ICTP Workshop Modelling and Quality Control for Advanced & Innovative Fuel Technologies

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**Opening Address** 

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International Atomic Energy Agency

# **NUCLEAR FISSION**



### **FISSILE & FERTILE ISOTOPES**

10-

10-2

10-3

LI4-MEV

FISSIO

Thermal Fission

**Fission Product** 

120

Heavy

(125-155)

Te, I, Xe, Cs

Ba, La, Ce,

140

160

RE

Light

(80-110)

Br, Kr, Rb

Mo, Tc

Sr, Y, Zr, Nb,

11111

11111

LILLI

180

### **URANIUM**

- 4 ppm in earth's crust, Klaproth, 1789

**Natural Uranium:** U<sup>238</sup>(fertile): 99.3%; 4.5x10<sup>9</sup>y U<sup>235</sup>(fertile): 0.7%; 8.0x10<sup>8</sup> y

U<sup>236</sup> <sub>92</sub>(fissile) + n<sup>1</sup><sub>0</sub> U<sup>236</sup> M<sub>1</sub>(95)+M<sub>2</sub>(139)+(2-3)n<sup>1</sup>+200 MeV U<sup>238</sup> 92 fertile) +  $n_0^1 U_{32}^{239} \beta^2 N p_{33}^{239} \beta^2 23.5 m$ (fissile) → (Pu<sup>23</sup> 94

### THORIUM

- 12 ppm in earth's crust, Berzellius, 1828



### PLUTONIUM

Does not occur in nature; man-made, Seaborg, Wahl and Kennedy, 1941 -

> Pu <sup>239</sup>	- :	oc ;	2.4 x 10 <sup>4</sup> y	
> Pu <sup>240</sup>	1	∞, <b>n</b> ;	6.58 x 10 <sup>3</sup> y	
> Pu <sup>241</sup>	:	β- ;	13y, Am241	strong γ emitte
> Pu <sup>242</sup>	1	∞, <b>n</b> ;	3.79 x 10⁵ y	
> Pu <sup>238</sup>		∞, <b>n</b> ;	86.4 y	

### **Maximum limits**

.....

body burden	:	0.18 – 0.65 μ g
concentration in air	:	<b>2 x 10</b> <sup>-12</sup> μ curie per c
concentration in water	:	10 <sup>-4</sup> μ curie per cc



# **NUCLEAR REACTOR**



# **NUCLEAR REACTORS & THEIR APPLICATIONS**

Source of intense heat energy & Source of neutrons

• Power Reactors:



- Production of Hydrogen

•

- District Heating
- Non -Power Reactors:



### Nuclear Energy for Tomorrow – to Combat CO<sub>2</sub> emission & Global Warming

**ENERGY PRESENTLY UTILISED:** 



Of today's 6 billion people in the world, one third has no access and another one third has limited access to electricity.

 Nuclear -66% global electricity is generated from Hydro fossil fuel, including 40% from coal, causing large release of CO₂ greenhouse gas to the environment and in turn global warming.

- Fossil-fuel would be significantly depleted by the end of this Century.

TRANSPORTATION: 75%

- Cars, trucks, trains and planes use 'fossil fuels': gasoline, diesel, jet fuel, etc. releasing CO<sub>2</sub> to the environment.

- Only 12% of the world's population has access to automobiles - when the other 88% decides to drive, the present 'fossilfuelled' vehicle, imagine the  $CO_2$  emission to the environment.

Fossil Fuels release 70 million tons  $CO_2$  every day or 800 tons/second to the environment causing Global Warming.

In the next 50 years - as world population expands to 9 billions - global energy consumption will double. How do we meet the ever-increasing demand minimizing  $CO_2$  emission that cause global warming?

NUCLEAR ENERGY IS THE INEVITABLE OPTION !

NUCLEAR POWER STATIONS (do not emit CO<sub>2</sub> & cause global warming)



- NUCLEAR FISSION HEAT ENERGY should be used for generation of electricity, desalination of sea water, district heating in cold countries & for production of hydrogen economically by electrolysis of water or by cracking of hydrocarbon.

- <u>HYDROGEN</u>, instead of GASOLINE, should be the energy source for 'land transportation' in tomorrow's mega cities for minimising CO<sub>2</sub> greenhouse gas emissions 8 other automobile pollution.



# Atoms for Peace – Civilian Nuclear Power programme

### **Commercial Phase**

- By the end of 1960s and early 1970s, PWRs, BWRs and RBMK were commercialized when several 1000 MWe units were commissioned in USA and Russia. The VVER-440 reactors were also commercialized during this period.
- The Nuclear power programme reached the commercial phase all over the world in the 1970s and 1980s when a large number of Light Water Reactors (LWR), including PWRs, BWRs, VVERs and RBMK type reactors, CANDU-PHWRs and Gas-Cooled Reactors (both Magnox and Advanced Gas-cooled Reactors) were commissioned in USA, Canada, USSR, Europe, Japan, South Korea and India. Several prototype and commercial LMFBRs were also commissioned in USSR, France and UK.
- As on March 2005, there are some 440 commercial nuclear reactors operating in 31 countries with over 360 GWe total capacity. They supply 16% of world's electricity.





