

Provenance, tectonic setting and geochemical maturity of the Early Miocene Pyawbwe Formation, Sakangyi –Thayet Area, Pyay Sub-Basin, Myanmar

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Abstract

The best exposed Early Miocene (820m.thick.) shales and interbedded silty sandstones beds of the Pyawbwe Formation at Sakangyi-Thayet area are investigated geochemically by using Siemens SRS- X Ray 303 AS XRF Spectrometer. Major and some trace element concentrations have been determined to achieve their provenance, tectonic setting paleoweathering, paleoclimate and sedimentation characteristics. The geochemistry of sediments is particularly valuable in the study of fine-grained rocks that are difficult to characterize through petrographic studies. Geochemical data revealed that the felsic granitic plutonic provenance of moderate relief of arc massif exposed on tectonically calm continental margin which is probably the Shan- Thai continental block and northeastern Myanmar. Average values of both Chemical Index of weathering (CIW) and Chemical Index of Alteration (CIA) (77.4 and 67.3 respectively) suggest a moderately chemical weathering condition prevailed during transportation and deposition on passive shallow margin as progressive mature sediments.

Keywords: *Pyawbwe Formation, Early Miocene, Central Myanmar Basin, Passive Margin.*

Introduction

A little studies have been carried out done on the sedimentation and sediment provenance in Central Myanmar Basin. Still the sedimentary supply to the Central Myanmar Basin is a matter of debate (Naing *et al.*, 2013, Licht *et al.*, 2013, Robinson *et al.*, 2014). The economic importance of the Miocene formations as reservoir rocks for the gas and oil in the Central Myanmar (Burman) Basin attracted the attentions of the several recent geologists (Wandrey, 2006, Harun *et al.*, 2014 and others), then the Pyawbwe Formation is chosen to this study. This study focuses on the provenance of the Pyawbwe Formation sediments of the Pyay embayment, Sakangyi – Thayet area within the central Myanmar Belt. The area under study is commonly located in the Thayet Saddle and northern margin of the Pyay Sub - Basin (Figs. 1 & 2). It is structurally complex trending NNW – SSE with many folds and fault systems (Fig. 3b).

It is majorly fallen in the Sakangyi anticline to north of the Thayet, bounded by Tokkaing syncline in the west and Thayet thrust fault to the east. This structure is cored by the Pyawbwe Formation and surrounded by middle Miocene to Pliocene sediments (Fig. 1). The aim of the present study is to understand and document the evolution of the sedimentary supply

and also to construct a pictorial sedimentation scenario of the Pyawbwe Formation for the first time. The present authors combine the litho- geochemical results to evaluate its source and tectonic depositional setting. The characterization of the provenance and tectonic setting of these units is important for understanding the geodynamic of a part of the Central Myanmar Basin of the western margin of the Shan Plateau (Fig. 2) during the Early Miocene.

Stratigraphically, the Pyawbwe Formation lies between the Okhmintaung Formation (Upper Oligocene) at lower and the Kyaukkok Formation (Middle Miocene) at upper level (Fig. 3C). In general, all the rocks belonging to the Oligocene – Miocene ages are mainly built up conglomerate, sandstone, mud, siltstone, shale and claystones. Lithologically, the Pyawbwe Formation consists of blue grey shales and clay members intercalated with few fine-grained sandstones and carbonaceous siltstones sediments (Fig. 4)

Regional Geological Setting

Located at the southern Range of the eastern edge of Himalayan Chain, the Central Myanmar Basin (CMB) is located between Shan Plateau to the east and the Indo-Burman Ranges to west which are composed of sedimentary, metasediments, intensive and volcanic rocks.

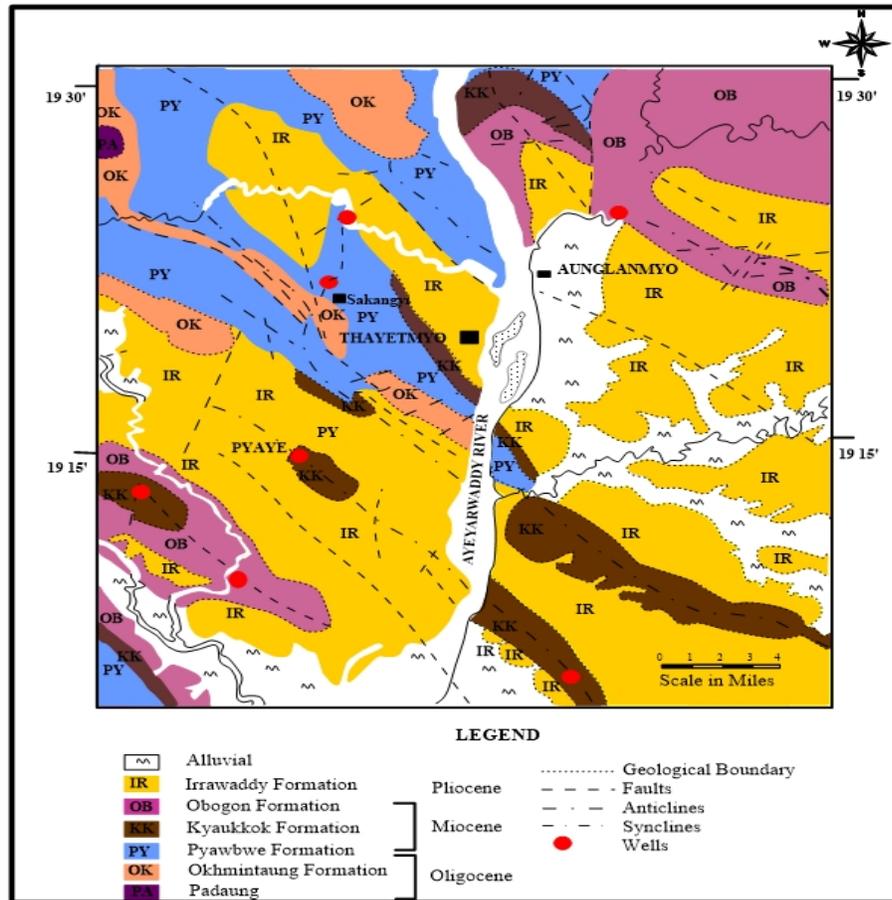


Figure 1. General geological map showing distribution of the Pyawbwe Formation around the Central Myanmar Basin, Pyay, Thayet, Sakangayi areas: (after Moc, 1985)

