# KANUPP – IAEA Training

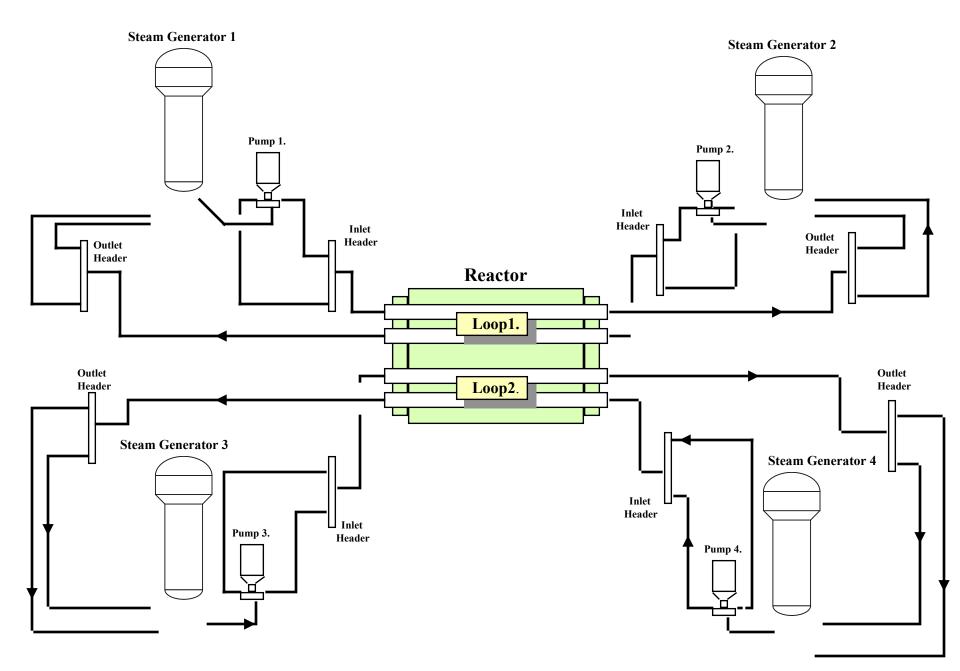
# Primary Heat Transport System Chemistry Control

#### System Purpose

- Transport heat from the fission of natural uranium fuel in the reactor fuel channels to steam generators, where the heat is transferred to light water to produce steam.
- Provide adequate cooling to the reactor fuel at all times including operation, shutdown, maintenance and loss of Class IV power.
- Provide overpressure protection for the system and system components.
- Minimize heavy water leakage.

#### **Major Components**

- 380 fuel channel assemblies
- four steam generators
- four electrically driven heat transport pumps
- one pressurizer
- four inlet and four outlet headers



#### **System Parameters**

System Volume 122400 Mg

#### • Temperatures

- Reactor Inlet Header 266°C
- Reactor Outlet Header 310°C

#### • Pressures

Reactor Inlet Header 11.33 Mpa
Reactor Outlet Header 9.99 MPa

#### • Flow Rates

•	<b>Total Core Flow</b>	7.7 Mg/s
•	Flow/channel (max)	23.9 Kg/s
•	Purification Flow (max)	24 Kg/s

 $(T_{1/2} = 60 \text{ min.})$ 

#### **Auxiliary Systems**

#### • Pressure and Inventory Control System

• maintains heavy water inventory and pressure in the main heat transport system

#### • Purification System

• removes ionic impurities (corrosion & fission products) as well as crud from the main heat transport system.

#### Shutdown Cooling System

• circulates and removes heat from the reactor coolant when the reactor is shut down, and when the reactor coolant is drained to the header elevation for equipment maintenance.

#### **Auxiliary Systems**

#### • Emergency Core Cooling

• provides light water flow to the primary heat transport system in the event of a Loss of Coolant Accident.

#### • D<sub>2</sub>O Storage, Transfer & Recovery System

- accommodate heat transport system swell when the reactor is warmed from cold to hot, stores D<sub>2</sub>O removed from the heat transport system when the coolant level is reduced for equipment maintenance, and recovers D<sub>2</sub>O spillage from the vault floor.
- D.N. & G.F.P. Monitoring Systems
  - the gaseous fission product (G.F.P.) monitoring system detects, and the delayed neutron (D.N.) monitoring system locates defective fuel in the reactor

#### **Chemistry Objectives**

- Minimize the corrosion of system components.
- Control the quantity of radioactive crud in the system by controlling the rates of crud production, movement and activation and by crud removal in the purification system.
- Control the concentrations of dissolved fission products by removing them by ion exchange or by degassing.
- Minimize the fouling of the Heat Transport surfaces by the crud principally by pH control and also through crud removal in the purification system.

#### **Materials of Construction**

#### **Metal/Alloy**

Carbon Steel

#### Stainless Steel

• Zirconium Alloys

#### System Component

- Piping
- Feeders
- Headers
- Bleed Condenser
- Bleed Cooler
- D<sub>2</sub>O Storage Tank
- IX Columns
- End Fittings (403/410)
- Instrument Tubing (304L)
- IX Column Inlet Nozzles
- Pressure Tubes
   (Zr-2.5% Nb)
- Fuel Cladding (Zr-4)

#### SUMMARY OF CHEMISTRY CONTROL PARAMETERS

Parameter	Permissible Range	Notes
pH (25°C)	10.2 to 10.8	Controls corrosion and activity transport
Li mg Li/Kg D <sub>2</sub> O	0.35 to 1.4	
Conductivity mS/m - 25°C	0.9 to 3.6	Conductivity, pH & Li must be considered together for effective control.
D <sub>2</sub> ml(STP)/Kg D <sub>2</sub> O	3 to 10	Oxidizing conditions if too low; possible hydriding of Zr alloys if too low or too high

#### SUMMARY OF CHEMISTRY CONTROL PARAMETERS

Parameter	Permissible Range	Notes
O <sub>2</sub> ug/Kg D <sub>2</sub> O	<10	Controlled by level of dissolved D <sub>2</sub>
Cl <sup>-</sup> mg/Kg D <sub>2</sub> O	<0.2	Stress corrosion cracking of stainless steels. Possible crevice and pitting corrosion.
Crud mg/Kg D <sub>2</sub> O	<0.1	High crud levels increase activity transport
I-131 MBq/L	500	OP & P Limit.

