

Phoscorite From Ultrabasic Rock in Carbonatite Complex of Tiruppattur, Tamil Nadu, India

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ABSTRACT

Phoscorite is a rare apatite and magnetite rock with accessories of olivine and its heteromorphic variants. The close association of phoscorite and carbonatite in ultrabasic alkaline complex of Tiruppattur (N 12o15'-12o 30' E 78o 15'-78o25') provides more information about volatile concentration of H₂O, CO₂, CH₄, SO₃, F, Cl, and P₂O₅ in mantle source and their roles on magmatic evolution of highly differentiated carbonatite complex of this area. Field evidences show that occurrence of crystallized high pressure temperature mineral (HPTM) at depth is seen in some rocks in this area. Early formed HPTM enveloped by series of layers of supercritical fluids derived by a small degree of partial melt of carbonated peridotite at mantle is stable on Earth's surface. Suddenly cracked conduit through which very rapid ascent of supercritical fluid of natrocarbonatite carrying HPTM from mantle at a speed ranging from 70 to 100 m/s within <10 hours of duration without any friction and energy loss, retains its structural stability to Earth's surface. Phoscorite and carbonatite fluids in peridotite are caused mineralization apatite and ilmenorutile in this complex.

Keywords: phoscorite, apatite, Magnetite, Silicate-perovskite Ultrabasic, Carbonatite complex of Tiruppattur, Tamil Nadu

INTRODUCTION

Phoscorite is a rare apatite and magnetite rock. It occurs in carbonatite ultrabasic alkaline complex of Tiruppattur (N 12o15'-12o 30' E 78o 15'-78o 25'). The rock is composed of olivine and any one of its heteromorphic equivalents of diopside, acmite, hornblende, phlogopite, magnetite, calcite, feldspar and foids / quartz as accessory mineral. It provides more information about mantle source and deep-seated magmatic processes controlled by volatile concentration of H₂O, CO₂, CH₄, SO₃, F, Cl, and P₂O₅. These volatiles are caused formation of ultralow viscous supercritical fluid of natrocarbonatite at lower mantle where ultralow velocity seismic zone is prevailed.

GEOLOGICAL SETTING

Carbonatite complex of Tiruppattur (N 12o15'-12o 30' E 78o 15'-78o25') is comprised with 3 carbonatite units located at Sevvattur, Jogipatti and Onnakarai villages [1- 6] in association with ultrabasic rocks and alkali syenites (Fig. 1). Conesheets of carbonatite of Sevvattur is exposed (1500 x 300 m) NW at contact between ultrabasic and syenite complex in Sevvattur basin. The carbonatite varies from ferrocarnatite, beforosite to sovite. Apart from the essential minerals [6] of calcite, dolomite, ferrocarnatite, accessory minerals of magnetite, perovskite, pyrochlore, zircon, apatite, aegirine augite, riebeckite, oligoclase and phlogopite are found as accessory minerals in the carbonatite. Concentration of pyrochlore is observed in ferrocarnatite. Lueshite (Sodium bearing reddish brown pyrochlore) shows expansion cracks in albitite exposed at south eastern periphery of carbonatite of Sevvattur. Semitransparent and semiprecious euhedral pale reddish brown zircon is found as accessory high pressure temperature mineral (HPTM) in carbonatite of Sevvattur and metamict dark brown euhedral zircon occurs in zircon syenite exposed about 1.5 km SW of Samalpatti village. An ultrabasic rock exposes at north western boundary of the carbonatite. Phoscorite occurs as diatremes up to 7500 x 10 cm at contact of a vermiculite body on NW part of Sevvattur carbonatite. It is the largest vermiculite mine in India formed by alteration of phlogopite-oligoclase pyroxenite.

Thin veins of sovite, beforosite, ferrocarnatite, barite and benstonite occur in ultrabasic rock exposing in Onnakarai area. The ultrabasic rock is intercalated ultramafic rock essentially composed of pyroxenite and peridotite. Phoscorites in ultrabasic rock are exposed at about 200 m SW of Onnakarai village. They occur as rootless thin diatremes of varying sizes from 100 x 5 to 5000 x 10 cm in ultrabasic rock near the village. At some places dunite and peridotite with pyroxenite dykes are seen Magnesite and kankar are released products at some places. Just south of the village barite

vein 100x15cm is found.

A skarn rock (1500 x 50m) extends from Garigaipalli to Onnakarai villages [7]. Near Garigaipalli a high grade skarn trends towards NNE-SSW and steeply dipping towards WNW. An intermittent shear zone trending WNW-ESE and shift it WNW-ESE direction. There it is attained low grade metamorphism towards south forming discontinuous arc like form dipping inwards. Ultra low viscous ferrocarnatite veins of <100x5 cm carrying euhedral magnetite (5mm) are seen along some shear zones trending in WNW-ESE direction. Monazite bearing pegmatite and galena bearing fine-grained albitite [8] occur at contact between syenite and skarn in SW part. Ultrabasic nodules of varying sizes between 1 and 50 cm dimensions [7] are seen in the skarn. The nodule occasionally contains native Cu and Ag in addition to chalcopyrite, covellite, cuprite, malachite and azurite, sphalerite, pyrite, pyrrhotite and galena [8]. The skarn and nodules are composed with tightly folded veins with alternate miniature swelled micro-basins and tightly compressed micro-domes. A single nodule comprises single basin and a dome. Wollastonite, diopside, garnet scapolite and calcite are seen as major minerals. Epidote and vesuvenite crystallize along some weak planes of the rock. Ultrapotassic syenite of microcline-melanite syenite [9] and garnet-syenite, anorthoclase-riebeckite potassic syenite occur as major rock types in Jogipatti basin.

NE-SW trending elongated small rings of carbonatite veins are seen in Jogipatti village along fold axial planes of the structural basin. The basin is located hardly 5 km SW of Sevvattur basin (Fig. 1). Both basins occur in NE-SW direction trending along Alangayam rift zone which is further extending over Amirthi fault near Vellore and further NE to Bay of Bengal, on SW it extends over 400 km into Arabian Sea [10]. The present width of the rift zone exceeds over 50 km at some places. Faulted Jogipatti basin including carbonatites and potassic syenites has uplifted 1600 m from NNE-SSW to NE-SW relative to Sevvattur basin which is sequentially filled with sodium syenites. Along a fault zone lying between the basins Jogipatti basin is rotated to 16o NE-SW [2, 5] Younger co-magmatic sequences are progressively differentiated and are emplaced in sequences in SW parts of these structural basins (2). There exists magmatic continuum between older sodium and younger potassium rich syenites [2, 5].

