MEASUREMENT OF THE ENERGY SPECTRUM OF THE NEUTRONS INSIDE THE NEUTRON FLUX TRAP ASSEMBLED IN THE CENTER OF THE REACTOR CORE IPEN/MB-01

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ABSTRACT

This paper presents the neutron energy spectrum in the central position of a neutron flux trap assembled in the core center of the research nuclear reactor IPEN/MB-01 obtained by an unfolding method. To this end, have been used several different types of activation foils (Au, Sc, Ti, Ni, and plates) which have been irradiated in the central position of the reactor core (setting number 203) at a reactor power level of 64.57 ± 2.91 watts . The activation foils were counted by solid-state detector HPGe (gamma spectrometry). The experimental data of nuclear reaction rates (saturated activity per target nucleus) and a neutron spectrum estimated by a reactor physics computer code are the main input data to get the most suitable neutron spectrum in the irradiation position obtained through SANDBP code: a neutron spectra unfolding code that use an iterative adjustment method. The adjustment resulted in $3.85 \pm 0.14 \ 10^9 \text{ n} \cdot \text{cm}^{-2} \text{ s}^{-1}$ for the integral neutron flux, $2.41 \pm 0.01 \ 10^9 \text{ n} \cdot \text{cm}^{-2} \text{ s}^{-1}$ for the thermal neutron flux, $1.09 \pm 0.02 \cdot 10^9$ n \cdot cm⁻² s⁻¹ for intermediate neutron flux and $3.41 \pm$ $0.02 \ 10^8 \text{ n} \cdot \text{cm}^{-2} \text{ s}^{-1}$ for the fast neutrons flux. These results can be used to verify and validate the nuclear reactor codes and its associated nuclear data libraries, besides show how much is effective the use of a neutron flux trap in the nuclear reactor core to increase the thermal neutron flux without increase the operation reactor power level. The thermal neutral flux increased 4.04 ± 0.21 times compared with the standard configuration of the reactor core.

1. INTRODUCTION

Several nuclear reactor physics parameters are obtained through the spectrometry gamma of targets irradiated inside the research reactor core. This is the case of the nuclear reaction rates measured by activation foils when are known parameters as irradiation time, time of counting, wait time for counting (decay), efficiency of the counting system, nuclear cross section, *etc*. Thus it is possible to determine the neutron flux in the very place where the foils are irradiated. The reaction rates depend on several factors. The most important one is the

cross section magnitude of the different materials (activation detectors) that cover different energy regions of the neutron spectrum. The experimental values obtained can be used to estimate important reactor physics parameters such as neutron flux (thermal, intermediate and fast), extension of the asymptotic region, spectral indices (cadmium ratio), buckling, etc. These experimental parameters when compared to the same parameters calculated by reactor physics codes enable check the accuracy and precision of the different calculation methodologies and their nuclear data libraries associated.

The knowledge of the energetic and spatial neutron flux distribution in the reactor core is very important because enable to estimate with good precision several of the reactors physics parameters such as nuclear reaction rates, fuel burn up and safety parameters like temperature distribution, peak factors, reactivity.

The neutron energy spectrum of research reactors can be obtained by nuclear reaction rates induced in activation detectors (metallic foils) irradiated inside of the reactor core always in the same position and the same reactor operational conditions such as power level, control rods positions, water temperature (moderator), etc [1,2]. For this purpose may be used unfolding codes such as SANDII, SANDBP [1], SPECTRA [3] and others. These computational codes work iteratively and the process start via a normalized initial spectrum usually calculated by a reactor physics code with an associated data libraries. In each iteration step an initial normalized neutron spectrum is modified by the "best fit" between the measured foils reaction rates and the computed activities. The process stop when occur a set of conditions defined initially. This study used the SANDBP which is a computer code developed by the Technical University of Budapest, Hungary. This code is a modified version of the code SAND II [4] and the main difference is that uncertainties of saturation activities per target nucleus are considered. Thus, it is possible to obtain the neutron energy spectrum by groups and its uncertainties after running the code SANDBP. The input neutron energy spectrum of the irradiation site is estimated by reactor physics codes. The adjustment process is iterative and step-by-step the flux values in the groups are obtained by the minimal deviation of the ratio between the nuclear reaction rates calculated in current iteration and the experimental values. By this process the initial neutron spectrum is modified in the several energy ranges where the detectors are activated. In this work, the initial neutron spectrum was calculated using the MCNP-4C code [5] in 640 energy groups in the same position in which activation foils were irradiated (central position of the neutron flux trap) computed in 250 cycles, 1.000.000 stories per cycle and an average error of 1% in each flux group. The nuclear data library used in SANDBP is the ENDF / BVII.0.

