

Geochemistry of Tikak Parbat Sandstones and Tipam Sandstones Occurring in and around Dilli Area, Sivasagar District, Assam, India

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Abstract

This study presents a geochemical investigation of Tikak Parbat and Tipam Sandstone Formations occurring in and around Dilli area, Sivasagar, Assam. Petrographically Tikak Parbat and Tipam sandstones are mainly quartzose arenite to sublitharenite types with their constituents being derived from recycled orogen provenance under sub-humid to humid climatic conditions. Geochemically, sandstones of both the formations range from sublitharenite to wacke. They indicate a recycled orogen source and influence of humid to arid condition. Source rocks of Tikak Parbat sandstones were more weathered than the Tipam sandstones. While Tikak Parbat sandstones show affinity towards passive margin, Tipam sandstones hint at active continental margin setup, where clasts were supplied from uplifted areas. Upliftment of provenance covering areas of Naga Patkai Range in the south east and Eastern Himalayas along the syntaxial bend during mid Miocene affected the sandstones. Tikak Parbat sandstones reflect a stable tectonic setup which later underwent a phase of volatility leading to deposition of the Tipam sandstones. Our study supports a sediment supply from the upper continental crust, largely of granitic composition, however, with a significant variation in their depth of source supply. Trace element analyses indicate depositional setup with low ventilation marked by both oxic and anoxic phases.

Keywords

Geochemistry, Provenance, Palaeoweathering, Tikak Parbat Sandstones, Tipam Sandstones, Dilli (Assam, India)

1. Introduction

Physical examinations of sedimentary rocks and subsequent provenance analyses have been priority research domains in the field of sedimentological studies. Such attempts mostly aim to reconstruct the pre-depositional history of a sediment or sedimentary rock which includes observations on the distance and direction of the provenance, type, size and setting of the source region, climate and relief in the source area, and the specific type of sedimentary rocks [1] [2] [3] [4]. Applications of sophisticated geochemical techniques provide robust support and have further widened the scope of sedimentological studies like predepositional evolution of the source terrain, source rock and depositional environment evaluation, palaeoweathering and redox analyses of sediments etc. [5]-[12]. Geochemical investigations are advantageous as they can be carried out on sedimentary rocks of variable grain size and even in mineralogically altered rocks. Geochemical data are usually sub-divided into four categories: major elements, trace elements, radiogenic isotopes and stable isotopes. The status and trends of these four categories throw some light on the geological background of the rocks.

In the present study an attempt has been made to mainly ascertain some geochemical attributes of two important lithostratigraphic units of the Assam-Arakan basin: Tikak Parbat sandstones and Tipam sandstones exposed in and around Dilli-Jeypore coalfield of Upper Assam. Assam-Arakan tectono-sedimentary basin is situated in the north eastern part of India and it is the largest onshore receptacle of Palaeogene-Neogene rocks in India. The major tectonic elements of the basin include Assam Shelf, Belt of Schuppen and the Assam Arakan fold belt. It may be mentioned that the two units: Tikak Parbat sandstones and Tipam sandstones belong to two different ages-Oligocene and Miocene, respectively and are very important hosts of economic resources like coal and petroleum. The area of interest is a part of the Upper Assam Shelf and is in proximity of the Belt of Schuppen. Further, it may be mentioned that the hinge zone of Assam-Arakan sedimentary basin which is a shelf-slope basinal system lies below the Belt of Schuppen. The major oxides' variations, trace element and rare earth elemental concentrations were utilised to meet the objectives of reconstructing tectonic setting of depositional basin, provenance and, reveal palaeo climatic conditions and palaeoweathering of the concerned rock units of the study area. Petrographic attributes of the investigated samples are also summarized here to give an insight into their mineralogical traits.

2. Geology of the Area

The present study were designed to investigate a part of Dilli-Jeypore coalfield of Upper Assam and located in the foot hills of Naga-Patkai hills along the Dilli river section (**Plate 1**). The area at large is traversed by a linear belt of overthrusts known as "Belt of Schuppen" where Tertiary strata have been folded and dispersed into a number of thrust blocks. The Barail (Oligocene) and Tipam



Plate 1. Panoramic view of the study area along with the coal mine (Lahkar, 2007).

(Miocene) Group of rocks are exposed in the area of study (**Table 1**). The study area is bounded by Lat. 27°04'N - 27°09'N and Long. 95°15'E - 95°22'E (**Figure 1**).

The Tikak Parbat Formation of Barail Group is well exposed in the area mostly in the hills adjoining the alluvial plains. These exposures overlie the Naga Thrust and are characterised by massive to well-bedded, fine to medium grained, yellowish to light grey and grayish white sandstones. Sandstones are characterised by two sets of joints and ripple marks. The massive sandstones are intercalated with shale and coal seams. The Naga thrust which is the north western margin of the Belt of Schuppen separates the Tikak Parbat Formation from the adjoining alluvial plains. Thickness of the Tikak Parbat sandstones range up to 20 mts. in the Dilli river section and strike is largely ENE-WSW to NE-SW dipping 35° to 65° towards SE. The Tipam Sandstone Formation unconformably overlies the Tikak Parbat Formation. The Tipam sandstones are characterised by fine to coarse grained texture. These sandstones which are pebbly and gritty in nature towards the base are also bedded, massive, micaceous, ferrugenous and grey coloured with salt and pepper appearance. Occasionally laminations are observed. The Tipam Formation rocks strike ENE-WSW to NE-SW and dip 35° to 55° towards SE (Plate 2). The regional structure of the Dilli-Jeypore coalfield is that of an anticline (commonly referred to as the Jeypore anticline). Attitude of both the units are also similar striking NE-SW and moderate to highly dipping SE.

3. Methods

Representative samples were collected systematically from Dilli area along traverses at regular intervals and points of lithological change from surface exposures. The exposures were mostly confined to Dilli river section, nallas and "Naga Baat" (regular walking routes of Naga tribes from neighbouring places).

Forty six representative thin sections were considered for detailed petrographic analyses where different quartz types were identified following Basu *et al.* [13]. Around 450 counts were made per section following the point counting technique of Chayes [14] for identifying the sandstone types.

Geochemical analyses of representative samples of Tikak Parbat sandstones and Tipam sandstones from the study area were considered to evaluate their chemical constitution with regard to major oxides and trace elements including rare earth elements. Twenty representative sandstone samples from both the

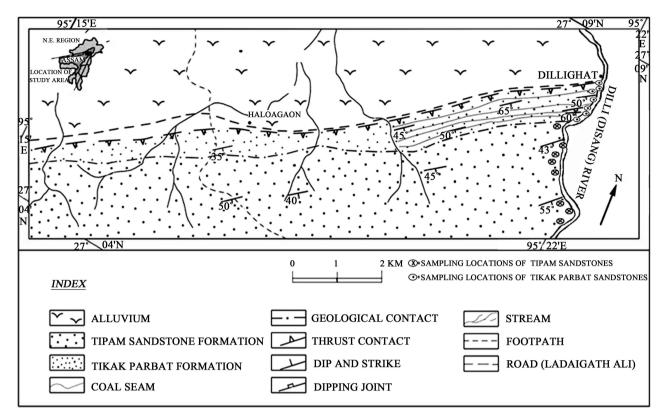


Figure 1. Geological map of the Dilli area, Sivasagar district, Assam (after Lahkar, 2007). The sample locations are marked in the map.

Age	Group	Formation	Lithology
Recent			Alluvial soil cover with terraces
			Hard grey sandstone, siltstone, shale
Miocene	Tipam	Tipam sandstone	
			Coarse to gritty hard sandstones with salt and pepper appearance.
			Unconformity
			Sandstone, carbonaceous shale and coal seams with shale and sandy shale.
Oligocene	Barail	Tikak Parbat	
			Shale, carbonaceous shale, sandy shale, sandstone and thick coal seam.
		Thrust	contact (Naga Thrust)

units were analysed by SIEMENS SRS 3000 XRF in Wadia Institute of Himalayan Geology, Dehradun, India for major oxides. The samples were further analysed for trace elements including rare earth elements by ICP-MS (Parkin Elmer DRC II) in National Geophysical Research Institute, Hyderabad, India.



Plate 2. (a) Ripple marks of Tikak Parbat sandstones; (b) Gritty nature of Tipam sandstones (Lahkar, 2007).

4. Petrographic Attributes of the Sandstones

Detailed petrographic analyses and the data reflected in Table 2 and Table 3 to understand the volumetric abundance of different constituents are from samples used in geochemical analyses. Table 3 is derived out of Table 2 and it presents a comparative statement of dominant petrographic constituents of the Tikak Parbat sandstones and Tipam sandstones of Dilli area. Overall, both Tikak Parbat and Tipam sandstones are mainly quartzose arenite to sublitharenite types [15].