METHODOLOGY

The phoscorite was analyzed using High Resolution Scanning Electron Microscope attached with EDAX probe in the Laboratory, Department of Metallurgical and Material Science, Indian Institute of Technology, Chennai. The elemental analyses were recalculated and Rittmann's normative proportions were calculated [11]. Iron ores were subjected X-ray powder diffractions analyses with Fe Ka radiations were made in the crystallography Department of Geology Faculty of Moscow State University, Russia. The minerals had mixed proportions of iron ores and they were difficult to distinguish individual components and indexing their faces. However, all the peaks and their intensities are given for future references (Table 1). Characteristic peaks of magnetite of were listed (Table 2) for correlation.

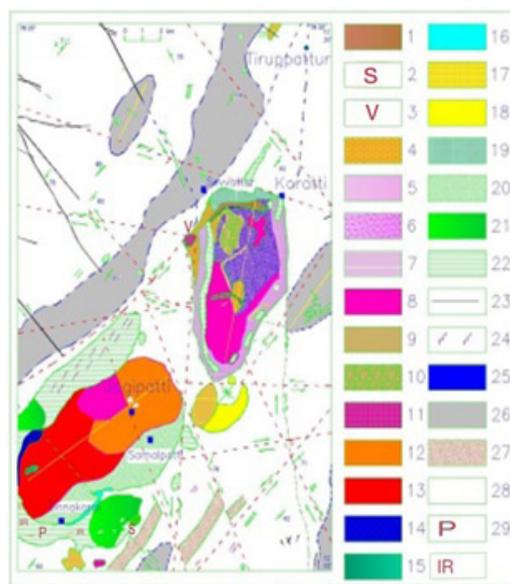


Fig1. Simplified Geological map of carbonatite complex of Tiruppattur, Tamil Nadu, India

1 Carbonatite, 2 S Shonkinite 3 V Vogesite 4 Oligoclasite 5 Hornblende syenite 6 Hornblende porphyritic syenite 7 Augite syenite 8 Porphyritic augite syenite 9 Foliated syenite 10 Foliated porphyritic syenite 11 Hornblende biotite syenite 12 Riebeckite syenite 13 Garnetiferous syenite 14 Microcline melanite syenite 15 Wollastonite syenite 16. Skarn 17 Granite 18 Granodiorite 19 Biotite oligoclase pyroxenite 20 Biotite pyroxenite 21 Ultrabasics 22 Ultramafics 23 Dolerites 24 Pegmatite / aplite 25 Albitite 26 Charnockite 27 Pyroxene granulite 28 Granite gneisses P Phoscorite IR Ilmenorutile

PETROGRAHY AND PETROCHEMISTRY

On hand examination in the field, phoscorite appears to be a feldspar-magnetite rock and it is subjected to wet-gravimetric analysis. Before wet-gravimetric analysis, spectroscopic analyse shows a high concentration of P, Fe, Ti, Y, Nb and Zr reveals that the rock is essentially composed of apatite nioborutile (ilmenorutile), zircon and magnetite Under polarizing microscope apatite exhibits euhedral form with strong relief and first order grey interference colour and opaque iron ore. The rock is termed as phoscorite. In Sevvattur basin it is a very coarse-grained rock with inequigranular porphyritic texture. Bluish yellow green euhedral apatite up to 4x2x1 cm is seen with blades or booklets of phlogopite amidst apatite. Large grain of magnetite up to 6 cm is seen.

On the other hand, it occurs 700 m NE of Mottusulakkarai village and 200 m SW of Onnakarai village and carries reddish brown massive apatite up to 2x0.5 cm and magnetite up to 1.5 x 0.5 cm with coarse-grained inequigranular porphyritic texture. Scanning electron microscope images show particle size of (Fig. 2) larger plates of euhedral prism and pyramid shaped apatite up to 180x60x3 µm form with length, breadth and thickness. Tabular apatite grains overlap one over the other as platelets. Apatite has transverse irregular gas cavities (Fig:-2). A globular gas cavity of 150 µm is filled with smaller but inequigranular prisms of apatite and other minerals. The gas cavities are aligned along peripheral portions and at the central portion of globular cavity. Around periphery of some irregular cavities, rims of very fine-grained minerals are developed. Cavities are represented as fluid phases. Particle size of magnetite 250x150x30µm is found. Edges of magnetite are very sharp and steep. Phlogopite flake of 200x5x1 µm is seen.

XRD analyses of apatite (Cu Ka) and magnetite (Fe Ka) from Sevvattur, Jogipatti and Onnakarai are also listed in the Table 1. The 2Theta values for iron ores on Fe Ka radiations show characteristic peaks for magnetite on faces of Miller's indices. The slight deviations from normal peaks might have been due to solid solution and entry of other elements, distortion of crystal faces or interference of mixed phases formed at ambient temperature and pressure conditions. Table-2 such variations are seen in magnetite. Magnetite (mt1) shows very low values of 2Theta for Fe Ka radiations might have been representing entry of silica perovskite molecules in magnetite in Table 1.

Table1. X-ray powder diffraction data for apatite Cu Ka and magnetite for Fe Ka radiation .

