# Hydrothermal alteration studies of gabbros from Northern Central Indian Ridge and their geodynamic implications

DWIJESH RAY<sup>1,\*</sup>, CATHERINE MEVEL<sup>2</sup> and RANADIP BANERJEE<sup>3</sup>

<sup>1</sup>National Centre for Antarctic & Ocean Research, Goa 403 804, India.
<sup>2</sup>Geosciences Marines, CNRS, IPGP, 4 Place Jussieu, F-75252 Paris Cedex 5, France.
<sup>3</sup>National Institute of Oceanography, Goa 403 004, India.
\*e-mail: dwijesh@rediffmail.com

Mylonitic gabbro and altered gabbro were recovered from off-axis high and corner high locations at ridge-transform intersection, adjacent to Vityaz transform fault of the slow spreading (32–35 mm/yr, full spreading) Northern Central Indian Ridge. Both the varieties show signatures of extensive alteration caused due to interaction with sea water. Mylonitic gabbro represents high temperature metamorphism ( $\sim$ 700–800°C) and comprised of hornblende mineral which exhibits well defined foliation/gneissic appearance along with dynamically recrystallised plagioclase grains frequently intercalated with magnetite-ilmenite. Altered gabbro from corner high generally includes low temperature greenschist grade ( $\sim$ 300°C) mineralogical assemblages: chlorite, albite, quartz and locally magnesio hornblende. Crystal plastic deformation resulted in mylonite formation and often porphyroclasts of plagioclase and clinopyroxene grains, while altered gabbro locally exhibits cataclastic texture. Presence of Vityaz transform fault and adjacent meganullion at the weakly magmatic ridge-transform intersection and off-axis high locations prompted the present scenario very much conducive for hydrothermal circulation and further facilitate the exhumation of present suite of gabbro.

#### 1. Introduction

Detailed petrologic studies of oceanic gabbros have demonstrated that metamorphic recrystallisation is related either to deformation and/or prolonged circulation of fluid during the progressive cooling of the gabbroic sequence (Honnorez et al 1984; Mevel 1987, 1988; Stakes et al 1991; Alt and Bach 2001). Magmatism in association with faulting plays a crucial role to understand deformation processes, magma migration paths and chemical differentiation processes, while hydrothermal circulation caused the post-crystallisation mineralogical changes within gabbro. Detailed studies of petrography and mineral chemistry of mylonitic gabbro and altered gabbro from Northern Central Indian Ridge (NCIR) have been undertaken to trace the hydrothermal alteration process of lower oceanic

crust at slow spreading ridges. Hydrothermally altered plutonic rock samples also provide information about both the primary igneous intrusions that supplied heat to drive hydrothermal circulation and the geochemical evolution of hydrothermal fluids during rock-fluid interaction. Thus, the alteration in layer 3 crustal section always provides rare opportunity to our understanding on hydrothermal processes at spreading ridge system.

Earlier studies on mineralogy and geochemistry of altered gabbro are mainly restricted with slow spreading MAR (ODP Leg 153, Agar and Lloyd 1997), fast spreading Hess Deep, EPR (Mevel and Stamoudi 1996) or ultraslow spreading SWIR (Hole 735B, Stakes *et al* 1991; Vanko and Stakes 1991; Alt and Bach 2001). In contrast, very less information on hydrothermal alteration of gabbro

Keywords. Mylonitic gabbro; altered gabbro; hydrothermal alteration; Northern Central Indian Ridge.

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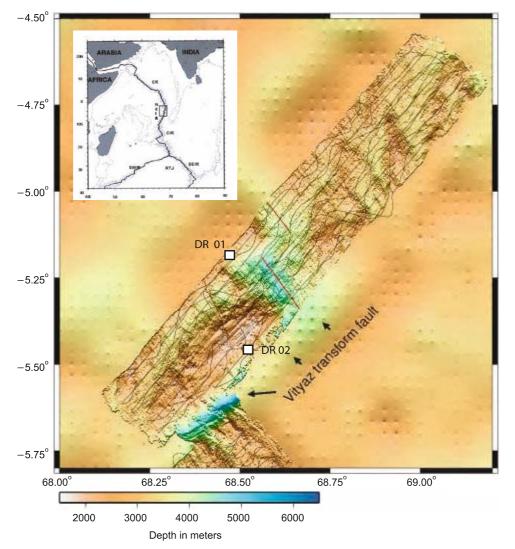


Figure 1. Map showing Vityaz megamullion and present gabbro locations (stations DR 01 and DR 02 respectively) along the Northern Central Indian Ridge (after Drolia and DeMets 2005). Inset map showing Indian Ocean Ridge System.

is available from NCIR – one of the least understood mid-ocean ridge system. Earlier reports on occurrences of gabbro from CIR were reported by Engel and Fisher in 1975, though the alteration characteristics of CIR gabbro were not discussed elsewhere. In order to characterize the extent of hydrothermal alteration in gabbros, we present the first report of petrography and mineral chemistry of the hydrothermally altered gabbros collected from NCIR (figure 1). The basic objective of this paper is to address the sequence and extent of alteration using texture, mineralogy and chemical compositional variations and also understand their geodynamic implications.

# 2. Methodology

Samples studied here were dredged using chain bag dredge from off-axis high and inside corner high (both the locations are adjacent to Vityaz 'megamullion') of Vityaz transform fault (TF) of slow-spreading NCIR (figure 1, Drolia and DeMets 2005). The full spreading rate of the present study area is 32-35 mm/yr (DeMets *et al* 2005). The first location (DR 01  $\sim 5^{\circ}11.02'$ S/68°28.10'E, figure 1) was dredged from an off-axis high, northern side of active 'Vitvaz megamullion' as described by Drolia and DeMets in 2005. Gabbros from this location include mainly mylonite gabbros. The second location (DR 02  $\sim 5^{\circ}26.92'$ S/ $68^{\circ}31.76'$ E, figure 1) is on the southern side of 'megamullion' represents an inside corner high of Vityaz transform fault, vice versa flank of 'Kurchatov seamount' as described by Russian researchers during their expedition (Baturin and Rozanova 1975). Rocks dredged from this site are mainly altered gabbro underwent greenschist facies alteration.

Electron probe micro-analyses of primary as well as the secondary minerals were conducted using

Station, position and depth	Type of gabbro	Tectonic setting	Petrography	Mineralogy (primary+secondary)	Grade of alteration
DR-01 5°11.02'S/ 68°28.10'E 3000 m	Mylonitic gabbro	Off-axis high	Mostly exhibits banded feature by recrystallised plagio- clase, amphibole and Fe-Ti oxide, plagio- clase and clinopyrox- ene as porphyroclasts	$\begin{array}{l} \mbox{Plagioclase (anorthite+}\\ \mbox{labradorite}) \sim 40\%\\ \mbox{Clinopyroxene} \sim 10\%\\ \mbox{Hornblende} \sim 20\%\\ \mbox{Fe-Ti oxide} \sim 15\%\\ \mbox{Chlorite} \sim 5\%\\ \mbox{Epidote} \sim 5\% \end{array}$	Mostly amphibolite facies, locally greenschist facies
DR-02 5°26.92′S/ 68°31.76′E 2260 m	Altered gabbro, brecciated	Corner high of Vityaz TF	Mostly comprised of angular, brecciated plagioclase. Albite, chlorite and quartz closely associated, cataclasic texture	$\begin{array}{l} \mbox{Plagioclase (labradorite+}\\ \mbox{albite})\sim 60\%\\ \mbox{Chlorite}\sim 10\%\\ \mbox{Amphibole}\sim 10{-}15\%\\ \mbox{Quartz}\sim 10{-}15\% \end{array}$	Mainly green- schist facies, locally amphi- bolite facies

Table 1. Location, tectonic setting and brief petrographic descriptions of Northern Central Indian Ridge gabbros.

both the Cameca SX 50 and SX 100 WDS electron microprobe analyzer at the CAMPARIS service of the IPG-Paris (University of Paris 6, France). Analytical conditions used were 15 kV accelerating voltage, 20 nA beam current and 20–40 s counting times. All analyses were performed in a point mode. A 2–3  $\mu$ m beam size was used for all minerals (plagioclase, amphibole, pyroxene, epidote). The obtained data were reduced using a ZAF and PAP correction procedure. Natural mineral standards were employed to check the precision and accuracy of the instrument (always better than 5%).