# KANUPP Primary Heat Transport System Chemistry Specification

•	pН	=	> 10.3
•	Conductivity	=	< 25 umho/cm
•	Chloride	=	< 0.1 ppm
•	Carbonate	=	ND
•	Copper	=	ND
•	Iron (dissolved)	=	ND
•	Lithium	=	1.2 ppm
•	Crud	=	< 25 ppb
•	Oxygen	=	< 70 ppb
•	Hydrogen	=	< 3 ml/Kg
•	Isotopic	=	< 99.0 % D20

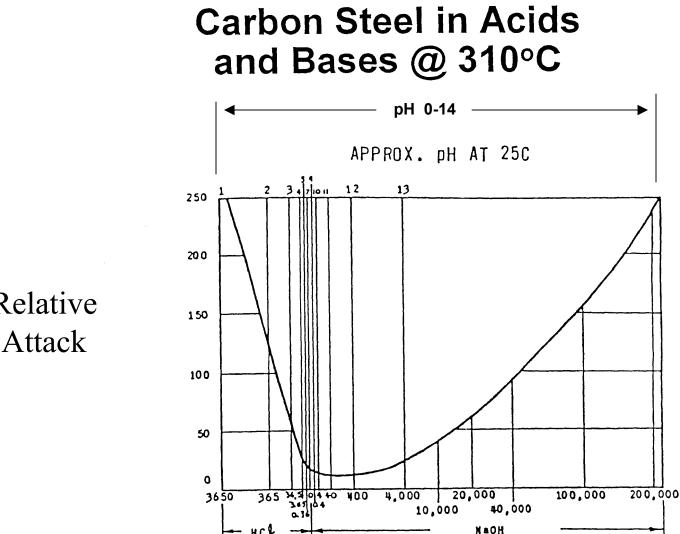
# pН

• Current Specification

#### 10.2 to 10.8

apparent pH  $pH_A = pH + 0.457$ 

- Minimize corrosion of carbon steel and zirconium alloys.
  - High pH optimum for preservation of the carbon steel magnetite layer.
  - Too high a pH can result in increased corrosion in localized areas where lithium hydroxide can concentrate (between fuel bearing pads & pressure tubes would be one location). The upper specification of 10.8 has been set as the point where it has been judged that corrosion even in localized regions will be acceptably low.
- Maintaining low activity transport throughout the system.
  - pH values above 10.2 will ensure low deposition of crud (magnetite) in the core at core temperatures, magnetite solubility increases with increasing pH.



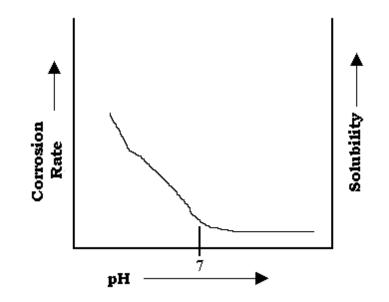
mg/kg

mg/kg

**Corrosion Rate of** 

Relative Attack

# **pH Versus Metal Corrosion**



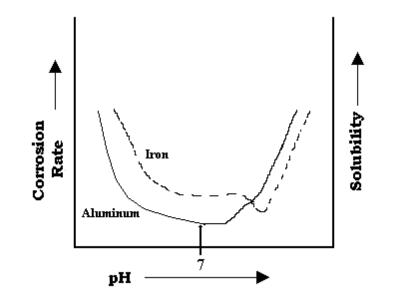
## **High Solubility in Acid**

### Examples:

Copper Nickel Chromium Cobalt

# pH Versus Metal Corrosion

### **AMPHOTERIC METALS**



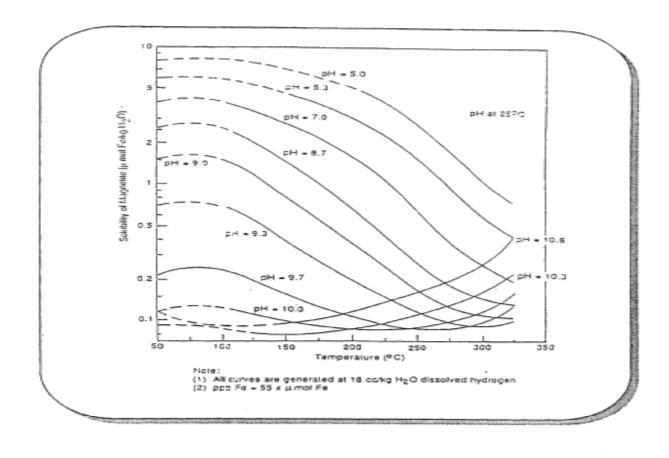
- High Solubility in Acid and Base.
- Characteristic pH where solubility is minium.

#### **EXAMPLES**:

Aluminum Iron Titanium Zirconium

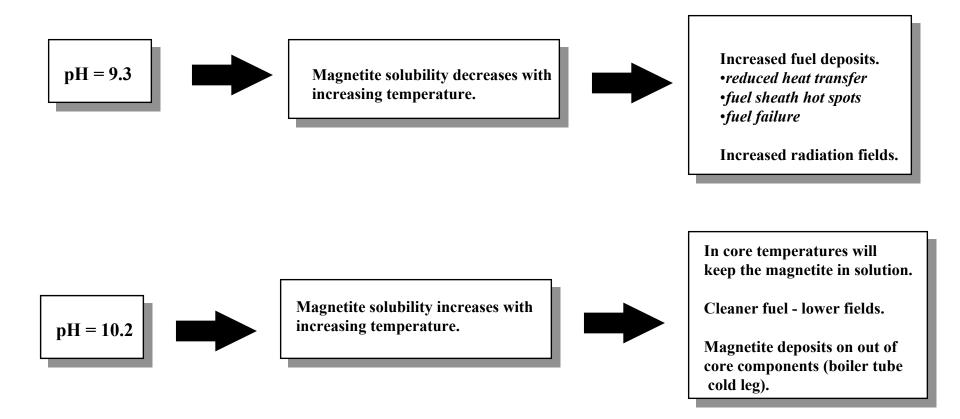
**Overhead 16** 

## **Magnetite Solubility**



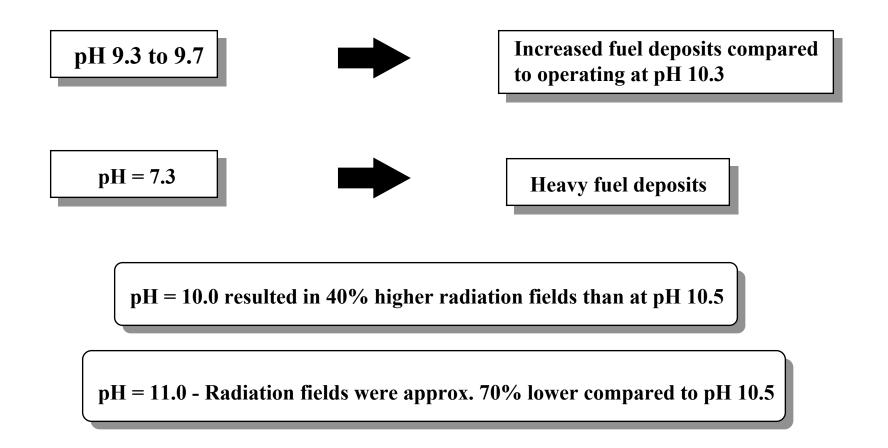
Overhead 76 AEC8 Conventional/Nuclear General

### **Effect of pH on Magnetite Solubility**



### **Effect of pH on Fuel Deposits & Radiation Fields**

**Results from AECL Loop Tests** 



# **Dissolved Oxygen**

• Current Specification

<10 ug/Kg

- Sources
  - Radiolysis of the D<sub>2</sub>O
  - $D_2O$  Makeup
- Concerns
  - Localized & General corrosion of zirconium pressure tubes and steam generator tubes.
  - Corrosion of the carbon steel piping loss of magnetite (conversion to hematite) higher crud levels increased radiation fields.

