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COMMERCIAL

Le présent document a été reproduit avec l'autorisation de CANCOPY. The Sixth Core revenimely lating solution, ulterieuro

Experimental Results

'by

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including work by

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Abstract

The results obtained during experiments on the sixth SGHW core in DIMPLE are presented in this report in the form of raw data. No attempt to apply theoretical corrections to allow comparison with theory has been made, since this will be included in a more general report covering the whole series of SGEW experiments carried out at Winfrith.

The report is intended to provide a record of all important results obtained on this core. For the sake of brevity it refers frequent)y to the reports on the first five cores (1), (2), (3), and is only a complete record when used in conjunction with these references. Less essential details are recorded in the original experimental log books.

A.E.E., Vinfrith.

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1. Introduction

This report describes the experiments carried out on the sixth of a series of cores built in the zero energy reactor DIMPLE in support of the general investigation of the reactor pbysios of Steam Generating Heavy Water (SOHW) reactors which has been undertaken by the Water Reactor Pbysios Division at A.E.E. Winfrith. The experiments in the earlier cores are reported in references (1) to (3) inclusive, and those in later cores will be described in further reports of this series. A companion series of papers covering the DIMPLE experiments and the work undertaken in subcritical assemblies will be issued to compare the experimental data with theoretical predictions based on the methods of calculation in current use at Winfrith.

Reference (1) , being the first of the series, described most of the experimental techniques in some detail, and reference (3) described the techniques for two additional experiments. For the sake of brevity, no such descriptions have been repeated in this report.

The previous core (3) consisted of a central region of 24 channels with mixed enrichment fuel oooled by light water, surrounded by 16 channels of fuel (identical to the second core (2)) acting as a driver region, the whole core then being surrounded by a D₂0 reflector. In this present core the only difference was in the coolant used in the central 24 channels, which was changed to $70.3 \pm 0.1\%$ D₂O and 29.7% H₂O by weight. This mixture has a value of $\boldsymbol{\xi} \boldsymbol{\Sigma}$ which corresponds quite closely to that of light water with a density in the region of 0.4 g m cm $^{-3}$, which is typical of the average density within a boiling channel of an SGHW power reactor. The central region under study was "driven" by an outer region of sixteen pressure tubes of the identical core design reported in the second report (2) of this series.

2. Description of DIMPLE as oonstruoted-for this experiment

Except for the amount and type of fuel, and the number of fuel channels, the reactor was precisely as described in reference (1).

The core in the present report differs from the third core (3) only in the ooolant used in the centre twenty four channels.

Figure 1 gives a plan view of the reactor tank and Figure 2 shows details of an inner zone lattice cell, with axial distances relative to search tubes and safety rods. Figure 3 is a detailed sketch of the inner zone fuel element and Figure 4. is a section through the reactor(a) in detail and (b) simplified for purposes of calculation (see Appendix I].

3. Approach to critical

At the completion of work on the third core, the fuel from the central twenty four channels was unloaded and dried, the pressure tubes were dried, and the fuel was then replaced. The H_2O/D_2O mixture (approx. 9.9. Kgm per tube) was then added, tube by tube, commencing at the centre. Throughout these operations a 10 ourie Po-Be source was installed near the centre of the tank bottom and the flux as indicated by the three installed BF3 chambers was recorded .at frequent intervals. There was little change at any time, as expected

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from experience with earlier cores, indicating that the multiplication of such cores in the absence of the main D₂O moderator was very small.

Taking advantage of the oonolusions reached in the earlier experiments (see in particular reference (1)) experimental BF₃ chambers were situated outside the fuelled part of the core and as remote from the source as was possible; the standard (1) approach to critical procedure was followed. The critical height was measured to be 151.00 cm (from the tank bottom) with all instruments removed from the core.

4. Reactivity measurements

All reactivity measurements have been normalised to a scale oalibrated. by the steady diverging period of the super-critical reactor. This was calculated by the method of reference *(1)* Appendix II.

In principle the reaotivity-doubling time relation for a multi-zone reactor may be calculated using a statistical weighting procedure. Since, however, the difference between the scales used in the first (air-cooled) and second (water-oooled) oores was everywhere less that 1% of the reactivity, and between the first and seventh (air-oooied 1.9 Co) cores ya everywhere less than 3% of the reactivity, the scale for the second core (2) was adopted for the fifth and sixth cores. Any error incurred is likely to be less than $+$ 1.5% of the reactivity measured. The table of values relating doubling time to reactivity is given in reference (2).

