

#### Outline

- Brief History
- Cermet sample testing during the NERVA/Rover era
  - Matrix/chart of samples tested & results
  - Comparison of approaches
- Important properties in context
  - Melting temperature
  - Vaporization rate
  - Chemical stability
- Engine performance
  - Location peak temperature
  - Heat deposition rate





# History of Cermet Sample Testing

- 1949- NEPA investigated Mo-UO<sub>2</sub> and W-UO<sub>2</sub>
- 1950's- Some further work
- 1961- Kennedy: "accelerate development of the Rover nuclear rocket"
- 1961- GE high-temperature materials program (HTMP & GE 710)
- 1962- Nuclear Propulsion Conference
  - LANL, LeRC, (GE) reported extensive testing results
  - UO<sub>2</sub> vaporization significantly reduced by thin tungsten cladding
  - UO<sub>2-x</sub> reduction issue, uranium hydride formation, and sample cracking
- 1960's- DOE's ANL, Pacific Northwest labs
- 1968- ANL 200/2000 engine design, 2500°C, <1% fuel loss, 10h, 25X
- 1968- Tighter budgets, terminal cermet fuel reports
- 1970~ Space race won: cancelled Apollo 18-20, manned Mars plans
- 1972- Rover/NERVA program cancelled
- Other cermet summaries: Haertling & Hanrahan, Lundberg & Hobbins



#### Performance of Historical Cermet Samples

A Broad Brush Painting of W-UO<sub>2</sub> Results

			Cracl	Unstable: Cracks or Forms Powder  Stable: Mass Loss > 5 %								: Mass < 5 %			
	Sample Group	1	2	3	4	5	6	7	8	9	10	11	12	13	14
	UO <sub>2</sub> Only	<b>V</b>						~							
	W-UO <sub>2</sub>		<b>V</b>	<b>/</b>	<b>V</b>	~	~		V	~	<b>V</b>	V	~	~	V
Partial Cla	ad (Not Edges)								V	~	~	~	~	~	
Coated	Full Clad Coated Fuel Particles			~									~		
	Stabilizers (Various)						~	~	V	V		~	V	~	~
	Temperature														
(C)	(K)														
2000	2273														
2300	2573														
2350	2623														
2500	2773														
2600	2873														
2650	2923														
2700	2973														
2800	3073														
Cycle	Cycles Tested								25	>25				<30	<10
Fuel Sam	ples Tested		29+14	46	19	2		25+	~30	~20	6	2	1	2	2
Refe	erence	[17]	[11]	[11]	[11]	[18]	[18]	[17]	[13]	[13]	[9]	[18]	[18]	[18]	[18]

- Considerable amount of cermet materials research in the early 1960's
- Over 200 W-UO<sub>2</sub> samples from five different labs: ANL, GE, LANL, LeRC, PNWL.
- Successes: full cladding, chemical stabilizers, coated particles
- Process improvement was done





# Melting Points & Vaporization Rates of NTP Reactor Fuels/Materials

Material/ Element	Melting Temperature (K)	Surface Vaporization Rate at 2800°K (mil/hr)					
Tungsten	3680	< 0.01					
Rhenium, Re	3453	0.1					
Graphite	3915 (sublimes)	10					
ZrC	3805	>>10					
Tantalum Carbide, TaC	4150	0.1					
Uranium Dioxide, UO <sub>2</sub>	3075	6×10 <sup>3</sup>					
Uranium Nitrides	Chemically Unstable						
Uranium Carbide, UC <sub>2</sub>	2835	10					
UC-40 ZrC  NERVA Composite	3050	2					

