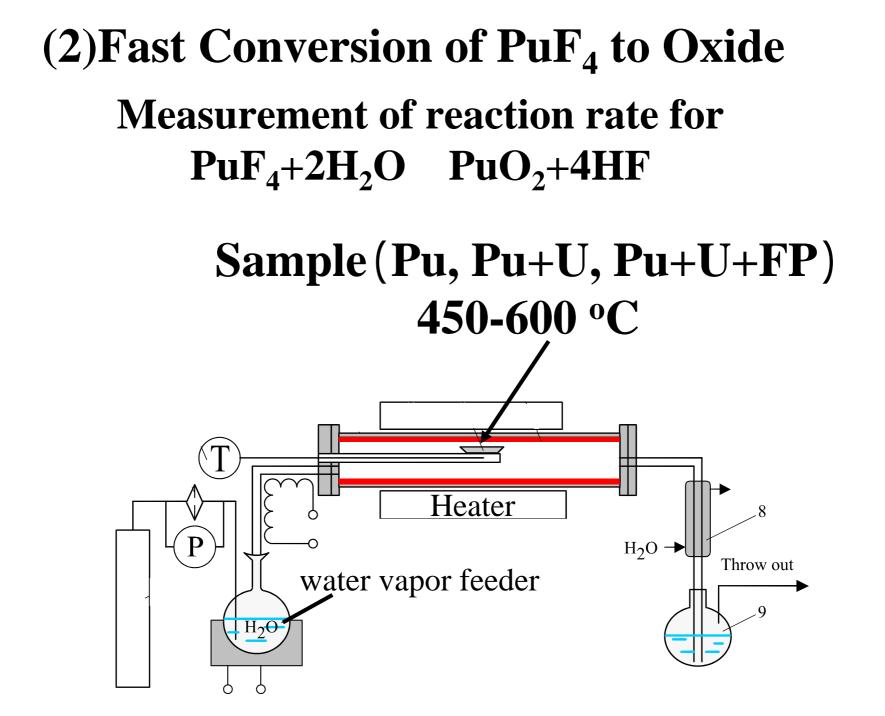


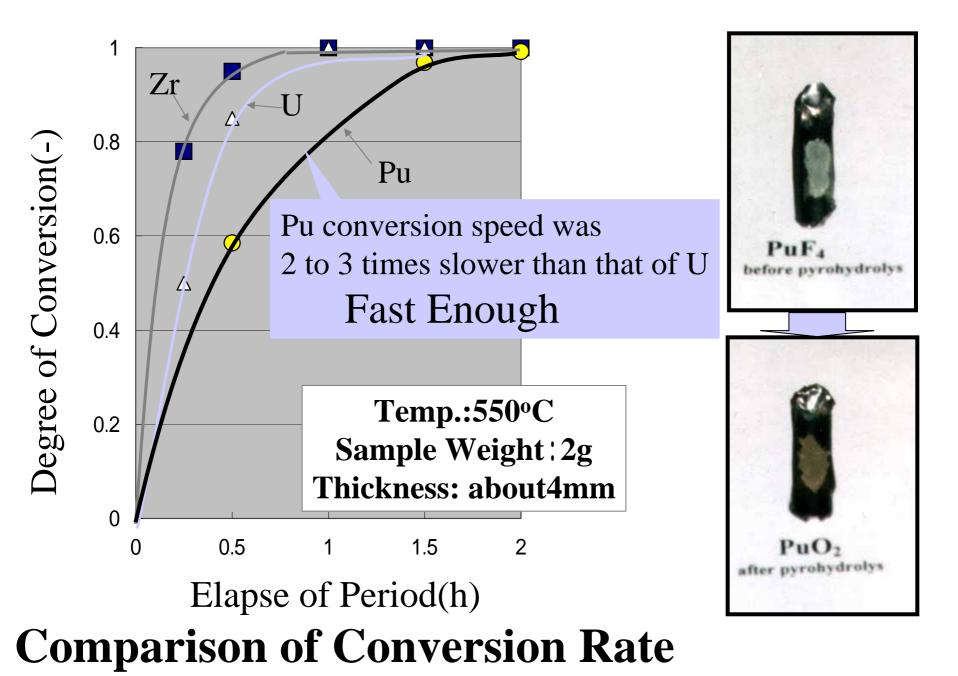
Equipment for Inside Decomposition Test

Gas purification system Flame reactor

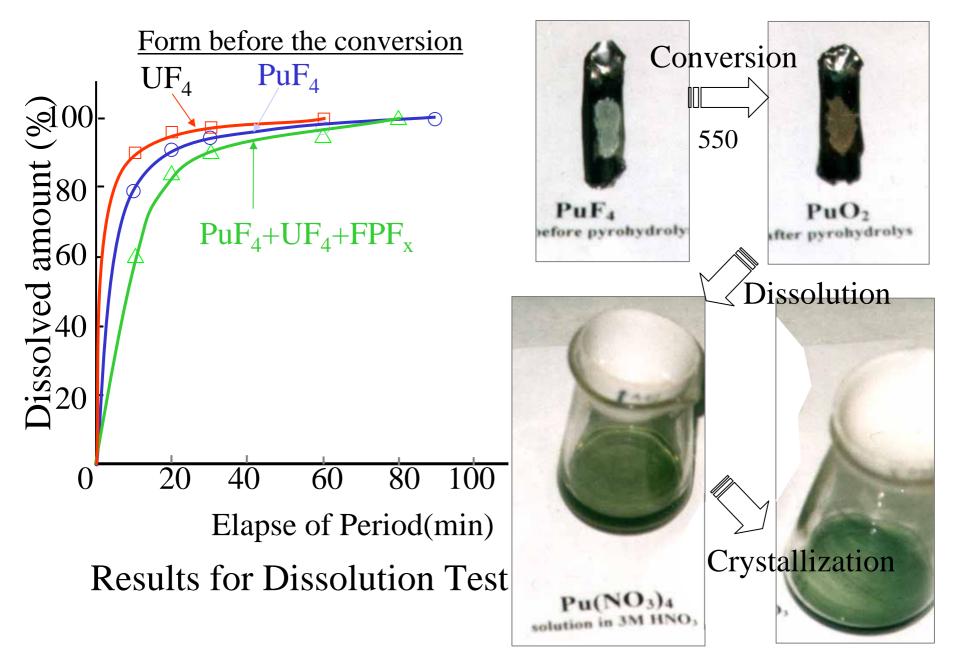


Experimental Results UO₂+PuO₂+FP 100U and Pu Supplier **80** Without 60 Decomposition Gas transferred **Reaction Chamber** 40With Decomposition 20^{-1} $UO_2 + 3F_2$ θ_{0} F_2 feed rate / Stoichiometric rate 1.6 $UF_6 + O_2$ $PuO_2 + 3F_2$ Filter $PuF_6 + O_2$ Recovered $PuF_4 + FPs$ Flame (1200 °C) **Decomposition** Area $PuF_4(solid) + F_2$ PuF₆





(3)Dissolution of Recovered Pu



CONCLUSION

New reprocessing technology, FLUOREX was proposed to meet the requirements for the near future LWR fuel cycle system.

The fundamental processes were proofed for the selective fluorination of uranium, fast conversion of Plutonium fluoride to oxide, and dissolution of the converted oxide.

ACKNOWLEDGMENT

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Supplements

Requirements for the future nuclear fuel cycle system

| Phase | LWR Period | Transition Period (LWR to FBR) | FBR Period |
|---------------------------------|--|---|---|
| Item | (LWR to LWR) | | (FBR to FBR) |
| Feature of Spent Fuel | Contents of $U > 95\%$ U enrichment > ~ 1% | Same as left | Driver: U: about 70% Pu: about 30% Blanket: U > about 95% |
| Usage of | U: Re-enrichment | U: Same as left | U: Fuel pellet or |