Mt	490 Sevvattur				450 Onnakarai				1 Sevvattur				2 Sevvattur			
	2Theta	I ₀	CuKa A	FeKa A	2Theta	I ₀	FeKa	AlKa	2Theta	I ₀	CuKa A	FeKa A	2Theta	I ₀	FeKa A	
	38.22	41.20	2.3547	3.00	11.10	9	7.9711	10.00	6.80	4	12.9989	16.20	22.90	5	3.8701	4.90
	40.60	25.00	2.2220	2.80	22.45	3	3.9903	5.00	7.40	4	11.9463	15.00	38.12	32	2.3907	3.00
	41.44	14.70	2.1789	2.70	26.40	32	3.3761	4.20	23.30	7	3.8080	4.80	45.12	100	2.0094	2.50
	42.10	23.50	2.1462	2.70	27.55	13	3.1523	4.00	30.68	6	2.8557	3.50	47.24	6	1.5241	2.40
	45.20	100.00	2.0060	2.50	28.76	11	3.1041	3.90	38.38	37	2.3465	2.90	55.10	54	1.6669	2.10
	53.80	16.20	1.7874	1.80	29.40	25	3.0380	3.80	42.36	34	2.1337	2.70	69.12	16	1.3590	1.70
	58.84	19.10	1.3038	1.70	32.40	100	2.7032	3.00	45.40	100	1.9577	2.50	72.88	61	1.3000	1.00
	74.00	27.90	1.2909	1.60	32.70	57	2.7369	3.40	47.32	7	1.9210	2.40	81.72	52	1.2000	1.50
	78.46	8.80	1.2189	1.50	33.53	50	2.6727	3.40	52.40	9	1.7461	2.20				
	80.26	11.80	1.1961	1.50	34.58	19	2.5996	3.20	55.40	63	1.6582	2.10				
	81.78	51.50	1.1776	1.50	35.80	6	2.5082	3.20	63.64	12	1.4021	1.80				
	84.10	10.30	1.1510	1.40	35.76	6	2.5060	2.80	65.18	17	1.3560	1.70				
					40.52	21	2.2263	2.90	70.10	17	1.3000	1.70				
					42.58	5	2.1182	2.70	74.20	70	1.3000	1.60				
					44.40	4	2.0403	2.60	81.90	70	1.2000	1.50				
					47.50	25	1.9218	2.40								
					48.76	11	1.8678	2.30								
					49.20	11	1.8483	2.30								
					50.08	31	1.8214	2.30								
					51.10	13	1.7874	2.20								
					51.56	11	1.7611	2.20								
					52.88	10	1.7395	2.20								
					53.52	21	1.7122	2.20								
					54.40	4	1.6806	2.10								
					55.56	6	1.6265	2.00								
					59.40	5	1.5326	1.90								
					62.08	6	1.4951	1.90								
					63.00	13	1.4830	1.80								
					64.56	11	1.4435	1.80								
					66.84	9	1.4223	1.90								
					70.40	5	1.3000	1.70								

Mt	3 Sevvattur				4 Sevvattur				5 Sevvattur				6 Sevvattur			
2θ (°)	Ipo	CaKa A	FoKa A	2θ (°)	Ipo	CaKa A	FoKa A	2θ (°)	Ipo	CaKa A	FoKa A	2θ (°)	Ipo	CaKa A	FoKa A	
23.10	8	3.8503	4.80	23.30	16	4.9554	8.20	38.24	21	2.7780	2.9568	38.28	24	2.9523	2.9559	
38.16	26	3.3584	3.00	38.30	6	3.8177	4.80	41.44	26	2.1790	2.7378	44.70	26	2.0273	2.5471	
41.40	12	2.7780	2.70	30.40	4	2.9403	3.70	42.22	35	2.1400	2.0892	45.22	100	2.0051	2.5192	
42.20	5	2.1415	2.70	38.32	20	2.3489	3.00	45.20	100	2.0080	2.5208	55.18	26	1.8845	2.0912	
45.17	100	2.0073	2.50	41.44	14	2.1790	2.70	52.38	14	1.7418	2.1955	69.00	13	1.3810	1.7102	
47.32	6	1.9210	2.40	42.32	13	2.1357	2.70	55.34	18	1.6900	2.0894	73.98	43	1.2815	1.6107	
51.48	3	1.7751	2.20	45.24	100	2.0044	2.50	63.64	18	1.4824	1.8373	81.78	54	1.1770	1.4800	
55.16	33	1.6051	2.10	47.36	4	1.9195	2.40	68.64	10	1.3657	1.7184	94.10	10	1.0532	1.3233	
69.18	11	1.3579	1.70	52.24	5	1.7511	2.20	73.56	16	1.3441	1.6867					
73.93	55	1.3000	1.60	55.16	31	1.6551	2.10	75.00	30	1.3000	1.6452					
80.45	5	1.2000	1.50	82.90	6	1.4776	1.90	82.00	31	1.2000	1.4764					
81.80	44	1.2000	1.50	85.55	4	1.4540	1.80	84.00	13	1.2000	1.4477					
				89.18	11	1.3580	1.70	95.44	11	1.1000	1.3093					
				70.00	7	1.3441	1.70	99.00	9	1.0000	1.2739					
				73.88	31	1.30	1.60									
				80.70	3	1.30	1.50									
				81.85	35	1.20	1.50									
				92.30	5	1.20	1.30									

Table 2. Comparative studies of 2 Theta and intensity of peaks for magnetite and apatite

mt1	mt2	mt3	mt4	mt5	mt6	mt7	mt8	mt222	mts498	mt380	mt381	mtc495	mts488
6.80													
7.40													
23.36		17.90	17.90			23.02					26.08		
36.36	36.12	36.32	36.32	36.24	36.26	36.54	36.16	37.64	36.22	37.44		34.64	38.22
		41.44	41.44			41.38	41.38		41.44	38.10	41.00	41.20	41.44
		42.32	42.32	42.22	44.70	42.24	42.18		42.10	42.00		44.20	42.10
45.40	45.12	45.24	45.24	45.20	45.22	45.18	45.18	44.84	45.20	45.18	45.20	45.80	45.20
55.40	55.10	55.18	55.18	55.34	55.18	55.14	55.22	54.80	68.84	55.12	55.22	56.00	68.84
74.20	72.88	73.80	73.80	72.40	73.90	73.92	73.92	73.44	74.00	73.94	74.02	73.10	74.00
81.90	81.72	81.85	81.85	84.40	81.78	81.80	81.80	81.42	81.78	81.80	81.84		81.78
						98.90			84.10	94.22	94.26	90.76	84.10