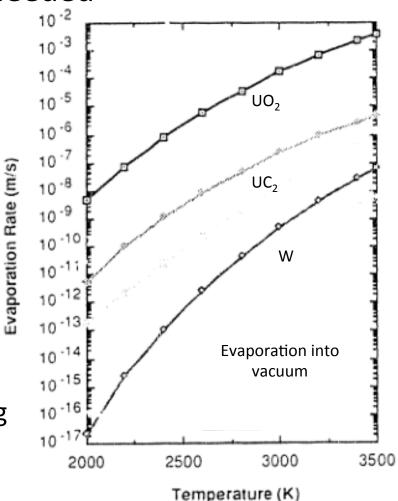




## Fuel Vaporization and Reactions

Coating/Cladding Needed

- Fuel vaporization is very high above 2000 K
- Cladding/particle coating needed
- At 1962 Nuclear Propulsion Conference
  - LANL (Lenz & Mundinger [9]): thin tungsten coatings reduce vaporization
  - LeRC (Saunders et al[13], McDonald[12]: fuel vaporization reduced 10X by cladding
- Face cladding is insufficient
  - Gluyas et al [13] demonstrated the need for full cladding



Evaporation rates of nuclear fuels and materials normalized to surface regression rates.

Lundberg, Hobbins EGG-M—92067



#### Fuel Vaporization Above 2000K

- Sample at 1900K for 30 minutes without significant mass loss
- At 2500K, sample experienced fuel evaporation

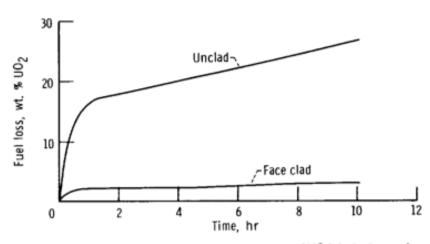
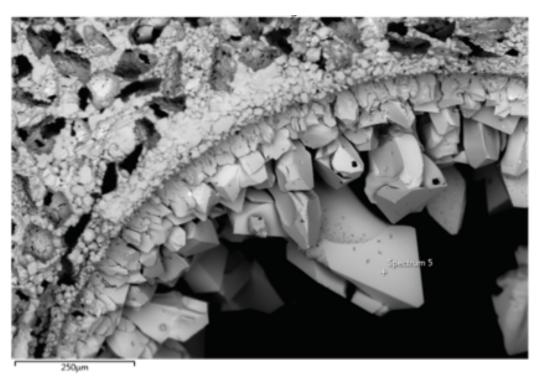


Figure 30. - Fuel loss as function of time at 2500° C in hydrogen for unclad and partially clad W - 20-volume-percent-UO<sub>2</sub> composites.





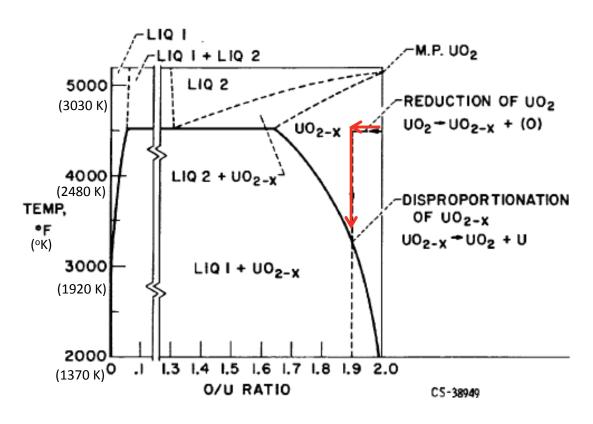
### Performance of Historical Cermet Samples

W-UO<sub>2</sub> Results

			Cracks or					e: Mass > 5 %				e: Mass < 5 %			
	Sample Group	1	2	3	4	5	6	7	8	9	10	11	12	13	14
	UO <sub>2</sub> Only	<b>V</b>						~							
D 11 1 01	W-UO <sub>2</sub>		~	<b>V</b>	<b>V</b>	V	V		<i>V</i>	V	<b>V</b>	V	~	V	<b>V</b>
Partial Cla	d (Not Edges)								~	~	~	~	~	V	~
Full Clad Coated Fuel Particles				~									~		
Stabilizers (Various)							~	~	~	~		~	~	~	V
Temp	erature														
(C)	(K)														
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Beals et al Baker et al							Bec	0/s et =	Sluyas,	Gen Gen	Mun	[18]			



# High Temperature Behavior of UO<sub>2</sub>



UO<sub>2</sub> structure Idealized

At temperatures above 2000°K, UO<sub>2</sub> becomes deficient in oxygen.

With cooling, free uranium forms.

