Petroleum Source Rock Properties of the Neogene Bhuban Shales, Bengal Basin, Bangladesh

(Pencirian Punca Batuan Petroleum di Syal Neogene Bhuban, Lembangan Bengal, Bangladesh)

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ABSTRACT

The present study evaluates the petroleum source rock generation potential of the Neogene Bhuban shales from Bangladesh. Organic geochemical and organic petrological methods were used for analyzing 11 drill core samples from 4 gas fields in the basin. Source rock potential, maceral composition, organic matter abundance, biomarker distribution, thermal maturity, hydrocarbon generation and depositional environment were evaluated. Kerogen in the studied shale samples is classified mainly as Type III with lesser amounts of Type II. Vitrinite is the dominant maceral group observed in the analyzed Bhuban samples followed by liptinite and inertinite. Vitrinite reflectance, T_{max} and biomarker parameters indicate the thermal maturity ranges from just pre-oil window to mid-oil window. Based on its total organic carbon (TOC), extractable organic matter (EOM) and hydrogen index (HI), the analyzed Bhuban shales are ranked as mainly poor to fair source rocks but with good gas generation potential. The dominant terrestrial environment prevailed during the deposition of the studied Bhuban shales while the condition was sub-oxic as indicated by cross-plots of pristane versus phytane and sterane versus pristane/phytane ratios.

Keywords: Bengal Basin; Bhuban shale; hopane; hydrocarbon potential; sterane

ABSTRAK

Kajian ini membincangkan penilaian batuan punca bagi petroleum sampel syal Bhuban Neogene dari Bangladesh. Geokimia organik dan petrologi organik telah digunakan untuk menganalisis 11 sampel teras gerudi daripada 4 medan gas di kawasan tersebut. Potensi batuan punca, komposisi maseral, kedapatan bahan organik, taburan petanda biologi kematangan terma penjanaan hidrokarbon dan persekitaran pengenapan dinilai. Kerogen dalam sampel syal yang dikaji kebanyakannya dikelaskan sebagai Jenis III dengan jumlah Jenis II yang lebih kecil. Vitrinit adalah kumpulan maseral yang dominan dalam sampel Bhuban yang dianalisis diikuti oleh liptinit dan inertinit. Pantulan vitrinit, T_{max} dan parameter bio penanda menunjukkan kematangan terma daripada hanya pra-minyak ke tingkap-minyak pertengahan. Berdasarkan data TOC, EOM dan HI, syal Bhuban yang dianalisis kebanyakannya dinilai sebagai miskin ke sederhana sebagai batuan sumber dengan potensi penjanaan gas yang baik. Persekitaran daratan dominan semasa pengenapan syal Bhuban dikaji manakala keadaan pengenapan adalah sub-oksik sebagaimana yang dicadangkan berdasarkan plot pristana melawan fitana dan sterana melawan nisbah pristana/fitana.

Kata kunci: Hopana; lembangan Bengal; potensi hidrokarbon; sterana, syal Bhuban

INTRODUCTION

The Bengal Basin covers the whole of Bangladesh and part of India. It is bordered with three sides by India and one small side (southeast) by Myanmar (Figure 1). The Bhuban is the most important geological unit of the Bengal Basin since majority of the discovered hydrocarbon resources of Bangladesh have been confirmed within this stratigraphic formation. It is believed that the Bhuban Formation is acting both as source rock (shale part) and reservoir rock (sandstone part) for the existing petroleum system of the Bengal Basin (Imam 2005). So far, there is no detail work published on organic geochemical and petrographical aspects of the Bhuban shales except Farhaduzzaman et al. (2014). Here, we present the data from both techniques: Organic geochemical and petrographical for Bhuban Formation shales. We also draw some conclusions considering kerogen facies, thermal maturity, depositional conditions and hydrocarbon generation potential.