## 3. Results

## 3.1 Petrography

Mylonite gabbro (mineralogical assemblages up to amphibolite facies) and altered gabbro (mineralogical assemblages up to greenschist facies), terminology were used on the basis of their texture as well as mineralogy and the petrographic observations (carefully chosen 20 samples on each location). Based on petrographic observations and presence and nature of secondary minerals, the samples are classified into two types: lowtemperature altered gabbro and high-temperature mylonite gabbro. The details of the microscopic characteristics of each group are described hereunder and also provided in table 1.

# 3.1.1 Mylonite gabbro from off-axis high, north of Vityaz TF

Samples from off-axis high (DR 01), north of Vityaz TF include principally mylonitic gabbro often gneissic in appearance. They generally display mylonitic texture often porphyroclast locally (figure 2a-d). Fe-Ti oxides (mainly magnetiteilmenite) often wraps against the amphibole veins (figure 2a) or poikilitically host the small clinopyroxene as well as plagioclase grains (figure 2b, d). Otherwise discrete veins of magnetite-ilmenite cut the dynamically recrystallised plagioclase. Plagioclase and clinopyroxene both occur as porphyroclasts even sometimes twins in plagioclase bend with tapering ends and curvy due to deformation (figure 2c) or sometimes plagioclase exhibit undulatory extinctions (figure 2d). Evidences of local shearing are ascertained due to distinct oxide-rich band that separate from dynamically recrystallised band of plagioclase and clinopyroxene. We note that clinopyroxene porphyroclasts also present with typical 'tail'-like feature. Porphyroclasts sometimes display a shape-preferred elongation parallel to the Fe-Ti oxide stringers. Fe-Ti oxides also show typical 'pressure shadow' feature around plagioclase porphyroclast (figure 2c). Amphibole is the principal high-temperature hydrous phase most common in MOR gabbros resulted due to crystallization of evolved hydrous silicate melt or as a product of high temperature water-rock reactions. Both the brown and green variety of amphibole (brown and green hornblende) are present within the mylonite or gneissic gabbro (figure 2a). Locally, they occur isolated or as a cluster or even as band alternate with recrystallised plagioclase, sometimes took the shape of lenses with tapering end (figure 2b). Alteration of clinopyroxene to amphiboles are noticed along the clinopyroxene grain margin or along the fractures developed within the grain (figure 2e). Amphibole shows variegated textures, e.g., locally clusters as subhedral grains, replacement of clinopyroxene and vein amphibole. Chlorite and epidotes are also present locally. Occurrences of epidote are volumetrically

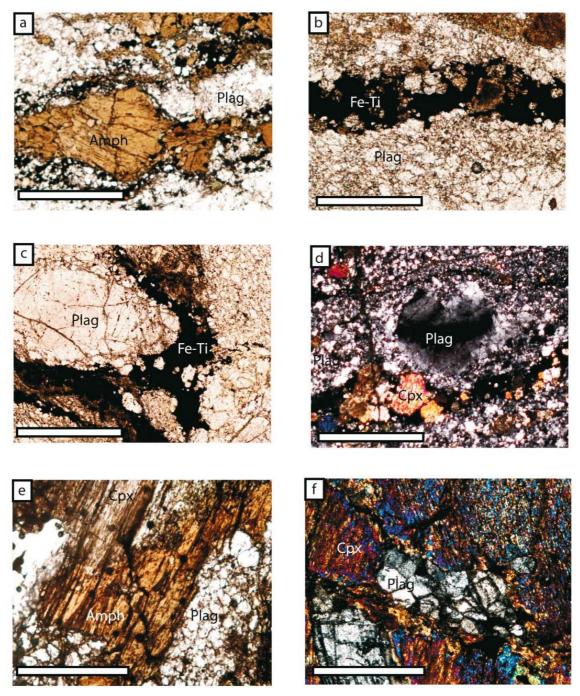


Figure 2. Photomicrographs of mylonitic gabbro from off-axis high (DR 01). (a) Brown amphibole (Amph) intercalated with Fe-Ti oxides (black) form alternating layers with recrystallised plagioclase grains (Plag, white) (PPL  $\times 5X$ ). (b) Fe-Ti oxide veins (black) to host clinopyroxene (Cpx) and amphibole with the recrystallised plagioclase matrix (PPL $\times 5X$ ). (c) Deformed twins within plagioclase porphyroclast (shape preferred orientation subparallel to the foliation). Matrix composed of fine grained plagioclase (PPL $\times 5X$ ). (d) Plagioclase showing undulatory extinction. A Fe-Ti vein in the lower half (black) also hosting Cpx grains (XPL $\times 5X$ ). (e) Alteration of Cpx to amphibole along the margin. Recrystallised plagioclase grains also seen in the matrix (PPL $\times 10X$ ). (f) Weakly recrystallised plagioclase grains showing triple junction (XPL $\times 10X$ ). Scale bar: 1 mm for figure 2(a–d) and 0.5 mm for figure 2(e–f).

not significant and occur mostly as a vein material or as a replacement of plagioclase in mylonitic gabbro. Plagioclase alteration noticed along fractures and chlorite took its place as fracture filling secondary minerals. Fe-Ti oxides (magnetite and ilmenite) mostly occur as thick veins or as thin stringers multibranching pattern with dynamically recrystallised plagioclase grains. Sometimes they also occur as big patches or as disseminated grains. Ilmenite

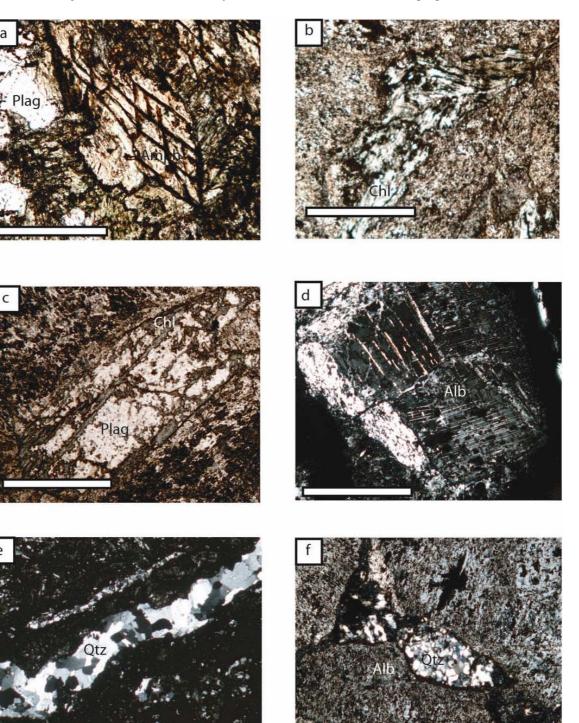


Figure 3. Photomicrographs of altered gabbro from corner-high (station DR 02). (a) Magnesio hornblende (Amph) with prominent cleavage trace (PPL  $\times$  10X). (b) Deformed chlorite (Chl) band within albitised matrix (PPL  $\times$  5X). (c) Chloritisation of plagioclase along fractures formed within the plagioclase phenocryst (PPL $\times$ 5X). (d) Twinning within albite (Alb). (XPL $\times$ 10X). (e) Late quartz vein (Qtz). (XPL $\times$ 5X). (f) Recrystallised Qtz grains enclosed by albite (Alb) phenocrysts (XPL $\times$ 5X). Scale bar: 0.5 mm for figure 2(a, d) and 1 mm for figure 2(b, c, e, f).

occasionally also preserved exsolution lamellae within the magnetite. Pyrite is also present, even though very rare. Recrystallisation in the weakly recrystallised texture is indicated by the distinct polygonal grains and enclosing grain boundaries with 120° triple junctions (figure 2f). They closely associated with brown amphibole band and the porphyroclasts of plagioclase and clinopyroxene.