Stabilizers (Gd<sub>2</sub>O<sub>3</sub>, Y<sub>2</sub>O<sub>3</sub>) interfere with this reduction.





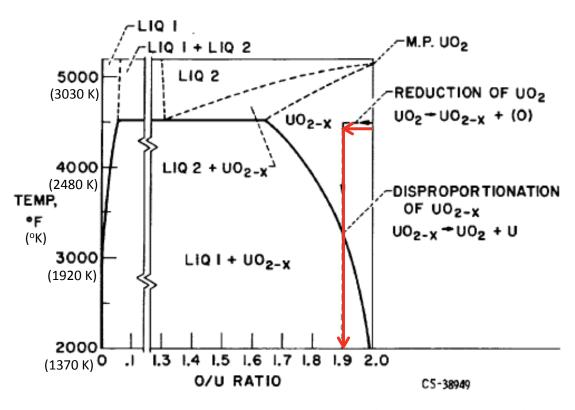
### Stability of UO<sub>2</sub>, and Chemical Stabilizers

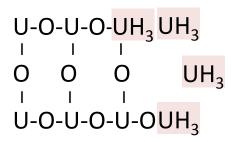
- 1960 Anderson: UO<sub>2</sub> reduces to UO<sub>2-x</sub>
- 1962 LeRC, LANL, GE: UO<sub>2-x</sub>, free uranium & UH<sub>3</sub> forms, sample cracking
- 1965 Beals, et al: Hydride, UH<sub>3</sub>, formation is accompanied by a disruptive volume change that destroys the integrity of the specimen
  - UO<sub>2</sub> was "heated to 2300 C in flowing dry hydrogen for 10 min. When cooled in hydrogen to below 500 C, the pellet disintegrated with sufficient force to shatter the (foil) crucible. The residue was a very fine black powder."
  - "crumbling or powdering of the specimen"
  - UH<sub>3</sub> forms between 370 K and 620 K; can cool in non-hydrogen environment
- Addition of rare-earth oxides improves stability, particularly gadolinium Gd<sub>2</sub>O<sub>3</sub>, also Y<sub>2</sub>O<sub>3</sub>
  - UO<sub>2</sub> at 2570 K crumbles
  - UO<sub>2</sub> 10 mol% Gd<sub>2</sub>O<sub>3</sub> heated to 2770K had 6-12 % weight loss
  - At 2920 K, needed 5 mol% GdO<sub>1.5</sub> + 5 mol% FeO<sub>1.5</sub> for stability





# High Temperature Behavior of UO<sub>2</sub>





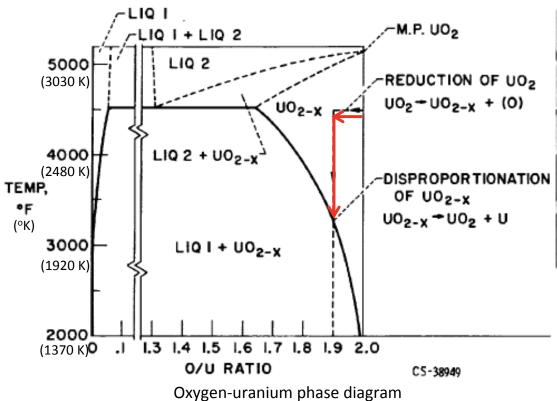
With cooling below 2000°K, free uranium forms.

Below 770°K, free uranium combines with hydrogen to form uranium hydride, UH<sub>3.</sub> This is hydrogen embrittlement





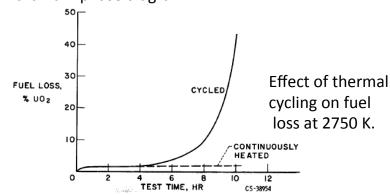
### Chemical Stability of UO<sub>2</sub> with Thermal Cycling



UO<sub>2</sub> Dispersoid

Uranium-Rich
Phase
Tungsten
Matrix

Micrograph of thermally cycled W-20 vol%  $UO_2$  cermet showing free U at grain boundaries. The specimen was heat treated for five 1-h intervals at 2770 K in  $H_2$  with cooling to room temperature between cycles.