# Pressure Vessel and Pressure Tube Type of Nuclear Power Reactors







# TYPICAL CALANDRIA VESSEL

# **NUCLEAR POWER REACTORS**

- Light Water Reactor (LWR): 87%
  - Pressurised Water Reactor (PWR). Russian type known as VVER
- Boiling Water Reactor (BWR) USA, France, Germany, Sweden, Belgium, Russia & CIS (WER), Japan, South Korea, Brazil & China
- Pressurised Heavy Water Reactor (PHWR):6% (also known as CANDU) Canada, India, South Korea, Romania, Argentina, Pakistan & China
- Light Water Graphite moderated Reactor (LWGR): 3% Russia and CIS known as RBMK type reactor.
- Gas Cooled Graohite Moderated Reactor (GCR): 3% Popular only in UK.

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 Liquid Metal Fast Reactor (LMFR):~1% Presently, only one prototype and commercial LMFR is in operation: Phenix 250 MWe (France) & BN-600 MWe (Russia). In India, a Fast Breeder Test Reactor (FBTR) is in operation. Prototype Fast Breeder Reactor is under construction in Japan (Monju) and India (PFBR)



REACTORS IN OPERATION 441 Nos. 3 67 496 MWe

# **Civilian Nuclear Power Programme in the 21<sup>st</sup> Century:**

### **On-going International Programmes:**

- IAEA-initiated INNOVATIVE NUCLEAR REACTORS & FUEL CYCLE PROGRAMME (INPRO)
- US-initiated Generation IV International Forum (GIF)
- In both INPRO and GIF, nuclear power reactors have been perceived to be utilised for the following purpose:
  - Generation of Electricity.
  - Production of Hydrogen Fuel
  - Desalination of Sea Water
  - District Heating



### **UNIQUE FEATURES OF NUCLEAR ENERGY & POWER**

### Relatively New

"Nuclear Fission Energy" is very recent (< 60 years) in the time scale of human civilisation. First Nuclear Power Reactor in the world : 1955

### High Energy Density

1 atom of 'C' on combustion releases ~ 4 eV 1 atom of U<sup>235</sup> on fission release ~200 MeV

Annual Fuel Requirement of a 1000 MWe Power Station: Nuclear : 30 tons; Coal : 2.6 million tons; Oil : 2.0 million tons

### • Environment Friendly:

Zero emission of  $CO_2$ ,  $SO_2 \& NO_x$  – no global warming and acid rain Annual Discharge from 1000 MWe Power Station: Nuclear : 3.5 tons spent fuel: Coal Fired : 6.5 million tons  $CO_2$ 

CO<sub>2</sub> emission per kWh: 0.967 kg Denmark ( 0.63 kg UK (4 0.064 kg France (

(82% Coal, 0% Nuclear) (49.5% Coal, 28% Nuclear) (77.36% Nuclear)

### · Generates man-made 'fissile' isotopes or fuels :

Nuclear fuel is made up of 'fissile' (U<sup>235</sup>, Pu<sup>239</sup> & U<sup>233</sup>) and 'fertile' (U<sup>238</sup> & Th<sup>232</sup>) isotopes. U<sup>235</sup> is the only 'fissile' isotope occurring in nature. The 'fission' process splits up 'fissile' nuclei, releases very high heat energy and generates extra neutrons which could convert naturally occurring 'fertile' isotopes, U<sup>238</sup> & Th<sup>232</sup> to man-made 'fissile' isotopes Pu<sup>239</sup> & U<sup>233</sup> respectively.



### High Safety & Security

From radiation safety point of view, natural uranium (U<sup>235</sup> & U<sup>238</sup>) and thorium (Th<sup>232</sup>) are mildly radioactive and have very little hazard from external radiation. However, Pu<sup>239</sup> (always associated with Pu<sup>240</sup>, Pu<sup>241</sup>, Pu<sup>242</sup> & Pu<sup>238</sup>), U<sup>233</sup> (always associated with U<sup>232</sup>) and fission products are highly radioactive and health hazardous & require proper containment, beta, gamma neutron shieldings and remote handling. In order to ensure safety from any 'criticality accident' (uncontrolled nuclear fission chain reaction), only limited and controlled quantity of 'fissile ' (U<sup>235</sup>, Pu<sup>239</sup> or U<sup>233</sup>) materials are permitted to handle at a time. The radioactive waste has to be properly treated, fixed and stored or disposed. From security point of view, physical protection of 'fissile' material is essential to avoid proliferation risk for non-peaceful purpose.



### Comparitive Hazards associated with external dose from radioactive material



### Alpha Radiation: (practically no hazard from external radiation)

Can be stopped completely by a sheet of paper – may just penetrate the surface of the skin.

Beta : (minimum hazard from external radiation) Radiation

Can be stopped by a sheet of aluminium a few mm in thickness.

Gamma: (hazardous-requires proper shielding) & X Rays

Are very penetrating and can pass right through human body – mostly absorbed by heavy elements like lead, which is normally used as shielding material for gamma x-rays.

