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RESEARCH MEMORANDUM

CALCULATED EFFECT OF URANIUM DISTRIBUTION ON REFLECTOR

CONTROL EFFECTIVENESS FOR A WATER-MODERATED

POWER REACTOR

By Thomas A. Fox and Michael F. Valerino

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NATIONAL ADVISORY COMMITTEE FOR AERONAUTICS

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RESEARCH MEMORANDUM

CALCULATED EFFECT OF URANIUM DISTRIBUTION ON

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SUMMARY

Two-group theory calculations were made to determine the effect of nonuniform uranium *loading* **as** cougared to uniform **loading** on the refIector control effectiveness attainable in a large thermal reactor of present interest **in** aircraft power application (the supercritical water reactor). The reflectors investigated were a 10-centimeter and an effectively infinite-thickness water reflector, which were considered to be practical for use in the particular reactor design considered.

The reflector-control mechanism considered employs a thermal neutron absorber that can be mved from a position in the reflector far **enough** from the cylindrical core to have negligible effect on the *reac*tivity to a position at the radial reflector-core interface where it could conceivably absorb all therm1 neutrons **trying** *to* leave or enter the radial boundary of the core.

The results **showed** that.nonuniform uranium loading **to** attain uniform radial power production doubled the reflector control effectiveness **over** that **with** the uniform uranium loading. Ebwever, this doubling of control effectiveness **was** still imufficient to provide the amount of control necessary for operation of the reactor. Even for the *most* favorable case considered, the change in reactivity obtained from reflector control **was** only 0.03 as compared to the **0.132** needed for complete control.

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Although the increase in reflector control effectiveness due to nonuniform uranium loading is not large enough to be of use for the reactor considered herein, it **may** provide the required margin to- permit use of reflector control for a smaller, more heavily uranium-loaded reactor, particularly if a more efficient reflector such as beryllium is used instead of water.

Although, for reactors loaded to give uniform radial power, reflector poisoning greatly distorted the power distribution, the resultant distribution was more favorable than that for the uniform uranium loading with or without reflector poisoning.

INTRODUCTION

Considerable interest exists **in** the use of the reflector as a means of controlling.the reactivity of power reactors for certain installations. By making use of a parasitic neutron absorber in the reflector and varying its position with respect to the reflector-care interface, it is possible to produce a definite change in reactivity in the reactor. **This** change in position or distance from the interface could be accomplished by several **means. For** a reflector of solid materials, a set of rods (made of the reflector material) coated with absorber on one aide and designed to be rotated **on** axes parallel to the core axis could be used. For water reflectors, rotating **drums or** just strips of the absorber could be utilized in a similar manner. In general, the reflectortype control, where usable, requires less space than the more conventional absorber-rod control. However, for water-moderated reactors of the size needed to accommodate the heat-transfer surface area and coolant-water flows required for power application, the change in re. activity attainable with reflector control is very **small.**

In reference 1, the manner **of** distributing the uranium over the reactor core volume to attain uniform power production is determined for **a** spherical water-moderated reactor for three thicknesses of water reflector. The uranium distributions obtained involve high uranium. concentrations near the reflector-core interface relative to the *con*centrations in the central portions-of the reactor core. To illustrate, for me of the reactor assemblies investigated in reference 1 **(having** an 8-cm reflector thickness) the uranium concentration near the reflector-core interface **was** of the order of three to four times that at the core center; the total uranium investment was about 15 percent higher than that for the **uniform** uranium dlstribution case. It **is** to be expected, then, that the action of the reflector in maintaining reactor criticality is much more important for the case of uniform power production (obtained by nonuniform uranium distribution Over the core production (obtained by nonuniform uranium distribution over the core
volume) than for the case of uniform-uranium-distribution; hence,
greater control effectiveness should be attainable with the reflector for the case of uniform power production than for the case of uniform uranium distribution. The question arises as to the magnitude of this increase in reflector control effectivenees.

In order to provide **an** indication of the magnitude of this effect, calculations were made for the supercritical water reactor described in reference 2 to determine the increase in reflector control effectiveness attainable by distributing the fissioasble material nonuniformly over the reactor cylindrical-core volume in a manner resulting in **uniform** radial power production. **This** reactor design is considered representative of the water-moderated reactors under consideration at present for power applications.

