Measurement of Uranium Concentration In different source (Milk powder) and countries By Using CR-39 Plastic Track Detector Eman Ibrahim Awad Department of physics, College of science, Kerbala University, Kerbala, Iraq

Abstract

The aim of this research is to measurement the alpha – emitter in milk powder samples taken from different source and countries using fission fragment track registration. The study includes collecting (13) sample.

The nuclear track technique by CR-39 plastic track detector sheets is used as detector in this research.

The nuclear reaction used as a source of nuclear fission fragments is U-235 (n , f) obtained by the bombardment of Uranium – 235 in sample with thermal neutrons from (Am - Be) sours which has flux of ($5*10^3$ n . cm⁻² .s⁻¹) the irradiation time was (7) days .

The concentration values were calculated through the comparison with standard samples which were prepared .

The results give ranged between lowest degree in type Al- Modhesh made in Sultanate oman and highest type Amies made in French as follows (0.4583 - 1.8273 ppm).

A Comparison of results to permissible limits of Uranium in milk also obtained.

Keywords

Depleted Uranium, CR-39 plastic, chemical etching, Solid State Nuclear Track Detectors (SSNTDs), Milk powder.

قياس تركيز اليورانيوم في مختلف المصادر (الحليب الباودر) والبلاد باستخدام كاشف الأثر البلاستيكي CR

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الخلاصة
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يهدف البحث الحالي إلى قياس بواعث (مطلقات) ألفا في نماذج الحليب الاصطناعي . الباودر)المأخوذة من مختلف المناشيء لأقطار مختلفة ، باستعمال تسجيل أثار شظايا الانشطار) . شملت الدراسة جمع ثلاثة عشر نموذج

. إنّ تقنيةَ عد الآثار النوويةِ للكاشف البلاستيكي هي المستعملةُ في هذا البحثِ ، قِيَم التركيزَ حُسِبتْ خلال المقارنةِ ' بالعينات القياسية المحضرة

تم تحديد تركيز اليورانيوم للنماذج المراد دراستها باستخدام عد الآثار النووية بواسطة كاشف الأثر وذلك النووي الناتجة من التفاعل U(n,f) بتسجيله لأثار شظايا الانشطار

الناتجة من قصف النماذج بنيوترونات حرارية منبعثة من المصدر النيوتروني

 10^3*5 . والمدة سبعة أيام بفيض نيوتروني ، $10^{-2}\,\mathrm{s}^{-1}$ ولمدة سبعة أيام ($10^{-2}\,\mathrm{s}^{-1}$

لَقد تم تحديد تركيز اليورانيوم بالمقارنة مع الحسابات المعتمدة الني تعطيها النماذج القياسية التي تم تحضيرها حيث كانت النتائج تتراوح بين أقل تركيز نوع المدهش صنع في سلطنة عمان وأعلى تركيز هو آميس صنع في : فرنسا وكانت كالآتي

(0.4583 - 1.8273 ppm)

أمن مقارنة النتائج مع المحددات العالمية تبين أن جميع النماذج هي ضمن المحددات العالمية

Introduction:

There are three isotopes of uranium present in nature , they are uranium -238 , uranium – 235 and uranium – 234 , with the abundance of 99.28 % , 0.71 , and 0.0058 % , respectively [1 , 2] . Nuclear power generation and nuclear weapon production calls for the enrichment of uranium (increase the content of uranium – 235) of up to 90 % . The enrichment process produce nuclear waste containing 0.2 - 0.3 % uranium – 235 and 99.8 % of uranium – 238 [3 , 4] U^{238} (half life of 4.468 x 10^9 y and specific activity of 12.4 Bq . mg⁻¹)

 U^{235} (half life of 7.038 x 10^8 y and specific activity of 80.0 Bq . mg⁻¹)

 U^{234} (half life of 2.445 x 10^5 y and specific activity of $2.3x10^5$ Bq .mg⁻¹) .

The major radionucclides present in U-238 decay chain are givin in table(1).

Natural Uranium emits ionizing alpha, beta, and gamma radiation [3].