The Central Cenozoic Belt or Central Myanmar Basin is divided into several Tertiary Sub-Basins (Fig.2) along its nearly 1100 km. length. The sub-basins have been almost filled since the Indo-Asian collision (Bender, 1983, Bertrand & Rangin, 2003, Searl *et al.*, 2007, Allen *et al.*, 2008, Licht *et al.*, 2014). These sub-basins may have formed as a series of en echelon pull-apart basins (Fig. 2) trending approximately NW-SE with about 50 km. wide in the Early Eocene (Fig. 3D) as the Burma Plate moved northward relative to the Asia Plate (Pivink *et al.*, 1998, Rangin *et al.*, 1999). A 15 km. thick succession of Cenozoic deposits was found in Central Myanmar Belt (Pivink *et al.*, 1998).

Materials and Methods

A total of 70 shale samples (Plate, 1) were collected mostly with an equal interval of space of 20 m. all along these sections (820 Thick.). Only 16 representative samples of visibly fresh shales and interbeds sandstones representative samples of

visibly fresh shales and interbedded sandstones were analyzed for their major and some trace element concentrations. They were examined with Shimadzu Model ED-720 energy dispersive XRF system with Standard Curves based on International Rock Standards at the Institute of Electron Optics, University Research Center (URC), Yangon. The obtained results as well as the calculated indices are tabulated in Table (1).

Results and Discussions

The obtained results of the major and some trace elements for both shales and the intercalated silty-sandstone beds of the Early Miocene Pyawbwe Formation are reflected by narrow range of variations.

Geochemistry

Major Elements

The range and averages of the major chemical concentrations and ratios are tabulated in Table (1).

Table 1. Major and some trace element concentrations (wt %) of the Pyawbwe Formation, Myanmar.

| Elements | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | Average |
|--|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|---------|
| SiO ₂ | 25.16 | 37.41 | 37.39 | 19.73 | 38.52 | 27.83 | 37.48 | 36.03 | 37.3 | 35.46 | 36.91 | 42.05 | 52.72 | 22.25 | 39.08 | 37.67 | 35.2 |
| Al ₂ O ₃ | 8.71 | 33.16 | 33.59 | 10.35 | 33.62 | 15.43 | 33.89 | 32.32 | 33.97 | 13.43 | 34.06 | 30.45 | 15.73 | 10.33 | 31.24 | 22.48 | 46 |
| Ti O ₂ | 0.47 | 0.58 | 0.54 | 0.53 | 0.5 | 0.25 | 0.52 | 0.53 | 0.51 | 0.75 | 0.56 | 0.48 | 0.18 | 0.62 | 0.52 | 0.42 | 0.5 |
| Fe ₂ O ₃ | 34.87 | 9.66 | 8.83 | 20.47 | 8.15 | 9.76 | 8.29 | 8.96 | 8.42 | 14.99 | 8.27 | 8.27 | 5.64 | 17.43 | 10.02 | 14.76 | 11.4 |
| MnO | 1 | 0.09 | 0.08 | 0.69 | 0.08 | 0.96 | 0.07 | 0.14 | 0.09 | 0.32 | 0.07 | 0.07 | 0.15 | 0.81 | 0.1 | 0.18 | 0.34 |
| CaO | 18.4 | 0.9 | 0.82 | 33.41 | 0.8 | 33.47 | 0.71 | 3.32 | 0.42 | 19.17 | 0.92 | 2.8 | 15.99 | 35.54 | 1.92 | 9.32 | 11.1 |
| MgO | 0.77 | 6.47 | 6.56 | 1.17 | 5.77 | 4.3 | 6.17 | 6.11 | 6.58 | 0.9 | 6.52 | 5.33 | 2.61 | 0.3 | 6.01 | 5.1 | 3.5 |
| Na ₂ O | 4.14 | 1.2 | 1.37 | 4.97 | 1.66 | 2.89 | 1.57 | 1.54 | 2.13 | 4.22 | 1.86 | 1.34 | 1.68 | 3.47 | 1.33 | 1.77 | 2.3 |
| K ₂ O | 6.11 | 10.44 | 10.75 | 3.33 | 10.34 | 5.08 | 11.21 | 10.59 | 10.47 | 10.45 | 10.75 | 9.14 | 4.01 | 9.1 | 9.53 | 7.34 | 9 |
| P ₂ O ₅ | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.22 | 0.03 | 0 | 0 | 0 | 1.15 | 0 | 0.14 | 0.82 | 0 |
| Cr ₂ O ₃ | 0.13 | 0.03 | 0.02 | 0.12 | 0.03 | 0.05 | 0.03 | 0.04 | 0.03 | 0.09 | 0.03 | 0 | 0.03 | 0 | 0.03 | 0.08 | 0.05 |
| SO ₂ | 0.1 | 0.01 | 0.01 | 0.09 | 0.01 | 0.02 | 0.02 | 0.17 | 0.02 | 0.12 | 0.01 | 0.02 | 0.06 | 0.06 | 0.03 | 0.02 | 0.05 |
| Zr O ₂ | 0.06 | 0.02 | 0.02 | 0.05 | 0.01 | 0.02 | 0.02 | 0.02 | 0.01 | 0.05 | 0.02 | 0.02 | 0.02 | 0.03 | 0.02 | 0.02 | 0.03 |
| ZnO | 0.03 | 0.01 | 0.01 | 0.01 | 0.01 | 0 | 0.01 | 0.01 | 0.01 | 0.02 | 0.01 | 0.01 | 0 | 0 | 0.01 | 0.01 | 0.01 |
| NiO | 0 | 0.01 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| CuO | 0 | 0 | 0.01 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Rb ₂ O | 0 | 0 | 0 | 0 | 0 | 0.01 | 0 | 0 | 0 | 0.01 | 0 | 0 | 0 | 0.02 | 0 | 0 | 0 |
| Y ₂ O ₃ | 0 | 0.01 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| ΣR ₂ O | 0.05 | 0.01 | 0.01 | 0.08 | 0.01 | 0.06 | 0.01 | 0.02 | 0.01 | 0.04 | 0.01 | 0.02 | 0.03 | 0.05 | 0.02 | 0.02 | 0.03 |
| Al ₂ O ₃ /SiO ₂ | 0.35 | 0.89 | 0.9 | 0.52 | 0.87 | 0.55 | 0.9 | 0.09 | 0.91 | 0.38 | 0.92 | 0.72 | 0.3 | 0.29 | 0.8 | 0.6 | 0.62 |
| SiO ₂ /Al ₂ O ₃ | 4.33 | 1.18 | 0.66 | 5.09 | 1.25 | 2.39 | 1.05 | 1.15 | 1.06 | 2.79 | 0.82 | 1.27 | 1.26 | 3.62 | 1.2 | 1.12 | 1.6 |
| K ₂ O/Na ₂ O | 1.48 | 8.73 | 7.85 | 1.68 | 6.55 | 1.76 | 7.14 | 6.88 | 4.91 | 2.48 | 5.77 | 6.8 | 2.39 | 2.62 | 7.18 | 4.15 | 5 |
| ICV | 8 | 8.7 | 0.85 | 6.7 | 0.81 | 3.7 | 0.83 | 0.95 | 0.83 | 3.7 | 0.83 | 0.89 | 1.9 | 6.5 | 1.2 | 1.7 | 3 |
| Fe ₂ O ₃ /MgO | 35.64 | 16.13 | 15.39 | 21.64 | 13.92 | 14.05 | 14.46 | 15.07 | 14.99 | 15.88 | 14.79 | 13.61 | 8.25 | 17.73 | 16.03 | 19.86 | 14.5 |
| Al ₂ O ₃ /TiO ₂ | 18.37 | 47.47 | 62.66 | 19.53 | 67.11 | 62.96 | 65.68 | 61.09 | 66.09 | 17.81 | 60.38 | 63.56 | 88.85 | 16.67 | 59.85 | 54.18 | 47 |
| K ₂ O/Al ₂ O ₃ | 0.7 | 0.32 | 0.32 | 0.81 | 0.32 | 0.33 | 0.33 | 0.33 | 0.31 | 0.78 | 0.32 | 0.3 | 0.26 | 0.88 | 0.31 | 0.33 | 0.43 |