### **Oxygen Production by Radiolysis**

$$D_2O \longrightarrow D_2, D_2O_2, e_{aq}, D, OD, DO_2$$

Molecules Radicals

 $OD + OD \longrightarrow D_2O_2$ 

 $D_2O_2 \longrightarrow D_2O + \frac{1}{2}O_2$ 

### **Oxygen Suppression by Hydrogen Addition**

$$OD + D_2 \longrightarrow D_2O + D \qquad (1)$$
$$D + D_2O_2 \longrightarrow D_2O + OD \qquad (2)$$

The rate of reaction 1 increases with increasing temperature. The reaction rate is (a) 200 times faster at reactor temperatures compared to room temperature.

## **Hydrogen Addition for Oxygen Suppression**

#### • Current Specification

10 to 25 ml/Kg

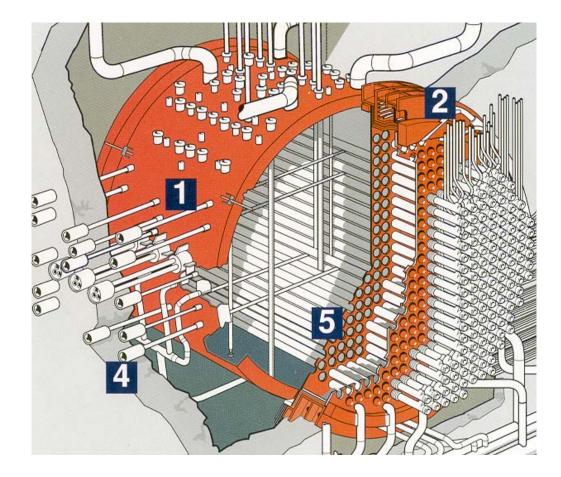
#### • Low Specification

- Low Specification based on early work performed by AECL in their NRU Reactor. Studies showed that 0.3 ml/Kg hydrogen (critical hydrogen concentration) required in non-boiling coolant to suppress oxygen formation (equal to or less than 2 ppb). In boiling coolant, some of the hydrogen/deuterium is stripped into the steam phase. Thus in boiling coolants, higher concentrations of hydrogen/deuterium is required to suppress oxygen formation ( 3% steam quality requires 1 ml/Kg H<sub>2</sub>. At 6% steam quality, about 1.5 ml/Kg is needed.)
- <u>The lower specification of 3 ml/Kg incorporates a small safety margin for dissolved</u> <u>deuterium concentrations</u>

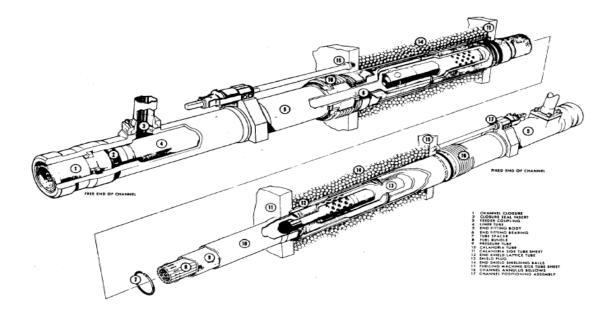
#### • High Specification

 Hydrogen levels in excess of 25-35 ml/Kg may cause cracking of any I-600 steam generator tubes (Primary Water Stress Corrosion Cracking - PWSCC). Maintaining the hydrogen levels below 10 ml/Kg ensures that the risk of tube cracking is minimized in I-600, but is not a restraint with Monel 400 tubes.

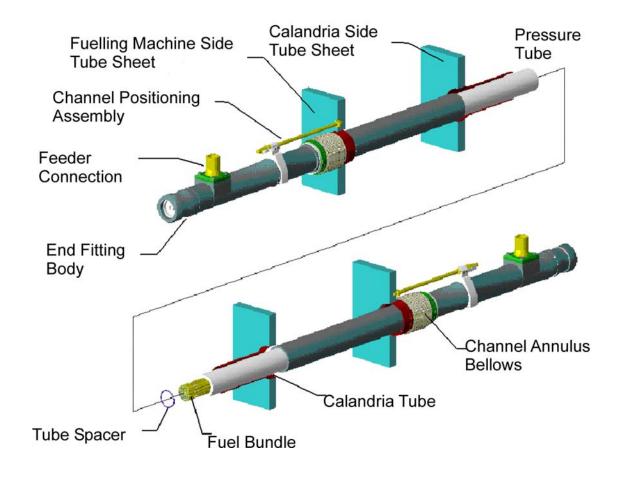
# **Fuel Channel Assembly**



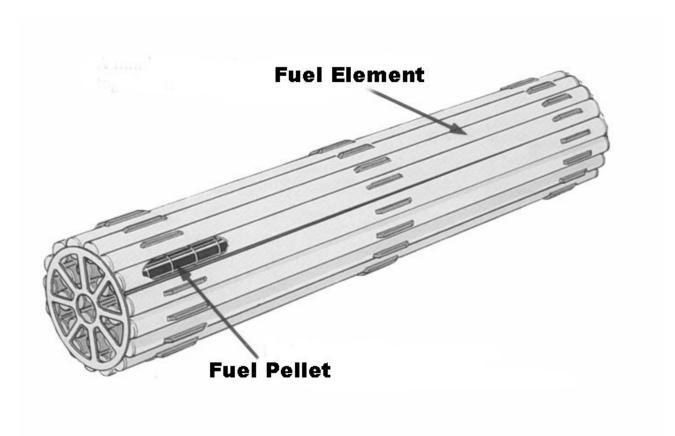
## **Calandria/Pressure Tube Arrangement**



### **Fuel Channel Features**



## **CANDU 6 Fuel Bundle**



### Typical Compositions of Zirconium Alloys (wt %)

<u>Element</u>	<u>Zr-2.5% Nb</u>	Zircoloy-2
Tin	-	1.1 to 1.7
Iron	-	0.07 to 0.20
Chromium	-	0.05 to 0.15
Nickel	-	0.03 to 0.08
Niobium	2.4 to 2.8	-
Oxygen	900 to 1300 ppm	1400 ppm max.
Zirconium	Balance	Balance

*Note : AECL Specification for Fr, Cr & Ni in Zr-2.5% Nb pressure tubes is:* 

*Fe - 1500 mg/g max. Ni - 70 mg/g max. Cr - 200 mg/g max.* 

**Overhead 28** 

### **Corrosion of Zirconium Alloys**

# $Zr + H_2O (or D_2O) \longrightarrow ZrO_2 + 2H_2 (2D_2)$

A percentage of the  $H_2(D_2)$  generated by the above reaction is picked up or absorbed by the zirconium metal. The amount of  $H_2(D_2)$  picked up will depend upon the composition of the zirconium alloy.