Neutrons: (hazardous - requires proper shielding)

Are very penetrating – in general, efficient shielding against neutron can be provided by water, perspex, etc. Concrete shielding (sometimes upto 1 m thickness) is used for neutron and gamma shielding.



Remote Operations using manipulators for handling highly radioactive materials (mainly high gamma and high neutron dose) inside concrete hot cells



### **CRITICAL MASS OF URANIUM & PLUTONIUM**

Fissile	Approximate		
Material	Critical Mass (kg)		
	Bare	Water reflected	
URANÍUM			
Density 18.8 g/cm <sup>3</sup>			
Natural 'U'	Cannot become	e critical alone	
$U^{235}$	47	20	
$U^{233}$	17	7	
90 % enriched $U^{235}$	53	24.5	
20 % enriched $U^{235}$	750	375	
PLUTONIUM		-	
Density 19.85 g/cm <sup>3</sup>			
$Pu^{239}$	10	5	
PuO <sub>2</sub>	24.5	12.2	
PuC	18	9	
$PuF_4$	56	25	

### Max. Permissible Conct. Of 'Pu' radionuclides

Plutonium Radionuclides	Body (MPBB) (µg)	Air (μ Curie/cm³)	Water (μ Curie/cm³)
Pu <sup>238</sup>	2.4 x 10 <sup>-3</sup>	2 x 10 <sup>-12</sup>	10-4
Pu <sup>239</sup>	0.65	2 x 10 <sup>-12</sup>	10-4
Pu <sup>240</sup>	0.18	2 x 10 <sup>-12</sup>	10-4
Pu <sup>241</sup>	8.2 x 10 <sup>-3</sup>	9 x 10 <sup>-11</sup>	7 x 10 <sup>-3</sup>
Pu <sup>242</sup>	12.8	2 x 10 <sup>-12</sup>	10-4

# MAJOR NUCLEAR FUEL FORMS A: Pellet-pin





(Vibratory compacted fuel pins have been utilized on a very limited scale for experimental purpose)

# **FUEL FORMS B:** Coated Fuel Particles

### 1. Coated fuel particles for HTGR

**Prismatic block** 

### US, Japan, Russia and France



Pyrolytic Carbon Silicon Carbide Porous Carbon Buffer UCO Kernel TRISO Coated fuel particles (left) are formed into fuel rods (center) and inserted into graphite fuel elements









FUEL ELEMENTS

2. Fuel particles (dry or wet route) for vibratory compacted fuel pins



Pebble Bed coated particle fuels embedded in spherical shape Germany, South Africa, China



Triso coated particle





For nuclear energy to be sustainable as a global source of emission – free energy, the reactor fuel cycle must also remain sustainable (DG-IAEA Scientific Forum 2004)

# **Workshop Structure – 3 Modules & Topics**

### **Module I: Nuclear Power Reactors and Fuels**

- Reactor systems
- Fuel cycle options
- INPRO and GIF
- Conventional, advanced and innovative fuels

### Module II: Fuel Design, Fabrication, QC, Modelling, Irradiation – Testing & PIE

- Fuel rod and assembly design
- Fabrication & QC of fuel pellets, rods and assembly
- Irradiation testing in research and power reactors and
- results of post irradiation examination (PIE)
- Fuel modelling and different codes

### Module III: Advanced Fuel Management

- Spent fuel management
- Use of plutonium and uranium-233 based fuels
- Proliferation-resistant fuels



ICTP Workshop Lecture 2 Nuclear Fuel Cycle Options

# **Chaitanyamoy Ganguly**

Trieste, 15 November, 2005

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For nuclear energy to be sustainable as a global source of emission – free energy, the reactor fuel cycle must also remain sustainable (DG-IAEA Scientific Forum 2004)

# **MAJOR CHALLENGE IN NUCLEAR FUEL CYCLE**

# Develop a fuel cycle that is economically viable, environmentally benign, proliferation-resistant, safe and sustainable



### **Permanent Repositories for Safe Disposal of Nuclear Wastes**



Monitored Geological Repository for High Level Radioactive Waste at Yucca Mountains, Nevada

**Basis of Selection:** 

- i) its remote location and long distance from a large population center—100 miles from Las Vegas, Nevada;
- ii) its very dry climate—less than 6 inches of rainfall a year
- iii) its extremely deep water table—800 to 1,000 feet below the level of the potential repository



Four Permanent Deep Repositories in Finland for Safe Disposal of Highly Radioactive Spent Fuels: 2 at Nuclear Power Plant sites, namely Olkiluoto at Eurojoki & Lovisa and other 2 at Kuhmo at Aanekoski.

The ONKALO facility at Olkiluoto is under construction at depths of 300, 400 and 500 meters with an access tunnel and an associated ventilation tunnel. The bed-rock at Olkiluoto is suitable for safe disposal. The radioactive waste would be packed in copper based canisters, which will be surrounded by compacted bentonite in the repositories. The Permanent Repository at Olkiluoto would be operational in 2010.