| Recovered U | or storage | MOX: Fuel pellet or | Vibro-packing |
| and MOX | MOX: Fuel pellet | Vibro-packing | MOX: Same as left |
| Required DF for Reprocessing | U: High DF MOX: High DF $(DF > \sim 10^7)$ | U: Same as left MOX: Low DF $(DF > \sim 10-10^2)$ | U: High/Low DF [*] MOX: Same as left |

* In the case that high DF U is used for blanket fuel (over aout 50% of FBR loaded fuels), process can be neglected in the FBR fuel production.

U should be reprocessed with high DF
MOX should be flexible from low to high DF

FEATURES OF THE PROPOSED SYSTEM (1/3)

| Economics | Fuel Cycle : UF₆ feed directly to re-enrichment (No conversion facility) Easy removal of daughter nuclides from UF₆ Reprocessing : Fast and Compact fluorination and U purification equipments Reduced scale of solvent extraction (less than 10%) Cost reduction in low-level waste treatment (by one digit) Flexibility in the choice of HLW geological disposal methods Ru/Rh, Tc, I and Np: easily removed in the U purification separated disposal from HLSW |
|-----------|--|
| Safety | Gaseous U inventory: only 2kg-U for 100t/y plant with reduced pressure equipments and cells Volatile FPs: fixed in suitable adsorbents in U purification treated in solid forms in the fluorination process. No gaseous Pu in the whole system Solvent extraction processes: less than 1/10 of the PUREX plant |

