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MONTHLY PROGRESS REPORT

ON

FUEL ELEMENT DISSOLUTION STUDIES

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May & June, 1958

CIRCULATING COPY RECEIVED 300 AREA AUG13 1958 RETURN TO TECHNICAL INFORMATION FILE

Nuclear Engineering Department Brookhaven National Laboratory

Upton, New York

AEC Budget No. 4301

Reviewed and Approved for Public Release by the NSAT Ilmed Brown PNNL ADD

Brookhaven National Laboratory

MEMCRANDUM

DATE: July 17, 1958

TC:

C. Will

FOM:

L. P. Hatch, J. J. Peilly, W. H. Regan

SUBJECT: Fuel Element Dissolution

Studies During May and Jume

Fluid Bed Hydrochlorination

A sample of a non-irradiated, sircaloy clad 302 fuel element has been decladded almost completely in fluidized bed hydrochlorinator. No filters of any kind were used in this experiment for de-entraining UO2 particles, since it was surmized that the UO2 core would remain relative intact. The total sample weight was 47.9 gms and it had a surface area of 44 cm2. The average reaction rate based upon the original surface area was 600 mgms cm2 hr. The final sample weight was 12.8 gms. The recovered plate of 302 appeared to be intact with no apparent dimensional change, although scaling was evident. The surface area of the recovered 302 was 25 cm2 out which about 3 cm2 appeared to be covered by a thin layer of unreacted cladding material. A sample of ZrCl4 product containing 10.3 gms of Zr contained 0.0253 gms of uranium; projecting this figure for the total amount of Zr rescted and assuming the product is homogeneous gives a value of approximately 1.2% of the total U present carried over with the product. A total of 2.06 gms of uranium was recovered from the reactor sand by leaching with HNO3. There was a small amount of insoluble black material recovered from the sand which was shown by spectrographic analysis to consist principally of silica.

As previously reported a sample of a zircalcy clad, Uranium-Zirconium alloy fuel element was hydrochlorinated in a system which included a sand filter for de-entraining Uranium particulate matter carried over with the ZrCl/. Analytical results have shown the amount of U which passed through the sand filter is at least as much as that which was recovered from the sand by leaching with nitric acid. Actual amounts were 3.17 mgms of uranium recovered from the sand filter while the gas scrubbing solution contained 3.4 mgms of uranium. Future experiments with allow type fuel elements will be concerned with determining the total uranium material balance.

Several preliminary experiments involving heat transfer between a heat source (500 watt strip heater) and a fluidized sand bed expanded about 20% over its settled volume gave values of 50-60 BTU/hr x ft² x f for heat transfer coefficients.

Equipment has been designed and is now being built for investigating the possibility of using an inert fluidized bed for heat removal in the highly exothermic reaction between metallic uranium and fluorine.

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