

Re-fueling the actinides ^{239}Pu , ^{241}Am and ^{243}Am as burnable absorbers in the MTR-22 MW research reactor

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□ ABSTRACT □

The main objective of this paper is the re-fueling of the MTR-22MW reactor by the $\text{U}_3\text{O}_8\text{-Al}$ Original Fuel Elements (OFEs) mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides as burnable absorbers. This procedure helps to consume some of the stored ^{239}Pu , and minimize the radiological impacts of the actinides ^{241}Am and ^{243}Am on the environment. The neutronic parameters such as: Reactivity ρ_{ex} , Control Plates Worth (CPsW), Shutdown Margin (SM), Shutdown Margin with first control plate failure (SM-1), Reactivity Safety Factor(RSF), neutron spectrum in the Central Neutronic Trap (CNT), in the Central Irradiation Box (CIB), in the Be reflector and Power Distribution Factors (PDFs) of the MTR-22MW reactor were re-evaluated using the MCNP4C transport code. The variation of the reactivity ρ_{ex} was studied as a function of the operation time of the reactor as well. The obtained results showed that the MTR-22 MW reactor can be operated with $\text{U}_3\text{O}_8\text{-Al}$ OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides within the safety conditions established in the design. Re-fueling of the MTR-22 MW reactor by $\text{U}_3\text{O}_8\text{-Al}$ OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides leads to reduce the enrichment with ^{235}U from 19.7% to 10.1% for every load, and to burn 1187.71 g ^{239}Pu , 220.04 g ^{241}Am and 96.89 g ^{243}Am per year. These results contribute to enhance the proliferation resistance of radioactive materials.

Key words: MTR-22 MW reactor, neutronic parameters, minor actinide, MCNP4C code.

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إعادة تحميل الاكتينيدات ^{239}Pu و ^{241}Am و ^{243}Am كاصات قابلة للاحتراق في مفاعل البحث MTR-22 MW

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□ ملخص □

إنَّ الهدف الرئيس لهذه الورقة هو إعادة تحميل المفاعل MTR-22MW بالوقود الأساسي $\text{U}_3\text{O}_8\text{-Al}$ الممزوج بالعناصر الاكتينية ^{239}Pu و ^{241}Am و ^{243}Am كاصات نترونية قابلة للاحتراق (الانشطار). ويساعد هذا الإجراء في استهلاك قسم من البلوتونيوم ^{239}Pu المخزن، وتخفيض التأثيرات السلبية البيولوجية الضارة للاكتينيدات ^{241}Am و ^{243}Am في الوسط البيئي المحيط. وجرى حساب الأوساط النترونية للمفاعل MTR-22 MW مثل: التفاعلية ρ_{ex} ، ووثوقية صفائح التحكم (CPSW)، وهامش الإغلاق (SM)، وهامش الإغلاق بوجود صفيحة تحكم خارج قلب المفاعل (SM-1)، ومعامل تفاعلية الأمان (RSF)، والطيف النتروني في المصيدة النترونية المركزية (CNT) وصندوق التشيع المركزي (CIB) وعاكس البيريليوم Be إضافة إلى توزيع الاستطاعة للمفاعل MTR-22 MW باستخدام الكود MCNP4C. وإضافة إلى ذلك، درس تغيير التفاعلية ρ_{ex} للمفاعل MTR-22 MW كتابع لزمن تشغيله. وقد أظهرت النتائج أن المفاعل MTR-22 MW يمكن أن يعمل بالوقود الأساسي $\text{U}_3\text{O}_8\text{-Al}$ الممزوج بالعناصر الاكتينية ^{239}Pu و ^{241}Am و ^{243}Am كاصات نترونية قابلة للاحتراق ضمن شروط الأمان المأخوذة في تصميم المفاعل، وإنَّ تحميل المفاعل MTR-22 MW بالوقود الأساسي $\text{U}_3\text{O}_8\text{-Al}$ الممزوج بالعناصر الاكتينية ^{239}Pu و ^{241}Am و ^{243}Am يؤدي إلى تخفيض الإغناء باليورانيوم ^{235}U من 19.7% إلى 10.1% لكل حمل، وإلى حرق 1187.71 جرام من ^{239}Pu ، و 220.04 جرام من ^{241}Am و 96.89 جرام من ^{243}Am في السنة. وتساهم هذه النتائج في تعزيز مبدأ مقاومة الوفرة من المواد المشعة.

الكلمات المفتاحية: المفاعل MTR-22 MW، الوسطاء النترونية، الاكتينيدات الثانوية والكود MCNP4C.

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Introduction

During nuclear energy generation highly radiotoxic by products such as the Minor Actinides (MAs) is formed, which accumulate in spent fuel. Minor actinides transmutation can take two forms: either the MAs nuclei undergo fission or yields fission products that are shorter lived or capture a neutron and are transmuted into another heavy nuclide. The main MAs that is produced in nuclear reactors are Np, Am and Cm. The problem of the Np, Am and Cm transmutation with the Pu surpluses is a part of the problem of nuclear power ecological safety and proliferation resistance of the fissile materials.

