

HW 79768

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CHEMICAL PROCESSING DEPARTMENT
MONTHLY REPORT
FOR

HW--79768

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NOVEMBER, 1963

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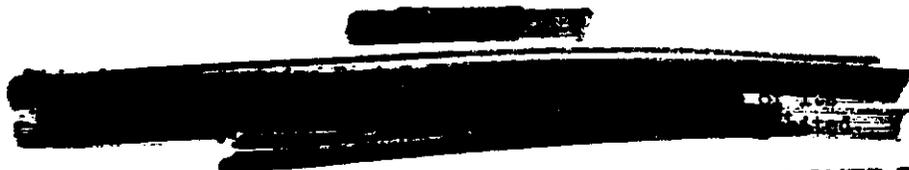
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OPERATION MANAGERS

December 20, 1963

HANFORD ATOMIC PRODUCTS OPERATION
RICHLAND, WASHINGTON

Work performed under Contract No. AT(45-1)-1350 between the Atomic Energy Commission and General Electric Company.



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I. SUMMARY

Production through November, as compared with the October 25, 1963 HAPO Production Forecast HW-79264, is summarized below:

	<u>Percent of Forecast Achieved</u>	
	<u>November</u>	<u>Fiscal Year To Date</u>
Separated plutonium nitrate	80.6	92.1
Separated uranium nitrate	83.6	90.9
Uranium oxide	78.8	94.2
Plutonium metal buttons	99.3	101.2
Fabricated parts	111.1	103.8

November production met or exceeded forecasted quantities for plutonium metal buttons and fabricated parts. With the Redox plant shutdown most of the month as a result of a fire in the 233-S Plutonium Concentration Building, forecasted quantities were not achieved for separated plutonium nitrate, separated uranium nitrate, and uranium oxide.

At approximately 1:15 a.m. on November 6, a fire of undetermined origin occurred in the Redox 233-S Building, damaging the plutonium anion exchange and concentration systems. No one was injured. The plant was immediately shut down and remained down for the balance of the month. Decontamination and rehabilitation work are now in progress, directed toward getting the final plutonium concentrator in operation so that Redox plant operation can be resumed.

The Purex plant resumed processing on November 4, following a scheduled shutdown which began October 27. Operation was then continuous at an average rate of 29.5 tons per day. Both the plutonium and uranium products met specifications except during the initial startup period.

A Purex plant process test was run to determine the effects of eliminating the feed adjustment step between the dissolvers and the HA first decontamination column feed. Improved HA column control resulted, with no adverse process effects noted.

The HAPO II-1 cask, loaded last month with approximately 140,000 curies of strontium-90, was released to the Commission on November 22 for shipment. Four SIT casks, containing an estimated 189,000 curies of cesium-137, and an ORNL cask, containing approximately 49,000 curies of promethium-147, were released to the Commission on November 27 for shipment. The 1 Kg of technetium-99, recovered last month, was loaded into a cask and shipped to Hanford laboratories, where it will be converted to the metal.

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A total of eleven direct PuO₂ calcination runs were made during November in the laboratory titanium screw calciner. Approximately 11 Kg of plutonium oxide were produced. The tests indicated that a readily chlorinated oxide is produced with high sulfate ratios. No substantial corrosion of the calciner was evident.

Funds for Project CAC-965, "In-Tank Waste Solidification - 200 East Area", were increased by an interim authorization of \$100,000, bringing the total to \$280,000.

Directive No. EQT-007 authorized \$75,000 to provide necessary facilities, in conjunction with the 242-Z Building Waste Treatment Facility, which will permit the recovery of americium from the solvent extraction waste stream.

A series of engineering studies, to establish a firm basis for Phase III Waste Management design, were completed. The equipment can be designed to accommodate either of two solvent extraction flowsheets. This will permit equipment design to be firmed up now, while preserving flexibility for better choice of the most economical waste management process.

J. S. Wasson
General Manager
Chemical Processing Department

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II. ACHIEVEMENTS

A. PRODUCTION OPERATION

1. <u>Production Statistics</u>	Fiscal Year	
	<u>November</u>	<u>to Date</u>
a. <u>Percent of Forecast⁽¹⁾ Achieved</u>		
Separated plutonium nitrate	80.6	92.1
Separated uranium nitrate	83.6	90.9
Uranium oxide	78.8	94.2
Plutonium metal buttons	99.3	101.2
Fabricated parts	111.1	103.8
b. <u>Purex</u>	<u>November</u>	<u>October</u>
Uranium nitrate produced (tons)	721.07	453.56
Average production rate during operation (T/D)	29.5	20.3
Total waste loss (%)		
Plutonium	0.34	0.36
Uranium	0.25	0.25
On-line efficiency (%)	89.7	88.7
c. <u>Redox</u>		
Uranium nitrate produced (tons)	10.49	38.41
Average production rate during operation (T/D)	5.3	7.4
Total waste loss (%)		
Plutonium	0.73	0.71
Uranium	0.34	0.29
On-line efficiency (%)	7	25
d. <u>Uranium Reduction (tons)</u>		
Normal UO ₃ loaded	577	541
Enriched UO ₃ loaded	10	55
Normal UO ₃ approved for shipment	551.70	648.79
Enriched UO ₃ approved for shipment	0	50.80
Normal UO ₃ shipped	451.82	648.95
Enriched UO ₃ shipped	0	101.66
Normal UNH backlog	370	226
Enriched UNH backlog	120	120

(1) HW-79264, HAPO PRODUCTION FORECAST, dated 10/25/63.

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e. Plutonium Metal Processing

	<u>November</u>	<u>October</u>
Reduction yield (%)	97.7	98.24
Product recovery output (Kgs)	79.02	213
Product recovery backlog (Kgs)	1387	1388
Waste disposal (grams)	260	241

f. Power

	<u>200-East</u>	<u>200-West</u>
Raw water pumped (gpm)	11,336	4,504
Filtered water pumped (gpm)	1,199	1,159
Maximum steam generated (lbs./hr.)	250,000	96,000
Average steam generated (lbs./hr.)	206,881	89,490
Total steam generated (M lbs.)	134,059	57,944
Coal consumed (tons)	7,150	3,116

November production met or exceeded forecasted quantities for plutonium metal buttons and fabricated parts. With the Redox plant shutdown most of the month as a result of a fire in the 233-S Plutonium Concentration Building, forecasted quantities were not achieved for separated plutonium nitrate, separated uranium nitrate, and uranium oxide.

E. E. Smith
for Manager, Production

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2. FISSION PRODUCT DATA

	<u>PUREX* RECOVERY</u>	<u>244-CR* INVENTORY</u>	<u>B-CANYON* INVENTORY</u>	<u>SEMWORKS PRODUCTION</u>	<u>PRODUCT INVENTORY</u>	<u>PRODUCT IN CASK</u>	<u>PRODUCT SHIPPED</u>	<u>CUSTOMER</u>	<u>REMARKS</u>
Sr-90 (Kilocuries)	56	300	1800	--	1830	--	140	ORNL	HAFO II-1 11-22-63
Cs-137 (Kilocuries)							189	ORNL	11-29-63
Ce-144 (Kilocuries)									
Pm-147 (Kilocuries)							49	ORNL	11-29-63

* Estimated figures based on Cs. calculations

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II. ACHIEVEMENTS (Continued)

B. PUREX OPERATION

1. Operating Continuity

Purex operation resumed November 4, 1963 after a plant outage which began October 27, 1963. By November 11, a capacity factor of 3.6 was reached and maintained the balance of the month.

