

# Chapt. 10 Fuel Element Performance

## 10.1) Important Features

### Oxide fuel characteristics

- **Temperature ( $T_{max} \approx 2800$  °C)**
  - **Sintering**
  - **Grain growth**
  - **Diffusion**
- $\frac{\Delta T}{\Delta x}$ 
  - **Stress**
  - **Pore closing**
  - **FP redistribution**

### 10.1.1) Oxide Fuels- (started $\approx$ 1955) (Fig/ Table)

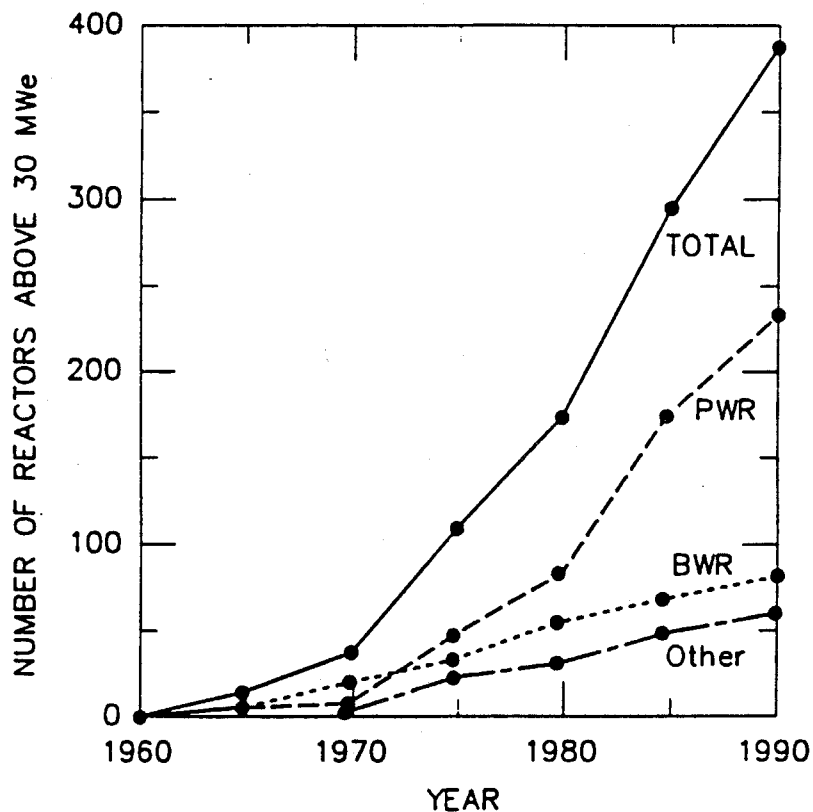
*Most important properties of mixed oxides depend on the O/M ratio*

$$q = \frac{N_{pu}}{\sum N_{\text{heavy metal}}}$$

$$x = \frac{N_{\text{excess.oxygen}}}{\sum N_{\text{heavy.metal}}}$$

for example.....  $(U_{1-q}Pu_q)O_{2+x}$

**Table 3-3.** Limiting factors in early LWR fuel performance (1965–1975)<sup>a</sup>.

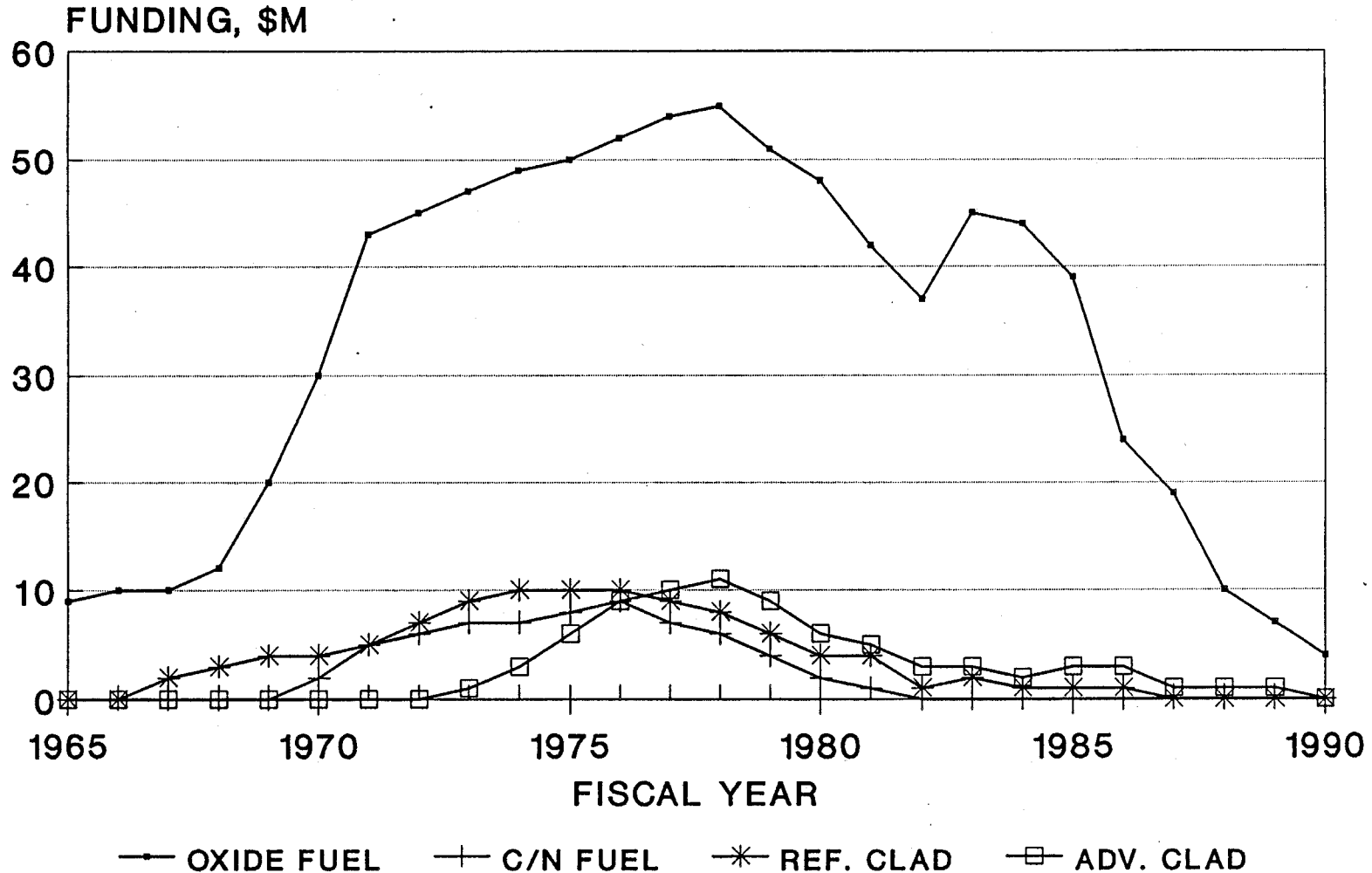


**Figure 3-4.** Worldwide growth for 1960–1990 in oxide-fueled power reactors above 30 MWe. Courtesy: *Nuclear News* (1990).

Factor	PWR	BWR	Remedies
Hydriding of zirconium	×	×	eliminate fuel moisture; add getters
Scale deposition		×	eliminate copper tubing from feedwater heaters
Enrichment errors	×	×	gamma-scan elements before shipment
Cladding collapse	×		prepressurize elements; use stable pellets
Pellet densification	×	×	stable fuel pellet microstructures
Manufacturing defects (e.g., faulty welds)	×	×	improve QC (now below 5% of in-reactor defects)
Cladding corrosion/fretting		×	rare; improve QA and cleaning; use element spacers
Fuel element growth/bowing	×		control texture, axial clearances, spacer design
Channel bulging		×	thicker channel walls. control residual stress
Pellet-cladding interactions (PCI)	×	×	(1) slow power rise (PWR) (2) power shape control (BWR) (3) fuel “preconditioning” phase (both) (4) pellet design changes (both)

<sup>a</sup> From Levenson and Zebroski (1976).

