

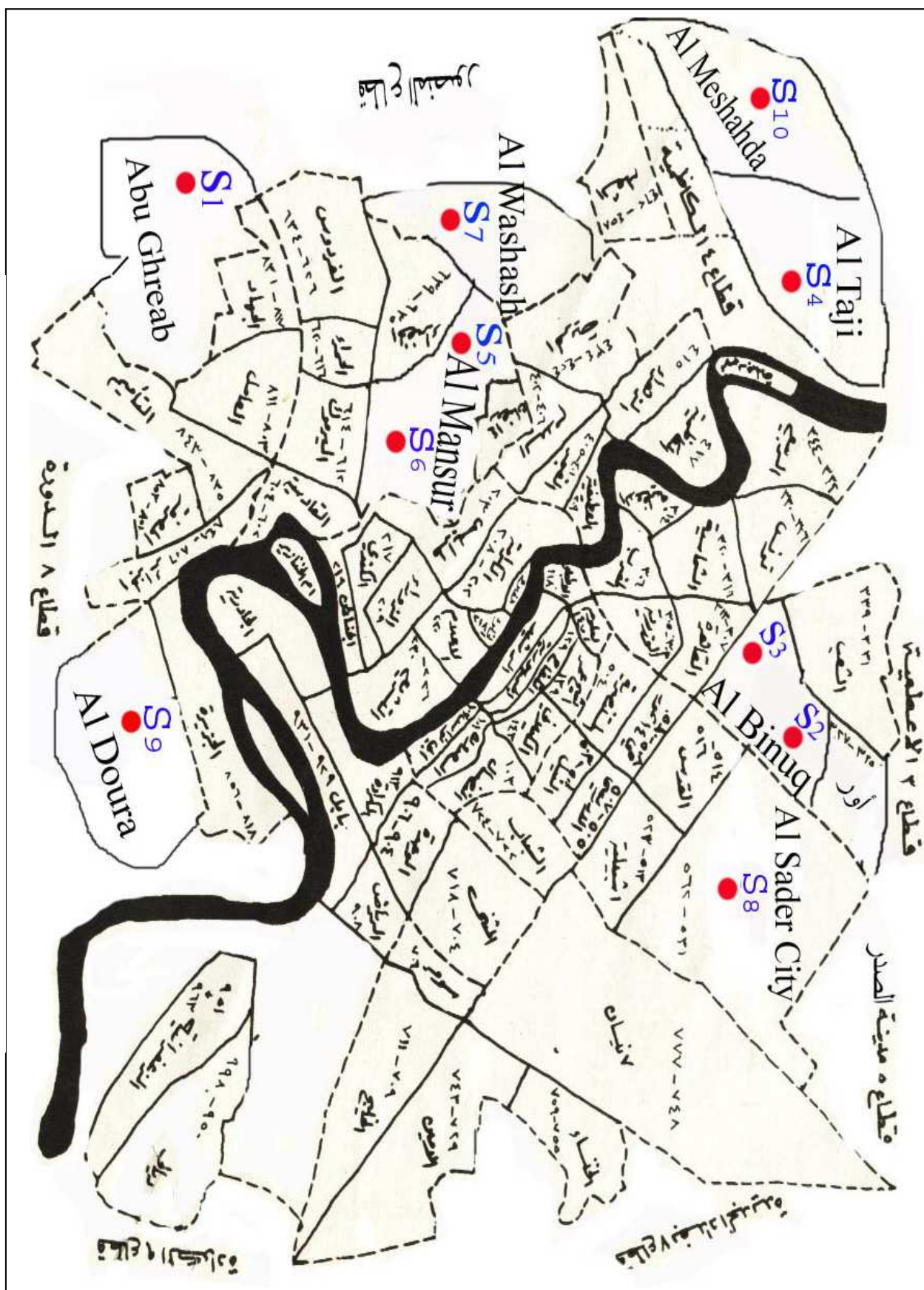


Fig. (3.1) The locations of collected soils samples from different sites in Baghdad city [92].

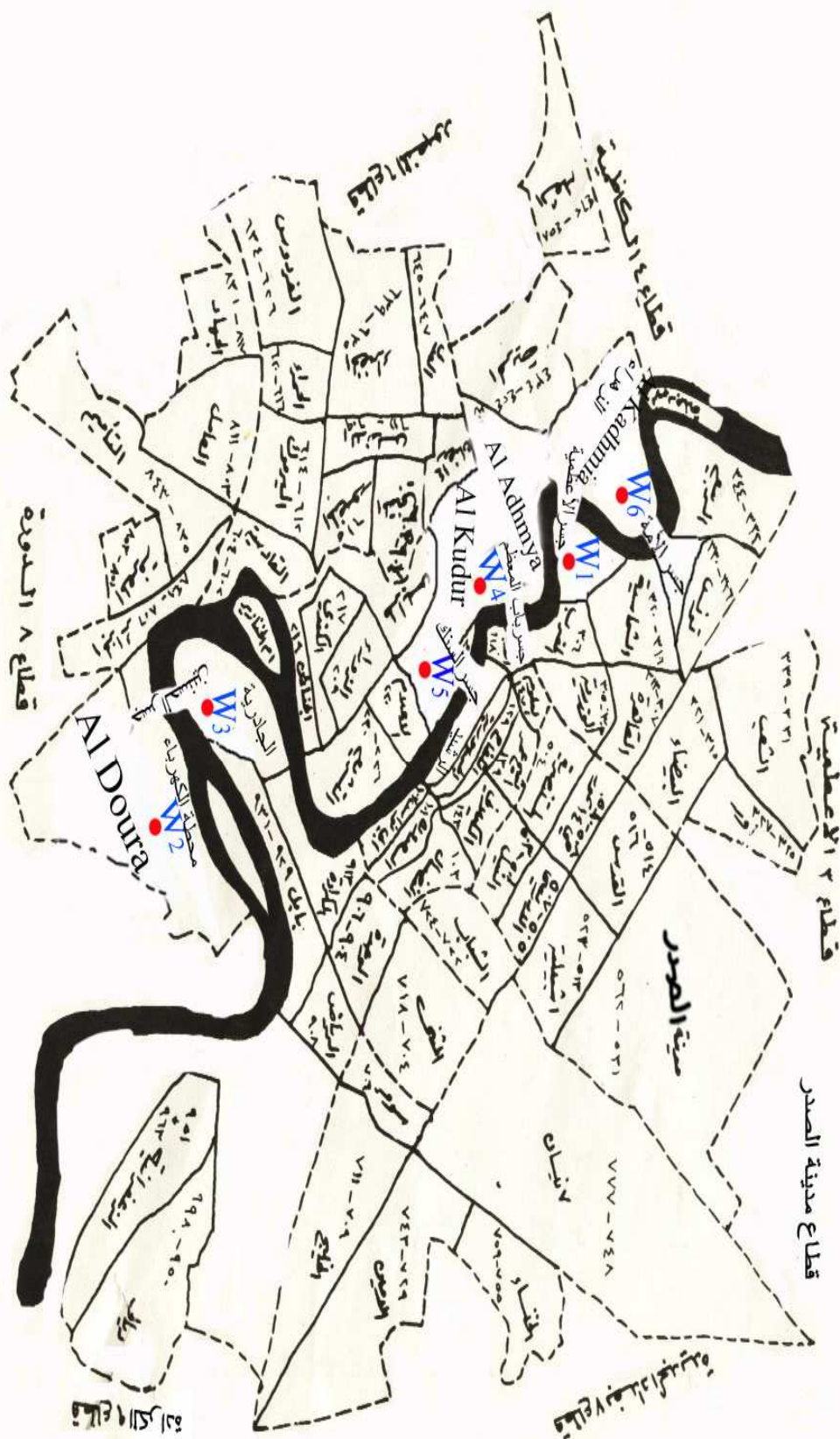


Fig. (3.2) The locations of collected water samples from different sites in Baehdad city (Tigris river) 1931 .



Fig. (3.3) The locations of collected soils samples from different sites in Karbala city [94] .