NASA X66 51413 NASA TM-X-1421





## Performance of Historical Cermet Samples

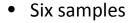
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Partial Cla	W-UO <sub>2</sub> ad (Not Edges)		~	<b>V</b>	<b>V</b>	~	<b>V</b>		V	<b>V</b>	~	V	~	V	<b>V</b>
Partial Cla	Full Clad									~	~		~	V	~
Coated	Coated Fuel Particles			~									~		
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		Beal	ls et al					[17] B <sub>C<sub>Q</sub></sub>	set al	Pluyas	$G_{e_{\alpha}}$			ANI	



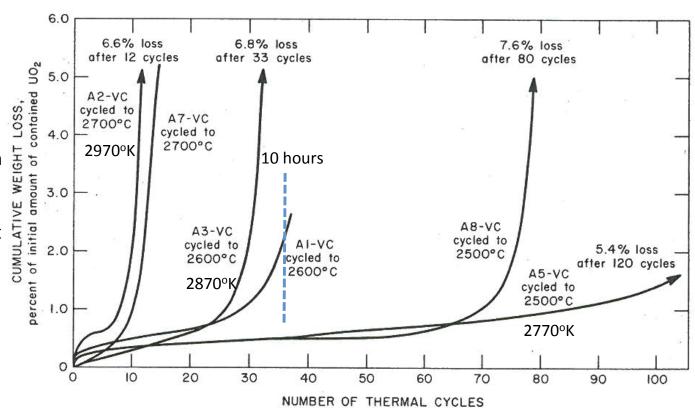


#### Pattern of Performance in Cermet Fuel Samples

Thermally cycled in furnace—not rocket/reactor conditions



- Low-pressure hydrogen,
- ANL-7150 suggests the hydrogen is static, or nearly so.
- Testing with flowing hydrogen at engine pressures would reduce performance.
- Comparable results found at 2500°C in reference 13, Y<sub>2</sub>O<sub>3</sub>, 20-35v% UO<sub>2</sub>, flowing hydrogen
- Each 100°C increase in temperature significantly decreases lifetime



Fuel loss behaviors of tungsten-clad W-66 v/o (10m/o GdO1.5-stabilized  $UO_2$ ) cermet samples (not fuel elements) thermally cycled to 2770 K, 2870 K, and to 2970 K.



ANL-7150





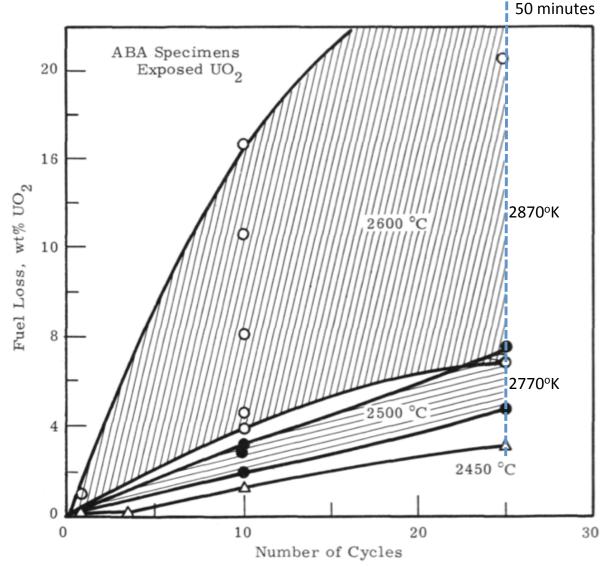
#### Pattern of Performance in Cermet Fuel Samples

Thermally cycled in furnace—not reactor conditions

Fuel particles coated with W, no cladding

• High-pressure, static hydrogen,

Thermal Cycling Behavior of Tungsten-Coated  $UO_2$ -W (13.3 vol%  $UO_2$ ) Cermets Under Accelerated Test Conditions in 68 atm Static Hydrogen.