Inner core of ultrabasic nodule free from any feldspar constituents in skarn rock is chemically analyzed. It is composed of SiO₂ 45.92, Al₂O₃ 6.12, Fe₂O₃ 5.52, FeO 5.33, MnO 0.18, MgO 9.60, CaO 22.46, Na₂O 1.08, K₂O 0.16, TiO₂ 1.26, P₂O₅ 0.19, CO₂ 1.02 and H₂O 0.50 wt% and has high content of MgO, CaO and SiO₂. The characteristic trace elements are Ba 2000, Zr 10, V 300, Cu 100, Zn 300, Cr 100, Ni 60, Co 30, and Pb 8 ppm. The normative proportion of the rock is cc 2.63, ap 0.46, il 1.63, mt 2.06, ol 16.64, hy 15.78, wo 37.85 and lab 22.76%. Microscopic studies reveal that presence of wollastonite, (Fig. 3) grossularite-andradite garnet and scapolite. It exhibits hypidiomorphic granular texture. Aplite and pegmatite of wollastonite-garnet sovite, wollastonite garnet syenite and melanite microcline ultrapotassic syenite have genetic link with skarn [7, 8]. Ferric iron are seen (Fig. 3) as opaque dusts [12, 15] in potassic feldspars in ultrapotassic melanite syenite, shonkinite and in ultrabasic rocks. Scapolite – arfvedsonite syenites are seen around carbonatites of Jogipatti.

Chemical analyses of magnetite from Sevvattur show progressively increasing silica content from 0.48 to 62.14%. The degree of oxidation (Fe³/Fe³+Fe²+Mn) of magnetites varies between 0.39 and 0.96. Maghemite phase is enriched with higher oxidation degree. This phase is found in all magnetite. Normative spinel is found in 498 and 13. Magnetite is polygenetic. Thin section studies reveal that it occurs as primary opaque cloudy dusts which are later oxidized and swept along grain boundaries, cleavage planes of calcite and they accumulated secondary crystals [12] in calcite of carbonatite. Scapolite – arfvedsonite syenites are seen around carbonatites of Jogipatti.

Some magnetite [13] have enriched with Ca ions on the basis of 32 (O, OH, F). Ca is present in 13 up to 3.205 ions (Table 4) indicating entry of perovskite. Si content of magnetite steadily increases from 498Si 0.141, 13Si 5.459, 21Si 8.717 and 47Si 11.840. Generally under alkaline condition porous silica is introduced to overall surface area in magnetite from 8.2 to 13.2 nm [14]. At HPT condition incorporation of Si in magnetite transits it into silicate perovskite under ambient HPT state.

Ca rich pyroxene [15] and Ti rich garnet show magmatic courses of differentiation [16] in this complex. Sr replaces Ca in apatite [13] on the basis of 26 (O, OH, H) Table 5. Fluorite and periclase enter in apatite. Si, Al and Mg (Table 6) substitute into apatite structure. Olivine content varies between 7 and 31%. Flour-apatite varies up to 90-97; fluorite up to 8; Ca bearing olivine 1 and portlandite has 5% (Table 5). Table 6 shows that 46-66% of apatite. It is enriched with Ca-olivine 23 portlandite 23, bytownite(ab₂₄ - an₇₆), 13% and nepheline 4 with or without kalsilite 1% are present to meet silica deficiency. Highly transparent and pale yellowish green phlogopite is found from phoscorite in Sevvattur. The ratio of Mg/(Mg + Fe) is 0.678 (Table 4) indicates it is phlogopite. It shows III or IV order interference colour. Structural

formula on the basis of 22 (O, OH, F), the phlogopite has lost considerable amount of Si, Al, Mg, K, and H₂O+. Ca exist 0.483 apfu which is higher than normal phlogopite (Table 4). Fluorine occurs in apatite up to 0.823 on the basis of 26 (O, OH, F) ions (Table 5). Normative fluorite reaches to 7 %. Pure apatite of Sevvattur is silica under saturated and has minute opaque inclusions of periclase up to 5%. SiO₂ varies up to 16.80% (Table 6) and it has normative nepheline 4 with or without kalsilite 1%. Apatite is present up to 55%. Locally, the ratio of apatite and iron ore is 79 : 21. There exists close relationship [17] between increase of apatite saturation and decreasing of silica content in mafic-alkali magma.

EDAX analyses (Table 7) of phoscorite of Onnakarai area show enrichment of FeO, TiO₂, Nb₂O₅, ZrO₂ and REE₂O₃. In Onnakarai the host rock of phoscorite is peridotite which is composed of Ca- olivine (Table 3) 45, clinopyroxene 32, calcic plagioclase 13 and apatite 6%. Phoscorite is composed of apatite 19-74, calcite 7, anhydrite 3 and iron ore 25% (Table 3 and 7). Olivine breaks down into pyroxene (451) and quartz at ambient temperature and pressure condition (Table 3). Table 7 shows that phoscorite is composed of apatite, ilmenorutile, perovskite as essential minerals [18]. The enrichments of TiO₂, Nb₂O₅, SiO₂, Y₂O₃ and other REE oxides are caused by presence of silicate perovskite. It is composed of SiO₂ 2.0, Al₂O₃ 1.5, FeO 26.0, MnO 2.0, MgO 0.5, CaO 15.0, Na₂O 0.5, K₂O 0.5, TiO₂ 25.0, P₂O₅ 13.0, F 0.5, SO₃ 1.5, Y₂O₃ 7.0, ZrO₂ 4.0 and Nb₂O₅ 24.0%.

Phoscorites and carbonatites appear to be formed at end stages [27] of magmatic differentiation and fractionation with intermittent liquid immiscibility [1-6, 8, 15, 16, 18 -26]. These nil silicate rocks superimpose one over the other in the field of Si-P-(Fe+Mn) for phoscorite (Fig.4). Large and dense ilmenorutile up to 15x10x8 cm are commonly found as floats around Onnakarai village.

