Mineralogical constraints of the Shergarh Sar amphibolites, Allai Kohistan, northern Pakistan

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ABSTRACT: Shergarh Sar amphibolites, a part of the southern amphibolite belt, are widely exposed in the area around Bana in Allai Kohistan. Epidote-amphibolite is the main unit with scattered patches of garnet-bearing variety. Amphibole, epidote and ,in some, garnet are the main phases with minor amount of quartz, plagioclase, clinopyroxene, opaque oxides and rutile. Amphiboles range in composition from tchermakitic-hornblende to actinolite with magnesio-hornblende as an intermediate phase. This transition is marked by decrease in "Al Ti, alkalies and Fe²⁺/Mg ratio and increase in Si. Epidote occurs as large crystals (both zoned and unzoned) with low Ps contents (9-15) and as small granular aggregates with high Ps contents (17-27). Garnet occurs sporadically in the form of porphyroblasts and is found in association with leucocretic veins. It is chemically unzoned and is predominantly made up of almandine-grossular-pyrope solid solution. Plagioclase ranges from An₂₇ to pure albite and clinopyroxene is chemically homogeneous diopside.

Petrographic observations and thermobarometry suggest that the rocks were initially equilibrated under amphibolite facies with overprinting of greenschist facies assemblage. This retrogression may have been facilitated by shearing during the obduction (uplift-cooling) of the rocks along the Indus suture following their Eocene collesion with the Indian plate.

INTRODUCTION

Amphibolites are the most voluminous rocks in the Shergarh Sar area north of Bana town in Allai Kohistan (Fig.1). These amphibolites are considered to be a part of the southern (Kamila) amphibolite belt of Kohistan (Jan, 1988). This belt of possibly late Jurassic to early Cretaceous age is composed mainly of volcanic and plutonic precursors with rare siliceous and calcareous metasediments (see Jan, 1979). It overlies ultramafic rocks and tectonic melange of the Indus suture zone, but in some places it thrusts over rocks of the Indian plate.

The Shergarh Sar amphibolites (Fig. 1) are massive to banded and mostly epidote-bear-

ing, with scattered patches of garnet epidoteamphibolites (Shah, 1986; Shah et al., 1992). These are characterized by dark-green to lightgray colour with linear fabrics defined by segregations of felsic and mafic minerals. The epidote-amphibolites are medium- to coarsegrained rocks having inequigranular porphyroblastic to subidioblastic texture. Their major mineral constituents are hornblende, epidote, and quartz. Chlorite, actinolite and ores occur as minor constituents, while rutile apatite and calcite as accessories. Small amounts of plagioclase and clinopyroxene occur rarely in these rocks. Mineralogically, the garnet-bearing epidote-amphibolites are similar to epidote-amphibolites except for the presence of garnet.



Fig. 1 Geological map of Shergarh Sar area, Allai Kohistan (after Shah, 1985). 1. Granite gneiss, 2. Limestone, 3. Schists, 4. Brecciated zone, 5. Greenschist, 6. Garnet epidote-amphibolites, 7. Epidote-amphibolites, 8. Serpentinite, 9. Peridotite, 10. Clinopyroxenite, 11. Lavas, 12. Overburden. MMT = Main Mantle Thrust; MKT = Main Karakoram Thrust. L = location of the area. Solid lines are thrust contacts, dashed where inferred.

Shah (1986) and Shah et al. (1992) described the geochemical characteristics of the Shergarh Sar amphibolites and suggested that these are derived from basic igneous rocks of tholeiitic character. These workers further suggested that plagioclase, clinopyroxene, and olivine fractionation played a major role in the evolution of igneous crystallization trend of



Fig. 2 Amphiboles from the amphibolites of Shergarh sar area on Mg/Mg+Fe vs Si diagram. Nomenclature is according to the International Mineralogical Association (Leake, 1978). A = actinolite, B = ferroactinolite, C = actinolitic-hornblende, D = ferro-actinolitic-hornblende, E = magnesiohornblende, F = ferro-hornblende, G = tchermakitic-hornblende, H = ferro-tchermakitic-hornblende, I = tchermakite, and J = ferro-tchermakite.

these amphibolites. Zahid (1991) and Hamidullah et al. (1991) presented mineralogical constraints of similar amphibolites from Gantar area, further eastward of the studied amphibolites. This paper presents the results of mineral-chemical investigations of the Shergarh Sar amphibolites; the main objective is to define the metamorphic history and its effect on the chemical and optical properties of index minerals.