Quartz in both the sandstones comprise of monocrystalline (unit and undulose) and polycrystalline (composite, schistose and pressure) varieties, while the feldspars are plagioclase, microcline and orthoclase. Rock fragments of metamorphic, igneous and sedimentary origin as well as protomatrix and epimatrix are also seen. Cement is commonly calcareous, ferrugenous and siliceous. Muscovite, biotite and chlorite reflect the micaceous components. A few secondary cherts are observed along with the detrital ones (**Plate 3**).

Monocrystalline quartz indicates their descent from intrusive igneous rocks while the composite, schistose and pressure units reflect their derivation from a terrain hosting medium to high grade metamorphic and intrusive rocks. Petrographic association of monocrystalline quartz, plagioclase and potash feldspar along with mica suggest derivation of the sediments from an uplifted crystalline basement of granitic to granodioritic composition and extensive low to high grade metasedimentary terrains [16]. Both opaque and non-opaque varieties of heavy minerals are observed. Tournaline, zircon, rutile, epidote, garnet, staurolite, kyanite, sillimanite and rare hornblende represent the non-opaques while iron oxide and ilmenite represent the opaques. Tectonic discrimination plots after Dickinson [2], Dickinson and Suczek [17] and Dickinson et al. [18] show the derivation of sediments from recycled orogen provenance. Climate plots after Suttner and Dutta [19] show concentration of plots mainly in the sub-humid to humid sector. Petrographic composition and heavy mineral suite of both the sandstone units along with the regional geological setting suggest their derivation from metamorphic, igneous and subordinate sedimentary source rocks transported mostly from Eastern Himalaya and Indo-Burma Range.

Table 2. Modal analyses data of the Tikak Parbat sandstones and Tipam sandstones of Dilli area (Comp. = Composite; Pres. =Pressure; Schist. = Schistose; Feld. = Feldspar; Cem. = Cement; Acc. Min. = Accessory Minerals).

						TIK	AK PA	RBAT SA	NDST	ONES						
SN	Sample	M	onocrystalli	ne Qua	rtz	Pol	ycrysta	lline Qua	artz	Total	D.1.1	Rock	Com	Madain	Miss	Acc.
21N	No.	Unit	Undulose	Vein	QMT	Comp.	Pres.	Schist.	QPT	Quartz	Feld.	Frag.	Cem.	Matrix	Mica	Min.
1	D9	16.66	25.12	-	41.78	11.28	7.17	14.1	32.55	74.37	1.53	7.17	8.71	5.12	2.3	0.76
2	D12	10.24	18.78	-	29.02	9.75	5.6	11.21	26.56	55.58	2.19	10.73	14.39	12.43	4.14	0.48
3	D16	16.04	22.55	1.39	39.98	7.9	6.51	14.65	29.06	69.04	1.39	8.6	10.69	7.9	2.09	0.23
4	D19	14.21	20.24	-	34.45	12.53	5.3	19.03	36.86	71.31	1.2	6.26	9.15	8.19	3.37	0.23
5	BP5	20.5	17.21	1.26	38.97	10.37	5.31	15.18	30.86	69.83	1.26	10.37	9.11	7.34	1.51	0.5
6	BP9	14.14	17.31	-	31.45	10.48	8.29	20.24	39.01	70.46	1.21	10.24	10	6.34	1.21	0.48
7	NB2	12.47	14.11	2.11	28.69	10.82	5.17	17.64	33.63	62.32	2.82	9.64	10.35	11.29	2.35	1.17
8	NB5	14.66	20.80	-	35.46	12.53	4.26	16.26	33.05	68.51	3.73	9.33	7.2	5.6	5.06	0.53
9	N4(1)	14.87	16.41	-	31.28	10.76	4.1	16.15	31.01	62.29	1.79	11.53	12.30	8.2	3.58	0.25
10	N5(2)	17.9	16.04	1.39	35.33	12.32	3.25	19.3	34.87	70.2	1.16	7.31	9.06	8.83	2.09	1.39
							TIPAN	4 SANDS	STONE	5						
SN	Sample	M	onocrystalli	ne Qua	rtz	Pol	ycrysta	lline Qua	artz	Total	Feld.	Rock	Cem.	Matrix	Mico	Acc.
214	No.	Unit	Undulose	Vein	QMT	Comp.	Pres.	Schist.	QPT	Quartz	reiu.	Frag.	Cem.	Watitx	Iviica	Min.
1	D25	10.54	10.81	-	21.35	18.37	2.43	12.16	32.96	54.31	8.64	10.54	14.32	7.29	4.32	0.54
2	D30	11.53	10.51	-	22.04	14.61	1.79	12.3	28.7	50.74	8.2	15.12	12.05	10.25	3.59	-
3	D34	12.47	10.11	-	22.58	12.7	1.17	14.58	28.45	51.03	7.52	10.11	15.05	9.17	5.41	1.64
4	D41	12.8	9.33	1.06	23.19	11.46	1.6	12.26	25.32	48.51	10.4	12.8	12.53	11.2	4	0.53
5	BP14	15.11	8.6	1.16	24.87	10.23	1.39	11.86	23.48	48.35	10	11.16	14.41	11.62	4.41	-
6	BP19	18.29	9.26	1.21	28.76	12.19	0.97	10.48	23.64	52.4	5.6	15.85	9.75	12.92	3.41	-
7	NB9	14.38	10.11	1.12	25.68	10.33	O.89	10.56	21.78	47.39	9.43	10.33	14.6	12.8	4.26	1.34
, 8	NB14	17.9	10.11	-	27.9	10.35	1.39	10.50	22.54	50.44	8.83	12.55	9.06	14.41	3.72	0.69
																0.09
9	B5	12.65	8.86	2.02	23.53	11.39	2.27	9.11	22.77	46.3	4.45	16.45	12.15	14.93	5.56	-
10	B12	18.29	9.75	-	28.04	15.6	1.21	8.53	25.34	53.38	8.29	10	15.36	9.51	3.41	-

 Table 3. Volumetric abundance range and comparative statement of dominant petrographic constituents of the Tikak Parbat sandstones and Tipam sandstones of Dilli area.

Constituents (in %)	Tikak Parbat sandstone	Tipam sandstone
Quartz	55.58 to 74.37	46.3 to 54.31
Feldspar	1.16 to 3.73	4.54 to 10.4
Rock fragments	7.17 to 11.53	10 to 16.45
Matrix	5.12 to 12.43	7.29 to 14.93
Cement	7.2 to 14.39	9.06 to 15.36
Mica	1.21 to 4.14	3.41 to 5.56



Plate 3. Photomicrograph (a) showing replacement of plagioclase by quartz in Tikak Parbat sandstones $(40\times)$ and (b) showing calcareous cement in Tipam sandstones $(40\times)$.

5. Geochemistry Findings and Discussion

5.1. Major Oxides

The abundance of major oxides and some commonly considered major oxide ratios of both Tikak Parbat sandstones and Tipam sandstones of the Dilli area are listed in **Table 4** and **Table 5** respectively. In terms of mean element abundance the values reflect the mineralogical character of the sandstones. Variations in major element geochemistry vide Harker diagrams were analysed in two groups-1) in relation to SiO₂ and, 2) variations between other radicals.

The Tikak Parbat sandstones have high SiO_2 concentration of 63.69 to 92.75 wt% with an average of 83.38 wt%, while the Tipam sandstones have moderate SiO_2 concentration of 51.55 to 69.33 wt% with an average of 59.74 wt%. It is interesting to note that SiO_2 has an average abundance of 66% for sandstones derived from upper continental crust [20]. In the present case, the average SiO_2 of Tikak Parbat sandstones is much higher at 83.38% while that of the Tipam sandstones, it is around 59.74% which means that sediment contributions were from deeper levels for the Tipam Sandstones.

In case of the Tikak Parbat sandstones, considering SiO_2 as a common factor, it is seen that Al_2O_3 , Fe_2O_3 , K_2O , Na_2O , MnO, CaO, TiO_2 and MgO show negative correlation with it indicating free silica being sequestered in the form of quartz. However, Tipam sandstones correlations between SiO_2 with Al_2O_3 , K_2O and Na_2O show positive relation while other radicals show a negative relation. There is a good amount of aluminosilicate represented by feldspars, mica and of course, the clays probably in case of the Tipam sandstones (**Table 3**).

The average Al_2O_3 content of Tikak Parbat sandstones is 7.16 wt%, while it is 9.78 wt% for the Tipam sandstones. The presence of more matrix component in the Tipam sandstones than the Tikak Parbat sandstones supports the higher content of Al_2O_3 in the latter. It might be due to alteration of Kfeldspar. Correlation between alumina and oxides of K_2O and Fe_2O_3 of both the sandstones shows negative nature suggesting their non association with phyllosilicates.