# PROGRAM ANNUAL COSTS



# ***PROGRAM DURATION AND COST***

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<i>Program Element</i>	<i>Period</i>	<i>Duration</i>	<i>Cost</i>
<b>Oxide Fuel</b>	<b>1965-1992</b>	<b>27 yrs</b>	<b>\$ 846M</b>
<b>Carbide/Nitride Fuel</b>	<b>1969-1982</b>	<b>13 yrs</b>	<b>64M</b>
<b>Reference Alloy</b>	<b>1966-1987</b>	<b>21 yrs</b>	<b>101M</b>
<b>Advanced Alloys</b>	<b>1973-1990</b>	<b>17 yrs</b>	<b>77M</b>
		<b>TOTAL</b>	<b>\$ 1,088M</b>

**Hyperstoichiometry**     $O/M > 2.00$

**Hypostoichiometry**     $O/M < 2.00$

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***Disadvantages of Oxide Fuels***

- 1.) Low U density (larger core)
- 2.) Low thermal conductivity  
( requires thin rods, produces high T)

Why not use UC, UN, UP, US, etc.?

**10.1.2 Fission Properties**

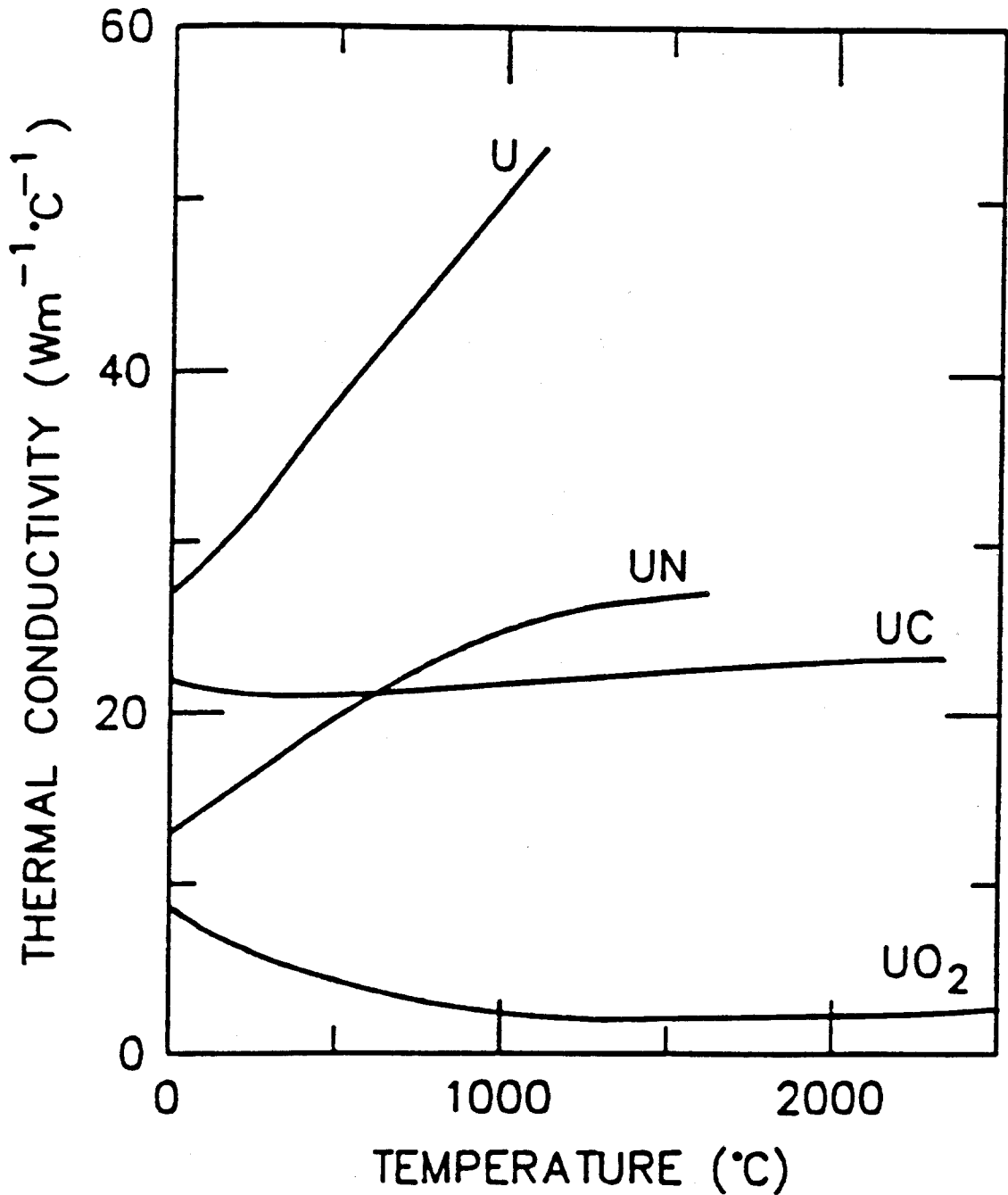
**Burnup**

$$\beta = \frac{\text{Number of Fissions}}{\text{Initial Number of Heavy Metal Atoms}}$$

**Rule of Thumb**

$$1 \text{ at\% B.U.} \approx 10,000 \frac{\text{MWd}}{\text{MtU}} \quad @ \quad 200 \frac{\text{MeV}}{\text{fission}}$$

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***Fissioning of 1 g of fissile material  $\approx$  1MWd***



**Figure 3-2.** Thermal conductivities of major nuclear fuels.

**Maximum burn up with no Pu credit;**

- *Natural U*                      **6650**  $\frac{\text{MWd}}{\text{MtU}}$
- *15% enriched*    **142,500**  $\frac{\text{MWd}}{\text{MtU}}$

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## Major Differences Between LWR & LMFBR Fuels

	<u>LWR</u>	<u>LMFBR</u>
<b>Ave neutron energy, eV</b>	<b>0.03</b>	<b>500,000</b>
<b>Fissile isotope</b>	<b><math>^{235}\text{U}</math></b>	<b><math>^{239}\text{Pu}</math></b>
<b><math>\sigma_f</math>, barns</b>	<b>550</b>	<b>1.8</b>
<b>Neutron flux, (rel)</b>	<b>1</b>	<b>300</b>
<b>Power density, (rel)</b>	<b>1</b>	<b>3 (why?)</b>
<b>Burn up, %</b>	<b>3</b>	<b>10</b>
<b>Enrichment, %</b>	<b>3</b>	<b>15</b>

