## <u>CHAPTER - VII</u> <u>PETROCHEMISTRE</u>

#### SCOPE

Only a limited chemical study has been undertaken, and the data obtained has been utilised for properly evaluating certain specific phenomena related to the metamorphism of the area. Chemical analyses of ortho-amphibolites, hornblende gneisses, biotite gneisses and pelitic schists, have been considered (Fig. VII.1) to understand the following aspects :

- 1) True nature of the magma which gave rise to ortho-amphibiotites.
- 2) Derivation of hornblende gneisses
- 3) Origin and nature of biotite gneisses
- 4) Nature of the original metasediments that gave rise to pelitic schists.

Fig.VII.1. 127



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Representative samples were analysed and the data plotted on various appropriate diagrams. The information obtained from the chemical data has been interpreted in conjunction with petrography and field details. Chemical analyses of the meta-dolerites and alkali-basalt have been given just for the same of information and to ascertain their probable parentage.

#### ORTHO-AMTHIBOLITES

These metamorphosed linear bodies occurring as fold cores represent the oldest rocks and have been interpreted as the oceanic basement (= plate) over which the Ajabgarhs were deposited. The chemical composition of 5 representative samples is given in the Table VII.1; and the percentage of major elements and their atomic proportions, computed from the major oxides, are given in Table VII.2. The overall oxide percentages point to the alkaline nature of the original-rock. The ratio between  $Fe_2O_3$  and Feo (Table VII.1) is more than 0.2, and that between Na and K (Table VII.2) is less than 10. These two are obvious indications of a basic magma of alkali basalt type (Engel et al., 1965, p. 719). The original alkaline nature of the parent rock also becomes clear on the graph showing weight percent of SiO<sub>2</sub> on arithmetic scale against that of K<sub>2</sub>O on logarithmic scale (Fig. VII.2); all the plots lie in the alkali basalt region (Engel et al., 1965, p. 728).

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	CHEMICAL ANALYSES OF OR THO-AM PHIBOLITES								
ange Gain Dino GRA Don A	1	2	3	4	5	Average of 5(from st- udy area)	Average* of 200		
	40.25	<u>БО Б</u> б	50 30	1:6. 03	57 10	19.51	50 30		
$Al_2O_2$	14.33	15.57	16.43	15.70	16.53	15.71	15.70		
Fe <sub>2</sub> 0 <sub>3</sub>	4.76	4.16	2.12	4•52/	2.96	3.70	3.60		
FeO	8.09	8.08	8.22	9.33	10.69	8.88	7.80		
MgO	7.95	6.37	7.89	6.91	6.22	7.07	7.00		
CaO	8.08	7.21	8.94	9•45	9.80	8.70	9.50		
Na <sub>2</sub> 0 <sup>′</sup>	3.20	3.92	3.30	2.97	2.77	3,•23	2 •90		
K20	1.82	1.72	1.93	0.81	1.62	1.58	1.10		
Total	97。48	97.59	99.13	95.72	101.99	98.38	97.90		

TABLE VII.1

A. Poldervaart, Geol. Soc. Am. Spec. Paper 62, p. 136; 1955 (in Walter T. Huang 1962 - P. 418) \*

The values of atomic proportions of the elements like Si and Al (Table VII.2) also furnish supporting evidence; a Si content that varies between 21.73 and 24.26, with an average of 23.37, when considered in conjunction with Al values averaging 8.32, also suggest the derivation of ortho-amphibolites from a magma of oceanic affinity. The Fe and Mg values further substantiate their oceanic setting. The FMA diagram (Fig. VII.3) after Ringwood (1975) provides an

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interesting tholeiitic differentiation trend, the FMA values from the study area being computed in Table VII.3. May be that the parent magma which gave rise to the alkali basalt, was tholeiitic. Of course, this is just a conjecture.