K₂O/Na₂O average ratios hovers around unity in Tipam sandstones and is attributed to the presence of nearly equivalent proportions of albitic plagioclase, K-feldspar, mica and illite [21]. Proportion of K₂O and Na₂O in the Tikak Parbat sandstones is relatively less than the Tipam sandstones suggesting an environ-

Major					SA	MPLE N	os.				
Oxides	D9	D12	D16	D19	BP5	BP9	NB2	NB5	N4 (1)	N5 (2)	Avg
SiO ₂	91.63	63.69	87.15	89.52	86.7	85.7	74.62	92.75	76.05	86.06	83.3
Al_2O_3	5.03	15.05	6.08	4.55	5.18	5.13	12.07	3.56	11.75	3.23	7.16
Fe ₂ O ₃	0.52	4.25	1.06	0.65	1.61	1	3.02	0.54	2.84	4.5	1.99
CaO	0.04	2.78	0.23	0.14	0.15	0.21	0.28	0.02	0.29	0.26	0.44
MgO	0.41	2.28	0.65	0.64	0.72	0.88	1.91	0.57	1.93	1.53	1.15
K ₂ O	0.53	1.16	0.92	0.6	0.72	0.66	1.36	0.45	1.54	0.41	0.83
Na ₂ O	0.52	1.72	0.32	0.26	0.31	0.28	2.09	0.25	2.35	0.18	0.82
TiO ₂	0.47	0.63	0.53	0.56	0.49	0.66	0.65	0.58	0.49	0.67	0.57
MnO	0.016	0.083	0.02	0.022	0.027	0.024	0.048	0.019	0.042	0.08	0.038
P_2O_5	0.033	0.112	0.061	0.052	0.052	0.058	0.077	0.047	0.078	0.052	0.062
LOI	1.41	7.41	1.77	1.7	1.96	1.85	3.64	1.72	3	3.85	2.83
Total	100.6	99.16	98.8	98.7	97.92	96.45	99.76	100.5	100.36	100.82	99.3
CIA	82.19	68.94	83.52	81.40	81.45	78.92	73.82	80.91	72.00	62.12	74.82
CIW	89.98	76.98	91.70	91.92	91.84	91.28	83.59	92.95	81.65	88.01	85.04
PIW	73.53	67.07	68.34	71.17	70.13	71.18	67.78	72.66	64.09	69.12	68.43
Major Oxide Ratio											
$Fe_2O_3 + MgO$	0.93	6.53	1.71	1.29	2.33	1.88	4.93	1.11	4.77	6.03	
Al ₂ O ₃ /SiO ₂	0.05	0.23	0.06	0.05	0.05	0.05	0.16	0.03	0.15	0.03	
K ₂ O/Na ₂ O	1.01	0.67	2.87	2.3	2.32	2.35	0.65	1.8	0.65	2.27	
$Al_2O_3/(CaO + Na_2O)$	8.98	3.34	11.05	11.37	5.95	4	2.36	4.5	2.26	0.69	
log (Na ₂ O/K ₂ O)	-0.0083	0.171	-0.458	-0.368	-0.365	0.372	0.186	-0.255	0.183	-0.357	
$log(Fe_2O_3/K_2O)$	-0.0083	0.563	0.061	0.034	0.349	0.18	0.346	0.079	0.265	1.04	
log(SiO ₂ /Al ₂ O ₃)	1.26	0.626	1.156	1.293	1.233	1.222	0.791	1.415	0.811	1.425	
Al ₂ O ₃ /Na ₂ O	9.67	8.75	19	17.5	16.7	18.32	5.77	14.24	5	17.94	
TiO ₂ /Al ₂ O ₃	0.09	0.04	0.08	0.12	0.09	0.12	0.05	0.16	0.04	0.02	
MnO/Fe ₂ O ₃	0.03	0.01	0.01	0.03	0.01	0.02	0.01	0.03	0.01	0.01	
$Al_2O_3 + K_2O + Na_2O$	6.08	17.93	7.32	5.41	6.21	6.07	15.52	4.26	15.64	3.82	
SiO ₂ /Al ₂ O ₃	18.21	4.23	14.33	19.67	16.73	16.7	6.18	26.05	6.47	26.64	

Table 4. Major oxides (in wt%) and major oxide ratios of the Tikak Parbat sandstones of Dilli area.

 Table 5. Major oxides (in wt%) and major oxide ratios of the Tipam sandstones of Dilli area.

Major		SAMPLE Nos.												
Oxides	D25	D30	D34	D41	BP14	BP19	NB9	NB14	B5	B12	Avg.			
SiO ₂	54.14	65.47	52.49	63.54	62.24	69.33	59.81	57.43	51.55	61.39	59.74			
Al_2O_3	9.02	13.58	8.38	9.35	9.14	12.41	8	8.4	9.57	9.99	9.78			
Fe ₂ O ₃	2.36	5.98	2.18	3.84	2.44	3.97	2.95	2.1	2.77	4.15	3.27			
CaO	16.16	1.33	16.29	8.39	8.61	0.86	12.23	14.66	17.01	4.21	9.97			

Continued											
MgO	2.02	3.96	1.92	2.37	2.8	3.08	2.56	1.79	2.23	3.55	2.02
K ₂ O	1.67	1.89	1.56	1.63	1.8	2.1	1.71	1.63	1.65	1.73	1.74
Na ₂ O	1.29	2.04	1.23	1.45	2.01	2.07	1.61	1.46	1.2	1.62	1.75
TiO ₂	0.38	0.69	0.38	0.62	0.25	0.54	0.36	0.34	0.38	0.71	0.46
MnO	0.601	0.069	0.699	0.24	0.518	0.059	0.494	0.663	0.61	0.153	0.41
P_2O_5	0.15	0.144	0.137	0.105	0.088	0.234	0.074	0.121	0.16	0.181	0.139
LOI	11.9	4.02	13.09	6.82	8.17	3.52	8.56	12.49	11.74	14.53	9.48
Total	99.69	99.17	98.36	98.35	98.07	98.18	98.36	101.08	98.87	102.21	99.23
CIA	32.05	72.08	30.52	44.91	42.39	71.16	33.97	32.12	32.52	56.92	42.08
CIW	34.08	80.12	32.36	48.72	46.26	80.90	36.63	34.26	34.45	63.15	45.49
PIW	26.12	62.05	24.84	37.08	34.04	59.12	26.71	25.89	26.91	47.07	34.60
Major Oxide Ratio											
$Fe_2O_3 + MgO$	4.38	9.94	4.1	6.21	5.24	7.05	5.51	3.89	5	7.7	
Al ₂ O ₃ /SiO ₂	0.16	0.2	0.15	0.14	0.14	0.17	0.13	0.14	0.18	0.16	
K ₂ O/Na ₂ O	1.29	0.92	1.26	1.12	0.89	1.01	1.06	0.11	1.37	1.06	
$Al_2O_3/(CaO + Na_2O)$	0.51	4.02	0.47	0.95	0.86	4.23	0.57	0.52	0.52	1.71	
log (Na ₂ O/K ₂ O)	-0.112	0.033	-0.103	-0.05	0.047	-0.0063	-0.026	-0.047	-0.138	-0.028	
$log(Fe_2O_3/K_2O)$	0.15	0.5	0.145	0.372	0.132	0.276	0.236	0.11	0.224	0.38	
$\log(SiO_2/Al_2O_3)$	0.778	0.683	0.769	0.832	0.833	0.747	0.873	0.834	0.731	0.788	
Al ₂ O ₃ /Na ₂ O	6.99	6.65	6.81	6.44	4.54	5.99	4.96	5.75	7.97	6.16	
TiO ₂ /Al ₂ O ₃	0.04	0.05	0.04	0.06	0.02	0.04	0.04	0.04	0.03	0.07	
MnO/Fe ₂ O ₃	0.25	0.01	0.32	0.06	0.21	0.01	0.16	0.31	0.22	0.03	
$Al_2O_3 + K_2O + Na_2O$	11.98	17.51	11.17	12.43	12.95	16.58	11.32	11.49	12.42	13.34	
SiO ₂ /Al ₂ O ₃	6	4.82	6.26	6.79	6.8	5.58	7.47	6.83	5.38	6.14	

ment detrimental for survival of feldspars. The age factor is also responsible for this compositional status. The low values of K_2O/Al_2O_3 (0.116 for Tikak Parbat and 0.178 for Tipam sandstones) probably suggest sedimentary recycling or increase in the degree of source area weathering.