MINERAL CHEMISTRY

The mineral phases in the amphibolites of the Shergar Sar area have been analyzed using the Cameca SX-50 electron microprobe at the University of South Carolina, Columbia. The operation conditions were 15 Kv accelerating potential and beam current of 25 nA. Natural and synthetic standards were used, with counting times of 20-30 seconds for each element. Three main mineral phases (i.e., amphibole, epidote and garnet) along with minor phases (i.e., plagioclase and pyroxene) were analyzed in selected samples in order to find out the variation in element abundance and change in color caused by metamorphic changes.

Amphiboles

On the basis of colour and pleochroism, the amphiboles are distinguishable into three varieties: (1) green to dark green, (2) blue-green to green, and (3) colorless to green. These have the following disposition in different rocks: entire grain made up either of (1), (2) or, rarely (3); zoned grains with cores of (1) and margins of (2), cores of (2) and margins of (3), and cores of (1), middle zone of (2) and margins of (3). At places the amphiboles are also rimed around the clinopyroxene. Textural evidence suggests that the dark green (locally olive) type has formed at peak metamorphic conditions, and the colourless to green variety during retrograde greenschist facies metamorphism with the transitional bluish green type probably during waning stages of epidote amphibolite facies metamorphism. Similar textural features are also observed further east of Shergarh Sar by Zahid (1991), Hamidullah et al. (1991) in the amphibolites of Gantar area.

Microprobe analyses of ten grains in six representative amphibolites are listed in Table 1. Total iron, determined as FeO, was partitioned into Fe₂O₃ and FeO, assuming the total number of cations (Ca+Na+K) = 13, on 23 (O) basis (Laired & Albee, 1981). All the analyses are calcic with Ca atoms in excess of 1.5 in half unit cell. According to IMA nomenclature (Leake, 1978), the dark green amphibole classifies as tchermakitic-hornblende, the bluish green as magnesio-horblende, and the colorless to green pleochroic as actinolite. The transition from tchermakitic hornblende to actinolite is marked, in addition to fall in "Al and increase in Si, by decrease in "Al, Ti, alkalies, A-site occupancy and Fe2+/ Mg ratios. These changes are suggestive of lowering temperature (Liou et al., 1974; Laird & Albee, 1981;

TABLE 1. CHEMICAL ANALYSES AND STRUCTURAL FORMULAE OF AMPHIBOLES FROM SHERGARH SAR AMPHI-BOLITES .

Sample No Grain No	Alk	46A 1	All	k40 2	Al	k33 3	All	k21 4	Alk 5	33	Al	k 8 6	Alk 9 7	92A	Alk	92A 8	All	<33)	Alk40 10
	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core
SiO ₂	44.61	44.66	44.74	44.01	42.86	42.76	43.32	43.39	45.69	47.50	43.86	48.70	48.54	47.78	49.12	54.67	53.78	53.22	54.17
TiO ₂	0.33	0.31	1.08	0.99	0.70	0.51	0.94	0.61	0.62	0.56	0.63	0.43	0.67	0.57	0.54	0.03	0.01	0.04	0.07
Al ₂ O ₃	14.97	14.83	12.21	12.88	16.30	16.18	15.21	15.39	14.21	11.50	14.47	9.22	9.81	10.02	9.02	2.31	1.54	1.32	2.40
FeO	12.37	12.84	15.15	16.09	15.63	15.64	15.89	15.12	10.64	11.43	14.51	11.94	12.62	12.15	12.69	13.26	12.73	15.64	13.79
MnO	0.06	0.12	0.17	0.13	0.23	0.32	0.20	0.33	0.20	0.26	0.13	0.06	0.06	0.08	0.04	0.25	0.30	0.30	0.25
MgO	12.13	12.03	11.00	10.17	8.63	8.80	9.15	8.77	13.22	13.50	10.69	13.40	13.08	12.75	13.75	14.98	.15.85	14.01	14.34
CaO	10.96	11.07	11.20	11.28	10.71	10.59	10.95	11.04	12.62	12.15	10.78	11.78	11.63	11.53	11.73	11.51	11.41	11.39	11.39
Na ₂ O	1.96	2.02	1.35	1.37	2.27	2.13	1.84	1.79	0.91	0.90	2.08	1.06	1.06	1.15	1.00	0.49	0.42	0.42	0.61
K ₂ O	0.02	0.04	0.28	0.25	0.27	0.22	0.24	0.22	0.07	0.06	0.07	0.00	0.01	0.02	0.01	0.03	0.02	0.00	0.00
Cl	0.00	0.00	0.02	0.00	0.06	0.04	0.07	0.01	0.04	0.01	0.01	0.02	0.03	0.16	0.02	0.00	0.00	0.00	0.00
TOTAL	97.41	97.92	97.20	97.17	97.66	97.19	97.81	96.67	98.22	97.87	97.23	96.61	97.51	96.21	97.92	97.53	96.06	96.34	97.02