# ACTIVITIES (PROGRAMME B) OF NUCLEAR FUEL CYCLE & MAT0ERIALS SECTION (NFC&MS), IAEA

B1: Uranium Production Cycle and Environment

B2: Water-Cooled Fuel Performance and Technology (Database: PIE)

B3: Management of Spent Fuel from Power Reactors

B4: Topical Nuclear Fuel Cycle Issues and Information Systems (Databases: NFCIS, UDEPO, MADB, VISTA)





L4: Technologies for Disposable Radioactive Waste (Pre-disposal & Disposal)

L7: Technologies for the Decommissioning of Installations and Restoration of Sites



International Atomic Energy Agency



# International initiatives on innovative reactors and fuel cycle

- INPRO: International Projects on Innovative Nuclear Reactors and Fuel Cycle
- GIF: Generation IV International Forum and Advanced Fuel Cycle Initiative (AFCI)



# **CURRENT TRENDS IN URANIUM DEMAND & SUPPLY**

**Demand:** 66,815 tons 'U'

**Production:** 40,251tons 'U' (60% of demand)

The shortfall of 40% is met from the following Secondary Supplies

- Excess Commercial Inventories
- LEU from Ex-Military HEU
- Re-enrichment of DU-Tailings (enrichment plants) & REU (reprocessing plants)
- Ex-Military & Civil Pu (to be used in the form of Mixed Uranium Plutonium Oxide (MOX)

THERE IS NO SHORTAGE OF URANIUM RESOURCES FOR NEXT 50 YEARS EVEN FOR ONCE-THROUGH FUEL CYCLE Countries with Major Uranium Resources & No. of Nuclear Reactors & Countries with Major Nuclear Power Programme & their U-Sources [World Uranium Resources (RAR<130US\$/kg 'U'): 3.17 million tons 'U']

(Pof: IAEA Pod Pook Dog 2002)

			CU DOOK DCC. 2003)
Country	Uranium Resources	Percentage	No. of Nuclear
	(Tons 'U')	of world	Power Reactors
		resource	(% Electricity)
Australia	735,000	23%	Nil
Kazakhstan	530,460	17%	Nil
Namibia	170,532	5%	Nil
Niger	102,227	3 %	Nil
Uzbekistan	79,620	2.5%	Nil
Mongolia	46,200	1.5%	Nil
USA	345,000	11%	104 (20%)
Canada	333,834	10.5%	20 (~12%)
South Africa	315,330	10%	2 (5.9%)
Russian Fed.	143,020	4.5%	30 (16%)
Brazil	86,190	3%	2 (4%)
France	100% from over	seas sources	59 (78%)
Germany	100% from over	seas sources	18 (30%)
Japan	100% from overseas sources		53 (39%)
Korea (R.O.)	100% from over	seas sources	19 (39%)
China (excl. Taiwan)	35,060	1%	9 (1.4%)





# **Uranium Resources Worldwide**

(Source: Uranium 2003 – 20th Edition of IAEA Red Book jointly with OECD-NEA)

1. Conventional Resources

**Reasonably Assured Resources (RAR) Estimated Additional Resources (EAR) category I & II)] Speculative Resources (SR)** 

a) Conventional Resources (RAR and EAR-1): ≤ US \$ 80 / kg U : 3,537,000 tons; ≤ US \$ 130 / kg U: 4,589,000 tons U

b) Undiscovered Conventional Resources (EAR-II and SR):9,794,000 tons U

Total Conventional Resource (a + b): 14,383,000 tons U

2. <u>Total Unconventional Resource</u> Uranium in Phosphates Uranium in Sea Water

22,000,000 tons U 4,000,000,000 tons U

()

(with improved adsorbent material, the latest estimated price of U from sea-water reported by Japanese is in the range of US\$ 300/kgU – IAEA Red Book 2001).

# **World Thorium Resources - economically extractable**



# **URANIUM MINING TRENDS**





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### FROM ZIRCON SAND TO ZIRCONIUM ALLOY INGOTS AT NFC, HYDERABAD



**Zircon Sand** 



Hf-free ZrO<sub>2</sub> Powder



**Nuclear Grade Zr Sponge** 

Compaction of Zr Sponge + alloying elements Briquettes



Zirconium Alloy Ingot Max. size: 350 mm dia x 2 m height







**Electron Beam Welding of Briquettes** to form Consumable Electrode



### **Major Activities of Zirconium Alloy Fabrication Plant at NFC**



Hot Extrusion of Seamless Zirconium Alloy Tubes



**Pilot Hole Expansion Press** 



Pilger Mill for Production of Zirconium Alloy Fuel Tubes







Hot Rolling of Zirconium Alloy Sheets



**Cold Swaging of Zirconium Alloy Bar** 



### FUEL ASSEMBLY FOR 540 MWe PRESSURISED HEAVY WATER REACTORS

### > ONE FUEL ASSEMBLY CONTAINS

• ZIRCALOY- 4 COMPONENTS

FUEL TUBES	(13.08 od x 0.38 t x 485.8	I)	37 Nos
END CAPS	(13.20 <b> </b>		74 Nos
SPACER PADS	(8.6 l x 2.5 w x 3.22 t)		12 Nos
	(8.6 l x 2.5 w x 0.87 t)		144 Nos
<b>BEARING PADS</b>	(33.5 l x 2.5 w x 1.3 t)		36 Nos
	(32.5 l x 2.5 w x 1.0 t)		18 Nos
END PLATES	(90.98 <b>¢</b> x 1.60 t)		2 Nos