A second B Plant oxalate run for Strontium-90 recovery was completed and preparations made for a peroxyacetate run for cerium-rare earth separation.

The C3 dissolver was reactivated on November 5. The unit contained three buckets of approximately 84-day-old metal and had been in standby since September 2.

2. Processing

Both uranium and plutonium have met product specifications except during the initial startup period.

Neptunium recovery during the month was satisfactory. No neptunium purification runs were completed.

Waste losses to underground storage during the period were .27 percent uranium and .37 percent plutonium.

The cask loading status is as follows:

<u>Cask</u>	<u>Quantity(Approx.)</u>	<u>Destination</u>	<u>Status</u>
ORNL Pm	49 Kc Pm-147	ORNL	Loaded in October. Released to AEC for shipment 11-27-63
STT's (4)	189 Kc Cs-137	ORNL	Loaded in October. Released to AEC for shipment 11-27-63
HAPO II-1	140 Kc Sr-90	ORNL	Loaded in October. Released to AEC for shipment 11-22-63
HL 40-Gal	~1025 Grams Tc-99	HL	Loaded in November. Sent to Hanford Labs on 11-18-63

HAPO I-B-1 and HAPO II-2 are off site.

Approximately 170 Kc of high nitrate strontium was reworked at Strontium Semiworks with very good results. Waste losses were about 2 percent.

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The elution of the technetium cask at Semiworks was completed and the 1025 grams of Tc-99 product was loaded into a Hanford Laboratories 40-gallon cask. Overall waste losses were about one percent. The cask was shipped to HL on November 18.

In connection with the sluicing program at 241-A Tank Farm, the supernate from Tank 102-A was transferred to Tanks 105-C, 106-C, and 101-C. Tanks 105-C and 106-C are now full and an approximate five foot pump heel was left in 102-A. The pump from 102-A was moved to 103-A and the supernate from this tank will be pumped to 101-C.

3. Equipment Experience

Two equipment failures were experienced on the east canyon crane in the Purex Plant. These failures involved the charging impact wrench and right hand optic head. Reconditioned replacement spares were installed.

The two cables on the railroad car puller, located in the Purex Building railroad tunnel, failed through normal wear and long service. New replacement cables were installed and the equipment returned to service.

Two Purex canyon pump failures were experienced during the current production run period. Reconditioned spare pumps were installed as replacements.

There were three canyon agitator failures this report period. These failures involved agitators F-7, F-13, and F-16. Cause of the failures is unknown at the present time. New replacements from Spare Equipment were installed.

The 6-1 cesium-rare earth storage tank in B Plant developed a leak in the coil on 11-6-63. The product was transferred to 7-1 tank. Piping modifications have been made to permit flushing tank 6-1 prior to replacement with a tank from U Plant.

4. Radiation Experience

The retention basins north of B Plant became contaminated due to the failure of the coil in the B Plant cesium-rare earth 6-1 storage tank on 11-6-63. General dose rates around the basins were 500 mrad/hr. with tumbleweeds reading up to 50 rads/hr. at surface. The wind carried a few highly contaminated tumbleweeds to the north perimeter fence. Contamination was also experienced around the basins, in the area northeast of the basins, and in the first 1000 feet of the drainage ditch leading to the swamp. The following work was accomplished to restore the area to near normal:

- a. The basins and adjacent areas were decontaminated.
- b. The inside of the north basin was coated with a sealer.
- c. A fence was erected around the basins.
- d. Approximately 1000 feet of the old ditch was backfilled and a new ditch was dug to replace this portion.

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WM Harty
Manager-Purex

WM Harty:OVS:gt

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II. ACHIEVEMENTS (Continued)

C. REDOX OPERATION

1. Operating Continuity

At approximately 1:15 a.m. on November 6, 1963, a fire of undetermined origin occurred in the 233-S Building and damaged the plutonium anion exchange and concentration systems. The Redox plant was immediately shut down and remained so for the balance of the month. Decontamination and rehabilitation work are now in progress and are directed toward getting the final plutonium concentrator in operation so that Redox plant operation can be resumed.

The Uranium Oxide plant operated at satisfactory rates throughout the month. Enriched uranium processing was discontinued following the 233-S fire at the Redox plant, and the calciner normally used for this material was converted to depleted uranium processing.

2. Processing Operations

a. Redox Processing

The fire in the 233-S Building occurred in the process area, near the ion-exchange contactor, but is believed to have started outside the process equipment enclosure part of which was rigid, transparent plastic panelling. The enclosure was breached by the fire and subsequent smoke "flashback", thus allowing the spread of alpha contamination from the interior of the grossly contaminated plutonium concentration "cell".

No one was injured. A total of thirteen people received skin or clothing contamination. Skin contamination, which to some individuals exceeded 4×10^4 d/m, was subsequently reduced to non-detectable during the shift on which the fire occurred. Nasal smears taken were negative.

Plutonium released from the processing cell grossly contaminated areas in the 233-S Building and nearby ground and building surfaces outside the building. Actual physical damage was confined to the viewing room and process area in the vicinity of the L-18 plutonium anion exchanger. Plastic hood panels and electrical wiring, including lights, were extensively damaged in the north end of the viewing room. The contactor loop itself does not appear to be damaged; however, gaskets, pumps, valves and electrical wiring to the equipment in the contactor enclosure are damaged. None of the process equipment other than the L-18 anion exchange contactor

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appears to have been damaged.

Action plans were formulated at once and all resources possible in men, materials, and services were directed toward decontaminating and rehabilitating the 233-S Building. Loose contamination was removed from the exterior surface of the building and the surfaces of the walkways and grounds immediately adjacent. Contamination which could not be removed from exteriors of the buildings and ground areas has been "fixed" with tar, paint, roofing compound, or masonry sealer, as appropriate to the surface.

Decontamination of all contaminated areas in the normal "cold side" areas of the building was completed, and such areas were restored to occupancy in less than two weeks.

Clean-up of the viewing room and stairwell proceeded less rapidly although good progress has been made. By month-end all viewing room and stairwell levels had been cleaned of loose contamination and given two coats of paint to "fix", as much as possible, any residual contamination.

Also by month-end, the plutonium solutions in the L-3 concentrator, the L-4 product receiver, the L-6 sampler and L-7 loadout tanks had been emptied into FR cans for storage. At the end of the month, plastic windows were re-installed in front of the L-18 contactor in preparation for the initial leach of the resin, and a complete barrier had been reestablished (on a temporary basis) between viewing room and the process cell. In general, good progress was made in decontaminating the service areas of the 233-S Building and concurrent action has been started to reactivate the concentrator and loadout equipment in this building. The goal date of December 15 was established for the start of limited production in the Redox plant.

b. Uranium Oxide Processing

Processing operations in the Uranium Oxide plant were started on November 6, as scheduled, following replacement of the jacket on the X-30 calciner feed tank (concentrated, depleted uranyl nitrate). Dilute feed plus a filter plugging problem curtailed initial production rates, but these problems were subsequently resolved and full rates were restored by November 11.

Due to the Redox plant shutdown on November 6, the G-cell calciner was converted to depleted uranium production and remained on this type feed for the balance of the month.