The solution of the poisoned condition in the reflector **was** accomplished by approximating the control-rod system with **a** cylindrical sleeve **of** poison at the radial core-reflector interface. This represents the ideal case or the maximum change in reactivity possible. Since most absorbers are not very effective in capturing fast neutrons, ¹¹⁰effect **on** the fast flux **was** considered other than the indfrect change caused by the difference in the thermal flux. The fast flux therefore **was** continuous at the reflector-core interface and dropped to zero at the extrapolated outer boundary of the reflector. The thermal neutrons were considered *to* be entirely taken up by the **small** layer of absorber; hence, the thermal flux went to zero at the interface.

In order to facilitate discussion of the calculations, the reactor conditions are defined *as* **follows:**

Condition (a): cold-clean. - Condition **(a)** is the startup condition at mom temperature, **with** no **fission** poison, and with sufficient fissionable material to allow for the contemplated burnup during the life **of** the reactor.

Condition (b) : cold-clean, poisoned reflector. - Condition (b) is identical to condition (a) except that the thermal neutron absorber is in position at the **radial** reflector-core interface.

Condition (c) : hot-burnup. - Condition (c) occurs at the end of the useful reactor life, herein taken *as* corresponding to **1.3-kilogram** burnup of U²³⁵. This includes equilibrium poisons. The calculations were made st operating temperatures, to be discussed later, and the condition is taken as critical.

Condition (a): cold-clean condition for **0.65-kilogram** burnup. - Condition (d) is a startup condition with everything the same as in condition (a) except for 0.65 -kilogram smaller loading of U^{235} .

 \mathbf{a}

For each of the reactor cases considered, a calculation **was** mde first for condition (c) in order to establish the proper uranium investments necessary **during** the life of one fuel **loading** of the reactor. CalcuLations for conditions **(a), (b),** and (d) were then performed to give the changes in reactivity present **under** the other conditfons.

SYMBOLS

 $\Delta\Delta\Delta\approx 0.5$

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*^V*average number of neutrons produced per fuel **atom** fissioned p density, g/cm^3 *2,* macroscopic neutron absorption **cross** section $\Sigma_{a,th}^{M}$ thermal value of Σ_n for moderator and structure macroscopic neutron fission cmsa section Σ_F $\boldsymbol{\Sigma}_{\mathbf{q}}$ macroscopic neutron cross section **for** *slowfng* **down** average value of the macroscopic neutron absorption cross $\Sigma_{\rm p,th}$ average value or the macroscopic neutron apsention for stable fission-product poisons $\Sigma_{u,th}$ macroscopic thermal neutron absorption cross section for U^{235} $\sum_{p,\text{th}}^{\text{T}}$ total macroscopic thermal neutron absorption cross section for all poisons *⁰*P,th average **value** of microscopic thermal-neutron absorption **=s** microscopic neutron scattering cross ection $\sigma_{\texttt{sm},\texttt{th}}$ average value of microscopic thermal-neutron absorption cross section for Sm^{149} $\sigma_{\rm tr}$ microscopic neutron transport cross section uxe,th average value of microscopic thermal-neutron absorption **cross** section for Xe¹³⁵ **CP** neutron flux Subscripts : f **fast** neutron **group** 0 refers to cases Kith **unlform-uranium** distribution *at* the hotburnup condition r **radial** position from **axls** of core - **sm** property of **sm149** th thermal neutron **group**

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u property of U^{235}

 x e property of $xe^{1.35}$

REACTIVITY CALCULATIONS

Description of Reactor

The reactor core is a 2.5-foot square cylinder with supercritical water (pressure, 5000 lb/sq in.) functioning as the combined coolantmderator and with the UL3\$ fuel contatned **in** stainless-steel-clad, sandwich-type plates. The core **is** reflected by supercritical water. At the hot conditions (corresponding to reactor full-power output), the average water temperature **in** the core is *620°* ^F(kT energy of 0.052 *ev)* and in the reflector is **480°** F *(3rT* energy of 0.045 ev) . **At** these conditions the average water density is 0.71 grams per cubic centimeter in the core and 0.83 grams per cubic centimeter **in** the reflector. For the cold conditions (prior to reactor startup), the temperature in the core and reflector is taken as 59⁰ F with the corresponding water density of 1 **gram** per cubic centimeter. Table I presents a tabulation of the core and ref lector compositions **for** the **hot** and cold conditions. The uranium contents are determined by the criticality calculations for **two** reflector thicknesses (10 cm **and** infinite) and for the cases of uniform **U235** distrlbution and **for** the distribution **giving** constant radial power production in the reactor core.