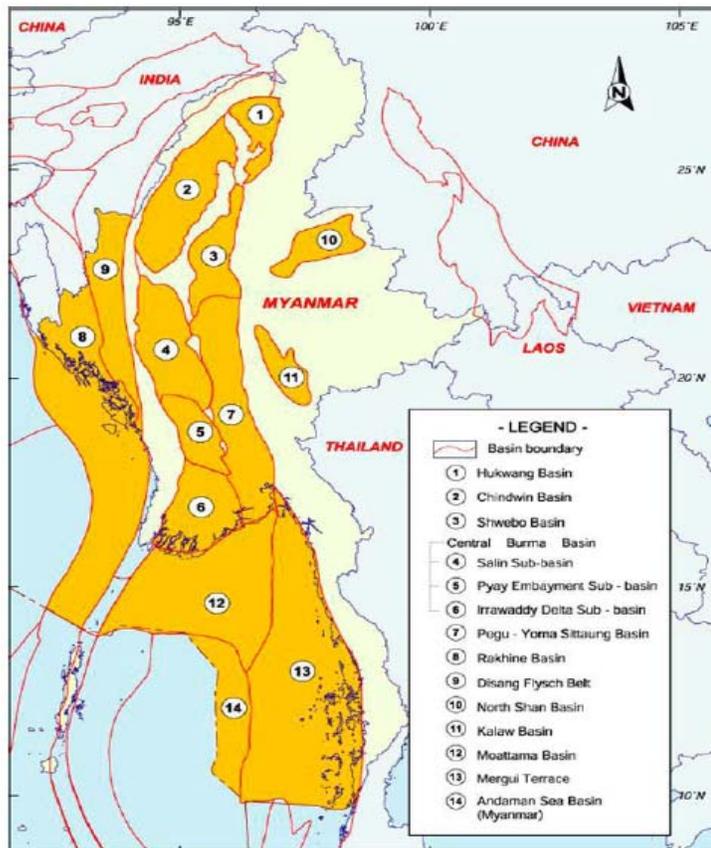


Figure 2. Basins of Myanmar, Pyay Embayment Sub-Basin no. 5, (after Utitsanet *et al.*, 2014).