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Table 1 con	ntinued																		
Cations on	23 (O)	basis																	
Si	6.37	6.37	6.52	6.45	6.28	6.27	6.32	6.41	6.48	6.76	6.37	7.04	6.96	6.96	6.99	7.81	7.76	7.77	7.63
AI	1.63	1.63	1.48	1.55	1.72	1.73	1.68	1.59	1.52	1.24	1.64	0.96	1.04	1.04	1.01	0.19	0.24	0.23	0.38
Site C																			
Al	0.90	0.86	0.62	0.68	1.10	1.07	0.94	1.09	0.86	0.69	0.84	0.61	0.61	0.68	0.51	0.20	0.02	0.00	0.11
Ti	0.04	0.03	0.12	0.11	0.08	0.06	0.10	0.07	0.07	0.06	0.07	0.05	0.07	0.06	0.06	0.00	0.00	0.00	0.00
Fe ³⁺	0.72	0.78	1.16	1.30	1.50	1.26	1.40	1.56	0.82	0.92	1.06	1.14	1.09	1.17	0.99	1.26	0.96	1.37	0.91
Fe ²⁺	0.76	0.75	0.68	0.67	0.41	0.66	0.54	0.31	0.44	0.47	0.70	0.30	0.42	0.31	0.53	0.33	0.57	0.54	0.52
Mn	0.00	0.01	0.02	0.02	0.00	0.04	0.03	0.04	0.02	0.00	0.02	0.00	0.00	0.01	0.00	0.03	0.04	0.04	0.04
Mg	2.58	2.56	2.39	2.22	1.88	1.92	1.99	1.93	2.79	2.86	2.31	2.89	2.79	2.77	2.92	3.19	3.41	3.05	3.42
Site B																			
Ca	1.68	1.69	1.75	1.77	1.68	1.66	1.71	1.75	1.92	1.85	1.68	1.83	1.79	1.80	1.79	1.76	1.76	1.78	1.78
Na	0.32	0.31	0.25	0.23	0.32	0.34	0.29	0.25	0.08	0.15	0.32	0.18	0.21	0.20	0.21	0.14	0.12	0.12	0.17
Site A																			
Na	0.22	0.25	0.13	0.16	0.33	0.19	0.23	0.26	0.17	0.10	0.26	0.12	0.08	0.12	0.07	0.00	0.00	0.00	0.00
К	0.00	0.00	0.05	0.05	0.05	0.05	0.05	0.04	0.01	0.01	0.01	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Fe/Mg	0.28	0.31	0.49	0.59	0.80	0.65	0.70	0.81	0.30	0.32	0.46	0.40	0.39	0.42	0.34	0.39	0.28	0.45	0.26
Mg/Mg+Fe	0.78	0.77	0.67	0.63	0.56	0.60	0.59	0.55	0.77	0.76	0.69	0.72	0.72	0.70	0.75	0.72	0.78	0.69	0.79
Mg#	63.61	62.55	56.41	52.98	49.60	50.07	50.65	50.83	68.89	67.80	56.77	66.67	64.88	65.16	65.89	66.82	68.94	61.49	64.96

1-4 = Unzoned green to dark green hornblende with brownish tint.

5&6 = Zoned hornblende having green core and bluish green rim.

7 = Unzoned bluish green hornblende.

8 = Zoned bluish green hornblende with marginal actinolite.

9&10 =Actinolites.

Spear, 1981; Jan & Howie, 1982) and confirm the petrographic observations.