General Method of **Analysis**

In order to assure proper Investment for the entire life of one .. fuel loading of the reactor, it was necessary first of all to make criticality calculations at the hot-burnup condition described previously. The reactor was considered to be at the full-power operating conditions at the end of its life. The reactor poisons considered were: (a) equilibrium concentrations of XeU5 **and** SmL4Y and (b) stable f ission-product poisons corresponding to approximately 10 percent **U235** burnup. The poisons were taken to be uniformly distributed over the core volume. **This** assumption is justiffed **by** the results of the investigation of reference 3. The stable poisons were specified **as** having *an* average thermal absorption cros8 section of 75 barns per fuel atom destroyed. This is the value at 0.025-ev energy, and a **l/v** variation is assumed. For the hot-burnup condition the uranium content required for reactor criticality **was** determined for each of the **following** cases:

I. Uniform U^{235} distribution, 10-centimeter reflector

11. Uniform radial power distribution, 10-centimeter reflector

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Uniform U²³⁵ distribution, infinite reflector III.

IV. Uniform radial power distribution, infinite reflector

For cases I and **III,** the radial power distribution is also obtained, while for caees.TI and **IVY** the uranium distribution is **also** obtained in the criticality calculations.

At the cold-clean condition, previously described briefly, no poi**sons** were present in the reactor core and the uranium content was larger than at the hot-poisoned condition by the amunt *of* fuel burnup, whfch **was** assumed to be **1.3** kilograms. This fuel burnup, which **is** somewbat less than 10 percent *of* the fuel investment, corresponds to the amount required for 300,000-kilowatt reactor parer output for a total **of 100** hours. For each of the cases I to IV, the 1.3 kilograms of U^{235} was distributed over the core **volume** so that the local fuel burnup *is* proportional to the **Local** power (or fission-rate} production existing at the hot-burnup condition (considered to be at the end **of** reactor life) . Although the relative **local power** production actually varies with time, this variation was small for the small burnups herein involved so that negligible error **was** intmducea by distribution of the fuel burnup in this manner. For each of the cases I to IV, two reflector configurations were considered for the startup condition, **namely, (a)** the **normal** (unpoisoned) water ref lector, and **(b)** the water reflector incorporating a sleeve **of** thermal-neutron poison sufficient to make 0th go to **zero** adjacent to the entire cylindrical bundary of the core.

From the foregaing calculations, the reactivity change from hot**burnup** to cold-clean and the reactivity change attainable with a thermally **poisoned** reflector were obtained for each of cases I to IV.

For comparative purposes, Calculations were also made for cases I to IT, cold-clean poisoned-reflector conditions, of the effect of initial loadings limiting the uranium burnup to 0.65 kilograms.

Reactor Calculations and Evaluation **of** Nuclear Constants

The two-group neutron-diffusion equations applicable to core and reflector in *a* critical reactor assembly are:

$$
\frac{\lambda_{\text{tr},\text{f}}}{3} \nabla^2 \varphi_{\text{f}} - (\Sigma_{\text{a},\text{f}} + \Sigma_{\text{q},\text{f}}) \varphi_{\text{f}} + k_{\text{th}} \Sigma_{\text{a},\text{th}} \varphi_{\text{th}} + k_{\text{f}} \Sigma_{\text{a},\text{f}} \varphi_{\text{f}} = 0
$$
\n(1)

$$
\frac{\lambda_{\text{tr,th}}}{3} \sqrt{2} \varphi_{\text{th}} - \Sigma_{\text{a,th}} \varphi_{\text{th}} + \Sigma_{\text{q,f}} \varphi_{\text{f}} = 0 \tag{2}
$$

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For a fully reflected cylindrical core (with both end and side reflectors), φ_f and φ_{th} are functions of two dimensions, radius r and height z (see fig. 1). Inasmuch as the effect of the side reflector on the reactivity of the reactor assembly **is** of interest here, the fully reflected assembly can, for this purpose, be suitably approximated **by** an equivalent reactor core, bare **at** the ends and reflected at the **sides;** this approximation leads to separation of the variables r **and z, in which case the flux** φ **is given by the product** $\varphi(\mathbf{r}) \psi(\mathbf{z})$ **where** $\varphi(r)$ is a function of r only and $\psi(z)$ is a function of z only.