Alloy	%H <sub>2</sub> /D <sub>2</sub> Pickup	
Zr-2	50%	
Zr-2.5% Nb	<5%	

### **Corrosion of Zirconium Alloys**

### Zr-4

The rate of hydrogen/deuterium pickup by zirconium alloys is directly related to the composition of the alloy.

The presence of nickel in the alloy will affect the  $H_2/D_2$  pickup rate. The higher the nickel content, the greater will be the  $H_2/D_2$  pickup rate.

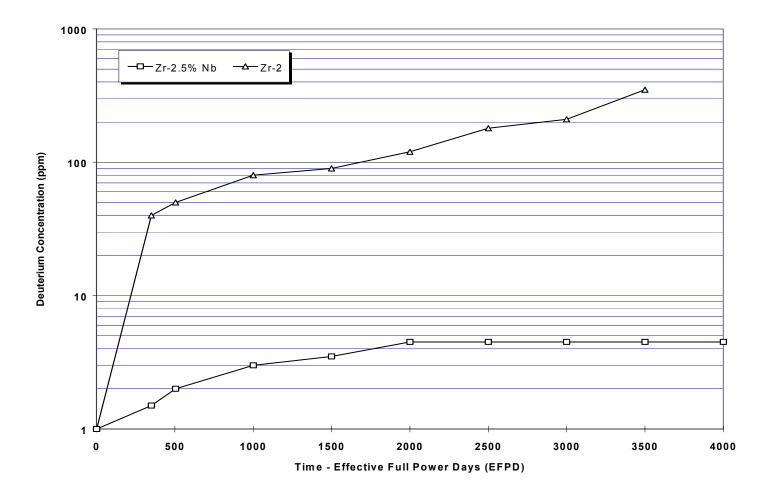
The reduction of nickel in zirconium alloys has resulted in alloys with much lower  $H_2/D_2$  pickup rates.

#### Zr-4 is basically a low nickel version of the Zr-2 alloy

Note: Zr-2 is used for calandria tubes; Zr-4 for fuel sheath

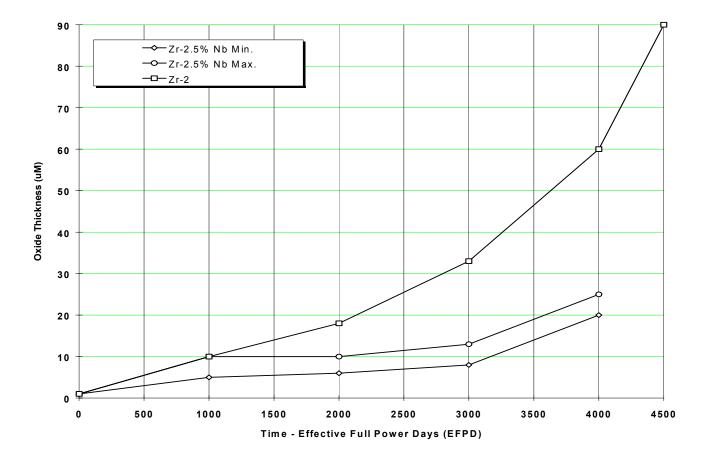
#### **Pressure Tube Deuterium Pickup From Corrosion**

Zr-2 & Zr-2.5% Nb Pressure Tubes

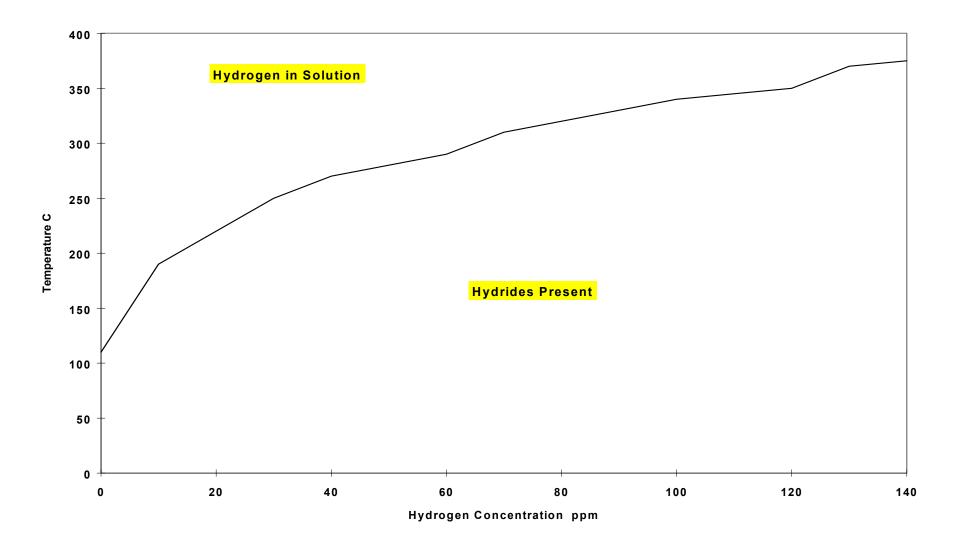


#### **Oxidation of Pressure Tubes**

Zr-2 & Zr-2.5% Nb Pressure Tubes



#### Effect of Temperature on the Solubility of Hydrogen in Zr-2.5 wt% Nb



### **Pressure Tube Operating Experience**

# Hydrogen Pickup

• Oxidation of the zirconium alloy surface in the presence of water results in the evolution of hydrogen (deuterium). The hydrogen released is very soluble in the Zr metal matrix where it tends to accumulate. Zr-2.5% Nd alloy demonstrates reduced rates of oxide growth and thus reduced pickup of hydrogen (deuterium) as compared to Zr-2.

### Delayed Hydrogen Cracking

Accumulation of hydrogen to the concentrations in the range of 20 to 50 ug per gram of Zr results in the precipitation of a zirconium-hydride (ZrH<sub>2</sub>). These hydrides can cause the metal to crack under large tensile stresses (thermal cycling).

### **Pressure Tube Operating Experience**

### • Blisters

• At very large concentrations of hydrogen ( about 350 ug/g Zr), blisters of ZrH<sub>2</sub> may form. When these blisters reach a critical size, cracks appear in the metal. One of the causes of the pressure tube failure (P2 G16) at Pickering Unit 2. Blister formation is accelerated when the pressure tube is allowed to come in contact with the calandria tube.

### • Creep & Growth

• Volumetric expansion of the pressure tubes will occur during warnup to operating range of 256 to 310°C, and from pressurizing the coolant inside the tubes to 10 MPa. In addition, prolonged exposure of the Zr-Nb material to neutron and gamma radiation results in a continuous additional expansion of the tube (less than 2.5% over 30 years). Recovery of the thermal and pressure effects will occur following reactor shutdown. Expansion due to radiation exposure will remain.

### **Pressure Tube Failures**

Factors leading to the failure of the Pickering Unit 2 G16 Pressure Tube

- Garder springs not in their design location.
- resultant contact between the pressure tube and calandria tube due to creep sag.
- high deuterium content in the pressure tube combined with the pressure tube to calandria tube contact led to the formation of hydride blisters, and the eventual cracking of the pressure tube material as a result of the stresses from blister growth.