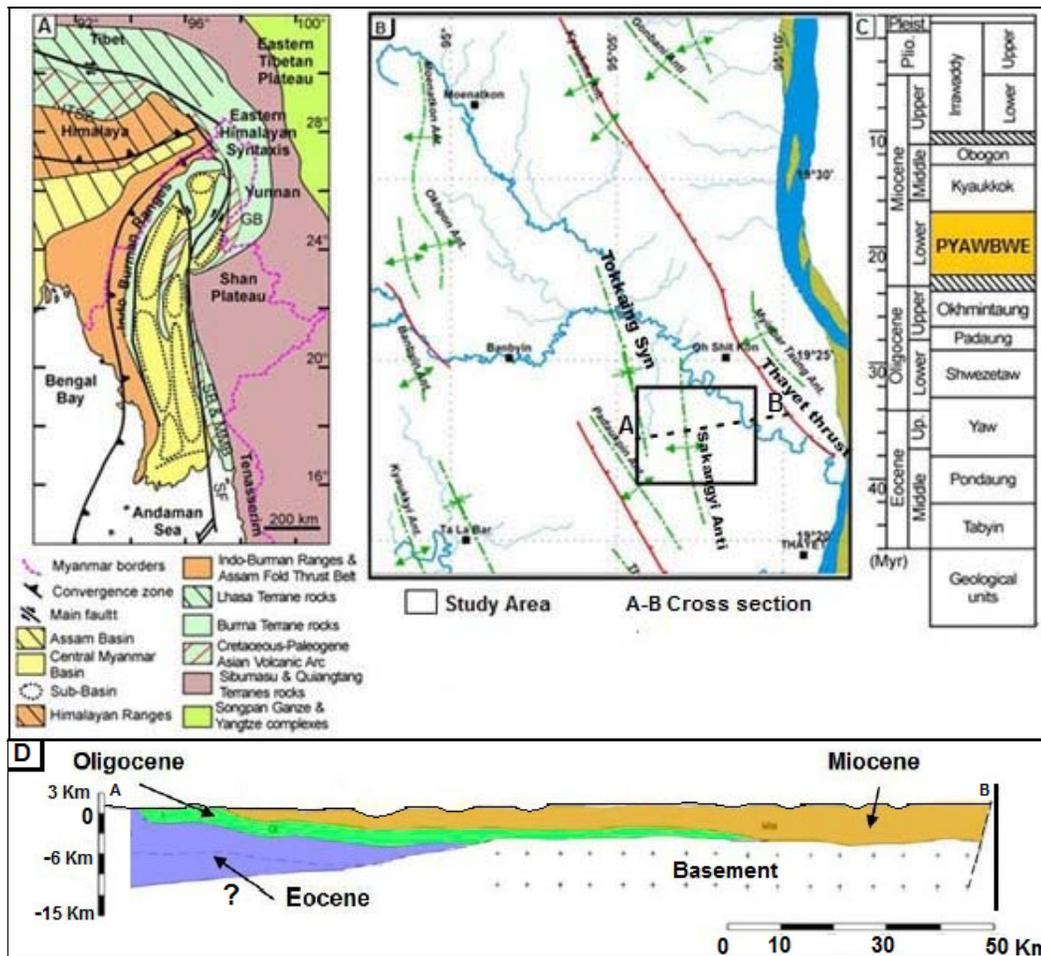


Figure 3. (A) Simplified geological map of central Myanmar Basin (after Mitchell *et al.*, 2012; Metcalfe, 2013). (B) Detailed structural map of the study area in Central Myanmar (partially Pyay sub-basin). (C) Schematic stratigraphic log of the Central Myanmar Basin (Licht *et al.*, 2013) showing the stratigraphic position of the Pyawbwe Formation. (D) Schematic cross section across the study area (Generally modified from Bertrand & Rangin, 2003).

K₂O and Na₂O contents and their ratios show a narrow range of differences (from 3.33 to 10.75 and from 1.2 to 4.97 respectively). This may be attributed to redistribution of alkali elements during post-depositional alterations (Bandopadhyay & Ghosh, 2015). The bivariate plotting K₂O wt. % versus Na₂O wt. % on the graph of Crook (1974) revealed that all the analyzed samples are plotted in the field of quartz-rich shales (Fig. 5). The composition of non-quartz component of the present samples can be evaluated from the values of Index of Compositional Variation (ICV) of Cox *et al.* (1995) where:

$$ICV = (Fe_2O_3 + K_2O + Na_2O + CaO + MgO + MnO + TiO_2) / TiO_2$$

ICV values of the present samples vary from 0.8 to 8.7 (average = 3). The values of ICV more than 1 indicate presence of less clay minerals and more rock-forming minerals such as

plagioclase, K-feldspar, amphiboles, pyroxenes and lithics (Cox *et al.*, 1995). It is known that the values of (K₂O/Al₂O₃) of clays are less than 0.3 and the values of the same ratio of feldspars range from 0.3 to 0.9 (Cox *et al.*, 1995).

The values of K₂O / Al₂O₃ ratio of the sandstones vary narrowly from 0.17 to 1.19 (average = 0.18, Pettijohn, 1957). The K₂O / Al₂O₃ values of the present samples averaged 0.4 indicate a significant quantity of alkali feldspar relative to other minerals in the original rocks. Also, average value of K₂O/Na₂O is 4.9, slightly high indicating presence of K-bearing minerals (McLennan *et al.*, 1983, Nath *et al.*, 2000, Osae *et al.*, 2006).

On the other hand, the present values of SiO₂ show negative correlations with major elements. All major elements except SiO₂ exhibit positive correlations between themselves. These correlations confirm

that the bulk of SiO₂ is present as quartz grains as indicated on Fig. (5), (Rahman & Suzuki, 200). Also, Al₂O₃ shows strong positive correlation with K₂O (r = 0.74) indicates that the bulk Al and K are primarily contributed by clay minerals as illite or smectite (McLennan *et.al.*, 1983). The negative correlation between SiO₂ and MgO (r = -0.5) and SiO₂ and (MgO + Fe₂O₃) (r = -0.64) rule out presence of biotite (Hayashi *et.al.*, 1997). The strong negative correlation between Al₂O₃ and (MgO + Fe₂O₃) (r = -0.46) rule out presence of biotite, chlorite and ferromagnesian minerals (biotite and hornblende) and confirm the presence of smectite (Fig. 6), (Hayashi *et al.*, 1997). The CaO content of the shale beds averaged 2.9 indicating the calcium resides in silicate phases while in

sandstone interbeds averaged 25 reflecting presence of carbonate cement. Also, MgO concentrations are higher than the global shale (Pettijohn, 1957) supporting carbonate association. The strong negative correlation coefficient between Al₂O₃ and CaO (r = 0.91, R² = 0.83) indicates that the carbonates are secondary rather than primary origin (Akarish & El-Gohary, 2011).

Trace Elements

The studied samples (Table, 1) have low concentrations and / or strongly depleted of the analyzed metals as Ni, Co, V, Rb, Y and others may indicate the negligible role of the mafic rocks in the source area.