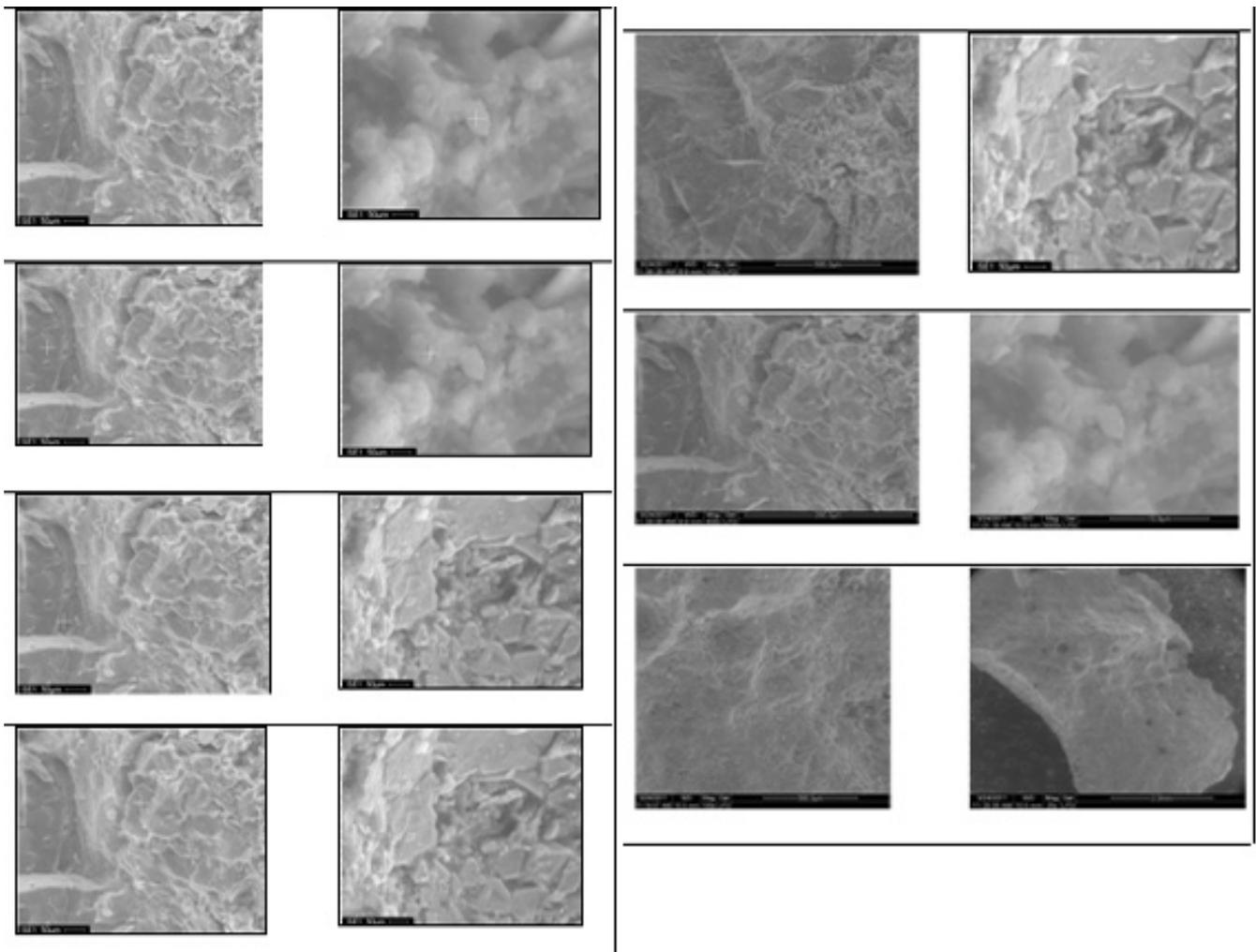


Fig2. The EDAX images of phoscorite show essential minerals of apatite and iron ore.

Table3. Wet-gravimetric analyses of magnetite (498, 13, 21, 47) from phoscorite from Sevvattur and phoscorite 451, apatite 450 and a host rock of peridotite 40 from Onnakarai are given below:

	451	450	40	498	13	21	47
SiO2	42.82	2.06	42.58	0.48	20.36	39.3	62.14
Al2O3	1.27	6.91	4.62	0.73	0.07	0.07	0.17
Fe2O3	21.86	2.89	6.32	54.17	27.6	49.84	33.58
FeO	2.51	2.06	5.25	18.07	37.85	10.36	1.22
MnO	0.04	0.07	0.15	0.49	0.46	0.02	0.01
MgO	0	1.46	10.62	0.73	0.01	0.01	0.01
CaO	14.76	41.36	25.08	6.66	11.16	0.01	1.07
Na2O	1.95	0.47	0.34	0.14	0.71	0.17	0.01
K2O	0.06	0.1	0.06	0.42	0.34	0.33	0.34
TiO2	1.56	10.01	0.93	13.2	1.57	0.14	1.3
P2O5	7.9	31.77	3.1		0.04		0.01
CO2	2.46		0.6				0.39
SO3	1.65		0.33	0.77			
LOI	0.24	0.26	0.6				
Retention's n	451	450	40	498	13	21	47
anh	2.65		0.41	1.57			
cc	7.08		1.44				1.18
ap	18.7	74.52	6.01		0.2		
il	2.53	15.58	1.23	25.92	2.72	0.28	2.1
mt	3.28		0.87		7.01		
oxe		5.67					
spil				69.21	81.89		
sippx						96.38	94.69
cl			45.1				
cpx	35.12		31.64				
ac		4.24			8.17	3.34	1.05
amcr							0.98
pl	7.9		13.3				
nc							
ks				3.3			
qr	22.74						
	100	100	100	100	100	100	100

Table4. Analysis of Magnetite and their structural formula (Magnetite 32 (O, OH, F), apatite 26 (O, OH, F) Phlogopite 22 (O, OH F)

	498	13	21	47	450 Phi	
SiO2	0.48	20.36	39.3	62.14	2.06	32.45
Al2O3	0.73	0.07	0.07	0.17	6.91	15.40
Fe2O3	54.17	27.6	49.84	33.58	2.89	14.86
FeO	18.07	37.85	10.36	1.22	2.06	0.00
MnO	0.49	0.46	0.02	0.01	0.07	0.05
MgO	0.73	0.01	0.01	0.01	1.46	15.85
CaO	6.66	11.16	0.01	1.07	41.36	3.46
Na2O	0.14	0.71	0.17	0.01	0.47	0.40
K2O	0.42	0.34	0.33	0.34	0.1	6.32
TiO2	13.2	1.57	0.14	1.3	10.01	3.37
P2O5	0	0.04	0	0.01	31.77	0.32
CO2	0	0	0	0.39	0	
SO3	0.77	0	0	0	0	
F						1.65
LOI	0	0	0	0	0.26	5.87
	498	13	21	47 Ap	Phi	
Si	0.141	5.459	8.717	11.840	0.353	4.234
Al	0.252	0.022	0.018	0.038	1.398	2.369
Fe3	11.958	5.571	8.322	4.817	0.373	1.459
Fe2	4.431	8.485	1.921	0.194	0.296	0.000
Mn	0.122	0.104	0.004	0.002	0.010	0.005
Mg	0.319	0.004	0.003	0.003	0.373	3.083
Ca	2.092	3.205	0.002	0.218	7.602	0.483
Na	0.080	0.369	0.073	0.004	0.156	0.102
K	0.157	0.116	0.093	0.083	0.022	1.052
Ti	2.911	0.317	0.023	0.186	1.292	0.330
P	0.000	0.009	0.000	0.002	4.614	0.035
CO2	0.000	0.000	0.000	0.101	0.000	
SO3	0.169	0.000	0.000	0.000	0.000	
F						0.682
OH						5.107