GEOLOGICAL CONTEXT

Bangladesh is situated in the northeastern part of south Asia. The first collision of the northward moving Indian Plate with the Eurasian Plate took place in Lower Eocene. Subsequent subduction led to the rising Indo-Burman Orogeny. The later finally separated the Burmese basins in the east from the Bengal Basin in the west (Reimann 1993). The identified stratigraphic formations of the Deep Basin unit (study area) include (from bottom upward) Tura Sandstone (Paleocene), Sylhet Limestone (Middle Eocene), Kopili Shale (Upper Eocene), Jenum



FIGURE 1. Location map of the study area showing major tectonic elements of the Bengal Basin (Farhaduzzaman et al. 2012a, 2013a, 2013c; Islam 2009; Reimann 1993)

(Oligocene), Renji (Oligocene), Bhuban (Miocene), Boka Bil (Early Pliocene), Tipam Sandstone (Middle Pliocene), Girujan Clay (Middle Pliocene), Dupi Tila (Late Pliocene), Madhupur Clay (Plio-Pleistocene) and Alluvium (Holocene) (Alam et al. 2003; Imam 2005). The studied unit is traditionally believed to have been deposited in a deltaic to shallow marine environment. Mostly all of the discovered gas and oil fields have been found within the Bhuban Formation of the Deep Basin unit. Estimated gas reserve (GIIP) in Bangladesh is 28.42 TCF with about 42 TCF undiscovered (Jamaluddin et al. 2001; Shamsuddin et al. 2004). Using 3D seismic survery (2011-2012), Petrobangla re-estimated the total recoverable oil reserves of 137 million barrels STOIIP.

SAMPLES AND METHODS

A total of eleven shale core samples of Bhuban Formation were collected from four different wells drilled of respective four gas fields (Fenchuganj, Titas, Kamta and Begumganj) located in Deep Basin unit. The samples were crushed into fine powder and analyzed using a Weatherford Source Rock Analyzer (equivalent of Rock-Eval equipment). Bitumen extraction was performed using Soxhlet apparatus and subsequently the extractable organic matter (EOM) was separated into aliphatic, aromatic and NSO fractions by column chromatographic techniques. The aliphatic hydrocarbon fractions were analyzed by gas chromatography (Agilent 6890N Series GC) and gas chromatography mass spectrometry (GCMS). The FID gas chromatograph with HP-5MS column, temperature programmed from 40 to 300°C at a rate of 4°C/min and then held for 30 min at 300°C, was used for GC analysis. GCMS experiments were performed on a V 5975B inert MSD mass spectrometer with a gas chromatograph attached directly to the ion source (70 eV ionization voltage, 100 mA filament emissions current, 230°C interface temperature). For organic petrographical study, the samples were prepared by mounting whole rock fragments in resin blocks and polished to a highly reflecting surface using progressively finer alumina suspension $(1, 0.3 \text{ and } 0.05 \mu m)$. Petrographical examination was carried out under oil immersion using a LEICA DM6000M microscope and CTR6000 photometry system equipped with fluorescence illuminators (ultraviolet light). Except for SRA from Weatherford (USA), all other analyses were carried out in the UM Geology Department, Malaysia.

RESULTS AND DISCUSSION

SOURCE ROCK PROPERTIES

The results obtained from Source Rock Analyzer (SRA) are shown in Table 1. All samples were organic lean (<1%) and hydrogen index (HI) values of Bhuban Formation range from 39 to 232 mg HC/g TOC. Most samples were plotted in the Type III range but one sample plotted within Type II band which implies an organic matter mixture of kerogen types (III/II) (Figure 2). The oxygen index (OI) varies from 64 to 168 mg CO₂/g TOC and the cross-plot of HI versus OI (modified van Krevelen diagram; not shown here) showed mostly Type III character. T_{max} values vary from 429 to 441°C and the mean vitrinite reflectance value ranges from 0.57 to 0.71 (%R_o) for the studied shales. The source rock potential of the analyzed Bhuban Formation shales was evaluated as mostly poor to fair based on classification of Peters and Cassa (1994).