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Table 2. Representative analyses of plagioclase from mylonite gabbro. Total iron calculated as  $FeO_t$ .

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Sample no.	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-2	DR-01-2
Oxides (wt%)								
$\mathrm{SiO}_2$	49.71	49.18	50.19	50.07	49.44	49.55	51.54	51.39
$\mathrm{TiO}_2$	0.06	0.01	0.07	0.01	0.02	0.00	0.08	0.04
$Al_2O_3$	32.04	32.13	31.60	31.81	32.20	31.86	30.04	30.23
$\mathrm{Cr}_2\mathrm{O}_3$	0.01	0.02	0.07	0.03	0.00	0.04	0.00	0.00
$\rm FeO_t$	0.11	0.16	0.19	0.12	0.19	0.16	0.42	0.49
MnO	0.00	0.07	0.01	0.00	0.05	0.01	0.01	0.00
MgO	0.03	0.00	0.00	0.03	0.00	0.01	0.04	0.05
CaO	14.82	14.94	14.62	14.59	14.57	14.26	12.85	12.59
$Na_2O$	2.89	2.82	3.23	3.13	3.07	3.08	4.05	4.25
$K_2O$	0.02	0.01	0.06	0.02	0.02	0.04	0.08	0.02
NiO	0.00	0.01	0.00	0.00	0.05	0.03	0.02	0.00
Cl	0.02	0.00	0.00	0.00	0.00	0.01	0.01	0.02
$P_2O_5$	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	0.05	0.00
Total	99.70	99.35	100.04	99.82	99.61	99.05	99.18	99.07
Cations based	on 32 oxygens							
Si	9.098	9.043	9.165	9.148	9.063	9.121	9.464	9.438
Al	6.911	6.962	6.801	6.849	6.957	6.911	6.500	6.543
Fe(II)	0.017	0.025	0.029	0.018	0.029	0.025	0.064	0.075
Ca	2.906	2.943	2.860	2.856	2.861	2.812	2.528	2.477
Na	1.025	1.005	1.144	1.109	1.091	1.099	1.442	1.513
Κ	0.005	0.002	0.014	0.005	0.005	0.009	0.019	0.005
Catsum	19.962	19.980	20.013	19.984	20.006	19.978	20.017	20.050
An	73.83	74.50	71.19	71.95	72.30	71.73	63.38	62.01
Ab	26.05	25.45	28.46	27.93	27.60	28.03	36.15	37.88

# 3.1.2 Altered gabbro from corner high, Vityaz TF

Samples collected from inside corner high of Vityaz TF (DR 02) are mainly broken and altered gabbro (figure 3a–f) often similar to cataclastics. The rocks look green coloured and frequently fractured. We note that extensive veins and vein networks of chlorite  $\pm$  epidote  $\pm$  quartz, fractured plagioclase and pyroxene grains set in a matrix. Texture is mostly brecciated. Signature of any plastic deformation or mylonitisation is totally absent within altered gabbro even though deformation features are well exemplified in mineral fabric. Plagioclase is the dominant phenocryst phase and are angular, fractured, twined. Plagioclase spectacularly shows microcline and albite twin (figure 3d). Relict plagioclase grains commonly exhibit weak undulose extinction and contain tapering twins often fractured also. Chlorite occurs along the fractures of the plagioclase or as deformed wavy bands or sometimes develop a schistosity (figure 3b, c). Recrystallised quartz vein occurs as secondary vein filling (figure 3e, f) or as mosaic aggregates. Quartz is common, although not voluminous;

occur as recrystallised vein closely associated with albite. Typically quartz is colourless to white and devoid of any inclusion. Hornblende mostly occurs as individual or as locally cluster at one place (figure 3a).

#### 3.2 Mineralogy and mineral chemistry

## 3.2.1 Plagioclase

Representative plagioclase compositions are given in tables 2 and 3 respectively. Anorthite content (An) of the plagioclase from mylonitic gabbro varies from relatively high An (An<sub>71-75</sub>) to low An (An<sub>62-66</sub>).

In contrast, gabbros from corner high include labradorite and albite both. Albite composition from altered gabbro varies from  $Ab_{91}$  to  $Ab_{96}$ . Labradorite plagioclase mostly display the intermediate Ab composition (Ab<sub>53</sub> to Ab<sub>62</sub>, figure 4). The intermediate member of plagioclase is missing. Secondary plagioclase is identified by its sodic nature and has a slightly higher K<sub>2</sub>O content than the associated magmatic plagioclase. However, K<sub>2</sub>O is always below 0.5 wt% (mostly 0.03

Table 3. Representative analyses of plagioclase from altered gabbro.

Sample no.	DR-02-1	DR-02-1	DR-02-1	DR-02-1	DR-02-1	DR-02-1	DR-02-1	DR-02-1	DR-02-1
Oxides (wt%)									
$SiO_2$	66.87	68.18	68.27	65.89	67.62	68.74	56.31	55.55	54.74
$\mathrm{TiO}_2$	0.03	0.04	0.00	0.05	0.00	0.02	0.03	0.06	0.06
$Al_2O_3$	21.01	20.15	19.96	19.56	20.64	20.04	28.11	27.81	28.09
$\mathrm{Cr}_2\mathrm{O}_3$	0.02	0.06	0.01	0.00	0.00	0.00	0.08	0.00	0.00
$\rm FeO_t$	0.06	0.05	0.13	0.49	0.03	0.05	0.39	0.34	0.17
MnO	0.04	0.00	0.00	0.06	0.00	0.03	0.00	0.08	0.06
MgO	0.02	0.04	0.01	0.91	0.00	0.01	0.03	0.00	0.05
CaO	1.66	0.85	0.69	0.75	1.39	0.82	9.54	9.86	10.33
Na <sub>2</sub> O	9.89	10.80	10.30	10.12	9.88	10.63	5.30	5.38	5.13
$K_2O$	0.07	0.04	0.05	0.07	0.05	0.01	0.05	0.03	0.01
NiO	0.03	0.00	0.04	0.00	0.09	0.00	0.00	0.05	0.05
Cl	0.01	0.04	0.01	0.03	0.02	0.00	0.01	0.01	0.00
Total	99.71	100.23	99.47	97.93	99.72	100.34	99.84	99.18	98.70
Cations based	on 32 oxyger	ns							
Si	11.735	11.894	11.958	11.884	11.834	11.947	10.139	10.080	9.991
Al	4.345	4.143	4.120	4.158	4.257	4.105	5.965	5.947	6.042
Fe(II)	0.009	0.007	0.019	0.074	0.004	0.004	0.005	0.052	0.026
Ca	0.312	0.159	0.129	0.145	0.261	0.153	1.840	1.917	2.020
Na	3.365	3.653	3.498	3.539	3.352	3.582	1.850	1.893	1.815
Κ	0.016	0.009	0.011	0.016	0.011	0.002	0.011	0.007	0.002
Catsum	19.782	19.865	19.476	19.815	19.719	19.793	19.810	19.896	19.897
An	8.45	4.16	3.56	3.92	7.19	4.09	49.71	50.23	52.64
Ab	91.12	95.61	96.13	95.65	92.50	95.85	49.98	49.59	47.30
Or	0.42	0.23	0.31	0.44	0.31	0.06	0.31	0.18	0.06