FEATURES OF THE PROPOSED SYSTEM (2/3)

| Utilization of Resources | High DF for U and MOX: enhance the nuclear energy utilization U: from 10⁷ to 10⁹ by chemical sorption and rectification MOX: 10⁷ (same as conventional PUREX system) |
|--------------------------------------|---|
| Minim. of Environmental Burden | U purification processes: small amount of radioactive waste (few volatile fission products accompanied with UF₆) Solvent extraction processes: less than 1/10 of PUREX Total amount of the waste: about 1/10 of the conventional one Heat generation nuclides like Cs and/or Sr: Able to be separated at decladding or conversion process. |
| Proliferation Resistance | Pu is always with U even in a purified form in FLUOREX solvent extraction processes. (higher proliferation resistance than PUREX, where Pu alone exists in its purification process) If Pu is stored with FPs for future, the resistance would be much higher. |

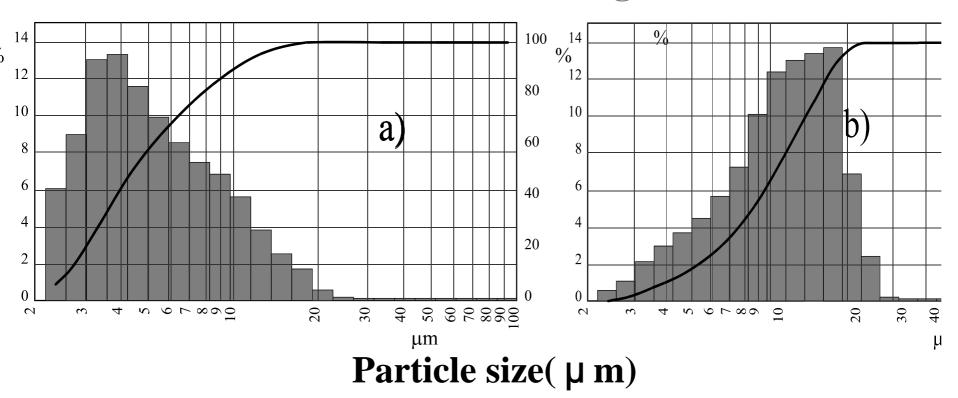
FEATURES OF THE PROPOSED SYSTEM (3/3)

| Maturity | Most techniques: already developed industrially in the past 1) Solvent extraction: Much experience in reprocessing industry 2) Fluoride volatility: Much industrial experience in conversion and U enrichment Pilot scale experience in U purification (Russia, USA, etc) | |
|----------|---|--|
| | The technologies developed in the past in the world can accelerate the development of FLUOREX reprocessing system, which indicates the importance of worldwide collaboration for the industrialization of the system and for the sustainable utilization of nuclear energy. The worldwide information exchange and collaboration is also indispensable to emerge from the present global nuclear shrinkage situation. | |

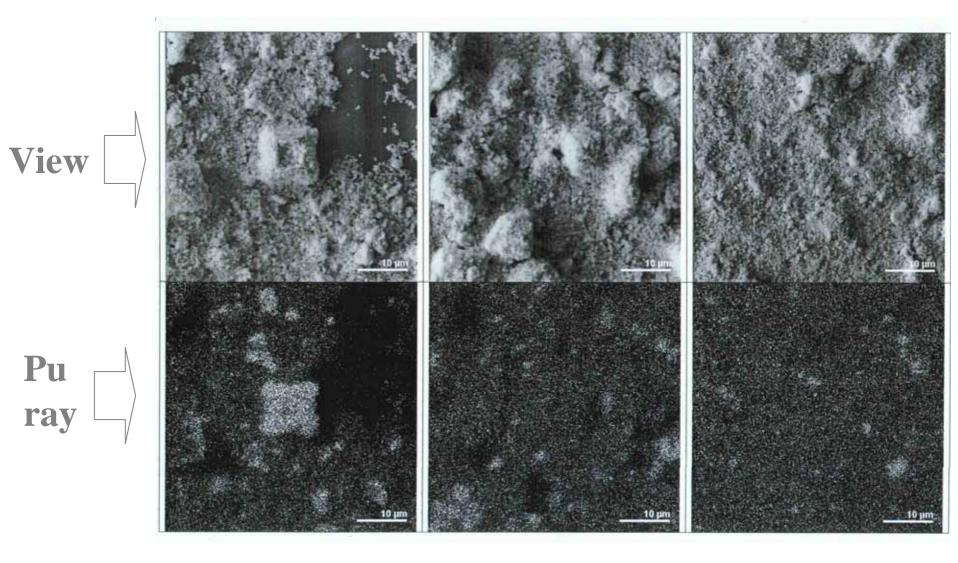
Sample Condition(Fluorination)