In the use of this approximation, the half-height $H_c/2$ of the equivalent reactor core is increased above that of the given **fully** reflected core **by** an amount equal to the reflector savings, **as** illustrated in figure 1. Reflector savings for water reflectors around watermderated cores are presented in reference **4 and** are substantially independent of core conrposition for water-mderated cores *that* are predominant **ly** therrnal.

Inasmuch *as* the ends &re bare **for** the equivalent reactor core, **Of** and φ_{th} must fall to zero at $z = \pm \frac{H_c}{2}$. If it is assumed that $\varphi = \varphi(r) \cdot \psi(z)$, this condition is satisfied by

$$
\psi(z) = A \cos \frac{\pi z}{H_C}
$$
 (3)

and, noting that **for** cylindrical geometry

$$
\nabla^2 = \frac{\partial^2}{\partial r^2} + \frac{1}{r} \frac{\partial}{\partial r} + \frac{\partial^2}{\partial z^2}
$$
 (4)

equations (1) and (2) reduce to equattons *in* the independent variable r only:

$$
\frac{\lambda_{\text{tr},\text{f}}}{3} \nabla_{\text{r}}^2 \varphi_{\text{f}} - \left(\Sigma_{\text{a},\text{f}} + \Sigma_{\text{q},\text{f}} + \frac{\lambda_{\text{tr},\text{f}}}{3} \frac{\pi^2}{\text{H}_c^2} \right) \varphi_{\text{f}} + k_{\text{th}} \Sigma_{\text{a},\text{th}} \varphi_{\text{th}} + k_{\text{f}} \Sigma_{\text{a},\text{f}} \varphi_{\text{f}} = 0
$$
\n(5)

$$
\frac{\lambda_{\text{tr,th}}}{3} \nabla_{\text{r}}^2 \varphi_{\text{th}} - \left(\Sigma_{\text{a,th}} + \frac{\lambda_{\text{tr,th}}}{3} \frac{\pi^2}{\pi^2}\right) \varphi_{\text{th}} + \Sigma_{\text{q,f}} \varphi_{\text{f}} = 0 \tag{6}
$$

where

$$
\nabla_{\mathbf{r}}^2 = \frac{\partial^2}{\partial \mathbf{r}^2} + \frac{1}{\mathbf{r}} \frac{\partial}{\partial \mathbf{r}}
$$

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and where $\varphi_f(r)$ and $\varphi_{th}(r)$ are designated, for convenience, as φ_f and φ_{th} . The terms involving the coefficient π^2/E_c^2 account for the net axial leakage of neutrons in the equivalent reactor core. Hence, if the proper value of H_c is used, it effectively accounts for the axial leakage in a fully reflected core. Equations (5) and (6) apply **to** either the'core or the side reflector by use of the appropriate nuclear constants characteristic of either the core or reflector cornposition. **In** the application of equations (5) and (6) for the uniform-radial-power cases (cases II and *N)* , account **was** taken of the variaradial-power cases (cases II and IV), account was taken of the varia-
tions of the fast as well as the thermal parameters with radial position r across the reactor core.

In the solution of the core **and** reflector equations, the radial boundary conditfons were taken **as** follows:

(1) For the normal reflector: $\varphi_f = \varphi_{th} = 0$ at the outermost (ex**^a**trapolated) boundary of the reflector; fast **and** thermal flux and current continuity were assumed at the core-reflector interface.

cu cu c (2) For the reflector incorporating thermal-neutron poison adjacent \overline{c} to the core boundary: $\overline{v}_P = 0$ at the outermost boundary of the reflector to the core boundary: $\Phi_P = 0$ at the outermost boundary of the reflector; fast flux and current continuity were assumed at the core-reflector interface; $\varphi_{th} = 0$ at the core boundary. Note that these boundary conditions imply that the fast flux is unaffected by the reflector poison except as indirectly affected by the thermal flux falling to zero at the core boundary.