### **Pressure Tube Failures**

#### **Conditions required for Delayed Hydride Cracking to occur**

- Hydrogen (deuterium) must be in the form of a hydride (i.e., if the hydrogen is all in solution, cracking will not occur).
- Sufficient stress must be present.
- There must be sufficient time for the above two conditions to occur. Initiation is enhanced by temperature cycling under stress.
- Time is needed for cracking to proceed once it has been initiated.

### **Preventing Pressure Tube Failures**

- Use Zr-2.5% Nb pressure tubes instead of Zr-2 (used in older reactors). This alloy displays a reduced tendency for oxidation and thus hydrogen/deuterium pickup (50% less than Zr-2).
- Increase the number of garder springs from 2 garder springs per channel to 4 garder springs per channel. Better support along the length of the pressure tube.
- Maintain reducing chemistry conditions. Maintain dissolved oxygen as low as possible by keeping a minimum of 3 ml/Kg of dissolved deuterium in the water.
- Maintain an oxide layer on the outside of the pressure tubes. Trace amounts of oxygen in the annulus gas system are sufficient to maintain this oxide layer.

## Crud

### **Concerns**

- Increased deposits in core.
  - activation of crud increased radiation fields
  - deposits on fuel decreased heat transfer fuel failures.
- Increased Deposits out-of-core.
  - increased radiation fields on out-of-core components
  - Deposition of crud on the inside of the Steam Generator tubes reduced heat transfer increased RIHT temperatures.

## Crud

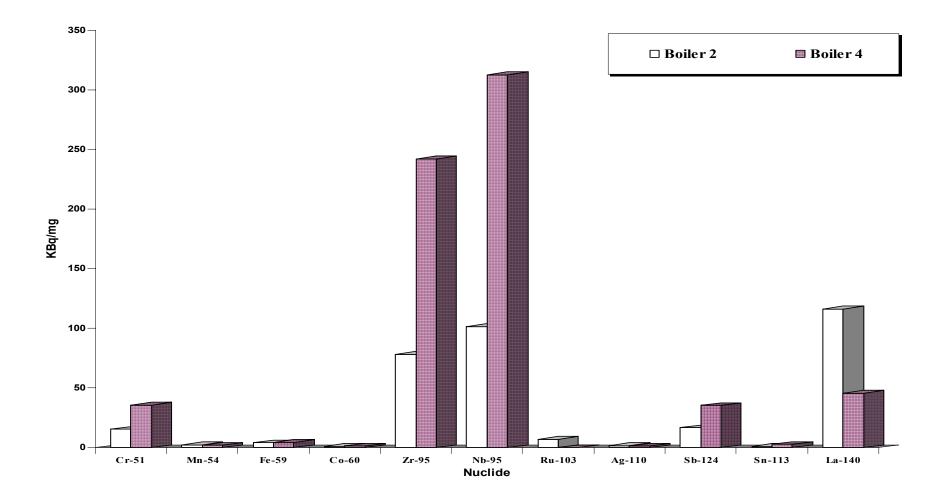
- Main corrosion product in the HTS is magnetite.
- Co-60 the main contributor to radiation fields on out-of-core piping and components. The Co-60 is in the magnetite. Transport of Co-60 is influenced by solubility and corrosion of iron in the system.
- Main method of reducing radiation fields is to reduce magnetite deposition in the reactor core.
- Deposition of magnetite in the reactor core is influenced by temperature and coolant pH.

## Crud

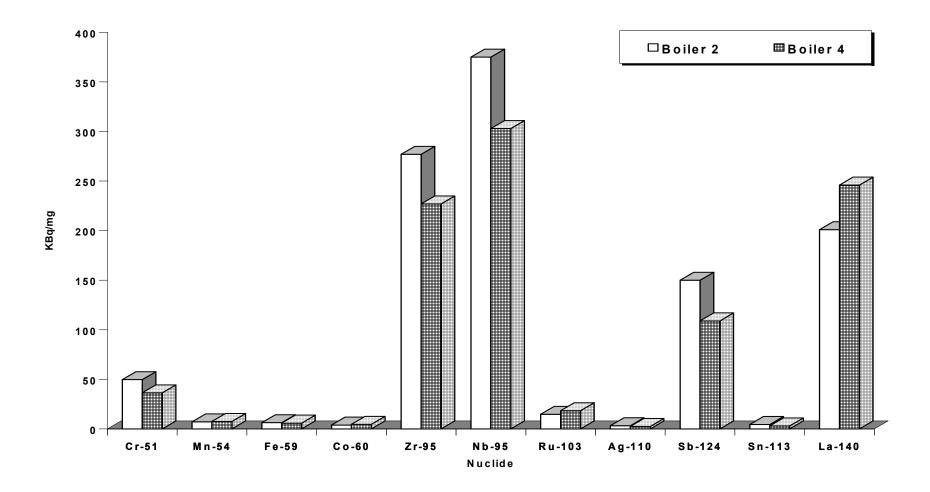
### **Reducing System Crud & Radiation Fields**

- Elimination of cobalt from materials of construction in the Primary Heat Transport System.
- Maintain proper system chemistry to reduce magnetite deposition in the reactor core. High pH and low dissolved oxygen required.

Comparison of Loop 1 & Loop 2 Crud Analysis at Point Lepreau January 1, 1994



Comparison of Loop 1 & Loop 2 Crud Analysis at Point Lepreau November 7, 1994



## Chloride

### Main Concern

- Stress Corrosion Cracking of Stainless Steel and Nickel based alloys. Examples include fuel channel end fittings, closure plugs, steam generator tubes.
- Pitting & General corrosion of carbon steel. This can occur in areas of low flow such as crevices where chloride can concentrate. Not as big of a concern as in the Secondary Side due to the fact that there are fewer crevices and low flow areas in the HTS.
- The presence of oxygen, greatly increases the effects of chloride in the system.