The method for measuring the steady doubling time of the divergent reactor was as described in reference (1). The sub-critical multiplication method adopted for measuring reactivities was also as described in reference (1), with the method for normalizing to the supercritioal soale as \cdot described in reference (2) .

Table I below summarises the reactivity changes measured by supercritical methods. The reaotivity worth of items has been computed by the change of critical height multiplied by the arithmetic mean of the values of $\frac{\partial P}{\partial h}$ at the two extreme moderator heights. All the values of $\frac{\partial P}{\partial h}$ are plotted against the arithmetic mean of the oritical and divergent heights in Figure 5, and the values of $^{\circ}$ P/3h shown in Table I are read off the dotted curve. Note that this curve has been drawn through the control rod results only, and, as in the earlier oores, the values of $^{op}/\delta h$ measured with fuel clusters missing lie well above it.

The errors quoted on the critical heights were deduced from repeated measurements on the BERRIOT MK I depth probe and represent the spread of. observations under steady conditions. The error on the absolute height was of order 0.05 cm. The errors quoted on the reactivity changes are almost entirely due to the random error on the measurement of doubling time, which is about 1%. An additional systematic error of at least $+$ 5%, due to uncertainties in delayed neutron data and knowledge of Peff (including photo-neutrons) must be taken into account before comparing these results with theory.

From Figure 5, the value of $\partial P/\partial h$ at the olean critical height was deduced to be $0.115 \pm .001\%$ om-1. The value of $\frac{\partial P}{\partial h}$ measured with a near central pressure tube removed was about $5 + 1.5$ percent higher than that measured at approximately the same critioal height but with the fine control rod inserted. This follows the same trend as in the earlier cores.

Table I

The effect was large in the first and second oore, but barely significant in the fifth gore, and it would appear to be larger the greater the magnitude of P/6h. Further theoretical investigation of this effeot is clearly necessary.

Table II summarises the measurements of negative reactivities using the sub-oritical multiplication technique. Figure 6 shows the counter positions during these measurements.

Table II

Reactivities deduced by sub-critical multiplication measurements

Examination of Table II and Figure 6 shows that the variation in apparent reactivity follows the same trends as were observed in all earlier experiments (see, in particular, reference (1)). The sub-oritioal estimate of fine control rod reactivity (O.655%) was in good agreement with the critical estimate (0.64%) of Table I. Due to the obvious limitations of the method the true reaotivities and their associated errors are indeterminable; as previously, the most pessimistic individual results were used to satisfy the safety criteria.

5. Macroscopic Reaotion Rate Distributions

The measurements were made using U235 and Pu239 fission chambers in precisely the manner as described in reference (I). The radial measurements were made with the active centre of the chambers \sim 75 cm above the tank bottom $(\sim 55$ om above bottom of fuel) - see Figure 2. Appendix II gives the results in detail and sections 5.1 and 5.2 summarise the spectrum and radial buckling results respectively. The axial measurements were made in the central search tube (K11) and an adjacent one (K13); the results are detailed in Appendix III 'and summarized in section *5.3.* Consistent with the earlier reports (see reference *(1)* in particular) we have allowed for a 0.1% counter drift error on each counter (0.14% on counts relative to monitor counts) in addition 'to the Poissonian variation of the number of counts recorded. All measurements have a counter dead time correction of 1.5 \pm .5 µsec applied, although count rates were such that the correction was less than 0.5% on each count and in general less than 0.1% on a ratio of counts'

5.1 Speotrum results

The plutonium 239 to uranium 238 reaotion rate ratio, uranium 235 cadmium ratio (R_5) and plutonium 239 cadmium ratio (R_9) were measured at most available radii inside D₂0 - filled search tubes. The results are summarised in Table 1 of Appendix II and plotted against radius in Figure 7. All three parameters were found to be constant, within their estimated errors of $+$.4% out to a radius of 48.3 Om (two lattice pitches), and the results were a7eraged, giving, for the centre core region:-

 $R_5 = 31.4 \pm .1$ $R_9 = 41.5 \pm .5$ Pu/U DIMPLE Pu/U NESTOR THERMAL COLUMN 1.152 + .004

This latter result was in good agreement with the value of $1.145 +$.046 obtained using Pu/Al and U235/Al foils (see section 9).

The results quoted above are uncorrected for attenuation of flux by the chamber wall and active coating, or the effect of displacing D₂₀ by the fission chamber.