Titanite (TiO_2) is persistently low in abundance. TiO_2 concentration is found to occur between 0.47 to 0.67 wt% for Tikak Parbat sandstones and 0.25 to 0.71 wt% for the Tipam sandstones. Titanium is characteristic of sediments that have undergone long period of consistent sub-aerial weathering. It is a stable mineral phase and might have come from metamorphosed argillaceous rocks or acid igneous rocks as detrital rutile or, from pre-existing sedimentary rocks and acid igneous rocks as detrital ilmenite rather than a soluble fraction. It is indicative of continental geochemical setting and, its content decreases with increasing distance from land [22].

The low average value of $\rm Al_2O_3/SiO_2$ for the sandstones (0.03 - 0.16 for the Ti-

kak Parbat and 0.13 - 0.18 for the Tipam sandstones) is also a hint of quartz enrichment [23]. Contrary to that is the ratio of SiO_2/Al_2O_3 for sandstones (4.23 - 26.05 for the Tikak Parbat and 4.82 - 7.47 for the Tipam Sandstones). Our analyses suggest the chemical maturity of the sandstones of Tikak Parbat Formation as relatively higher and it is also reflected by the TiO_2/Al_2O_3 ratio [24] and $SiO_2/Al_2O_3 + K_2O + Na_2O$ plot (**Figure 2**) of Suttner and Dutta [19]. While Tikak Parbat Sandstones reflect the influence of both arid and humid climatic conditions, Tipam sandstones seem to have been deposited under arid conditions.

5.2. Palaeoweathering

Alteration of minerals due to chemical weathering mainly depends on the mineral chemistry, intensity and the duration of weathering as well as climate. One of the dominant processes during weathering in the upper crust is the degradation of feldspars and formation of clay minerals. The degradation of feldspar, which is very sensitive to chemical weathering, increases the mobility of many elements like K, Na, Ca [20] [25] only to reside in the clay minerals and soil profile. Alteration of rocks during weathering results in depletion of alkalis and alkaline earth elements and preferential enrichment of Al_2O_3 , [26]. The amount of these chemical elements surviving in the soil profiles and in the sediments derived from them is a sensitive index of the intensity of weathering [27]. The influence of weathering on sedimentary rocks can be calculated by using Chemical Index of Alteration (CIA) value [28]. The CIA value which indicates the degree of weathering of source rocks is determined by the equation:

$$CIA = \left[Al_2O_3 / (Al_2O_3 + CaO sil + Na_2O + K_2O) \right] \times 100$$

The CIA gives a measure of the ratio of original/primary minerals and

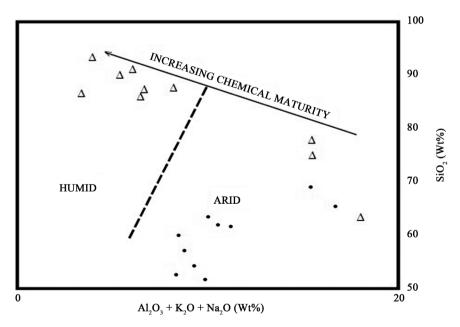


Figure 2. Scatter plot of $Al_2O_3 + K_2O + Na_2O$ vs. SiO_2 of Tikak Parbat sandstones (Δ) and Tipam sandstones (•), after Suttner and Dutta, 1986.

secondary products such as clay minerals. CIA values ranges from almost 50 in case of fresh rocks to 100 for completely weathered rocks, consisting entirely of secondary minerals. High CIA and PIA values (75 - 100) indicate intensive weathering in the source area whereas low values (≤ 60) indicate low weathering in the source area. The CIA is a good measure of palaeoweathering conditions and it essentially monitors the progressive weathering of feldspars to clay minerals [7]. The present study shows that K-feldspar is less in Tikak Parbat sandstones compared to Tipam sandstones. Hence CIA value of Tikak Parbat sandstones is high (62.12 to 83.52; ave.: 74.82) indicating high intensity of chemical weathering in the source areas and lower CIA value (32.52 to 72.08; ave.: 42.08) of Tipam sandstones is an indication of low rate of chemical weathering in the source areas.

Apart from the Chemical Index of Alteration (CIA), Chemical Index of Weathering (CIW) put forward by Harnois [29] and Plagioclase Index of Alteration (PIA) proposed by Fedo *et al.* [30] are also good reflectors of the extent of weathering at source. PIA and CIA can be worked out by the following equations:

$$CIW = 100 [Al_2O_3 / (Al_2O_3 + CaO + Na_2O)]$$
$$PIA = 100 [(Al_2O_3 - K_2O) / (Al_2O_3 + CaO + Na_2O - K_2O)]$$

The CIW concept is similar to CIA except that it eliminates K_2O from the equation. CIW does not account for Al_2O_3 associated with K-feldspar. PIA on the other hand may be used as an alternative to CIW as because plagioclase is abundant in silicate rocks and dissolves rapidly. PIA is generally used when plagioclase weathering needs to be monitored.

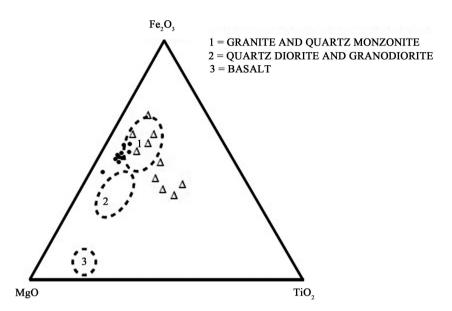
CIW and PIA patterns in the present case are seen to support the CIA trend. CIW value of Tikak Parbat sandstone varies from 76.98% to 92.95% (ave.: 85.04%) while the same for Tipam sandstones range from 32.36% to 80.90% (ave.: 45.49%). Similarly, PIA value of Tikak Parbat sandstone varies from 64.09% to 73.53% (ave.: 68.43%) while the same for Tipam sandstones vary from 24.84% to 62.05% (ave.: 34.6%). CIA, PIA and CIW values greater than 60% as shown by Tikak Parbat sandstones indicate moderate to high weathering either at the source or during transport before deposition [30].

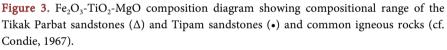
5.3. Provenance

Source rock composition is commonly thought to be the dominant factor that controls the composition of sediments [20]. Secondary processes like weathering and diagenesis can also variably impact the chemical composition [31] and as such, one can best rely on elements that show little mobility under geologic conditions. However, composition of clastic rocks like sandstone and shale is also influenced by the nature of sedimentary processes within the depositional basin, the kind of dispersal paths that link provenance to the basin and plate tectonics [17].

Condie [32] proposed a ternary diagram of Fe₂O₃-MgO-TiO₂ for determina-

tion of the source rock of the sediments which in the present case reflects a granite-granodiorite-quartz monzonite provenance for the Tikak Parbat sandstones and granite-granodiorite provenance for the Tipam sandstones (**Figure 3**). The CaO-Na₂O-K₂O ternary diagram plots after Le Maitre [33] largely shows a granitic provenance for the Tikak Parbat sandstones and andesitic to granitic sources for the Tipam sandstones (**Figure 4**).





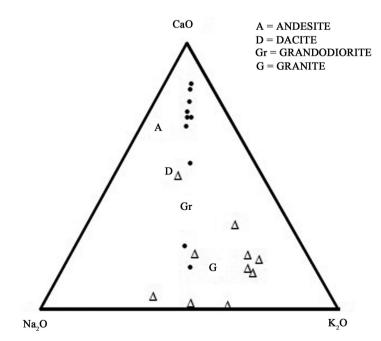


Figure 4. CaO-Na₂O-K₂O plot of Tikak Parbat sandstones (Δ) and Tipam sandstones (•), (cf. Bhatia, 1983), (Also shown are the average of Andesite (A), Dacite (D), Grandodiorite (Gr), Granite (G) (after LeMaitre, 1976).

Association of Tipam sandstone unconformably overlying Barail Group of rocks has been encountered in the Noa Dihing upstream of Miao in Arunachal Pradesh [34]. The same association has also been encountered in Upper Brahmaputra plains in Upper Assam [35]. The presence of coniferous pollen Podocarpidites in the Tipam sandstones also point towards a Himalayan source for the sediments which is also indicated by the heavy mineral assemblage [36]. Abor Volcanic Formation which is considered coeval with lower Permian Panjal Volcanic Formation of N.W. Himalayas describes mafic volcanic constituents of Eastern Himalayas in Arunachal Pradesh. Studies of Talukdar and Mazumdar [37] shows Abor Volcanic suite to comprise of basalt, andesite, acidic tuffs, lapillis and agglomerates. Reflectance of granodioritic and andesitic provenance in some provenance analyses may be due to contribution of clasts from the Abor Volcanics as well as granodiorites of Lohit Complex [38]. On a regional scale these sediments have been derived from metamorphic, igneous and subordinate sedimentary rocks transported mainly from the Eastern Himalayas and the Indo-Burman Ranges during Oligocene to Miocene time [39]. While Tikak Parbat sandstones were deposited under deltaic to near shore framework, Tipam sandstones were laid down under fluviatile regime [40].