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MAJOR ELEMENTS AND THEIR ATOMIC PROPORTIONS (ORTHO-AMPHIBOLITES)

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					•		
Si	23.25	23.86	23.74	21.73	24.26	23:•37	23.74
Al	7.59	8.24	8.70	8.31	8.75	8.32	8.31
Fe <sup>+3</sup>	3.33	2.91	1.48	3.16	2.07	2.59	2.52
$Fe^{+2}$	6.29	6.₀28	6.39	7.25	8.31	6.90	6.06
Mg	4.080	3.84	4.76	4.17	3.75	4.26	4.22
Ca	5•77	5.15	6.39	6.75	7.00	6.22	6.•79
Na	2 • <b>3</b> 7	2.91	2.45	2.20	2.05	2.40	2.15
K	1.29	1.22	1.37	0 <b>。57</b>	1.15	1.12	0.78
Total catio	n <sup>54,69</sup>	54.41	55.28	54.14	57•34	55.18	54•57
Total oxyge	n <sup>42</sup> •79	43.18	43.85	41.58	44.65	43₀20	4333
Total cation +oxyge	⊧97 <u>∘4</u> 8 n	97.59	99.13	9 <b>5.7</b> 2	101.99	98.38	9 <b>7</b> .90



### FMA DIAGRAM SHOWING CHARACTERISTIC THOLEIITIC AND CALC-ALKALINE TRENDS (After Ringwood, 1975).



Wt %

Curve S represents the skaergaard differentiation trend and Curve T the Thingmuli.

Shaded calc-alkaline band represents the differentiation trends displayed by magmas from the Cascade, Aleutian, and New Zealand calc-alkaline provinces.

----- Curve shown by the orthoamphibolites of the study area.

TABLE VII.3 (ORTHO-AMPHIBOLI FMA VALUES 1 2 35 4 5 (FeO+Fe<sub>2</sub>O<sub>3</sub>) 49•77 50.47 56.44 56.27 44.07 M (MgO) 30.79 26.27 33.63 28.16 25.64 Alk (A) ( $K_2$ 0+Na<sub>2</sub>0) 19.44 23,.26 22.30 15.40 18.09

For the purposes of comparison, the author has given in Table VII.4, the average values of weight percentage of oceanic alkali basalts from the main oceans of the world (Manson, 1968)

TAB	LE.	VII	•4
		-	-

AVERAGE VALUES OF THE OCEANIC ALKALIC BASALT						
Major oxides	Pacific, Atlantic and Indian oceans (After Manson, 1968)	Study area				
sio <sub>2</sub>	46 ₀ 9	49•51				
A1203	15.5	15.71				
Fe203	- 3.1	3 <b>.7</b> 0				
FeO	8.6	8.88				
MgO	6. 9	7.07				
CaO	10.4	8.70				
Na20	<b>3.</b> 0	3.23				
K20 ~	1.3	1.58				

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Considering the above facts, it can be concluded that the amphibolites of the study area comprise metamorphosed derivatives of oceanic basalt of alkaline nature.

#### HORNBLENDE GNEISSES

These streaky hornblendic rocks consist of alternating layers rich and poor in hornblende. The leucocratic hornblende poor layers comprise quartz, oligoclase and microcline. On the other hand, hornblendic layers contain in addition to hornblende, and sine and quartz. The chemical data of hornblendic bands only are given in Table VII.5.

	CHEMICAL AN	ALY SES OF HOR	NBLENDE GNEIS	<u>SSES</u>
Sample No	o. 6	.7	8	9
-				
sio <sub>2</sub>	56.9	9 61.46	60.40	66.14
Al203	15.8	2 16.72	15.07	13.12
Fe203	4.3	2 3.85	3.96	5.28
FeO	3.8	4 2.64	3.79	1.72
MgO	4.4	3 4.15	- 3-75	3.15
CaO	2.1	0 3.08	1.54	1.40
Ma20	4.3	5 3.82	3 <b>.7</b> 1	3.087
K20	5•4	3, 4.25,	4.31	4.87
Total	97.2	8 99.97	96.53	99 • 55
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TABLE VII.5

It is observed that the chemical composition is broadly comparable with that of the amphibolites, and no significant chemical change is revealed. This leads to an important conclusion that the leucocratic layers definitely represent the granitic material injected lit par lit in the amphibolites, the granitic constituents having been derived from the overlying granitised metasediments. Actual granitisation transformation of amphibolite to biotite gneiss is nowhere seen either in the field as is revealed in the chemical data.

#### BIOTITE GNEISSES

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The chemical analyses of the biotite gneisses are most revealing (Table VII.6), and typically point to their derivation from the pelitic schists (Table VII.7).