5.2 Radial component of buckling β^2

A summary of all results obtained in the radial scan is presented in Table 2 of Appendix II. As in the earlier cores a statistical analysis was carried out to determine the region of constant spectrum and to check the symmetry of the core.

Examination of Table 2 of Appendix II shows a marked symmetry between the flux ratios in K09 (0.966 \pm 0.001) and K13 (0.974 \pm .002), which are both at radius 24.13 cm from the core centre. Section *5.4* describes a further investigation into this asymmetry. The four results at a larger radius of 48.26 cm i.e. K07 (0.860 + .002), K15 $(0.854 + .001)$, G11 $(0.856 + .001)$ and 011 $(0.856 + .002)$ were much more symmetrical

Table 3 of Appendix II gives the results of the computation of radial buckling. The error on β was calculated in precisely the manner of the earlier **reports** (1), (2), (3). Results are summarised in Table III below, KO9 and K13 being treated separately.

Table III

Estimates of B

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In Table 4 of Appendix II weighted mean values of β , together with the "goodness of fit" parameter x^2 , are tabulated for various combinations of the above results. Examination of Table III above indicates that K13 is in error, and some doubt of the validity of G13 exists since this might be outside the region of constant spectrum illustrated in Figure 7. The final value of β was calculated, neglecting these two positions, to be $1.596 + .019$ m⁻¹, the standard deviation quoted being scoaled up to be consistent with the fit of the two values used. On this basis the radial buckling was $2.55 \pm .06$ m⁻², but in view of the asymmetries present it is recommended that the error quoted should be increased by about a factor of two, giving

 β^2 = 2.55 + .12 m⁻²

It is interesting to note that, whereas in the $E_20 -$ cooled version of the oentre zone (3) the speotrum was apparently constant out to a radius of 72.4 om, this was not true of this core. This gives further justification to the decision not to use the point at 72.4 cm in computing the radial buckling of core 5.

The radial distributions obtained with the bare and cadmium covered U235 and Pu239 chambers are plotted in Figure 8, together with the deduced J_{Ω} (β r).

5.3 Axial component of buckling

Measurements were made using a 10' long U235 fission chamber, connected to an extension scale, which was moved manually from above the top biological shield in precisely the manner used in earlier cores (1). Scans were made with oadmium covered and bare chambers in K11, and merely cadmium covered in K13.

The cadrium ratio obtained is plotted in Figure 9. As in the

previous core (3) , the ratio falls at points near to the centre plate of the 28" long clusters, but rises at the join of the clusters. Figure 3 shows quite clearly that the main difference between the centre and end plates was that the 1.8% (28" long) fuel in the centre 23 pencils was continuous through the oentre plate, whereas there was a 2.35 cm gap in the fissile material of the outer twenty pencils in this centre plate. Because of this behaviour only very few points oould be fitted to a oosino to produce an estimate of the axial buckling. These points are shown in Figure 10. Table IV below summarises the results, which were corrected where required for the effect of the cadmium covered fission chamber in the manner described in reference (1). The complete experimental scans in search tube K11 (corrected for dead time) divided by the fitted cosine are given in Figure 10. and details of all three scans are given in Appendix III.

Table IV

Results of cosine fitting to axial scans

The behaviour of the total and epi-cadmium reaction rates at the ends and centres of the clusters are quite different. Whereas the total reaction rate rises at the end of a 28" cluster, it falls at the end and rises at the centre plate. The major difference between end and centre plate geometry is that only the outer ring, of fuel has a gap in the latter whereas all fuel has a gap in the former.

One possible explanation of the observed phenomena would be that removal of moderating coolant but not fuel, (i.e. replacement of coolant by aluminium) causes the epi-cadmium neutron rlux to rise and the thermal flux to fall, whereas removal of moderating coolant and fuel causes a nett fall in epithermal flux (because fewer fast neutrons are produced) and a nett rise in thermal flux (since fewer neutrons are absorbed with no fuel present).

Alternatively one could argue that the epi-cadmium peek at the oentre plate is associated with the reduction of U238 absorptions in the outer ring (since there is a fuel gap in this ring). At the top plate removal of all fuel removes fast sources and reduces the local epi-oadmium flux, overriding the effect observed in the centre plate. Which, if either, of the above explanations is correct will only be clear after considerable theoretical investigation.