## Differences Between LWR and LMFBR Fuel Assemblies

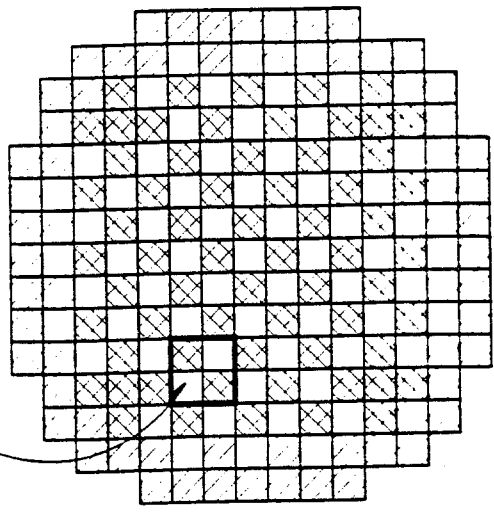
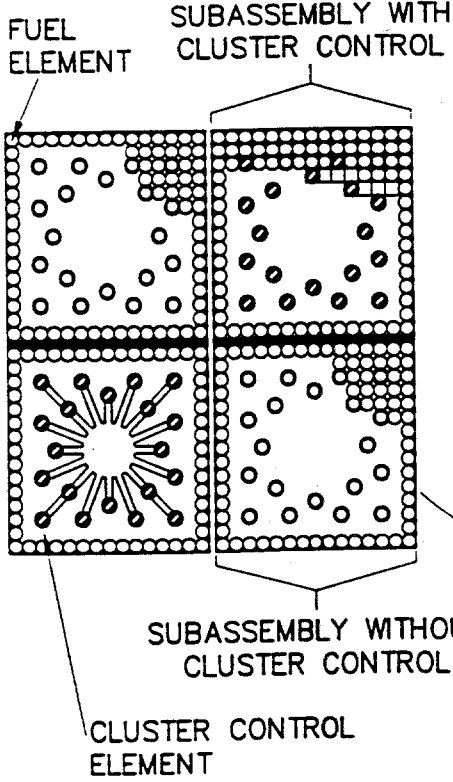
	<b>LWR</b>	<b>LMFBR</b>
<b>Damage to clad</b>	<b>1</b>	<b>100</b>
<b>Fuel pin diameter, mm</b>	<b>11</b>	<b>6</b>
<b>Core fuel fraction</b>	<b>-</b>	<b>Higher</b>
<b>Cladding</b>	<b>-</b>	<b>Hotter</b>
<b>Size of Core</b>	<b>-</b>	<b>Smaller ( no moderator needed)</b>

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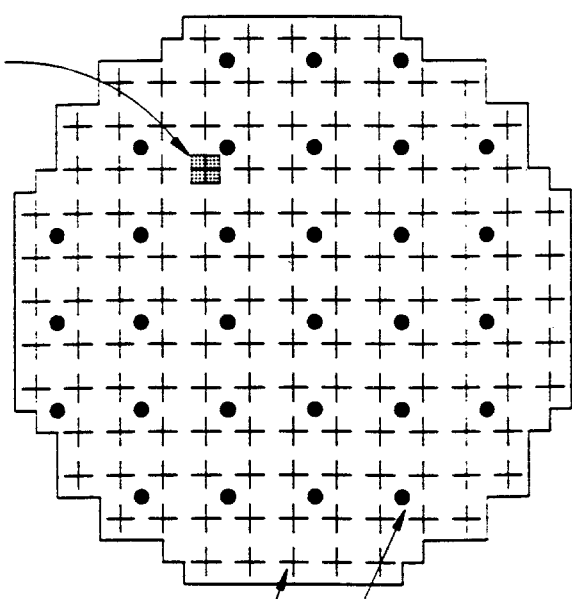
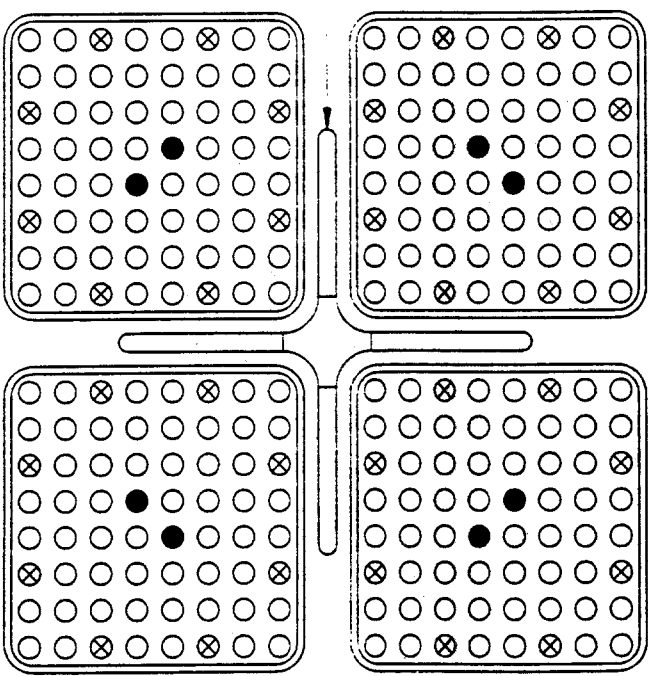
**See Table 10.2 and figures 10.1 thru 10.4**



<b>Table 10.2 Comparison of Fuel Element Characteristics</b>		
	<b>Thermal Reactor</b>	<b>Fast Reactor</b>
<b>Fuel</b>	<b>UO<sub>2</sub></b>	<b>(U,Pu)O<sub>1.96</sub></b>
<b>Fuel Pellet Density (% of theoretical)</b>	<b>92</b>	<b>90</b>
<b>Max. fuel centerline temperature (overpower condition)°C</b>	<b>2450</b>	<b>2800</b>
<b>Cladding</b>	<b>Zircaloy-4</b>	<b>316 Stainless Steel</b>
<b>Max. cladding mid- wall temperature °C</b>	<b>380</b>	<b>660</b>
<b>Coolant temperature, °C</b>	<b>H<sub>2</sub>O, 280-320</b>	<b>Na, 470-650</b>
<b>Maximum rod linear power, W/cm</b>	<b>620</b>	<b>550</b>
<b>Fuel wrapper assembly</b>	<b>Square, 30x30</b>	<b>Hexagonal, 13 cm across flats</b>
<b># of pins in assembly</b>	<b>200</b>	<b>220</b>
<b>Fuel-rod outside diameter, mm</b>	<b>10.7</b>	<b>6.3</b>
<b>Cladding thickness, mm</b>	<b>0.6</b>	<b>0.4</b>
<b>Initial fuel-cladding radial gap, mm</b>	<b>0.08</b>	<b>0.07</b>
<b>Length of fueled portion, cm</b>	<b>365</b>	<b>90</b>

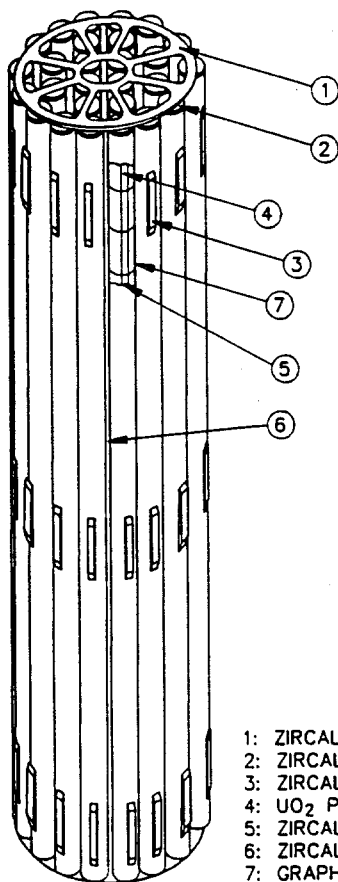


— CONTROL BLADE



- FUEL ELEMENT
- WATER ROD
- ⊗ TIE ELEMENT

Figure 3-8. Schematic of BWR fuel subassembly and reactor core.



- 1: ZIRCALOY END PLATE.
- 2: ZIRCALOY END CAP.
- 3: ZIRCALOY BEARING PADS.
- 4:  $UO_2$  PELLETS.
- 5: ZIRCALOY CLADDING.
- 6: ZIRCALOY SPACERS.
- 7: GRAPHITE COATING.

Figure 3-9. Schematic of CANDU fuel bundle. Courtesy: Atomic Energy of Canada.