In the last years, several studies were investigating the use of UO₂ and MOX fuel mixed with MAs in Light Water Reactors (LWRs) and fast reactors to both enhance the economy of the fuel cycle and to enhance the proliferation resistance of fissile materials, and to convert their nuclei to non-radioactive nuclei or to nuclei with shorter decay times [1], [2], [3], [4].

The mentioned Mas play an important role in the physics of the LWRs and fast reactors through their effect on the properties of the fuel cycle. Therefore, all experimental and researches carried out in this field focused on using these actinides in LWRs and fast reactors [1], [2], [3].

Some of the new actinide isotopes resulting from fuel burning in the reactor can undergo the fission process as well. This means that re-fueling the MA has its effect not only on the initial reactivity but also on the duration of the fuel cycle. Therefore, it is very important from my standpoint of view to study the effect of these actinides on the neutronic parameters and fuel cycle in Research Reactors (RRs) having medium power such as the MTR-22MW reactor [4], [5].

The main problem of this research

The radioactive properties and the toxicity of the MAs make them very harmful to human beings and the environment. Therefore, many studies to re-cycle the MAs in the fuel cycle of LWRs and fast reactors have been performed [1], [3], [4], [5], [6]. The results of these studies have showed the following:

- The enrichment with ²³⁵U and the required amount of the fuel decrease significantly with increasing the number of fuel assemblies charged with plutonium and MAs,
- Re-cycling of plutonium and MAs together with discharging uranium can reduce the radio-toxicity of discharged heavy metal (HM) waste to become less than that of the loaded uranium, and to reduce the spent fuel for storage and improve the proliferation resistance of the radioactive materials,
- re-cycling MAs and UO₂ fuel together can serve as burnable absorber to reduce the initial excess reactivity.

Since re-cycling RR by MOX mixed with MAs seems not to be investigated, therefore, it is very important to investigate the re-cycle and effect of the MAs on the neutronic parameters of the RRs. This helps to improve the life of the fuel cycle and re-design the RRs using fuels with MAs and using closed fuel cycle in the future; this leads the burn the MAs resulting from these reactors and remove their risks.

The importance of this research

The major benefit of beginning minor actinide recycling in existing thermal reactors and in the RRs having medium power is that it would remove the

long-lived radiotoxic nuclides from the fission product waste, helps in saving energy and enhancing the proliferation resistance of radioactive materials and serving as a burnable absorber to improve the fuel cycle performance. In addition, the advantage of re-cycling MOX fuel with MAs in RRS is to help in:

- reducing the volume of the spent fuel for storage and enhancing the proliferation resistance, especially the length of the fuel cycle in these reactors is short in comparison with that of LWRs which means burning more plutonium and MAs,
- contributing to extend the spreading of RRs which use MOX fuel with minor actinides or original fuel with some fuel elements of MOX fuel mixed with minor actinides,
- increasing plutonium isotopes utilization,
- reducing the enrichment of the fuel with ^{235}U loading in the reactor core,
- transmuting to less hazardous and possibly more useful forms.
- using a closed fuel cycle for RRs in the future.

The main objective of this paper was discussing the effect of the ^{239}Pu , ^{241}Am and ^{243}Am actinides on the neutronic parameters and the fuel cycle of the MTR-22 MW reactor (The Egyptian Second Research Reactor (ETRR-2) as example). This study was carried out changing only the fuel composition. This means, the dimensions of the fuel material, the Fuel Plate (FP), the Fuel Element (FE), the Reactor Core (RC) or any component of the reactor were not changed.

The MCNP4C code [7] and PECOM system [8] were used to provide comparison between the neutronic characteristics for the proposed core re-fueling by $\text{U}_3\text{O}_8\text{-Al}$ OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides and the initial $\text{U}_3\text{O}_8\text{-Al}$ OFEs core.

2. Methodology

2.1 The ETRR-2 reactor

The general characteristics of the ETRR-2 such as: the fuel types, the fuel material, the absorber material and the active zone dimensions are detailed in Table 1 and Table 2 [9], [10], [11], [12].

| Parameter | Weight percentage % | | |
|------------------------------------|---------------------|-----------|-----------|
| | SFE | FE Type 1 | FE Type 2 |
| ^{235}U | 12.377 | 6.598 | 8.398 |
| ^{238}U | 50.450 | 26.894 | 34.230 |
| ^{27}Al | 25.91 | 60.504 | 49.730 |
| ^{16}O | 11.263 | 6.004 | 7.642 |
| Density (g/cm^3) | 4.802 | 3.299 | 3.655 |

Table 1: Composition of the fuel in a typical ETRR-2 core.