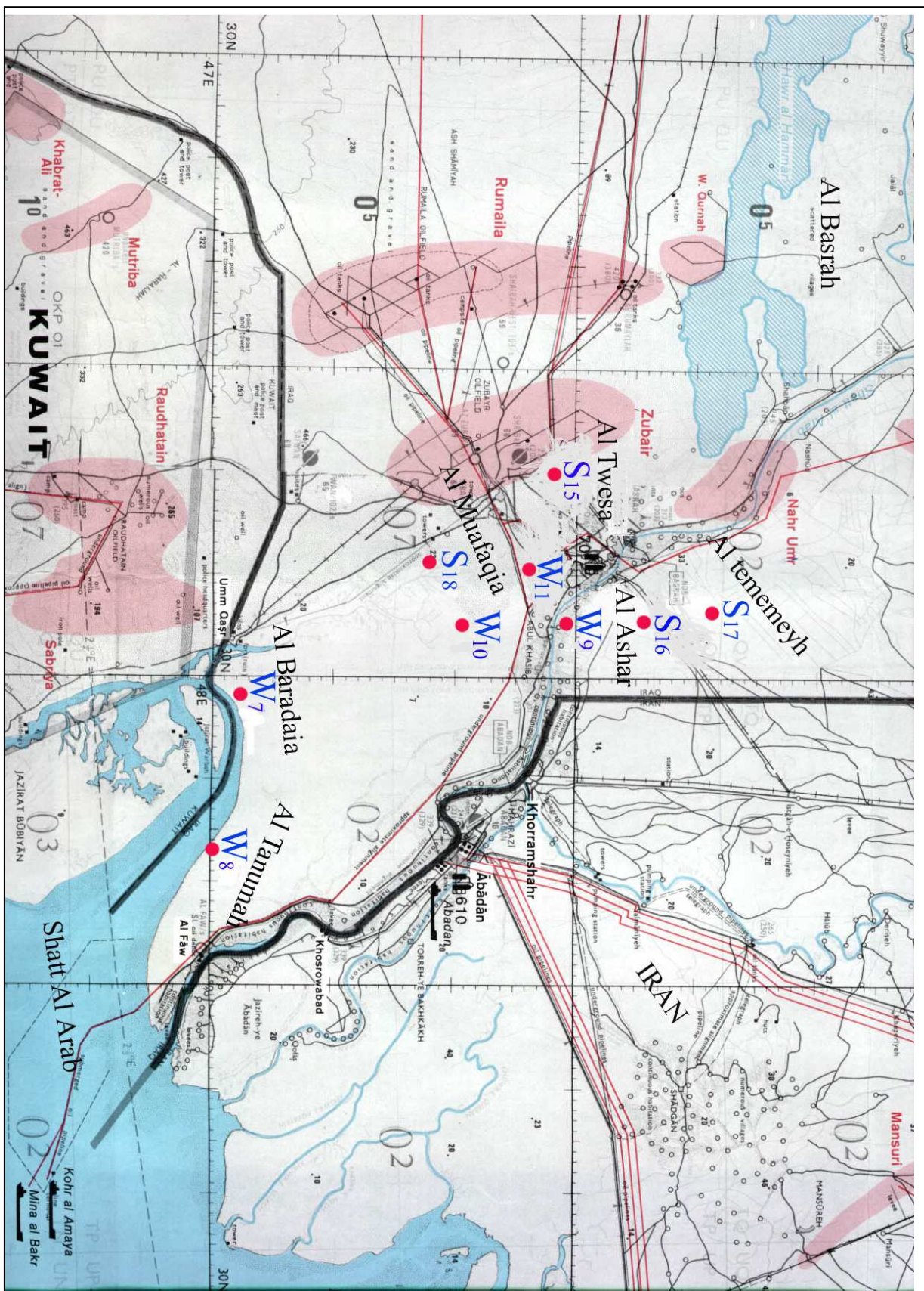


Fig. (3.4) The locations of collected soils and water samples from different sites in Basrah city [951].

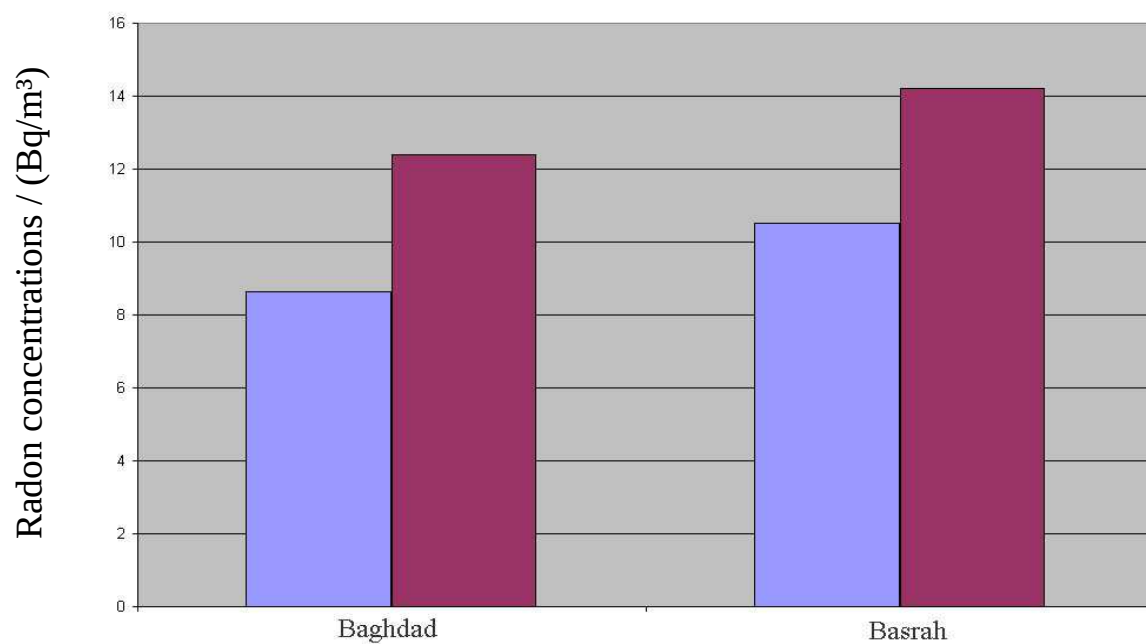


Fig. (4.8) Min. and Max. of radon concentrations in water samples.

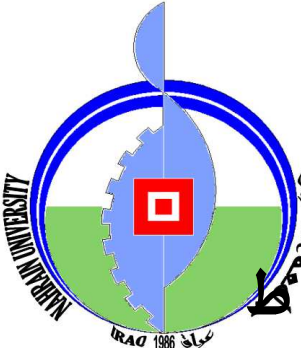
4.3. Conclusions

The discussion of the results, which are obtained from this study leads to the following conclusions :

- 1- The uranium concentration varies from 1.57 to 2.73 ppm and from 0.67 to 2.12 ppm in soil and water samples respectively and less than the allowed limit (11.7 ppm).
- 2- The radon gas concentration varies from 226.89 to 665.56 Bq/m³ and from 8.64 to 14.20 Bq/m³ in soil and water samples respectively and less than the limit recommended (100 Bq/m³).
- 3- The maximum uranium concentration in soils was (2.73 ppm) and in water (2.12ppm) in the Al-Taji and Al-Ashaar regions respectively .
- 4- The maximum radon gas concentration in soils was (665.56 Bq/m³) and in water (14.20 Bq/m³) in the Al-Taji and Al-Ashaar regions respectively .
- 5- Radionuclides (²²²Rn and ²³⁸U) had high concentration in site (S₄) and in site (W₉) because these sites are a field of a military operations during the war against Iraq in 2003.

4.4. Suggestions and Future Orientations

- 1- Study the radioactivity in soil and water by using γ -ray spectroscopy technique.
- 2- Study the uranium concentrations in biological samples for people living at these regions.
- 3- Study the uranium and radon gas concentrations for some weapons materials were using in military operations.