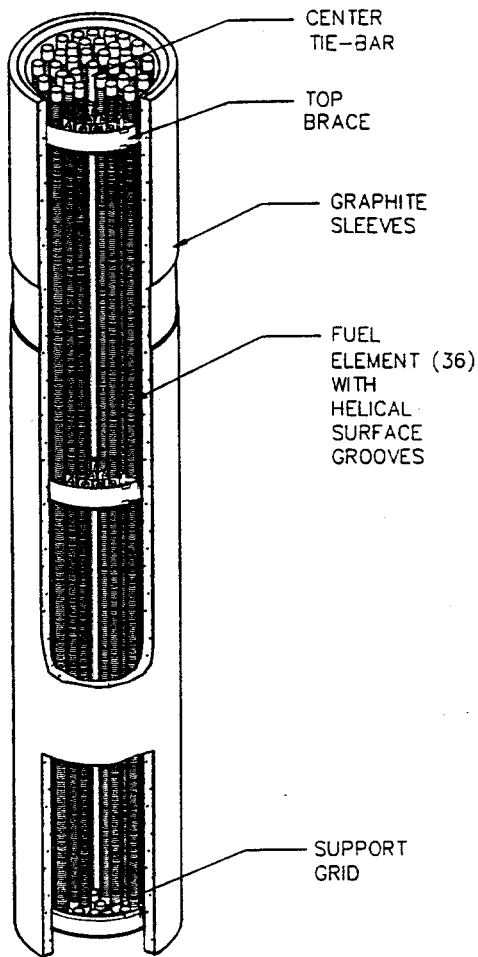


Figure 3-10. Schematic of an AGR subassembly and fuel stringer. Courtesy: United Kingdom Atomic Energy Authority.

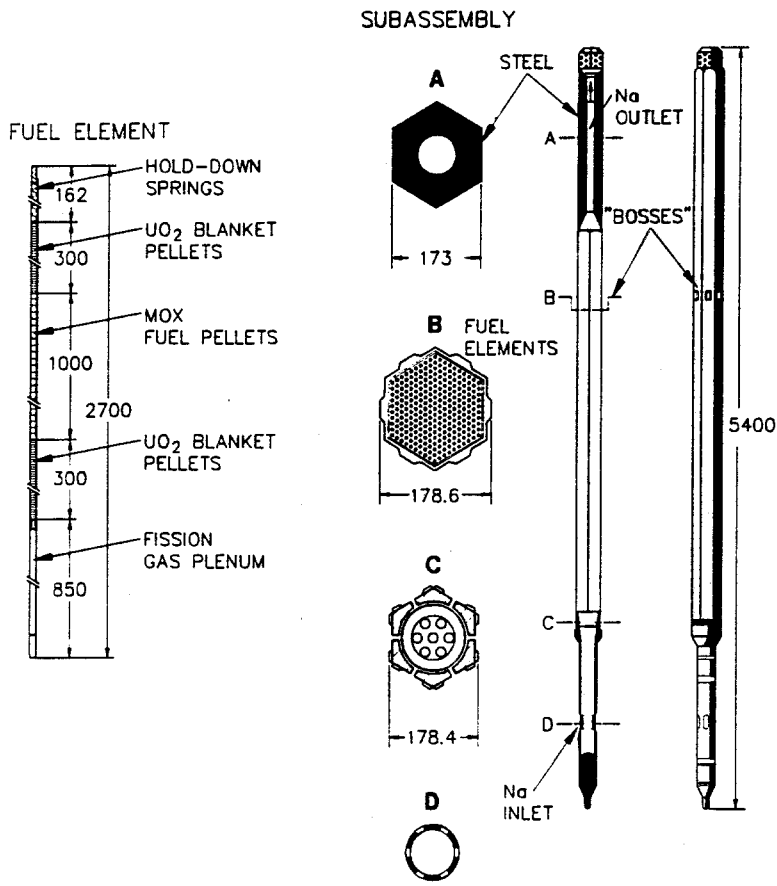


Figure 3-11. Schematic of an FBR fuel element and subassembly for the Superphenix-1 reactor. Courtesy: Commissariat à l'Énergie Atomique.