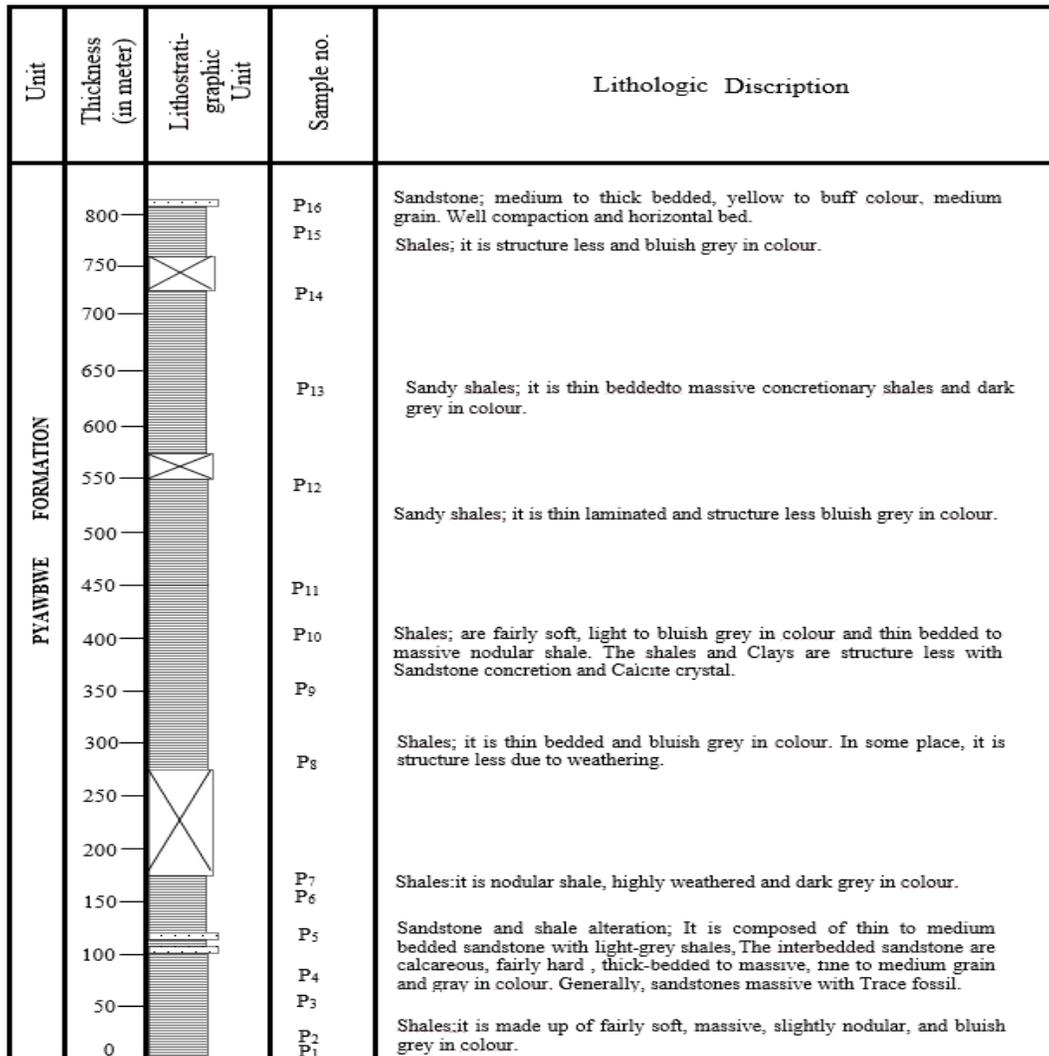


Figure 4. Stratigraphic section showing the different lithologic beds of the Pyawbwe Formation, Central Myanmar Basin.



Plate 1. (1): Photograph showing light-gray, thin-bedded to massive concretionary shales in the middle part of the Pyawbwe Formation. (2): Photograph showing bluish -gray, thin-bedded to massive concretionary shales exposed in the Upper part of the Pyawbwe Formation. (3): Field photograph showing massive, light gray, slightly mottled clay exposed in the Lower part of the Pyawbwe Formation. (4): Photograph showing massive, light gray, slightly mottled clay exposed in the Lower part of the Pyawbwe Formation. (5): Photograph showing massive, light gray, slightly mottled clay exposed in the Lower part of the Pyawbwe Formation. (6): Photograph showing massive, light gray, slightly mottled clay exposed in the Lower part of the Pyawbwe Formation.

Zn shows weak negative correlation with SiO_2 ($r = -0.19$), Al_2O_3 ($r = -0.09$) and very weak positive correlation with TiO_2 ($r = 0.44$) and K_2O ($r = 0.16$). Sr shows negative correlation with SiO_2 ($r = -0.71$), Al_2O_3 ($r = -0.89$), TiO_2 ($r = -0.13$) and with K_2O ($r = -0.77$). Also, Cr shows negative correlations with SiO_2 ($r = -0.48$), Al_2O_3 ($r = -0.6$), K_2O ($r = -0.52$) and positive correlation with TiO_2 ($r = 0.04$), Mn ($r = 0.52$), CaO ($r = 0.41$), Na_2O ($r = 0.73$) and Fe_2O_3 ($r = 0.76$). The negative correlations indicate presence of the trace elements in the clay fraction, while the positive correlations confirm their occurrence in the mafic minerals of the shales. According to Hallberg (1976), low values ($= 1$ in present samples) of

Cu/ Zn ratio suggest oxidizing conditions of deposition indicating very shallow marine conditions. Zn shows very weak negative correlation with MgO ($r = -0.18$), with SiO_2 ($r = -0.19$) and with Al_2O_3 ($r = -0.09$) indicating that Zn is endemic to the mafic minerals of the shales. The present shales have low content of Cr ranges from 20 to 130 ppm (average = 50 ppm) and depleted both Ni, Rb, Y and Cu. Cr and Ni values suggest that ultramafic and even mafic rocks were hardly present or even not widespread at the source region (Garver et al., 1996). Low Sr content of our samples is generally related to low content of CaO probably due to the lack of calcic plagioclase (Akarish &

ElGohary, 2011) Long *et.al.*(2008) showed that the relatively high Rb concentration (>40 ppm) and low Rb/Sr (0.04 – 3.24) ratio indicative of acidic intermediate igneous source rocks that had undergone weak chemical weathering. The present Rb/Sr ratio (average = 0.7) is very low indicating felsic igneous source rocks and reflecting weak to moderate chemical weathering. The present Rb/Sr ratio (average = 0.7) is very low indicating felsic igneous source rocks and reflecting weak to moderate chemical weathering(Rashid, 2002) . This interpretation is consistent with that inferred from the major element interpretation and the framework compositions.