## Iodine-131

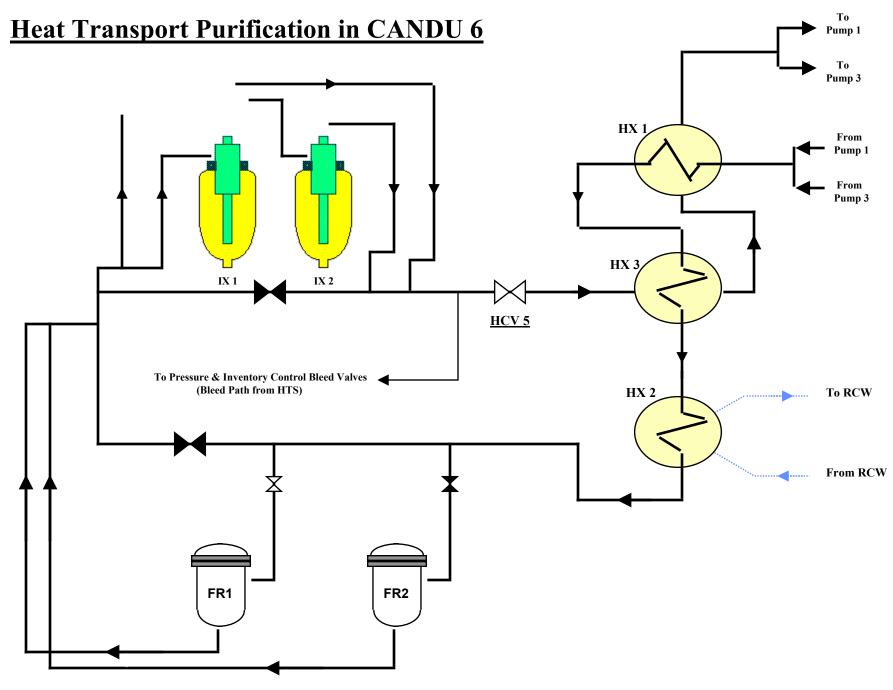
- Radiodines (I-131) levels are controlled in the HTS to a limiting value to minimize the radiation dose received by plant staff and the population in the event of an accident loss of coolant accident.
- Part of the Station Operating License OP & P. Shutdown limit of 500 MBq/Kg.
- There are no chemistry or corrosion issues related to I-131 in the HTS because the chemical concentrations are extremely low. For example, 500 MBq/Kg corresponds to a concentration of about 0.12 ug/Kg (ppb).

### **System Purpose**

- To minimize the build-up of activated corrosion products on PHTS components which are outside the core.
- To control the concentration of fission products, iodine in particular, released from fuel defects.
- To assist in controlling the coolant pH and coolant chemistry.
- To provide a source of clean water for Heat Transport bleed requirements.

## **Major Components**

- Purification Intercoolers (2)
- Purification Cooler
- Filters (2)
- Ion Exchange Vessels (2)



### **Purification Intercoolers - HX1 & HX3**

• D<sub>2</sub>O leaving the HTS flows through the tube side of each of the Intercoolers and is cooled by colder D<sub>2</sub>O returning to the HTS. Cools water entering the purification System and also heats it back up again prior to re-entering the main system.

#### **HXI Conditions**

<u>Tube Side</u>		Shell Side		
Inlet Temperature Outlet Temperature	265°С 194°С			
		HX3 Conditions		
<u>Tube Side</u>		Shell Side		
Inlet Temperature Outlet Temperature	194ºC 103ºC	Inlet Temperature Outlet Temperature	65°С 159°С	

## **Purification Cooler - HX2**

- Cools the D<sub>2</sub>O leaving the Interchangers (HX3) to 65°C. Water must be cooled to this temperature in order to prevent resin damage.
- RCW supplied to the shell side of the HX. Temperature adjusted by regulating the RCW flow.
- Purification is isolated if HX2 outlet temperature reaches 85°C.

## Filters -FR1 & FR2

- Removes insoluble matter (crud) from the D<sub>2</sub>O before it enters the Ion Exchange columns.
- Nominal Filter Rating 5 uM
- Normal operation has one filter in service and the other on standby.

### Ion Exchange Vessels - IX1 & IX2

- Two carbon steel vessels.
- Each vessel contains 1080 liters of mixed bed ion exchange resin (cation resin Li<sup>+</sup> based).
- Normal operation is with both vessels in service.
- Normal purification flow rate is 24 Kg/sec (maximum flow).

## **Design Criteria**

- Original design of the HTS Purification System was based on minimizing the build-up of activated corrosion products on HTS components outside the core (reducing out or core radiation fields).
- Crud activated in-core will re-deposit on out of core components.
- If the purification system is to reduce activity build-up, its cleanup half time must be equal to the crud re-deposition half-time.

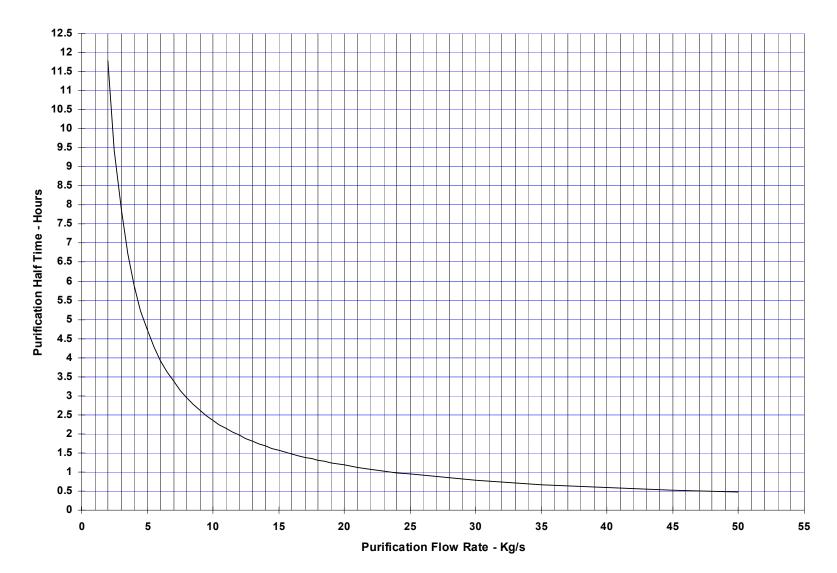
## Design Criteria

- Original design of the HTS Purification System (CANDU 600) was based on experience at Douglas Point (had the most EFPD's). Crud deposition studies were conducted at this plant and the crud deposition half-time determined to be 8-12 minutes.
- Based on the above deposition half-time, purification flows of 70.7 Kg/sec are required ( $T_{1/2} = 0.34$  hours).
- The design of the Purification System on future CANDU 600 Stations was based on this experience at Douglas Point.

## Design Criteria

- Later experience at Pickering "A" and Bruce "A" indicated that radiation fields (from activated corrosion products & fission products) were kept under control with purification half-lives between 2 to 3 hours.
- On this basis, the recommended purification half-life for CANDU 600 Plants is one hour. This corresponds to a purification flow rate of 24 Kg/sec.
- Purification flow rate should normally be operated at 24 Kg/sec (maximum flow rate). This will ensure low crud levels & radiation fields throughout the system.

#### **CANDU 6 - Primary Heat Transport System** <u>Purification Half Time as a function of Purification Flow Rate</u>



#### **Storage Tank Cover Gas**

Parameter	High Specification	<b>Upper Control Limit</b>
D <sub>2</sub>	4.0%	2.0%
$O_2$	1.0%	0.5%
$N_2$	6.0%	4.0%

#### **Storage Tank Cover Gas**

#### Deuterium

- Sources
  - Dissolved Deuterium is produced from the radiolytic decomposition of  $D_2O$ .
  - System Corrosion.
  - Addition of Hydrogen to the Main System for dissolved oxygen control.
  - $D_2O$  from the main system exchanges with the  $D_2O$  in the storage tank. Dissolved  $D_2$  transfers from the water into the storage tank Helium Cover Gas. The reduction in pressure in going from the main system to the storage tank decreases the solubility of the dissolved deuterium thus increasing the degassing process.

#### **Storage Tank Cover Gas**

#### Deuterium

#### • Upper Control Limit

 Derived from the flammability limits for hydrogen/helium/oxygen mixtures and is set at 2% providing a safety factor of four below the lower flammability limit of 8%.