5.4. Tectonic Setting

Plate tectonic settings of provenance and depositional basins leave huge impressions on various attributes of siliciclastic sedimentary rocks. Bhatia [23] and Roser and Korsch [6] used composition as a tool to decipher the tectonic setting of sediment accumulation. However, to infer tectonic setting of provenance of ancient siliciclastic sedimentary rocks, several major-trace-rare earth element based discrimination diagrams have been proposed. According to McLennan *et al.* [41] the tectonic setting discrimination diagrams can provide reliable results for siliciclastic rocks that have not been strongly affected by post-depositional weathering/metasomatism/metamorphism. In the present case, tectonic setting discrimination, plots suggested by Blatt *et al.* [1], Bhatia [23] and Roser and Korsch [6] were used.

Bhatia [23] recognized four types of settings namely Oceanic Island Arc (OIA), Continental Island Arc (CIA), Active Continental Margin (ACM) and Passive Margin (PM). The parameters that Bhatia considered to unearth the tectonic setup of deposition were mainly based on major oxide ratios comprising of elements of maximum and minimum mobility and, crustal and oceanic affinity. The plots in the binary diagrams for Tikak Parbat sandstones are close to ACM and PM Fields while those of Tipam sandstones are mostly close to CIA and ACM (**Figures 5(a)-(d)**). In the binary discrimination diagram of Roser and Korsch [6] the Tikak Parbat Sandstones plots mostly in the PM while Tipam Sandstones, plots are mainly seen falling in ACM field (**Figure 6**). Sandstones of PM type are derived from stable continental areas while those of ACM type are derived from uplifted areas [6]. Rifted continental margin, intracratonic and rift

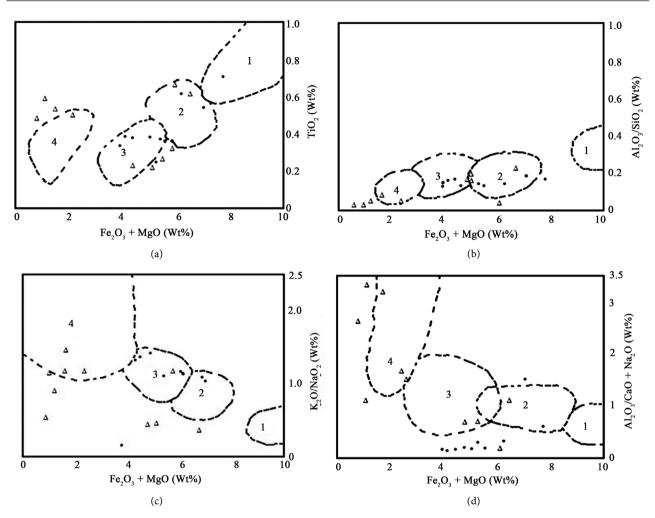


Figure 5. Scatter plot of: (a) $Fe_2O_3 + MgO$ vs. TiO_2 ; (b) $Fe_2O_3 + MgO$ vs. $Al_2O_3 + SiO_2$; (c) $Fe_2O_3 + MgO$ vs. $K_2O + Na_2O$, and (d) $Fe_2O_3 + MgO$ vs. $Al_2O_3/CaO + Na_2O$ of Tikak Parbat sandstones (Δ) and Tipam sandstones (•). (1 = Field of Oceanic Island Arc, 2 = Continental Island Arc, 3 = Active Continental Margin, 4 = Passive Margin, cf., Bhatia 1983).

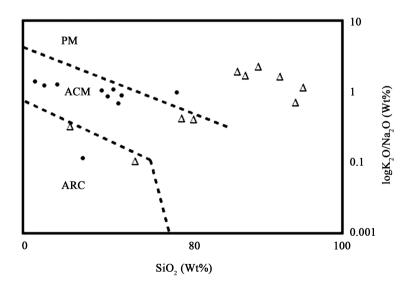


Figure 6. Satter plot of SiO₂ vs. log (K_2O/Na_2O) of Tikak Parbat sandstones (Δ) and Tipam sandstones (•) (after Roser and Korsch, 1986).

bounded basins are generally reflected by unstable mineralogy. Tectono-province plots based on petrographic investigations of the present sandstones point towards their derivation form a recycled provenance [15]. Low Al_2O_3/SiO_2 ratio of sandstones is further characteristic of quartz source deposited in a Passive Continental Margin Setting [42]. However, overlapping plots between passive and active continental margin settings reflect a complex provenance and history of recycling of these sediments [23].

5.5. Geochemical Classification of Sandstones

Geochemical classification of terrigenous sedimentary rocks has been proposed by many authors based on major element composition [1] [43] [44]. Using the indices of SiO_2/Al_2O_3 and Na_2O/K_2O ratios Pettijohn *et al.* (1972) in Herron [44] proposed a classification for terrigenous sands based on a plot of log (Na₂O/ K_2O) versus log (SiO₂/Al₂O₃). Herron [44] modified the diagram of Pettijohn *et al.* (1972) using log (Fe₂O₃/K₂O) instead of (Na₂O/K₂O).

The plots after Herron [44] for Tikak Parbat sandstones fall mostly in "sublitharenite" to "wacke" field. The plots for Tipam sandstones fall mostly within "wacke" field (**Figure 7**). However, petrographically both the sandstones have been identified as quartzose arenite and sublitharenite types. This difference in petrographic and geochemical nomenclature arises from the fact that finer populations are ignored in modal analyses petrographically. These finer fragments are an integral part in whole rock analyses and they contribute Al₂O₃. This shifts the distribution mostly to "sublitharenite" and "wacke" field.

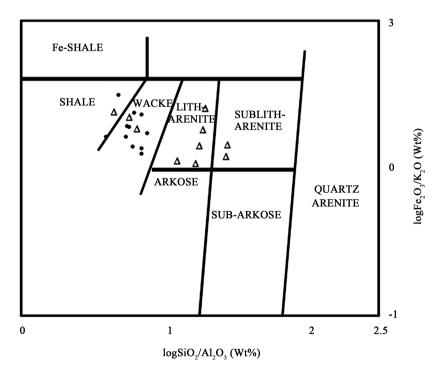


Figure 7. Geochemical classification for terrigenous sands (after Herron, 1988) of the Tikak Parbat sandstones (Δ) and Tipam sandstones (•).

5.6. Trace Elements

A few trace elements and trace elemental ratios help ascertain geochemical attributes of sandstones. Trace element concentrations in sediments result from the competing influences of the provenance, weathering, diagenesis, sediment sorting and the aqueous geochemistry of the individual elements. These may also be controlled by the concentration of heavy minerals. As such, trace element geochemistry facilitate in distinguishing source rock composition and depositional environment [43] [45]. They can also be used to deduce the palaeotectonic setting of sedimentary packages [5] [46]. Low mobility of the trace elements during sedimentary processes and their low residence time in sea water makes them suitable for provenance and tectonic setting determinations of clastic sediments. These elements are in fact transported quantitatively into clastic sediments or sedimentary rocks during weathering and transportation and as such they carry the impressions of parent materials [5] [47].

Distribution of the trace elements for the Tikak Parbat sandstones (**Table 6**) and Tipam sandstones (**Table 7**) does not show much difference from the UCC [48]. There are some exceptions too. Copper and Zirconium are less in abundance relative to UCC. The presence of each trace element suggests some background information about the sandstone. For example, sufficient amount of Co, Cr and Ni in the sandstones suggests that these elements were derived from ferromagnesian minerals of metamorphic rocks. Geochemical analyses of Abor volcanic [49] and serpentinites of Tidding Formation of Lohit Complex [50] of the Eastern Himalayas which are presumed to be source rock(s) also report the presence of elements like Cr, Co and Ni. Similarly Pb is also derived from source rocks which contributed Fe to the sediments.

The Ba:Al₂O₃ relationship of Tikak Parbat sandstones is strongly positive ($R^2 = 0.700$) and the same for Tipam sandstones is weakly positive ($R^2 = 0.402$). Similarly, the Rb:Al₂O₃ relationship of Tikak Parbat sandstones is strongly positive ($R^2 = 0.860$) and the same for Tipam sandstones is of lower much value ($R^2 = 0.229$) compared to Tikak Parbat sandstones. These correlations suggest that phyllosilicate minerals have more distinct control on the distribution of Ba and Rb in case of the Tikak Parbat sandstones compared to the Tipams. A similar finding comes out in case of the relationship Ba:K₂O which is otherwise positive. In case of Tikak Parbat sandstones R^2 is 0.961 while it is 0.058 in case of the Tipams. Feldspar is a major host of Ba and Rb in terrigenous sedimentary rocks. Feldspars and Illites had more control in the distribution of Ba in Tikak Parbat sandstones. Analogous findings have come up from the ration Ti:Nb as well.