جمهورية
العراق
وزارة
التعليم
البحوث
والعلمي
جامعة
النهرين
كلية العلوم

دراسة النشاط الأشعاعي لنماذج من التربة والماء في بغداد وكربلاء والبصرة

رسالة
مقدمة الى كلية العلوم، جامعة
النهرين كجزء من متطلبات
نيل شهادة الماجستير علوم في
الفيزياء
من قبل

مصطفى عرب بداي
البيضاني
بكالوريوس - جامعة النهرين
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جمادي الأولى
حزيران

Examination Committee Certificate

We certify that we have read the thesis entitled “***Determination of Radioactivity in Soil and Water in Baghdad, Karbala and Basrah Samples***” and as Examining Committee, examined the student ***Mustafa Arab Badai Al-Baidhani*** in its contents and what is related to it, and that in our opinion it is adequate as standard of thesis, with ***Excellent*** standing of degree of Master of Science in Physics.

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List of Contents

<i>Contents</i>		
Abstract		i
List of Contents		ii
List of Figures		v
List of Tables		vii
List of Symbols		viii
<i>Chapter One "Introduction"</i>		
1.1	Radioactivity	1
1.2	Natural Radioactivity Sources	1
1.2.a	U-238 Series	2
1.2.b	U-235 Series	3
1.2.c	Th-232 Series	4
1.2.d	Np-237 Series	5
1.3	Previous Studies	6
1.3.1	Previous Studies of Uranium	6
1.3.2	Previous Studies of Radon	8
1.4	The Aim of the Present Work	9
<i>Chapter Two "Theoretical Part"</i>		
2.1	Solid State Nuclear Track Detectors SSNTDs	10
2.2	Application of the SSNTDs	11
2.3	Track Geometry	11
2.4	Revelation of Particle Tracks in Solids by Selective Chemical Etching	14
2.5	The Chemical Form of CR-39	16
2.6	The Chemical Etching	17

2.7	Track Affecting Parameters	19
2.7.1	The Track Etch Rate Velocity (V_T)	19
2.7.2	The Bulk Etch Rate Velocity (V_B)	19
2.8	The Etching Efficiency	20
2.9	Critical Angle for Etching	21
2.10	Man –Made Radioactivity Sources	23
2.11	Uranium	23
2.12	Depleted Uranium	24
2.13	Properties of Depleted Uranium	25
2.13.1	Radioactivity	26
2.13.2	Toxicity	26
2.13.3	Pyrophorosity	26
2.14	Applications of Depleted Uranium	27
2.14.1	Commercial	27
2.14.2	Military	27
2.15	Uses of Depleted Uranium in 2003 War Against Iraq	27
2.16	Health Effects of Depleted Uranium	29
2.16.1	Inhalation	29
2.16.2	Ingestion	29
2.17	Direct Radiation	30
Chapter Three “Experimental Part”		
3.1	Introduction	31
3.2	Collection of Samples	31
3.2.1	Soil Samples	31
3.2.2	Water Samples	31
3.3	Materials and Apparatus	38
3.3.1	The Track Detectors	38

3.3.2	The Etchant Solution	38
3.3.3	The Water Bath	39
3.3.4	The Optical Microscope	39
3.3.5	The Irradiation Source	39
3.4	Experimental Details for Uranium Concentration Measurements	40
3.4.1	Experimental Procedure for Soils Samples	40
3.4.2	Experimental Procedure for Water Samples	44
3.5	Experimental Details for Radon Concentration Measurements	47
3.5.1	Experimental Procedure for Soils Samples	47
3.5.2	Experimental Procedure for Water Samples	49
<i>Chapter Four “Results and Discussion”</i>		
4.1	Introduction	52
4.2	Results and Discussion	52
4.2.1	Uranium Concentrations in Soil Samples	52
4.2.2	Uranium Concentrations in Water Samples	55
4.2.3	Radon Concentrations in Soil Samples	57
4.2.4	Radon Concentrations in Water Samples	60
4.3	Conclusion	63
4.4	Suggestions and Future Orientations	63
References		64

List of Figures

Figure No.	Captions	Page No.
(2.1)	A track geometry for particle penetrates a detector material normally	13
(2.2)	A growth of track during chemical etching, from one side only	15
(2.3)	The chemical form of CR-39 plastic	16
(2.4)	Track etching process	18
(2.5)	A track etching a particle incident obliquely on the SSNTD	22
(3.1)	The locations of collected soils samples from different sites in Baghdad city	34
(3.2)	The locations of collected water samples from different sites in Baghdad city (Tigris river)	35
(3.3)	The locations of collected soils samples from different sites in Karbala city	36
(3.4)	The locations of collected soils and water samples from different sites in Basrah city	37
(3.5)	The irradiation of the detectors and samples to the neutron source	41
(3.6)	The relation between track density and uranium concentration (ppm) for standard geological soil samples	43
(3.7)	Evaporation of the water droplet and the formation of a thin deposit	45
(3.8)	The relation between track density and uranium concentration (ppm) for standard water samples	46
(3.9)	The apparatus of Radon gas (^{222}Rn) estimation by using (CR-39) detector for soil sample	48
(3.10)	The relation between track density and radon concentration (Bq/m^3) for standard geological soil samples	49
(3.11)	Radon gas (^{222}Rn) estimation by using (CR-39) detector for water sample	50
(3.12)	The relation between track density and radon concentration (Bq/m^3) for standard water samples	51

(4.1)	Uranium Concentrations in Soils Samples	54
(4.2)	min. and max. of Uranium Concentrations in Soils Samples	54
(4.3)	Uranium Concentrations in Water Samples	56
(4.4)	min. and max. of Uranium Concentrations in Water Samples	57
(4.5)	Radon Concentrations in Soils Samples	59
(4.6)	min. and max. of Radon Concentrations in Soils Samples	59
(4.7)	Radon Concentrations in Water Samples	61
(4.8)	min. and max. of Radon Concentrations in Water Samples	62

List of Tables

Table No.	Caption	Page No.
(1-1)	Radioactive decay series	2
(1-2)	U-238 Decay Series	3
(1-3)	U-235 Decay Series	4
(1-4)	Th-232 Decay Series	5
(2-1)	Isotopic composition	25
(2-2)	Armor-piercing projectiles among those in the U.S. military arsenal that use DU	28
(3-1)	Locations of soils samples in Baghdad and south regions	32
(3-2)	Locations of water samples in Baghdad (Tigris river) city and south regions (Shatt Al-Arab)	33
(4-1)	Uranium Concentrations in Soils Samples of Baghdad City and South Regions of Iraq	53
(4-2)	Uranium Concentrations in Water Samples of Baghdad City and South Regions of Iraq	55
(4-3)	Radon Concentrations in Soils Samples of Baghdad City and South Regions of Iraq	58
(4-4)	Radon Concentrations in Water Samples of Baghdad City and South Regions of Iraq	60

List of Symbols

Symbol	Description
Am-Be	Americium –Beryllium Source
Bq	Bequerel
°C	Degree Centegrate
C _S	uranium concentration of standard sample
C _X	uranium concentration of unknown sample
D	diameter of the etch pit
DU	Depleted uranium
E _B	Activation Energy of the Bulk Etching
E _T	Activation Energy of the Track Etch
EPA	Environmental Protection Agency
h	thickness of the surface removed by etching
IAEA	International Atomic Energy Agency
ICRP	International Committee of Radiation Protection
k	Boltzman constant = 1.38×10^{-23} j mol /K
KOH	Potassium Hydroxide
L	length of the latent track
LSA	Low Specific Activity
n	neutron
N	Normality
NaOH	Sodium Hydroxide
ppm	Part Per million

R	Length of track at the removed surface
R.O.	Reverse Osmosis
S	Soil sample
SSNTDs	Solid State nuclear track detectors
t	etching time
T	Temperature of the etching solution
UNEP	United Nation Environment Program
UNSCEAR	United Nation Scientific Committee on the Effects of Atomic Radiation
V	volume of distilled water
V_B	bulk etch rate
V_T	track etch rate
w	the weight of NaOH
W	Water sample
w_{eq}	equivalent weight of NaOH
WHO	World Health Organization
Z	Atomic Number

Symbol	Description
α	Alpha particle
$\emptyset$	Incident angle
$\emptyset_C$	Critical angle
η	Etching efficiency
ρ	Track Density
ρ_s	track density of standard sample
ρ_x	track density of unknown sample
μm	micrometer
μg	microgram

3.1. Introduction

This chapter describes the methods for estimation of depleted uranium and radon gas concentrations for the different samples of soil and water (natural and standards) from different sites in Iraqi regions .The materials and the apparatus which used included.