The military establishment in the United States found depleted uranium very useful in manufacturing ammunitions against armer targets due its high density,pyrophorisity, and availability in very large quantities and cheapness [5]. Although depleted uranium ammunition were available sice the Seventies of the 20^{th} century [6] , they were used for the 1^{st} time in open environments against Iraq in second Gulf war in 1991 [7] . Consequently , awide spread environmental contamination with depleted uranium had occurred causing elevations in exposure rates and uranium – 238 decay products in air , food , soil , and water adjacent to the destroyed target [8] .

Table (1) show U-238 decay chain [9]

Sample	Half - life	Radioactive Emissions
	Sample	Sample Half - life

Uranium	U-238	4.5*109	y.	α
Thorium	Th-234	24.1	d.	β,γ
Proactinium	Pa-234	6.75	h	β,γ
Uranium	U-234	$247*10^3$	y.	α,γ
Thorium	Th-230	$80*10^3$	y.	α,γ
Radium	Ra-226	1600	y.	α,γ
Radon	Rn-222	3.823	d.	α, γ
Polonium	Po-218	3.05	m.	α, β
Lead	Pb-214	26.8	m.	β,γ
Astatine	At-218	2	S.	α
Bismuth	Bi-214	19.7	m.	β,γ
Polonium	Po-214	1.64*10-4	S.	α,γ
Thallium	Ti-210	1.3	m.	β,γ
Lead	Pb-210	21	y.	β,γ
Bismuth	Bi-210	5.01	d.	β
Polonium	Po-210	138.4	d.	α
Thallium	Ti-206	2.19	m.	β
Lead	Pb-206	stable		•••••

Goal of the Research

The main goals of this research is to measurement the Depleted Uranium in milk powder samples taken from different source to different country using fission fragment track registration.

In this study CR-39 plastic track detector used due to its high sensitivity and availability in Iraq, using Can technique the most widely used technique.

Experimental Details

1. Milk Samples

The study includes collecting (12) samples of powder milk which were collected from different source to different country as follows in table (2), hence taken powder sample measurement uranium concentration (concentration unknown) and to press on form pellet (weight 0.5g, thickness 1.5 mm, diameter 2 cm).

2. Track Detector

Sheets of 250 μ m thick CR-39 plastic (supplied by per shore molding LTD Co.UK) were used . these sheets were cut into small piece each of 1cm x 2cm area . then stored at normal laboratory conditions [10] .

3. Etchant solution

Sodium hydroxide solution with (6.25~N) has been used for the etching process . This process was performed at ($70~^{\circ}C$) .

4. Water Bath

An etching bath of the type (Memmert) Germany was used .It includes a thermostat which can be operated over arrange of ($20-120~^{\circ}\text{C}~$) . A distilled water was used as the bath liquid . The accuracy of regulation of temperature was better than $\pm\,0.1~^{\circ}\text{C}$.

5. Optical Microscope

The counting of all the chemically etched tracks was carried out using an optical microscope (Bausch & Lamb , Japan) . It is capable of giving magnifications of up to $x\ 400$.

6. Neutron Irradiation Source

The nuclear reaction used as source of nuclear fission fragments is U-235 (n,f) obtained by the bombardment of Uranium -235 in sample with thermal neutrons from (Am - Be) source which has flux of ($5*10^3$ n.cm⁻².s⁻¹) the irradiation time was (7days) as shown in figure (1) [11] .