> The two-group equations for core and reflector subject to the fore**going** boundary conditions were solved by use of an electrical-analog simulator at the **NACA** Lewis laboratory. **This** nuclear-r-&actor simulator and the general procedure in its **use** to solve reactor criticality prob**lems** have been described in **detail in** references 1, 5, **and** *6.*

> The proceaure for evaluating the nuclear constants for use in **equa**tions (5) **and** (6) is described **in** reference *7.* In the evaluation of the constants, use is made of the following definitions:

$$
L_{f}^{2} = \frac{\lambda_{tr,f}}{3(\Sigma_{a,f} + \Sigma_{q,f})}
$$

$$
p_{th} = \frac{\Sigma_{q,f}}{\Sigma_{a,f} + \Sigma_{q,f}}
$$

$$
k_{f} = \frac{\nu \Sigma_{F,f}}{\Sigma_{a,f}}
$$

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$$
k_{th} = \frac{v\Sigma_{F, th}}{\Sigma_{a, th}}
$$

The procedure is patterned after that successfully used in reference 8 to predict the criticality of water-moderated reactors. For the reactors of reference 8, $p_{th} \approx 1$, whereas for the reactors herein considered, p_{th} \approx 0.80 to 0.90; hence, the fast-fission contribution should be accounted for. The general procedure of reference 8 is used in accounting for this effect, as well as for the calculations of the cross section values. The procedure is briefly outlined *as* **follows:**

 $\lambda_{\text{tr},f}, \Sigma_{\text{a},f}, \Sigma_{\text{F},f}, \text{p}_{\text{th}}$. - The quantities $\lambda_{\text{tr},f}, \Sigma_{\text{a},f}, \Sigma_{\text{F},f}$, and Pth were obtained by weighting **local** energywise values according to the energy distribution **of** neutron **flux, a8** indicated by age theory, in an infinite medium of the same composition. The dependence of this distribution on the fission spectrum is included.

 L^2_{f} . – For water, L^2_{f} is based on the experimental value of 33 square centimeters **at** room temperature **(p** = **1** g/cc) and **1s** taken as inversely proportional to the square of the water density at higher water temperatures. For the given core comgosition, this **value is** increased by 2 square centimeters to account for the 11.6 volume percent of stainless steel in the core. $\begin{array}{c}\n \text{energy of:}\n \text{infini}\n \text{in} \n \text{in} \n$ where of the water density at higher
en core composition, this value is
is to account for the 11.6 volume per-
pre.
and of reference 9 to account for the
experimental values of σ_g for

 $\lambda_{\text{tr.th}}$. - By use of the method of reference 9 to account for the chemical binding of hydrogen, the experimental values of $\sigma_{\rm g}$ for hydrogen are used to calculate the local values of $\sigma_{t,r}$ of hydrogen. The quantity $\lambda_{tr.th}$ is then evaluated by weighting $\lambda_{tr.}$ = according to the neutron **flux** in **a** Maxwellian distribution. $\mathbf{1}$ $\mathbf{z_{N_1}}$ $\sigma_{\mathbf{tr}^{\mathbf{\mathbf{\tau}}}}$

 $\frac{\Sigma_{\text{a},\text{th'}}\Sigma_{\text{F},\text{th}}}{\Sigma_{\text{a},\text{th'}}\Sigma_{\text{F},\text{th}}}$. The fission poisons are treated separately in the $\frac{\Sigma_{\text{a},\text{th'}}\Sigma_{\text{F},\text{th}}}{\Sigma_{\text{F},\text{th}}}$. The following description applies, however, for all materials in the reactor excepting Xe^{135} and $Sm^{\overline{1}49}$. The terms $\Sigma_{a,th}$ and $\Sigma_{\rm F, th}$ are obtained by assuming the local values of $\Sigma_{\rm a, th}$ and $z_{\rm F,th}$ are obtained by assuming the local values of $z_{\rm a,th}$ and $z_{\rm F,th}$ to obey the $1/v$ law and by then weighting the local values according to the neutron flux in a Maxwellian distribution. **For** this varfation with energy, $\Sigma_{\text{a.th}}$ (or $\Sigma_{\text{F-th}}$) equals 0.886 times the value of Σ_{a} - .. . - . . " " ecription applies, however, for all mate-
 xe^{135} and Sm^{149} . The terms $\Sigma_{a,th}$
uming the local values of $\Sigma_{a,th}$ and
by then weighting the local values accord-
xwellian distribution. For this variation with energy, $z_{a,th}$ (or $z_{F,th}$) equals 0.000 thes the value of z_{a}
(or \bar{z}_{F}) corresponding to the most probable energy (kT) of the thermal