## TABLE VII.6

			· · · ·
Sample No.	10	1.1	12
Si0 <sub>2</sub>	61.94	62.45	<b>7</b> 0.00
A1203	14.32	14.011	14.76
Fe203	3.14	3.41	3 <b>.7</b> 0
FeO	1.48	0.88	0.80
MgO	4.36	4.35	2.12
CaO	3.36	3•49	1.35
Na <sub>2</sub> 0	419	4.99	4.24
к <sub>2</sub> 0	4.024	5.62	2.75
Total	97.03	99.30	99 <b>.7</b> 2

CHEMICAL ANALYSES OF BIOTITE GNEISSES

As compared to the coarse non-foliated biotitic granite (Erinpura granite proper), these (which were originally supposed to be foliated varieties of the former) show a relatively high percentage of MgO and FeO (Table VII.12). On the other hand, the chemical composition of gneisses compares well with those of biotite schists etc.

A critical appraisal of the chemical data of all the three varieties leads to the following conclusions :

- 1. The biotite gneisses have chemical composition almost comparable to that of the overlying meta-sediments.
- 2. No significant addition of any constituent is seen in the gneisses, and the transformation of schists to granite gneisses appears to be a transformative process of internal readjustment of constituents only.
- 3. The relatively high percentage of MgO in gneisses reflects the original magnesian richness of meta-sediments.

#### PELITIC SCHISTS

The chemical composition of biotite schists and related rocks (Table VII.7) throws some light on the original composition and nature of the meta-sediments.

#### TABLE VII.7

CHEMICAL ANALYSES OF PELITIC SCHISTS

Sample No	13,	14	15;	16.	17	18	19
SiO <sub>2</sub>	70.10	65.34	65.85	66.14	67.69	72.86	67.29
Al203	12.31	12.46	12.35	18.01	13.67	14.21	12.56
Fe203	3.97	5.68	3.63	2.88	5,07	1.46	4.04
FeO	1.28	1.00	0 <b>.</b> 96	1.00	1.72	1.20	1.04
MgO	1.72	1.10	2.17	1.45	1.77	1.99	3.65
CaO	0.56	4.35	3:•50	0.28	0.70	0.28	3•93
Na <sub>2</sub> 0	4.08	3.19	3.82	3₀98	3.13	2.92	5.04
К <sub>2</sub> 0	5.43	4.87	6.05	5.99	5.68	4.68	3.87
Total	99.45	97.99	98 <b>.33</b> )	99•73	99.43	99 <b>.6</b> 0	101.42

The data shows that the sediments were mostly graywackes to subgraywackes; with a fair MgO content. Pettijohn (1975, P.228) has given the chemical composition of representative graywackes (Table VII.8) from which it will be seen that the composition of the metasediments of the study area also point to their graywacke affinity.

The Al<sub>2</sub>O<sub>3</sub>:SiO<sub>2</sub> variation diagram (Fig. VII.4) based on the molecular percentages (Table VII.9) shows that the chemical composition of the pelitic schists as well as biotite gneisses is identical to that of graywackes - subgraywackes,