NASA CR-54840, p. 48





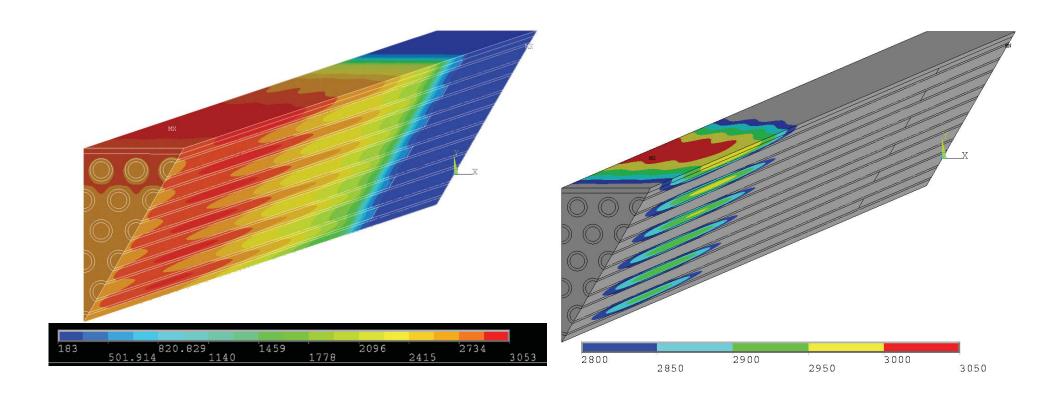
# What Does a Fuel Element Designer Do with Material Performance Data?

- Engine/Fuel designer must:
  - Highest possible propellant outflow temperature
  - Minimum peak fuel temperature
  - Nuclear criticality & control
  - Engine system performance (turbopump, nozzle)
  - Acceptable fuel loss, maintain fuel integrity
- High fidelity simulations help understanding
  - Neutronics simulations predict criticality
  - High-Fidelity fluid / thermal / structural simulations
  - Can simulate materials and performance





#### Where is the Hottest Cermet Fuel?

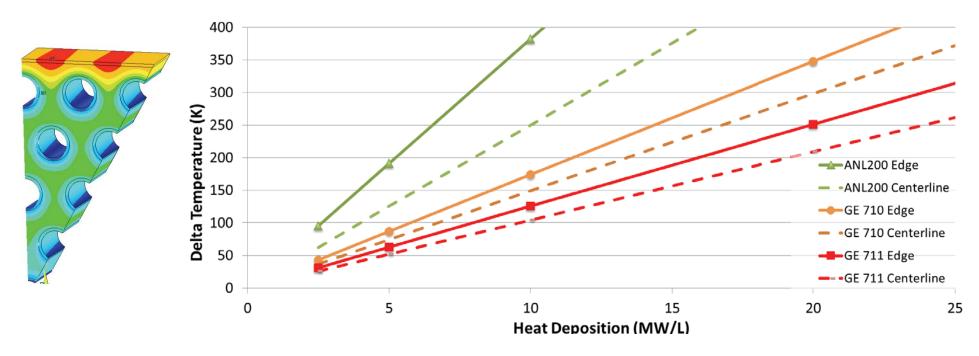


Predicted temperature distribution through a GE 711 cermet fuel element (left) and detail of the hottest 250K region of the fuel element (right).





# Fuel Temperature Differences Versus Heat Deposition Rate



Predicted temperature difference, fuel peak at edge to coolant channel (solid) and fuel centerline to coolant channel (dashed) for several cermet fuel geometries.