Table 2: Specifications of the fuel material, fuel element, absorber material, active zone dimensions and water gap between plates of the ETRR-2 core for the $\text{U}_3\text{O}_8\text{-Al}$ OFEs and the $\text{U}_3\text{O}_8\text{-Al}$ OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides.

| Parameter | Fuel material | |
|---------------------------------|--|---|
| | $\text{U}_3\text{O}_8\text{-Al}$ original fuel | $\text{U}_3\text{O}_8\text{-Al}$ mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides |
| Fuel meat | $\text{U}_3\text{O}_8\text{-Al}$ | $\text{U}_3\text{O}_8\text{-Al} + ^{239}\text{Pu}$, ^{241}Am and ^{243}Am |
| Weight percentage (%) | 100 | $98.2 \text{ U}_3\text{O}_8\text{-Al} + 1.3 ^{239}\text{Pu} + 0.5 (^{241}\text{Am} + ^{243}\text{Am})$ |
| Enrichment ^{235}U (%) | 19.70 | 10.10 |

| | | | | |
|--|-----------------------|------------------|-----------------------|----|
| Initial mass of ^{235}U loading in the reactor core (g) | 6944.5 | 6332.956 | | |
| Saved mass of ^{235}U in the beginning of the load (g) | | 611.543 | | |
| Density of fuel (g/cm^3) | See Table 1 | 4.928 | | |
| Fuel element | | | | |
| Number of the fuel materials | 29 | 29 | | |
| Number of fuel plates in the fuel material | 19 | 19 | | |
| Dimensions of the fuel plate (cm) (length x width x thickness) | 80 x 6.40 x 0.07 | 80 x 6.40 x 0.07 | | |
| Absorber material | | | | |
| Composition | Ag – In – Cd | | B_4C | |
| | Weight percentage (%) | | Weight percentage (%) | |
| | Ag | 80 | ^{10}B | 5 |
| | In | 15 | ^{11}B | 75 |
| | Cd | 5 | C | 20 |
| Density (g/cm^3) | 10.18 | | 2.52 | |
| Active zone dimensions | | | | |
| Active length (cm) | 80 | | 80 | |
| Clad length (cm) | 80 | | 80 | |
| External section of fuel element (cm^2) | 8x8 | | 8x8 | |
| Section in grid to house the fuel element (cm^2) | x 8.18.1 | | x 8.18.1 | |
| Plate thickness (cm) | 0.15 | | 0.15 | |
| Meat thickness (cm) | 0.07 | | 0.07 | |
| Meat width (cm) | 6.40 | | 6.20 | |
| Side plate thickness (cm) | 0.50 | | 0.50 | |
| Side plate width (cm) | 8.00 | | 8.00 | |
| External distance between frames (cm) | 8.00 | | 8.00 | |
| Internal distance between frames (cm) | 7 | | 7 | |
| Cladding material | Al- 6061 | | Al- 6061 | |
| Water gap between plates | | | | |
| of single fuel element (cm) | 0.27 | | 0.27 | |
| of different fuel element (cm) | 0.39 | | 0.39 | |

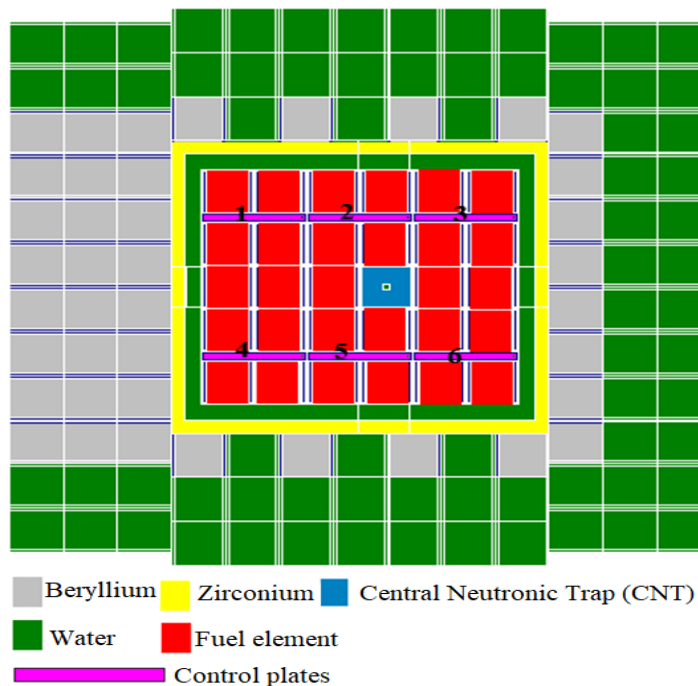


Figure 1: A schematic horizontal cross section of the ETRR-2 core coming

from the MCNP4C code with six control plates.

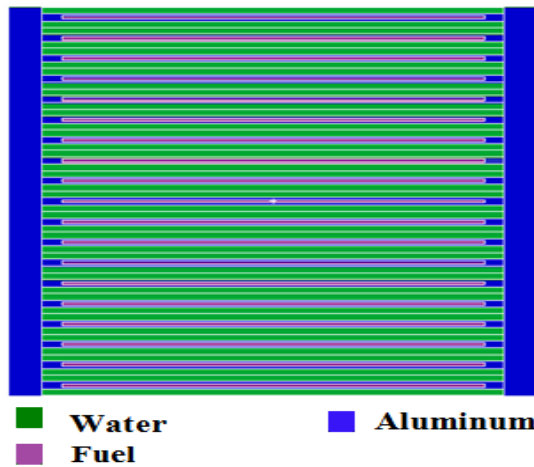


Figure 2: A cross section of the fuel element used in the ETRR-2 by using the MCNP4C code