NATURAL URANIUM DIOXIDE

PELLETS	(12.20 ¢ x 13.29 l)	1295 Nos.
WEIGHT OF PEI	LETS	21.65 kg

> ONE PHWR 540 CORE CONTAINS

FUEL ASSEMBLIES	5096 Nos
URANIUM DIOXIDE	110 Tons
FUEL Assly. WELD JOINTS	31,69,712 Nos

EIECTRIC POWER FROM ONE FUEL ASSEMBLY .... 10,44,000 units

DIMENSIONS IN MM

A typical Calandria Vessel for CANDU-PHWR 500 Unit (392horizontal fuel channels - each channel contain13 nos. of37 element fuel bundles)

(**102.36 ♦** x 495.3 I)





Main long-lived or parent Radionuclides present in Irradiated Fuel		
Radionuclide	Half-life (Year)	
Uranium		
U-234	$2.46 \times 10^5$	
U-235	$7.04 \ge 10^8$	
U-236	$2.34 \times 10^7$	
U-238	<b>4.47</b> x 10 <sup>9</sup>	
Actinides (α emitters)		
Pu-238	87.7	
Pu-239	24,100	
Pu-240	6,560	
Pu-241	14.35	
Pu-242	$3.74 \times 10^5$	
Nn-237	$2.14 \times 10^6$	
Am-241	432.7	
Am-243	7,368	
Cm-245	$8.5 \times 10^3$	
Cm-246	$4.73 \times 10^3$	
Fission Products (β/γ emitters)		
Se-79	6.5 x 10 <sup>5</sup>	
Zr-93	$1.5 \ge 10^6$	
Тс-99	$2.13 \times 10^5$	
Pd-107	$6.5 \ge 10^6$	
Sn-126	$1 \times 10^{5}$	
I-129	$1.57 \times 10^7$	
Cs-135	$2.3 \times 10^6$	
Activation Products (8/4 cmittors)		
Activation Froducts (p/y enniters)		
C-14	5,715	
Ni-59	$7.6 \times 10^4$	
Ni-63	100	
Zr-93	$1.53 \times 10^6$	
Nb-94	$2.03 \times 10^4$	



Fast Reactors in the world and their driver fuels+				
Country	Name₽	Туре₽	Power₽	Driver Fuel
United	DFR₽	Experimental+	+	U-Mo(HEU)+
Kingdom₽	PFR₽	Prototype 🕫	250 <u>MWe</u> ₽	(U, Py)O2+
Germany	KNK-II+	Experimental+	+	$(U, Py)O_2$ (HEU) +
	SNR-300#	Prototyppe ↓	300 <u>MWe</u> ≁	(U, Py)O2+
		(not operated)+		
India#	FBTR₽	Experimental+	40 <u>M Wi</u> +	(Pu <sub>0.7</sub> , U <sub>0.3</sub> )C+
	PFBR₽	Prototype⇔	500 <u>MWe</u> ₽	(U, Py) O2+
China₽	CEFR	CEFR +	4	(U, ₽y)O₂ (HEU)₽
		(under construction)+		
Korea	KALIMER₽	Demonstration +	Ð	(U, Py)O₂(HEU)₽
(Republic of)•	-	(under construction)+		
Russia₽	BR-5/BR-10+	Experimental +	5/10 <u>MWt</u> +	PuO <sub>2</sub> /UC/UN +
e.	BOR-60+	Experimental 🕶	60 <u>MW</u> +	UO2 (HEU) +
	BN-600₽	Commercial₽	600 <u>MWe</u> +	UO2 (HEU) +
	BN-350 (Kazakhstan)+	Prototype ≁	350 <u>MWe</u> ↓	UO2 (HEU) +
	BN-800#	Planned#	800 <u>MWe</u> +	$UO_1/(U_Pu)O_2^{+2}$
USA₽	EBR-1, EBR-II, 🛩	Experimental₽	+	U-Fs & U- <u>Pu-Zr</u> ↓
	FFTF₽	Experimental+	₽	(U, Pu)O2+
France₽	Rapsodie₽	Experimental₽	÷	$(U, P_{\mathcal{U}})O_2(HEU) \downarrow$
	Phenix+	Prototype≁	250 <u>MWe</u> +	$(U, Py)O_{2\downarrow}$
	SuperPhenix-I+	Commercial	1200 <u>MWe</u> ₽	(U, Py)O <sub>2</sub> -
		(shutdown)₽		
Japan₽	JOYO₽	Experimental+	÷	(U, Pu)O2 (HEU) +
	Monju₽	Prototype 🕫	230 MWe₽	(U.Pu)O₂₽