The total perturbation of the epi-cadmium reaction rate was $+2\%$ to -2% , and of the total reaction rate was $+$ $.8\%$ to -1.4% . Having corrected the cadmium covered scans in the manner recommended in reference (1) , the overall mean effective height was calculated by weighting results by the inverse of their deviations from the cosine, and was as follows

 $H = 156.0 \pm 0.6$ cm $= 2.014 \pm .008 \text{ m}^{-1}$ α = 2.014 + .008 m⁻²
 α ² = 4.06 + .03 m⁻²

Top extrapolation distance $= 8.4$ cm combined error $+ 0.6$ cm Bottom extrapolation distance = 17.0 cm

The errors in c^2 quoted above are those deduced from the consistency between one bare and two cadmium covered scans. Due to the extremely limited region used in the fit and the fact (see Figure. 8) that there is only a small axial region of constant cadmium ratio, it would be imprudent not to add a large systematic error to the random errors deduced from these measurements. In the absence of any theoretical evidence it is recommended that the value of α^2 to be compared with theory should be $4.1 \pm .3$ m⁻². (i.e. we should arbitrarily increase the error by a factor ten).

5.4 Radial asymmetry in the core

Examination of Table 2 of Appendix II indicates that a noticeable asymmetry of 0.8 *+* .21 existed between search tubes K09 and K13 situated at radii of 24.13 cm either side or the core centre. This asymmetry did not, apparently, extend beyond this region. The effect was independent of search tube or fission chamber angular orientation, or the nature of the reaction rate measured.

Just prior to dismantling the core an experiment was carried out to investigate this effect, as follows:-

- (a) Two fission chambers were placed, one in each of these search tubes, and a series of counts taken. The fission chambers were reversed and the count repeated.
- (b) The search tubes were exchanged and (a) was repeated.
- (c) The fission chambers were covered with cadmium sleeves and (a) was repeated.
- (d) The 4 pressure tubes complete with fuel surrounding K09 were exchanged for those surrounding K13.

Table V below summarises the results. The errors quoted are due to counting statistics plus $0.1%$ per count for counter drift.

State	Ratio of K13/K09					
	Fission chamber 1	<i>i</i> Fission chamber 2				
a ъ o d	1.0047 \pm .0015 1.0047 \pm .0015 $1.0046 \pm .0015$ $0.9959 \pm .0015$	1.0048 \pm .0015 $1.0090 \pm .0015$ 1.0060 \pm .0015 0.9972 $\frac{1}{2}$.0015				

Table V

Comparison of cases (c) and (d) shows that the effect of exchanging the pressure tubes and contents surrounding one search tube with those from the other reversed the asymmetry. Two independent fission chambers indicated a change in the ratio of .0087 \pm .0024 and 0.0082 \pm .0024 respectively. Since this change, in both cases, was beyond the 30 limit (with errors which were overestimated in all probability), we conclude that the asymmetry observed was associated with the fuel clusters and/or the pressure tubes surrounding the two

search tubes. The coolant in each pressure tube was analysed for D_2O content and all were found to be within the quoted experimental error of \pm 0.3%. The clusters were examined and found to be visually identical. The pressure tubes were examined for bowing but none was found. Due to shortage of time the investigation was concluded at this point with no obvious reason for the discrepancy apparent.

6. Microscopic reaction rate distributions using manganese foils

One-half inch diameter by 0.005 inch thick manganese foils were placed, in two planes 6.7 cm apart, between fuel pellets in special telescopic fuel pencils, on the outside of the pressure tube, on the inside of the calandria tube, and inside a $D₂O$ filled search tube. Full details of the equipment used for positioning and counting the foils is included in reference 3.

The DIMPLE collapsible foil machine used in the earlier cores was abandoned for this and subsequent experiments (since it was found to give poor foil positional accuracy), and thus only one position in the moderator (that in a search tube) was measured. The foils were positioned accurately in a vertical plane in the search tube by means of a 0.375 inch diameter by .039 inch walled aluminium tube; accurate height registration was given by slots in this tube at its lower end. Around the foils the tube was extensively cut away to reduce the amount of aluminium/unit length to about 20% of normal.

Three irradiations, each with two layers of foils in planes 52.4 and 59.1 cm above the bottom of the fuel, were carried out. Examination of Figure 10 shows .that the foils were in the region of constant spectrum between the oentre plate and top of the lower 28 inch cluster.

Appendix IV gives details of the irradiations in Table 1 and a summary of the results is given in Table 2. Standard deviations were computed from the run-to-run consistency in Table 2 in the manner described in reference 1, and were between I and 2% depending on the position of the foil. The results are plotted in Figure 11.

