



# **Distribution and speciation of uranium in Jordan phosphate rocks and their phosphate fertilizer industry**

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# OUTLINE

**1- Introduction**

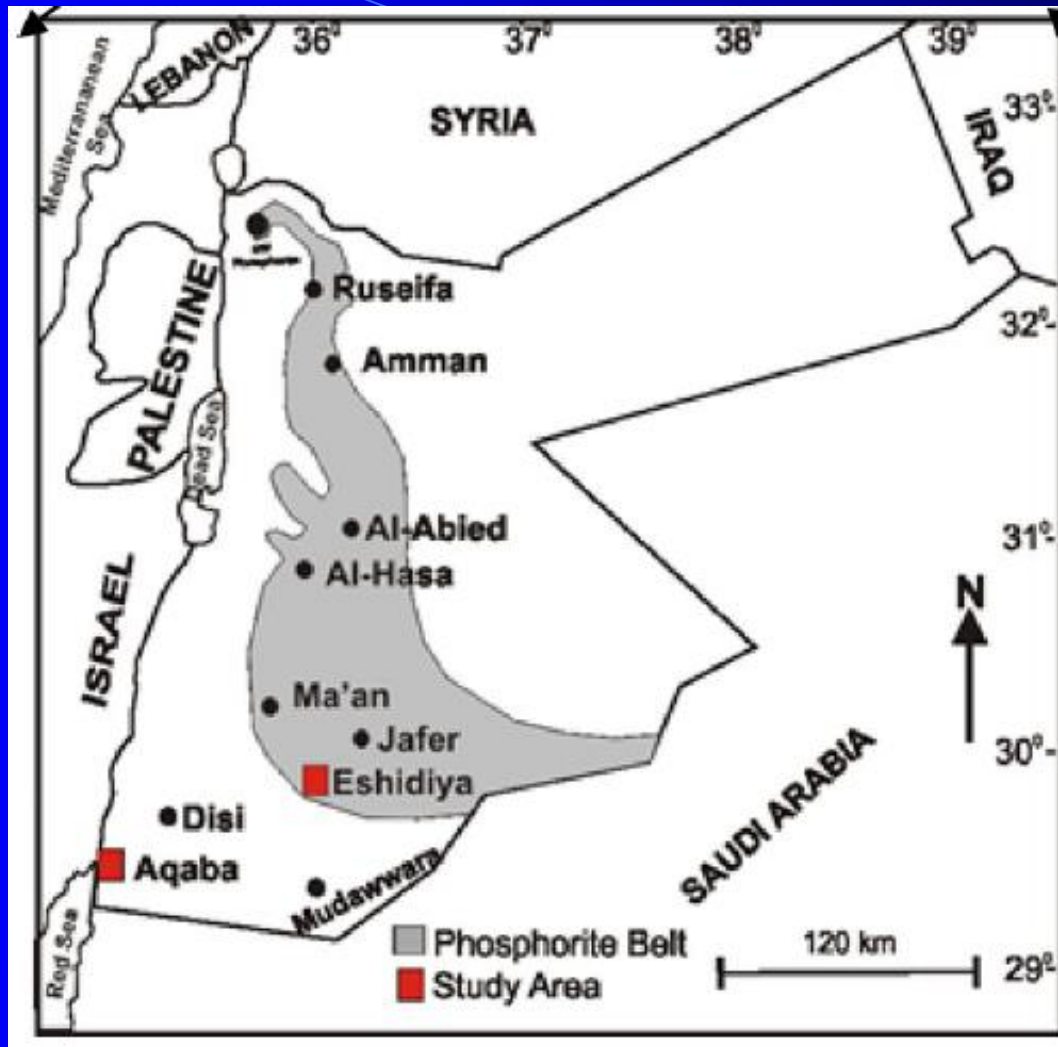
**2- Objectives**

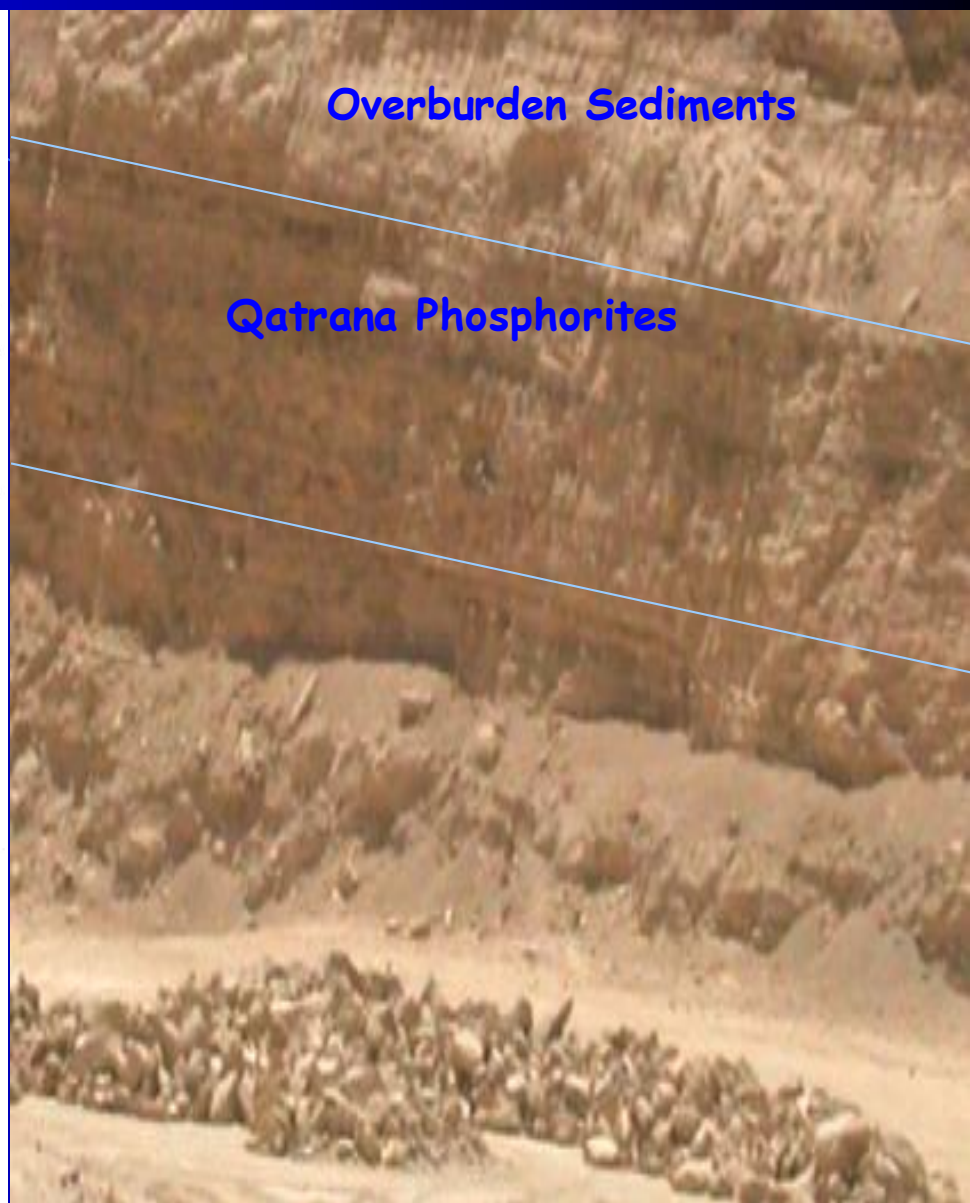
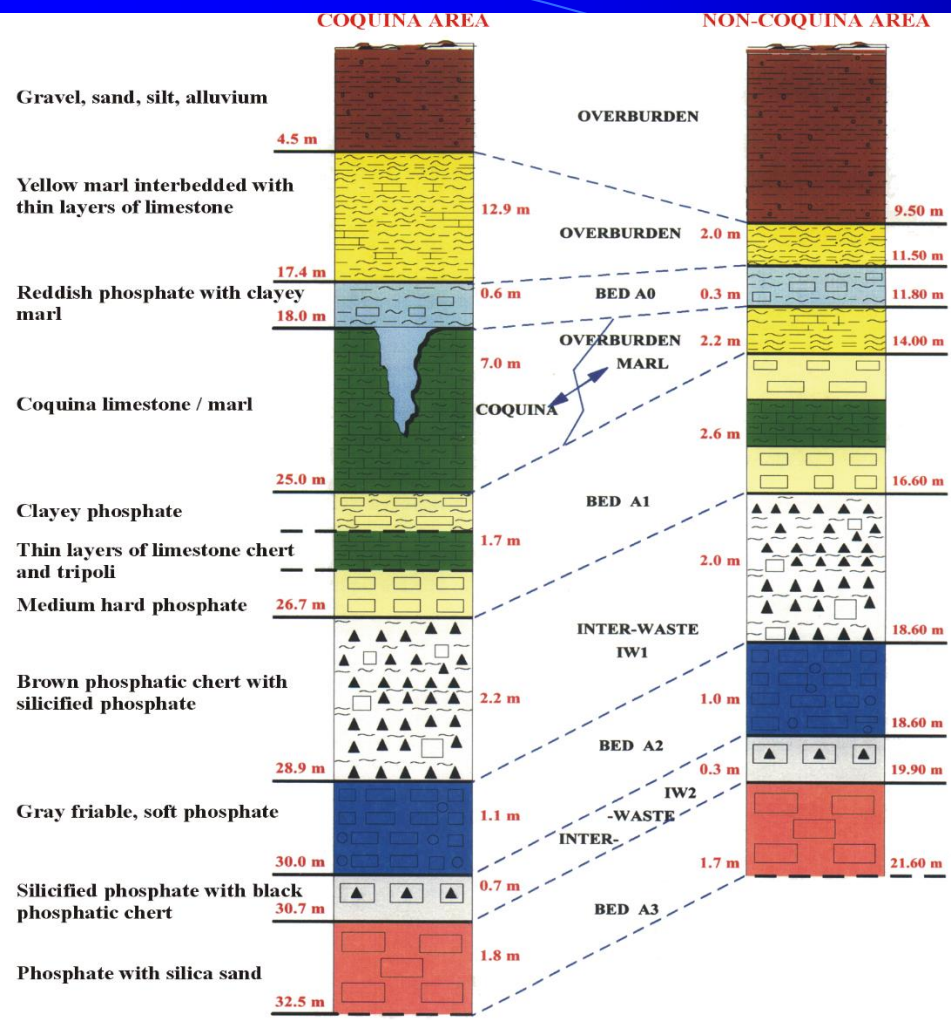
**3- Methods and Techniques**

**4- Uranium fate during different processing steps**

**5- Plant availability of U**

**6- Conclusions**





Overburden Sediments