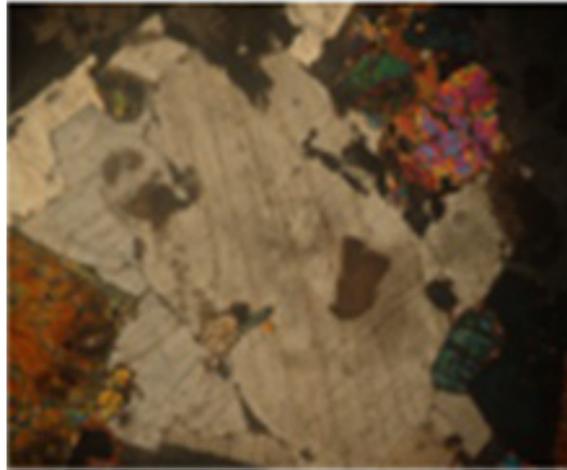


Fig3. 1 Opaque fine dusts in potash feldspars in panidiomorphic shonkinite 2 Lueshire with expansion cracks in albite 3 Garnet bearing pyroxenite 4 Garnet syenite 5 Secondary magnetite in sovite 6 Skarn shows garnet, wollastonite, diopside scapolite and plagioclase.

It is essentially composed apatite 40% and ilmenorutile 60%. The distribution of Si, P and Fe + Mn in [27] an equilateral triangle in Fig. 4 shows that phoscorite is differentiated from carbonatite into apatite – magnetite end. Similar feature is seen in Fig. 5. This may be further illustrated in Fig. 6. Fig. 6 classifies without adding heteromorphic equivalents to olivine. Therefore fields of olivine shrinks hence, olivine rich or olivine poor phoscorites are absent [27]. A linear variation (Fig. 7) is present between K₂O vs CaO/ Al₂O₃. Most rocks the ratios of CaO/Al₂O₃ are limited to <10 (Fig. 7) indicating [27] low to medium degree of partial melting of carbonated peridotite at depth. SiO₂ vs MgO shows a linear positive variation [27] indicating differentiation. Nb₂O₅ + Ta₂O₅ vs ZrO₂ + Hf₂O₃; La₂O₃ vs Sc₂O₃; Ta₂O₅ vs Nb₂O₅ and HfO₂ vs ZrO₂ (Fig. 7) is showed similar variations. Only La₂O₃ vs Y₂O₃ shows negative correlation since Y₂O₃ is much abundant and heavier than La₂O₃.

Table5. Wet-gravimetric analyses of apatite from phoscorite, of Sevvattur

	aps1	aps2	aps3	aps4
SiO ₂	0.05	0.22	0.00	0.00
Al ₂ O ₃	0.18	0.06	0.00	0.00
Fe ₂ O ₃	0.00	0.00	0.00	0.00
FeO	0.00	0.00	0.00	0.00
MnO	0.00	0.00	0.00	0.00
MgO	0.52	0.17	1.51	1.51
CaO	54.25	54.15	55.73	53.28
Na ₂ O	0.30	0.29	0.00	0.00
K ₂ O	0.00	0.00	0.00	0.00
TiO ₂	0.00	0.00	0.00	0.00
P ₂ O ₅	39.80	40.03	40.85	37.96
SrO	0.80	1.51	0.00	0.00
CO ₂	0.00	0.00	0.00	0.00
SO ₃	0.00	0.00	0.00	0.00
F	0.00	0.00	1.51	0.80
2E(O, OH)				
Si	0.009	0.040	0.000	0.000
Al	0.038	0.013	0.000	0.000
Fe ₃	0.000	0.000	0.000	0.000
Fe ₂	0.000	0.000	0.000	0.000
Mn	0.000	0.000	0.000	0.000
Mg	0.140	0.048	0.388	0.415
Ca	10.472	10.425	10.294	10.530
Na	0.105	0.101	0.000	0.000
K	0.000	0.000	0.000	0.000
Ti	0.000	0.000	0.000	0.000
P	6.071	6.090	5.962	5.928
Sr	0.084	0.157	0.000	0.000
CO ₂	0.000	0.000	0.000	0.000
SO ₃	0.000	0.000	0.000	0.000
F	0.000	0.000	0.823	0.467
	aps1	aps2	aps3	aps4
caf2			7.00	4.03
ap	95.66	96.42	91.22	91.30
peric	4.04	2.58	1.78	4.67

Increase of pressure with increasing temperature to the extent of a given density of 8.8 g/cm³ at CMB sudden cracking and creeping develops at lower mantle. Volatiles fill voids initiate widening with developments supercritical fluids. Such supercritical plasmatic bubbles inflate and weaken creeps reduces static friction across their walls. Continental rifting leads direct eruption of natrocarbonatitic supercritical fluids from mantle. These fluids are caused cracking and creeping at mantle for direct effusion of supercritical natrocarbonatitic fluid [32-36]. The Eastern Ghats Paleo Rift system is a continental rift zone [32-35] along which several carbonatites are emplaced in India.