Complex substitutions in the composition of the amphiboles were studied graphically by plotting the analyses in various ways. Figure 3 shows the position of several amphibole end members on "Al vs "Al + 2Ti + Fe3+ diagram. The amphibole analyses plot along a well-defined trend and clearly show substantial tchermakite and subordinate glaucophane substitution from actinolite to tchermakitic-hornblende. A similar picture emerges when the analyses are plotted on (Na), vs Al + 2Ti +Fe³⁺ diagram (Fig. 3). In Figure 4 the analyses are plotted in terms of A-site vacancy-Aliv-Alvi components Again a substantial tchermakitic substitution and minor edenitic substitution from actinolite to tchermakitic-horblende, resulting in a trend towards pargasite, are observed.









Epidote occurs in two habits: (a) as long subhedral to anhedral grains up to 4x2 mm in size, and (b) as small granular aggregates with bright interference colors. The type "a" epidote crystals are weakly zoned with cores displaying an anomalous bluish gray interference color and margins showing normal gray to yellow interference color. In some cases an additional highly birefringent rim of type "b" epidote surrounds the type "a" epidote. The type "a" epidote oc--curs in close association with hornblende and both minerals are commonly intergrown. In the banded epidote-amphibolites, the dark bands consist of type "a" epidote and hornblende, whereas the white bands generally contain quartz and feldspar. In such rocks, the type "a" epidote may occur as "augen porphyroblasts" surrounded by relatively finer grains of prismatic hornblende.

Chemical composition of both the varieities of epidote are given in Table 2. It is clear from this table that the change in interference colour reflects the change in chemistry (especially iron contents) in these epidotes. The bluish gray cores of the type "a" epidotes are

Sample Nos Grain Nos	Alk46A 1		Alk92A 2		Alk33 3		Alk8 4		Alk33 5	Alk8 6	Alk46A 7	A1k37 8
	Core(B)	Rim(Y)	Core(B)	Rim(B)	Core(Y)	Rim(Y)	Core(Y)	Rim(Y)	B 1	B2	B 3	B4
SiO ₂	38.52	37.64	38.20	37.56	38.22	38.03	36.71	36.69	36.48	38.25	37.59	35.17
TiO ₂	0.09	0.17	0.04	0.26	0.18	0.15	0.19	0.30	0.22	0.11	0.01	0.07
Al ₂ O ₃	29.85	27.85	29.97	27.12	28.29	28.27	26.96	26.27	25.26	27.39	25.16	25.27
Fe ₂ O ₃	5.02	7.94	4.99	8.00	7.56	7.70	8.78	8.99	10.00	8.71	10.70	14.21
MnO	0.02	0.07	0.11	0.03	0.20	0.30	0.16	0.17	0.21	0.29	0.18	0.26
CaO	23.84	23.31	23.61	23.24	22.43	22.06	23.02	22.84	23.05	23.23	23.71	22.02
TOTAL	97.34	96.98	96.92	96.21	96.88	96.51	95.82	95.26	95.22	97.98	97.35	97.00
Cation on 13	(O,OH) b	asis										
Si	3.11	3.08	3.10	3.10	3.12	3.11	3.06	3.08	3.08	3.11	3.10	2.95
^{iv} Al	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.05
^{vi} A1	2.84	2.68	2.86	2.64	2.72	2.73	2.65	2.60	2.51	2.62	2.45	2.45
Ti	0.01	0.01	0.00	0.02	0.01	0.01	0.01	0.02	0.01	0.01	0.00	0.00
Fe	0.30	0.49	0.30	0.49	0.46	0.47	0.55	0.57	0.63	0.53	0.67	0.90
Mn	0.00	0.00	0.0	0.00	0.01	0.02	0.01	0.01	0.01	0.02	0.01	0.02
Ca	2.06	2.04	2.05	2.06	1.96	1.93	2.05	2.05	2.09	2.02	2.10	1.97
PS	9.55	15.45	9.49	15.65	14.47	14.69	16.93	17.20	20.06	16.83	21.47	26.86

TABLE 2. CHEMICAL COMPOSITION AND STRUCTURAL FORMULAE OF TYPE "A" AND TYPE "B" EPIDOTES FROM SHERGARH SAR AMPHIBOLITES.

1&2 = Zoned type "a" epidote.

3&4 = Unzoned type "a" epidote.

5,6,7&8 = Type "b" epidote.

Ps = Pistacite contents.