Qatrana Phosphorites

- Notes :
- a. Vertical and horizontal not to scale.
  - b. Typical thicknesses vary.
  - c. The coquina limestone south and east changes to marl.

Lithostratigraphical sections of coquina and non-coquina areas (modified after Sofremines, 1984).

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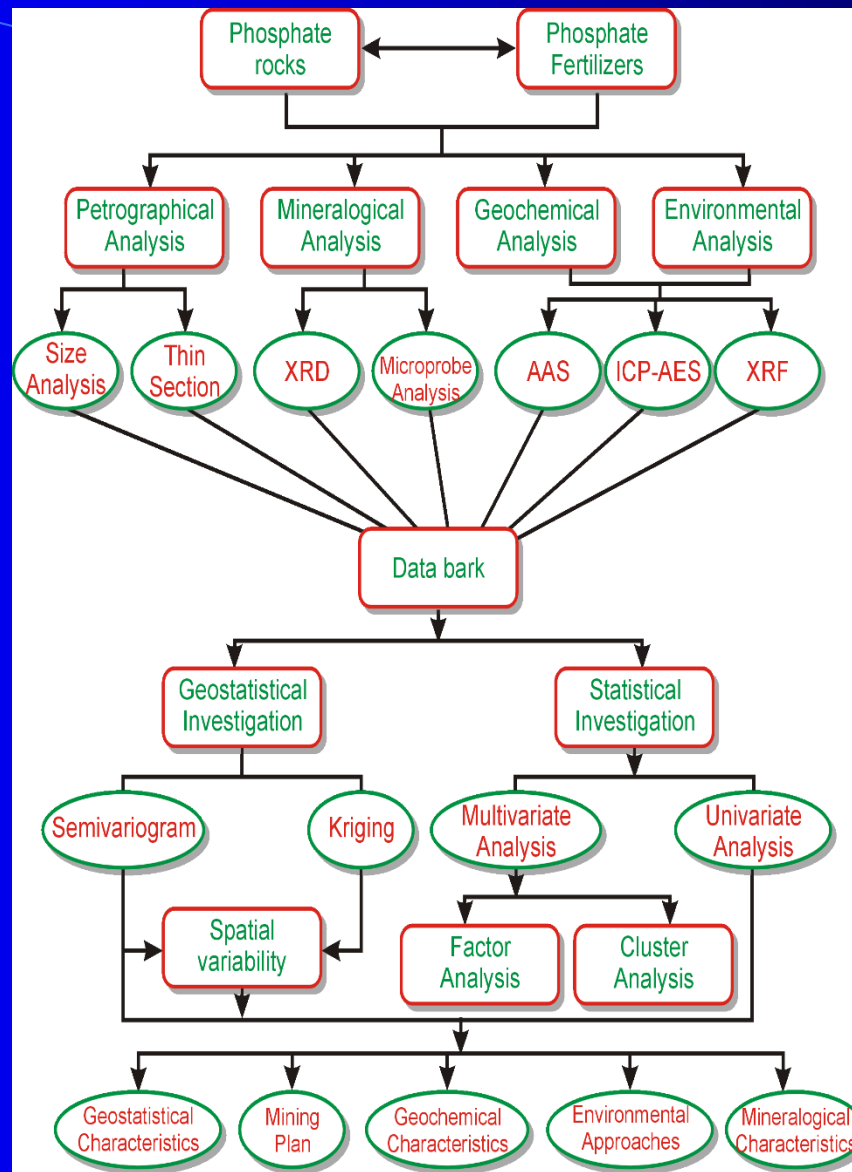
# Aims of This Study