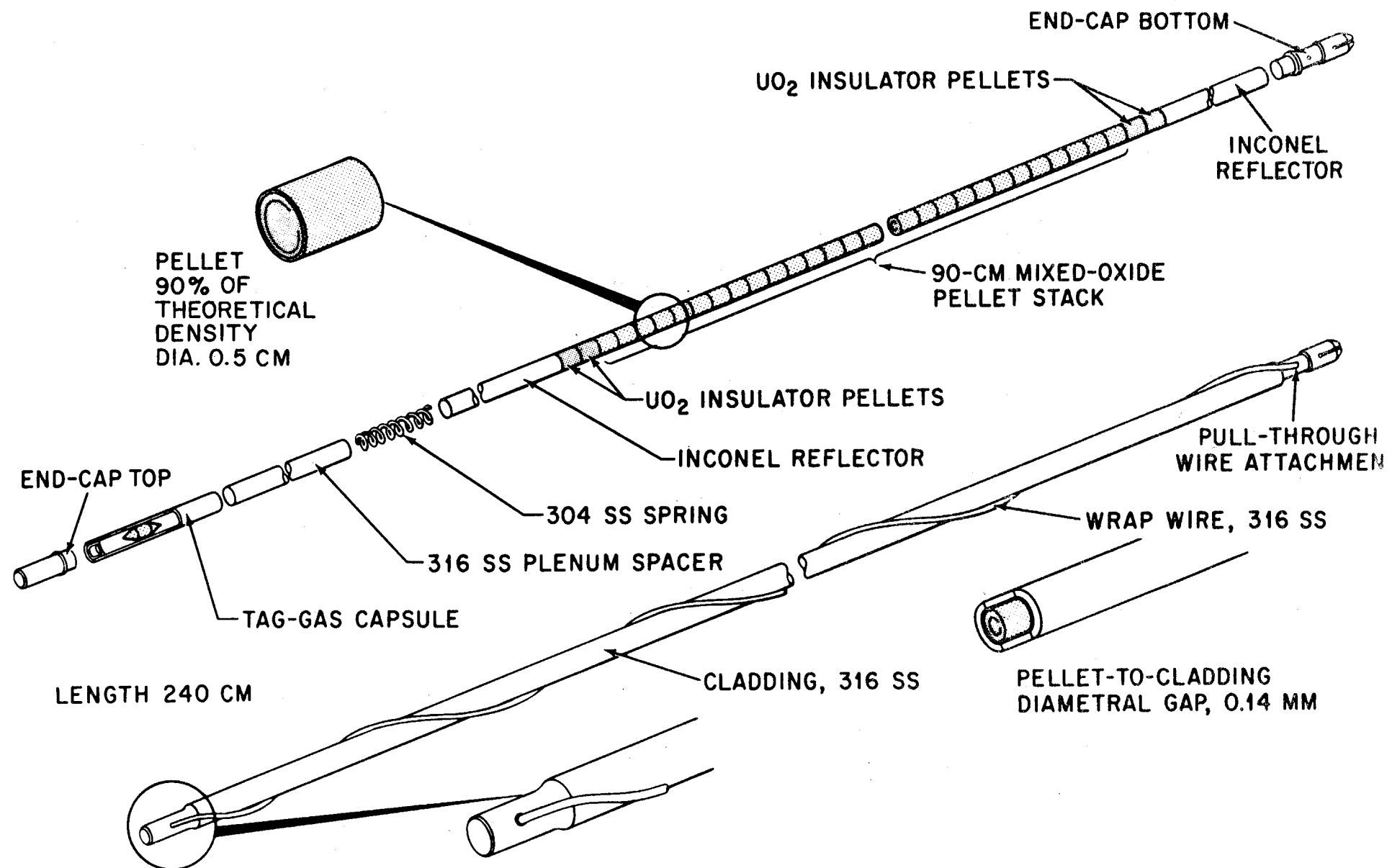
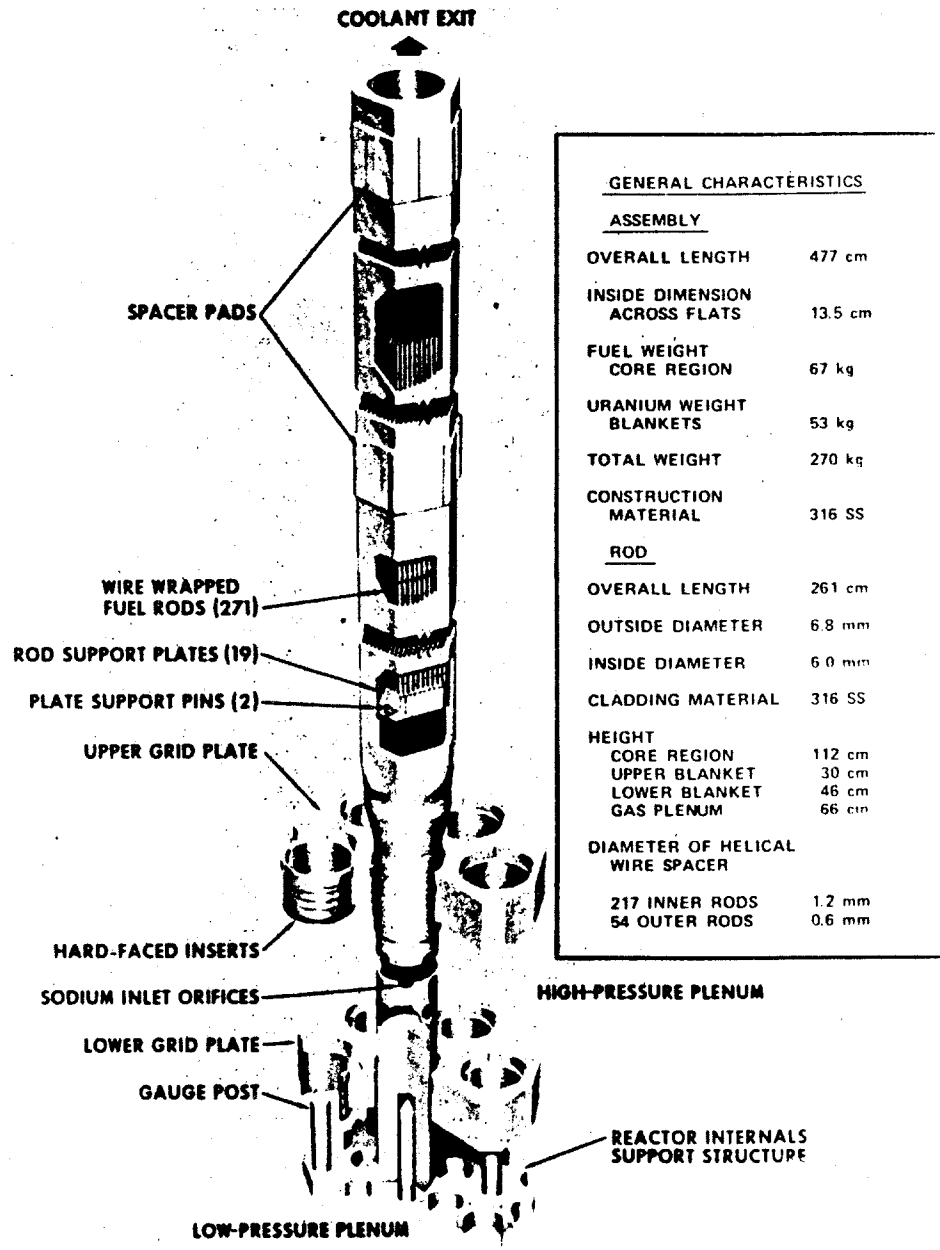
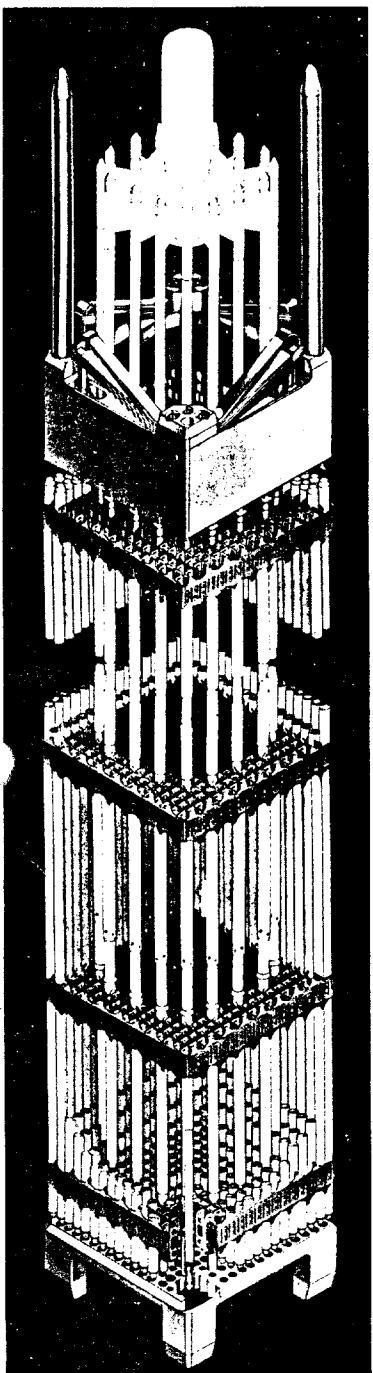


Fig. 10.2 Fuel pin of the Fast Test Reactor. (Courtesy C. Burgess, Hanford Engineering Development Laboratory.)



GENERAL CHARACTERISTICS	
<u>ASSEMBLY</u>	
OVERALL LENGTH	477 cm
INSIDE DIMENSION ACROSS FLATS	13.5 cm
FUEL WEIGHT CORE REGION	67 kg
URANIUM WEIGHT BLANKETS	53 kg
TOTAL WEIGHT	270 kg
CONSTRUCTION MATERIAL	316 SS
<u>ROD</u>	
OVERALL LENGTH	261 cm
OUTSIDE DIAMETER	6.8 mm
INSIDE DIAMETER	6.0 mm
CLADDING MATERIAL	316 SS
HEIGHT	
CORE REGION	112 cm
UPPER BLANKET	30 cm
LOWER BLANKET	46 cm
GAS PLENUM	66 cm
DIAMETER OF HELICAL WIRE SPACER	
217 INNER RODS	1.2 mm
54 OUTER RODS	0.6 mm

Fig. 10.4 LMFBR fuel assembly. (Courtesy L. Bernath, Atomics International.)

Fig. 10.3 Pressurized-water-reactor fuel assembly. (Courtesy Westinghouse Company.)