Table6. Wet-gravimetric analyses of Phoscorite from Sevvattur Phoscorite from carbonatite of Sevvattur

	p1	p2	p3	p4
SiO ₂	16.58	9.72	3.78	16.80
Al ₂ O ₃	5.40	2.62	0.74	5.01
Fe ₂ O ₃	0.00	0.00	0.00	0.00
FeO	2.73	2.98	4.68	7.08
MnO	0.00	0.00	0.00	0.00
MgO	1.46	2.36	0.98	5.16
CaO	45.37	58.16	60.10	54.52
Na ₂ O	1.18	1.01	0.58	0.68
K ₂ O	0.87	0.36	0.00	0.71
TiO ₂	0.00	0.00	0.00	0.00
Nb ₂ O ₅	0.00	0.00	0.00	0.00
P ₂ O ₅	26.42	21.35	29.13	8.80
CO ₂	0.00	1.44	0.00	1.24
Total	100.00	100.00	100.00	100.00
Rittmann's norm				
ap	58.59	46.55	65.81	18.74
cc	0.00	3.82		3.17
mt	0.41	0.46	0.78	1.13
ol	20.35	19.28	7.45	30.97
port		21.31	23.26	31.03
or	5.60			
ab	2.48			
an	7.26	1.45		8.66
ne	5.31	5.73	2.70	3.74
ks		1.39		2.55
	100	100	100	100

Apatite in carbonatite show rarely primary fluid inclusion carrying ore surrounded by liquid CO₂, CO₂, brine solutions and water vapour with decrepitating temperature of 750oK. Triple phase inclusions are seen inclusions carrying gypsum brine and water vapour and similar phases occur for halide and sylvite [36]. HPTM appears to be enclosed more than 4 layers of supercritical fluids of natrocarbonatite at mantle horizon. These volatile constituents are associated with alkalis. Globular primary melt inclusions of calcite in clinopyroxene in peridotite and in feldspar in ultrapotassic syenite indicate liquid immiscible separation of carbonatite magma from silicate magma at ambient temperature and pressure at hydrous condition [4].

Table7. EDAX analyses of Phoscorite from Onnakarai.

	AR1	AR2	AR3	AR4
SiO2	1.65	1.56	2.38	2.05
Al2O3	1.54	1.10	1.24	0.00
FeO	14.37	26.23	16.01	9.21
MnO	1.68	2.74	1.69	0.91
MgO	0.67	0.38	0.00	0.00
CaO	1.86	1.72	3.93	15.35
Na2O	1.56	0.37	0.00	0.00
K2O	0.60	0.62	0.39	0.00
TiO2	25.29	36.03	29.00	17.49
P2O5	4.54	3.93	7.79	12.89
CO2	0.00	0.00	0.00	0.00
F	0.48	0.19	0.40	0.00
Cl	0.29	0.24	0.11	0.00
SO3	1.71	1.71	1.68	1.04
Sc2O3	0.79	1.03	0.44	0.22
V2O3	0.14	0.21	0.23	0.00
Cr2O3	0.33	0.00	0.00	0.00
CoO	0.57	0.34	0.20	0.18
NiO	0.32	0.16	0.19	0.17
ZnO	0.28	0.44	0.25	0.41
CuO	0.27	0.00	0.21	0.28
PbO	0.06	0.10	0.13	0.30
Y2O3	4.20	3.60	6.92	9.20
HfO2	0.03	0.02	0.02	0.02
ZrO2	3.92	2.74	3.33	5.08
Ta2O5	0.13	0.15	0.00	0.16
Nb2O5	20.55	13.51	22.94	24.55
La2O3	0.48	0.65	0.38	0.30
Ce2O3	0.32	0.18	0.14	0.00
Nd2O3	0.05	0.07	0.00	0.00
Eu2O3	0.03	0.00	0.00	0.00
Dy2O3	0.03	0.00	0.00	0.00
Yb2O3	0.03	0.00	0.00	0.00
Lu2O3	0.05	0.00	0.00	0.19
zr	5.78	3.44	4.36	5.61
il	30.14	63.17	39.87	23.25
ap	15.43	11.57	13.16	38.33
Nb3P5			6.38	1.07
naf	1.08			
nacl	1.44			
(nak)2SO4	6.68			
MSO4	3.43			
Al2(SO4)3		4.3		
(Alscv)2(SO4)3			6.94	
Y2(SO4)3	1.35	2.74		
Sccrvy	4.33	2.74		4.45
cor	2.26			
Tinbca				27.29
Tinbsi		12.04		
TinbySi			29.3	
TinbRee	28.07			

The southern part of Tamil Nadu, a number of carbonatitic lava exposures are seen with air fall tephra, lava flows, tuff, lapilli, pisolites and ashes [41-44].

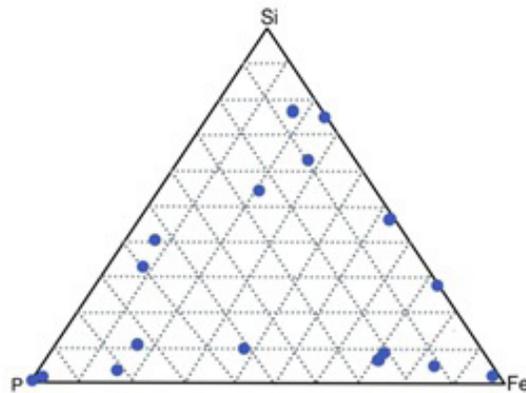


Fig4. Distributions of Si, P and Fe+Mn are seen in equilateral triangle. It includes magnetite 5, apatite 5 and phoscorite 9 and a peridotite 1.

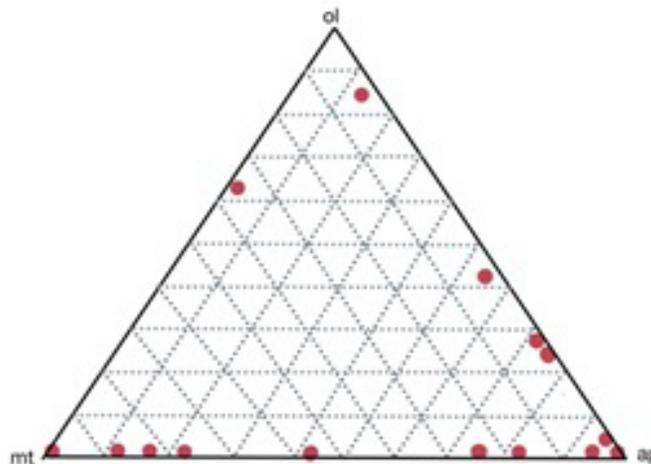


Fig5. Distribution of ol, mt, and ap is shown but ol is not included with its heteromorphic variants.