3.2. Collection of Samples

Two types of samples (soils and water) were collected from different sites in Baghdad city and other regions of Iraq ,as shown in Fig. (3.1),(3.2),(3.3) and (3.4) .

3.2.1. Soil Samples

Samples of soils were collected from different sites in Iraq regions (bombardment regions during the war against Iraq in 2003) from depth (5 cm),as shown in Table (3.1) and Fig. (3.1),(3.3) and (3.4) from Baghdad, Karbala and Basrah cities respectively .Then the samples dried and cleaned from the doping grinds by using special sieve (0.27 mm in diameter) .

3.2.2. Water Samples

Samples of water were collected from different sites in Iraqi regions . One litter volume of water was collected from some projects of water refinement on the rivers (Tigris and Shatt Al –Arab) , also collected from different houses (drinking water), as shown in Table (3.2) and Fig. (3.2) and (3.4) from Baghdad and Basrah cities respectively .

**Table (3 –1) Locations of soils samples in Baghdad city
and south regions .**

Date of samples	Number of samples	Symbol	City	Location
22-11-2004	1	S ₁	Baghdad	Abu Ghreab
22-11-2004	2	S ₂	Baghdad	Al Binuq
22-11-2004	3	S ₃	Baghdad	Al Binuq (Civilian Advocate ship)
25-11-2004	4	S ₄	Baghdad	Al Taji
28-11-2004	5	S ₅	Baghdad	Al Mansur
28-11-2004	6	S ₆	Baghdad	Al Mansur (Al Mamoon communications)
28-11-2004	7	S ₇	Baghdad	Al Washash
15-12-2004	8	S ₈	Baghdad	Al Sader City (Destroyed Houses)
5-1-2005	9	S ₉	Baghdad	Al Doura (Electrical Station)
14-1-2005	10	S ₁₀	Baghdad	Al Meshahda (Al Ishaqi Company)
20-2-2005	11	S ₁₁	Karbala	Al Abbas Street
20-2-2005	12	S ₁₂	Karbala	Al Abbas Street
20-2-2005	13	S ₁₃	Karbala	Industrial Region
20-2-2005	14	S ₁₄	Karbala	Al Quzwinia
12-3-2005	15	S ₁₅	Basrah	Al Twesa
13-3-2005	16	S ₁₆	Basrah	Al Ashaar
13-3-2005	17	S ₁₇	Basrah	Al Tememeyh
13-3-2005	18	S ₁₈	Basrah	Al Muafaqia (Residential Region)

Table (3 –2) Locations of water samples in Baghdad city (Tigris river) city and south regions (Shatt Al-Arab) .

Date of samples	Number of samples	Symbol	City	Location
4-1-2005	1	W_1	Baghdad	Al Adhmya (Al Adhmya Bridge)
5-1-2005	2	W_2	Baghdad	Al Doura (Electrical Station)
5-1-2005	3	W_3	Baghdad	Al Hasanian Bridge
4-1-2005	4	W_4	Baghdad	Maqam Al Kudur region (Bab Al muadham Bridge)
4-1-2005	5	W_5	Baghdad	Al Sinaq Bridge
4-1-2005	6	W_6	Baghdad	Al Kadhmia (Al Aemaa Bridge)
13-3-2005	7	W_7	Basrah	Al Baradaia (Shatt Al Arab)
13-3-2005	8	W_8	Basrah	Al Tanumah (Shatt Al Arab)
13-3-2005	9	W_9	Basrah	Al Ashaar (Shatt Al Arab) (near the Ports)
14-3-2005	10	W_{10}	Basrah	Al Muafaqia (Drinking Water of Houses)
14-3-2005	11	W_{11}	Basrah	Petrochemical Factory (Drinking Water of R.O.)

3.3. Materials and Apparatus

3.3.1. The Track Detectors

Commercially available sheets of CR-39 plastic which are presently known to be the most sensitive SSNTD and also characterized by low background were used in the present work.

These detector sheets of 250 μm thick were cut into small pieces each of $1\text{cm} \times 1\text{cm}$ area. The present sheets of CR-39 were made by Pershore Moulding LTD Company, U.K. The detector sheets were stored at normal laboratory conditions.

3.3.2. The Etchant Solution

Sodium hydroxide solution with 6.25 normality has been used for the etching process. The preparation of normality is using [86]:

$$w = w_{\text{eq}} \times N \times V \quad \dots(3.1)$$

where:

w = the weight of NaOH needed to prepare the given normality.

w_{eq} = equivalent weight of NaOH = addition of the atomic weight of Na, O and H = 40.

N = normality = 6.25.

V = volume of distilled water = 250 ml.

The etchant compartment has a volume of about 250 ml contains the NaOH solution with 6.25 N. This apparatus is closed assembly, except for small vent at the top of the condenser tube, which prevents any change of etchant normality (concentration) during the experiment due to evaporation. The etching was performed at 60°C while the etching time was 6 hours.

3.3.3. The Water Bath

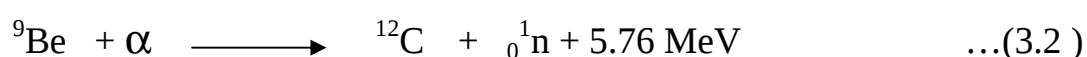
An etching bath of the type “Labsco” (Germany) was used in the present work. It includes a thermostat, which can be operated over a range of 20°C to 110°C. However, in this study the chemical etching was carried out at 60°C as, suggested previously [21]. The accuracy of regulation of temperature is better than $\pm 0.1^\circ\text{C}$.

3.3.4. The Optical Microscope

The counting of all the chemically etched tracks was carried out using an optical microscope. It is a standard transmission type binocular research microscope (Bausch & Lomb, Japan). It is capable of giving magnifications of up to 400 x .

3.3.5. The Irradiation Source

For the irradiation test, an (Am–Be) source with flux $5 \times 10^3 \text{ n/cm}^2.\text{s}$ was used. It emits fast neutrons from the (α , n) reaction such as:



This source consists of a rod of (Am–Be) surrounded by a paraffin wax. This wax is usually used for moderating the fast neutrons to thermal energies [87].

3.4. Experimental Details for Uranium Concentration Measurement

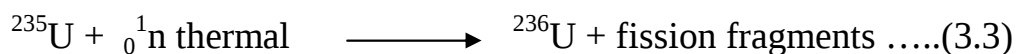
In the present work, the methods for measurement of uranium concentrations of the samples (soil and water), which is called the irradiation methods are :

3.4.1. Experimental Procedure for Soils Samples

Soil samples were taken from locations in Iraqi regions , as shown in Table (3-1) and Fig. (3.1),(3.3) and (3.4) . Soil samples were dried and cleaned from the doping grinds using special sieve (0.27 mm in diameter).

0.5g of soil samples were mixed with 0.1g of methylcellulose powder ($C_6H_{10}O_5$)_n used as a binding material . The mixture was pressed into a pellet of 1cm diameter and 1.5mm thickness.

The pellets were covered with (CR-39) detector and put in a plate of paraffin wax at a distance of (5 cm) from the neutron source (Am-Be) as shown Fig. (3.5), with flounce of thermal neutron (3.024×10^9 n.cm⁻²) and flux (5×10^3 n cm⁻² s⁻¹), to obtain induced fission fragments from the equation (3.3) [88] :



After the irradiation time (7 d) , (CR-39) detectors were etched in (6.25N) NaOH solution at temperature of 60 °C for (6 h) , then the induced fission tracks density were recorded using the optical microscope.