B = Blue, Y = Yellow in crossed polars

compositionally Al-epidotes (Holdaways, 1972) with pistacite contents [Ps = 100 Fe³⁺/(Fe³⁺+Al)] of 9.5 to 14.5. These epidotes are considered to be stable at epidote-amphibolite/amphibolite facies conditions attained by the host rock. The rims are yellow in color with epidote compositions having a Ps content greater than 15. This zoning can be attributed to retrogressive metamorphism (see Miyashiro & Seki, 1958) and is observed only in the large crystals (2-4 mm in size) of type "a" epidote. The smaller type "a" epidote grains (<2mm in size) are unzoned and match well the rims of the zoned epidotes in interference colour and composition. The type "b" epidotes occur as fine-grained granular aggregates. They have greater iron contents (Ps = 17-27) and range from Al-epidotes to Fe-epidotes. Such compositions are typical of greenschist metamorphism (Holdaway, 1966; Cooper, 1972) and are considered here to imply lower equilibration temperatures than the type "a" epidote. The type "b" epidotes also have higher contents of MnO than the type "a" epidotes.

The abundant epidote in the Shergarh Sar amphibolites appears to reflect the calciumrich nature of the parent rock. Its occurrence is attributed to the release of Ca (mainly from calcic- plagioclase) during the regional metamorphism of epidote-amphibolite facies conditions, which is supported by the presence of albite in the rock. The attainment of the large size of these epidote grains may suggest the prevalence of these conditions for a considerable time or rapid reaction velocities.

Garnet

The occurrence of garnet in the epidote-amphibolites of the area is sporadic. It is generally confined to those places where leucocratic veins and bands are prominent. The garnet commonly forms porphyroblasts up to 6 cm across. It is colourless to pinkish in thin section, and usually found along the contact of hornblende and type "a" epidote. These features suggest that the garnet may have formed at the expense of hornblende and type "a" epidote. Quartz surrounding garnet crystals and occupying microveins, diverging from reaction spots, is especially common.

Five grains in different samples were analvzed from core to rim and the analyses are presented in Table 3. The structural formulae, recalculated on the basis of 24 oxygens, closely correspond to the ideal formula X2+, Y3+, Z,O,,. No distinct variation in chemical composition is present from core to margin in single grain analyses. However, different grains display noticeable variation in Fe, Mg, Mn, and Ca within a thin section. The lack of zoning in garnet in these rocks suggests that it may have escaped retrogressive metamorphism, however, variation from grain to grain within the domain of a thin section suggests local equilibration. The garnet is predominantly made up of almandine component with substantial pyrope and grossular solid-solution (Alm44.60 Pyr14.39 Gros11.26). These compositions are broadly within the range of previous analyses from amphibolite belt discussed by Jan and Howie (1987), except for the rather low grossular + andradite contents of the analyses in sample Alk 49A. The average CaO (8wt.%) is within the range of garnet from metabasites of medium to high pressure (Miyashiro, 1973).

All the analyses (core to rim) have been plotted on the pyrope-(almandine + spessertine)-(grossular + andradite) triangular diagram (Fig. 5) of Dobretsov et al. (1972). Four of the analyses plot within the overlapping field of eclogite and granulites while the rest of the analyses plot in the amphibolite garnet field. No analysis falls in the field of garnet from epidote-amphibolites.

The development of sporadic garnet in amphibolites has been attributed to variation