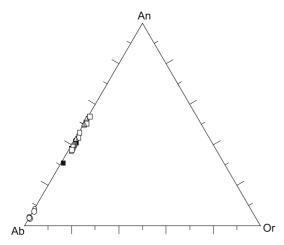


Figure 4. An–Ab–Or ternary plot of plagioclases from mylonite gabbro (DR 01) and altered gabbro (DR 02). Open triangles and open circles represent plagioclase and albite from altered gabbro. Filled and open squares represent plagioclases from mylonite gabbro. An: Anorthite, Ab: Albite, Or: Orthoclase.

to 0.07 wt%), similar to secondary plagioclases reported from other oceanic altered gabbros (Mevel 1984; Marion *et al* 1991). Secondary plagioclases range from albite in some samples to oligoclase and andesine in others. It has also been noticed that secondary plagioclase in altered gabbros dredged from the Mid-Cayman Rise, altered under upper greenschist to lower amphibolite facies conditions (Ito and Clayton 1983). Thus, it is highly unlikely that the albite-rich alteration assemblage in the altered gabbro could have formed at temperatures much greater than 300°C, which is similar to that have been described earlier from Hole 504B, Mid-Atlantic Ridge. Also the associated mineral assemblage, presence of chlorite and epidote suggest the low temperature paragenesis for albite.

# 3.2.2 Clinopyroxene

Compositions of clinopyroxene from both in the altered gabbro and mylonitic gabbro are given in table 4. Present clinopyroxene composition varies from  $Wo_{42}En_{38}Fs_6$  to  $Wo_{48}En_{49}Fs_{14}$  and plotted in Wo:En:Fs ternary diagram (figure 5). The clinopyroxene from mylonite zone are more diopsidic compared to clinopyroxene away from the mylonite zone and they plot away from the diopside-end in the same diagram (figure 5). Al<sub>2</sub>O<sub>3</sub> content of clinopyroxene porphyroclast is comparatively low (as low as 0.16 wt%, table 4). Similar relationships between magmatic and secondary

Sample no.	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-2	DR-01-2
Oxides (wt%)							
$SiO_2$	53.65	53.15	52.39	53.92	54.32	52.37	52.43
$\mathrm{TiO}_2$	0.22	0.22	0.16	0.19	0.01	0.40	0.42
$Al_2O_3$	1.98	1.16	1.64	1.43	0.16	2.80	2.51
$\mathrm{Cr}_2\mathrm{O}_3$	0.23	0.15	0.14	0.23	0.06	0.18	0.18
$\rm FeO_t$	5.25	8.10	4.03	4.24	5.98	6.06	6.11
MnO	0.36	0.19	0.12	0.13	0.27	0.14	0.22
MgO	16.75	15.32	17.07	16.76	15.23	16.16	16.27
CaO	19.89	21.90	23.10	23.25	24.44	21.60	21.91
Na <sub>2</sub> O	0.47	0.28	0.35	0.16	0.12	0.30	0.33
$K_2O$	0.04	0.02	0.03	0.00	0.01	0.00	0.00
NiO	0.07	0.01	0.00	0.00	0.01	0.00	0.03
Cl	0.65	0.02	0.06	0.05	0.03	0.00	0.00
$P_2O_5$							
Total	99.56	100.51	99.09	100.37	100.63	100.00	100.42
Cations based of	n 6 oxygens						
Si	1.976	1.964	1.939	1.965	1.996	1.925	1.924
Ti	0.006	0.006	0.004	0.005	0	0.011	0.012
Al	0.086	0.05	0.072	0.062	0.007	0.121	0.109
$\mathbf{Cr}$	0.007	0.004	0.004	0.007	0.002	0.005	0.005
Fe	0.162	0.25	0.125	0.129	0.184	0.186	0.188
Mn	0.011	0.006	0.004	0.004	0.008	0.004	0.007
Mg	0.92	0.844	0.942	0.91	0.834	0.885	0.89
Ca	0.785	0.867	0.916	0.908	0.963	0.851	0.862
Na	0.034	0.02	0.025	0.012	0.009	0.021	0.024
Κ	0.002	0.001	0.001	0	0.001	0	0
Catsum	3.989	4.012	4.032	4.002	4.004	4.009	4.021
Wo	42.05	44.21	46.19	46.64	48.61	44.28	44.43
En	49.28	43.04	47.50	46.74	42.10	46.05	45.88
Fs	8.68	12.75	6.30	6.63	9.29	9.68	9.69
Mg#	85.03	77.15	88.28	87.58	81.93	82.63	82.56

Table 4(a). Representative analyses of magmatic clinopyroxene.

Mg# mole  $Mg^{2+}/mole (Mg^{2+} + Fe^{2+})$ .

pyroxene have been documented in oceanic gabbros elsewhere (Mevel 1987; Vanko and Stakes 1991; Gillis *et al* 1993; Tartarotti *et al* 1998). A few of the present clinopyroxene porphyroclast are also enriched in MnO (up to 0.64 wt%). Clinopyroxene is comparatively rich in iron in the mylonite gabbro (up to 10.5 wt%). A binary diagram has been incorporated to display the variation of FeO<sub>t</sub> and MnO (figure 6). Clinopyroxene in mylonitic gabbro (mostly porphyroclasts) are consistently high in FeO<sub>t</sub> and MnO content as compared to clinopyroxene away from the mylonite zone.

The composition of the clinopyroxene porphyroclast suggests their formation at a restricted temperature ( $600^{\circ}-900^{\circ}$ C) which is comparatively less than the formation temperature of clinopyroxene away from mylonite zone (figure 5, Bird *et al* 1986). Clinopyroxenes exhibiting different textural pattern fall in the different range of temperatures have also been well reported from MARK area (Gillis *et al* 1993) as well as in the gabbros from Hole 735B in the Indian Ocean (Stakes *et al* 1991). Orthopyroxene is not observed in the present samples. Olivine is rare constituent for either altered or mylonitic gabbros.

#### 3.2.3 Amphibole

Amphiboles from corner-high altered gabbro are magnesio hornblende in the terminology of Leake, 1978 replaced clinopyroxene, while the vein hornblendes (brown hornblende) of mylonitic gabbro from megamullion are mostly edinitic hornblende. In amphibole binary diagram ( $Al^{IV}$  vs. (Na + K)<sub>A</sub> diagram), magnesio hornblende mostly fall under actinolitic hornblende field, are generally Ti poor (<1 wt%) as compared to edinitic hornblende of mylonitic gabbro (Ti-rich, maximum TiO<sub>2</sub> up to 3 wt%, figure 7, table 5). However, the

Table 4(b). Representative analyses of vein clinopyroxene.