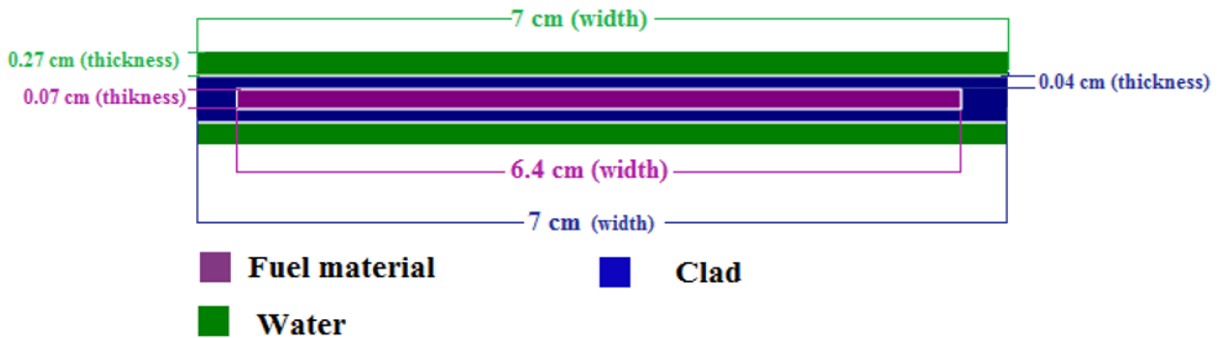


Figure 3: A cross section of the FP by using the MCNP4C code.

2.3 Required conditions for conversion from the U_3O_8 -Al OFEs to the U_3O_8 -AlOFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides as burnable absorbers.

In assessing the practicability of converting the existing ETRR-2 core to the use of U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides, three factors have to be taken into account:

1. The criticality safety parameters of the fuel should not be lower than those for the current design based on the U_3O_8 -Al OFEs,
2. Major modifications to the reactor should not be required, nor should the costs increase,
3. The essential capabilities of the reactor should be conserved requiring the neutron flux in the CNT and the CIB to be conserved,

Therefore, the modifications, which are brought to the ETRR-2 core are related only to fuel material and absorber material in the control plates, where the absorber material Ag-In-Cd alloy was changed into B_4C material for the ETRR-2 core re-fueled by U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides being more effective without changing neither the dimensions nor the sites [13], [14], [15].

The specifications of the fuel material, the fuel element, the absorber material, the active zone dimensions and the water gap between plates of the U_3O_8 -Al OFEs and the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides are listed Table 2. The weight percentage of the ^{239}Pu , ^{241}Am and ^{243}Am actinides used is tabulated in the Table 3 and Table 4 too [16].

| Parameter | Weight percentage % |
|-------------------|---------------------|
| ²³⁸ Pu | 0.001 |
| ²³⁹ Pu | 95.01 |
| ²⁴⁰ Pu | 4.51 |
| ²⁴¹ Pu | 0.47 |
| ²⁴² Pu | 0.01 |

Table 3: Composition of the ²³⁹Pu plutonium.

| Parameter | Weight percentage % |
|-------------------|---------------------|
| ²⁴¹ Am | 60.655 |
| ²⁴³ Am | 39.345 |

Table 4: Weight percentage of the ²⁴¹Am and ²⁴³Am isotopes.

3. Calculation of the power distribution and criticality parameters for the ETRR-2 core refueled by the U₃O₈-Al OFEs and by the U₃O₈-Al OFEs mixed with ²³⁹Pu, ²⁴¹Am and ²⁴³Am actinides.

The MCNP4C code was used to calculate the neutronic and criticality parameters of the ETRR-2 core. The following data:

- nuclear data of the fissile and non-fissile materials were taken from ENDF/BVI libraries,
- 400×10⁶ neutron histories were used to run the code,
- 400 cycles with 50 passive cycles before the active cycles begin,
- initial value of the k_{eff} multiplication factor is 1,
- fuel elements in the ETRR-2 core were used as fission source points,
- fission source locating in the middle of each fuel element,
- $S(\alpha, \beta)$ was used to treat the thermal scattering in beryllium and hydrogen (See MCNP4C code manual [7]).

- energy range of neutrons was divided to three groups as: <0.625eV for thermal, (0.625eV to 5.53KeV) for epithermal and up to 20 MeV for fast.

were taken to run the input file of the ETRR-2 core by the MCNP4C code and calculate the neutronic and criticality parameters, and the power distributions as well as.

3.1 Criticality parameters

Criticality parameters include the calculation of factors such as:

- The effective multiplication factor, k_{eff}, and the corresponding excess core reactivity, ρ_{ex},
- The Shutdown Margin (SM) of the control system,
- The Shutdown Margin with the first control plate is failure (SM – 1),
- The Control Plates Worth (CPsW),
- The Reactivity Safety Factor (RSF). Whereas the calculated value of the RSF was obtained using this relation:

$$RSF = CPsW / \rho_{ex}$$

Criticality calculations were performed by the KCODE criticality card (See the MCNP4C code manual [7]). The obtained results are listed in Table 5 and Table 6.