## 10.2 Thermal Properties

### 10.2.1 Melting Points

$$T_{MP} \text{ of } UO_2 = 2865 \text{ }^\circ\text{C}$$

( 2847 °C in some papers)

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### Several Factors Influence the Melting Point

- *Stoichiometry (phase diagram)*
  - *Mixed Oxide ,UO<sub>2</sub> -PuO<sub>2</sub> , (2 figs see correc.)*
  - *Burn -Up (figure + table)*
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### 10.2.2 Thermal Expansion

*Problems -Cladding stress, Poor heat transfer*

**Stoichiometry Effect-Figure 10.8**

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### 10.2.3 Specific Heat ( See Chapter 1)

(Important for dynamic behaviour,  $\frac{k}{\rho C_p}$ )

$$C_p = \left( \frac{\delta H}{\delta T} \right)_p = C_v + \left( \frac{\alpha^2 V}{\beta} \right) T$$

thermal expansion    Molar Volume

compressibility

Figure 10.10

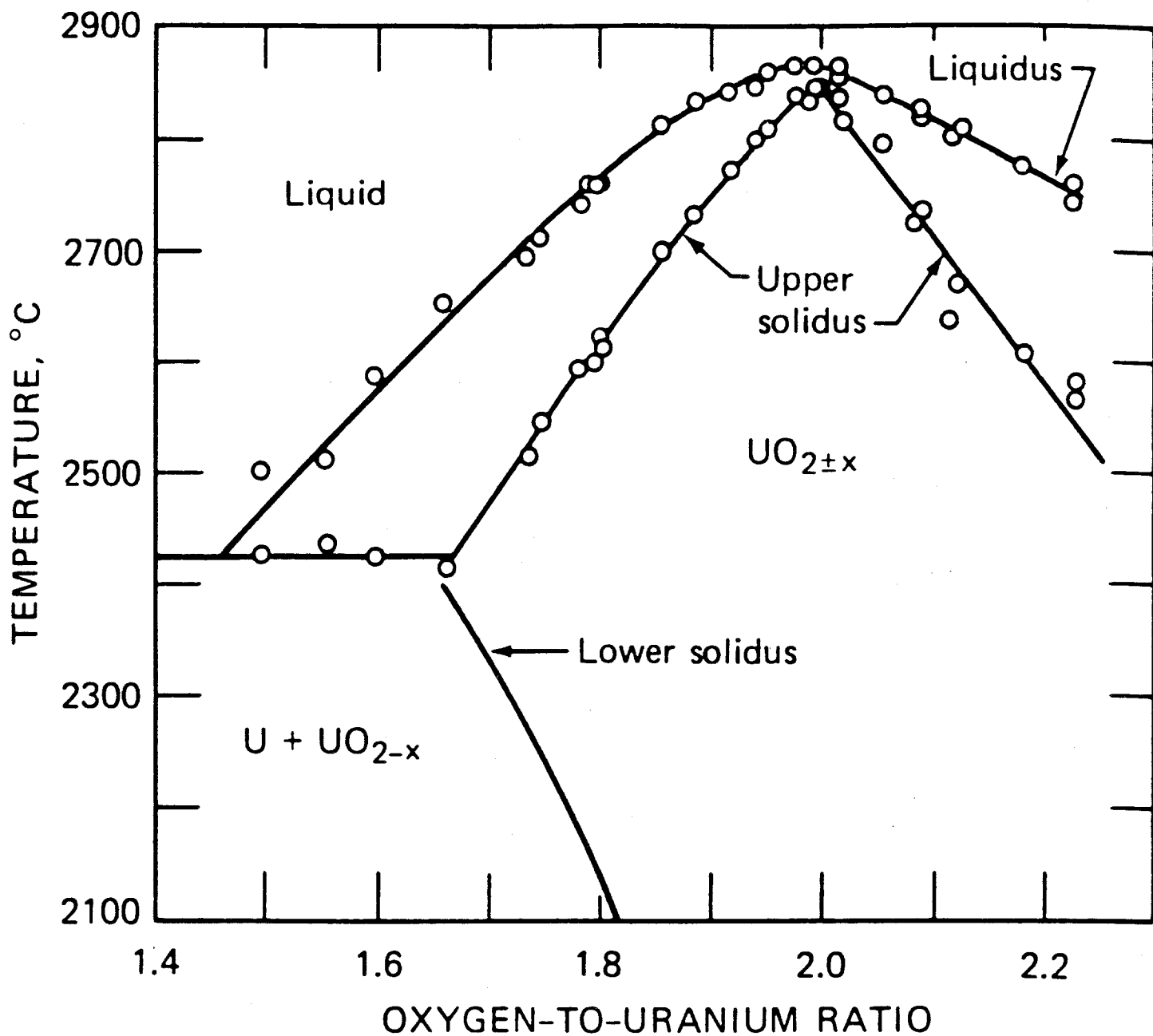
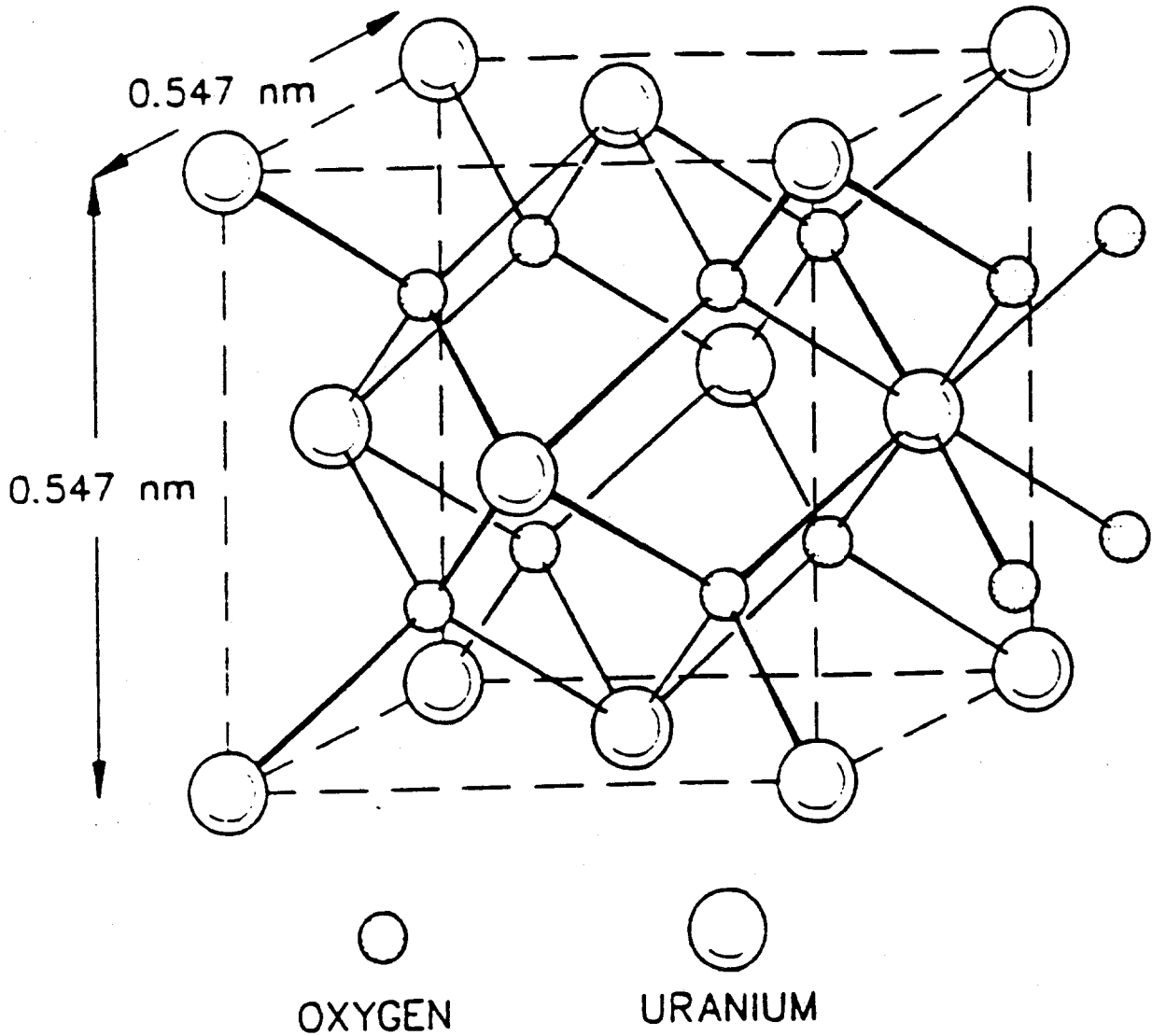


Fig. 10.5 Partial phase diagram for urania from  $\text{UO}_{1.5}$  to  $\text{UO}_{2.23}$ . The separation of the peaks of the liquidus and solidus curves at  $\text{O}/\text{U} = 2.0$  is undoubtedly due to measurement errors. The  $\text{UO}_2$  melts congruently; thus, the curves should coincide for  $\text{UO}_{2.0}$ . Similarly, the lower solidus curve should intersect the corner of the upper solidus and horizontal lines. [From R. E. Latta and R. E. Fryxell, *J. Nucl. Mater.*, 35: 195 (1970).]