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TABLE VII.8

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	CHEMI(	CAL CON	IPOSITI	LON OF	REPRES	SENTAT:	IVE GR.	AYWACKI	<u>ES</u>	
الانتاز بلقاع مربور ججو الانتبر اللقاء الأست			(Afte	er Peti	tijoh <b>n,</b>	1975) /				
Consti- tuent	A	В	C	D	E	F	G	H	I	J
Si02	60.51	66.24	76.84	69.11	6 <b>8.</b> 85	74.43	70.60	71,10	68.84	65.05
TiO <sub>2</sub>	00.87	00.64	-	00.60	00.74	00 <b>.83</b> )	00.70	00.50	00.25	00 <b>.4</b> 6
A1203	15.36	15.28	11.76	11.38	12.05	11.32	13.50	13.90	14.54	13.89
Fe203	00.76	00.70	00.55	1.41	2.72	00.81	`2.20	Trace	00.62	00.74
FeO	7.63	4 • 53	2.88	4.64	2.03	3.88	1.60	2.70	2.47	2 .60
MnO	00.16	00.06	Trace	00.17	00.05	00.04	00.10	00.05	Nil	00.11
MæO	3.39	2.74	1.39	2.06	2.96	1.30	1.60	1.30	1.94	1.22
CaO	2.14	1.70	00.70	1.15	00,50	(j.17	1.30	1.80	2.23	5 •62
Na <sub>2</sub> 0	2.50	3.12	2.57	3.20	4.87	1.63	2.90	3.70	3 ,88	313
К <sub>2</sub> 0	1.69	1.91	1.62	1.76	1.81	1.74	1.60	2.30	2.68	1.41
⊞ <sub>2</sub> 0 <sup>+</sup>	3 <b>.3</b> 8	2.49	1.87*	+ 4.13	2.30	2.15	2.80	1.90	1.60	2.30
H <sub>2</sub> 0	0.15	00.08	3 -	00.05	00.77	00.20	00.3;0	00.26	00.35	00.28
P205	00.27	00.12	-	00.03;	00.06	00.18	00.20	00.10	00.15	00.08
Zr02	-		-	-	-	-	~		00.05	-
CO2	1.01	00.30	-		00.08	00.48	00.60	00.12	00.14	2.835
50 <sub>3</sub>	-			-	-	~	·	~ <b>an</b>	00.15	
S	00,42		<b></b>	<b></b>	00.08	00.12		Trace		00.05
BaO	-	-			Trace		-	- 、	00.04	~
<b>G</b> <sup>s</sup>	-		<b></b>	-	00.07	00.17	-	00.09	-	
Total	100.24	99•99	109 <b>.1</b> 8	99.69	99.94	100.4 5	00.00	99.80	99•93	99.77

\* Loss on igniton

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Table VII.8 Contd.

- A Archean, Manitou Lake, Ontario. B. Brunn, analyst (Pettijohn, 1949, p. 250)
- B. Average of three Burwash Formation graywackes (Archean), Yellowknife Bay, Northwest Territories, Canada. L. Seymour, analyst (Henderson, 1972, p. 890)
- C Tyler Slate (Animikean), Hurley, Wisconsin. H.N. Stokes, analyst (Diller, 1898, p. 87)
- D (?) Ordovician, Rensselaer, near Spencertown, New York, H.B. Wiik, analyst (Balk, 1953, p. 824).
- E Tanner Graywacke (Upper Devonian Lower Carbonifierous), Scharzfeld Germanry, R. Helmblod, analyst (Helmbold, 1952, p. 256).
- E Carboniferous graywacke from Stanley Shale, near Mena, Arkansas. B. Brunn, analyst (Pettijohn, 1957, p. 319)
- G Average of six Kulm graywackes from Oberharz, Germany (Mattiat, 1960, Table 14)
- H Lower Mesozoic composite sample prepared by using equal parts of 20 graywackes exposed along shoreline between Palmer Head and Hue-Te-Taka, Wellington, New Zealand. J.A. Richie, analyst (Reed, 1957, p. 16).
- I Franciscan (? Jurassic) Quarry. Oakland Paving Co., Piedmont, California. J.W. Howson, analyst (Davis, 1918, p. 22)
- J Eocene, near Solduc, Olympic Mountains, Washington. B. Brunn, analyst (Pettijohn, 1949, p. 250).





Points p,g and a represent the compositions of average pelite, greywacke and arkose respectively (after Pettijhon, 1957)

mostly all plotted points falling within the gray,wacke field of Pettijohn (1957). The rocks thus plotted, also include the so called "Schistose quartz porphyries" of Sharma (1931) and obviously they do not represent a eruptive rock.

## TABLE VII.9

		MOIE	CULAR	PERCEN	VT AGES	OF SiC	2 and	Al203		
		( <u>BÌ</u> C	OTITE G	NEISSE	S AND	PELITI	CSCH	(STS)		
		•								
Samp- le No	10	11	12	13)	14	15;	- 16	17	18	19
si0 <sub>2</sub>	69.02	68.29	76.68	77•36	73j•03j	7523,	74.42	75•93	79.26	71.45
A1203	9.41	9.10	9.51	7.99	8.23)	8.34	11.94	9.06	9.12	° 7.87
term frames into stree is										

In this connection, the diagram (Fig. VII.5) showing Kohler Raaz values for pelitic schists (Table VII.10) clearly demonstrates the sedimentary parentage of these rocks.