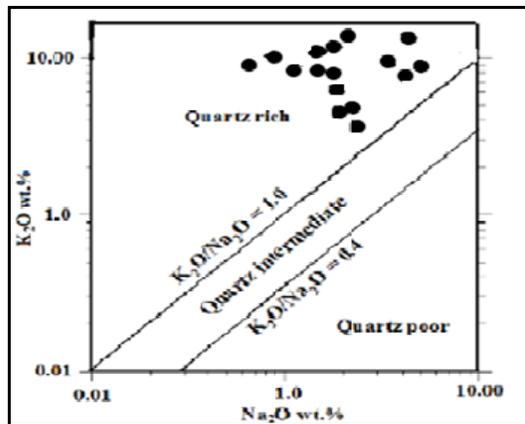


Figure 5. Geochemical classification, bivariate diagram of Pyawbwe Fm., Data reveal quartz rich sediments.

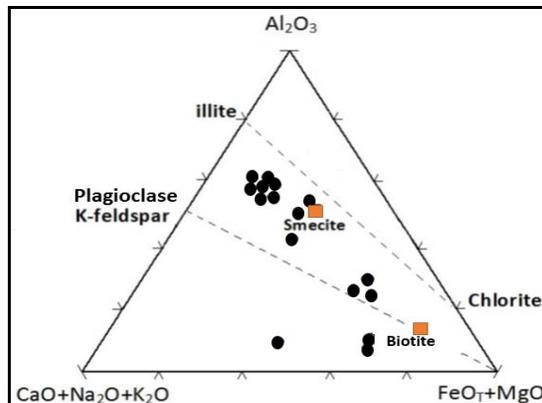


Figure 6. Provenance indicating ternary diagram. The lines are after Hayashi *et al.* (1997).

Source Rocks of the Pyawbwe Formation Sediments

With respect to Al_2O_3/TiO_2 ratios range of 3 to 8 for mafic igneous rocks, from 8 to 12 for intermediate rocks and from 21 to 70 for felsic igneous rocks (Hayashi *et.al.*,1997) . The average

Al_2O_3/TiO_2 ratio of our analyzed samples of the Pyawbwe Formation is 47 suggested felsic igneous rocks as being probably source rocks (Fyffe & Pickerill, 1999). Also, low content of average TiO_2 (= 0.5) indicates presence of phyllosilicates in minor amounts (Dabard, 1990, Condie*et.al.*, 1992, Ferdous& Farazi, 2016).

As indicated (Fig.7),owing to low solubility of their oxides and hydroxides in low temperature aqueous solutions (Stumm& Morgan, 1981, Yamamoto *et.al.*,1986, Sugitani ,1996), the values of Al / Ti ratios of residual soils can be considered to very close to those of the parent igneous rocks. Then, the residual soils contain kaolinite, illite or smectite. The presentation of our data (Fig.7) confirm the most of clays represented by the smectite type. Several studies on clay mineralogy have shownthat during fluvial transportation of Al and Ti involve insignificant fractionation (Yamamoto *et.al.*, 1986).

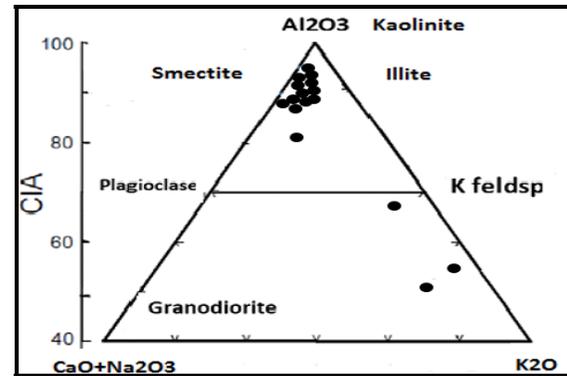


Figure 7. CIA ternary diagram (Al_2O_3 , K_2O , $CaO+Na_2O_3$) of Pyawbwe Fm., (diagram after Nesbitt & Young, 1984).

Plotting the data (Fig. 8) on the tectonic discriminating diagram of Maynard *et.al.* (1982) indicates the sediments of the Pyawbwe Formation is derived from the evolved arc setting, felsic plutonic detritus. Also, their plotting (Fig. 9) on diagram of Roser & Korsch (1986) modified by Murphy (2000) and on McLennan *et al.* (1980), (Fig. 10) confirm the same interpretation.

Paleoweathering and Paleoclimate

The upper crustal rocks are composed of feldspars and volcanic glass. By chemical weathering of these materials ultimately results in the formation of clay minerals (Nesbitt& Young, 1984, 1989; Taylor & McLennan, 1985, Fedo*et.al.*, 1995). Ca,Na,and K are largely removed from the source rocks. These elements are surviving in soil profiles (Fig.11).

Owing to the presence of $K_2O + Na_2O$ (average = 9.9 \approx 10) and the K_2O / Na_2O ratios (average = 4.9 \approx 5) and contents of Na_2O and K_2O of the present samples do not suggest intense weathering of the source area. By applying the data of Lindsey (1999) and Harnois (1988) on the present data, it can be concluded that the provenance of the Pyawbwe sediments were subjected to low to moderate intense chemical weathering in spite of presence of chemical weathering on source rocks (K-feldspars, granitic felsic composition), another chemical weathering of detritus during fluvial transportation and chemical weathering in depocenters and diagenetic processes.