**1- Determining the mode of occurrences of U in EShidiya phosphorites.**

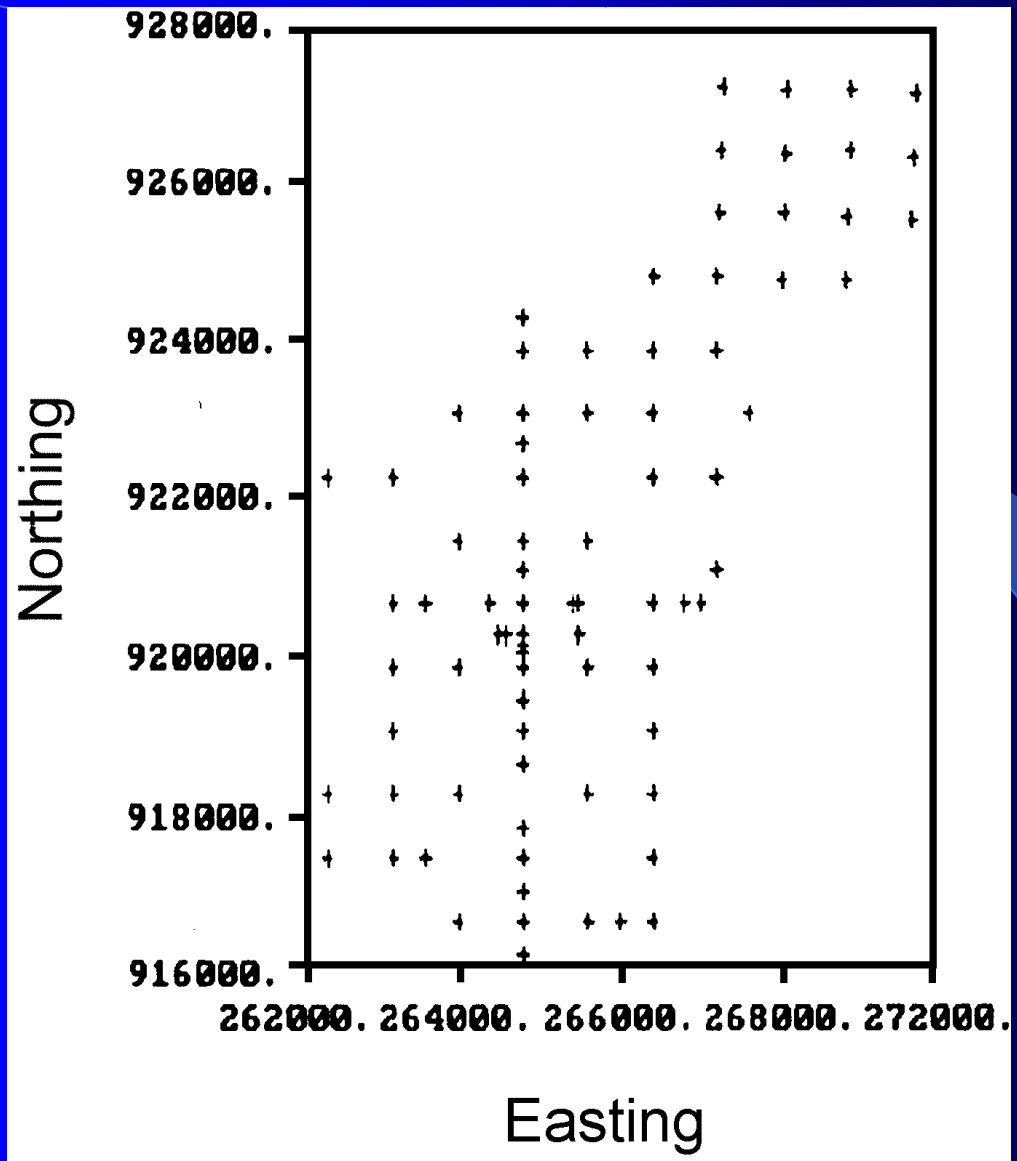
**2- Determining the redistribution pathway of U during different processing steps (e.g. **Crushing and Screening Processes**) the **beneficiation processes** (e.g. **Washing and Flotation**) the **drying processes in all beds A1, A2 and A3** and the **fertilizer processes** (e.g. **DAP**).**

# Aims of This Study

***3- Determining the plant available toxic heavy metals like U the might be enter the soil through fertilizers produced from these phosphates.***



**Flow chart showing the systematic methods of Uranium investigation.**



**Plot of the samples locations.**



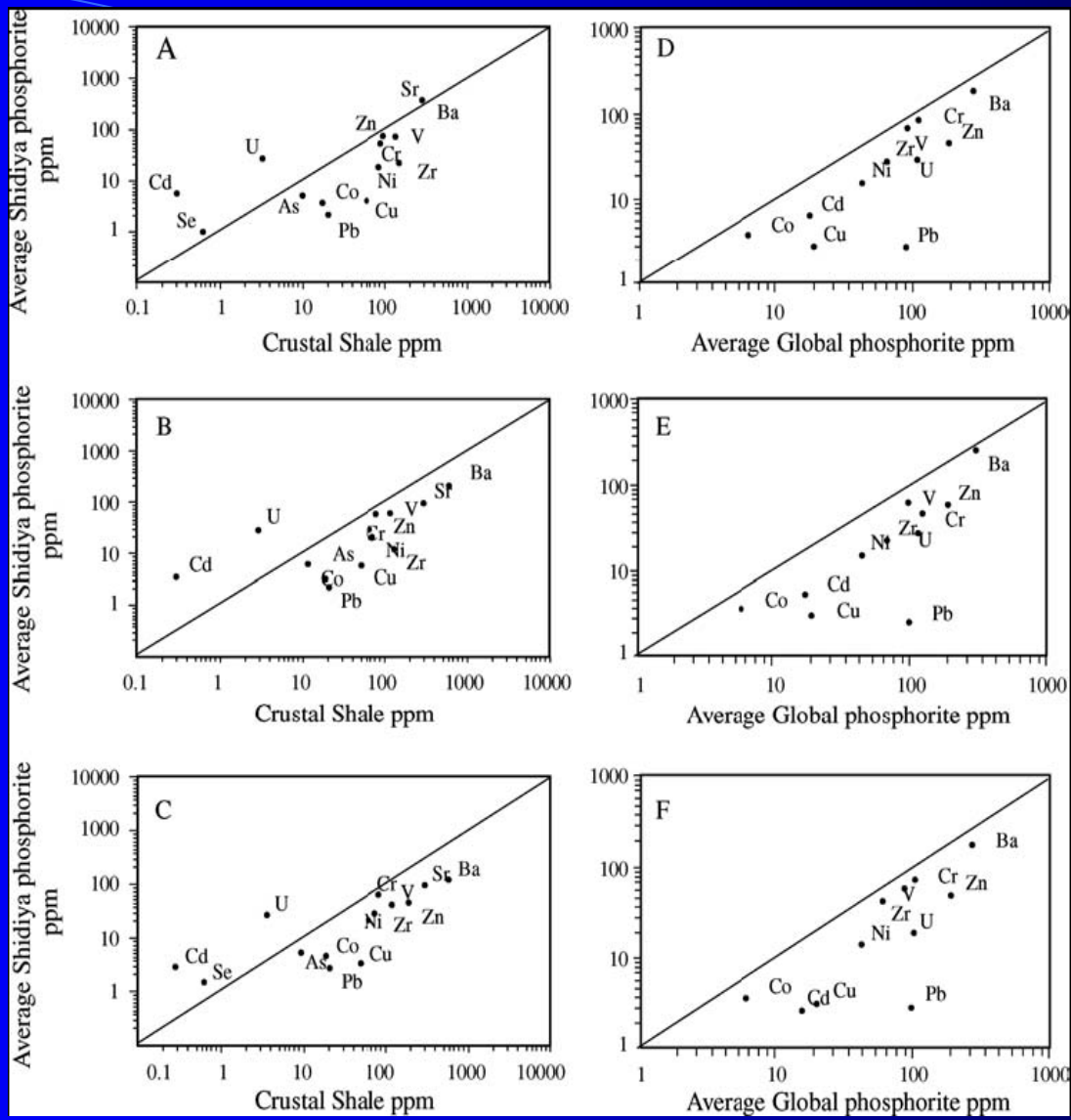
## Mean concentrations of major oxides (%) of phosphate from Eshidiya Mine