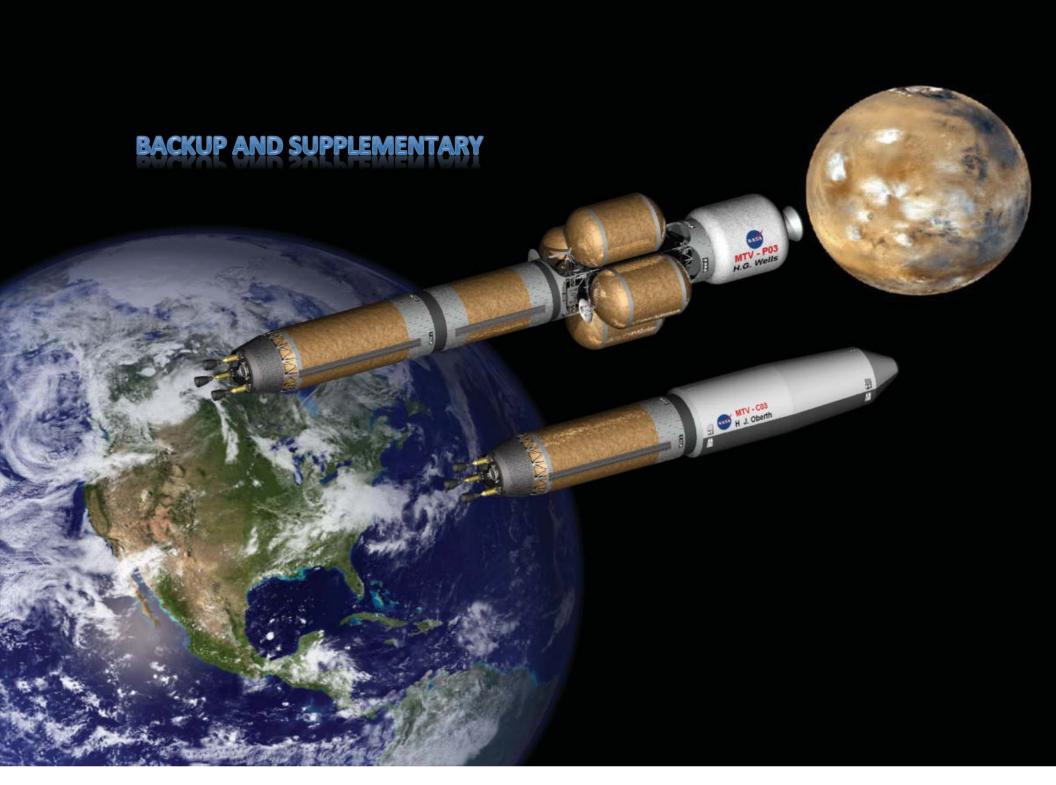


## Summary and Conclusions

- To better understand Cermet engine performance, examined historical material development reports
- Two issues:
  - High vaporization rate of UO<sub>2</sub>
  - High temperature chemical stability of UO<sub>2</sub>
- Cladding and chemical stabilizers each result in large, order of magnitude improvements in high temperature performance
- Some long duration, low mass-loss, samples at 2770°K
- Few samples were tested above 2770°K
- Contemporary testing may clarify performance

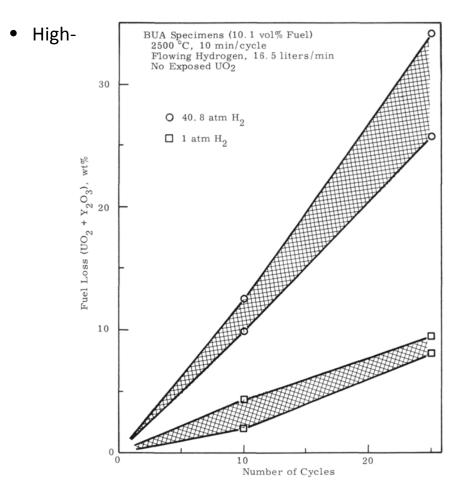




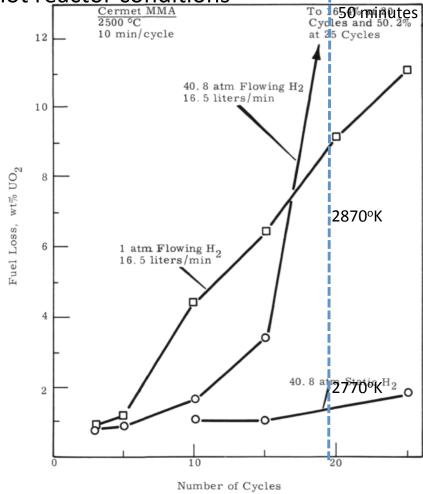


#### Pattern of Performance in Cermet Fuel Samples

Thermally cycled in furnace—not reactor conditions



Thermal Cycling Behavior of W-UO $_2$  Coated Particle Cermets Containing 10 Mole%  $\rm Y_2O_3$  in UO $_2$  Solid Solution.