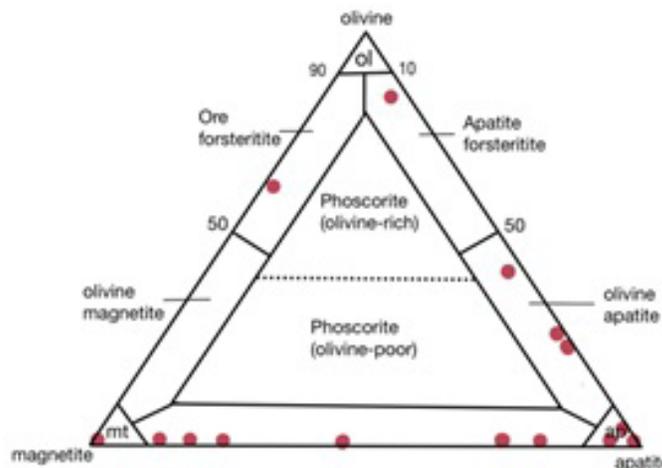


Fig6. Classification of Phoscorite is shown. Heteromorphic components of olivine are not included with olivine, If they were olivine rich or poor phoscorite are identified.

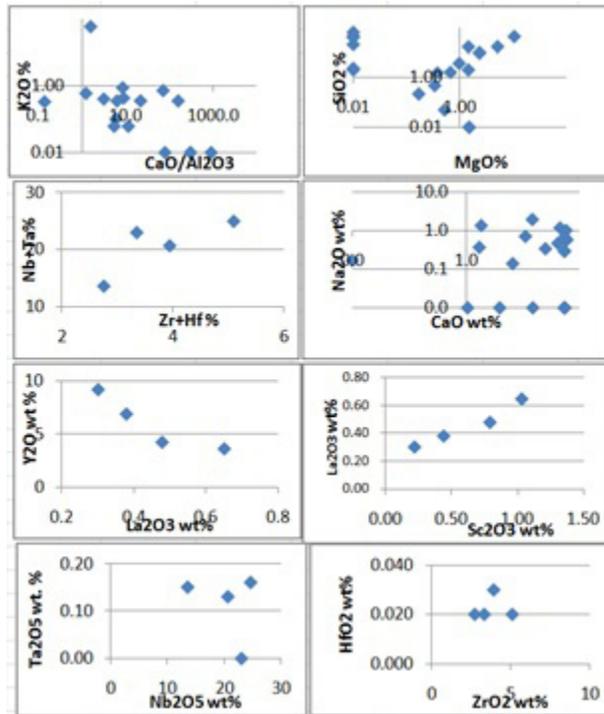


Fig7. Bi-component variations of some characteristic chemical components are shown.

DISCUSION

The petrogenesis of carbonatites and phoscorites at mantle source remain as an unsolved problem. The cause ultralow velocity seismic zone is detected near core - mantle boundary (CMB) is not clearly known [27,]. Recent discovery of presence of primary natrocarbonatite melt inclusions in diamond from mantle source has revised origin of supercritical natrocarbonatite. Occurrence of sodium fluorite, sodium chloride and sodium potassium sulphate (Table 7) indicates that the primary fluid is natrocarbonatite from lowermost mantle. SO₃ is present in syenites and ultramafic rocks up to 1.75 wt%, in this complex. HPTM [28, 29] inclusions enclosed by multiple layers of fluids to maintain its stability. A large quantity of potential/latent energy is consumed or released for the formation / deformation of layers enclosed or around HPTM as that of homogenization in fluid inclusion of multiple phases. Such behaviour preserves the stability of ascend of HPTM. During course of magmatic differentiation volatiles and alkalis are concentrated in the residual magma. The released H₂O, C, O, F, S, P silica and alkalis are consumed by younger crystallizing minerals under alternate miaskitic and apgaitic environment [18, 20, 21].

Studying C-O-H system CH₄, C₂H₆ and other hydrocarbons are possible to generate under HPT conditions [31] at deep mantle. These hydrocarbons and H₂, He, H₂O, N, CF₄, C₄F₈ and SF₆ exist as supercritical fluids above their critical temperatures of 300°C in wide range of pressure conditions from 300 MPa to 300 GPa [32]. Solvent power of supercritical fluid of CH₄ and H₂O increases at this state.

The discovery of natrocarbonatite inclusions in diamond [28-30] is illustrated by presence of ultralow viscous supercritical natrocarbonatite fluid in lowermost mantle near CMB. Small degree of partial melting of dense carbonated peridotite releases [38] primary natrocarbonatite fluids energized with enrichment of elements of low atomic numbers of H, He, C, O, F, Na, Mg, Al, Si, P, S, Cl, K, Ca, Sc, Ba, Ti, Mn. LREE and Fe at HPT state excerpts pressure inducing creeping and cracking. Through such conduit supercritical fluids of natrocarbonatite carrying HPTM enveloped by multi-layered fluids ascends at a speed of from 70 to 100 m/s within 10 hours duration to Earth's surface retains its structural stability [30, 45, 46]. There is not enough time for structural change.

CONCLUSION

The carbonatite complex of Tiruppattur is unique of its differentiated and fractionated co-magmatic sequences of alkaline and carbonatite rocks [1-6]. The ultralow viscous supercritical natrocarbonatite fluid at lower mantle may be developed by low degree of partial melting of carbonated peridotite mantle at depth. HPTM of Si-perovskite, Ca-per-

ovskite, zircon, periclase apatite and ilmenorutile are crystallized at its early stage of development of supercritical fluid [40]. Geophysicists believe that mantle horizon near CMB is highly heterogeneous because of development of ultralow velocity zones for seismic waves due to presence of ultralow viscous primary supercritical natrocarbonatite fluid. The supercritical natrocarbonatite fluid carrying HPTM surrounded by HPT layers ascends very rapidly at a speed of 70-100m/s without any friction and loss of potential energy [39, 45, 46] through suddenly developed creep / conduit in lower mantle. HPTM does not any structural change near surface. The natrocarbonatite fluid is the parent magma for differentiated phoscorite in this carbonatite complex of Tiruppattur.

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