Sample no.	DR-01-3	DR-01-3	DR-01-3	DR-01-3	DR-01-3	DR-01-3	DR-01-3	DR-01-3	DR-01-3
Oxides (wt%)									
$SiO_2$	51.20	51.43	50.87	51.94	52.64	52.48	51.78	51.40	51.53
$\mathrm{TiO}_2$	0.46	0.55	0.36	0.39	0.28	0.19	0.32	0.39	0.45
$Al_2O_3$	1.13	1.27	1.15	1.12	1.16	0.99	1.18	1.19	1.14
$\mathrm{Cr}_2\mathrm{O}_3$	0.00	0.05	0.00	0.04	0.03	0.00	0.03	0.00	0.00
$\rm FeO_t$	14.01	13.70	12.44	11.79	12.06	12.40	12.92	14.04	16.49
MnO	0.64	0.55	0.39	0.50	0.41	0.35	0.42	0.55	0.55
MgO	12.17	11.80	11.92	12.42	12.25	12.94	12.41	12.11	12.80
CaO	19.11	19.22	19.93	20.16	20.43	19.98	19.85	20.24	16.81
$Na_2O$	0.35	0.45	0.36	0.36	0.31	0.34	0.32	0.35	0.28
$K_2O$	0.00	0.01	0.00	0.02	0.00	0.01	0.04	0.03	0.02
NiO	0.03	0.11	0.05	0.00	0.00	0.00	0.00	0.07	0.15
Cl	0.00	0.00	0.00	0.00	0.03	0.01	0.00	0.00	0.00
$P_2O_5$	0.00	0.00	0.04	0.00	0.02	0.03	0.04	0.02	0.04
Total	99.09	99.14	97.50	98.73	99.61	99.72	99.29	100.39	100.26
Cations based	on 6 oxygens	3							
Si	1.965	1.971	1.976	1.982	1.991	1.984	1.974	1.955	1.964
Al	0.051	0.057	0.052	0.05	0.052	0.044	0.053	0.053	0.051
Ti	0.013	0.016	0.01	0.011	0.008	0.005	0.009	0.011	0.013
$\mathbf{Cr}$	0	0.002	0	0.001	0.001	0	0.001	0	0
Fe	0.45	0.439	0.404	0.376	0.381	0.392	0.412	0.447	0.526
Mn	0.021	0.018	0.013	0.016	0.013	0.011	0.013	0.018	0.018
Mg	0.696	0.674	0.69	0.706	0.691	0.729	0.705	0.686	0.728
Ca	0.786	0.789	0.829	0.824	0.828	0.81	0.81	0.824	0.687
Na	0.026	0.033	0.027	0.027	0.022	0.025	0.024	0.026	0.021
Κ	0	0.001	0	0.001	0	0.001	0.002	0.001	0.001
Catsum	4.008	4	4.001	3.994	3.987	4.001	4.003	4.021	4.009
Wo	39.74	41.48	43.11	43.23	43.58	41.95	42.03	42.11	35.39
En	35.2	35.44	35.88	37.04	36.37	37.75	36.59	35.05	37.51
$\mathbf{Fs}$	23.74	23.08	21.01	19.73	20.05	20.30	21.38	22.84	27.10
Mg#	60.73	60.56	63.07	65.25	64.46	65.03	63.12	60.55	58.05
	2+ / 1 / 1	2+ · · · · · · · · · · · · · · · · · · ·							

 $Mg\# mole Mg^{2+}/mole (M^{2+} + Fe^{2+}).$ 

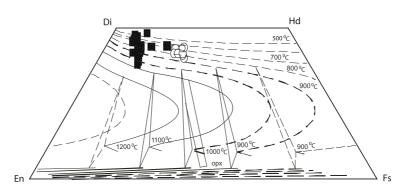


Figure 5. Di-Hd-En-Fs quadrilateral of clinopyroxene (Cpx). Filled squares and open circles both represent clinopyroxene from mylonitic gabbro. Open circle data represents Cpx porphyroclast. Temperature curves drawn after Bird *et al* (1986). Di: Diopside, Hd: Hedenbergite, En: Enstatite and Fs: Ferosilite.

colour of edenite in veins vary from greenish tint to more brown as the Ti content increases. Their characteristics are high in TiO<sub>2</sub> (>1 wt%) and Al<sub>2</sub>O<sub>3</sub> (9–11 wt%) contents, sometimes have a low Cl content (< 0.10 wt%). They are comparable to the

brown high aluminum hornblendes reported from oceanic gabbros elsewhere (Vanko and Stakes 1991; Gillis 1996). These amphiboles are generally considered to be late-magmatic phases directly crystallised from magma or formed by late-magmatic

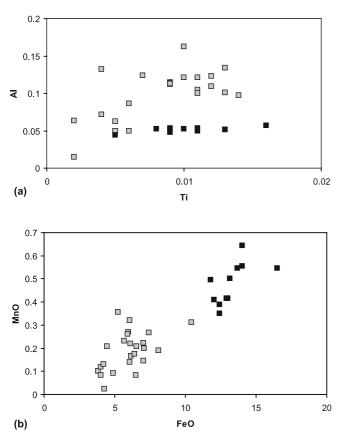


Figure 6. (a) Binary plot of Al (a.p.f.u) vs. Ti (a.p.f.u) of magmatic and vein Cpx, (b) Binary plot of MnO (wt%) vs. FeO<sub>t</sub> (wt%) of magmatic and porphyroclast of clinopyroxene (Cpx). Open squares represent magmatic Cpx and filled squares represent Cpx porphyroclast. Both the samples are from mylonite gabbro.

replacement of pyroxene (e.g., Miyashiro and Shido 1980; Mevel 1984; Kagiamanidou 1986; Vanko and Stakes 1991; Marion et al 1991; Gillis 1996). However, present vein amphiboles are Cl-rich and the chlorine concentrations, reach up to 0.62 wt% Cl, probably represent the products of interaction by anomalously saline hydrothermal fluid. Vein amphiboles show uniformly high Cl content as compared to the hornblende those have replaced clinopyroxene (figure 8). Also the high Cl amphiboles show higher Na and K content (figure 9). A graph showing Ti vs. tetrahedral Al  $(Al^{IV})$ in amphiboles can, in many cases be used to ascertain qualitatively the metamorphic grade of the host rock. Low-grade actinolitic hornblendes have lower Ti and  $Al^{IV}$ , and higher grade hornblendes have higher Ti and Al<sup>IV</sup>. Actinolitic hornblendes are mostly Si-rich and Al-poor and define a distinct field in Si vs. Mg/Mg + Fe binary diagram (figure 10). As the  $Al^{IV}$  content of amphibole is essentially temperature dependent (Moody et al 1983; Blundy and Holland 1990), the difference in concentration of Al among actinolites and edenite

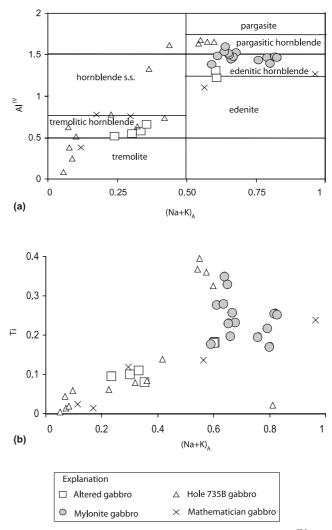


Figure 7. Representation of amphiboles in (a)  $Al^{IV}$  vs.  $(Na + K)_A$  diagram and (b) Ti vs.  $(Na + K)_A$  diagram. Altered gabbro (DR 02) and mylonite gabbro (DR 01) are from present data. Hole 735B, SWIR and Mathematician gabbro data taken from Stakes *et al* (1991) and Stakes and Vanko (1986), respectively.

thus allow to estimate the temperatures. They also do not vary to each other but Al and Na covary. CaO is nearly constant (10–11 wt%) and  $Al^{VI}$  is low for all, especially for actinolitic hornblende (DR 02, table 5).

#### 3.2.4 Epidote

Representative epidote compositions are given in table 6. The range of  $\text{Fe}^{3+}/(\text{Fe}^{3+} + \text{Al}^{3+})$  ratios (pistacite content) for individual grains is <0.23 (mostly < 0.17). The MnO content of all epidote varies from 0.02 to 0.11 and TiO<sub>2</sub> varies from 0.002 to 1.81; there is no difference in these minor elements with respect to their occurrence or alteration type. The epidote has nearly constant SiO<sub>2</sub> and CaO but varies significantly in iron and alumina. Weak iron enrichment within the epidote is also

Table 5. Representative analyses of amphibole.