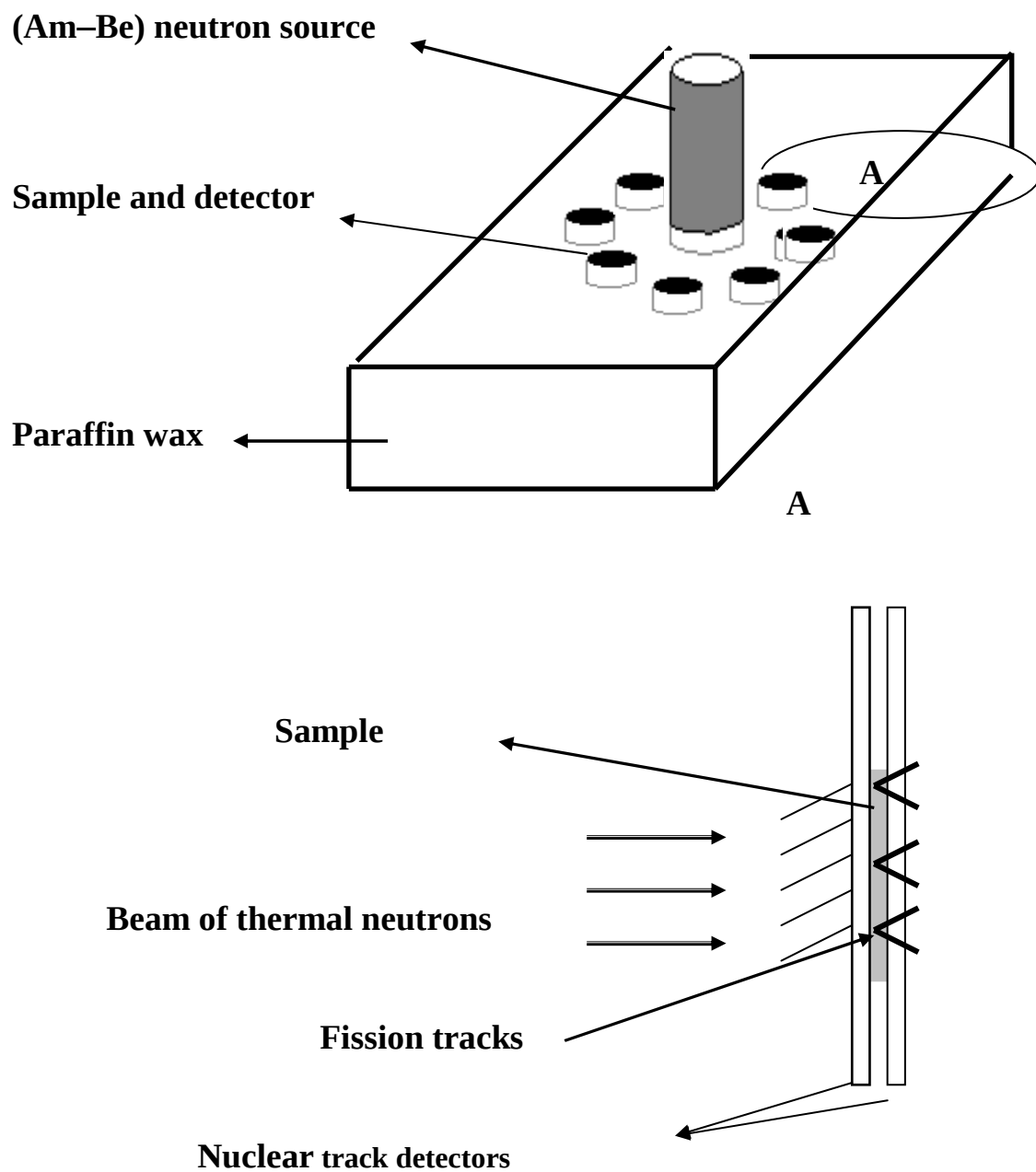


Fig. (3.5) The irradiation of the detectors and samples to the neutron source [89].

The density of the fission tracks (ρ) in the samples was calculated according to the following relation [90] .

$$\text{Track density } (\rho) = \frac{\text{Average number of total pits (tracks)}}{\text{Area of field view}} \quad \dots(3.4)$$

The uranium concentrations in the soil samples were measured by comparison between track densities registred on the detectors of the sample pellet and that of the standard geological sample pellets from the relation [6,7,91] :

$$C_X (\text{sample}) / \rho_X (\text{sample}) = C_S (\text{standard}) / \rho_S (\text{standard}) \quad \dots(3.5)$$

$$C_X = C_S \cdot (\rho_X / \rho_S) \quad \dots(3.6)$$

Fig. (3.6) shows this relation , when (slope = ρ_S / C_S) .

Where :

C_X : uranium concentration in unknown sample (ppm).

C_S : uranium concentration in standard sample (ppm).

ρ_X : track density of unknown sample (tracks/mm²).

ρ_S : track density of standard sample (tracks/mm²).

.

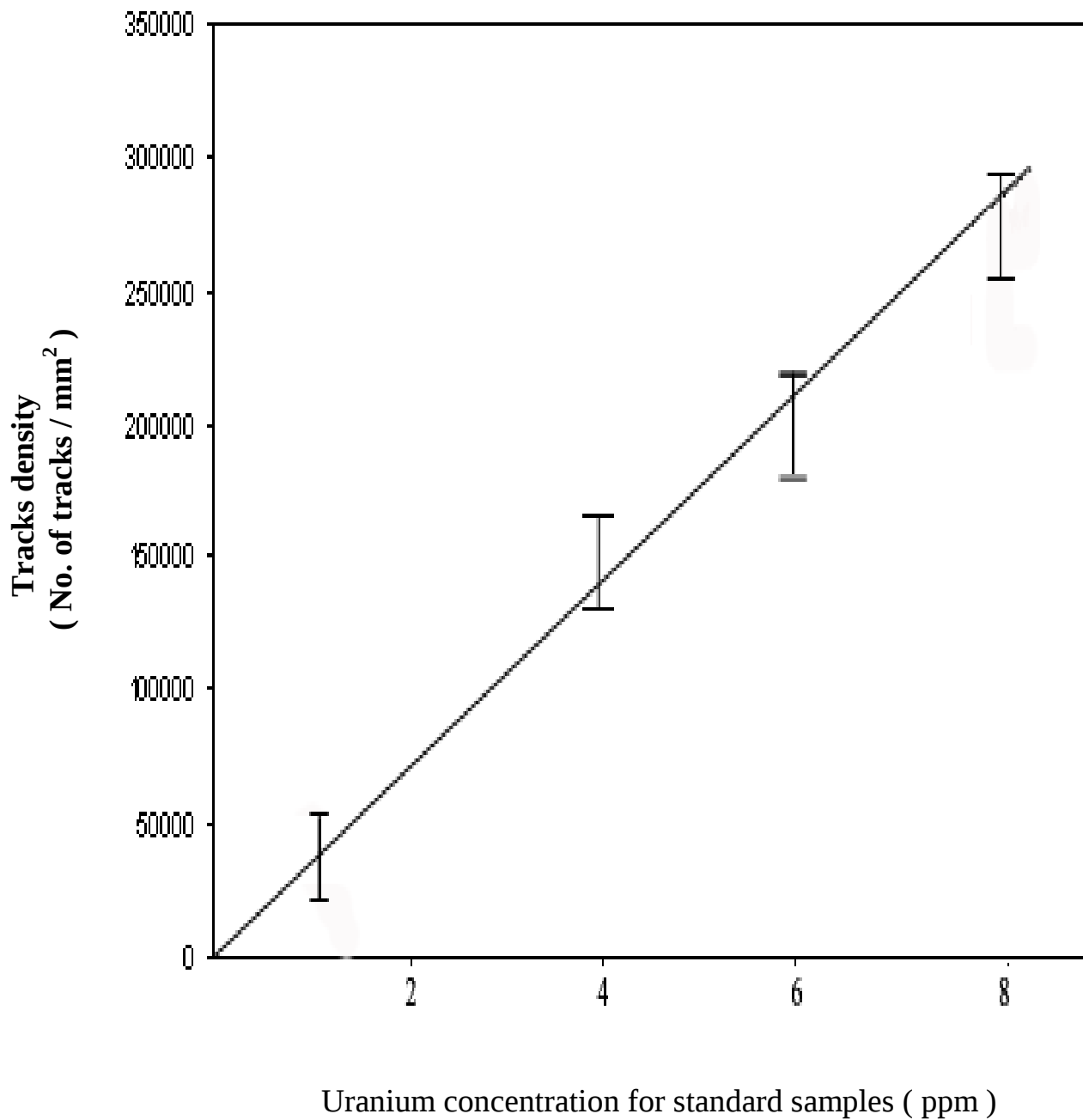


Fig.(3.6) The relation between track density and uranium concentration (ppm) for standard geological soil samples.