**Figure 3-1.** Unit cell of stoichiometric  $\text{UO}_2$ .



FUEL-ELEMENT THERM

CHANGE IN MELTING POINT, °C

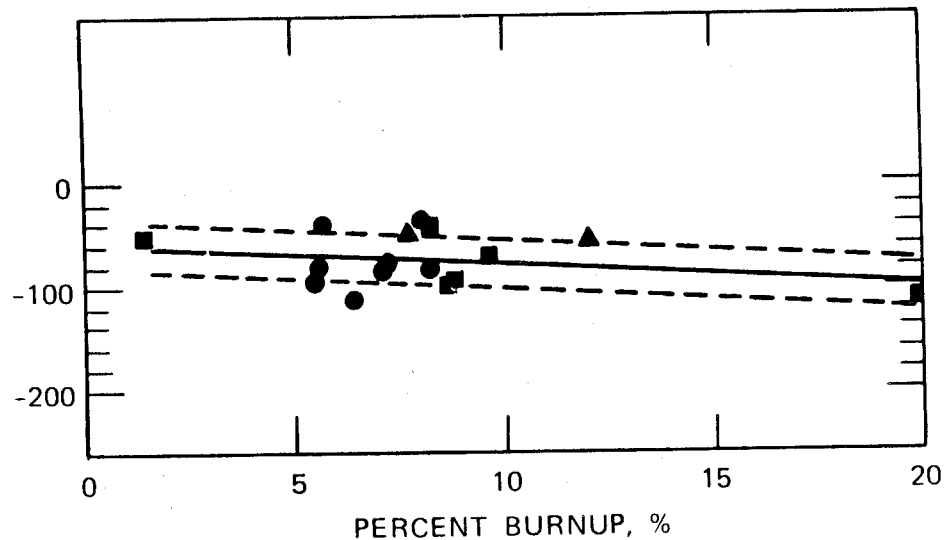


Fig. 10.7 Effect of burnup on the melting point of mixed-oxide fuel material. The dashed lines delineate a band  $\pm$  one standard deviation wide. ■, 25% PuO<sub>2</sub>, O/M = 2.00. ▲, 25% PuO<sub>2</sub>, O/M = 1.96. ●, 20% PuO<sub>2</sub>, O/M = 2.00. [From A. Biancheria, U. P. Nayak, and M. S. Beck, in *Proceedings of the Conference on Fast Reactor Fuel Element Technology*, R. Farmakes (Ed.), p. 361, American Nuclear Society, Hinsdale, Ill., 1971.]

TEMPERATURE, °C

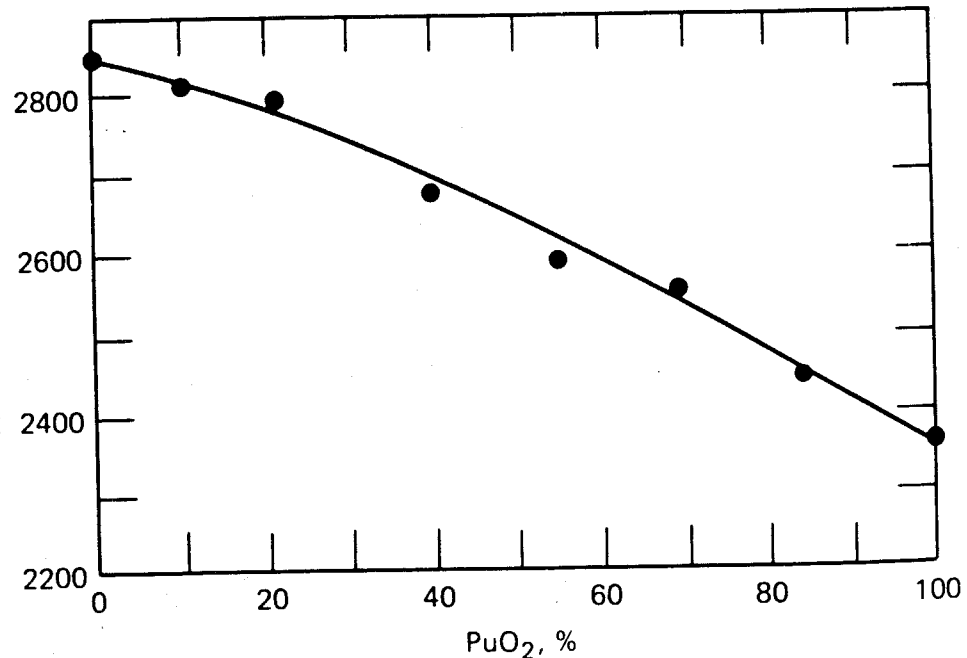


Fig. 10.6 Melting points of mixed uranium-plutonium oxides. (From E. L. Zebroski, W. L. Lyon, and W. E. Bailey, Effect of Stoichiometry on the Properties of Mixed Oxide U-Pu Fuel, in *Proceedings of the Conference on Safety, Fuels, and Core Design in Large Fast Power Reactors*, Oct. 11-14, 1965, USAEC Report ANL-7120, p. 382, Argonne National Laboratory, 1965.)

**Corrections to Olander Book**  
**from Nuclear System Handbook (NSH)**

**Melting Point, °C**

<b>Mole Fraction PuO<sub>2</sub></b>	<b>Solidus</b>	<b>NSH Liquidus</b>	<b>Liquidus Olander</b>
<b>0.0</b>	<b>2847</b>	<b>2847</b>	<b>2865</b>
<b>0.2</b>	<b>2728</b>	<b>2767</b>	
<b>0.4</b>	<b>2632</b>	<b>2685</b>	<b>≈2700</b>
<b>0.6</b>	<b>2553</b>	<b>2600</b>	<b>≈2600</b>
<b>0.8</b>	<b>2487</b>	<b>2530</b>	
<b>1.0</b>	<b>2428</b>	<b>2428</b>	<b>≈2360</b>

**Density:**

$$\rho = \frac{\text{MW metal atoms}}{\text{MW nat. U}} \left[ 5875.5 + 4.97(\% \text{PuO}_2) + 2540 \frac{\text{O}}{\text{M}} \right] \frac{\text{kg}}{\text{m}^3}$$

# Effect of Burnup on MP of Mixed UO<sub>2</sub>-PuO<sub>2</sub>

For 20-25% PuO<sub>2</sub>  
from NSH

<u>Burn up</u> <u>MWd/MtU</u>	<u>Change in</u> <u>Melting Point °C</u>
15,000	-62
25,000	-64
50,000	-69
75,000	-74
100,000	-80