Sample no.	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-02-1	DR-02-1	DR-02-1	DR-02-1
Oxides (wt%)									
$SiO_2$	43.53	43.61	43.37	43.58	43.23	50.06	50.74	50.92	48.52
$\mathrm{TiO}_2$	1.72	1.92	2.26	1.49	2.22	0.97	0.91	0.87	0.69
$Al_2O_3$	10.20	10.50	10.30	9.73	10.20	5.08	4.83	4.40	5.17
$Cr_2O_3$	0.05	0.05	0.09	0.14	0.01	0.00	0.00	0.00	0.00
$\rm FeO_t$	14.63	14.25	14.25	14.68	15.37	14.07	13.78	14.37	12.95
MnO	0.20	0.20	0.14	0.17	0.12	0.23	0.22	0.35	0.21
MgO	12.06	11.98	11.95	11.98	11.23	13.75	14.31	14.13	14.20
CaO	10.92	11.00	10.84	11.09	10.89	10.26	10.27	10.62	10.56
$Na_2O$	2.20	2.45	2.56	2.43	2.43	1.14	1.03	0.83	1.15
$K_2O$	0.60	0.42	0.37	0.46	0.60	0.04	0.06	0.01	0.09
NiO	0.00	0.03	0.00	0.00	0.01	0.05	0.07	0.00	0.07
Cl	0.49	0.15	0.09	0.58	0.51	0.07	0.04	0.00	0.02
Total	96.60	96.55	96.21	96.32	96.82	95.72	96.26	96.49	93.61
Si	6.572	6.55	6.535	6.614	6.544	7.422	7.459	7.484	7.348
Al(IV)	1.428	1.45	1.465	1.386	1.456	0.578	0.541	0.516	0.652
Т	8	8	8	8	8	8	8	8	8
Al(VI)	0.387	0.408	0.365	0.354	0.363	0.309	0.295	0.246	0.27
Ti	0.195	0.217	0.256	0.17	0.253	0.109	0.101	0.096	0.079
$\operatorname{Cr}$	0.005	0.01	0.01	0.016	0.002	0	0	0	0
Fe(III)	0	0	0	0	0	0	0	0	0
Fe(II)	1.846	1.789	1.795	1.863	1.945	1.744	1.693	1.766	1.64
Mn	0.025	0.025	0.018	0.022	0.015	0.029	0.027	0.044	0.026
Mg	2.715	2.682	2.684	2.711	2.534	3.038	3.136	3.096	3.206
Ca	1.767	1.77	1.751	1.802	1.766	1.63	1.617	1.672	1.714
C + B	6.942	6.902	6.879	6.937	6.878	6.859	6.87	6.92	6.935
Na	0.645	0.713	0.749	0.714	0.712	0.327	0.292	0.236	0.339
К	0.115	0.08	0.071	0.089	0.117	0.008	0.012	0.002	0.017
А	0.760	0.793	0.82	0.803	0.829	0.335	0.304	0.238	0.355
$X_{Ab}$	0.46	0.49	0.59	0.55	0.60	n.a.	n.a.	n.a.	n.a.
$T^{\circ}C$	825	818	806	801	796	n.a.	n.a.	n.a.	n.a.

especially one grain shows iron content (FeO<sub>t</sub>) up to 9 wt%. Present epidote composition show similar range in Al-Fe<sup>3+</sup> substitution (figure 11a). In fact the NCIR gabbroic epidote compositional range overlaps the field of epidote trend in metagabbro of SWIR or MARK (figure 11a).

# 3.2.5 Chlorite

 $X_{Mg}$  (mole Mg/mole Mg+Fe) of chlorite varies from 0.76 to 0.85 (table 7). FeO<sub>t</sub> content of present chlorite samples are variable (8.36 to 14.32 wt%). Chlorite of present metagabbro also exhibits high  $X_{Mg}$  as compared to chlorite from MARK metagabbro (figure 11b).

## $3.2.6 \quad Quartz$

As expected the quartz is composed of nearly pure SiO<sub>2</sub>, occasionally accompanied by trace amounts

of  $Al_2O_3$ ,  $TiO_2$ ,  $FeO_t$  or CaO. The nonsilica components never exceed 1 wt% and do not vary with the mode of occurrence of quartz.

## 3.2.7 Ilmenite

Ilmenite contains 1.4-8.5% MnO and 0.07-0.09% Cr<sub>2</sub>O<sub>3</sub>. Titanite also present locally and probably replaces titanomagnetite.

## 4. Discussion

Gabbros from NCIR represent a range of mineral assemblages resulted under variable physical-chemical conditions, high temperature amphibole-rich mylonite gabbros to highly altered greenschists. The gabbros from off-axis high, north of Vityaz TF, after cooling and crystallisation, underwent ductile deformation. A late magmatic stage was associated with intrusion of Fe-Ti rich magma to form the oxide rich layers (figure 2).

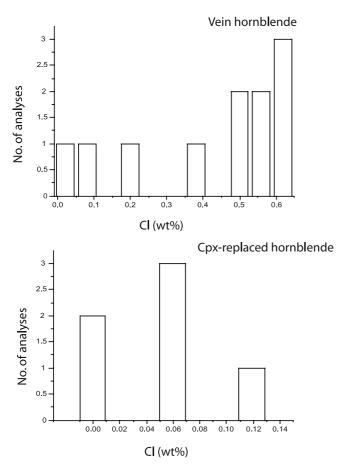


Figure 8. Cl-histogram of vein amphibole and clinopyroxene (Cpx)-replaced amphibole.

In addition, a later stage of subsolidus deformation, produced strongly foliated rocks resulting in the formation of porphyroclastic and mylonitic textures. In contrast, altered gabbro from corner high, adjacent to Vityaz TF, underwent low temperature greenschist grade alteration only with typical greenschist mineralogical assemblages (figure 3). The low temperature metamorphic minerals observed in late veins correspond to greenschist facies conditions and lower temperatures formed during cooling of lithosphere once shearing of the rocks (caused due to the presence of active TF) ceased. The brittle deformation of the altered gabbro can be ascertained by the presence of bending of twins, highly fractured and undulatory extinction of plagioclase grains.

# 4.1 Condition of mineralogical changes in gabbro

Mineralogical characters of present gabbro reflect difference in their alteration style due to temperature constraints and mode of emplacement due to tectonic evolution in the present study area. Among the minerals studied, amphibole compositions suggest evolutionary processes of present

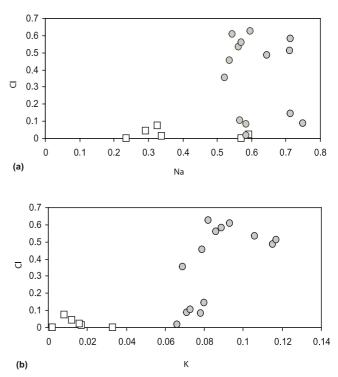


Figure 9. Representation of amphiboles in binary diagrams of Cl (wt%) vs. (a) Na (a.p.f.u) and (b) K (a.p.f.u). Filled circles represent vein amphibole and open squares represent amphiboles replaces Cpx.