MACERAL COMPOSITION

Vitrinite (60-80 vol.%) is the dominant maceral group found in the analyzed Bhuban shales followed by liptinite (20-25 vol.%) and inertinite (8-15 vol.%). The important liptinitic macerals include sporinite, cutinite, resinite, amorphous organic matter, liptodetrinite and alginite (trace amount). The significant presence of these liptinitic macerals along with the solid bitumen (staining) contribute a minor oil-prone character to the dominantly vitrinitic assemblages (Figure 3).

EXTRACTABLE ORGANIC MATTER AND BIOMARKER CHARACTERISTICS

The total soluble extract varies from 132 to 2814 ppm in the analyzed samples. The concentration of aromatic hydrocarbons (57-548 ppm) is higher than that of aliphatic hydrocarbons (9-300 ppm) in the studied shales. The total soluble hydrocarbon yield ranges from 21 to 139 mg HC/g TOC. The TIC (total ion current), m/z 191 and 217 chromatograms of the analyzed aliphatic fractions were used for interpretation and biomarker distributions (Figure 4). The unimodal distributions of n-alkanes from C_{10} to C_{35} with the maxima standing at C_{16} (mostly) or C_{18} have been observed in the gas chromatograms of the analyzed Bhuban shale samples (Figure 4(a)). The calculated CPI values are close to unity from 0.99 to 1.37 in Bhuban shales. In most of the analyzed samples, the odd carbon homologs dominate over the even carbon homologs. Nonetheless,

The $C_{30}\alpha\beta$ -hopane is the dominant member of the abundant pentacyclic triterpanes (hopanes and moretanes) in all the analyzed Bhuban shale samples (Figure 4(b) and 4(c)). Homohopanes are lower in concentration and dominated by C₃₁-hopane in the studied samples. The S-isomers are dominant over R-isomers in some of the analyzed samples (Figure 4(b)) which indicate the samples are thermally mature for hydrocarbon generation. On the other hand, R-isomers are dominant over the S-isomers among the homohopanes $(C_{31} - C_{33})$ for a few of the analyzed samples which indicate the samples are thermally immature for hydrocarbon generation (Figure 4(c)). In general, $\alpha\beta$ -hopping are more dominant than the $\beta\alpha$ -hopanes (moretanes). The Ts/Tm ratio of the studied samples ranges from 0.26 to 0.87. C_{30} moretane/ C_{30} hopane and $C_{32} 22S/(22S + 22R)$ ranges from 0.09 to 0.44 and 0.47 to 0.63, respectively, for the analyzed Bhuban Formation shales. Considerable abundances of 18a(H)-oleanane (higher plant marker) have been found in all the studied samples. C₂₉ sterane is the most dominant component in the m/z 217 mass fragmentograms which are dominated by regular steranes compared to diasteranes (Figure 4(d)). The most commonly used sterane parameters include C_{29} sterane, $C_{27}/(C_{27}+C_{29})$ sterane and diasterane/sterane ratios and these values range from 36 to 64%, 0.21 to 0.48 and 0.05 to 0.42, respectively, in the analyzed Bhuban samples.

THERMAL MATURITY

Thermally immature to mature oil window has been appraised for the investigated Bhuban Formation shales as evidenced by randomly measured mean vitrinite reflectance and T_{max} values. The randomly measured vitrinite reflectance value ranges from 0.57 to 0.71 (%R) in the analyzed Bhuban samples. It implies that the thermal maturity of the analyzed shale samples varies from just pre-oil window to mid-oil window thermogenic condition for hydrocarbon generation (Peters & Cassa 1994). The T_{max} value obtained from SRA ranges from 429 to 441°C indicates the studied Bhuban samples are of thermally immature to mature for oil generation which is in a good agreement with the interpretation based on vitrinite reflectance values. The recorded production index (PI) value ranges from 0.17 to 0.25 in the analyzed shales and it also suggests the immature to mature oil window condition (Peters & Cassa 1994).