7. Standard Samples

The standard geological samples of Uranium concentration recommended by IAEA were prepared . The pellet were covered with (CR-39) on both sides and enclosed in aluminum capsules and put in aplate of paraffin wax at adistance of (5cm) from the neutron source (Am-Be) . was measured by comparison between track densities register on the detectors from the relation : [12 , 13]

$$C_{ppm}(Sample) = \frac{\rho(Sample)}{\rho(S \tan dard)} * C_{ppm}(S \tan dard) \dots (1)$$

Were (p) is the induced fission track density and C_{ppm} denotes the Uranium content [14, 15, 16], Figure (2) show this relation.

$$C_{ppm}(Sample) = \frac{\rho(Sample)}{Slope} \dots (2)$$

Table (2) the producer name and the origin

Country of origin	Producer name	Sample code
SULTANATE OF	Al-Modhesh	S_1
OMAN	7 II Wodilesii	
EGYPT	Pure	S_2
KSA	Crystal	S_3
FRENCH	NIDO	S_4
FRENCH	Lacto	S_5
INDONISIA	AL- Aham	S_6
OMAN	Al-Atiaf	S_7
SULTANATE OF	Fresh	S_8
OMAN		
FRENCH	Dielac	S_9
RUSIA	Dolce	S_{10}

Measurement of Uranium Concentration

HOLANDA	Two Cows	S ₁₁
INDONISIA	Milgro	S_{12}
FRENCH	Amie [,] s	S_{13}

Result and Discussion

Table (3) show the results of uranium concentration in (ppm) in the (13) sample (milk powder).

By comparison between the results obtained and the permissible limit of EPA for milk, there are two samples of higher level, which are samples (13) with (1.8273) ppm and sample (12) with (1.4841) ppm.

Conclusions:

The low uranium concentration in milk sample measurement in this work (Al – Moudhesh , made in sultanate Oman) while high concentration in ($\,$ Amie's , made in French) .

Table (3) Track density (P) and Concentration of U-238

Sample code	No. of track	Track density (P) (track/mm²)	Concentration Of Uranium (ppm)*10 ⁻⁷
S_1	0.1059	169.59 ± 5.00	0.4583
S_2	0.1441	230.56 ± 7.30	0.6231
S_3	0.1783	285.34 ± 3.00	0.7711
S_4	0.2085	333.67 ± 6.53	0.9018
S_5	0.2314	370.24 ± 1.05	1.0006
S_6	0.2488	398.19 ± 4.16	1.0761
S_7	0.2649	423.93 ± 3.60	1.1457
S_8	0.2690	430.41 ± 4.60	1.1632
S_9	0.3129	500.77 ± 4.44	1.3534
S_{10}	0.3239	518.33 ± 0.15	1.4009
S_{11}	0.3320	531.34 ± 0.25	1.4360
S_{12}	0.3432	549.12 ± 1.46	1.4841
S ₁₃	0.4225	676.12 ± 0.83	1.8273

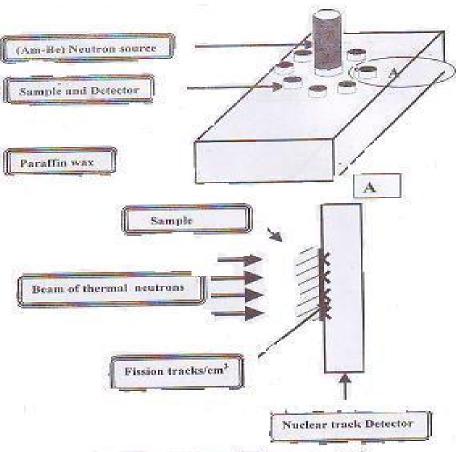
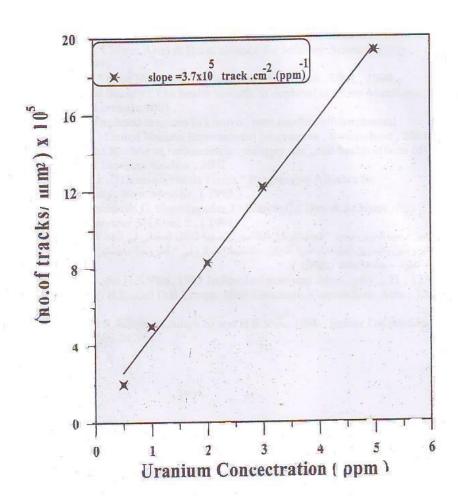


Figure (1) The irradiation of the detectors and samples to the neutron sours

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re (1) the density of track and Uranium concentration in the standard ple

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