All three irradiations were carried out with the foils placed on a radius from the centre of the reactor passing through the centre of the fuel element J10 (see Figure 1). For the first two irradiations foils were placed between the centre of J10 and the core centre. In the third irradiation the pressure tube was rotated through 180 degrees and the foils were placed (still in 10) between the centre of J10 and the centre of H8. Thus the macroscopic corrections $1/Jo(\beta r)$ [r is the radius of the foil from the core centre] applied were different in this third irradiation and the first two irradiations. Examination of Table 2 of Appendix IV shows no systematic differences between these irradiations outside the experimental errors of \pm 1%. Thus it would appear that the procedure of making a macroscopic correotion is valid within the precision of these results.

7. Mioroscopic reaotion rate ratios using Indium foils

Two irradiations were carried out, each with a single layer of foils, in the fuel and moderator only. Results are given in detail in Appendix V Table I, summarised in Appendix V Table II, and plotted, together with the manganese results, in Figure *11.*

8. Luteoium to manganese reaction rate ratios in the lattice cell

Three irradiations were carried out using foils from the same batch as were used in the previous core (reference $\overline{3}$). The technique was unaltered from reference 3, and results are given in detail in Appendix VI. As before, the manganese activity was separately analysed (Tables 3 and 4) and plotted in Figure 11. The agreement between these measurements and the manganese measurements was very satisfactory. The luteoium to manganese ratio is given in detail in Table I of Appendix VI, summarised in Table 2, and plotted in Figure 12. Run-to-run consistency was within the orrors of \pm 2% estimated in the analysis of results from the activity counting.

9. Plutonium to uranium reaotion rate ratios in the lattice cell:

Two irradiations were carried out in lattice cell J10 using plutonium 239-aluminium and uranium 235-aluminium foils. The experiment was carried out in an identical manner to that described in reference 3. Detailed results are recorded in Appendix VII Table 1, and are summarised in Appendix VII Table 2.

Examination of these results indicated that the errors .produoed by analysis of the counting sequence (those quoted in Table I are based on the consistency of the counts obtained during the experiment) were quite insufficient to explain the differences in ratios of up to 10 % obtained in separate irradiations. Up to that point in time the importance of placing the wrapped plutonium foils the same way up in the counting equipment has not boon appreciated. A subsequent experiment showed that about a 3% difference existed between results counted one way up and the reverse way due to the foil overlap on one side produced by the wrapping process. This 3% error goes some way to explaining the observed disorepancies between irradiations. Results obtained in later cores, where careful note of the foil orientation was made, point to this as being the most probable explanation.

Results from both irradiations are averaged in Table 2, and the error on the mean at each position computed from the average range of the two irradiations. The results are plotted in Figure 13.

10. Uranium 238 to uranium 235 Fission Ratio and Relative Conversion Ratio

The technique has been fully described in references 4 and 5 and its application.to DIMPLE is described in the previous report in this series (Referenoe 3).

Three irradiations were carried out, and the results are given in Appendix VIII. The chemical separation technique was not used in this oore. In Appendix VIII Table 1 gives details of individual runs, and Table 2 summarises the RCR and fission ratio results which are plotted in Figure 14. Finally Table 3 of Appendix VIII gives the results of U238 capture, U235 fission, and U238 fission in each ring of fuel, and these are plotted in Figure 15. The sharp rise of the U238 fission rate in the outer ring of fuel pencils compared with the smoother variation of U235 fission (see Figure 15) explains the somewhat discontinuous variation of the 1238 to U235 fission ratio (see Figure 13).

There are three types of error associated with the RCR and fission. ratio results of Appendix VIII, namely

- (1) Estimated errors, based on run-to-run consistenoy of results.
- (2) Relative errors, which are oalculated from the estimated errors plus foil holder calibration errors.
- (3) Total errors, which are calculated from the relative errors plus known sources of systematic errors.

For the RCR measurements a correction of $+$ 3.7% to the measured value was necessary to correct the DIMPLE reference spectrum to that in NESTOR. The method of making this correction by measuring the U238 cadmium ratio in the reference search tube (015) introduced a systematic error of up to -2% , that is, the correction of + 3.7% might be too large by up to 2% . In addition a correction of up to - 1% (for 2.5 Co) and - 0.3% (1.28 Co) was applied in the RCR to allow for the experimental foils being respectively .010" and .004" smaller in diameter than the normal fuel pollets. The smaller diameter was necessary to allow the experimental foil packs to be wrapped in aluminium foil to ensure alignment. The systematic error due to this effect was \pm .2%. Finally a systematic error of - 0.3% allowed for the bowing of the depleted (metal) foils used in the fission ratio, since the fission ratio result was fed into the RCR calculation.