3. Mechanical Experience

a. Redox Plant

The major portion of the maintenance activity this month was in

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direct support of the accelerated program to restore the Redox plant to operational status.

Concurrent with the 233-S Building decontamination, work was started to return the Redox 202-S canyon system to an earlier process flow scheme employing three solvent extraction cycles for plutonium decontamination and accumulating neptunium for recovery and decontamination on a campaign basis. Major work in the canyon involved the installation of a tower and 15 new jumpers for the E-4 tank. A mock-up of this installation was made in the 200-W shops and work in the cell was nearing completion at month end. Related work in F and G cells was also essentially completed.

Re-routing of the vessel vent system (from 233-S Building ventilation discharge to the 202-S Building ventilation discharge) was completed during the latter part of the month.

Other significant 233-S Building rehabilitation work included: 1) replacement of electrical wiring leading into the viewing room to replace that burned out, and reestablishing receptacle circuits; 2) repair and calibration of all process instruments needed for the revised process flow scheme.


Manager - Redox


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II. ACHIEVEMENTS (Continued)

D. WEAPONS MANUFACTURING OPERATION

1. Operating Continuity

Fabrication of weapons models 1807 and 74-C continued during the month. Unfabricated plutonium production was limited by the availability of feed from the primary plants. The plutonium dissolvers were shut down for a portion of the month because of the lack of primary plant feed which could be used for blending dissolved scrap. The incinerator was operated when manpower was available.

2. Processing Operations

a. Plutonium Fabrication

Information on fabrication activities is presented in Document HW-79922 (Atomic Weapon Data).

b. Plutonium Reduction

Unclassified plutonium and scrap powder processing occupied the Button Line during the early part of November, with weapons grade plutonium being processed during the remainder of the month. At first, the Pu-240 content of the latter material was above the specification, but a gradual decreasing trend resulted in an acceptable Pu-240 concentration being reached by month end.

The negative MUF (material unaccounted for) across the Button Line for the month of October was under investigation during the entire month. Principal items still under study at month end are: possible overstatement of receipts from primary plants; possible overstatement of slag and crucible account (based on neutron count); calculations of reduction yields; and vessel holdup.

c. Plutonium Reclamation

The dissolvers operated satisfactorily on normal weapon grade material.

Incinerator operation was also satisfactory with 27 boxes of scrap being processed.

In the new recovery facility, primary calibration of the tanks was completed. However, rechecks will be required at a later date when instrument facilities become available. Panel board activation has

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c. Plutonium Reclamation (Continued)

been delayed pending completion of modifications required to correct sub-standard work by the Phase II contractor. Delay resulting from this is currently estimated at two weeks.

Removal of sludge from the floor of the RB (reception and blending) hood of the old Recuplex facility was started during the month.

3. Mechanical Performance

The fabrication equipment on both the RMA and RMC lines operated well during November.

The Button Line equipment continued to require considerable maintenance attention. Principal difficulty involved plugging of the vacuum drum filter by oxalate from the processing of metal recovery solutions and ion exchange product solution. Improved operation was experienced when the processing of normal feed was resumed at mid-month.

The new double-ball valve between the calciner and fluorinator has been removed for further development work. It has been replaced temporarily by one of the former "B" valves.

4. Radiation Experience

Radiation and contamination control statistics revealed satisfactory performance in November.

During the extinguishment of a fire in a process hood in Room 227, the hood was pressurized during the use of a dry chemical extinguisher. As a result, the room was contaminated but no plutonium deposition to the three employees involved was experienced as they were all equipped with proper respiratory protection.

5. Analytical Experience

	<u>November</u>
Number of Samples Received	2,292
Number of Determinations	17,715

The Weapons Manufacturing Analytical Laboratory, in cooperation with the Purex Analytical Laboratory, is engaged in a study to define the accuracy of the plutonium concentration determination for Purex L-10 solutions.

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5. Analytical Experience (Continued)

	November		
	<u>Weapons Grade</u>	<u>Special**</u>	<u>Unclass.</u>
Total Impurities, Buttons (Avg.)	3100	3660	3250
Buttons Rejected for Impurities	12%	20%	10%
Pu-240 Content*	6.095%	6.515%	8.54%***

*Based on Neutron Counter Method.

**Above weapon grade specifications for Pu-240.

***Average for October applies.

W. J. Gartin
Manager
Weapons Manufacturing

WJ Gartin: csj

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II. ACHIEVEMENTS (Continued)

E. POWER AND CRAFTS OPERATION

1. Operating Continuity

There were no interruptions of steam, water or electrical services to the production facilities that affected continuity of operation during this report period.

2. Inspection, Maintenance and Repair

Ten electrical housing castings were machined, on a priority basis, for Spare Parts Records Operation. These parts are normally purchased off-site from a vendor whose business is closed down at this time. The subject castings are used on electrical call pipe jumpers and on process pumps and agitators.

Special emphasis was placed on modifications to the conveyor rollers in Weapons Manufacturing Operation's sprue removal hood. The urgency was necessary to meet scheduled project completion dates.

Fabrication of a spare D-12, D-14 Concentrator for the Redox facility continued to progress and was an estimated 55% complete at month's end.

A feed tank (F-1), complete with coil, was mocked up and made ready for service at the Redox facility.

New plexiglass panels were installed in hoods 41, 45 and 46 at Weapons Manufacturing Operation, as replacements for panels which were no longer transparent.

A temporary air exhaust system, including a blower, ductwork, exhaust filters, etc., was installed as required in the decontamination and reactivation of the 233-S Building, following the November 6 fire at that location.

Semi-permanent air locks were fabricated and installed at the 233-S Building for entry into and exit from certain areas in the building, as an aid to contamination control.

Alterations to the Weapons Manufacturing Operation's comparator hood are in progress. Work on the indicator holders, face plates and six jaw chucks was complete at month's end.

Thirty-eight pipe jumpers were fabricated during this report period. All were replacement requirements, 23 for Purex and 15 for Redox.

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Craft assistance was provided Hanford Laboratories personnel at 231-2 in placing in operation the "Instron," a mechanical testing apparatus for use in tensile compression and hardness testing of plutonium pieces. Included in the craft work was the checking out of a furnace in the new hood and a new instrument console located in close proximity to the hood.

An open circuit in the 200-West Area fire alarm loop was traced out and repaired, on an emergency call-out basis, November 14.

A 3,000' ditch was excavated to accommodate the B-Plant settling basins. A portion of the original drainage ditch for this facility had become contaminated as the result of a leaking coil in a process vessel at the Fission Products Processing Operation, 221-B. The original ditch was backfilled to prevent further spread of contamination.

Renovation of the 200 Area elevated water tanks by Jones forces is in progress. The 2901-B tank is out of service at this time, for installation of a new standpipe, heating element, frost box, etc.

At the request of IPD Engineering, 4" orifices have been installed in the bypass lines in both 200-East and West raw-water inlet houses. According to test data, this particular size orifice should prevent excessive surge pressures.

Assistance rendered other departments included balancing work on the ventilation systems in the N-Reactor Department's 105-N Building. The work was an estimated 55% complete at month's end.