Variable	Bed-A1	Bed-A2	Bed-A3
<b>P<sub>2</sub>O<sub>5</sub></b>	<b>21.09</b>	<b>30.43</b>	<b>11.67</b>
<b>CaO</b>	<b>32.06</b>	<b>42.28</b>	<b>17.61</b>
<b>SiO<sub>2</sub></b>	<b>29.95</b>	<b>12.11</b>	<b>57.40</b>
<b>Al<sub>2</sub>O<sub>3</sub></b>	<b>2.40</b>	<b>1.81</b>	<b>1.37</b>
<b>Fe<sub>2</sub>O<sub>3</sub></b>	<b>1.77</b>	<b>1.44</b>	<b>1.32</b>
<b>MgO</b>	<b>0.31</b>	<b>0.19</b>	<b>0.20</b>
<b>Na<sub>2</sub>O</b>	<b>0.34</b>	<b>0.34</b>	<b>0.33</b>
<b>K<sub>2</sub>O</b>	<b>0.10</b>	<b>0.07</b>	<b>0.06</b>
<b>TiO<sub>2</sub></b>	<b>0.11</b>	<b>0.08</b>	<b>0.07</b>

## Mean concentrations of trace metals (ppm) of phosphate from Eshidiya Mine

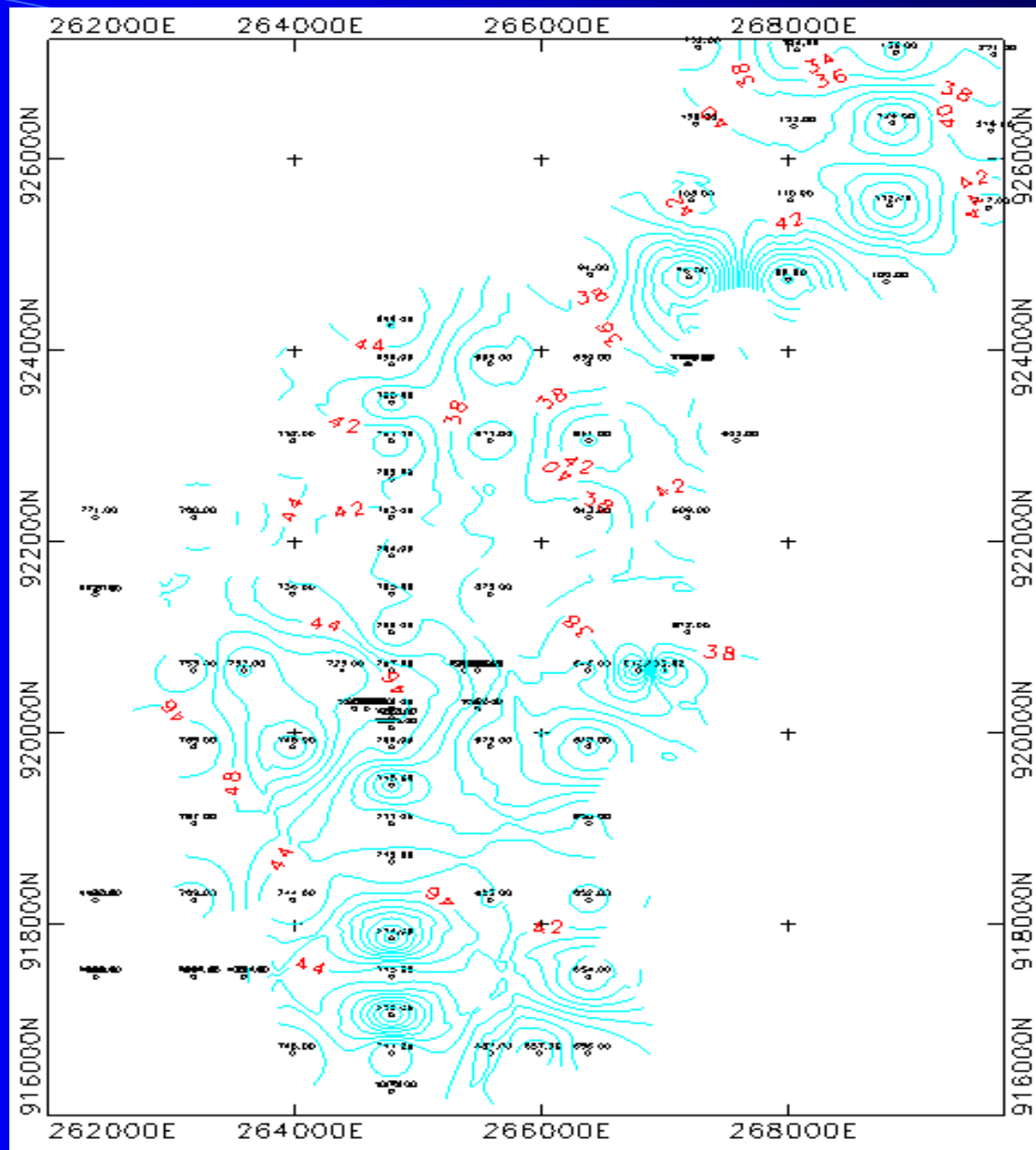
<b>Variable</b>	<b>Bed-A1</b>	<b>Bed-A2</b>	<b>Bed-A3</b>
<b>As</b>	<b>7</b>	<b>9</b>	<b>7</b>
<b>Cd</b>	<b>6</b>	<b>5</b>	<b>3</b>
<b>Co</b>	<b>4</b>	<b>4</b>	<b>3</b>
<b>Cu</b>	<b>4</b>	<b>5</b>	<b>3</b>
<b>Cr</b>	<b>91</b>	<b>57</b>	<b>81</b>
<b>Mn</b>	<b>572</b>	<b>662</b>	<b>524</b>
<b>Ni</b>	<b>22</b>	<b>20</b>	<b>17</b>
<b>Th</b>	<b>3</b>	<b>3</b>	<b>4</b>
<b>U</b>	<b>33</b>	<b>42</b>	<b>23</b>
<b>V</b>	<b>87</b>	<b>87</b>	<b>63</b>
<b>Zn</b>	<b>52</b>	<b>61</b>	<b>48<sup>10</sup></b>