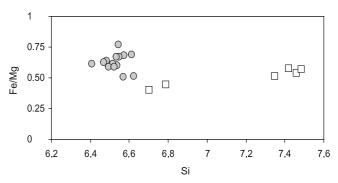


Figure 10. Representation of amphiboles in Fe/Mg vs. Si (a.p.f.u.) binary diagram. Filled circles represent vein amphiboles from mylonite gabbro and open squares represent amphibole from altered gabbro.

gabbro samples. Hornblende most likely to be formed at temperature above  $550^{\circ}$ C (Liou *et al* 1974) to  $600^{\circ}$ C (Spear 1981). The gabbro mineralogy from off-axis high adjacent to Vityaz TF is consistent with sites 921 to 924 of MARK area, Hole 735B at the Atlantis fracture Zone along the SWIR. However, characteristic features of the deformation fabrics and metamorphic assemblages in the site 735 gabbros suggest that deformation at Hole 735B was started at hypersolidus conditions and continued from granulite to amphibolite facies (550°–850°C, Stakes *et al* 1991). The formation of brown hornblende and its mineralogy (high Ti

1	b	0 1			-
Sample no.	DR-01-1	DR-01-1	DR-01-1	DR-01-1	DR-01-1
Oxides (wt%)					
$\mathrm{SiO}_2$	38.79	38.45	39.84	38.82	38.48
$\mathrm{TiO}_2$	0.00	0.08	0.02	1.81	0.00
$Al_2O_3$	26.35	26.29	21.98	25.41	26.87
$Cr_2O_3$	0.00	0.05	0.06	0.01	0.01
$\rm FeO_t$	7.62	7.60	9.09	6.95	7.75
MnO	0.10	0.03	0.09	0.11	0.11
MgO	0.33	0.22	2.61	0.49	0.04
CaO	23.05	23.06	20.56	23.24	23.24
Na <sub>2</sub> O	0.04	0.04	0.09	0.06	0.03
$K_2O$	0.00	0.00	0.01	0.00	0.01
NiO	0.00	0.01	0.00	0.00	0.00
Cl	0.00	0.00	0.02	0.00	0.02
Total	96.28	95.83	94.37	96.91	96.55
Cations based of	on 25 oxygens				
Si	6.222	6.2	6.537	6.185	6.164
Ti	0.001	0.01	0.003	0.217	0
Al	4.982	4.999	4.252	4.774	5.074
Cr	0	0.006	0.007	0.001	0.002
$\mathrm{Fe}^{2+}$	1.022	1.025	1.248	0.926	1.038
Mn	0.014	0.004	0.013	0.014	0.015
Mg	0.078	0.053	0.638	0.115	0.01
Ca	3.961	3.984	3.615	3.968	3.99
Na	0.013	0.014	0.03	0.018	0.008
К	0.002	0	0.003	0.001	0.002
Catsum	16.295	16.295	16.346	16.219	16.303

Table 6. Representative analyses of epidote. Total iron calculated as  $FeO_t$ .

content, table 5) suggest the formation of present mylonitic gabbro initiated in a rock-dominated system at 700°C or higher. In addition, amphibole chemistry also helps to decipher the condition of alteration and chemistry of interacting fluids. Vein amphibole (figure 2a) is a major sink for Cl and characteristically show Cl-enrichment. Chlorine, a major constituent of sea water thus enters the structure of amphibole during sea water-rock interaction. Thus, Cl enrichment within the amphibole is totally structurally controlled (table 5). The chlorine composition can also provide useful information on fluid composition. Some of the present aluminous edenitic hornblendes contain significant amount of chlorine (up to 0.62 wt%) suggesting formation of sea water-rock interaction (Ito and Anderson 1983; Batiza and Vanko 1985; Vanko 1986; Mevel 1988). Also the restricted occurrences of Cl-rich amphibole (in mylonitic veins only) suggests sea water of a concentrated brine substantially influences its formation at the time of mylonitisation (figure 8). Additionally, the Cl-rich amphibole are uniformly Na and K-rich  $(Na_2O > 2 \text{ and } K_2O > 0.4 \text{ wt\%}, \text{ table 5, figure 9}).$ Though, interaction of high temperature magmatic Cl-rich fluids in gabbroic rocks well reported from Kane fracture Zone, Mid-Atlantic Ridge (Kelley and Delaney 1987), Cl-rich amphiboles from present study seems to be the result of sea water circulation. The Na and K vs. Cl diagrams suggest the interacting fluid phase particularly enriched in Na, K and Cl.

There is difference in compositions between hornblende replaces primary phases (hence clinopyroxenes) and those filling veins. The Al content of amphibole increases with increasing metamorphic grade (Deer et al 1992), as noticed in mylonitic gabbro and the low Al actinolitic hornblende ( $\sim Al_2O_3$  value goes down up to 4.83 wt%) of altered gabbro from corner high also supports their low temperature metamorphic mineralogical assemblages (figure 7). In addition, low Ti (TiO<sub>2</sub> always less than 1 wt%) favourably suggest their formation at low temperatures. The typical mineralogical assemblages of corner high gabbro suggest that altered gabbroic rocks underwent brittle deformation and associated hydrothermal alteration to greenschist assemblages. Based on metamorphic grade and the pattern of elemental enrichments, we infer that the present gabbro was altered in the deeper levels by the hydrothermal systems. At shallow levels (< 1 km) and recharge

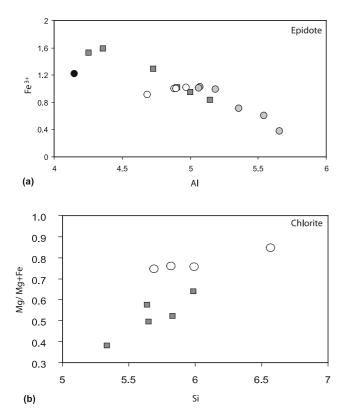


Figure 11. Representation of epidote and chlorite in binary diagrams. (a) Al vs.  $Fe^{3+}$ . Filled squares epidote from MARK area (Gillis *et al* 1993), grey circles from SWIR (Stakes *et al* 1991) and open circles and black circles epidote from NCIR. Black circle represents vein epidote. (b) Mg/Mg+Fe vs. Si. Filled squares represent chlorite from MARK area (Gillis *et al* 1993) and open circles are chlorite from NCIR, respectively.

zones, oceanic rocks commonly have a tendency to gain Na and Mg, and loss of Ca (Alt et al 1986; Harper et al 1988; Lecuyer et al 1990). In contrast, oceanic rocks in discharge zones, however are commonly epidotized, and show a gain of Ca,  $\pm$ Fe,  $\pm$ Al and loss of Mg (Harper *et al* 1988; Nehlig et al 1994). The abundance of albite with close association of chlorite especially within the altered gabbro from corner high (figure 3b, f) suggests interaction of present gabbro with fluids enriched in Na/Mg at 300°C temperature under greenschist facies condition (Seyfried et al 1978). Continuous supply of a Mg-depleted, Si-rich fluid, high in Na/Ca molal ratio, enhances the albitisation processes (Ab content reaches up to  $96\,\mathrm{wt\%})$  of the primary mineralogy. Such a condition generally prevails where Si-rich, Mg-free and Ca-depleted fluids which form during high temperature processes at depth are cooled both adiabatically and conductively during ascent (Von Damn *et al* 1985). Reaction of these fluids with fresh or previously altered basalt or gabbros are likely to be responsible for extensive albitisation. Seyfried *et al* (1978) and Seewald (1987) inferred

that the removal of Mg from sea water at the pillow exterior (owing to chlorite formation) may provide high Na/Mg ratio fluids for pillow-core alteration and albitisation. Mineralogy of present altered gabbro also exhibits similar kind of fluid interaction to cause pervasive albitisation once they have emplaced the sea bottom.

Clinopyroxene in mylonitic gabbro which crystallized from circulating hydrothermal fluid and also inherited the chemical signatures of fluid may be considered as secondary (table 4b). MnO enrichment (up to 0.64 wt%) suggests that these veins are affected by circulation of hydrothermal fluids with evolving compositions (table 4b). Their iron enrichment trend also suggests that the fluids from where they crystallized must be iron-rich (figure 6). Also the difference in the temperature range for clinopyroxenes within the mylonite zone and outside of it suggests difference in their temperature of formation (figure 5). The progressive Fe-rich composition at late stage fluids is also exemplified by relatively high Fe concentration in vein epidote as compared to those replacing plagioclase (figure 11a).