T. S. LaFollette
Manager-Power & Crafts

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CHEMICAL PROCESSING DEPARTMENT
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II. ACHIEVEMENTS (continued)

F. FACILITIES ENGINEERING OPERATION

1. Purex

a. Process Design Engineering

Multipurpose Dissolvers

The program to increase processing flexibility at Purex has advanced with completion of the following designs: "A" Cell Arrangement, Side Elevation of "A" Cell, Annular Dissolver Assembly, Annular Dissolver Details, and Plan of Scrubber Solution Catch Tank. Scope design of a critically safe 2AF Tank (feed for 2AF Column) for Purex use is nearing completion. The engineering flow diagram was completed.

X-Ray Photometer

Performance of the X-ray photometer used in the Purex laboratory for final product analysis has declined beyond permissible limits. Failure has been attributed to the thyrite non-linear resistor in the amplifier. Since this HAPO-developed item is not available commercially, a replacement is being fabricated in the Instrument Development Shop.

b. Project Engineering

CAC-965 - In-Tank Waste Solidification - 200 East Area

Additional interim funds of \$100,000 authorized by Directive No. EQT-004, Mod. No. 1, dated November 12, have been allocated to the J. A. Jones Company (\$95,000) and to Vitro Engineering Corporation (\$5,000 for Title III work). Interim authorizations now total \$280,000.

CAC-970 - Purex Essential Waste Routings System

Bid opening for the fixed-price work, consisting of the piping systems and a diverter station, is scheduled for December 3.

c. Manufacturing Engineering

F-6 Concentrator

A remotely-operated television was used to view the Purex "F" Cell area adjacent to the QQ Nozzle, which has been leaking and

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causing some corrosion to the dunnage. The TV inspection permitted an evaluation of damage to supporting structures and it has been concluded that the structure has not been significantly affected. However, the TV inspections are being planned for 6-month intervals to discover any damage which might affect the integrity of the unit.

2. Redox

a. Process Design Engineering

233-S Incident

A decision has been reached to start the Redox Plant up using a three cycle flow sheet. In support of this decision, 18 new jumpers have been designed for the installation of a new ozone stripper and process re-routings.

Several additional items of engineering assistance were provided for safe entry and clean-up of the 233-S Building. Decontamination work has progressed to the stage of painting within the building viewing room.

Filter Box Survey

The ducts and filters of the 233-S Building exhaust were surveyed for evidence of plutonium accumulations. The readings taken, using the 380 KEV gamma method showed the sludge in the bottom of the filter box to contain approximately one gram of plutonium per pound. Measurements have been resumed since the fire.

b. Project Engineering

CGC-122 - Exhaust Ventilation System - 233-S Building

Directive No. HW-553, dated November 5, authorized interim funds of \$14,400 to the General Electric Company for the new exhaust ventilation facility at the 233-S Building

3. Weapons Manufacturing

a. Process Design Engineering

Recoverable Plutonium Partitioning Facilities

A design criteria document and drawings were prepared on the basis of a rough draft flowsheet, the feasibility of which has been demonstrated in the laboratory. Design criteria will now await completion and issuance of an approved flowsheet, tentatively set for February 1, 1964.

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"A" Fabrication Line

Installation of equipment and services in the blank preparation glove box (HA-29-EP) is being completed. The target date for "going hot" is December 2. Drawings for modifications of the optical comparator and its glove box were released to the 200 West Shops. Design criteria were completed for the plutonium thermal treatment facilities. This equipment is planned for hot-and-cold cycle treatment of the machining blanks to promote metal movement before machining and thereby improve dimensional stability. The facility will replace the temporary one now installed on the "A" Fabrication Line.

Survey of Vacuum Lines for Plutonium

Certain of the vacuum lines in the 234-5 Building were surveyed by the 380 KEV method to determine any increase of plutonium deposits in the past year. The quantity in the regions checked has not increased significantly.

b. Project Engineering

CAC-880 - Plutonium Reclamation Facility - "Z" Plant

The service contractor has begun making necessary piping and electrical modifications required before the plant can start up. Work is being directed toward completion in early January, 1964.

CAC-987 - Metal Stabilization Facility - 234-5 Building

The design criteria have been issued to the Architect-Engineer who has begun work on Title I Design. The isostatic press and its associated equipment is being fabricated.

CAC-102 - New Standards Laboratory - "Z" Plant

The Commission has stated that funds are not available in the budgets for FY-1964 and FY-1965, but can be provided for in the FY-1966 Budget. The Commission will hold the proposal for action, however, if funds should become available before that time.

CAC-121 - Americium Recovery Facility - "Z" Plant

Directive No. EQT-007, dated November 15, authorized \$75,000 for providing necessary facilities in conjunction with the Waste Treatment Facility, 242-Z Building, which will permit the recovery of americium from the solvent extraction waste (CWW) stream. The Architect-Engineer was given an interim authorization for \$31,000 to begin design. Work Authorization No. CAC-121(1), dated November 18, authorized \$4,000 to the Company

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c. Manufacturing Engineering

Power-rolled Model

Recent efforts have been directed toward improved design of the administrative forms and faster process information feedback. The following forms have been designed and approved: (1) 74-C - Machining, (2) 74-C - Stabilization, (3) Ingot Melting for 74-C and 1807, (4) 74-C Power-rolled Forming, and (5) 1807 Inspection Forms.

4. General

a. Process Design Engineering

Waste Management - Phase III

A series of engineering studies to establish a firm basis for Phase III Waste Management design have been completed. These studies have resulted in the development of a very attractive concept. The equipment can be designed to accommodate either of two solvent extraction flowsheets. This will permit equipment design to be firmed up now, while preserving flexibility for better choice of the most economical waste management process. In addition, the concept offers the potential for reducing later investments which might be required to add fission product purification facilities in B-Plant. Specific changes would include: (1) extraction equipment that could be operated on the flowsheet of either CSREX or D₂EHFA extraction, (2) Cs precipitation equipment, and (3) a storage tank for virgin Cs solution. These changes and additions increase the estimated construction cost from \$3,900,000 to \$4,200,000, but provide a potential savings of approximately \$1,000,000 on the future isotope packaging plant. The estimated scoping time will be extended approximately two months, or to June 1964.

244-AX Circulator Prototype

A prototype of the 244-AX Tank Farm waste circulator was completed by the Shops early in November. Preliminary calibration runs have been completed. Further design verification tests are planned to determine the hydraulic, solid carrying, and solid suspension characteristics.

"Plastic Man" Suits

A jacket assembly which employs the vortex cooling tube, and which has been used in the General Electric Lamp Department, was adapted for use with the tunnel-type plastic suit. Tests on-site have indicated that personnel can work several hours in the suit without overheating. Two jacket cooling assemblies are in use, and six more have been ordered.

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b. Project Engineering

Project Cost Information

Total Authorized Funds - 19 Active Projects	\$ 4 208 000
Total Cost-to-date	3 117 000
Commitments and Oper. Work Releases	318 000
Unencumbered Balance	773 000
Costs Charged to Above Projects (10/20/63 to 11/17/63)	65 000

CAC-117 - Modification of Steam Lines

Directive No. AEC-229, dated November 15, authorized \$200,000 for modification of steam lines in the 200 East and West Areas, of which \$4,700 was allocated to the Company. The Commission is preparing the bid package for this work.


Manager - Facilities Engineering

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CHEMICAL PROCESSING DEPARTMENT
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II. ACHIEVEMENTS (Continued)

G. RESEARCH AND ENGINEERING OPERATION

1. Purex Process Engineering

(a) Dissolver Charging Tests

Proposals have been made to use longer enriched fuel elements in the Hanford reactors. Simulated charging tests were made in a mocked-up dissolver annulus that has an OD of 70 inches and an annulus width of 10.5 inches. One thousand steel slugs measuring 1.5" x 12.565" were charged with no jamming or other difficulties noted.