<b>Average chemical analysis of U abundance in Worldwide sedimentary phosphate rocks.</b>			
<b>Country</b>	<b>Deposits</b>	<b>U</b>	<b>Refernces</b>
<b>Algeria</b>	<b>Djebel Onk</b>	<b>25</b>	<b>IFDC (1997)</b>
<b>Australia</b>	<b>Duchess</b>	<b>92</b>	<b>IFDC (1997)</b>
<b>China</b>	<b>Kaiyang</b>	<b>31</b>	<b>IFDC (1997)</b>
<b>Egypt</b>	<b>Abu Tartur</b>	<b>80</b>	<b>IFDC; TVA</b>
<b>Israel</b>	<b>Arad</b>	<b>150</b>	<b>IFDC (1997); TVA</b>
<b>Jordan</b>	<b>Central Jordan</b>	<b>65</b>	<b>Abed et al. (2008)</b>
	<b>Shidiya</b>	<b>46</b>	<b>IFDC (1997)</b>
<b>Eshidiya</b>	<b>A<sub>1</sub></b>	<b>33</b>	<b>This study</b>
	<b>A<sub>2</sub></b>	<b>42</b>	<b>This study</b>
	<b>A<sub>3</sub></b>	<b>23</b>	<b>This study</b>
<b>Morocco</b>	<b>Khouribga</b>	<b>88</b>	<b>Baechle and Wolsteen (1984)</b>
<b>Peru</b>	<b>Sechura</b>	<b>72</b>	<b>IFDC; TVA</b>
<b>Senegal</b>	<b>Taiba</b>	<b>64</b>	<b>IFDC</b>
<b>Syria</b>	<b>Khneifiss</b>	<b>75</b>	<b>IFDC (1997)</b>
<b>Togo</b>		<b>94</b>	<b>IFDC (1997)</b>
<b>Tunisia</b>		<b>44</b>	<b>IFDC; TVA.; Altschuler (1980)</b>
<b>United State</b>	<b>Central Florida</b>	<b>141</b>	<b>IFDC;TVA</b>
	<b>North Florida</b>	<b>81</b>	<b>IFDC;TVA; Wakefield (1980)</b>
	<b>Idaho</b>	<b>107</b>	<b>IFDC; TVA; Altschuler (1980)</b>
	<b>North Carolina</b>	<b>65</b>	<b>IFDC; TVA</b>



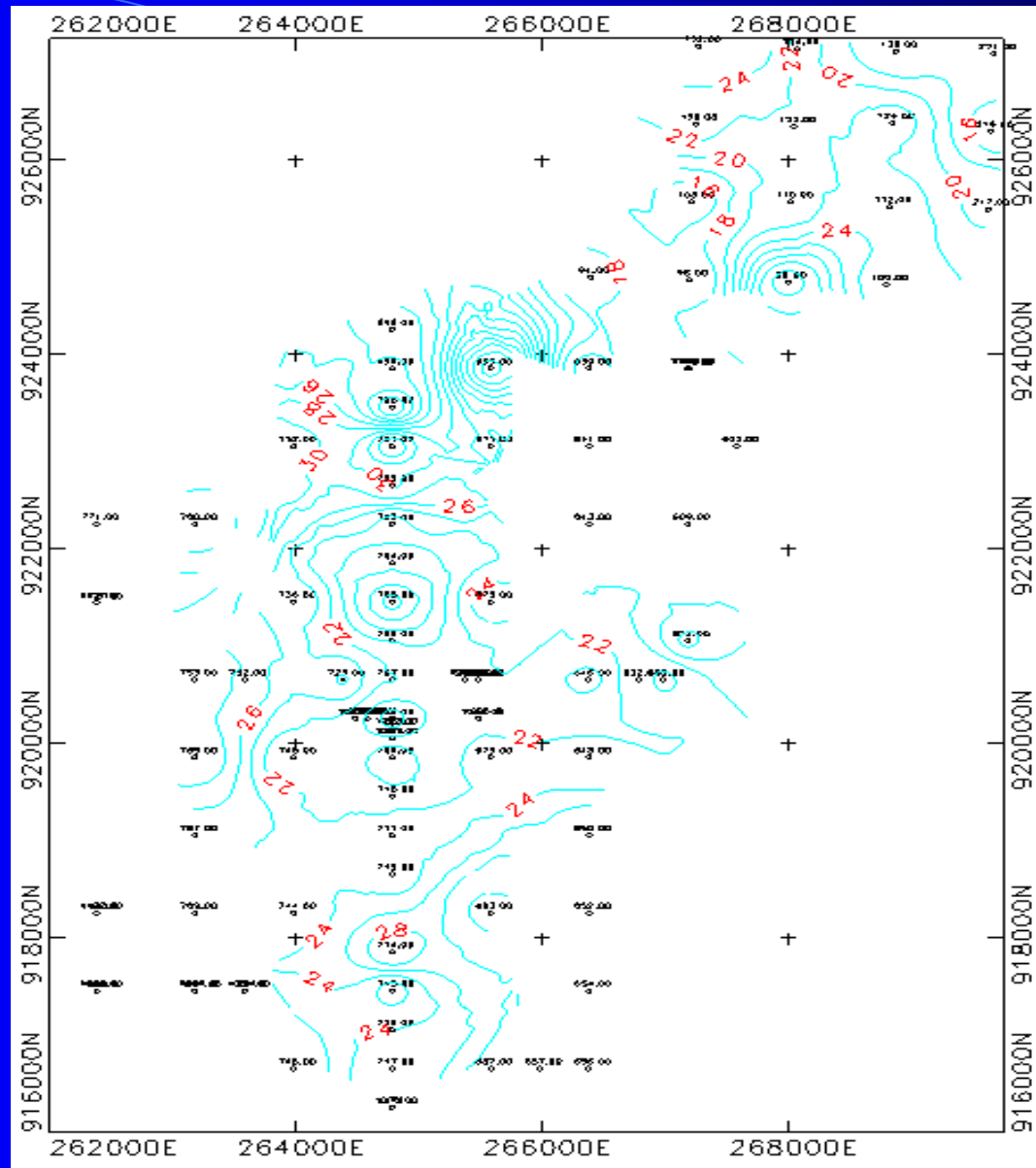
**A, B, C** Trace element composition of Eshidiya phosphorites beds A1, A2, and A3 respectively compared with shale (after Turekian and Wedepohl 1961). **D, E, and F** Trace element composition of Eshidiya phosphorites beds A1, A2, and A3 respectively, compared with average global phosphorites (after Altschuler 1980)