For the fission ratio measurements a systematic error of \pm 10% arises from the uncertainty in the calibration factor relating fission product activity ratio to actual fission ratio.

11. Conclusions

- (1) Poorer spectral matching between centre $(D_2O/H_2O$ mixture cooled) and driver $(H₂O \text{ cooled})$ regions in this core compared with the fifth core (3) (where both regions were H₂0 cooled) has reduced . the radial region of constant cadmium ratio to 48.3 cm or slightly greater. This is, however, still considerably greater than that of the first two cores (1) , (2) , The radial component of buckling was deduced to be $2.55 \pm .12$ m⁻², the error quoted being double the random error to allow for systematic effects caused by the known, but little understood, asymmetry.
- (2) An unmistakable radial asymmetry existed in the centre of the core. This was shown to be a function of the fuel wurrounding the relevant search tubes, but examination of the fuel and coolant revealed no obvious cause. An axial scan in the relevant search tube agreed well with another in the core centre.
- (3) The axial flux shape was once again seriously perturbed by the gaps in the fuel and the presence of aluminium spacer plates, and only six measuring positions out of twenty six were used in the final fit, since the regions of apparently constant spectrum between plates (themselves about 30 cm apart) was 10 cm or less. The axial component of buckling was deduced to be 4.06 ± 0.03 m⁻², the' standard deviation quoted being based on internal consistency only. It is recognised that the method of analysis use& cannot eliminate entirely the possibility of some systematic error, and for this reason we have arbitrarily increased the above error by a factor of ten in our recommendation.
- (4) Internal oonsistenoy of microscopic reaction rate distributions were very satisfactory with the one exception of the plutonium to uranium ratio. It is thought that the poor consistency here was mainly due to non-appreoiation of the importance of counting the foils with one particular face upwards, since the method of foil wrapping was not uniform on both sides of the foil.
- (5) .During.the manganese miorosoopic reaction rate measurements it was demonstrated that macroscopic correction, to the observed. reaction rates by $1/J_{\Omega}(\beta r)$ yields consistent results when the macroscopic correction to be applied differs appreciably between irradiations.
- (6) Application of experience gained on earlier cores with respect to detector positioning has improved the shape of the approaohto-oritioal curves and reduced the differences between estimates of reactivity by the sub-oritioal mulitplication technique.

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Aoknowledgements

The Authors wish to acknowledge the co-operation of the DIMPLE Operations and Maintenanoe Staff, without whose efforts much of the work reported here could never have been oompleted.in the time allotted (the experiment was completed in two and a half weeks)

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Appendix I

Details of core materials

The details were identical to those of reference 3 with the one exception that, for region A (see Figure 4 (b) of this report), the E20 was replaced by a mixture of 29.7% E20 and 70.3% D20 by weight.

Reference 3 should be used to obtain all neoessery data.

Appendix II

Radial scan results

Table 1

Speotrum results

Absolute Rs in K11 Absolute Ro in K11 Mean Pu/U ratio (out to radius of 48.5 cm) Pu/U ratio in ESTOR Thermal column (reference 1) Pu/U DIMPLE/Pu/U NESTOR $- 31.4 \pm .1$ - 41-5 + -5
- 0.4047 + .0006 $= 0.3514 \pm .0009$ -1.152 ± 0.004

> (compared with $1.145 \pm .046$ using Pu and U foils - see Appendix VII) \mathcal{L}

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Table 2

Summary of all radial scan results

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Radial buckling

*Based on theoretiosl error of 0.4% on each of three measurements inoreased by factor 2 to all allow for possible asysmetry.

Table 4

Measurements used	ß	Var β	x^2	Corrected $S.D. on \overline{B}$	
A11		1.575 0.6 x 10 ⁻⁴ 51		032ء	
All except K13		1.582 0.6 \times 10 ⁻⁴	24	027ء	
All except K13 and G13		1.596 0.6×10^{-4} 5.7		.019	

 \mathcal{A}_1

Arial scan results

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Apenix IV

Manganese foil irradiation results

Table I

Basic Results

Details of irradiation

Run No:- I *Note incorrectly placed foil - macroscopic correction includes axial factor Date:- 13/11/62.