(b) Solvent Extraction

The plant process test to determine the effect of eliminating the feed adjustment step between the dissolvers and the HA Column was continued during the month. The HAF uranium concentration was diluted by extensive rework of plutonium product solutions, but averaged ten per cent higher than the former adjusted feed concentration. Improved HA Column control has resulted because of more uniform concentration feed and no adverse process effects have been noted.

(c) Product Treatment

The resin lost from the neptunium purification unit to the waste tank last month was discarded and the unit charged with fresh resin. Miscellaneous neptunium rework solutions were combined with a small batch of virgin feed and processed through the unit. The run was unsuccessful with loss of essentially all of the neptunium to the backcycle system. The poor process performance is believed due to neptunium valence control difficulties. At month end, about two kilograms of neptunium was transferred from the in-canyon recovery system to the purification unit for the next purification run.

(d) Waste Processing

The addition of sugar to the IWF tank for ruthenium control in the high level waste concentrator continued. A special sampling and analytical program was initiated to determine changes in the amount and spectrum of fission products in the acid absorber overheads resulting from sugar additions.

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The effect of Demister water in the acid absorber acid activity was studied. A two-fold increase in acid gamma activity occurred without water flow to the Demister. Activity levels returned to normal when water flow was restored.

A new direct-drive slow speed agitator was tested for mixing performance in a system simulating triaurylamine treatment of high level waste. Excellent mixing was achieved and the unit has since been installed in the plant for additional triaurylamine extraction studies.

About 560,000 gallons of supernatant solution was transferred from TK-102-A in the 241-A Tank Farm to non-boiling storage in 241-C Farm in preparation for sludge-mixing tests in TK-103-A. Water is being added to TK-102-A to soften the sludge in the tank and serve as a hot water supply for a sludge nozzle assembly in TK-103-A. Installation of special pumps and transfer lines for the test are underway.

(e) Fission Product Recovery

The modified Oak Ridge cesium cask containing technetium recovered from the October cesium loadout reported last month was transferred to the Strontium Semisworks for elution. The anion exchange bed in the cask was washed with four bed volumes of 0.5 M sodium nitrate, 1.5 bed volumes of 0.25 M nitric acid and eluted with eight bed volumes of 6.0 M nitric acid. The eluted nitric acid stream was steam stripped and concentrated to a volume of 25 gallons and shipped to Hanford Laboratories. The cask contained slightly in excess of one kilogram of technetium.

A coil leak occurred in a B-Plant vessel used for storing concentrated cerium-rare earth oxides. The tank was a reclaimed vessel from U-Plant and the leak was noted three days after it was placed in service. The leak was detected promptly by the cooling water header scintillation counter and the tank contents transferred to another vessel. Contaminated water was segregated in the retention basin and later routed to a litch that was subsequently back-filled. Pond water radiation levels were as high as 2 R/hr at 2 feet above the liquid. Approximately 30 curies of cerium-144 and 0.05 curies of strontium-90 were contained in the contaminated water.

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2. Redox Process Engineering

a. Fire in the 233-S Facility

A fire occurred in the 233-S Facility on November 6, 1963. The fire was restricted to the viewing room and process area in the vicinity of the L-18 Plutonium Anion Exchanger. Plastic hood panels and electrical wiring including lights were extensively damaged in the north end of the viewing room. The four-inch pipe which makes up the contactor loop does not appear to be damaged. However, the gaskets at the flanges leak; the pumps, valves, and instruments are charred; and all of the electrical wiring to the equipment in the contactor enclosure is shorted out. None of the process equipment other than the L-18 Contactor appears to be damaged. While the viewing room, stairwells, and load-out room were grossly contaminated, the contamination was confined to the 233-S Facility and immediate vicinity.

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3. Plutonium Process Engineering Operation

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a. Button Line

The sintered stainless steel calciner blow-back filter elements that were found destroyed by corrosion after two months of service were replaced with the same kind of filter elements. These new elements were inspected twice this month at approximately 2-week intervals. No corrosion was detected on either inspection. Inspections will continue at the same frequency.

The difference between the physical inventory and the book inventory at the end of October was approximately 33 kilograms of plutonium. In order to establish that this amount of plutonium had not accumulated in the button line and created a critical mass problem, the button line was shut down for a thorough inspection of the system, both visually and with the aid of gamma and neutron detection instruments. In addition, another physical inventory was performed. No unusual accumulation was found, nor did the inventory indicate any change. Operation was resumed, but the discrepancy still could not be explained at month's end.

b. PuO₂ Teflon-Lined Dissolvers

The prototype teflon-lined plutonium oxide dissolvers were provided with Kel-F coated spool pieces connecting the dissolver with a new Kel-F coated reflux condenser. Since the installation of these spool pieces, the dissolution rate in these dissolvers has decreased to 1/3 the maximum rate (from 300 to 100 g/hr) obtained previously. It appears that considerable vapor escaped into the hood through the badly corroded stainless steel spool pieces, which changed the solution composition and resulted in the higher dissolution rates.

c. Plutonium Reclamation Facility - Project CAC-880

HW-79534, "Plutonium Reclamation Facility Operating Specification", was issued during the month. The document defines process conditions and limits for initial operation of the Plutonium Reclamation Facility.

HW-79586, "Start-Up Flowsheets for the Plutonium Reclamation Facility (CAC-880) - Edition I", also issued this month, provides the basis for "cold" and initial "hot" run plans. Initial cold runs will simulate flowsheets given in this document.

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4. Separations Chemistry Laboratory

a. Purex Process Improvement

Several types of additives including mono and di-ketones, mono and di-oximes, and dibasic acids have been evaluated in the laboratory in a search for materials to increase decontamination of ZrNb across both the HA and the 2D columns. Although many of the compounds tested show some improvement, oxalic acid addition was best. The ZrNb DF was improved by a factor of 3 across the HA column when the HAF was 9×10^{-4} in oxalate and the 2D ZrNb DF was improved by a factor of 2 when the ZDS was made 9×10^{-4} in oxalate. Increases in Ru DF during these tests were not considered to be significant. The side effects of adding oxalic acid to the process streams are being quantitatively evaluated.

b. Redox Process Improvement

Mesityl oxide was evaluated in the laboratory as a possible solvent in a thorium recovery process. Multiple batch counter current extraction runs under acid deficient conditions gave thorium distribution ratios in mesityl oxide that were a factor of 4 higher than those achieved using hexone. However, the ruthenium arithmetic decontamination factor was three compared to nine for hexone. Decontamination from ZrNb and rare earths was only slightly lower in the mesityl oxide system.

A gas chromatographic method was developed for determining the presence of hexone in nitric acid and plutonium process solutions down to 0.005 v/o. The method has been applied to several Redox samples, some of which are representative of streams at the time of the 233 incident. In no case was a significant amount of hexone detected in the feed to the 233 Bldg. The method is outlined in HW-79703.

c. Fission Product Process Improvement

Laboratory studies on methods of removing lead from the rare earths and sulfate crudes led to a recommendation that the crude solution be first precipitated as the hydroxide which removes 88% of the lead from the product stream. The rare earths and strontium can then be dissolved in a reasonable volume of nitric acid, and oxalate precipitation to separate the rare earths and strontium can proceed normally. Under the old method the early oxalate precipitation resulted in a precipitate loaded with lead which responded poorly to caustic metathesis with the result that dissolution and hydroxide precipitation of the rare earths was still required. The recommended method had a measured rare earths waste loss of only 0.3%.