Contour maps showing dispersion patterns of U in bed A2 14

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Contour maps showing dispersion patterns of U in bed A3

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# Sampling Scheme and Results for Bed A1

Matrix weighs 1000 tons

Crushing and Screening

-12.5 mm (Product)

+12.5 mm (Reject)

**Soft Ph. Weighs (502.4 tons)**

Element	in ppm	in tons
U	59	0.03

**Hard Ph. weighs (497.6 tons)**

Element	in ppm	in tons
U	45	0.02

Beneficiation (Washing)

**-5+1 mm reject weighs (27.7 tons)**

Element	in ppm	in tons
U	61	0.002

**-12.5+5 reject weighs (107.5 tons)**

Element	in ppm	in tons
U	19	0.002

**-1+53μ final product weighs (139 tons)**

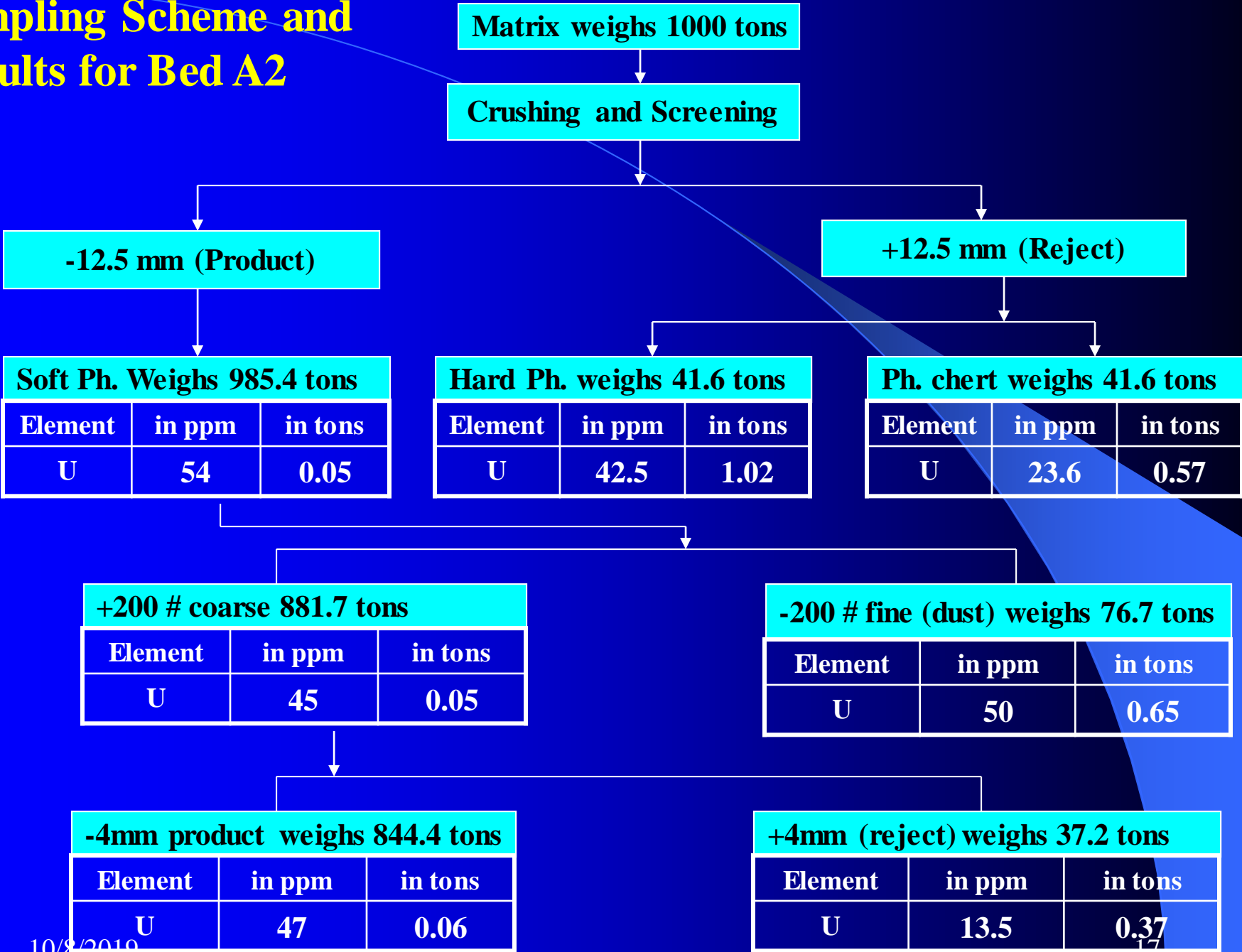
Element	in ppm	in tons
U	80	0.01

**-53μ slime weighs (147.4 tons)**

Element	in ppm	in tons
U	38	0.006



# Sampling Scheme and Results for Bed A2



**Soft Ph. Weighs 985.4 tons**

Element	in ppm	in tons
U	54	0.05

**Hard Ph. weighs 41.6 tons**

Element	in ppm	in tons
U	42.5	1.02

**Ph. chert weighs 41.6 tons**

Element	in ppm	in tons
U	23.6	0.57

**+200 # coarse 881.7 tons**

Element	in ppm	in tons
U	45	0.05

**-200 # fine (dust) weighs 76.7 tons**

Element	in ppm	in tons
U	50	0.65

**-4mm product weighs 844.4 tons**

Element	in ppm	in tons
U	47	0.06

**+4mm (reject) weighs 37.2 tons**

Element	in ppm	in tons
U	13.5	0.37

# Sampling Scheme and Results for Bed A3

Matrix weighs 1000 tons

Crushing and Screening

-12.5 mm (Product)

+12.5 mm (Reject)

Soft Ph. weighs (615.3 tons)

Element	in ppm	in tons
U	50	0.08

H. Ph. weighs (234.4 tons)

Element	in ppm	in tons
U	41	0.17

Ph. Chert weighs (41.6 tons)

Element	in ppm	in tons
U	24	0.58

Beneficiation (washing and Flotation)

-12.5+2 mm reject weighs (253 tons)

Element	in ppm	in tons
U	35	0.14

-53# slime weighs (42.3 tons)