Table 5: Criticality parameters of the ETRR-2 core re-fueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides as a burnable absorber actinides and design value for U_3O_8 -Al OFEs.

| Parameter | Design value for U_3O_8 -Al OFEs (pcm) with Ag-In-Cd alloy as absorber material in the control plates | U_3O_8 -Al mixed with ^{239}Pu , ^{241}Am and ^{243}Am (pcm) with B_4C alloy as absorber material in the control plates | Differences % | Design value of safety criteria (pcm) |
|-----------------------------------|---|---|---------------|---------------------------------------|
| ρ_{ex} at BOC ⁽¹⁾ | 8220 | 8234.99 ± 6.80 | 0.18 | > 3000 |
| CPsW ⁽²⁾ | 14120 | 14213.22 ± 18.95 | 0.66 | > 1000 |
| CPsW ⁽³⁾ #1 | 10390 | 10630.01 ± 17.70 | 2.26 | >1000 |
| CPsW #2 | 11030 | 10791.73 ± 18.87 | 2.20 | >1000 |
| CPsW #3 | 11050 | 11074.39 ± 18.25 | 0.22 | >1000 |
| CPsW #4 | 10810 | 10656.23 ± 18.84 | 1.44 | >1000 |
| CPsW #5 | 11330 | 10881.21 ± 19.04 | 4.12 | >1000 |
| CPsW #6 | 11290 | 11294.84 ± 19.34 | 0.04 | >1000 |

(1) ρ_{ex} at BOC- is the reactivity at Beginning of Cycle,

(2) (CPsW) - is the Control Plates Worth,

(3) CPsW#1 - is the Control Plates Worth without first control plate (See Figure 1)

Table 6: Calculated values of the RSF, SM and SM-1 of the ETRR-2 core re-fueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides and design value for U_3O_8 -Al OFEs.

| Statement | Design value for U_3O_8 -Al OFEs (pcm) and Ag-In-Cd alloy as absorber material in the control plates | U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am (pcm) and B_4C material as absorber material in the control plates |
|------------------|--|--|
| RSF >1 | 1.7177 | 1.7259 ± 0.0032 |
| SM > 3000 pcm | 5900 | 5978.23 ± 15.45 |
| SM -1 > 1000 pcm | 2170 | 2395.02 ± 6.16 |

3.2 The power distribution parameters

To calculate the power distribution parameters, the Average Thermal, Epithermal and Fast Neutron Fluxes (ATNF, AENF and AFNF) in the CNT, in the CIB and in the Be beryllium reflector were to be calculated. The calculated values of the ATNF, AENF and AFNF in the CNT, in the CIB and in the Be beryllium reflector with reference values are given in Table 7 for the ETRR-2 core fueled by U_3O_8 -Al OFEs and re-fueled by U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides.

Table 7: Calculated values of the neutronic parameters of the ETRR-2 core re-fueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides and reference values for U_3O_8 -Al OFEs.

| Parameter | Reference value | Calculated values for U_3O_8 -Al original fuel | U_3O_8 -Al mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides |
|--|----------------------|--|--|
| ATNF in the CIB ($n/cm^2.s$) $\times 10^{14}$ | 4.230 ^(a) | 4.215 ± 0.017 | 4.276 ± 0.018 |
| AETNF in the CIB ($n/cm^2.s$) $\times 10^{14}$ | | 1.018 ± 0.031 | 1.037 ± 0.032 |
| AFNF in the CIB ($n/cm^2.s$) $\times 10^{14}$ | | 1.533 ± 0.014 | 1.563 ± 0.025 |
| ATNF in the CNT ($n/cm^2.s$) $\times 10^{14}$ | 2.700 ^(b) | 2.700 ± 0.007 | 2.723 ± 0.008 |
| AETNF in the CNT ($n/cm^2.s$) $\times 10^{14}$ | | 0.836 ± 0.008 | 0.865 ± 0.007 |
| AFNF in the CNT ($n/cm^2.s$) $\times 10^{14}$ | | 1.450 ± 0.007 | 1.491 ± 0.009 |

| | | | |
|---|----------------------|---------------|---------------|
| ATNF in the Be reflector (n/cm ² .s)×10 ¹⁴ | 1.000 ^(b) | 0.988 ± 0.014 | 1.003 ± 0.017 |
| AETNF in the Be reflector (n/cm ² .s)×10 ¹⁴ | | 0.227 ± 0.024 | 0.232 ± 0.021 |
| AFNF in the Be reflector (n/cm ² .s)×10 ¹⁴ | | 0.297 ± 0.019 | 0.311 ± 0.025 |

(a) Reference value was taken from reference (Hussein et al., 2011).
 (b) Reference value was taken from reference (Hussein et al., 2011).

References values were taken from the references (www.etr2-aea.org/eg/Data Sheet about MTR-22MW.html and Main Core Data of MTR Reactor.html).

The values of the Power Distribution (PD) in the modified ETRR-2 core were calculated to know whether the re-fueled ETRR-2 core by the U₃O₈-Al OFEs mixed with ²³⁹Pu, ²⁴¹Am and ²⁴³Am actinides causes a big change in the values of the peaking power factors compared with the ETRR-2 core fueled by U₃O₈-Al OFEs, and to predict the temperature change resulting from fission in the FEs.

The power peaking factors are the:

- radial P_{rp} and axial P_{apf} peaking factor of the Hottest Fuel Element (HFE) [17],

- P_{max} maximum and P_{min} minimum power releasing in the FEs,

- P_{avr} average power released in the ETRR-2 core.