The effect of peroxide addition on the pH during peroxy-acetate precipitation was also measured and found to have only a slight downward effect once the pH has been adjusted to the desired 4.1. Therefore, it would be desirable to work on the high side during pH adjustment.


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The extraction and stripping characteristics of semiworks solvent were evaluated during the month and found to be comparable to fresh solvent. The used solvent analyzed 0.16 curies/l ZrNb⁹⁵, 0.19 curies/l CePr¹⁴⁴, and 0.51 curies/l Y⁹¹.

d. Laboratory Improvement

A spectrophotometric method has been developed to detect traces of tri lauryl amine in Purex solvent. The method utilizes the high molar absorptivity of TLA at the 259 mμ maxima. TBP interference is removed by the use of a blank and nitrate interference is removed by carbonate washing. The lower detection limit was found to be 0.035 v/o.

The X-ray emission spectrometer was calibrated for iron, chromium, and nickel and applied to the determination of these elements in UO₃ powder samples after dissolution in nitric acid and TBP separation. The lower limits of detection are in the 1-2 ppm range. Work is continuing on the application of the X-ray unit to the measurement of tungsten, tantalum, and titanium in plutonium metal.

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5. Plutonium Chemistry Laboratory

a. Direct Calcination Of Plutonium Nitrate

A total of 11 runs were made in the titanium screw calciner and approximately 11 kilograms of plutonium oxide were produced. Use of a high sulfate-plutonium ratio (0.3 - 0.5) produced a readily-chlorinated oxide. Visual inspection of the calciner and chemical analysis of the oxide produced failed to indicate any substantial corrosion.

A series of runs was made at various sulfate - Pu ratios with all other conditions being held as constant as possible. Chlorination results indicate that a readily-chlorinated oxide is produced with high sulfate ratios.

TABLE I

CONVERSION VERSUS SULFATE RATIO

<u>SO₄⁼/Pu</u>	<u>% Oxide Chlorinated</u>
0.3	76
0.04	80
0.07	84
0.30*	96
0.50	90
0.50*	97

(*) High screw speed to prevent sticking, resulting in lower residence time.

Work is now under way to determine more accurately the effect of SO₄⁼/Pu and residence time on reactivity.

Thermobalance studies are now under way in the Separations Chemistry Laboratory to measure reactivity. Initial results on duplicate samples indicate a reliability of about ± 3 percent for the instrument. Results to date check very closely with results obtained in the batch chlorination work.

The calciner operated satisfactorily during the month with only minor operating problems. Some leakage of solution was encountered through certain gaskets, but this should be eliminated when the gaskets are replaced. Wear on the bearings is becoming excessive and the bearings probably will have to be replaced in the near future. To date there has been no indication of excessive corrosion of the calciner. Analysis of the oxide produced shows only nominal amounts of titanium in the product. No corrosion was detected by visual observations after each calcination run.

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b. Chlorination Of Plutonium Oxide

Work with a stirred-bed batch reactor was continued and chlorination of direct calcined oxide was begun on a large scale.

The stirred-bed batch reactor consists of a filter funnel to which a top has been attached and fitted with a stirrer. The gas comes up through the frit and reacts with the bed of agitated oxide powder. About 500 g feed PuO_2 are used and reaction occurs over a 100-minute period at 450 C. Only a slight excess of phosgene (or $COCl_2$) is used.

The work was started with a low sulfate ratio (0.05) based on earlier results. Various changes were made in the chlorination parameters (bed temperature, rate of chlorination, length of chlorination, and chlorinating agents) without improving the percent conversion appreciably from the 70 - 80 percent range.

After observing very little difference with the low-level sulfate, the higher ratio 0.5 was tried. This made a marked improvement as shown by conversions of 90 and 91 percent. Two runs were then made to see if the sulfate requirements could be reduced to 0.3 or 0.15. At present it appears as though 0.3 mol ratio is about right.

Tap densities for some typical oxide feeds and product chlorides were determined and are shown in Table II.

TABLE II
EFFECT OF SULFATE IN OXIDE FEED ON CHLORIDE
DENSITY AND CHLORINATION EFFICIENCY

<u>Sulfate/Pu</u>	<u>Tap Densities</u>		<u>% Conversion</u>
	<u>Oxide</u>	<u>Chloride</u>	
0.05	3.9	3.9	76
0.15	2.3	2.8	92
0.3	1.7	2.9	96
0.5		2.3	97

Sulfate removal has been essentially 100 percent except for one run. In this case about 20 percent of the original sulfur remained in the chloride. This was attributed to a low bed temperature (440 C) during chlorination.

The product chloride from the above runs is a granular free-flowing powder.

On the basis of the limited number of runs that were made, the following conclusions are drawn:

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- (1) A mol ratio of sulfate-to-plutonium of 0.3 with a minimum residence time in the calciner is required for adequate chlorination.
- (2) Bulk density is the criterion that determines percent total conversion.
- (3) Bed temperature in the chlorinator in excess of 450 C and less than 500 C is required for adequate sulfate removal.
- (4) The reaction is essentially instantaneous; very high rates should be obtainable in the proper continuous equipment.

c. Americium Recovery

Distribution measurements have been made to aid in defining the americium recovery flow sheet.

Stripping of americium from 30 % dibutyl butyl phosphonate - 70 % CCl_4 with water results in an K_D of 0.08. The equilibrium aqueous phase has an acidity of 0.6 M H^+ at an L/V of one. Impurities in the strip solution in ppm will approximate 115 Ca, 130 Fe, and 120 Mg, relative to the solution. These, together with plutonium and aluminum, are the principal impurities to be separated.

Contact time variations from one-half to five minutes had no effect on extraction or stripping coefficients. Likewise, the use of vacuum-distilled DBBP vs. the commercial product resulted in no differences.

A steady-state extraction stripping flow sheet has been demonstrated. With one stage of extraction and one of stripping with equal volumes, over 95 percent recovery was demonstrated.

The use of 50 % DBBP was briefly tested. Lower than expected distributions for americium were observed in extraction stages, apparently due to the higher tendency of this solvent to complex with nitric acid.

J. S. Orank

Manager
Research and Engineering

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II. ACHIEVEMENTS (Continued)

H. FINANCIAL OPERATION

I. Production Cost Accounting

The Company recently determined that a substantial over-accrual of pension costs currently exists and has passed a pro rata share of this credit to HAPO of which approximately \$500,000 is applicable to CPD. This amount is being credited to cost over the balance of FY 1964. The midyear budget is being reworked to reflect this change and recent organization changes. (The former Power and Crafts Operation was dissolved on December 1, 1963 with most of its functions being transferred to Facilities Engineering Operation.)

Following the 23-B incident, coding and instructions were issued for segregation of these costs from normal production. Informal approval was obtained from AEC for use of a shutdown account in which to accumulate costs of the incident.

During the month, the AEC advised HAPO of substantial cuts in Isotope Inventory balances allocated as of June 30, 1964 and June 30, 1965. Also, a reduction of \$2,000,000 was made in the Sr-90 shipping schedules for these two periods.