The Sr:Al₂O₃ relationship is positive in Tikak Parbat sandstones ($R^2 = 0.59$) while it is negative in Tipam sandstones ($R^2 = -0.509$). The Sr content of sedimentary rocks is variable because of many influences on Sr in low temperature depositional environments [51]. Sr for example can be affected by the presence of Ca. Additional Sr can be incorporated in diagenetic carbonate, and fractionation of Sr can result from the weathering of feldspars, particularly plagioclase. In

Trace		SAMPLE Nos.												
elements	D9	D12	D16	D19	BP5	BP9	NB2	NB5	N4(1)	N5(2)				
Sc	4.66	13.6	4.7	4.34	4.68	4.04	9.19	4.03	7.73	8.46				
V	34.64	106.38	81.73	93.6	72.67	70.36	87	27.7	77.38	74.73				
Cr	64.04	81.95	25.03	30.63	26.81	21.53	130.97	32.14	102.05	26.72				
Со	117.22	45.12	85.81	123.9	124.44	82.41	43.4	133.5	54.38	98.55				
Ni	20.1	71.45	17.84	24.41	13.65	13.59	84.84	13.37	85.78	15.94				
Cu	13.88	21.2	11.87	13.23	11.71	12.67	14.25	12.71	15.37	10.31				
Zn	50.04	39.76	26.03	35.55	22.08	27.01	41.63	27.81	29.42	24.762				
Ga	5.31	13.19	5.98	5.09	5.17	4.34	11.79	3.75	11.26	4.03				
Rb	23.39	45.69	28.87	23.56	26.17	21.12	51.78	18.07	55.73	17.93				
Sr	176.33	381.97	152.48	142.54	164.25	136.94	191.37	186.44	191.25	147.18				
Y	12.4	27.75	18.23	16.52	16.6	15.61	21.3	18.68	17.64	24.15				
Zr	123.9	207.34	118.37	115.46	113.05	133.06	188.11	243.27	163.65	133.93				
Nb	6.6	8.26	7.22	7.49	7.08	8.21	9.39	7.575	7.09	8.82				
Cs	1.08	3.24	1.22	0.95	1.03	0.84	2.72	0.75	2.63	0.73				
Ba	151.28	196.61	146.82	133.72	135.81	120.59	248.7	117.71	282.73	120.11				
Hf	3.49	6.79	3.45	3.14	3.35	3.9	5.62	7.25	5.02	4.21				
Та	1.3	0.79	1	1.15	1.08	1.02	0.82	1.44	0.82	1.18				
РЬ	17.33	15.47	11.04	10.71	11.28	10.17	14.8	15.67	13.03	10.38				
Th	3.41	8.49	5.16	5.97	4.95	5.24	7.39	7.25	6.08	8.69				
U	2.81	5.07	8.22	8.75	15.84	15	21.15	0	43.15	49.55				
Trace element ratios														
Cu/Zn	0.27	0.53	0.45	0.37	0.53	0.46	0.34	0.45	0.52	0.41				
Sc/Cr	0.07	0.16	0.18	0.14	0.17	0.18	0.07	0.12	0.07	0.31				
V/Cr	0.54	1.29	3.26	3.05	2.71	3.26	0.66	0.86	0.75	2.79				
Ni/Co	0.17	1.58	0.2	0.19	0.1	0.16	1.95	0.1	1.57	0.16				
Cr/Th	18.79	9.64	4.84	5.13	5.41	4.11	17.71	4.43	16.79	3.07				
Th/Sc	0.73	0.06	1.03	1.37	1.05	1.29	0.8	4.02	0.78	1.02				
Th/Co	0.02	0.18	0.06	0.04	0.03	0.06	0.17	0.05	0.11	0.08				
Th/U	1.21	1.67	0.63	0.68	0.31	0.35	0.35	0.00	0.14	0.18				
Zr/Y	9.99	7.47	6.49	6.99	6.81	8.52	8.83	13.02	9.28	5.55				
Zr/Hf	35.50	30.54	34.31	36.77	33.75	34.12	33.47	33.55	32.60	31.81				
Cr/Ni	3.19	1.15	1.40	1.25	1.96	1.58	1.54	2.40	1.19	1.68				

Table 6. Trace element concentration (in ppm) and some trace element ratios of the Tikak Parbat sandstones of Dilli area.

Table 7. Trace element concentration (in ppm) and some trace element ratios of the Tipam
sandstones of Dilli area.

Trace	SAMPLE Nos.												
elements	D25	D30	D34	D41	BP14	BP19	NB9	NB14	B5	B12			
Sc	7.79	11.33	8.18	8.81	6.9	9.26	7.08	7.68	8.33	10.2			
V	79.55	118.28	58.94	70.92	48.54	68.46	47.82	51.09	51.19	72.6			
Cr	49.29	91.98	56.86	58.05	59.3	58.49	79.74	49.78	52.17	76.6			
Со	46.29	55.47	60.24	53.39	59.38	42.88	38.63	41.08	3485	38.1			
Ni	37.64	49.1	36.11	30.38	37.4	48.3	31.91	35.05	37.81	42.7			
Cu	11.31	13.12	9.41	9.97	10.81	12.1	10.49	9.86	10.47	12.9			
Zn	27.98	32.5	25.79	31.95	21.52	30.26	24.98	23.45	26.71	55.2			
Ga	11.25	15.21	10.29	11.87	10.65	14.29	9.53	10.49	11.06	12.6			
Rb	61.52	71.99	56.11	59.66	6.49	79.89	53.34	58.26	59.47	62.2			
Sr	227.92	181.33	223.3	178.1	255.13	145.39	229.9	226.94	219.62	177.4			
Y	23.54	30.36	22.85	32.02	25.44	23.81	27.29	21.77	24.35	33.1			
Zr	79.34	357.84	69.47	86.52	58.71	105.5	61.53	95.96	90.72	101.			
Nb	8.12	11.44	8.49	12.23	5.13	9.06	7.63	7.51	8.42	12.1			
Cs	2.79	3.04	2.6	2.28	1.99	3.72	1.57	2.66	2.76	2.7			
Ba	311.07	414.72	294.96	321.5	425.19	423.4	371.77	301.97	308.66	353.			
Hf	2.84	3.8	2.72	3.29	2.11	3.17	2.13	3.43	3.38	3.7			
Та	0.86	1.22	1.1	1.03	0.78	0.9	0.82	0.7	0.82	1.1			
Pb	10.34	11.88	11.94	13	17.29	19.59	14.55	12.7	12.25	15.5			
Th	8.06	16.2	9.24	11.53	7.65	9.78	11.56	7.96	8.97	15.0			
U	49.45	66.92	66.74	28.09	66.07	25.26	69.92	56.42	65.45	72.1			
Trace element ratios													
Cu/Zn	0.4	0.4	0.36	0.31	0.5	0.4	0.42	0.42	0.39	0.2			
Sc/Cr	0.15	0.12	0.14	0.15	0.11	0.15	0.08	0.15	0.15	0.1			
V/Cr	1.61	1.28	1.03	1.22	0.81	1.17	0.59	1.02	0.98	0.9			
Ni/Co	0.81	0.88	0.59	0.56	0.62	1.12	0.82	0.85	1.08	1.1			
Cr/Th	6.11	5.67	6.15	5.03	7.74	5.97	6.89	6.24	5.81	5.0			
Th/Sc	1.03	1.42	1.12	1.3	1.1	1.05	1.63	1.03	1.07	1.4			
Th/Co	0.17	0.29	0.15	0.21	0.12	0.22	0.29	0.19	0.25	0.3			
Th/U	0.16	0.24	0.14	0.41	0.12	0.39	0.17	0.14	0.14	0.2			
Zr/Y	3.37	11.79	3.04	2.70	2.31	4.43	2.25	4.41	3.73	3.0			
Zr/Hf	27.94	94.17	25.54	26.30	27.82	33.28	28.89	27.98	26.84	27.4			
Cr/Ni	1.31	1.87	1.57	1.91	1.59	1.21	2.50	1.42	1.38	1.7			

the present case, Al_2O_3 content in Tikak Parbat sandstones is more and CaO is very less. It means that in this distribution, Sr content is controlled by the feldspars. In case of the Tipam sandstones, CaO is more. Here distribution of Sr is controlled more by the carbonates. Admittance in place of calcium is a dominant process of removal of strontium from magma. Calcite is common cement (**Plate 3(b**)) in the Tipam Sandstones [52].