Sample No. Grain N	А	Alk8 1		lk8 2	А	1k9 3	Alk9	92A	Alk	49A 5
	Core	Rim	Core	Rim	Core	Rim	Core	Rim	Core	Rim
SIO ₂	38.25	38.52	38.71	38.80	39.00	38.65	38.13	38.14	39.59	39.79
TiO ₂	0.05	0.19	0.06	0.08	0.05	0.03	0.04	0.32	0.01	0.00
Al_2O_3	21.25	21.38	21.80	21.87	22.06	21.94	21.03	21.60	22.42	22.65
CrO ₃	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Fe ₂ O ₃	0.55	0.43	0.32	0.19	0.14	0.12	0.69	0.00	0.00	0.00
FeO	26.66	26.14	22.39	23.53	24.20	24.45	26.93	27.75	21.66	20.84
MnO	1.51	1.47	0.46	0.54	0.85	0.45	0.92	0.99	1.15	1.35
MgO	3.97	4.24	6.46	5.77	6.84	6.17	3.52	4.04	10.27	10.15
CaO	8.17	8.37	9.78	9.72	7.97	8.62	9.00	9.01	4.81	5.49
TOTAL	100.41	100.71	99.98	100.50	100.84	100.43	100.35	100.85	99.91	100.27
Cations on the	e 24 (O) bas	is								
Si	6.00	6.00	5.97	5.98	5.97	5.96	5.99	5.96	6.00	6.00
Al	0.00	0.00	0.03	0.02	0.03	0.04	0.01	0.04	0.00	0.00
Al	3.93	3.93	3.96	3.95	3.95	3.95	3.90	3.94	4.00	4.02
Ti	0.00	0.02	0.01	0.01	0.00	0.00	0.00	0.04	0.00	0.00
Cr	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Fe ³⁺	0.06	0.05	0.04	0.02	0.01	0.01	0.08	0.00	0.00	0.00
Fe ²⁺	3.50	3.41	2.89	3.03	3.10	3.15	3.54	3.63	2.74	2.63
Mn	0.20	0.19	0.06	0.07	0.07	0.06	0.12	0.13	0.15	0.17
Mg	0.93	0.99	1.48	1.33	1.56	1.42	0.82	0.94	2.32	2.28
Ca	1.37	1.40	1.62	1.60	1.31	1.43	1.52	1.34	0.78	0.88
AL	58.35	57.03	47.78	50.31	51.32	52.09	58.99	60.28	45.80	44.40
PY	15.47	16.51	24.57	21.99	25.85	23.45	13.74	15.62	38.73	38.21
GR	21.13	21.67	25.65	25.84	21.08	23.09	23.12	21.33	13.00	14.8
SP	3.34	3.25	0.98	1.17	1.24	0.97	2.04	2.18	2.46	2.80
AD	1.71	1.54	1.02	0.69	0.49	0.40	2.10	0.57	0.02	0.00
Mg/Fe	0.27	0.29	0.51	0.44	0.50	0.45	0.23	0.26	0.85	0.87

TABLE 3. CHEMICAL COMPOSITION AND STRUCTURAL FORMULAE OF GARNET FROM SHERGARH SAR AMPHI-BOLITES

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Fig. 5 Garnet of the garnet epidote-amphibolites from Shergarh Sar area on Pyrope-(Grossular + Andredite)-(Almandine + Spesertine) triangle showing the fields of garnet from (1) eclogites, (2) granulites, (3) amphibolites, and (4) epidote-amphibolites (after Dobretsove et al., 1972).

in bulk chemistry and/or changes in temperature and pressure (Engel & Engel, 1962; Leake, 1963; Buddington, 1966; Dewaard, 1967; Manna & Sen, 1974; Jan, 1988). There can be various causes of garnet growth in amphibolites currently studied: (1) low Mg/Fe²⁺ ratio (see also Jan, 1988), (2) low Fe₂O₃/FeO ratio and (3) local increase in temperature due to the intrusion of leucocretic veins.

Plagioclase

Plagioclase is generally sparse, having been replaced almost completely by type "a" epidote. When present, it is cloudy due to subsequent alteration. Analyses of two grains in two different samples are unzoned (Table 4). In one of these, the plagioclase is pure albite (An_1) while in the other it is more calcic (An_{27}) and is closely associated with bluish-green amphibole. It is possible that the former has grown in greenschist facies retrogression and the latter during epidote-amphibolite facies condition.

Clinopyroxene

Found only in a few rocks, the clinopyroxene varies from minor in amount up to 30% in rare

cases. It most commonly occurs as cores within large blue-green hornblende, but is also found as small interstitial grains, along with quartz, magnetite and rutile.

Analyses of four clinopyroxenes presented in Table 5 show that they are diopsidic (WO48.56 En37.40 Fs3.14). The core to rim analyses suggest that they are very homogeneous in nature. The clinopyroxene can be divided into two groups. Those of sample Alk40 and Alk92A are normal diopsides whilst those in Alk33 are strongly aluminous (Al,O, up to 7.07%) and enriched in Ca. The former have lower TiO, and higher FeO, MnO, and Na,O. Several workers (e.g. Jan, 1988) have shown that the growth of clinopyroxene in amphibolites can be controlled by bulk composition (notably high CaO) and temperature. In Shergarh Sar area, there does not appear any compositional control on its growth. It is likely that the clinopyroxene grow locally in response to high temperature.