The values of the factors P_{rp}, P_{apf} and the P_{avr} are calculated as:

- P_{rp} is the ratio between the P_{max} and the P_{avr},

- P_{apf} is the ratio between the maximum power released in the segment of the HFE and the average power released in the HFE [13], [14], [15],

- P_{avr} is the ratio between the calculated value of the power releasing in the reactor core and the number of the FEs.

The obtained results of the PD and the power peaking factors are given in Figure 4 and Table 8.

Table 8: Power peaking factors of the ETRR-2 core fueled by U₃O₈-Al OFES and re-fueling by the U₃O₈-Al OFEs mixed with ²³⁹Pu, ²⁴¹Am and ²⁴³Am actinides.

| Parameter | U ₃ O ₈ -Al original fuel | U ₃ O ₈ -Al mixed with ²³⁹ Pu, ²⁴¹ Am and ²⁴³ Am actinides |
|---|---|---|
| Maximum power per element P _{max} (MW/element) | 1.050±0.004 | 1.040 ±0.003 |
| Minimum power per element P _{min} (MW/element) | 0.584±0.004 | 0.576±0.004 |
| Average power per element P _{ave} (MW/element) | 0.731±0.003 | 0.755±0.004 |
| P _{max} /P _{min} | 1.798±0.010 | 1.788±0.012 |
| P _{rp} = P _{max} /P _{ave} | 1.432±0.011 | 1.377±0.039 |
| A _{apf} (MW) | 1.428 ±0.010 1.355 ^(a) | 1.424±0.015 |
| Calculated and reliability value of the power (MW) | 21.801±0.004 22 ^(b) | 21.900±0.004 22 ^(b) |

(a) Reference value was taken from reference (Mohamed, M.F., 2011. Analysis of Reactivity Accidents in MTR for Various Protection System Parameters and Core Conditions, Thesis, Master of Science. Faculty of Engineering, Alexandria University, Egypt).

(b) Reliability value of the reactor power.

| | | | | | |
|-------|-------|-------|-------|-------|-------|
| 1 | 2 | 3 | 4 | 5 | 6 |
| 0.670 | 0.584 | 0.660 | 0.674 | 0.635 | 0.774 |
| 0.680 | 0.676 | 0.638 | 0.653 | 0.583 | 0.822 |
| 1 | | 2 | | 3 | |
| 7 | 8 | 9 | 10 | 11 | 12 |
| 0.616 | 0.761 | 0.694 | 0.755 | 0.821 | 1.000 |
| 0.670 | 0.645 | 0.768 | 0.742 | 0.852 | 0.985 |
| 13 | 14 | 15 | 16 | 17 | 18 |
| 0.645 | 0.623 | 0.751 | 0.000 | 0.715 | 0.723 |
| 0.688 | 0.597 | 0.730 | 0.000 | 0.730 | 0.750 |
| 19 | 20 | 21 | 22 | 23 | 24 |
| 0.622 | 0.620 | 0.709 | 0.778 | 0.859 | 1.050 |
| 0.640 | 0.792 | 0.631 | 0.752 | 0.834 | 1.040 |
| 4 | | 5 | | 6 | |
| 25 | 26 | 27 | 28 | 29 | 30 |
| 0.651 | 0.584 | 0.682 | 0.714 | 0.959 | 0.829 |
| 0.693 | 0.586 | 0.679 | 0.727 | 0.877 | 0.733 |

Black color - is the number of the FEs in the ETRR-2 core.
Site 16 - denotes the Central Neutronic Trap (CNT).
Red and blue color - is the calculated values of the PD for the ETRR-2 core refueled by the U_3O_8 -Al OFEs and refueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides, respectively.

 Control plates

Figure 4: Calculated values of the PD for the ETRR-2 core refueled by the U_3O_8 -Al OFEs and re-fueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides.

In the calculation the PD in the reactor core and the power peaking factors, and the ATNF, AENF and AFNF in the CNT, in the CIB and in the Be beryllium reflector for the U_3O_8 -Al OFEs and the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides, the F_4 tally, the FS, the SD and the FM cards in the MCNP4C code were used [13], [14], [15].

5. Study the reactivity as a function of the operation time of the ETRR-2.

The reactivity ρ_{ex} of the ETRR-2 core re-fueling by U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides was studied as a function of the operation time of the reactor for the 10 MW power. This procedure is necessary to safe operational conditions accomplishment in the design of the ETRR-2 core as an initial U_3O_8 -Al OFEs. In this case,

to calculate the reactivity ρ_{ex} we used the PECOM system and taken the main actinides and major fission products [13]. The obtained results are plotted on Figure 5 for the U_3O_8 -Al OFEs and U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides.