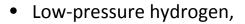
Effect of Pressure and Flow Rate on the Thermal Cycling Behavior of a Tungsten-Coated UO<sub>2</sub>-W Cermet Containing 13.3

vol% UO NASA CR-34840, p. 48

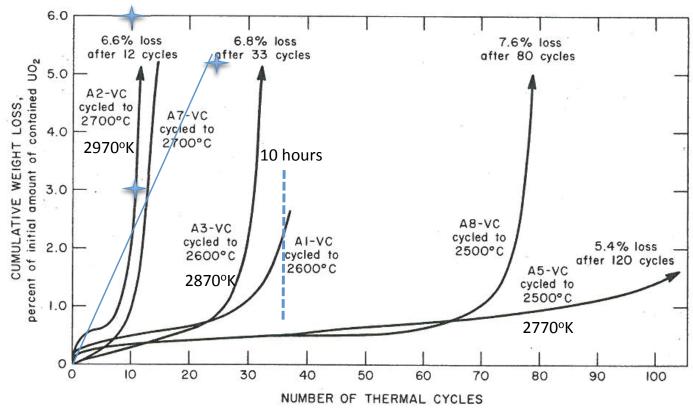


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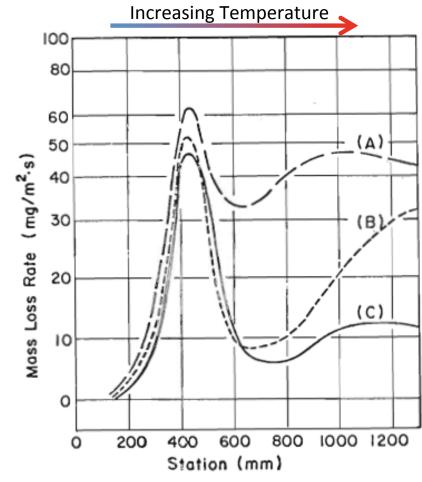
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#### Mid-Band Erosion in NERVA Composite Fuel Elements

- High CTE composite fuel elements were crack-free as fabricated
- Hot end fuel loss agreed with predictions
- Midrange losses were unexpected, 2/3 of total
- Cold end coating cracks caused midrange loss, H<sub>2</sub> + C -> CH<sub>3</sub> & C<sub>2</sub>H<sub>2</sub>



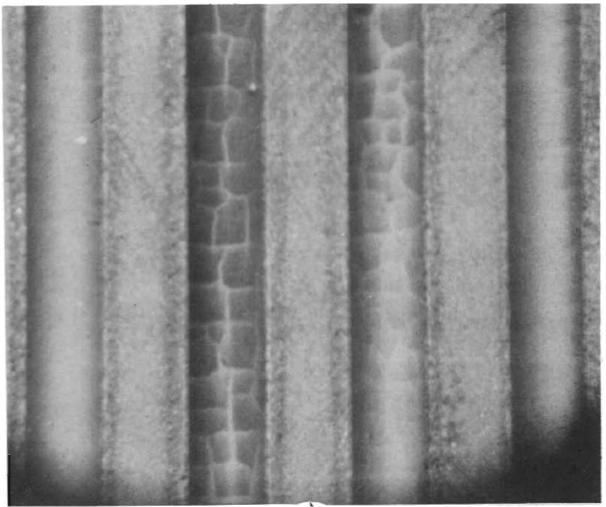
Mass loss rates per unit surface area of coolant channel versus station for graphite and composite elements. (A) average for 102 Pewee-1 graphite fuel elements coated with NbC, (B) average for 12 Pewee-1 graphite fuel elements coated with ZrC, (C) average for 23 NF-1 high-CTE composite fuel elements coated with ZrC, adjusted to the Pewee-1 test temperature.

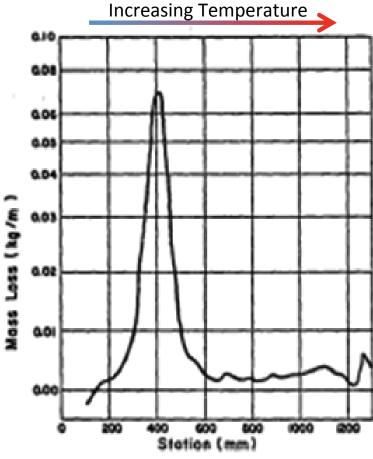
LA-5398-MS





#### Pattern of Cracking / Erosion in NERVA Fuel





Mass loss versus length for (U,Zr)C-graphite fuel element in NF-1 test. CTE > 6.5  $\mu$  m/mK.