DISCUSSION

Petrographic observation, supplemented by mineral chemistry suggest that the Shergarh sar amphibolites passed through a thermal history of amphibolite to greenschist facies metamorphism. The paragenesis representing the peak metamorphic conditions realised in the area is well-preserved and abundant in most of the rocks. This consists of green tchermakitic hornblende and Al-epidote , with or without garnet and clinopyroxene. Garnet-hornblende geothermometry (estimates for average composition in three samples) suggests temperatures of 600 to 650°C following the model of Perchuk (1967) and 685 to 715°C following Powell (1985). A clinopyroxene-hornblende pair gives a temperature of 670°C (Perchuk, 1967). Maximum temperature in excess of 650°C are consistent with the presence of local granitic bands and veins which may be a partial

Sample No	А	lk21	Alk92A				
	Core	Rim	Core	Rim			
SiO ₂	67.39	65.82	63.17	61.84			
A ₂ O ₃	21.12	21.36	25.13	24.72			
MgO	0.01	0.04	0.00	0.00			
CaO	0.30	0.68	5.18	5.20			
MnO	0.00	0.02	0.01	0.00			
FeO	0.29	0.92	0.03	0.02			
BaO	0.00	0.00	0.02	0.00			
Na ₂ O	9.76	10.20	7.52	7.79			
K ₂ O	0.05	0.02	0.03	0.03			
TOTAL	98.92	99.06	101.09	99.60			
Cation on 8 (O)) basis						
Si	2.96	2.91	2.75	2.74			
Al	1.09	1.11	1.29	1.29			
Fe	0.01	0.03	0.00	0.00			
Mg	0.00	0.00	0.00	0.00			
Mn	0.00	0.00	0.00	0.00			
Ca	0.01	0.03	0.24	0.25			
Ba	0.00	0.00	0.00	0.00			
Na	0.83	0.87	0.64	0.67			
K	0.00	0.00	0.00	0.00			
Or	0.00	0.00	0.00	0.00			
Ab	99.00	97.00	73.00	73.00			
An	1.00	3.00	27.00	27.00			

TABLE 4. CHEMICAL ANALYSES AND STRUCTURAL FORMULAE OF FELDSPAR FROM SHERGARH SAR AMPHIBOLITES

S. No	A	k40	Alk	92A	Alk	33	Alk33		
<u> </u>	Core	Rim	Core	Rim	Core	Rim	Core	Rim	
SiO,	50.75	49.69	50.46	50.17	47.68	47.56	47.08	47.33	
TiO,	0.25	0.35	0.23	0.31	0.50	0.51	0.59	0.60	
Al ₂ O ₃	3.58	3.75	3.41	3.63	6.85	6.66	7.07	6.87	
Cr ₂ O ₃	0.06	0.04	0.04	0.00	0.02	0.01	0.00	0.03	
Fe ₂ O ₃	2.90	3.24	2.62	2.91	3.66	4.18	4.45	3.90	
FeO	7.22	7.06	7.70	7.41	2.53	2.11	2.08	2.57	
MnO	0.24	0.29	0.32	0.30	0.08	0.06	0.08	0.08	
MgO	12.22	12.07	12.26	12.07	12.86	12.91	12.70	12.76	
CaO	22.45	21.74	21.55	21.90	24.73	24.90	24.81	24.55	
Na ₂ O	0.62	· 0.65	0.66	0.64	0.05	0.05	0.06	0.06	
TOTAL	L100.29	98.88	99.25	99.34	98.96	98.95	98.92	98.74	
Cations	s on 6 (O)	basis				•			
Si	1.90	1.88	1.90	1.90	1.78	1.78	1.77	1.78	
Al	0.10	0.12	0.10	0.10	0.22	0.22	0.23	0.22	
Al	0.06	0.05	0.05	0.06	0.08	0.07	0.08	0.08	
Ti	0.00	0.01	0.00	0.01	0.01	0.01	0.02	0.02	
Fe	0.08	0.09	0.07	0.08	0.10	0.11	0.13	0.11	
Fe	0.22	0.22	0.24	0.23	0.08	0.06	0.06	0.08	
Mn	0.01	0.01	0.01	0.01	0.00	0.00	0.00	0.00	
Mg	0.68	0.68	0.69	0.67	0.72	0.72	0.71	0.71	
Ca	0.89	0.88	0.87	0.88	0.99	0.99	0.99	0.99	
Na	0.04	0.05	0.05	0.05	0.00	0.00	0.00	0.00	
WO	47.34	46.81	46.28	47.06	52.38	2.66	52.38	52.38	
EN	36.17	36.17	36.70	35.83	38.10	38.30	37.57	37.57	
FS	16.49	17.02	17.02	17.11	9.52	9.04	10.05	10.05	