Fission Poisons and Burnup

The equilibrium concentration of Xe¹³⁵ is given by

$$
N_{xe} = \frac{Fy_{xe}}{\lambda_{xe} + \overline{\sigma}_{xe,th} \varphi_{th}}
$$
 (7)

where $\vec{\sigma}_{\text{xe},\text{th}}$, obtained from reference 10, is given by weighting local values of σ_{xe} according to the neutron flux in a Maxwellian distribution.

For purposes of calculating **poison** concentrations, the reactor is assumed to be nearly thermal in which case $F = \Sigma_{F, th} \phi_{th}$ so that equation (7) can be written as

$$
N_{xe} \overline{\sigma}_{xe,th} = \overline{\Sigma}_{xe,th} = \frac{Fy_{xe}}{\lambda_{xe} + \frac{F}{\Sigma_{F,th}}} \tag{8}
$$

The equilibrium concentration *of* **S,149 is** given by

$$
N_{\rm sm} \overline{\sigma}_{\rm sm,th} = \overline{\Sigma}_{\rm sm,th} = \frac{F y_{\rm sm}}{\varphi_{\rm th}} \tag{9}
$$

or, for a thermal reactor,

$$
\overline{\Sigma}_{\text{sm,th}} = y_{\text{sm}} \Sigma_{\text{F,th}} \tag{10}
$$

The remining poisons, which are lumped together, are specified as having an average thermal absorption cross section $\bar{\sigma}_{p,th}$ of 75 barns per fuel atom destroyed at a temperature of 59⁰ F and as following the $1/\nu$ **law.** For 10-percent fuel burnup (11.1 percent **of** the fuel **left** in **the** reactor **at** the end of **its life),** the absorption **is** given by

$$
\overline{\Sigma}_{\text{p,th}} = 0.111 \left(\frac{75}{549} \right) \Sigma_{\text{F,th}}
$$

where 549 is the value of the U^{235} fission cross section at 0.025 ev.

The pertinent constants used to evaluate the foregoing poison cross sections are: $y_{xe} = 0.063$; $\overline{\sigma}_{xe,th} = 2.32 \times 10^6$ barns at 0.052 ev.; λ_{xe} = 2.103×10⁻⁵ sec⁻¹; y_{gm} = 0.014; average F = 2.7×10¹³ fissions per **second** per cubic centimeter 'based **on** 300,000-kilowatt reactor total **power output** at 200 MeV **per fission.**

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The core and reflector parameters for case I (uniform 0^{235} distribution, 10-cm reflector) are tabulated in table **II** for the cold-
clean and hot-burnup conditions. The parameters N_u, P_{th}, k_f, Z_{u,th}, and $\Sigma_{\text{F},\text{th}}$ vary with uranium concentration while the remaining parameters are essentially constant. The reflector parameters were the same for all cases, differing only for the change in temperature conditions **as** listed.

RESULTS AND DISCUSSION

(1) Reactivities and investments. - Table **I11** presents keff and Akeff due to reflector poisoning and also the uranium investments **for** all cases and conditions considered. The reactor is critical $(k_{eff} = 1.00)$ **for** the hot-burnup condition for each case. For the cold-clean condition various amounts of excess reactivity are present. **For** the uniform-uranium cases, k_{eff} is 1.154 for the 10-centimeter reflector and 1.147 for the infinite reflector; the amounts of excess reactivity to be controlled are approximately 0.162 and 0.148, respectively. Similarly, for the uniform power cases (cold-clean) k_{eff} is 1.150 for the 10centimeter reflector **and** 1.l32 **for** the infinite reflector, which means excessive reactivities of 0.150 and 0.132 , respectively. Since the introduction of thermal-neutron poison in the reflector causes changes of only 0.014 and 0.016 for the uniform-uranium cases, and 0.029 and 0.030 for the uniform-power cases, reflector control is inadequate in this type of reactor. Certain interesting observations can be made, however. Slightly greater control **was** possible **with** the better reflector. More important, nearly twice the change in reactivity **was** found when the fissionable material **was** distributed for uniform power production **as** compared to a uniform distribution of fuel. Condition (d) of all cases gives keff for the cold-clean reactor with **a** poisoned reflector but for an assumed **burnup** of 0.65 kilogram instead of **1.3-kilogram burnup** as in the previous cases. As expected, k_{eff} is reduced, but the reactor is still supercritical **by** 9 to **12** percent in the **various** cases.