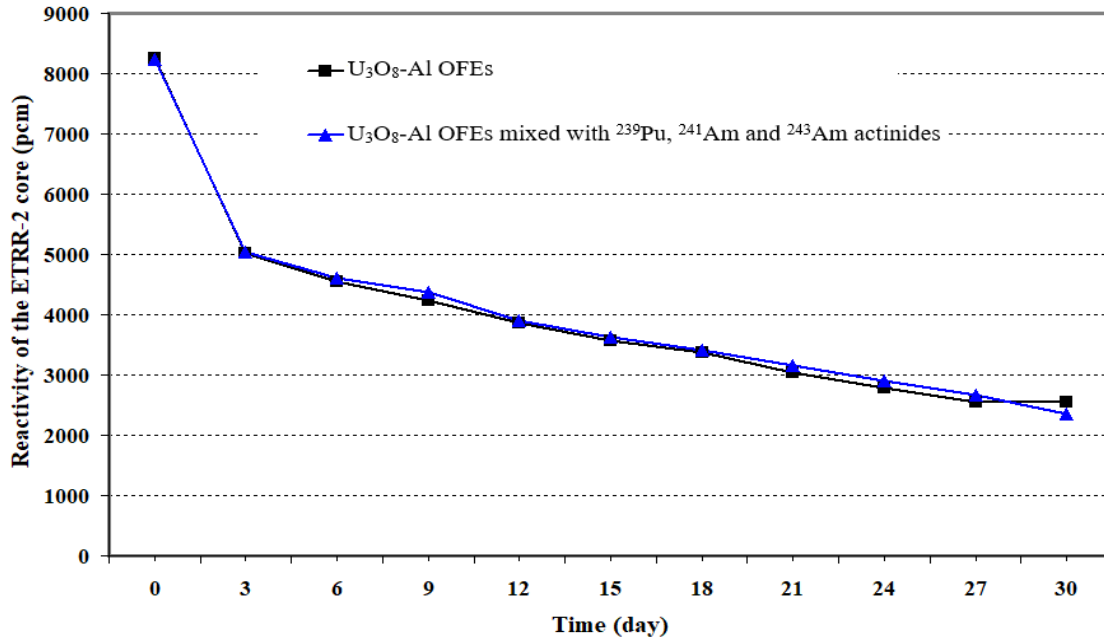


Figure 5: Variation of the reactivity as a function of the operation time of the ETRR-2 core refueled by the U_3O_8 -Al OFEs and re-fueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides.

Results and discussion

From Table 5 it results that the maximum difference between the design values of the CPsW for the ETRR-2 core refueled by the U_3O_8 -Al OFEs and the ETRR-2 core re-fueled by U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides does not exceed 4.12%. This result indicates that the used U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides does not have any negative effect on the criticality safety of the reactor within the conditions given in Table 5.

In the criticality calculations:

- the reactivity ρ_{ex} was calculated from this equation [18]:

$$\rho_{ex} = (k_{eff} - 1)/k_{eff}$$

- the values of the SM and the SM-1 were calculated with the control plates are fully inserted in the ETRR-2 core and the first control plate is failure (See Figure 1), respectively.

- the calculated values of the CPsW were calculated using the following equation:

$$CPsW = \rho_{ex} + SM.$$

Table 6 shows the calculated value of the RSF, SM and SM-1. Noticeably, from this table the differences between the design values of the RSF, SM and SM-1 for the ETRR-2 core refueled by U_3O_8 -Al OFEs and the same parameters for the ETRR-2 core re-fueled by U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides are 0.48%, 1.30% and 9.39%, respectively. This result enhances that the

use of U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides in the ETRR-2 does not break the safety conditions established in the design of the ETRR-2 (See Table 5, Table 6 and Table 7).

From Table 7 it can be seen also that:

1. the difference between the calculated values of the ATNF, AETNF and AFNF in the CIB and in the CNT, and in the Be reflector are 1.423%, 1.832% and 1.919%, 0.845%, 3.352% and 2.749%, 1.495%, 2.155% and 4.501% respectively. This result emphasizes the fact that re-fueling the ETRR-2 reactor by a U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides will not significantly affect the thermal neutron flux in the CNT and in the CIB, and in the Be reflector. This result leads to say that:

- re-fueling the ETRR-2 by a U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides will not affect so much the neutronic parameters of the ETRR-2 core,

- the availability of the scientific applications that were available with U_3O_8 -Al OFEs will still be available now with the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides,

- The ATNF in the CNT and in the CIB is higher than the 2.0×10^{14} n/cm²s. This value is sufficient to produce the radioisotopes such as: ^{99}Mo , ^{14}C , ^{32}S , ^{51}Cr , ^{60}Co , ^{89}Sr , ^{153}Sm , ^{169}Yb , ^{170}Tm , ^{192}Ir [1], [19], [20]. It results from Table 8 that the maximum difference between the calculated values of the parameters P_{max} , P_{min} , P_{ave} , P_{rpf} , A_{apf} , P_{max}/P_{min} is about 3.84% for the ETRR-2 core refueled by the U_3O_8 -Al OFEs and re-fueling by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides. This result leads to say that re-fueling of the ETRR-2 reactor by the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides does not create a big change in the power peaking parameters.