Axial foil positions:- 59.14 cms **and** 52.44 **cms** above bottom of fuel.

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Table I (contd.)

Basic Results

Details of **irradiation**

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Run **Not-** II

Date:- 14/11/62.
Axial foil positions:- 59.14 cms and 52.44 cms above bottom of fue

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Table 1 (contd.)

Basic Results

Details of irradiation

 \mathcal{A}_\bullet

Dates:- 16/11/62.

Axial foil positions:- 59.14 ems and 52.44 **cms** above bottom of fuel

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Run No:- III

Table 2

Summary of results

 \overline{C}_{11} = Mean ratio of activity in run i to that in run 3 in each position measured.

 \overline{c}_{11} = 0.95265 \overline{c}_{21} = 1.01508 \overline{c}_{31} = 1.00000 \overline{c}_{41} = 0.99928 \overline{c}_{51} = 0.98976 \overline{c}_{61} = 1.01202

Remarks:

The omissions in columns x_1 and x_2 were results which differed from the mean by 6 standard deviations, probably arising through foil positioning errors.

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Appendix V

Table I

Basic Results

Details of irradiation

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Run No:-Date: 15/11/62 Axial foil positions:- 49.54 **cms** above bottom of fuel

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Table I (contd.)

Basio Results

Details of irradiation

 \mathcal{A}_\bullet

Run No $t-$ I Date i - 14/11/62.

Diagram:- See Fig.

Axial foil positions:- 52.44 cms above bottom of fuel

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Summary of results

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 C_{21} = Mean ratio of activity to run 2 to that in run 1 for each position measured.

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-0.98862

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Appendix VI

Manganese/Intecium foil irradiation results

Table 1

Basic Results

Em For-Date of irradiations-Time of start of irradiations-Langth of irradiations-77 Time of start of Mn countings-
Time of start of Dn countings-. Foil details:-

 \mathbf{r} 13/11/62.
19 Hrs. 15 N. 21 Secs.
1 Hrs O N. 8 Secs. The U ma C Souster on 13-11-62. The 12 Ers. 53 M. 4 secs. on 15-11-62. 9 Ers
12 Ers. 42 M. 22 Secs. on 16-11-62. 9 Ers
Dis. 0.480" In content 13.74% by veight
m. content 5.76% by weight

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II 15/11/62.
14 Ers. 18 N. 48 Secs. 4 mass of the second 15-11-62.
17 Hrs. 23 M. 20 Secs. on 15-11-62.
9 Hrs. 53 M. 1 Sec. on 19-11-62. III
16/11/62.
8 Hrs. 38 X. 5 Secs. 60 Mins. 10 Hrs. 43 M. 45 Secs. on 16-11-62.
11 Hrs. 41 M. 9 Secs. on 19-11-62.

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Table 2

Summary of results relative to the NESTOR Thermal Column

Lu/Mn ratios

 \mathbb{R}^2

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 \mathcal{L}

M' %O

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Summary of results of Manganese activity

- Mean ratio of activity in run i to that in run 1 for each position ii measured

Radius	from cell	τ_1	τ_2	τ_3	Noan	Standard Deviation	
position	centre	τ_1	τ_2	τ_3	τ_3	Noan	Standard Deviation
52.5	0	0.25722	0.25074	0.25186	0.25327	0.0019	
52.5	1.803	0.26995	0.26742	0.26990	0.26909	0.0019	
52.5	3.708	0.33517	0.34186	0.33435	0.33713	0.0019	
52.5	5.525	0.51197	0.52287	0.51694	0.51726	0.0019	
52.5	17.062	1.00000	1.02129	1.01448	1.01192	0.0063	

 \overline{c}_{21} = 1.02129 \overline{c}_{31} = 1.01448

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Appendix VII

Plutonium/uranium foil irradiation results

Table 1

Basic results

In Dimple $14/11/62$. Run No. 1 In Nestor Thermal Column tor Instant
11/12/62.
15.00 hrs.
30 mins.
16.33 hrs. Date of irradiation 13.32 hrs. Time of start of irradiation Length of irradiation 30 mins. Time of start of counting 15.09 hrs.

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Table I (contd.)

Run No. Date of irradiation . Time of start of irradiation Length of irradiation	In Dimple 15/11/62. 12.00·hrs. 30 mins.	In Nestor Thermal Column 6/12/62. 14.39 hrs. 30 mins.	
Time of start of counting	14.12 hrs.	16.47 hrs.	