# Liquid Metal-cooled Fast Reactor Fuel Cycle with multiple recycling of U, Pu and Minor Actinides



ICTP Workshop Lecture 3 Conventional & AdvancedFuels (including inert matrix fuels)

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Trieste, 16 November, 2005

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# **Natural Resources for Nuclear Fuel, Fissile and Fertile Materials**

### NUCLEAR FUELS are:

- made of fissile isotopes U<sup>235</sup>, Pu<sup>239</sup> or U<sup>233</sup> mostly with judicious combination of fertile isotopes U<sup>238</sup> or Th<sup>232</sup>.
- used in the form of metals, alloys, oxides, carbides, nitrides, cer-met or cer-cer of the above fissile and fertile materials.
- used in the geometric shapes of rod, pin, plate or microsphere.
- encapsulated or clad with Zr-alloys (water-cooled reactor), SS (sodium-cooled fast reactor) or multilayer coatings of pyrolytic carbon and silicon carbide (high temperature gas-cooled reactor).







# **Conventional & Advanced Fuels For Nuclear Power Reactors**

Reactors	Conventional Fuels	Advanced/Alternative Fuels
Light Water Reactor (LWR): BWR, PWR & VVER		
Fuel (pellets)	LEU(U-235 ≤ 5 %) as UO2	LEU (U-235 5-10%) Mixed Uranium Plutonium Oxide (\$10% PuO2) [LEU+Minor Actinide (MA)] oxide for large grain size and controlled porosity 'Proliferation Resistant' spent fuel PuO2 in Inert Matrix for burning 'Pu'
Cladding	Zircaloy 2 (BWR) Zircaloy 4 (PWR) Zr-1% Nb (VVER)	Zr-Sn-Nb-Fe & Zr-Nb-O alloys
Burning up	20 000-30 000 MWD/t	High : up to 60 000 MWD/t Ultra High : up to 80 000 MWD/t
Pressurized Heavy Water Reactor (PHWR)		
Fuel (pellets)	Natural UO2	REU, SEU in the form of UO2, (U,Pu)O2 (Th,Pu)O2 & (Th,U233)O2, containing up to 2% fissile material. Large grain size and controlled porosity PuO2 in Inert Matrix for burning 'Pu'
Cladding	Zircaloy 4	Zircaloy 4
Burnup	6 700 MWD/t	15 000 – 20 000 MWD/t



# **Conventional & Advanced Fuels For Nuclear Power Reactors**

Liquid Metal-cooled Fast Breeder Reactor (LMFBR)		
Fuel (pellets/particles/pins)	HEU in the form of UO2 & (U,Pu)O2 (≤25% Pu) He-bonded pins	Na-bonded (U,Pu)C, (U,Pu)N & U-Pu-Zr, (≤25% Pu) fuel with/without MA He-bonding also for carbide/nitride (PuO2+ThO2) for burning 'Pu' He-bonded vibratory compacted oxide, carbide and nitride fuel pins 'Pu and (Pu,MA) in inert matrix for burning (U/Th+MA) in blanket for 'Proliferation Resistance" in irradiated blanket
Cladding	Stainless Steel D-9	Stainless steel (type ferritic HT-9 or Oxide dispersed ODS)
Burnup	100 000 MVVD/t	Up to 200 000 MWD/t >1.00 up to 1.5
Breeding ratio	1.0 – 1.2	1.2-1.6
High Temperature Gas Cooled Reactors (HTR) (coated microspheres)	Multi-layer (pyrolytical carbon & SiC-coated) Uranium Oxide fuel particles (BISO or TRISO) embedded in graphite	Multi-layer (pyrolytical carbon& ZrC coated) Uranium Oxide, Mixed Uranium Plutonium Oxide, Mixed Uranium Thorium Dicarbide, embedded in graphite





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Figure 9. Three-stage nuclear power programme in India involving 'closed fuel cycles'





### **Key Features**

- Thorium bearing fuel  $[(Th + Pu)O_2 Mox, (Th$  $+ ^{233}UO_{2}Mox$
- Hollow cylindrical (ZrO<sub>2</sub>-Dy<sub>2</sub>O<sub>3</sub>) displacer rod.
- Emergency core cooling water injected into the cluster through the holes in displacer rod.
- Low pressure drop design.

AEA









### **AHWR FUEL CLUSTER (D5)**



# **Advanced fuel cycle schemes**



### COMMERCIAL SPENT URANIUM OXIDE FUEL REPROCESSING PLANTS IN OPERATION AND UNDER CONSTRUCTION IN THE WORLD

Country / Company	Facility / Location	Fuel Type	Capacity (tHM/year)
France, COGEMA	UP2 and UP3, La Hague	LWR	1700
UK, BNFL	Thorp, Sellafield	LWR, AGR	1200
UK, BNFL	B205 Magnox	Magnox GCR	1500
Russian Federation, Minatom	RT-1 / Tcheliabinsk-65 Mayak 400	VVER	400
Japan, JNC	Tokai-Mura	LWR, ATR	90
Japan, JNFL	Rokkasho-Mura (under construction)LWR		800
India, BARC	PREFRE-1, Tarapur PREFRE-2, Kalpakkam	PHWR PHWR	100 100
China, CNNC	Diowopu (Ganzu)	LWR	25-50