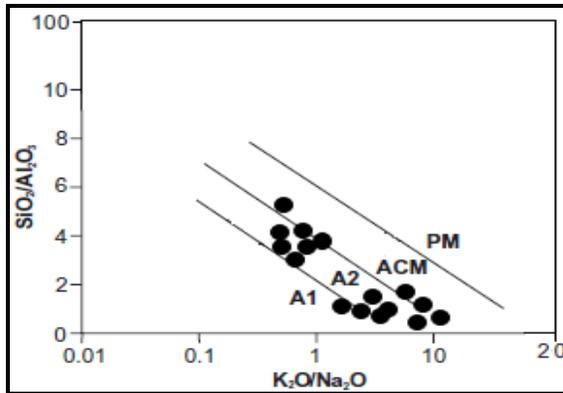


Figure 8. Tectonic setting bivariate discrimination diagram of Pyawbwe Fm. (after Maynard *et al.*, 1982). A1= arc setting, basaltic and andesitic detritus. A2=evolved arc setting, felsic plutonic detritus. ACM=active continental margin. PM= passive margin

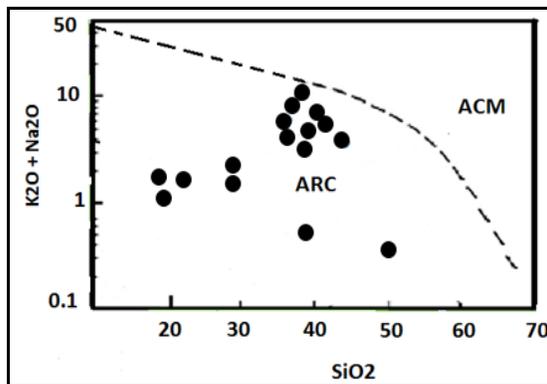


Figure 9. Tectonic setting discrimination diagram of Pyawbwe Fm. The lines are modified after Roser and Korsch (1986) and Murphy (2000). ACM=active continental margin. ARC=oceanic island arc margin.

This may probably due to preponderance of arid to semi-arid paleoclimate (Sutter and Dutta, 1986), (Fig. 11). By applying the calculated average of CIW (Chemical Index of Weathering) and CIA (Chemical Index of Alteration) (Fig.12), according to Nesbitt & Young (1982) and Fedo *et al.* (1995), where: $CIA = Al_2O_3 / (Al_2O_3 + CaO + Na_2O + K_2O) \times 100$, $CIW = Al_2O_3 / (Al_2O_3 + CaO + Na_2O) \times 100$.

(Chemical Index of Alteration) (Fig.12), according to Nesbitt & Young (1982) and Fedo *et al.* (1995), where: $CIA = Al_2O_3 / (Al_2O_3 + CaO + Na_2O + K_2O) \times 100$, $CIW = Al_2O_3 / (Al_2O_3 + CaO + Na_2O) \times 100$.

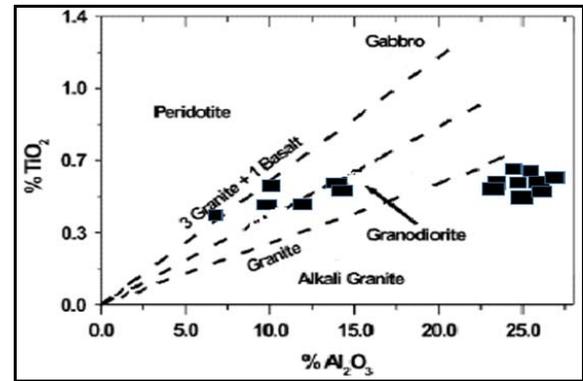


Figure 10. Provenance discrimination diagram of Pyawbwe Fm. $TiO_2\%$ versus $Al_2O_3\%$ bivariate plot, (diagram after McLennan *et al.*, 1980). The granite line and $3\text{ granite} + 1\text{ basalt}$ line are after Schieber (1992).

It can be concluded that the source rocks were exposed to intermediate silicate weathering either in the original source terrane or during transportation by fluvial streams before deposition on passive margin of shallow marine environment (Fig. 13).

Moreover, high averages of CIA values suggest derivation from a stable cratonic source (Hossain *et al.*, 2010, Akarish & El-Gohary, 2011). The present studied sediments commonly form a trend of almost parallel to K-feldspar to smectite clay type.

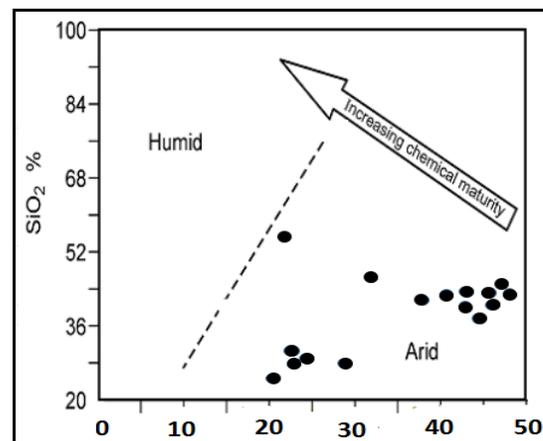


Figure 11. Bivariate paleoclimate discrimination diagram, boundary lines after Suttner & Dutta, 1986, showing the chemical maturity of the Pyawbwe Fm. sediments.

The smectite samples are more than that of K-feldspars with increase of Al_2O_3 concentrations. The presence of smectite mineral improves the

conclusion of passive marine margin as site of deposition with fluctuations (El-Gammal, 1985).

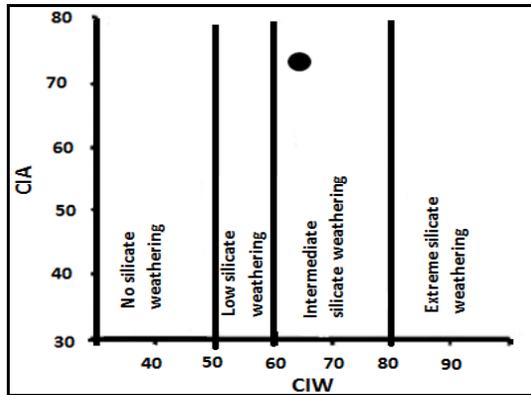


Figure 12. Plot of averages of chemical index of alteration (CIA) vs. chemical index of weathering (CIW), showing intermediate silicate weathering of Pyawbwe Formation.