3.4.2. Experimental Procedure for Water Samples

Water samples were taken from locations in Iraqi regions , as shown in Table (3-2) and Fig. (3.2) and (3.4) .

Measurement of alpha-emitters in the environmental surface water can be complicated by the presence of a mass of suspended uranium bearing particle [13]. The procedure is to place known volume of water in the form of few drops (2 drops = 0.04 cc) on a track detector materials , water droplets are allowed to evaporate at room temperature , as sketched in Fig. (3.7), this leaves a thin residue of non-volatile constituents, the detector was then covered with a second piece of the detector. Subsequently exposure to (Am-Be) neutron source with flux of thermal neutrons (5×10^3 n/cm².s) for (7 d). Fig. (3.5) presents the irradiation of the detectors and the samples to the neutron source.

The detectors base or cover were etched in (6.25N) NaOH solution at temperature of 60 °C for (6 h) and washed by distille water , dried and the number of fission tracks produced was then counted by using the optical microscope .

This procedure was done to all samples including the standard samples

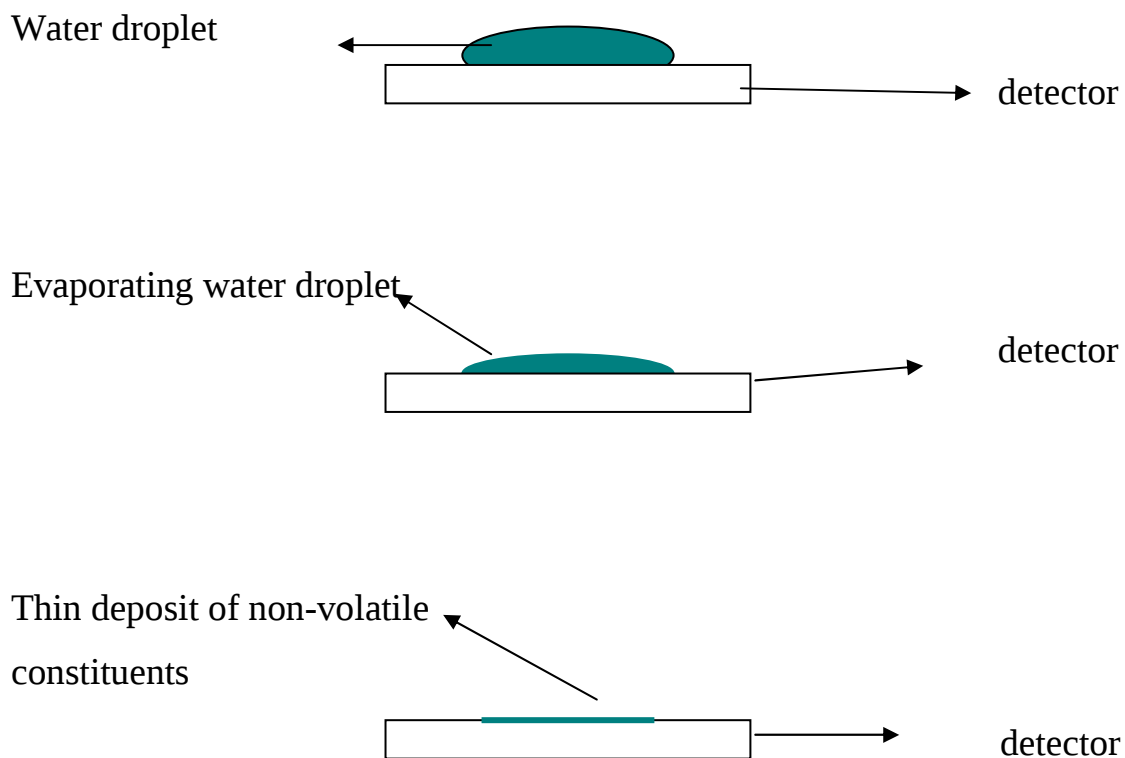


Fig. (3.7) Evaporation of the water droplet and the formation of a thin deposit.

The uranium concentrations in the water samples was measured by comparison between track densities of the water samples and that of the standard solutions from the equation (3.6).

The concentration – track density relationship for the present standard samples are illustrated in Fig.(3.8) .

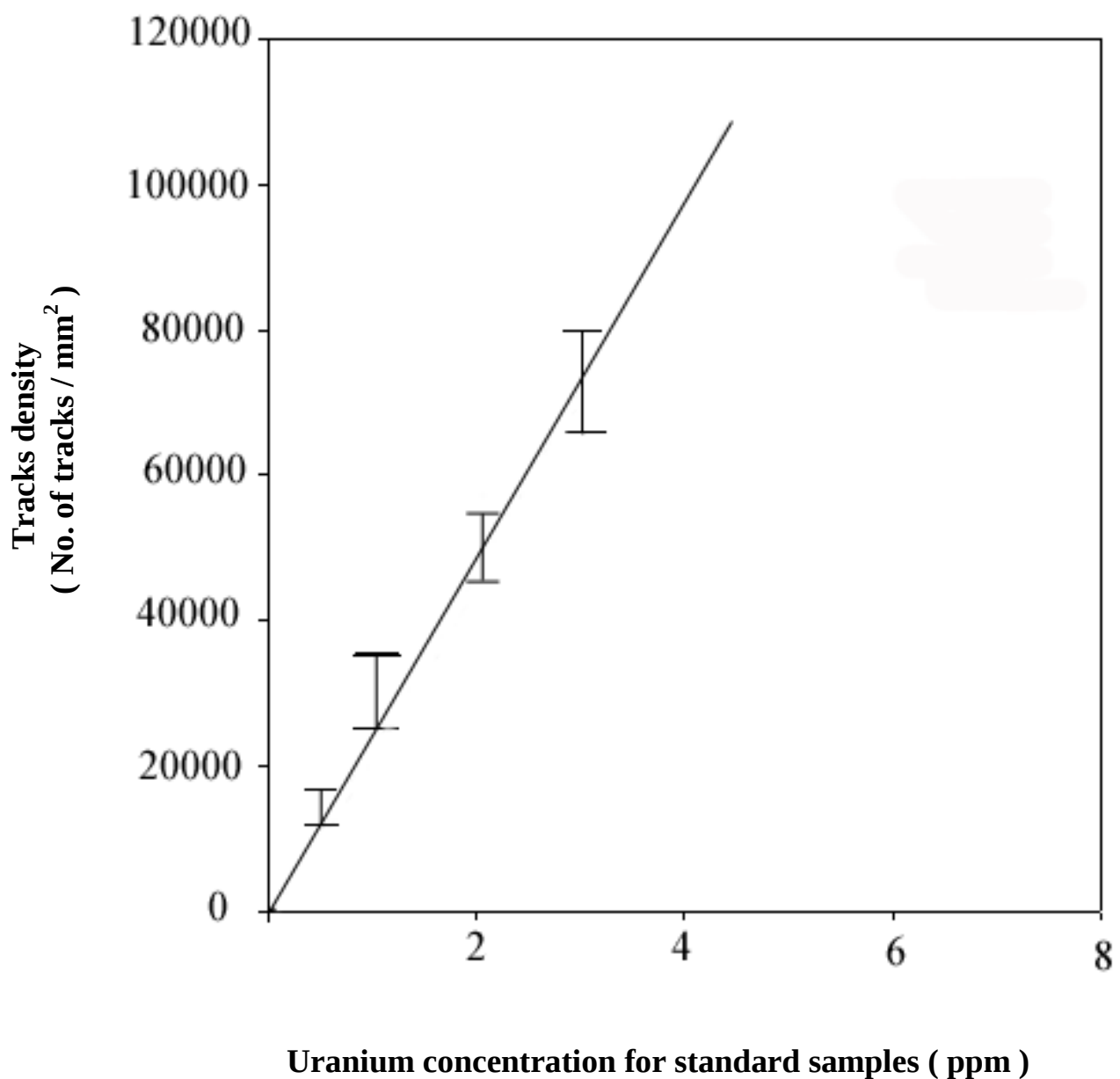


Fig.(3.8) The relation between track density and uranium concentration (ppm) for standard water samples.