Special requests handled during the month included:

Services provided APED by R. E. Tomlinson and E. O. Swain.

12 grams Pu nitrate sent to APED.

Preparation of blueprints for Bechtel Corporation.

Plutonium scrap reclamation activity for other sites during the month included receipt of 2546 grams from Hanford Laboratories (Project Whitney).

CPD's investment in inventories as October 31, 1963, compared with budgeted amounts, is as follows:

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(in thousands)	Balance 10-31-63	Control Allocation	Surplus (Deficit)
<u>Inventories</u>			
Essential Materials	\$ 911	\$ 815	\$(96)
Spare Parts & Standby	1 621	1 615	(6)
Special Materials	85	103	18
Yttrium	<u>153</u>	<u>150</u>	<u>(3)</u>
Gross Inventories	<u>2 770</u>	<u>2 683</u>	<u>(87)</u>
<u>Reserves</u>			
Essential Materials	42	59	(17)
Spare Parts & Standby	562	404	158
Yttrium	<u>153</u>	<u>150</u>	<u>3</u>
Total Reserves	<u>757</u>	<u>613</u>	<u>144</u>
<u>Net Investment</u>	<u>\$2 013</u>	<u>\$2 070</u>	<u>\$ 57</u>

2. General Accounting

As of October 31, 1963, seventeen active projects had incurred costs of \$3,078,036 against authorized funds of \$4,096,100. Outstanding commitments totaled \$182,852.

During November one work authority and one directive was received from the AEC: Directive No. EQT-007, Work Authority No. 1, Project CAC-121, Americium Recovery Facility-Z Plant, interim authorization of \$44,000; Directive No. HW-553, Project No. CGC-122, Exhaust Ventilation Facility - 233-S Building, interim authorization of \$14,400.

During the month five appropriation requests were processed authorizing expenditures of \$17,000.

3. Business Programs

A cost distribution model was developed and issued during the month which more adequately displays the financial impact of various types feed on chemical separations. This was prepared in collaboration with representatives of Contract Accounting and was reviewed by them with other financial organizations before issuance.

Manager - Finance

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III. PERSONNEL ACTIVITIES

A. FORCE SUMMARY

<u>Operation</u>	<u>Monthly Salaried</u>		<u>Weekly Salaried</u>		<u>Total</u>	
	<u>10-27-63</u>	<u>11-30-63</u>	<u>10-27-63</u>	<u>11-30-63</u>	<u>10-27-63</u>	<u>11-30-63</u>
General Manager's Group	10	9	2	2	12	11
Financial	13	13	13	13	26	26
Research & Engineering	67	69	30	30	97	99
Facilities Engineering	71	71	23	23	94	94
Power & Crafts Operation	36	36	230	231	266	267
Production	6	6	4	4	10	10
Redox	63	62	222	223	285	285
Purex	70	71	258	262	328	333
Weapons Manufacturing	<u>55</u>	<u>56</u>	<u>227</u>	<u>239</u>	<u>282</u>	<u>295</u>
Total	<u>391</u>	<u>393</u>	<u>1009</u>	<u>1027</u>	<u>1400</u>	<u>1420</u>

B. PERSONNEL CHANGES

W. M. Harty, formerly Manager, Process Design, N-Reactor Department, was appointed to the position Manager, Purex Operation, effective November 15.

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3. TRIPS

<u>Visitor</u>	<u>To</u>	<u>Nature of Discussion</u>
<u>To Other G.E. Components</u>		
P. H. Reinker	Palo Alto, California	Business Review. (11/6 - 7/63)
J. J. Shefcik C. W. Smith	Shelbyville, Indiana	Heaters for in-tank solidification systems. (11/12/63)
J. W. Kolb	Evandale, Ohio	Electron beam welding. (11/13 - 14/63)
P. H. Reinker	San Jose, California	Business Review. (11/13 - 14/63)
<u>To AEC and Other AEC Contractors</u>		
A. E. Barber	Rocky Flats Plant Denver, Colorado	Follow-up on 74 shipment (11/7/63)
R. Y. Lyon	Mallinckrodt Chemical Works Weldon Springs Plant St. Charles, Missouri	Uranium and thorium de- nitration tests. (11/15 - 19/63)
S. J. Beard W. L. Godfrey W. C. Schmidt P. W. Smith S. G. Smolen	Phillips Petroleum Co. Idaho Falls, Idaho	Fuel processing and tour ICPP and WPF facilities. (11/18/63)
B. M. Dobbs	Albuquerque, New Mexico	03 Budget Program (11/18 - 20/63)
H. A. Clark	U. S. AEC New York Operations Office New York, New York	Filter testing workshop. (11/18 - 22/63)
<u>To General Industry</u>		
J. J. Shefcik C. W. Smith	Watlow Electric Co. St. Louis, Missouri	Heaters for in-tank solidification systems. (11/11/63)
	W. L. Wiegand Co. Pittsburgh, Pennsylvania	" " (11/13/63)
	Vulcan Electric Co. Danvers, Massachusetts	" " (11/14/63)

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C. TRIPS (Continued)

<u>Visitor</u>	<u>To</u>	<u>Nature of Discussion</u>
<u>To General Industry</u> (Continued)		
A. J. Waligura	Union Carbide Nuclear Div. Sterling Forest, N. Y.	Tour radiochemical lab and Isotope service facilities. (11/19/63)
	Isotopes, Inc. Westwood, New Jersey	Tour high and low radio- chemical labs and counting system facilities. (11/21/63)
<u>To Conventions and General Meetings</u>		
E. M. Johnston	Philadelphia, Pennsylvania	Attend American Society of Mechanical Engineers meeting. (11/14/63)
A. J. Waligura	New York, New York	Attend ANS meeting and Hot Lab and Equipment Conference. (11/18 - 21/63)

D. VISITORS

<u>Visitor</u>	<u>From</u>	<u>Nature of Discussion</u>
<u>From Other G.E. Components</u>		
Howard Zellinger	St. Louis, Missouri	Cooling Vest for Decontami- nation Work. (11/12 - 14/63)
Dr. J. V. Grimaldi	New York, New York	Accident prevention. (11/20 - 22/63)
<u>From AEC and Other AEC Contractors</u>		
W. P. Bebbington	E. I. duPont de Nemours Savannah River Plant Aiken, South Carolina	Fission products and waste management. (11/7/63)
Dr. M. T. Kelley L. T. Corbin	Analytical Division Oak Ridge National Lab. Oak Ridge, Tennessee	To discuss high-level cell fission product analysis and measurements. (11/8/63)
T. R. Workinger	Division of Production US-AEC Headquarters Washington, D. C.	Purex Tour. (11/19/63)

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D. VISITORS (Continued)