Th, U and Th/U ratio each of them reveal significant information about arenaceous distribution. Concentration of Th for example may be an estimate of the degree of weathering in sedimentary rocks. Both Th and U are relatively immobile during weathering. However, in reworked sediments under aerobic conditions U may be removed from the system thereby increasing the Th/U ratio and the value may be higher than upper crustal igneous values. The Th/U ratio in most upper continental rocks typically ranges between 3.5 and 4.0 (McLennan *et al.*, 1993). As such, in sedimentary rocks, Th/U values higher than 4.0 may indicate intense weathering in the source areas or sedimentary recycling. In the present case, low Th/U ratio hint at low intensity weathering. Relatively, weathering was more intense in Tikak Parbat sandstones. Low Th/U ratios on the other hand are commonly seen in active margin settings where rapid accumulation and burial of sediments can occur.

Positive correlations have been obtained for TiO_2 with Y, Zr and Nb. In case of Tikak Parbat sandstones $R^2 = 0.4$ for Y, 0.109 for Zr and 0.843 for Nb. For Tipam sandstones $R^2 = 0.618$ for Y, 0.344 for Zr and 0.907 for Nb. These patterns suggest that the abundance of these components were controlled by the detrital heavy mineral fraction in case of Tipam sandstones while there were some other contributors for Tikak Parbat sandstones.

Vanadium often occurs in organic matter and is best preserved under low pH and high reducing conditions [53]. Organic matter derived from marine planktons is known to concentrate vanadium. The high vanadium level in most of the samples of Tikak Parbat sandstone and Tipam sandstones suggests organic derivation too. Dypiv [54] opines that the V/Cr ratios of sediments are indicators of the ventilation conditions in the depositional environment. The values for the Tikak Parbat and Tipam sandstones indicate low ventilation conditions in the depositional basin which resulted in poor to moderate sorting of these sediments. The V/Cr ratio of Tikak Parbat sandstone varies from <2 to >2 indicating occurrence of anoxic and oxic depositional condition. In Tipam sandstones the ratio is <2 which indicates an oxic depositional condition [55]. The Ni/Co ratio helps interpret Eh conditions in the depositional basin which may have been a mixed environment in the present case. Similar observations were noticed in Disang sandstones of Nagaland [56]. The low values of Cu/Zn ratio suggest increased oxidising condition.

The plots of Zr versus TiO_2 after Hayashi *et al.* [57], show that the source rocks of both the sandstones are felsic igneous rocks (**Figure 8**). According to Bhatia and Crook [5] the triangular plots of the elements Th-Sc-Zr/10 show

excellent discrimination of tectonic settings. In the triangular diagram of Th-Sc-Zr/10 for Tikak Parbat Sandstone most of the plots concentrate in PM to CIA while the Tipam Sandstone plots are close to CIA and ACM (**Figure 9**).

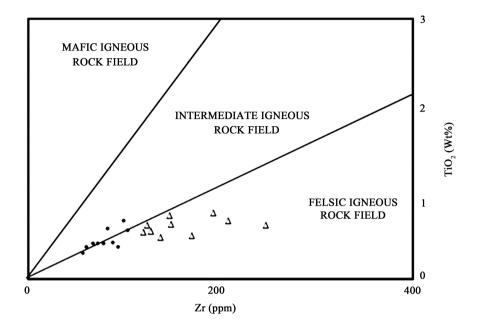
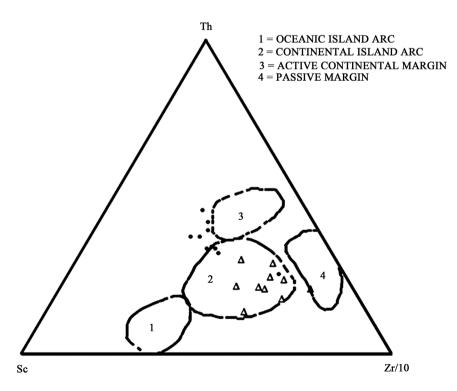
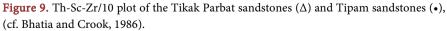


Figure 8. Zr (ppm) vs TiO₂ (wt%) plot of the Tikak Parbat sandstones (Δ) and Tipam sandstones (•) showing Mafic, Intermediate and Felsic Igneous rock fields after Hayashi *et al.*, 1997.





5.7. Rare Earth Elements

Rare earth elements (REE) behave as an unusually coherent group of elements and are amongst the least soluble elements that are relatively immobile during weathering, low-grade metamorphism and hydrothermal alteration. The rare earth elements have very low residence time in sea water and their concentration in the ocean is exceedingly low. Most common sedimentary processes do not significantly affect the rare earth element distribution in sedimentary rocks [58]. Rare earth elemental distribution in the upper continental crust is widely assumed to be similar to that of clastic sediments [59]. The abundance of REE and some elemental ratios of both Tikak Parbat sandstones and Tipam sandstones of the study area are listed in **Table 8** and **Table 9**, respectively.

The Tikak Parbat sandstones show a large variation in REE content (66.02 to 174.04 ppm) while the Tipam sandstones show less variation in REE content

Table 8. Rare earth element concentration (in ppm) and some REE ratios of the TikakParbat sandstones of Dilli area.

Rare Earth	SAMPLE Nos.												
Elements (REE)	D9	D12	D16	D19	BP5	BP9	NB2	NB5	N4(1)	N5(2)			
La	12.61	22.93	19.84	23.58	18.5	20.08	21.39	29.81	16.68	32.31			
Ce	26.92	47.27	45.58	53.58	40.76	46.62	43.61	65.72	35.38	72.11			
Pr	3.16	5.85	5.54	6.73	4.99	5.55	5.45	8.1	4.33	8.87			
Nd	12.01	22.08	21.47	25.29	18.83	21.02	20.48	30.4	15.98	34.13			
Sm	2.47	4.67	4.71	5.27	4.06	4.51	4.35	6.06	3.38	7.1			
Eu	0.67	1.13	1.12	1.18	0.95	0.99	1.09	1.27	0.89	1.42			
Gd	2.32	4.46	4.14	4.43	3.73	4.03	4.02	5.33	3.2	6.08			
Tb	0.42	0.83	0.71	0.7	0.63	0.64	0.75	0.84	0.62	0.97			
Dy	2.42	5.16	3.66	3.48	3.34	3.35	4.27	3.98	3.48	5.04			
Но	0.44	0.94	0.6	0.57	0.59	0.54	0.71	0.67	0.63	0.85			
Er	1.19	2.68	1.64	1.57	1.56	1.49	2.12	1.85	1.87	2.45			
Tm	0.18	0.43	0.22	0.21	0.27	0.21	0.34	0.25	0.29	0.36			
Yb	1.05	2.59	1.29	1.18	1.28	1.22	2	1.51	1.71	2.06			
Lu	0.16	0.38	0.19	0.18	0.19	0.18	0.29	0.22	0.26	0.29			
REE ratios													
ΣREE	66.02	121.4	110.71	127.95	99.68	110.43	110.87	156.01	88.7	174.04			
ΣLREE	57.84	103.93	98.26	115.63	88.09	98.77	96.37	141.36	76.64	155.94			
ΣHREE	8.18	17.47	12.45	12.32	11.59	11.66	14.5	14.65	12.06	18.1			
ΣLREE/ΣHREE	7.07	5.95	7.89	9.39	7.60	8.47	6.65	9.65	6.35	8.62			
Eu/Eu*	0.09	0.08	0.08	0.08	0.08	0.08	0.08	0.07	0.09	0.07			
La/Yb	12.01	8.85	15.38	19.98	14.45	16.46	10.70	19.74	9.75	15.68			
La/Y	1.01	0.82	1.08	1.42	1.11	1.28	1	1.59	0.94	1.33			