#### • Specification

- Probability of an explosion occuring in the Storage Tank is much lower than in the Moderator Cover Gas since there are no recombiners or compressors - the main potential ignition sources. Specification set at 4% providing a safety factor of two below the flammability limit.
- There is no shut down specification associated with Storage Tank  $%D_2$  (as compared to the Moderator Cover Gas). This is because shutdown does not immediately reduce the production of deuterium (as in the moderator) and also due to the lack of ignition sources.

#### **Storage Tank Cover Gas**

#### Oxygen

#### • Sources

- Primarily from air in-leakage into the storage tank.

#### • Specification

- Set at 1%. Ensures that the concentration required to create a flammable mixture is not reached (4% needed given enough hydrogen). The 1% limit provides a large safety factor.
- The limit of 0.5% minimizes oxygen mass transfer from the gas into the liquid. Increases in the dissolved oxygen content of the storage tank  $D_2O$  and indirectly the main system through mixing are minimized.

#### **Storage Tank Cover Gas**

#### Nitrogen

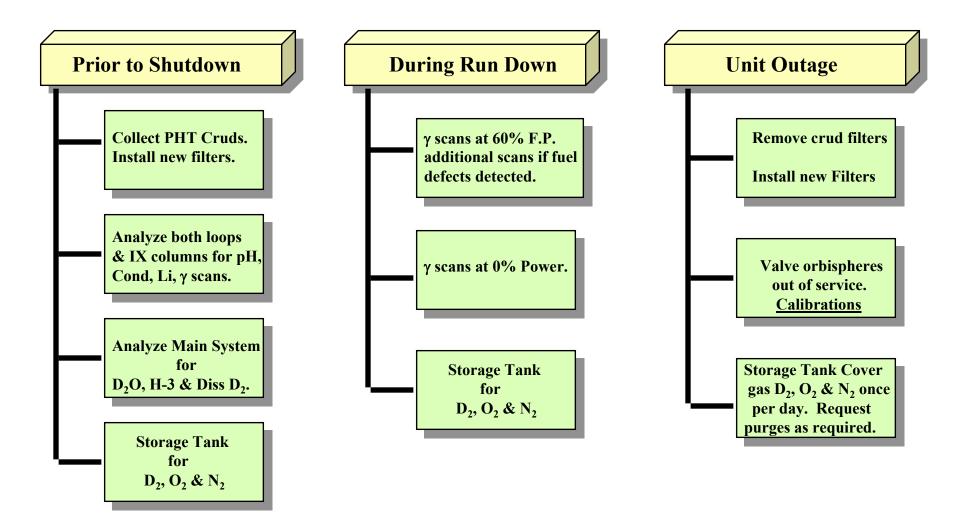
#### • Sources

- Primarily from air in-leakage into the storage tank.

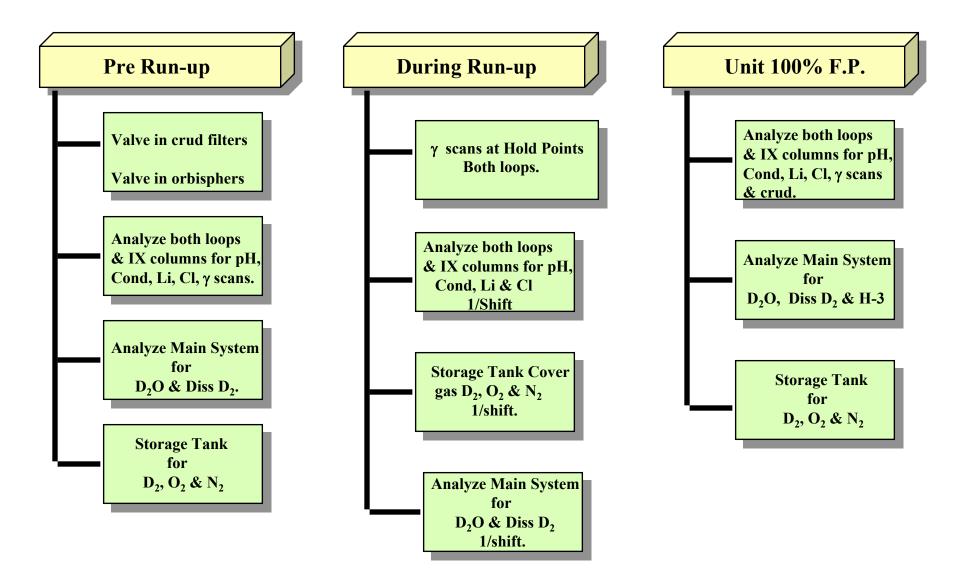
#### • Specification

- Nitrogen can result in the production of nitrogen oxides by radiolysis which can then react with water producing nitric acid.
- High levels of nitric acid increase the load on the heat transport ion exchange columns.

#### **Chemistry Monitoring Requirements**



#### **Chemistry Monitoring Requirements**



#### **Typical Activation Products**

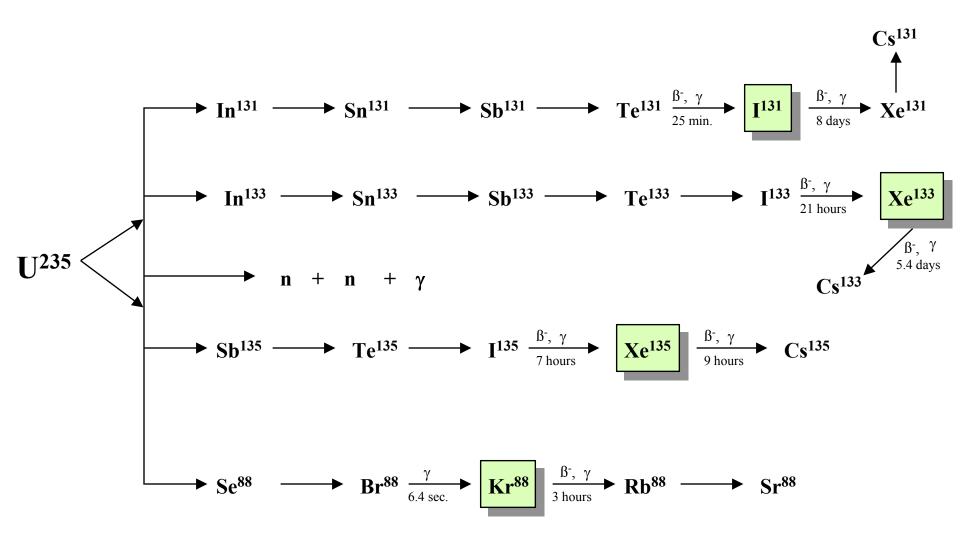
N U C L I D E	HALF-LIFE	MAJOR SOURCE
Н -3	12.26 years	D <sub>2</sub> O
N -16	7.1 seconds	D <sub>2</sub> O
N -17	4.2 seconds	D <sub>2</sub> O
0 - 19	29 seconds	D <sub>2</sub> O
A r - 4 1	1.83 hours	A ir ingress
C r - 51	27.8 days	Steel & Incoloy
M n - 54	312 days	Steel & Incoloy
F e-59	45 days	Steel & Incoloy
C o -58	71 days	Nickel Alloys
C o - 6 0	5.24 years	Steel & Incoloy
Z r - 95	65 days	Zircoly
S b - 1 2 4	60 days	Carbon Steel