In the absence of quartz, the temperature of amphibole-plagioclase equilibration may be quantified through the extent of  $[NaSi]_{-1}[CaAl]$  exchange among the phase components (Holland and Blundy 1994):

$$= Na_2 CaMg_5 Si_8 O_{22} (OH)_2 + CaAl_2 Si_2 O_8 anothing$$

They used nonlinear least squares regression and a well constrained plagioclase solid solution model to derive within site and cross site parameters for amphibole solid solutions. This allows the calculation of temperature for a particular amphibole and plagioclase composition. Calculated temperatures for present coexisting amphibole and plagioclase pair varies from 796° to 825°C which certainly fall under high temperature metamorphism in ocean floor setting. More analyses of coexisting plagioclase-amphibole pairs are required to assess the accuracy of calculated temperatures. It can be concluded that the metamorphism of off-axis gabbro (mylonite gabbro) of the present study may have started about 700°–800°C. Typical mineral assemblages include plagioclase, Fe-Ti oxide, green and brown amphibole. These amphibole gneisses are texturally and mineralogically similar to dredged samples from Vema fracture zone (Honnorez *et al* 1984) and the Mid-Cayman rise (Ito and Anderson 1983). Comparison with amphibole chemistry with other

Sample no.	DR-02-1	DR-02-2	DR-02-3	DR-02-4
Oxides (wt%)				
$SiO_2$	33.06	29.41	28.18	28.79
$\mathrm{TiO}_2$	0.03	n.d.	0.04	0.01
$Al_2O_3$	15.92	17.37	19.93	19.40
$\mathrm{Cr}_2\mathrm{O}_3$	0.04	0.02	n.d.	0.02
$\rm FeO_t$	8.36	13.83	14.32	13.39
MnO	0.27	0.18	0.19	0.29
MgO	25.84	24.19	23.34	23.51
CaO	0.67	0.10	0.05	0.08
Na <sub>2</sub> O	0.12	0.01	0.02	0.01
$K_2O$	0.07	0.01	0.00	0.03
NiO	0.00	0.12	0.04	n.d.
Cl	0.06	n.d.	n.d.	n.d.
$P_2O_5$	n.d.	0.03	0.02	0.04
Total	84.42	85.27	86.13	85.57
Cations based of	n 28 oxygens			
Si	6.567	5.993	5.69	5.819
Ti	0.004	n.d.	0.006	0.002
Al	3.728	4.17	4.744	4.622
Cr	0.007	0.003	n.d.	0.004
Fe	1.388	2.356	2.419	2.263
Mn	0.046	0.03	0.033	0.05
Mg	7.649	7.343	7.024	7.082
Ca	0.142	0.023	0.01	0.017
Na	0.045	0.004	0.008	0.005
Κ	0.017	0.002	n.d.	0.008
Catsum	19.593	19.924	19.934	19.872
$\rm X_{Mg}$	0.85	0.76	0.74	0.76

Table 7. Representative analyses of chlorite. Total iron calculated as  $FeO_t$ .

known occurrences of oceanic gabbro has shown that amphibole from mylonite gabbro is similar in composition with high temperature gabbro from Hole 735B, SWIR (figure 7, Stakes *et al* 1991). Ti and  $Al^{IV}$  content of present amphibole (from mylonite gabbro) is comparatively high from amphibole of Mathematician Gabbro (figure 7, Stakes and Vanko 1986). However, amphibole from altered gabbro is compositionally similar with low temperature gabbro of SWIR as well as gabbro from Mathematician Ridge (figure 7).

## 4.2 Geodynamic implications

Metamorphosed mylonitic gabbroic rocks from the off-axis high, north of Vityaz TF setting record high temperature deformation metamorphism and recrystallisation. Fractures and microfractures in the gabbroic rocks that are filled with lower amphibolite to greenschist facies mineral assemblages suggest that the plutonic sequence has undergone brittle deformation during progressive cooling throughout the subsolidus regime, following the high temperature plastic deformation. The abundance of amphibole in the zones of crystal-plastic deformation indicates these zones acted as the major conduits for hydrothermal fluids (figure 2, sample no. DR 01). Strong foliations as shown by amphiboles suggest stress during recrystallisation. The nearly homogeneous amphibole compositions from pargasite to actinolite in altered gabbro reflect progressive unroofing in the footwall of a brittle-ductile normal fault. The active detachment surface of the Vityaz megamullion facilitates the extension across an amagmatic spreading segment triggered the formation of altered gabbro (Drolia and DeMets 2005). Detachment faulting associated with crystal-plastic deformation at off axis high location would have thinned the crust and allowed the penetration of sea water.

The formation of amphibole veins in selective samples and alteration of clinopyroxene rim by amphibole in other samples suggest that the crystal-plastic deformation might be localised. Offaxis high set up, adjacent to Vityaz TF, as well as the presence of Vityaz megamullion probably represent extensional fault domains where lithospheric stretching of the lower crust occurred via brittle-ductile deformation (Karson and Dick 1983; Mevel et al 1991). Presence of Vityaz TF, however, facilitates intense shearing helped to develop immense stress made the rock foliated during recrystallisation. Gabbros from corner high set up had solidified and cooled and subsequently fractured which facilitate to get the access of deep penetration of sea water, finally led to become metamorphosed. Rapid cooling due to the presence of Vityaz TF regulates the temperature of metamorphism and restricts the grade of metamorphism up to greenschist facies (figure 3, sample no. DR 02). The mineralogical assemblages further suggest declining fluid temperature of hydrothermal veins.

#### 5. Conclusion

The present study confirms that the gabbros from NCIR record both the high temperature metamorphism and low temperature hydrothermal alteration in a segment of lower crust formed at a slow spreading ridge. The amphibole in mylonite gabbro is closely related with plastic deformation and its mineral chemistry suggests that they have formed at about 800°C. The result of present study on mylonite gabbros thus provide a typical high temperature amphibolite facies assemblage of calcic plagioclase and hornblende. Based on their textural relations, we infer that the monomineralic amphibole veins from off-axis formed essentially at deep crustal level prior to its emplacement at present level which is common everywhere in the ocean basins (Phipps Morgan and Chen 1993). The temperature estimates for the amphibole veins in the present study are comparatively high as reported by Manning et al (1999) for amphibole + plagioclase veins in gabbros from Hess Deep  $(687^{\circ}-745^{\circ}C).$ 

Altered gabbros from corner high adjacent to Vityaz TF formed exclusively under greenschist facies condition (300°C), resulted the formation of greenschist mineralogical assemblages. Occurrences of actinolitic hornblende suggest locally the temperature had attained up to 450°C. Evidences of decreasing fluid temperatures as indicated by precipitation of albite, chlorite and deposition of late-stage quartz veins/mosaic in the groundmass. Present study also suggests that low temperature alteration probably occurred once gabbro tectonically emplaced to the sea bottom. Progressive Feenrichment trend is also noticed in the late stage fluid.

The proactive role of tectonic activity (hence fault displacements and presence of megamullion) over magmatism in the present slow spreading NCIR is well exemplified by the present finding of gabbroic rocks and further enhanced by postcrystalline mineralogical and textural changes due to rigorous hydrothermal circulation. Presence of the major detachment fault (hence Vityaz megamullion), which had a long deformation history, unroofed the lower crust and presumably caused rapid cooling. Thus, detachment faulting played a major role not only in the tectonic evolution of this crustal segment of NCIR but also in the timing and extent of alteration of the present lower ocean crust. We further conclude that deformation is necessary at present off-axis environment for prolonged alteration of ocean crust at slow spreading environment.

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