Rare Earth					SAMPL	E Nos.						
Elements (REE)	D25	D30	D34	D41	BP14	BP19	NB9	NB14	B5	B12		
La	22.14	38.84	26.41	35.18	22.83	25.14	39.84	23.13	26.91	35.27		
Ce	44.92	77.21	53.21	71.35	47.24	51.84	77.26	45.94	55.24	71.49		
Pr	5.36	8.87	6.31	8.47	5	6.09	8.11	5.49	6.57	8.29		
Nd	19.39	31.22	23.19	30.43	18.04	22.4	27.08	19.63	23.82	29.71		
Sm	3.96	6.13	4.59	6.09	3.57	4.47	4.76	3.92	4.67	5.86		
Eu	0.96	1.33	1.07	1.38	1.11	1.16	1.13	0.98	1.15	1.35		
Gd	3.96	5.76	4.21	5.78	3.94	4.33	4.96	3.83	4.69	5.73		
Tb	0.71	1.01	0.76	1.07	0.74	0.79	0.84	0.7	0.79	0.99		
Dy	4.22	5.69	4.09	6.03	4.23	4.47	4.74	3.9	4.57	5.88		
Но	0.79	1.07	0.76	1.11	0.83	0.85	0.89	0.72	0.85	1.13		
Er	2.26	3.26	2.2	3.18	2.45	2.44	2.64	2.15	2.41	3.22		
Tm	0.35	0.49	0.34	0.48	0.38	0.38	0.39	0.32	0.36	0.52		
Yb	2.01	3.08	1.94	2.83	2.26	2.22	2.42	1.92	1.99	2.96		
Lu	0.31	0.43	0.31	0.39	0.33	0.34	0.36	0.28	0.31	0.45		
REE ratios												
ΣREE	111.34	184.39	129.39	173.77	112.95	126.92	175.42	112.91	134.33	172.85		
ΣLREE	96.73	163.6	114.78	152.9	97.79	111.1	158.18	99.09	118.36	151.97		
ΣHREE	14.61	20.79	14.61	20.87	15.16	15.82	17.24	13.82	15.97	20.88		
ΣLREE/ΣHREE	6.62	7.87	7.86	7.33	6.45	7.02	9.18	7.17	7.41	7.28		
Eu/Eu*	0.08	0.07	0.08	0.08	0.1	0.09	0.08	0.08	0.08	0.08		
La/Yb	11.01	12.61	13.61	12.43	10.10	11.32	16.46	12.05	13.52	11.92		
La/Y	0.94	1.27	1.15	1.09	0.89	1.05	1.45	1.06	1.1	1.06		

Table 9. Rare earth element concentration (in ppm) and some REE ratios of the TipamSandstones of Dilli area.

(111.34 to 184.39 ppm). A strong assumption on the sorting processes during sedimentation is that fine grained sediments tend to have the relative abundance of REE as found in their source region [45] [60]. Sand fraction tends to have low REE content.

The Σ LREE/ Σ HREE ratio of Tikak Parbat sandstones varies from 5.95 to 9.65 with an average of 7.76 and that of Tipam sandstone varies from 6.45 to 9.18 with an average of 7.42 which are almost similar to that of the UCC. The high ratio of Σ LREE/ Σ HREE in both the sandstones suggest source rock area to be granitic [61].

The chondrite normalized REE patterns of the Tikak Parbat sandstone (Figure 10(a)) and the Tipam sandstone (Figure 10(b)) are similar in nature and bears resemblance with the UCC for most rare earth elements. The chondrite normalized REE distribution pattern of the sandstones exhibits a mod-

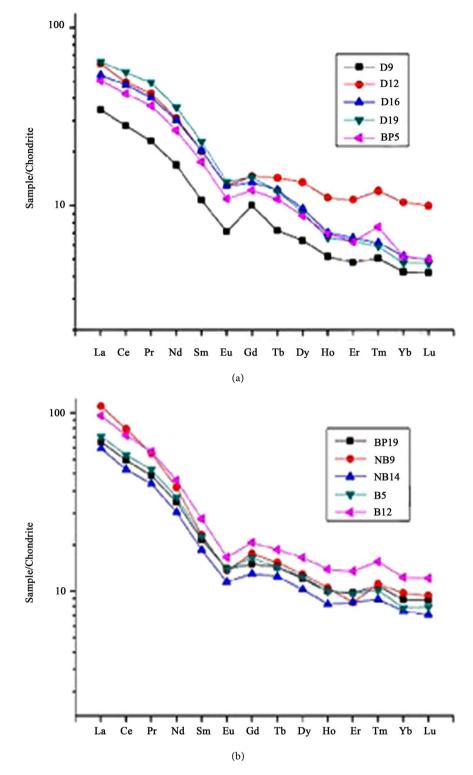


Figure 10. REE distribution pattern of: (a) Tikak Parbat sandstone; and (b) Tipam sandstone.

erately steep slope, which suggests enrichment in LREE and depletion of HREE. These patterns also show negative Eu anomaly. Intracrustal fractionation and separation of plagioclase impart the negative Eu anomaly in the upper crust [20].

Such chondrite normalized REE distribution patterns and negative Eu anomaly indicate their derivation from granitic and granodioritic type plutonic source rocks [20] [59] [62]. The negative Eu anomalies in the sediments indicate preferential removal of plagioclase feldspar due to weathering. Plagioclase is known to be more rapidly destroyed than either quartz or K-feldspar in weathering profiles [63].

Variable La/Yb ratio of the sandstones indicates differing proportions of felsic rocks in the source. Bhatia [64] worked out the discriminating rare earth elements characteristics of different sedimentary basins based on REE content and La/Yb ratio. The average La/Yb ratio of Tikak Parbat sandstones is 14.3 which is very close to passive margin (15.9) tectonic setting. The average La/Yb ratio of Tipam sandstones is 12.5 which is same as active continental margin (12.5) tectonic setting. The La-Th-Sc ternary discriminating plots after Bhatia and Crook (1986) which otherwise is a very useful indicator of tectonic environment shows plots of both the sandstones spread out in the ACM and PM fields. However, a few plots of Tikak Parbat Sandstones fall in the CIA field as well (**Figure 11**).

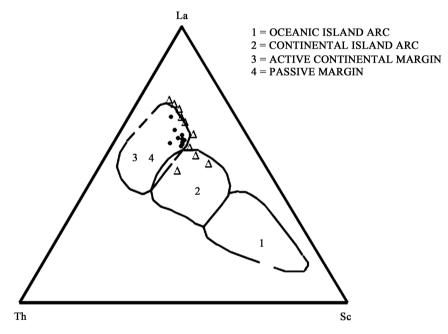


Figure 11. La-Th-Sc plot of the Tikak Parbat Sandstones (Δ) and Tipam Sandstones (•) (cf. Bhatia and Crook, 1986).

6. Conclusions

The detailed geochemical studies conducted in the present study provide a new understanding about the evolution of the Tikak Parbat and Tipam sandstones of Sivasagar district of Assam. The salient findings from our study stem out along the following concluding observations.

1. Analysis of the major oxides indicates the older Tikak Parbat sandstones to be chemically more matured than the Tipam sandstones, a fact also reflected from petrographic investigations. Sediments of both the sandstones were derived from upper continental crust and had undergone low levels of recycling. Relatively, sediment contributions were from deeper levels for the Tipam sandstones which indicate tectonic rejuvenation at later stages. Climatic conditions were largely humid during the deposition of Tikak Parbat sandstones, while it was arid during the deposition of Tipam sandstones. Variable palaeoclimatic signatures as reflected in petrographic and geochemical plots are mainly due to different parameters involved and framework of the discriminatory plots.

- 2. Tikak Parbat sandstones are found to be more weathered than the Tipam sandstones. The weathering impressions on the distributions have been mod-ified during the transportation stage.
- 3. Both the sandstones have been largely derived from granitic (felsic) rocks. Andesitic to granitic sources also contributed towards buildup of Tipam sandstones.
- 4. Tectonically, Tikak Parbat sandstones are found to have more affinity towards passive margin (PM) setup wherein sediments were mostly contributed from stable continental areas whereas Tipam sandstones express a strong likeness towards active continental margin (ACM) setup to where clasts were supplied from uplifted areas.
- 5. Both Tikak Parbat and Tipam sandstones are classified as sublitharenite to wacke from geochemical point of view. Finer populations in both the distributions contributed Al₂O₃ which led to the concentration of plots in the ""sublitharenite" and "wacke" fields.
- 6. Trace element composition of both the distributions does not differ much from that of the upper continental crust (UCC) and hint at felsic igneous source rocks. Phyllosilicates and feldspars had more control on the distribution of elements like Ba, Rb and Ni in case of the Tikak Parbat sandstones than the Tipam sandstones. CaO controlled the distribution of Sr in case of the Tipam sandstones. Depositional setup had low ventilation and the temporal span was marked by both oxic and anoxic settings.
- 7. The ΣLREE/ΣHREE ratio and chondrite normalized REE patterns of both the sandstones are almost similar to that of the UCC and suggest compositionally granitic type of source rocks. This study shows the LREE as the more enriched entity, while HREE is found to be depleted. A negative Eu anomaly further hints at sediment derivation from upper crust. Affinity of Tikak Parbat sandstones to PM and Tipam sandstones to ACM is also supported by REE.
- 8. The Tipam sediments were transported over a high gradient topography and rapidly deposited in the area. The sediments were affected by the tectonic upliftment of provenance covering the areas of Naga Patkai Range in the south east and Eastern Himalayas along the syntaxial bend during mid Miocene. Contribution of Mishimi Hills composed mainly of granites and granodioritic rocks in the Tipam sandstone is probably less considering the lower popu-

lation of Zr in the Tipam unit. Comparatively the older Tikak Parbat sandstones reflect a stable tectonic setup which later on underwent a phase of volatility leading to the deposition of the Tipam sandstone.

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