### MIXED URANIUM PLUTONIUM OXIDE (MOX) FUEL FABRICATION FACILITIES

			Capacity
<b>Country / Company</b>	Facility / Location	Fuel Type	(tHM/year)
France, COGEMA	Cadarache	Cadarache LWR, FBR	
France, COGEMA	Marcoule-Melox	Marcoule-Melox LWR	
Belgium, Belgonucleaire	Dessel	LWR	40
UK, BNFL	Sellafield SMP	LWR	120
UK	Sellafied MDF	LWR	8
Russian Federation, Minatom	Chelyabinsk	FBR	60
Japan, JNC	Tokai-Mura	ATR	10
Japan, JNFL	, JNFL Rokkasho		130
India, AFFF, BARC	Tarapur	LWR, PHWR & FBR	

# The GANEX concept : <u>Group ActiNides EXtraction</u>





### **Objectives of advanced methods of fabrication of ceramic nuclear fuel pellets**

Safety	Economics	Performance
• Avoid generation and handling of powder of fuels	• Minimise process steps	• Tailor make fuel microstructure for higher burn up
for minimising :	• Reduce fuel synthesis & sintering temperatures	<ul> <li>High density (≥96% T.D.), closed</li> </ul>
<ul> <li>radiotoxic dust hazard</li> <li>fire hazard (for carbide &amp;</li> </ul>	• Reduce gas cost during synthesis and sintering	"porosity" and large (>25µ) grain size for LWR & PHWR
nitride fuels)	- gas purification and recirculation	<ul> <li>Low density (&lt;85% T.D.) "open" porosity and small (&lt;5µ) grain size</li> </ul>
• Fabrication flow sheet should be amenable to automation & remotisation	- alternative less expensive gas	for LMFBR - Excellent micro-homogenity of
- for minimising personnel	• Reduce process losses and rejects	fissile material in fuel
exposure to radiation		<ul> <li>avoid fine pores (&lt;1µ) for minimising in-pile densification</li> </ul>



### "Sol-Gel-Microsphere Pelletization (SGMP)" and "Vibro-Sol" Processes for Manufacturing Mixed Uranium Plutonium Oxide (MOX), Monocarbide (MC) and Mononitride (MN) Fuels for LMFR



![](_page_55_Picture_0.jpeg)

![](_page_56_Picture_0.jpeg)

# <section-header> Microstructure & Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysis of ThO2-2%UO2 prepared by Sol-Gel Microsphere Pelletisation (SGMP) Process Image Analysi

**As-Polished Microstructure** 

0031 100.0 µm

**Porous Microsphere (SEM Picture)** 

25 kV 120×

![](_page_57_Figure_2.jpeg)

![](_page_57_Figure_3.jpeg)

**Etched Microstructure** 

# **OBJECTIVES OF INERT MATRIX FUEL**

- Minimizing "proliferation risk" of plutonium (~ 200 tons of weapon-grade and ~ 1000 tons civilian grade) by using them in nuclear power reactors in operation
- Minimizing "Minor Actinides" (MA: Np, Am & Cm) and in turn radiotoxicity in waste
- In some cases minimizing 'proliferation risk' of weapon-grade (> 90 % <sup>235</sup>U) uranium (though conventional process is down-blending)

![](_page_58_Picture_4.jpeg)

### Inert Matrix

- > Neutron (very low capture and absorption cross-sections)
- > Chemical compatibility with
  - $\rightarrow$  Fuel
  - $\rightarrow$  Cladding
  - $\rightarrow$  Coolant
- Consideration of direct disposal after use

### Fuel

- > 'Plutonium form' alloys and compounds
- > Utilization of Minor Actinides together with plutonium
- Weapon-grade HEU (<sup>235</sup>U > 90 %) –alloys or compounds

![](_page_59_Picture_11.jpeg)

# **EXAMPLES OF INERT MATRIX**

**Inert Matrix type** Element Inter-metallics Alloy Carbide **Nitrides Binary oxide Ternary oxide** Oxide solid solution

Inert Matrix formula C, Mg, Al, Si, Cr,V, Zr, Mo, W AlSi, AlZr, ZrSi Stainless steel, zirconium alloys SiC, TiC, ZrC AIN, TiN, ZrN, CeN, MgO, Y<sub>2</sub>O<sub>3</sub>, ZrO<sub>2</sub>, CeO<sub>2</sub> MgAl<sub>2</sub>O<sub>4</sub>, Y<sub>3</sub>Al<sub>5</sub>O<sub>12</sub>, ZrSiO<sub>4</sub>  $Y_{v}Zr_{1-v}O_{2-v/2}, Mg_{(1-x)}Al_{(2+x)}O_{(4-x)}$ 

![](_page_60_Picture_3.jpeg)