TABLE 5. CHEMICAL ANALYSES AND STRUCTURAL FORMULAE OF PYROXENE FROM SHERGARH SAR AMPHIBOLITES.

melting product of the amphibolite (cf. Whyllie, 1977). Such temperature have already been recorded from the amphibolite belt in Swat and adjacent Indus valley (Bard, 1983; Jan, 1988). However, a lower temperature range (500-530°C) is reported for the similar rocks at Gantar area of Allai Kohistan (see Hamidullah et al., 1991; Zahid, 1991).

The transition from amphibolite to greenschist facies metamorphic conditions in the rocks is documented by the overgrowths of bluish-green magnesio-hornblende on the tchermakitic hornblende and more ferric epidote around the Al-epidote. These appear to have grown at intermediate temperatures. The composition of a magnesio-hornblende and an adjacent calcic-oligoclase yield a temperature estimate of about 520 °C both by the model of Spear (1980) and Plyusnina (1982). Overprinted on these paragenesis in most rocks is typical greenschist facies assemblage consisting of actinolite, Fe-epidote, albite and local chlorite. The former two of these minerals occur in independent grains and as outer most zones around homblende and Fe-poorer epidote, respectively.

Quantitative estimates of operating pressures during the metamorphic evolution of the rocks have not been obtained, however, there are indications that these may have been high. Laird and Albee (1981, Fig. 8) showed that amphiboles from low-, medium- and high-pressure metamorphic terrains plot in distinct fields on "Al vs "Al + Ti + Fe3 + diagram. The hornblendic amphiboles from the area fall in overlapping fields of medium and high-pressure fields. The plagioclase-hornblende thermobarometer of Plyusnina (1982) suggests a pressure of 5 kb for the one analysed pair referred to above. The "Al vs "Al plots of the hornblendic amphiboles (Fig. 6) suggest pressure in excess of 5 kb. The "garnet-epidote" thermobarometers of Dobretsov et al. (1972) gives rough esti-



Fig. 6 Relationship of "Al to "Al in the analysed amphiboles. D limit is after Doolan et al. (1978) and L limit is after Leake, (1971). Fields of igneous Ca-amphiboles and the boundary of high and low pressure Ca-amphiboles are after Fleet and Barnett (1978).

mates of 690°C and 6.2 kb for epidote with Ps contents of 15. Pressure in excess of 5 kb have already been suggested for peak metamorphism in the amphibolite belt (Bard, 1982; Jan, 1988, Treloar et al., 1989).

Radiometric data suggest that metamorphism in southern Kohistan coincides with collision and overthrusting of Karakoram plate onto the Kohistan magmatic arc. The entire amphibolite belt is a major shear zone (Treloar et al., 1989, 1990). The amphibolite facies metamorphism, thus, occurred in response to thrust stacking and crustal thickning. The initial suturing (collision) of the Kohistan magmatic arc with the Indian plate is considered to have taken place during the Eoceine (Patriate & Achache, 1984; Treloar et al., 1989). The uplift and cooling of the amphibolite belt occurred, probably, during ensuring obduction of the Kohistan arc and its emplacement onto the Indian plate. During this episode of upliftcooling, overprinting of lower grade assemblages took place along grain boundaries. The process was enhanced along shear zones due to granulation and availability of fluids, mainly H₂O. The Shergarh Sar amphibolites nicely document these events. It would be fruitful to perform additional work on the mineral chemistry and radiometric ages to construct a P-T path for the rocks, beginning with their basaltic and gabbroic protoliths. Jan (1988) and Hamidullah et al. (1990, 1991) have reported several gabbroic relics in the amphibolites of Indus, Swat and Allai Kohistan valleys, and a careful search may also reveal such relics in the area under discussion.

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