A fundamental activity of the research reactors is the production of radioisotopes. To maximize the thermal neutron flux responsible for most of the radioactivity induced in the irradiated material, use is made of so-called neutron flux traps that are essentially spaces within the core of the reactor filled with a moderator with a very high moderation ratio that in our case is light water, but could be other materials such as beryllium or heavy water. This paper investigates the thermalization of the neutrons energy spectrum and the increased of the thermal neutron flux obtained from the use of neutron flux trap.

2. EXPERIMENTAL METHODOLOGY

The IPEN/MB-01 is a zero power nuclear reactor designed to measure a variety of nuclear reactor physics parameters with more than 3,000 operations realized where were done several experiments and some of them have international status of the OECD / NEA (benchmarks). The standard reactor core consists of fuel assembly (26x28 rods rectangular configuration). This configuration has 680 fuel elements (rods) and two (2) safety bars and two (2) control bars. Each control bars and safety bars contains 12 rods of materials with high neutron absorption cross section. The safety rods are made by Boron Carbide B4C and the control rods are made by a mixture of chemical elements Ag-In-Cd. The fuel is UO2 enrichment of 4.3%. The fuel element is clad with 304 stainless steel. A complete description of IPEN/MB-01 can be viewed through a wide literature [5].

Research reactors can be used for the production of radioisotopes and radio-pharmaceuticals of broad social application in industry, agriculture and medicine. The most important radioactive elements produced are obtained from materials of very high capture cross section in the thermal neutron spectrum region. So if the magnitude of the thermal neutron flux increase this results in the production of radioactive elements with higher activities. There are two ways to increase the thermal neutron flux in a research reactor: increase the operating power or assembly neutron flux trap inside the core. In this work were assembled several core configurations containing inside a neutron flux trap, and the most effective was the configuration number 203, which are sketched in Fig.1. To assemble this configuration was withdrew a total of 32 fuel rods from the center reactor core. To minimize the neutron leakage from the system was assembled a hexagonal cross section core (approximated cylindrical cross section). See Fig. 1.

2.1 Irradiation Conditions

To obtain the neutron energy spectrum in the center of the IPEN/MB-01 reactor core, five (5) different foils have been irradiated with their respective eight (8) nuclear reactions. The characteristics of the activation foils irradiated are shown in the Table 1.

Irradiated		Irradiation	Cross	Half-	Foil	Nominal
Activation	Nuclear Reaction	Time	Section	Life	Mass	Thickness
Foil		(hours)	(barns)	(hours)	(g)	(cm)
1%Au-Al	$^{197}Au(n.\gamma)^{198}Au$	1	99.57	64.56	0.02394	0.02
1%Au-Al *	$^{197}Au(n,\gamma)^{198}Au$	1	15630**	64.56	0.02478	0.02
Sc	$^{45}Sc(n,\gamma)^{46}Sc$	2	27.21	2011.92	0.01843	0.00127
Sc*	$^{45}Sc(n,\gamma)^{46}Sc$	2	11.18**	2011.92	0.01848	0.00127
Ti	$^{47}\text{Ti}(n,p)^{47}\text{Sc}$	2	1.76.10 ⁻²	80.16	0.14463	0.00254
Ni	⁵⁸ Ni(n,p) ⁵⁸ Co	2	$62.4.10^{-3}$	1728	0.28500	0.00254
In	115 In(n,n') 115m In	1	183.10 ⁻³	4.50	0.04046	0.00127
In*	115 In(n,n') 115m In	2	183.10-3	4.50	0.03926	0.00127
* cadmium covered (0.5 mm Thickness): ** Resonance integral						

Table 1: Foils Data: Nuclear reactions, irradiation time, cross section, half-life of formatted radionuclide, mass and thickness.

cadmium covered (0.5 mm Thickness); Resonance integral The time of irradiations were estimated from knowledge of the half-lives of the different radionuclides and the magnitude of its cross section. Some foils were covered by cadmium boxes to prevent interference of thermal neutron flux. Each irradiation was made at power of 64.7 watts and in the same experimental conditions (control rods positions, positions Foils). The foils were placed in the central position of the core by means of an articulated device consisting of an acrylic support. The foils are fixed in acrylic support and then the apparatus was inserted into the space between channels 14 to 15 shown in Fig.1.