NERVA fuel element interior coolant channels experienced coating cracks in the NF-1 test, while edge channels retained their coatings. Mid-passage erosion region.

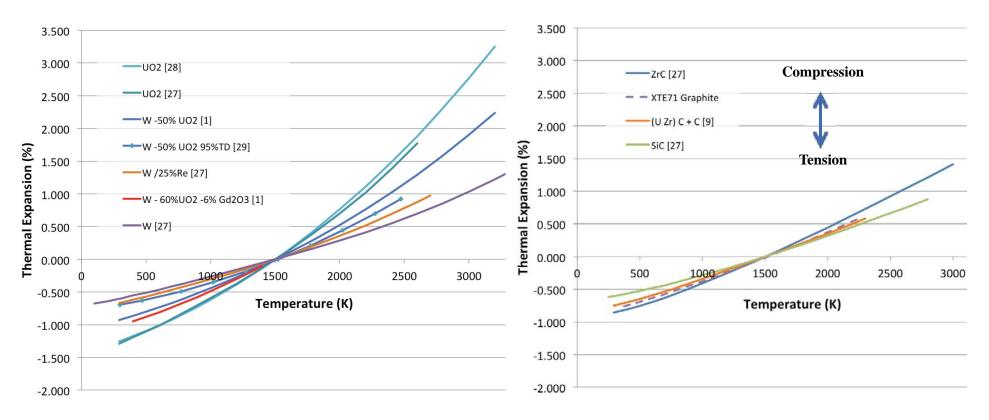
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#### Stress Sources: Differential Thermal Expansion

Cermet and Graphite-Based Fuels



**Cermet Fuels** 

Coatings in *compression* on cool down

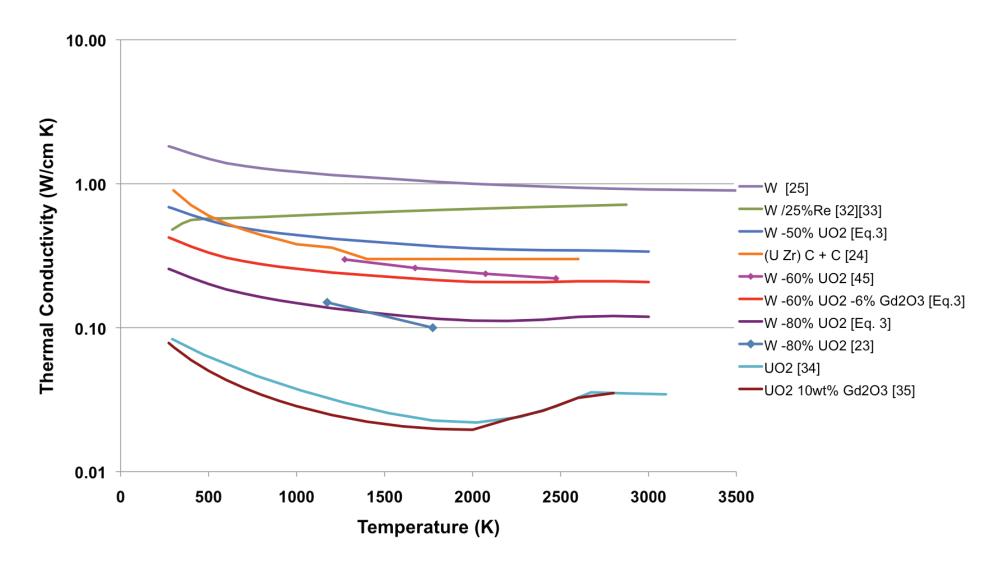
**Graphite-Based Fuels** 

Coatings in *tension* on cool down





#### NTP Fuel Elements: Thermal Conductivity







#### NTP Fuel Elements: Modulus of Elasticity