#### META-DOLERITES AND ALKALI BASALTS

The dolerite sills intruded into the metasediments prior to the coasegrained Erinpura granite, as also the narrow basaltic Post-Erinpura dykes, reveal alkaline nature in their chemical composition (Table VII.11). It is most likely that they have been intruded by the remobilization of the oceanic crust during successive deformational events related to the upheaval.

# KÖHLER-RAAZ DIAGRAM



Pelitic schist

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### TABLE VII.10

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Sample No.	Case	qZ	Ē	fm		
13;	II	46.07	35.39	18.54		
14	I	39.17	34.51	26.31		
15	II	29.89	39.71	30.40		
16	III	34.•38	36.29	29•33		
17	III	41.22	36.77	22.01		
18	III	60.15	20.07	19.77		
19.	° I	24.82	39.20	35.98		

# KOHLER RAAZ VALUES FOR PELITIC SCHISTS

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TABLE VII.11

	CHEMICAL ANALYSES OF BASIC ROCKS							
	20+	21 <sup>+</sup>	22+	23+	24*			
Si0 <sub>2</sub>	53.04	50.36	51.00	52.00	48.70			
Al203	18.02	18.42	18,28	16.37	19.50			
Fe203	3.28	4.12	3.96	2.76	3.45			
FeO	8.53	9.00	9.38	9.83	9.24			
Mg,O	6.10	6.80	6.82	6.36	6.49			
CaO	8.80	6.24	6.66	7.43	7.38			
Na20	3.14	3.46	3.09	3•45	3.29			
К <sub>2</sub> 0	0.95	1.01	0.81	0.91	0.80			
Total	101.86	99.41	100,00	99.11	98.85			

+ Post-Delhi (Pre-Erinpura)

\* Post-Erinpura.

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CHEMICAL ANALYSES OF GRANITES

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التوقع المحق المحق التاريخ المحق	25	26	27	28	29	3,0
SiO <sub>2</sub>	66.89	72•47	72.13	71.66	64.29	70 <b>.4</b> 2
<sup>A1</sup> 2 <sup>0</sup> 3 Fe <sub>2</sub> 0 <sub>3</sub>	1.81	0.61	0.55	14 •22 0 <b>•</b> 29	1.70	1.39
FeO	3.24	3.24	2.12	1.80	3. <b>.</b> 80	2.16
Mg0 Ca0	0.29 2.21	0.92 2.60	0.59 2.63	2.08	0.36 2.63	0,34 2,49
Na20	3.34	2.61	2.71	2.60	2.60	2.87
K20	5.93	5.31	5.56	5.12	5.24	5•37
Total	100.12	100.97	99 <b>•</b> 55	98.27	98.38	98.60

The chemical analyses of Calcareous rocks in Table VII.13 and 14 have been given to ascertain the composition of the original sediments and to calculate the A, C, & F values plotted in Fig. VII.1.

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# TABLE VII.13

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Sample No.	31, i	32	33	34	35
SiO2	0.7	48	1.0	1.0	9.3
A1203	0.3	2.8	0.2	0.6	1.9
₽e₂03	0.4	0.2		0.4	1.6
FeO	-	-		integr	
MgO	0.7	3.3	0.7	0.8	0 <b>.</b> 4
CaO	54.9	. 50.2	55.2	549	50.2
Na20	· 🛥		·	8000	***
K <sub>2</sub> O	-	-		400	
loss on ignition	42.6	38.9	42.7	42.6	35.6
Total	99.6	100.2	99.8	100.3	99.0

CHEMICAL ANALYSES OF MARBLES

TABLE VII.14

CHEMICAL ANATY	SES OF CAL	-SCHIST AND PARA	AMHIBOLITES
Sample No.	36 <sup>@</sup>	3,7*	3;8 <sup>*</sup>
Si0 <sub>2</sub>	48.81	47.50	57.78
Al <sub>2</sub> 0 <sub>3.</sub>	19.13	18.13	21.51
Fe <sub>2</sub> 0 <sub>3</sub>	6.29	13.79	0.62
FeO	5.12	1.96	3.40
MgO	5.60	5.00	2.15
CaO	4.91	3:•22	2.10
Na <sub>2</sub> 0	5.25	472	3.18
K20	0.56	2.75	4.37
Total	95.67	97.07	95.11

@ Calc-schist

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\* Para-amphibolite