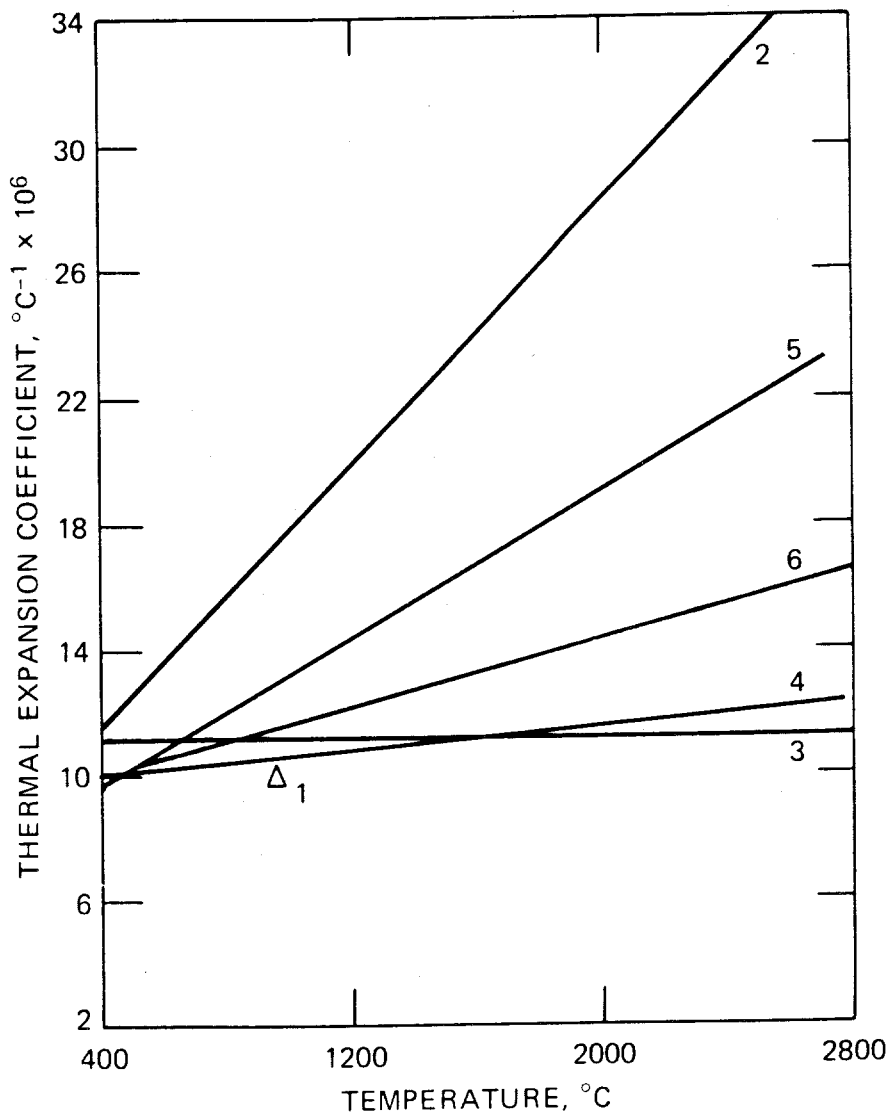


Fig. 10.8 Thermal expansion coefficients of mixed-oxide fuels.

1.  $(U_{0.8}Pu_{0.2})O_2$ , R. P. Nelson, USAEC Report BNWL-473, 1967.

2.  $(U_{0.8}Pu_{0.2})O_{2.10}$ , J. Roth and E. K. Halteman, USAEC Report NUMEC-2389-9, 1965.

3.  $UO_{2.24}$ , *ibid.*

4.  $UO_{2.08}$ , *ibid.*

5.  $(U_{0.95}Pu_{0.05})O_{2.11}$ , *ibid.*

6.  $(U_{0.85}Pu_{0.15})O_{2.13}$ , *ibid.*

(From F. J. Homan, Parametric Analysis of Fuel-Cladding Mechanical Interactions, USAEC Report ORNL-TM-3508, p. 13, Oak Ridge National Laboratory, August 1971.)

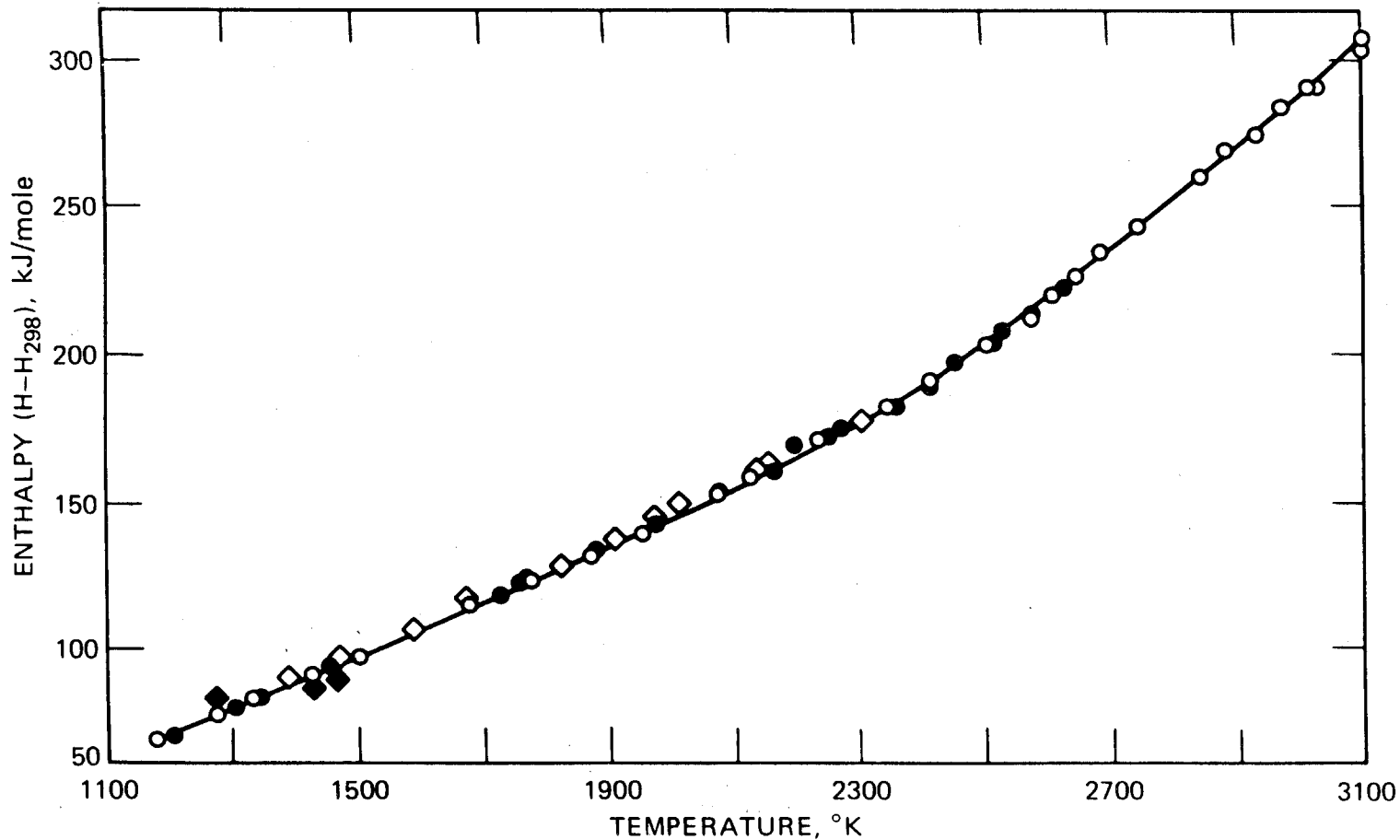


Fig. 10.10 Enthalpy—temperature data for stoichiometric  $\text{UO}_2$ . [From R. A. Hein, L. H. Sjodahl, and R. Szwarc, *J. Mater.*, 25: 99 (1968).]