The investment of uranium required for a critical assembly is less when distributed **uniformly than** when distributed for uniform radial power production. For the LO-centimeter reflector thickness, the uranium investment is increased **from** 16.0 to 21.05 kilograms when the fissionable material *is* distributed nonuniformly to attain constant radial power production; the corresponding increase **is** from **15.5 to 18.24** kiloable material is distributed nonuniformly to attain constant radial
power production; the corresponding increase is from 15.5 to 18.24 kilo-
grams for the infinite reflector.

(2) Ursnium distributions. - **In** figure 2 are presented the uranium distributions as functions of care radius for cases I through **IV** in the cold-clean and the hat-burnup conditions (for **1.3-kg** burnup); figure 2(a) is for the 10-centimeter reflector **(cases** I and TI} and figure 2(b) is for the infinite reflector (cases **I11 and IV)** .

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The ordinate in figure 2 is $N_u/N_{u,0}$ where N_u is the local-uranium concentration and $N_{u,0}$ is the concentration required for the uniform-uranium cases at the hot-burnup condition (case I(c) in fig. 2 (a) and case III(c) **in** fig . **2(b)**) . Figure 2 shows the typically **'high** uranium concentrations near the core-reflector interface relative to the concentrations in the central. portion8 of the core required to attain uniform radial power production. For the case of uniform uranium distribution in the hot-burnuy condition, the fuel burnup varzes over the reactor core **volume;** hence, the uranium *Loading,* of necessity, must vary over the core volume in the cold-clean condition. This variation, although slight for the burnup assumed, is evident in figure 2. For the cases of uniform power, the burnup is essentially constant over the core volume; hence, the uranium loading for **the** cold-clean conditfon is greater, **by** a constant amount, over that for the hot-burnup condition. The total uranium requirements for the uniform-power *cases* are 31.6 and 17.7 percent higher than for the corresponding uniform-uranium cases for the 10-centimeter and infinite reflector, respectively.

(3) Power distributions. - The power-production distributions within the reactor core are presented in figure **3** as plots of **E/&** ver-(3) <u>Power distributions</u>. - The power-production distributions with the reactor core are presented in figure 3 as plots of $\frac{H}{H_{\text{max}}}$ ve sus radius r, where H is the local power production and H_{max} is the meas the maximum power production. The ratio of the average *to* the maximum sus radius r, where H is the local power production and H_{max} is
the maximum power production. The ratio of the average to the maximum
power density H_{av}/H_{max} is also indicated for each of the cases treated. Figure 3(a) is for the 10-centimeter reflector (cases I and **11)** and figure 3(b) is **for** the infinite reflector (cases IU: **and** IV) . **For** each case, the power distribution and average- to maximum-power production is given for: **(a) the** cold-clean condition **with** unpoisoned reflector, **(b)** the cold-clean condition with **poisoned** reflector, and (c) the hot-burnup condition.

Figure 3 illustrates the large spatial variations in power obtained
for uniform uranium loading; for example, in figure $3(a)$ for case $I(c)$,
the nexter drama to 37.5 persont of maximum near the noflector. Companifor uniform uranium loading; for example, in figure 3(a) for case I(c), the power drops to 37.5 percent of maximum near the reflector. Compari**son** of the hot-burnup and the cold-clean unpoisoned-reflector conditions for each case gives an indication of the power variations with fuel burnup. In figure 3(a), comparison of II(a) and II(c) shows that for uniform power in the hot-burnup condition, the power distribution in the cold-clean condition is distorted resulting in $E/E_{max} = 0.83$ near the reflector and = 0.93 at the center of the core. In figure $3(b)$, case $IV(a)$ shows a more severe power distortion resulting in H/H_{max} = 0.76 near the reflector.