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Sumimary of results

*From spread of results mean range w .0734

 \cdot . Mean S.D. = \cdot 0734/1.13

S.D. on mean of 2 observations

 $\frac{.0649}{\sqrt{2}}$.046

 \bullet .0649

Appendix VIII Relative modified conversion ratio (RCR9) and fission ratio results

TABLE I Basic Results

 $Run No.: D5$

Date irradiation: 12/11/62

Time of start of irradiation: 12.33 hours

Length of irradiation: 2 hours

Location of thermal foil: DIMPLE reflector (SEARCH TUBE $c.15$) U²³⁸ Cd. Ratio = 27.9 Height.correction depleted foil: -0.21%

Remarks: Foil rotator in C15 stopped during the irradiation

Run No.: D6

Date irradiation: 13/11/62

Time of start of irradiation: 0917 hours

Length of irradiation: 2 hours

Location of thermal foil: DIMPLE reflector (SEARCH TUBE C.15) $U^{2,38}$ Cd Ratio = 27.9 Height correction depleted foil: -0.21%

Table I (Contd.)

Run No.: D7

Date irradiation: 15/11/62

Time of start of irradiation: 0831 hours

Length of irradiation: 2 hours

Location of thermal foil: DIMPLE reflector (SEARCH TUBE C.15) U^{238} Cd Ratio = 27.9 Height correction depleted foil: -0.21% \sim

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Summary of RCR* and fission ratio results

(a) Relative modified conversion ratio (RCR*) measured by coincidence counting method

'Radius (\mathtt{cm})	Relative error	Total error	Means corrected for smaller foil O.D.
	0.8	$+0.8 - 3.1$	2.694
1.803	0.8	$+0.8 - 3.1$	2.595
3.708	0.8	$+0.8 - 3.1$	2.278
$5 - 525$	0.8	$+0.8 - 3.1$	1.856

(b) 238/235 fission ratio per atom

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Summary of σ^{238} capture, σ^{235} fission and σ^{238} fission distributions (All in J10)

mean ratio of run y to run x $c_{\mathbf{x} \mathbf{y}}$ \bullet

(a)
$$
\overline{v}^{238}
$$
 Captive distribution (\overline{v}^8)

Table 3

(b) u^{235} Fission distribution (F^5)

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Radius from cell centre (cm)	F_1^8 C_{11}	$\begin{array}{ c c c c c } \hline \end{array}$ $\begin{array}{ c c c } \hline \end{array}$ $\begin{array}{ c c c c } \hline \end{array}$ $\begin{array}{ c c c $		Mean	Macros. Corr	Corr. Mean	Stand. Dev.
	.0277	.0272	.0275	.0275	0.98464	.0271	.0002
1.803	.0287		.0287	.0287	0.98778	.0283	.0002
3.708	.0277	.0283	.0283	.0281	0.99063	.0278	,0002
5.525	.0455	.0453	.0448	.0452	0.99299	.0449	.0003

(c) \overline{u}^{238} Fission distribution (\overline{r}^8)

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 $\mathcal{L}^{\text{max}}(\mathbf{r})$

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FIG.I PLAN VIEW OF TANK TOP

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CHANNEL $I \sim$ 170 cm FROM TANK BOTTOM ~80 cm. FROM TANK BOTTOM

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FIG.b SOURCE AND DETECTOR POSITIONS FOR SUB-CRITICAL REACTIVITY MEASUREMENTS

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CADMIUM RATIOS AS FUNCTIONS OF RADIUS

 $\mathcal{P}^{\text{max}}_{\text{max}}$

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FIG.9 AXIAL CADMIUM RATIO DISTRIBUTION IN KIT

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LATTICE CELL

 $\label{eq:3.1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{i=1}^{N-1} \frac{1}{\sqrt{2}}\sum_{$

FIG. 12 LUTECIUM TO MANGANESE RATIO IN THE LATTICE CELL

 $\mathbf{A}^{(n)}$ and $\mathbf{A}^{(n)}$

 $\mathcal{F}^{\text{max}}_{\text{max}}$ and $\mathcal{F}^{\text{max}}_{\text{max}}$

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FIG.15 U235 FISSION, U238 FISSION, AND U238 CAPTURE IN THE FUEL (RANDOM ERRORS TOO SMALL TO PLOT)