

#### Risk Assessment of Uniform Corrosion and Localized Corrosion of Alloy 22

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> Materials Science and Technology (MS&T) '03 Chicago, IL, November 9-12, 2003

> > 1



## **OBJECTIVES**

Conduct Sensitivity Studies of Localized Corrosion and Nonpassive Uniform Corrosion of Alloy 22 with Nuclear Regulatory Commission (NRC)'s Total-system Performance Assessment (TPA) Code

- severe chloride solution
- high temperature deliquescence point of salts
- fabrication-induced microstructural alteration
- inhibitor effects
- controlled release through pits
- anodic sulphur segregation and long-term development of surface roughness



#### REPOSITORY AND ENGINEERED SYSTEM NRC'S TOTAL-SYSTEM PERFORMANCE ASSESSMENT (TPA) CODE



Schematic Illustration of the Emplacement Drift with Cutaway Views of Different Waste Packages (DOE, 2002)







### LOCALIZED CORROSION

- Sensitivity study of the effect of critical relative humidity Some salts deliquesce at a low RH.
- If RH<sub>environment</sub> > RH<sub>lowerCriticalAqueousCorr</sub> localized corrosion occurs.

	Range of Critical RH for Aqueous Corrosion	
<b>Base</b> (Drip Shield Included)	0.60 – 0.65	
<b>Modified</b> (High Temperature Deliquescence of Salts	0.35 – 0.60	



Temp effect on the repassivation potential crevice corrosion of Alloy 22 in Clsolutions data was integrated into modified corrosion potential equation parameter set in TPA.

Data from [NO<sup>3</sup>]/[Cl<sup>-</sup>] experiment was used to empirically determine new values for the repassivation potential equation (the inhibitor effect)



## LOCALIZED CORROSION (Modified Criteria)

- $E_{repass} = E_o crit (T) + B(T) log[Cl<sup>-</sup>]$ 
  - $E_{o} crit (T) = A_{1} + A_{2}T$ 
    - OuteroverpackErpIntercept, A<sub>1</sub> (mV<sub>SHE</sub>)
    - TempCoefOfOuterPackErpIntercept, A<sub>2</sub>, (mV/°C)
  - $B(T) = B_1 + B_2 T$ 
    - OuterOverpackErpSlope, B<sub>1</sub> (mV)
    - TempCoefOfOuterPackErpSlope, B<sub>2</sub> (mV/°C)
      Critical Corrosion Equation Values
- Nearly constant with sufficient concentration of inhibitors are present



## LOCALIZED CORROSION

	Repassivation Potential Parameters			
	A <sub>1</sub> (mV <sub>SHE</sub> )	A <sub>2</sub> (mV/°C)	B <sub>1</sub> (mV)	B <sub>2</sub> (mV/°C)
BASE	2006.0	-15.2	-590.7	4.3
MODIFIED (more data added and fitting refined)	1541.0	-13.1	-362.7	2.3



#### **Base Case**



Time, yrs

Dose from corrosion/juvenile failure – manufacturing defects, human error, etc.



Dose, mrem/yr

#### Base Case, Lowered Critical RH (High Temperature Deliquescence of Salts), No Nitrates



Time, yrs

Dose from corrosion/juvenile failure – manufacturing defects, human error, etc.



## Modified Equation, Base Case, No Nitrates



Time, yrs

Dose from corrosion/juvenile failure – manufacturing defects, human error, etc.





Time, yrs

Dose from corrosion/juvenile failure – manufacturing defects, human error, etc.



#### LOCALIZED CORROSION CONTROLLED RELEASE THROUGH PITS

• Overall Factor = 10<sup>-9</sup>, 1" log uniform distribution

(Pits)

- Size: (10<sup>-4</sup> 10<sup>-1</sup>) cm<sup>2</sup>
- Density: (0.1 100)/cm<sup>2</sup>
- Fraction:
  - $(10^{-4} \text{ cm})^2 \text{ x } 0.1/\text{cm}^2 = 10^{-9}$  $(10^{-1} \text{ cm})^2 \text{ x } 100/\text{cm}^2 = 1$

(Stress Corrosion Cracks)

- Size: (25 x 1.02) cm<sup>2</sup> = 25.5cm<sup>2</sup>
- Fraction: 25.5cm<sup>2</sup>/WP / WP surface area 2.3 x 10<sup>5</sup> cm<sup>2</sup> = 1.1x10<sup>-4</sup>

From Ahn (1994), Esh (2002)



#### LOCALIZED CORROSION CONTROLLED RELEASE THROUGH PITS



# JUCLEAR REGULATOR

#### DEVELOPMENT OF LONG-TERM STATISTICAL SURFACE ROUGHNESS AND SULFUR SEGREGATION

- Cyclical process of degradation
- Periods of slow (passive) corrosion due to passive layer formation are followed by anodic sulfur segregation and sloughing of the passive layer
- Then fast (non-passive) corrosion occurs in those areas before they repassivate.



#### DEVELOPMENT OF LONG-TERM STATISTICAL SURFACE ROUGHNESS AND SULFUR SEGREGATION

- Penetration Depth =  $\Sigma[CR_f * Ct_f + CR_p * Ct_p]$
- CR<sub>f</sub> = fast corrosion rate Varied ~ 10<sup>-4</sup> – 10<sup>-2</sup> cm/yr
- Ct<sub>f</sub> = fast corrosion time <sup>(1)</sup>
  Varied ~ 0.000119 1.19 yr
- CR<sub>p</sub> = passive corrosion rate <sup>(1)</sup> 10<sup>-4</sup> cm/yr
- Ct<sub>p</sub> = passive corrosion time <sup>(2)</sup>
  1.8 yr
- Process repeated until 2cm of container thickness corroded.

<sup>(1)</sup> From *Repassivation Kinetics*, Brossia et al., 2001
 <sup>(2)</sup> From *Sulfur Segregation Time*, Jones, 2002



#### DEVELOPMENT OF LONG-TERM STATISTICAL SURFACE ROUGHNESS AND SULFUR SEGREGATION

Fast Corrosion Time vs Lifetime for 3 Different Fast Corrosion Rates





#### CONCLUSIONS

- Lowering the critical RH for the Base Case (no nitrates) and Modified case for the repassivation potentials resulted in an increase in dose.
- The modified repassivation equation with nitrates incorporated resulted in a lower dose.
- WP failure by sulfur segregation is not expected based on the unrealistic combination of fast corrosion rates and recurrence frequencies necessary to breach the outer container.



## References

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