Figure 3 shows the distortions in power distribution caused by the use of reflector poisoning (cases I(b), II(b), III(b), and **IV(b)** in figs. $3(a)$ and $3(b)$. For the uniform-power cases (note that uniform power is achieved for hot-burnup condition **with** unpoisoned reflector), the distorted power distribution due to reflector poisoning is still

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more favorable, insofar as total power output for limitfng heat **flux** is concerned, than the power distributions for any of the conditions of the uniform-uranium cases. To illustrate, in figure $3(a)$, $H_{av}/H_{max} = 0.82$ for case II(b) compared to 0.61 for case I(c). Case II(b) is for the poisoned reflector, whereas case I(c) is for the unpoisoned reflector. The same-general result is indicated in figure $3(b)$ wherein $H_{av}/H_{max} = 0.79$ for case IV(b) compared to 0.65 for case $III(c)$. It appears, then, that if the uranium is distributed nonuniformly to achieve uniform power **during** normal reactor operation with unpoisoned reflector, the distorted power distribution resulting from the use of reflector poison is nevertheless more favorable than that for the uniform-uranium case with or without reflector poison.

CONCLUSIONS

Nonuniform **uranium** loading in the core of **a** large therm1 reactor . ." . . ." . ? $(2.5-ft)$ square cylinder with water moderation; resonance escape probability, \bullet 0.90) to attain uniform radial power production resulted in a doubling of the reflector control effectiveness over that obtainable for uniform uranium loading. A smaller further increase in effectiveness **was** also obtained by using a more efficient reflector. However, the reactivity changes were still much too **small** compared to the amount required, **For** the best case, the *change* in over-all **neutron** multiplication factor Δk_{eff} was 0.03 as compared to the 0.132 required. The uranium,investments required **for** the uniform-power cases were 31.6 and *17.7* percent higher than that for the comparable uniform uranium cases. The power distribution was better for the cases with the uranium distributed for uniform power, even after being distorted by the reflector poison (ratio of average to **max~mum power** deasity in reactor $reflector$ poison $(H_{av}/H_{max} = 0.61)$. $H_{\text{av}}/H_{\text{max}} = 0.82$) than it was for the uniform uranium case without

Although the increase **Fn** reflector control effectiveness **was** not sufficient to be of **use** for the reactor considered herein, *It* **may pro**vide the required margin to permit use of reflector control for a smaller, more heavily loaded reactor employing a more efficient ref lector.

Lewis Flight Propulsion Laboratory National Advisary Committee **for** Aeronautics Cleveland, Ohio, September 9, 1953.

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TABLE II. - TWO-GROUP THEORY REACTOR CONSTANTS FOR

CASE I (UNIFORM URANIUM LOADING, 10-CM REFLECTOR)

(b) **Reflector.**

***Representative values applying** only **to case** I. Al presentative values applying only to case 1. All other parameters are the same for all four cases.

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TABLE III. - REACTIVITY CHANGES DUE TO REFLECTOR POISONING (REACTOR CRITICAL

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AT HOT-BURNUP CONDITION, REFLECTOR POISONING

INTRODUCED IN COLD-CLEAN CONDITION)

reflector

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Fully reflected reactor core

Equivalent core with bare ends

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Figure 1. - Conversion of fully reflected core to an equivalent core with bare ends by application of end reflector savings.

 $\alpha = 1.4$

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(a) Reflector thickness, 10 centimeters.

Figure 2. - Variation in uranium loading for hot-hurnup and cold-clean conditions for cases where uranium is adjusted to give uniform loading and uniform power in hot-burnup condition.

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cold-clean conditions for cases where uranium *is* **adJusted to give** uniform **loading and uniform power in hot-burnup condition.**

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(b) Infinite **water reflector.**

Figure 3. - Concluded. Variation in power production and ratio of average **to** maximum parer **density** wer **reactor** volume for **cases where uranium Le addusted to give uniform loading and** uniform **pawer in hot-burnup condition.**

NACA - **Longley Fidd.** *VL.*

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