Element	in ppm	in tons
U	28	0.66

-1+0.053mm product (320 tons)

Element	in ppm	in tons
U	32	0.10

-2+1mm reject weighs(48.6 tons)

Element	in ppm	in tons
U	35	0.72

-1+0.50 mm coarse weighs (200 tons)

Element	in ppm	in tons
U	35	0.18

-0.5+0.053 μ fine weighs (120tons)

Element	in ppm	in tons
U	30	0.25

Final Product

Element	in ppm	in tons
U	44	0.14

# Sampling Scheme and Results for phosphate fertilizer

**Matrix weighs 1000 tons**

Feed		
Element	in ppm	in tons
U	57.5	0.058

Grinding		
Element	in ppm	in tons
U	57.5	0.058

**React with sulphuric acid**

Phosphoric acid 30% weighs (950 tons)		
Element	in ppm	in tons
U	44	0.05

Phosphoric acid 54% weighs (560 tons)		
Element	in ppm	in tons
U	109	0.19

Phosphogypsum weighs (3800 tons)		
Element	in ppm	in tons
U	2	0.0005

**React with Ammonia (HNO3)**

DAP Fertilizer weighs (560 tons)		
Element	in ppm	in tons
U	126	0.2319

## Elements fate in bed A<sub>1</sub>.

Crushing and Screening		Beneficiation (Washing)	
Element	Fate	Element	Fate
As	Feed	As	Reject
Cd	Reject	Cd	Reject
Cr	Even	Cr	Reject
Cu	Feed	Cu	Reject
Mn	Feed	Mn	Reject
Ni	~ Feed	Ni	Reject
Pb	Feed	Pb	Reject
<b>U</b>	<b>~ Feed</b>	<b>U</b>	<b>Final product</b>
V	Even	V	Reject
Zn	~ Feed	Zn	Reject

## Elements fate in bed A<sub>2</sub>.

Crushing and Screening		Drying	
Element	Fate	Element	Fate
<b>As</b>	Reject	<b>As</b>	Final product
<b>Cd</b>	Reject	<b>Cd</b>	Final product
<b>Cr</b>	Even	<b>Cr</b>	Final product
<b>Cu</b>	Feed	<b>Cu</b>	Final product
<b>Mn</b>	Feed	<b>Mn</b>	Final product
<b>Ni</b>	Feed	<b>Ni</b>	Even
<b>Pb</b>	Reject	<b>Pb</b>	Even
<b>U</b>	<b>Reject</b>	<b>U</b>	<b>Final product</b>
<b>V</b>	Even	<b>V</b>	Final product
<b>Zn</b>	Even	<b>Zn</b>	Final product

## Elements fate in bed A<sub>3</sub>.

Crushing and Screening		Beneficiation (Washing and Flotation)	
Element	Fate	Element	Fate
<b>As</b>	Even	<b>As</b>	Reject
<b>Cd</b>	Reject	<b>Cd</b>	Reject
<b>Cr</b>	Feed	<b>Cr</b>	Reject
<b>Cu</b>	Feed	<b>Cu</b>	Reject
<b>Mn</b>	Feed	<b>Mn</b>	Reject
<b>Ni</b>	Feed	<b>Ni</b>	Reject
<b>Pb</b>	Reject	<b>Pb</b>	Reject
<b>U</b>	<b>Feed</b>	<b>U</b>	<b>Final product</b>
<b>V</b>	Feed	<b>V</b>	Reject
<b>Zn</b>	Feed	<b>Zn</b>	Final product

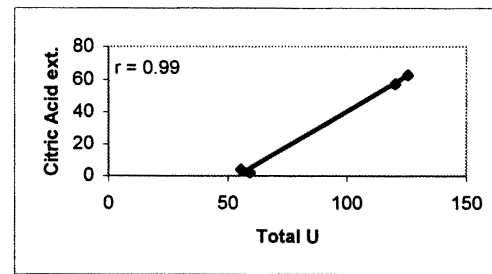
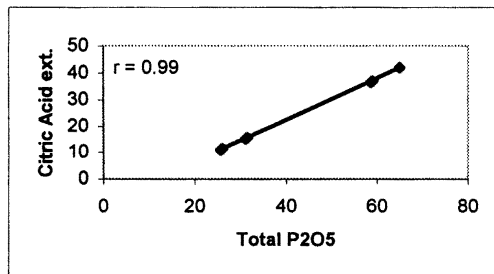
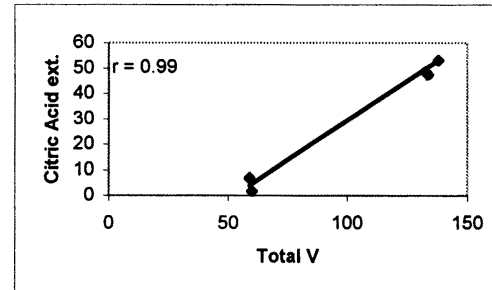
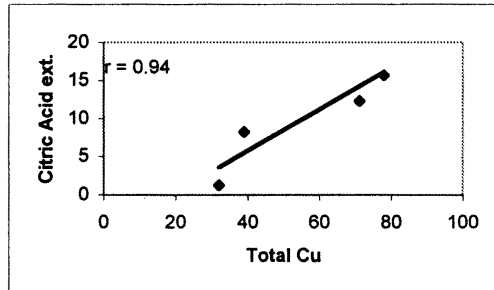
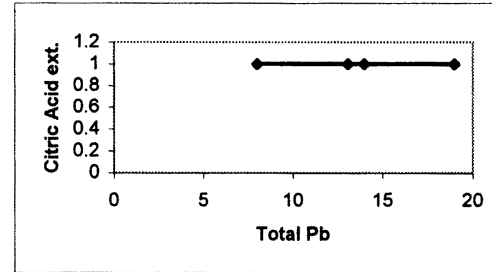
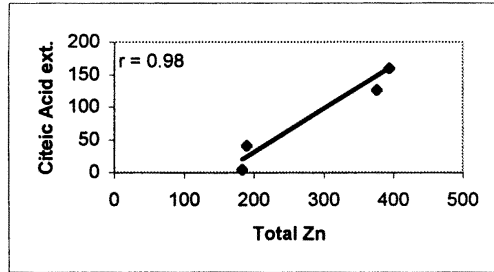
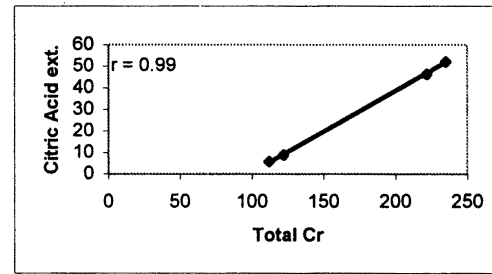
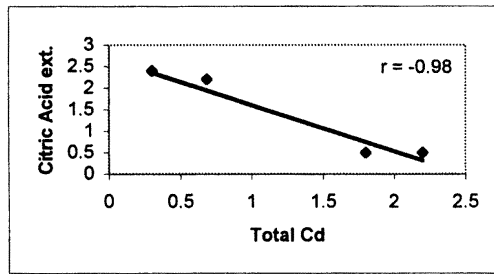
# Elements fate in phosphate fertilizer