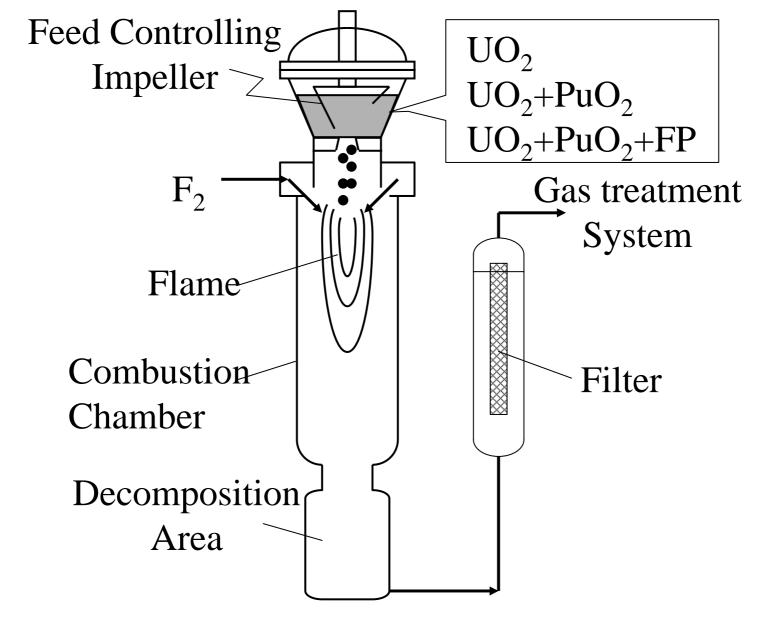


Weight Distribution

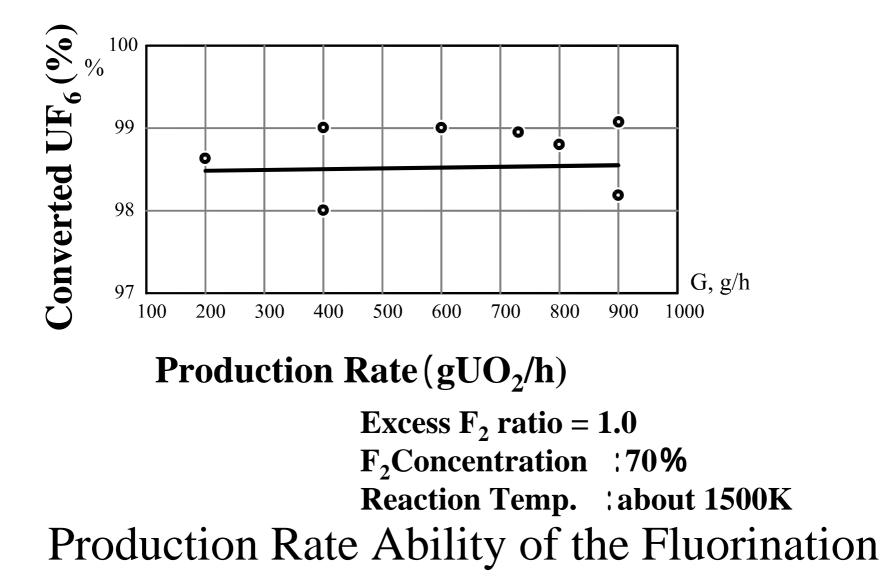


Sample view for U+Pu mixture before fluorination





Small flame reactor for fluorination



| Compd. | Rate | Compd. | Rate | _ |
|-----------------|------|------------------|-------|---|
| UF ₄ | 1 | CeF ₃ | 0.2 | |
| ZrF_4 | 2 | NdF ₃ | 0.05 | |
| PuF_4 | 0.3 | SrF_2 | 0.005 | |

Relative Reaction Rate of Fluorides Conversion

| Dissolution test sample before conversion | | | |
|---|-----------------|-----------------------------------|---------------------------------------|
| FORM | Contents (%) | | |
| | UF_4 | UF ₄ +PuF ₄ | UF ₄ +PuF ₄ +FP |
| UF ₄ | 100 | 80 | 40 |
| $Pu\dot{F}_4$ | - | 20 | 10 |
| SrF_2 | - | - | 3.6 |
| $La\tilde{F_3}$ | - | - | 5.15 |
| CeF_3 | - | - | 9.8 |
| NdF_{3} | - | - | 21.15 |
| RbF | - | - | 9.78 |
| GdF_3 | - | _ | 0.52 |