The C_{32} -homohopanes ratio (22S/(22S+22R)) rises from 0 to about 0.65 while 0.57 to 0.62 is the equilibrium range commonly observed during maturation (Seifert & Moldowan 1986). The calculated ratio values of 0.47-0.63 for the studied Bhuban samples fall within and outside the equilibrium range. It is thus demonstrating the thermal maturity condition has been reached partially, i.e., it represents immature to mature oil window. The calculated C_{30} -moretane/ C_{30} -hopane ratio of the studied Bhuban

Sample	Gas Filed	Well no.	Depth (m)	TOC	Tmax	S1	S2	S3	IH	IO	Ιd	%Ro
KSH2	Kamta	1	3138	0.26	429	0.05	0.22	0.45	85	139	0.19	0.57
KSH3	Kamta	1	3139	0.27	433	0.06	0.26	0.39	96	143	0.19	0.60
KSH4	Kamta	1	3377	0.53	435	0.11	0.33	0.42	84	107	0.25	0.63
BSH5	Begumganj	1	3100	0.28	435	0.05	0.22	0.29	80	105	0.19	0.71
BSH6	Begumganj	1	3572	0.32	439	0.05	0.24	0.25	75	78	0.17	0.71
FSH7	Fenchuganj	7	3142	0.29	431	0.06	0.23	0.48	81	168	0.21	0.57
FSH8	Fenchuganj	7	3772	0.36	440	0.05	0.23	0.44	63	121	0.18	09.0
FSH9	Fenchuganj	7	3773	0.40	440	0.05	0.24	0.36	60	90	0.17	0.71
TSH54	Titas	11	2714	0.61	434	0.33	1.41	0.81	232	133	0.19	0.58
TSH65	Titas	11	2783	0.34	439	0.03	0.13	0.02	39	64	0.19	0.62
TSH68	Titas	11	2792	0.30	441	0.10	0.38	0.27	128	91	0.21	0.66
Range			2713-3773	0.26-0.61	429-441	0.03-1.41	0.13-0.81	0.02-0.81	39-232	64-168	0.17-0.25	0.57-0.71
Note: TOC= to	otal organic carbon (wt.%	5). S1= free HC	(mg HC/g Rock). S2	= hvdrolvsable HC	(mg HC/g Rock).	S3= CO2 generated	from pvrolvsis. Tma	x= maximum tempera	ture at top S2 pe	ak (deø C). HC=	= hvdrocarbon. HI=	hvdrogen index

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FIGURE 2. Cross-plot of hydrogen index vs T_{max} indicates the analyzed samples are mostly of Type III kerogen. The organic matters are of mainly thermally immature to mature for hydrocarbon generation (Farhaduzzaman et al. 2013b; Koeverden et al. 2011)



FIGURE 3. Photomicrographs of the analyzed Bhuban shale samples. (a) Common occurrence of Type III kerogen represented by grey vitrinite (vt) maceral, depth 3377 m of the well Kamta 1, (b) Type II kerogen represented by brownish thin cutinite (ct) maceral, depth 2792 m of the well Titas 11, (c) Type II kerogen represented by brownish amorphous materials (ap), depth 3100 m of the well Begunganj 1 and (d) Type II kerogen represented by bright yellow color resinite (re) maceral, depth 3773 m of the well Fenchuganj 2 [Photomicrograph (a) is taken under normal reflected white light while others are under ultraviolet light]