# **Example of heterogeneous materials as IMF's**

Design	Composition
Solid solution	An <sub>z</sub> Y <sub>y</sub> Pu <sub>x</sub> Zr <sub>1-y</sub> O <sub>2-ψ</sub> *
Cercer	MgAl <sub>2</sub> O <sub>4</sub> - Y <sub>y</sub> Pu <sub>x</sub> Zr <sub>1-y</sub> O <sub>2-y/2</sub> *
Cermet	Zr - Y <sub>y</sub> Pu <sub>x</sub> Zr <sub>1-y-x</sub> O <sub>2-y/2</sub> *
Metmet	PuAl <sub>4</sub> *-Al

Examples of	f Inert M	Aatrix design	and additives
-------------	-----------	---------------	---------------

Additive type Additive formul
-------------------------------

Burnable poison

Resonance additive

Stabiliser

B, Gd, Dy, Ho, Er, Eu

Fe, W, Th, U

 $Y_2O_3$ , CaO in ZrO<sub>2</sub> Al<sub>2</sub>O<sub>3</sub> in SiC

![](_page_61_Picture_10.jpeg)

![](_page_62_Figure_0.jpeg)

# Irradiation behavior of rock-like oxide fuels

Pin	Maximum temperature (K)	$\Delta \Phi/\Phi$ (%)	$\Delta V/V$ (%)	Xe FGR (%)
ZM-7	1040	_	_	63
ZM-6	1270	_	_	61
ZM-4	980	10.8	_	63
SD	1850	2.7	5.5	38
SH	2080	5.0	10.2	22
Z	1580	2.0	<4.0	2.2
CD	1930	2.1	4.3	22
СН	1830	2.8	5.7	7.8

Pin	Compositi	Composition (mol%)					Fissile density
	YSZ <sup>a</sup>	PuO2 <sup>b</sup>	UO2 <sup>c</sup>	$MgAl_2O_4$	$Al_2O_3$	size	$(10^{20}/\text{cm}^3)$
ZM	16.7	11.1	_	11.1	61.1	2–10 µm	21.10
SD	20.0	_	37.1	42.9	_	250 µm	13.00
SH	20.0	_	37.1	42.9	_	10–50 µm	13.10
Z	80.0	_	20.0	_	_	Solid solution	8.62
CD	16.5	_	30.6	_	52.9	250 µm	13.54
СН	16.5	_	30.6	_	52.9	10–50 µm	13.17

Dimensional variation and fractional gas release of ROX fuels

<sup>a</sup> YSZ = 79.9 mol% ZrO<sub>2</sub> + 20.1 mol% YO<sub>1.5</sub>. <sup>b</sup> Pu isotopic composition (at%) was 94.3, 5.3 and 0.4 for <sup>239</sup>Pu, <sup>240</sup>Pu and <sup>241</sup>Pu, respectively.

° 19.6% enriched UO<sub>2</sub>.

Ref; Yamashita et.al., Journal of Nuclear Materials 320 (2003) 126–132

![](_page_63_Picture_7.jpeg)

# R&D on Advanced LMFR Fuels and Advanced Methods of Fuel Fabrication

- Ceramic Nuclear Fuels Conventional: (U, Pu)O<sub>2</sub> Advanced: (U, Pu)C & (U, Pu)N with/without Minor Actinides
- Advanced methods of fabrication of ceramic fuels: Dust-free advanced fabrication processes like vibratory compaction, vibro-sol & sol-gel microsphere pelletization
- Metallic Fuels: U-Pu-Zr, Th-U-Pu-Zr & U-Pu (for high breeding)
- Fuel Cladding, Hexcans & Other Fuel Assembly components: Ferritic stainless steel HT9 & Oxide dispersed stainless steel with minimum radiation damage and void swelling
- Advanced fabrication processes should be amenable to secured automated fabrication, real-time accounting of special nuclear material and proliferation resistance

![](_page_64_Picture_6.jpeg)

# **Conventional and Advanced Methods of Spent Fuel Reprocessing**

### **AQUEOUS PROCESS :**

Dissolution of spent fuel in Nitric acid followed by purification by solvent extraction by adapting the PUREX process, using TriButyl Phosphate (TBP) as solvent, is being used on an industrial scale for reprocessing of spent UO2 and MOX fuels. The PUREX process is not suitable for mixed carbide fuel but could be utilized for reprocessing mixed nitride and metallic fuels.

Modifications are being incorporated in PUREX process to make it proliferation resistant and economic.

### **PYROPROCESSING**:

### - Pyroprocessing involving electrolytic reduction

This route has been initially developed on a pilot plant scale for reprocessing of spent metallic fuels (U-Zr & U-Pu-Zr) in USA and was successfully extended on a laboratory scale for reprocessing of carbide and nitride fuels. The pyroprocessing route is yet to be adapted on an industrial scale.

In recent years, the Russian Federation has successfully demonstrated the pyroprocessing route for reprocessing of spent oxide fuels on a pilot plant scale.

### - Pryoprocessing involving fluoride volatilization

The process includes fluorination followed by distillation. The method has so far been demonstrated on a laboratory scale only.

![](_page_65_Picture_10.jpeg)