Figure 5 shows the reactivity of the ETRR-2 core as a function of the operation time for the U_3O_8 -Al OFEs and the U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides. Noticeably, there is a good agreement between these curves. Analysis of these curves shows that the,

1. ^{239}Pu is compensating the reactivity loss that results from the absorption of neutrons by the ^{241}Am and ^{243}Am actinides,

2. ^{242}Am forms as a result of the absorption neutrons by the ^{241}Am (See Figure 6) that has high cross section of fission (2100×10^{-24} cm²),

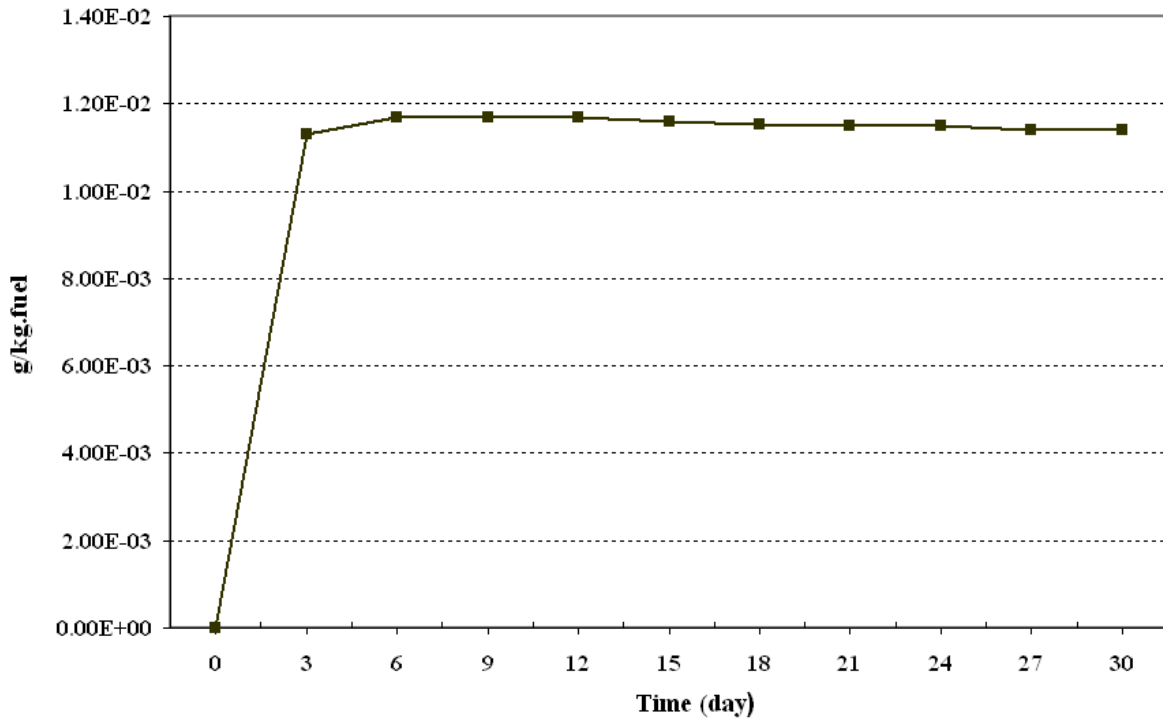


Figure 6: Concentration of the ²⁴²Am for the ETRR-2 core re-fueling by the U₃O₈-Al OFEs mixed with ²³⁹Pu, ²⁴¹Am and ²⁴³Am actinides.

3. ²⁴²Am increases the fuel burn-up beginning on day15 (See Figure 6) as a result of fissioning the ²⁴²Am nuclei. This fact can be used:
- to improve the fuel cycle in RRs,
 - to help the conversion from an open to a closed fuel cycle in the future,
 - to lower the ²³⁵U mass in the fuel or decrease the number of the FEs which are required for a reactor which comes out from enrichment reduction of uranium (See Table 9)
 - to compensate the reactivity loss at the end of the fuel cycle.

Table 9: Saved and burning mass of the ²³⁵U and ²³⁹Pu, ²⁴¹Am and ²⁴³Am per year.

| Type | Mass of actinide per year (g) |
|-----------------------------------|-------------------------------|
| Saved mass of ²³⁵ U | 8561.616 |
| Burning mass of ²³⁹ Pu | 1187.714 |
| Burning mass of ²⁴¹ Am | 220.040 |
| Burning mass of ²⁴³ Am | 96.890 |

Table 9 shows the saved and burnt mass of the ²³⁵U, ²³⁹Pu, ²⁴¹Am and ²⁴³Am actinides per year. From this table it can be seen that the burnt mass of the ²³⁹Pu, ²⁴¹Am and ²⁴³Am actinides per year is small, but this result is very important because it is open this field to:

- increase researches and enhance the development of a distinguished fuel cycle in RRs by using the minor actinides,
- help in enhancing the proliferation resistance of the radioactive materials.

- prove the possibility of designing RRs with high or medium power with closed fuel cycles in the future,
- burn a bit of the ^{239}Pu in the MTR reactors,
- use of ^{239}Pu , ^{241}Am and ^{243}Am actinides as additional sources of energy,
- contribute in reducing the risks of these actinides in the environment.

Conclusion

The U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides was proposed as a new fuel of the ETRR-2 reactor. Results indicate that the re-fueling of the ETRR-2 reactor by a U_3O_8 -Al OFEs mixed with ^{239}Pu , ^{241}Am and ^{243}Am actinides will not have any negative effect on the criticality and neutronic characteristics of the ETRR-2 reactor. Using the ^{239}Pu , ^{241}Am and ^{243}Am actinides leads to reduce the ^{235}U mass loaded in the ETRR-2 core. I hope these results will encourage the researchers in this field to development the fuel cycle in the RRs.

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