3.5. Experimental Details for Radon Concentration Measurements

The radon concentrations in two types of samples (soil and water), were measured using the natural exposure method.

3.5.1. Experimental Procedure for Soils Samples

Soil samples were taken from locations in Iraqi regions, shown in Table (3-1) and Fig. (3.1), (3.3) and (3.4).

Soil samples were dried and cleaned from the doping grinds by using special sieve (0.27 mm in diameter). 10 g of soil samples were placed in plastic can. The dimensions of the can minimize the effect of gas thoron. Pieces of (CR-39) track detectors were fixed under the cover of plastic can, which contain the soil samples. The exposure time was (30 d) as shown in Fig. (3.9).

After the exposure time, (CR-39) detectors were etched in 6.25 N (NaOH) solution at temperature of 60 °C for (6 h) and washed by distilled water, dried. The tracks density were recorded using the optical microscope with magnification (400 x).

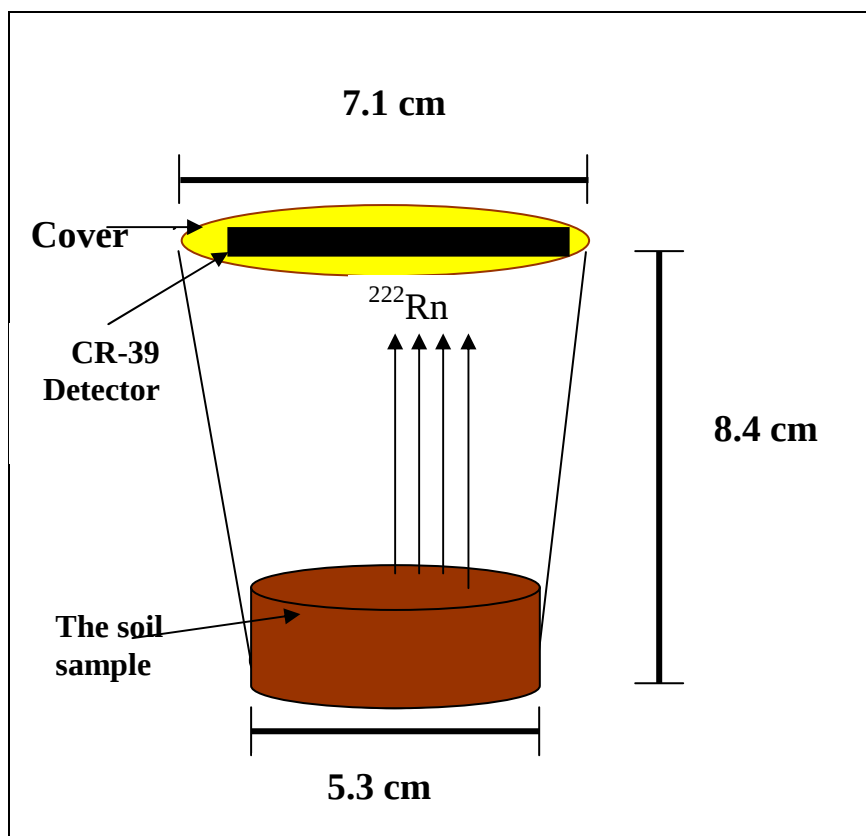


Fig.(3.9)The apparatus of Radon gas (^{222}Rn) estimation by using (CR-39) detector for soil sample .

At the same way in determination of the uranium concentrations , radon gas (^{222}Rn) concentrations in the soil samples was measured from the equation (3.6) by comparison between tracks densities registred on the detectors around the sample and that of the standard geological sample .

Fig. (3.10) shows the relation between the tracks density and radon concentration for the standard samples .

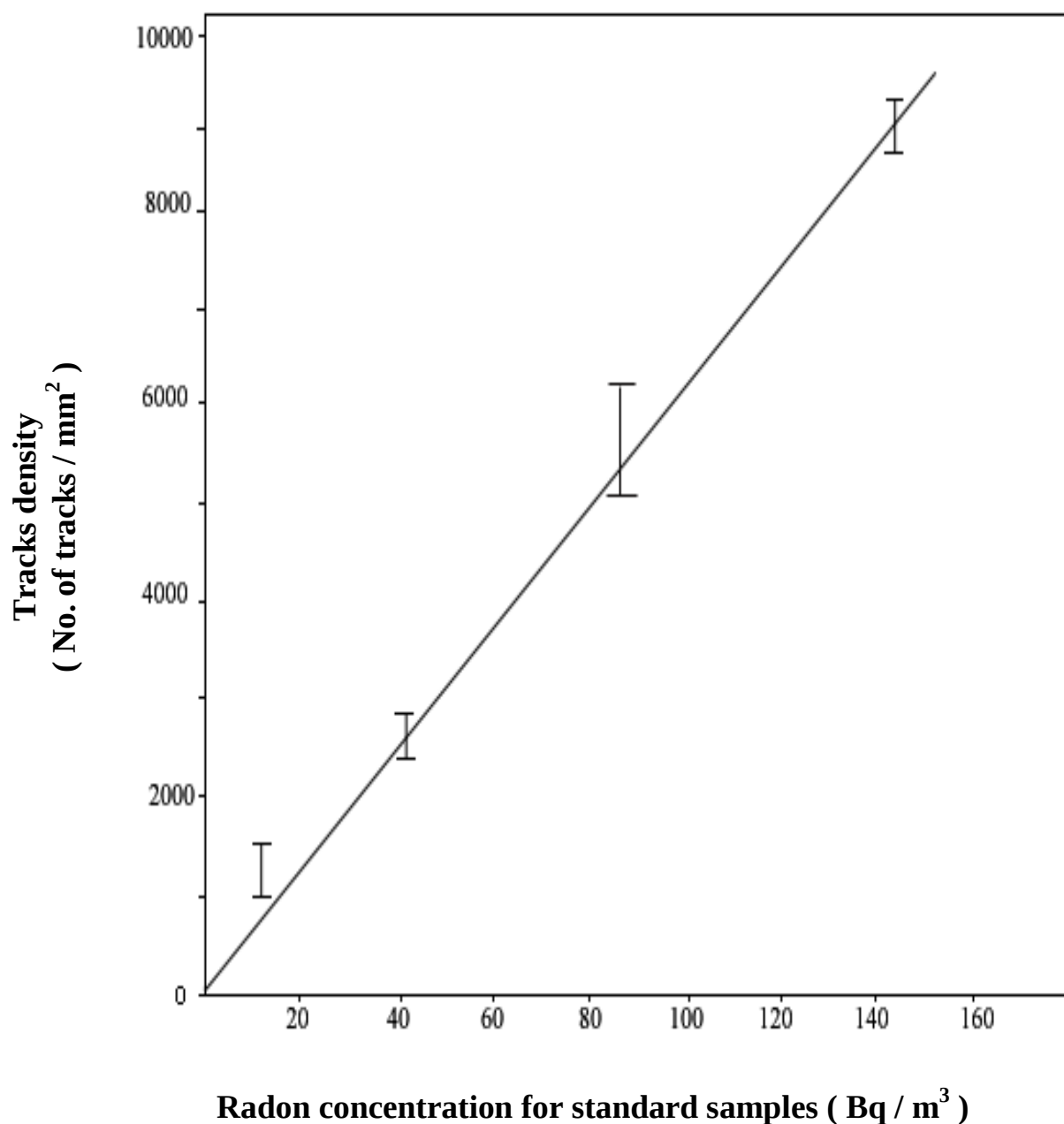


Fig.(3.10) The relation between track density and radon concentration (Bq/m³) for standard geological soil samples.

3.5.2. Experimental Procedure for Water Samples

Water samples were taken from locations in Iraqi regions , as shown in Table (3-2) and Fig. (3.2) and (3.4) .

The method includes exposing the detectors to the water samples directly for certain period of time.

(250 ml) volume of water samples were placed in plastic can at room temperature . Pieces of (CR-39) track detectors with dimensions ($1 \times 1 \text{ cm}^2$) were fixed under the cover of plastic can . The exposure time were (30 days) and the detection geometry was almost 4π , and the experimental set up is shown in Fig. (3.11).