<u>Visitor</u>	<u>From</u>	<u>Nature of Discussion</u>
<u>From AEC and Other AEC Contractors (Continued)</u>		
Dale Singleton	E. I. duPont de Nemours Savannah River Plant Aiken, South Carolina	Repair of "D" Cell - Redox. (11/21/63)
J. Daniledes W. H. Johnson	Lawrence Radiation Lab. Livermore, California	Plutonium specifications. (11/21/63)
C. W. Barrett	Rocky Flats Plant Denver, Colorado	Plutonium specifications. (11/21/63)
<u>From General Industry</u>		
T. F. D'Muhala	General Dynamics Groton, Connecticut	Decontamination. (11/5 - 6/63)
Norman Hart	Columbia-Geneva Steel Seattle, Washington	Tube bundle fabrication and performance. (11/5/63)
M. M. Braidech	National Bureau of Fire Underwriters New York, New York	Investigate 233-S incident. (11/11 - 12/63)
S. W. Silverman	Boeing Company Seattle, Washington	Discuss waste management, fission products and encapsulation. (11/19/63)
R. Wischow	Martin-Marietta Co. Nuclear Division Baltimore, Maryland	Fission product recovery. (11/22/63)
R. L. Meyer	Betz Laboratories Seattle, Washington	Filming amine; inert gas system. (11/26/63)
<u>From Other Government Agencies</u>		
Major J. E. Wassiac Captain R. J. Thomas	Department of Defense	Tour facilities. (11/19/63)

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IV. SAFETY AND SECURITY

<u>Operation</u>	<u>General</u>	<u>F&O</u>	<u>Finance</u>	<u>WMO</u>	<u>P&CO</u>	<u>Purex</u>	<u>Redox</u>	<u>Prod.</u>	<u>R&E</u>	<u>Total</u>	<u>YTD</u>
Dis. Injuries										-	6
Ser. Accidents										-	3
Med. Treat. Inj.		2		4	12	10	7			35	451
Rad. Occur.				3		5	1			9	131
Contam. Wds.										-	20
Pu Depositions										-	20
Fires				2*			1**			3	9
Sec. Viol.				1						1	21

- * - November 4, Fire 62-8 - Rags and resin in hood in Room 227 - Damage, \$1686.40.
- November 10, Fire 62-9 - Material in chopper hood in 232-Z Bldg. - Damage, \$10.00.

** - November 6, Fire 62-7 - 233-S Bldg., believed to be started from causes external to the process. Preliminary estimate of damage, \$310,000.00.

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V. REPORTS

A. PREPARED AND ISSUED

- HW-79147, Unclassified, "Critical Mass Control Specification - 232-Z Contaminated Waste Recovery Facility", dated November 8, 1963, by R. J. Sloat.
- HW-79219 RD, Secret, "Product Specifications - Hanford Uranium Trioxide (From Depleted E-Metal) for Off-Site Shipment", dated November 6, 1963, by R. J. Sloat.
- HW-79241, Official Use Only, "Increased Processing Flexibility - Purex, Project CGC-124", dated November 14, 1963, by L. W. Finch.
- HW-79276, Unclassified, "Recommended Improvements to the Dry Air System - 234-5 Building", dated October 23, 1963, by D. D. Wodrich.
- HW-79363, Secret AWD, "Model 74-C Shipping Problem Status", dated October 24, 1963, by A. E. Barber.
- HW-79369, Unclassified, "Design Scope, Americium Recovery Facility", dated October 31, 1963, by D. E. Braden and R. A. Ciccarelli.
- HW-79384, Secret AWD, "Quarterly Summary Report and Associated Data for 234-5 Building Final Product Shipped During the Third Quarter of Calendar Year 1963", dated October 28, 1963, by A. E. Smith.
- HW-79385, Secret AWD, "Statistical Statements and Associated Data for 234-5 Building Final Product Shipped During the Third Quarter of Calendar Year 1963", dated October 28, 1963, by A. E. Smith.
- HW-79428, Confidential AWD, "FPT-63-27, Dimensional Stability, Model 1807 Shape Castings", dated October 18, 1963, by C. M. Walker.
- HW-79470 RD, Secret, "Purex Plant Production Schedule - November 1963", dated November 4, 1963, by D. McDonald.
- HW-79471 RD, Secret, "Redox Plant Production Schedule - November 1963", dated November 4, 1963, by D. McDonald.
- HW-79472 RD, Secret, "UO₂ Plant Production Schedule - November 1963", dated November 4, 1963, by D. McDonald.
- HW-79473 RD, Secret, "234-5 Plant Production Schedule - November 1963", dated November 4, 1963, by D. McDonald.

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- HW-79496, Secret AWD, "Progress Report, Weapons Process Engineering, October, 1963", dated November 5, 1963, compiled by Members of the Weapons Process Engineering Operation, Research and Engineering.
- HW-79534, Confidential, "Plutonium Reclamation Facility Operating Specifications", dated November 11, 1963, by L. E. Bruns
- HW-79577, Secret AWD, "Fabrication Process Test Evaluation, Shipping HI-74-SL", dated November 8, 1963, by A. E. Barber.
- HW-79578, Unclassified, "Critical Mass Control Specification - The Storage and Handling of Plutonium Solutions in the PR Room (Room 19) of the 236-Z Building", dated November 15, 1963, by R. J. Sloat.
- HW-79586, Confidential, "Start-Up Flowsheets for the Plutonium Reclamation Facility (CAC-880) Edition I", dated November 18, 1963, by L. E. Bruns.
- HW-79645, Secret, "Process Design Scope Criteria, Purex Multi-purpose Dissolvers", dated October 15, 1963, by R. D. Ehrlich and J. R. LaRiviere.
- HW-79703, Unclassified, "Quantitative Determination of Hexone in Aqueous Solutions by Gas Chromatography", dated November 25, 1963, by C. A. Colvin and R. J. Sorenson.
- HW-79714, Official Use Only, "Redox Operation Planning - Task Force Interim Report", dated November 20, 1963, by J. B. Fecht, M. K. Harmon, B. F. Judson and J. H. Warren.
- HW-79733, Unclassified, "Critical Mass Control Specification - PR and SN Can Transportation", dated November 26, 1963, by R. J. Sloat.
- HW-79745, Secret AWD, "A-Line Product Characteristics, October 1963", dated November 26, 1963, by A. E. Smith.

B. PREPARED FOR SIGNATURE AND ISSUANCE

- HW-79479, Secret, "Production - October 1963", dated November 1, 1963, by W. E. Johnson.
- HW-79605, Secret, "Production of Zirconium-Niobium-95 and Scandium-46 at Hanford - Addendum to HW-79404", dated November 15, 1963, by W. E. Johnson.

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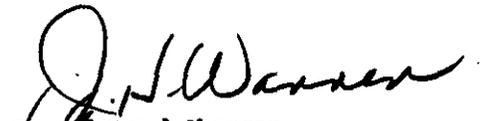
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VI. PATENT SUMMARY

All persons engaged in work that might reasonably be expected to result in inventions or discoveries advise that, to the best of their knowledge and belief, no inventions or discoveries were made in the course of their work during the period covered by this report, except as listed below. Such persons further advise that, for the period therein covered by this report, notebook records, if any, kept in the course of their work have been examined for possible inventions or discoveries.

<u>INVENTOR</u>	<u>TITLE</u>
R. W. Lambert	The Use of Sugar for Suppressing Ruthenium Volatilization in Nitric Acid Concentrators
J. S. Ruckingham	The Destruction of Citric Acid with Hydrogen Peroxide
A. W. Hildebrandt	Automatic Compression Clamp For Holding Pieces For Presentation For Gaging
W. P. Ingalls (CFD) and	A Design For a Neutron Shielded (Class I) Shipping Container For Unirradiated Fissile Materials.
C. L. Brown (HLO)	(The neutron shield precludes a nuclear chain reaction between the fissile materials in two or more adjacent containers by preventing the exchange of neutrons between containers..)


General Manager
Chemical Processing Department


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