Phosphoric Acid 30% and Gypsum		Phosphoric Acid 54% and DAP	
Element	Fate	Element	Fate
<b>As</b>	<b>Gypsum</b>	As	<b>DAP</b>
<b>Cd</b>	<b>Acid</b>	<b>Cd</b>	<b>Acid</b>
<b>Cu</b>	<b>Acid</b>	Cr	<b>DAP</b>
Cr	<b>Acid</b>	<b>Cu</b>	<b>DAP</b>
<b>Ni</b>	<b>Acid</b>	Ni	<b>Even</b>
<b>U</b>	<b>Acid</b>	<b>U</b>	<b>DAP</b>
<b>V</b>	<b>Acid</b>	V	<b>Even</b>
Zn	<b>Acid</b>	<b>Zn</b>	<b>DAP</b>

# Leaching experiments from raw phosphate to DAP

Leaching agent	Material	P <sub>2</sub> O <sub>5</sub>			U		
		T %	cx %	% T	T %	cx %	% T
0,1m EDTA pH = 4.5 0,1m EDTA pH = 4.5	Raw ph.	25.4	11.5	45	53	-	-
	DAP	63.2	36.5	58	128	-	-
	DAP/Raw ph.	2.5	3.2	1.3	2.4	-	-
Leaching agent	Material	P <sub>2</sub> O <sub>5</sub>			U		
		T %	cx %	% T	T %	cx %	% T
1 % Citric acid pH = 2.4	Raw ph.	25.4	11.0	43	53	17	32
	DAP	63.2	40.3	64	128	96	75
	DAP/Raw ph.	2.5	3.2	1.5	2.4	5.6	2.3





**Relationship between the total of heavy metals content and their metal extracted using citric acid from phosphate fertilizers.**

## Correlation coefficients as a measure of the degree of mobility

<b>Ranges (r)</b>	<b>Degree of Mobility</b>
0.01-0.30	low mobility (L. M)
0.30-0.59	Intermediate (I. M)
0.60-1.00	high mobility (H. M)

## Degree of elements mobility from fertilizer basd on EDTA and Citric acid extraction

Element	Techniques	(r ) Values	Degree of Mobility	Overall assessment
<b>P<sub>2</sub>O<sub>5</sub></b>	<b>EDTA</b>	-	-	
	<b>Citric acid</b>	<b>0.99</b>	<b>H. M.</b>	<b>H. M.</b>
<b>Cd</b>	<b>EDTA</b>	<b>0.29</b>	<b>L. M.</b>	
	<b>Citric acid</b>	<b>0.98</b>	<b>Immobile</b>	<b>L. M.</b>
<b>Zn</b>	<b>EDTA</b>	<b>0.95</b>	<b>H. M.</b>	
	<b>Citric acid</b>	<b>0.98</b>	<b>H. M.</b>	<b>H. M.</b>
<b>Cu</b>	<b>EDTA</b>	<b>0.96</b>	<b>H. M.</b>	
	<b>Citric acid</b>	<b>0.94</b>	<b>H. M.</b>	<b>H. M.</b>
<b>Cr</b>	<b>EDTA</b>	<b>0.80</b>	<b>H. M.</b>	
	<b>Citric acid</b>	<b>0.99</b>	<b>H. M.</b>	<b>H. M.</b>
<b>Pb</b>	<b>EDTA</b>	<b>0.07</b>	<b>L. M.</b>	
	<b>Citric acid</b>	<b>0.00</b>	<b>Immobile</b>	<b>L. M.</b>
<b>V</b>	<b>EDTA</b>	<b>0.90</b>	<b>H. M.</b>	
	<b>Citric acid</b>	<b>0.99</b>	<b>H. M.</b>	<b>H. M.</b>
<b>U</b>	<b>EDTA</b>	-	-	
	<b>Citric acid</b>	<b>0.99</b>	<b>H. M.</b>	<b>H. M.</b>

Abbreviation: L.M.= low mobility, H.M.= high mobility and I.M.= intermediate mobility

# CONCLUSIONS AND RECOMMENDATIONS

# CONCLUSIONS

## I

# Geochemical findings

### **III Geochemical findings:**

**1- Based on the  $P_2O_5$ , CaO and fluorine contents the dominant Eshidiya phosphorite mineral phase is francolite.**

## CONCLUSIONS

### III Geochemical findings:

**2- U is accommodated in the francolite mineral phase**

**3- U is enriched in the raw phosphates when compared with their abundances in shale, while they depleted when compared with average global phosphorites.**

# **II**

## **Redistribution and mobility findings**



**1. Uranium fates during crushing and screening process of the three phosphatic beds are variable, whereas during beneficiation and drying they exhibit some consistency.**

**2. U redistribution pathways during the phosphate processing steps were determined in the feed and in the final product. During crushing and screening process, U measured elements got enriched in the feed bed A1 and A3, whereas U got enriched in the reject bed A2. During beneficiation process, most U elements got enriched in the final product of three beds A1, A2 and A3**

**3- Uranium (ppm) fate during crushing and screening and beneficiation processes allow drawing the following conclusions:**

	<b>Final product</b>	<b>Hard Phosphate</b>
<b>A1</b>	<b>80</b>	<b>45</b>
<b>A2</b>	<b>47</b>	<b>43</b>
<b>A3</b>	<b>44</b>	<b>41</b>

**4. Uranium redistribution pathways during the phosphate fertilizer processing steps were determined in the feed and in the final product. U measured elements got enriched in the phosphoric acid (30%), phosphoric acid (54%) and DAP product. It was found that 93% of the U is partitioned in the phosphoric acid while the remainder is precipitated with PG waste materials.**

**5- Uranium (ppm) fate during phosphate fertilizer  
allow drawing the following conclusions:**

	<u><b>U (ppm)</b></u>
<b>Phosphoric acid (30%)</b>	<b>44</b>
<b>Phosphoric acid (54%)</b>	<b>109</b>
<b>DAP</b>	<b>126</b>
<b>Phosphogypsum</b>	<b>2</b>

**6. The assessment of the availability (mobility) of U elements from DAP to plant through soil using Citric Acid techniques allow drawing the following conclusions:**

**a- Based on citric acid, U elements with exhibit high mobility in acidic soils ( $\text{pH} \geq 2.4$ )**

**b- The mobility of U elements will decrease strongly in alkaline soils ( $\text{pH} \geq 7$ ).**

**7. The assessment of the variability of toxic elements to plants using Citric Acid leaching techniques allowed to show the following conclusion: U is going to be available for plants even in acidic soils, and will be available upon direct application of phosphate to soils as well.**



***THANK YOU FOR  
YOUR  
ATTENTION***