After the exposure period, all the track detectors were etched in (NaOH) solution with 6.25 N at 60°C for (6 h). The detectors were then washed in distilled water, dried and were scanned under the optical microscope to record the tracks density .

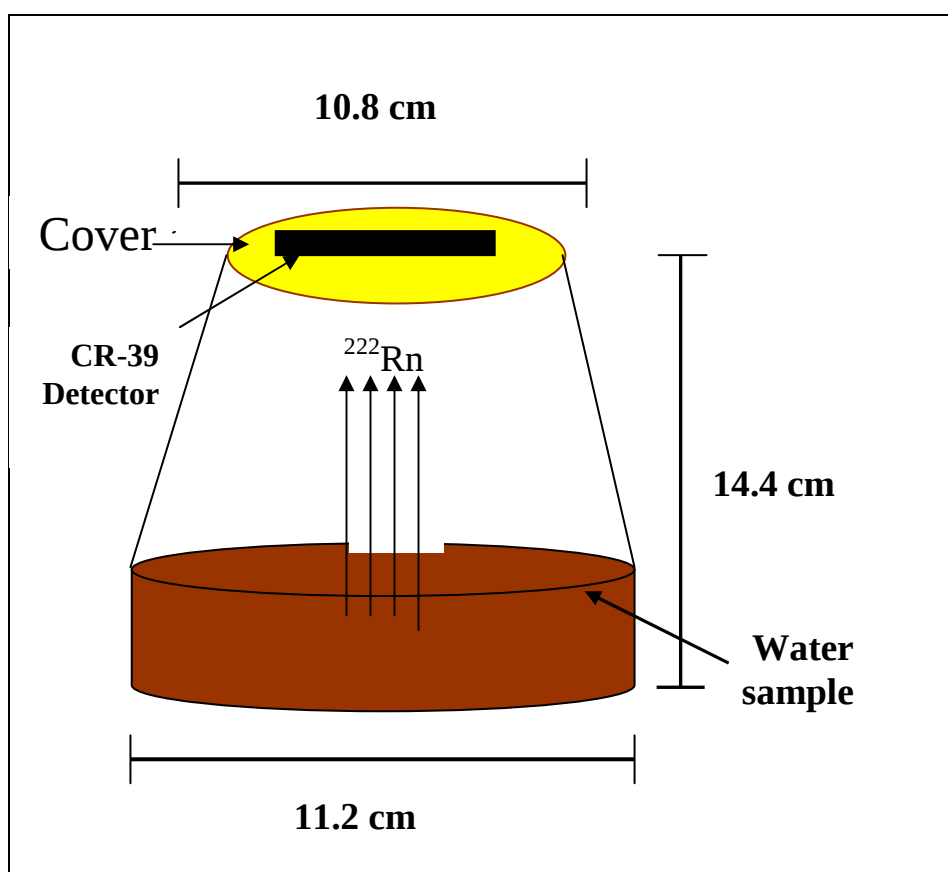


Fig.(3.11) Radon gas (^{222}Rn) estimation by using (CR-39) detector for water sample .

Radon gas (^{222}Rn) concentrations were determined by comparing mean track densities recorded in CR-39 detectors of water samples and with the standard samples from the equation (3.6) .

Fig. (3.12) shows the relation between the track density and radon concentration (Bq/m^3) for standard sample .

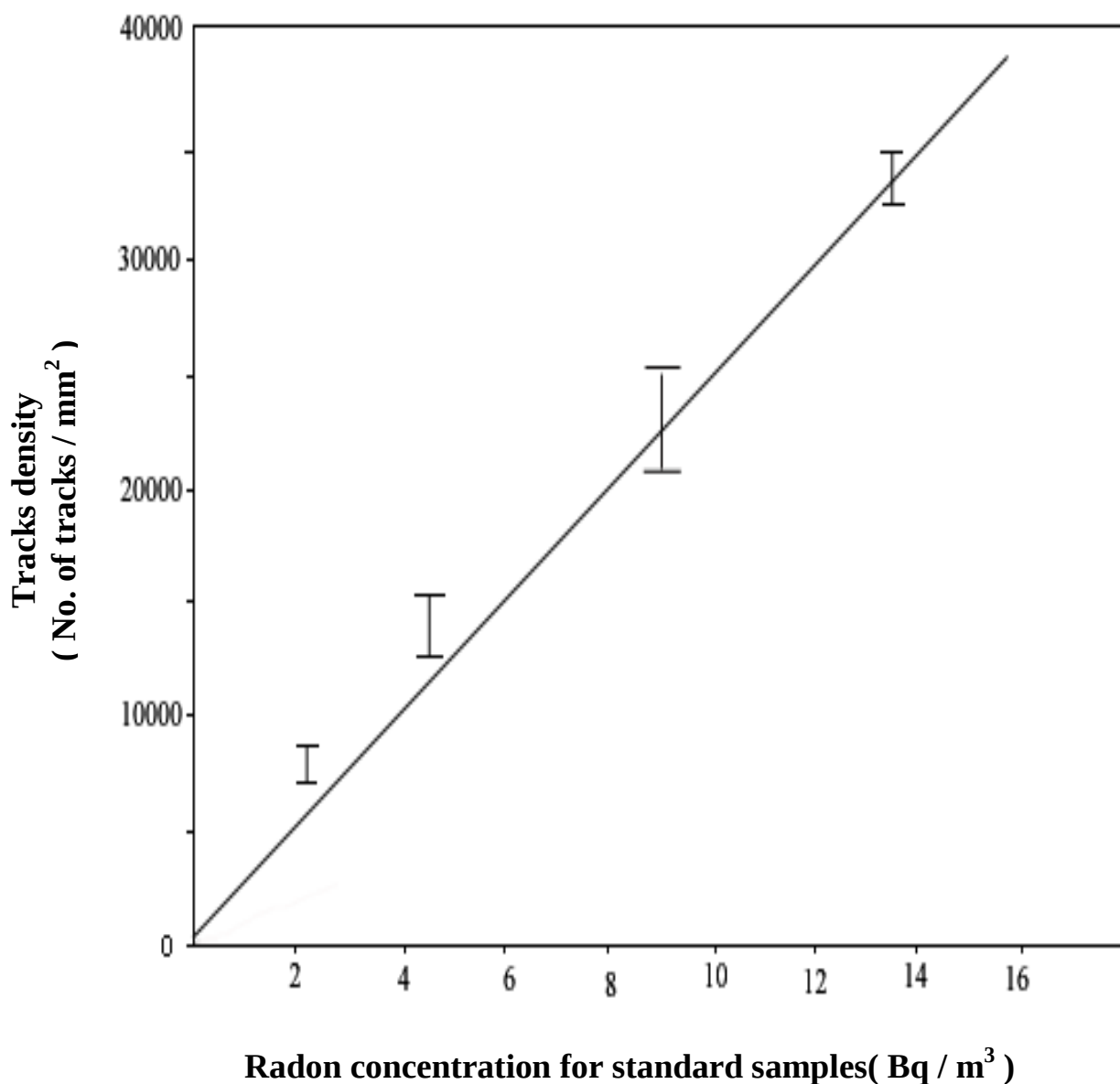


Fig.(3.12) The relation between track density and radon concentration (Bq/m^3) for standard water samples.

Examination Committee Certificate

We certify that we have read the thesis entitled “***Determination of Radioactivity in Soil and Water in Baghdad, Karbala and Basrah Samples***” and as Examining Committee, examined the student ***Mustafa Arab Badai Al-Baidhani*** in its contents and what is related to it, and that in our opinion it is adequate as standard of thesis, with ***Excellent*** standing of degree of Master of Science in Physics.

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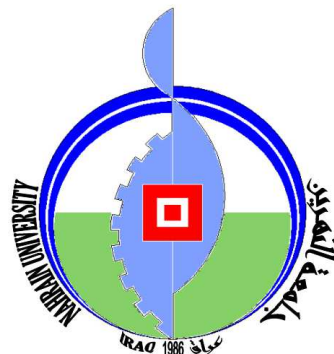
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