The Nuclear Channel number 10 (B-10 detector) is the detector further away from the core and is on east side of the reactor core (~40 cm). Nuclear Channel 10 has been used to normalize the little difference of the operation power level between each irradiation. The control rods always were the same positions (90.2 % withdrawn) during all irradiations to avoid neutron flux disturbance.



Figure 1: Reactor Core (Configuration Number 203) – View of the Neutron Flux Trap at the Central Position and irradiation position of the activation foils.

2.2 Gamma Spectrometry

The activation foils were sent to the laboratory for gamma spectrometry to determine radioactivity induced after irradiation. This parameter is proportional to the saturation activity of the target nucleus. A high pure germanium detector (HPGe) with efficiency of 45% percent was used to make the gamma spectrometry. Fig.2 shows the radioactive decay curve of an infinity dilute foil compound with 1% Au-198 and 99% Al.

Figure 2 – Decay Curve to 1% Au (Au-198) and 99% Al Foil

The net counting (C) is obtained by gamma ray spectrometry of activation foil irradiated in the center of the core. The counting of photopeaks gamma emitted by foils made possible to obtain the nuclear reaction rate which is numerically equal to the saturation activity. This requires prior knowledge of the counting efficiency for the energy of the gamma photopeak range that is counting, as well as the irradiation times: (ti) = irradiation time, (te) = wait time and (tc) = counting time. Equation (3.1) calculates the saturation activity (A^{∞}).

(3.1)
$$A^{\infty} = \frac{\lambda \cdot e^{\lambda \cdot t_e} \cdot (C - BG)}{\varepsilon \cdot I \cdot (1 - e^{-\lambda \cdot t_i})(1 - e^{-\lambda \cdot t_c})} \cdot CF$$

CF is a total correction factor and correspond to three different factors: normalization factor that correct little fluctuation power level between the several irradiations, ramp factor that diminish the value of detector activity obtained during ramp power elevation until the steady state radiation level and self-absorption factor. The saturated activity per target nucleus (A_s) is given by the equation (3.2), where (W) is the atomic weight of the target nucleus, m the mass foil, (f_{iso}) the isotropic fraction of the target nucleus, and (N_A) the Avogadro number.

$$A_{s} = \frac{A^{\infty} \cdot W}{m \cdot N_{A} \cdot f_{iso}}$$
(3.2)

3. ANALYSIS OF THE RESULTS

The experimental saturation activities and the calculated activity obtained from SANDBP code after three (3) iterations for each foil are given in Table 2. The total standard deviation is 5.44 % between the measured activities and those adjusted by the most appropriated neutron spectrum obtained through SANDBP code. Table 3 presents the neutron fluxes values given by SANDBP code at different ranges of energy. Fig. 3 shows a comparison between the adjusted spectrum computed by SANDBP code and the initial spectrum calculated by MCNP-4C code.

Table 2: Measured and calculated saturated Activity per target nucleus after 3 iterationsby SANDBP code. The MCNP-4C code has been used how input spectrum.

Foil	Daughter Nuclide	Gamma Photo- peak (keV)**	Saturated Measured Activity*** (DPS/Nucleus)	Calculated Activity by SANDBP Code (DPS/Nucleus)	
Au-Al	¹⁹⁸ Au	411.8	$3.011.10^{-13} \pm 2.38\%$	$2.919.10^{-13}$	
Au-Al*	¹⁹⁸ Au	411.8	$8.626.10^{-14} \pm 2.38\%$	8.850.10 ⁻¹⁴	
$\operatorname{Sc}^{\Delta}$	⁴⁶ Sc	889.20	$4.978.10^{-14} \pm 5.22\%$	5.611.10 ⁻¹⁴	
Sc ^Δ ,*	⁴⁶ Sc	889.2	$1.076.10^{-15} \pm 2.34\%$	1.01210 ⁻¹⁵	
Ti	⁴⁷ Sc	159.4	$8.294.10^{-17} \pm 4.62\%$	8.12610^{-18}	
Ni	⁵⁸ Co	810.8	$4.917.10^{-16} \pm 5.22\%$	$4.787.10^{-16}$	
In	^{115m} In	336.2	$8.021.10^{-17} \pm 5.54\%$	8.238.10 ⁻¹⁷	
In*	^{115m} In	336.2	$8.321.10^{-17} \pm 5.53\%$	8.143.10 ⁻¹⁷	
Total Standard Deviation				5.44% (2 Sigma)	
* and minimage variable $**$ not nearly counts during the same spectrometry $^{\Lambda}$ Corrected to self shielding					

* cadmium covered; ** net peak counts during the gamma spectrometry; $^{\Delta}$ Corrected to self-shielding effect.