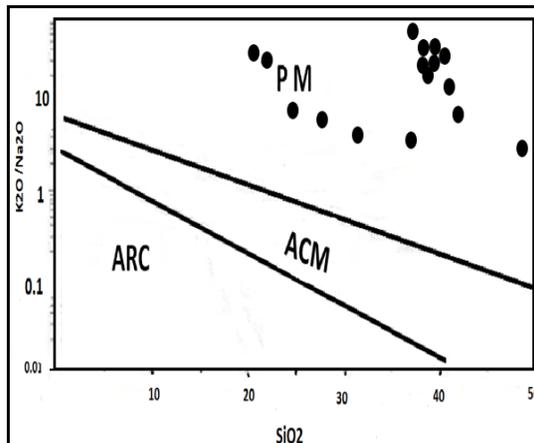


Figure 13. Tectonic Discrimination diagram. Boundaries after Roser and Korsch (1986). PM=Passive Margin. ACM=active continental margin. ARC=ARC=oceanic island arc margin.

Maturity and Tectonics

Suttner & Dutta (1986) proposed a bivariate diagram to constrain the climatic conditions during the sedimentation of the siliciclastic rocks. Plotting our data on that diagram (Fig. 11) indicates that our shales were deposited under arid to semi-arid climate and may be oxic conditions (Jones & Manning, 1994). Average $\text{SiO}_2/\text{Al}_2\text{O}_3$ ratios in unaltered igneous rocks range from ≈ 3.0 (basic rocks) to ≈ 5.0 (acidic rocks). Our data reached to 1.6 (i.e. < 5) indicating of progressive maturity (Roser *et al.*, 1996) as siliciclastics of predominant shales and silts confirmed mature nature (Bahatia, 1983), where the samples are enriched in SiO_2 but depleted in Na_2O , TiO_2 , MnO , and CaO (except the sandstone samples) indicating fluctuating in fluvial sediment input on passive margin. Increase

of degree of chemical weathering may reflect the decrease in tectonic activity and / or change of climate towards warm and humid conditions (Jacobson *et al.*, 2003). Figure (14) confirms that our data imply maturity low than those shales of UC (Upper Crust, after Taylor & McLennan, 1985) and NASC (average North American Shales, after Gromet *et al.*, 1984).

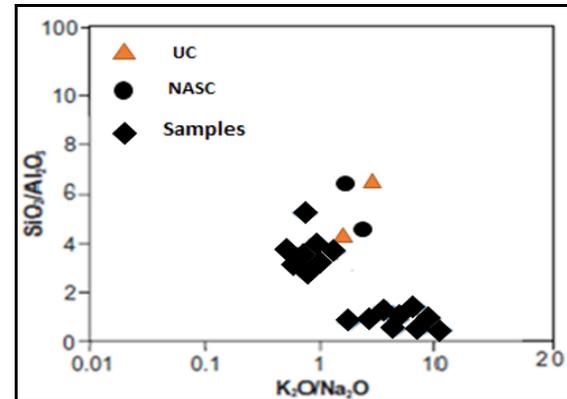


Figure 14. Bivariate diagram of the Pyawbwe shales maturity degree compared with UC and NASC shales.

Provenance

On the major elements and some trace elements – based discrimination function discussed before, indicated the sediments were derived from weathered felsic –granitic –gneissic terrane (Laird, 1972). The type of source terrane as indicated have been located to the eastern and northeastern Myanmar where the Pre-Cambrian to the Paleozoic, intermediate to felsic igneous rocks of the Shan-Thai Block (Mitchell, 1993). A moderate range of weathering inferred in this study indicates the source region was a moderate relief and tectonically calm terrane exposed to arid to semi-arid paleoclimate during weathering.

The weathered debris was transported by fluvial streams to alluvial fans deposited on passive marine margin with relatively shallow fluctuating marine conditions (Fig. 15). On an unconformable Late Oligocene surface marking the top of the Okhintaung Formation, the Pyawbwe Formation was deposited in shallow marine Central Burman Basin which was opened to present Gulf of Martaban (Fig. 15) since Early Eocene times (Wandrey, 2006). The eastern portion of the Central Burman Basin which is part of the Asian Plate was emergent during the Early Eocene (Wandrey, 2006, Searly *et al.*, 2007, Lich *et al.*, 2014). This final interpretation is consistent with the other micropaleontological

and paleoecological research on the same collected samples by the same present authors and also, in consistent with the regional geology of the study area and the basin configuration of the Central Myanmar Basin.

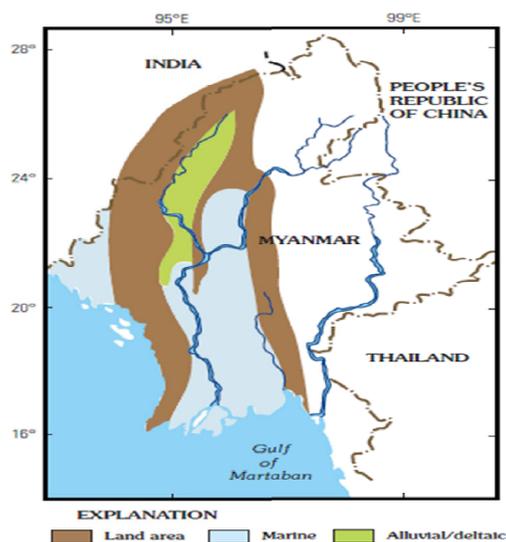


Figure 15. Generalized Early Miocene paleogeography (modified from Khin and Win, 1968)

Conclusion:

The exposed Early Miocene Pyawbwe Formation of Thayet Suddle, northern margin of Pyay sub-basin is an argillaceous unit, which is mainly composed of thick clays and shales with sandstone interbeds. The clays and shales are fairly soft, bluish gray in colour and thin bedded, massive in the base of the formation. The interbedded sandstones are calcareo-

us, fairly hard, and thick bedded to massive, fine to medium grained, and gray in colour. Geochemically, the shales are smectite clay type rich in quartz grains and interbedded with silty arkosic sandstones. Major and some trace elements ratios and diagrams suggest that the source rocks are moderate relief of arc massif of felsic igneous rocks exposed to moderately intensive chemical weathering. The weathered debris were transported by fluvial streams to alluvial fans as progressive maturity sediments and redeposited on passive shallow marine margin with fluctuating silty silty sandstone alluvial loads in arid to semi-arid paleoclimate. Such lithologies and indices suggest that the Shan-Thai block of the eastern and northeastern Myanmar continental region as the dominant source terrane.

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