#### Heat Transport Standard Operating Conditions

SYSTEM	PARAMETER	SPECIFICATION	VALUE	CONTROL METHOD
	рН	* 10.3 - 10.7	10.3 - 10.5	LiOD IX resins
Main HTS	Dissolved D .	* 3-10 oo/kg	7	H, addition
	cı.	< 0.2 mg/kg	* ALARA	1X resins. Bleed cooler temperature over-nde
	I-131	(1)	* ALARA	IX resins. Removing delective fuel.
HTS Storage Tank Cover Gas	D O N .	D, 5 2% O, 5 2% N, 5 8%	ALARA	Purging with Helium.
(1) Pickering is required to shut a unit down if an inventory of 1000 curies is reached. At the Bruce and Dartington stations, the shutdown limit is ≥ 500 curies for ≥ 8 hours.				

#### Overhead 78 AECB Conventional/Nuclear General

Fission of U<sup>235</sup>



#### **Sources of Fission Products**

### • Uranium Contamination

• Uranium contamination in the fuel cladding and other zirconium structural materials in the core. Total approximate uranium contamination present 3.5 mg/kg (Tramp Uranium).

### • Fuel Failures

• Penetration of the fuel bundle cladding.

### **Control of Fission Products**

### • Location of failed fuel bundles in the core

- Gaseous Fission Product Monitor
- Delayed Neutron Monitor
- Removal of soluble ionic fission products by the HTS Purification System
  - Iodine's and Cesium

## • Removal of fission product gases by degassing the HTS

- Noble Gases Xenon, Krypton
- Abnormal mode of operation

## **Failed Fuel Detection & Location Systems**

- Detection Using:
  - Gaseous Fission Product Monitor on-line

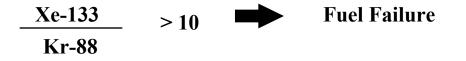
- Location Using:
  - Delayed Neutron Monitoring

### **Gaseous Fission Product Monitor**

- The Gaseous Fission Product Monitor (GFP) is a computer controlled, high resolution gamma ray spectrometer. It operates continuously and measures Xe-133, Kr-88, Xe-135 and I-131 present in continuous sample flows from each of the two Heat Transport loops.
- The relative concentrations of the various nuclides will indicate the presence of failed fuel as well as give information on the size of the defect.

### **Fuel Failures**

- Main indicator of a fuel failure is an increase in the Xe-133 concentration.
  - Normal Xe-133 levels in the HTS coolant are 1-2 MBq/Kg (core free from fuel defects).
  - When a small leak develops, an increase in Xe-133 & Kr-88 will be observed (Xe-133 will increase the most).
  - Confirmation of a defective fuel bundle can be done by comparing the ratio of Xe-133 to Kr-88. A ratio of > 10 would confirm the presence of a leak.



## **Half-Lives of nuclides monitored by the GFP**

- Xe-133 5.4 days
- Xe-135 7 hours
- Kr-88 3 hours
- I-131 8 days

### <u>Fuel Failures</u>

#### • No Leaks

- Fission products are contained within the fuel sheath.

#### • Small leak

 Fission product gases are allowed to escape from the small hole in the fuel sheathing and enter the coolant. Water must enter the fuel sheath in order to permit Iodine release. Water is needed to dissolve the iodine's and make them available for release. Very little water will enter the fuel sheath with small defects, thus very little increase in I-131 concentrations will be observed.

#### • Large Leak

 Coolant can enter the fuel element dissolving the iodine's in the process. This type of failure will result in large increases in coolant iodine's & fission gas concentrations.

## **Types of Fuel Failures**

- Type 1
  - A pinhole defect in new fuel. Results in xenon & krypton releases. Low iodine releases.
- **Type 2** 
  - A pinhole defect in fuel at one-half burnup. Results in release rates about 2.5 times higher than Type 1. Fission products are in the same ratio as in a Type 1 leak (high xenon & krypton low iodine's).
- Type 3
  - A small crack in the fuel cladding at one-half burnup. Results in similar Xe-133 release rates as a Type 2. Much higher Kr-88 and especially I-131 release rates. Thus the main indicator is the isotopic proportions.

## **Gaseous Fission Product Monitor**

- Xe-133
  - Long lived fission product (5.4 day half-life). It has a high release rate from defective fuel.
  - Its concentration when compared to the short lived fission gas Kr-88, provides information about the source, the extent of fuel sheath damage and the buildup of Tramp uranium in the core. Main indicator of failed fuel.
- Kr-88
  - High release rate from fuel defects. Lower concentration in coolant due to its low half-life.

## **Gaseous Fission Product Monitor**

#### • Xe-135

 Provides information about the iodine release rates when high purification rates are removing fission products. I-135 decays to Xe-135 in the IX system. The Xe-135 is not retained by the resin, it becomes a secondary source of Xe-135.

#### • I-131

• Not a reliable indicator for assessing fuel damage due to the fact that it is removed by the purification system. Monitored for public safety reasons due to its biological hazard (OP& P limitations on its concentration in the coolant).

## **Delayed Neutron Monitor**

- Design Basis for CANDU 6
  - Based on KANUPP design
  - Uses samples extracted from outlet feeders
  - Monitors for presence of delayed neutrons

### **Delayed Neutron Monitor**

- Used to locate the fuel channel containing the defective fuel.
- Locate the position of the defective fuel within the fuel channel.

Sample lines from each of the 380 fuel channels carry coolant to the DN Rooms (RI-303 & RI-304). Each sample then flows through sample coils which are contained in cylinders called sample coil dry wells. Each of these dry wells are located in a moderator tank filled with light water. The light water is used for cooling and also to thermalize the delayed neutrons which is required for accurate detection.

Above the moderator tank is located a carriage which is used to carry and position the thermal neutron detectors  $(BF_3)$  into the sample coil dry wells. During fuel sheath failure, I-137 & Br-87 are released into the coolant. Both are delayed neutron emitters and will be detected by the  $BF_3$  detectors.

## **Delayed Neutron Monitor**

Isotope	Half-Life	Source
N-17	4 seconds	Activation
N-16	7 seconds	Activation
I-137	22 seconds	Failed Fuel & Tramp Uranium
<b>Br-87</b>	57 seconds	Failed Fuel & Tramp Uranium

The four neutron emitting isotopes are always present in the core. N-17 & N-16 are activation products. I-137 & Br-87 are due to the presence of Tramp Uranium. Sample lines are sized as to provide a 50 second delay time before the sample reaches the detector. This is to allow for decay of the N-17 & N-16 in the sample.

During a fuel sheath failure, I-137 & Br-87 will be released into the coolant and thus their concentrations will increase above background concentrations.