Energy (MeV)	Integral neutron Flux $(n \text{ cm}^{-2}\text{s}^{-1})$	Standard Deviation
>1.10 ⁻¹⁰	3.8472 10 ⁹	0.35%
< 0,20.10 -6	2.2753. 10 ⁹	0.51%
< 0,56.10 ⁻⁶	2.4139 10 ⁹	0.47%
> 0,1	6.1479 10 ⁸	0.51%
>0,4	4.954 10 ⁸	0.42%
>0,5	$4.0905 \ 10^8$	0.41%
>1,0	3.410 10 ⁸	0.39%

Table 3 – Neutron Flux values obtained by SANDBP code [6]

Figure 3. Neutron Spectrum at the Central Position of the IPEN/MB-01 obtained by SANDBP Code (3 iterations-) to 50 energy groups [6] compared with calculate spectrum by MCNP-4C collapsed in 50 groups .

The most appropriated energy neutron spectrum adjusted by SANDPB in 50 groups is showed in Fig. 4.

Figure 4: Energy spectrum adjusted by SANDPP in 50 groups of energy obtained with initial spectrum calculated by MCNP-4C in 640 groups of energy

The neutron flux collapsed in 3 energy groups, thermal (below 0,56 eV), intermediate (0.56eV until 0,5 MeV) and fast (above 0,5 MeV) at the central position of the configuration core number 203 (Fig.1) is given below.

 $\Phi \text{ (Thermal)} = (2.4139 \pm 0.0142) \ 10^9 \text{ n cm}^{-2}.\text{s}^{-1}$ $\Phi \text{ (Intermediate)} = (1.0943 \pm 0.0552) \ 10^9 \text{ n cm}^{-2}.\text{s}^{-1}$ $\Phi \text{ (Fast)} = (3.41 \pm 0.02) \ 10^8 \text{ n cm}^{-2}.\text{s}^{-1}$

Table 4 shows a comparison between the thermal neutron flux in the central position of the flux trap and the rectangular standard core configuration in the same position and same power level .

The thermal neutron flux increased by 4.04 ± 0.21 times. This result can be view in more detail at Mura [6].

Table 4: Thermal neutron flux obtained by SANDBP code at central position of thereactor core to two different core configuration of the IPEN/MB-01 reactor operating atthe same power level.

Thermal Neutron Flux Adjusted at Experimental values by SANDBP Code [1] (n cm ⁻² .s ⁻¹) at Central position of the neutron flux trap - Core configuration 203	Thermal Neutron Flux Adjusted at Experimental values by SANDBP Code [1] (n cm ⁻² .s ⁻¹) at Central Position: of the rectangular core configuration 28x26 fuel rods	Increase of the Thermal neutron Flux in the same position and power level (64,57 watts) by utilization of the neutrons flux trap
$(2.41 \pm 0.01) \ 10^9$	$(5.97 \pm 0.30) \ 10^8$	(4.04 ± 0.21)

4. FINAL COMMENTS

The total standard deviation between the measured activities and those adjusted by the most appropriated neutron spectrum obtained through SANDBP code is 5.44 % . Only 3 iterations were used in the unfolding code SANDBP to obtain the neutron energy spectrum. The assembling of a neutron flux trap in the IPEN/MB-01 reactor core center (configuration number 203) generate an increase of about 304% percent in the thermal neutron flux when compared with that obtained using the standard rectangular configuration at the same power level. This work shows the importance to assemble a neutron flux trap in a region of the reactor core to increase the thermal neutron flux without increase the standard reactor power level and consequent increase the production of radioisotopes to medicinal and industrial purposes. The next work step is to investigate the increase of the thermal neutron flux caused by assembling a box with heavy water inside the neutron flux trap using the same core configuration 203.

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