Note: Pr = pristane, Ph= phytane, nC13-32 = normal alkanes; C24/4= tetracyclic terpane, Ta= trisnorneohopane, Tm= trisnorhopane, C28aβ= bisnorhopane, bcT & bcR= bicadenanes T & R, C29aβ= norhopane, C30aβ= hopar C29fa & C30fa= moretanes, C31-33aβ 225 & 22R= homohopanes; C27aaa 20R & 20S = 5a(H).14a(H),17a(H)-cholestane (20R & 20S) (steranes), C28aaa 20R & 20S = 24-methyl-5a(H).14a(H),17a(H)-cholestane (20R & 20S)(steranes), C29aaa 20R & 20S = 24-methyl-5a(H).14a(H),17a(H)-cholestane (20R & 20S) (steranes), C29aaa 20R & 20S = 24-methyl-5a(H).14a(H),17a(H)-cholestane (20R & 20S) (steranes), C29af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steranes), C28af 20R & 20S = 24-methyl-3a(H),17b(H)-cholestane (20R & 20S) (steraneS), C28af

FIGURE 4. (a) The total ion current (TIC) of a studied sample (BSH6) which displays unimodal distribution in the n-alkane members,
(b) An example of a m/z 191 mass fragmentogram of a mature sample (BSH6), (c) An example of a m/z 191 mass fragmentogram of a minimature sample (KSH2), (d) The m/z 217 mass fragmentogram of the studied sample (KSH2) showing the dominance of C₂₉ regular sterane compared to C₂₇ or C₂₈ steranes

shales varies from 0.09 to 0.44 which correspond again to the range of immature-mature thermal maturity condition (Mackenzie et al. 1980). Nonetheless, the solid bitumen or bitumen stain is considered as 'free' or expelled heavy petroleum. This type of bitumen stain has been observed in the analyzed shale samples under the microscope and it also suggests the organic matter of the studied shales has already expelled hydrocarbon in parts within the associated petroleum system of the Bengal Basin, Bangladesh.

HYDROCARBON GENERATION POTENTIAL

The cross-plot of $T_{_{max}}\ (^{\circ}C)$ and production index (PI) depicts that the organic matter of the analyzed Bhuban shales have already started to generate hydrocarbon (Figure 5). This is supported by the earlier interpretation on the basis of vitrinite reflectance and T_{max} values. The measured SRA T_{max} of the analyzed shale samples varies from 429 to 441°C while the hydrocarbon generation usually starts at the maturity level of 435°C (Peters & Cassa 1994; Peters & Moldowan 1993; Peters et al. 2005). The studied Bhuban shales with low to fair TOC (0.26-0.61%), low to fair S2 values (0.13-1.41 mg HC/g TOC), low to moderate total extract yield (132-2814 ppm), low to fair production index (0.17-0.25), low to medium hydrocarbon yield (21-139 mg HC/g TOC) and the presence of common liptinitic materials suggested poor to fair potential for hydrocarbon generation. The mean vitrinite reflectance value ranged from 0.57 to 0.71 (%R) in the analyzed Bhuban shales (equilibrium value is 0.60 as stated by Philp 1985 and Peters & Cassa



FIGURE 5. The cross-plot of T_{max} and production index shows that the investigated Bhuban shale samples fall within and outside the petroleum generation zone (c.f. Farhaduzzaman et al. 2013b)

1994) again supports the organic matter of the analyzed samples have partially achieved the thermal maturity level for hydrocarbon generation. The dominancy of short chain n-alkanes (C_{16} - C_{20}) in the gas chromatogram (TIC) of Bhuban Formation indicates the generation of light hydrocarbons together with some condensates. However, the liquid hydrocarbon potential of the analyzed shale samples is most likely attributed to the contents of liptinitic macerals. The entire petroleum discovery is natural gas with little amount of oil and condensate in the

Bengal Basin, Bangladesh (Imam 2013). This is in good agreement with the present interpretation of hydrocarbon generation potential for the Bhuban Formation.

DEPOSITIONAL ENVIRONMENT

The depositional environment of shale and its condition of deposition is often considered as very complex (Wan Hasiah 1999). The modeled steranes triangular plot (Figure 6) indicates the analyzed Bhuban shales deposited mostly in terrestrial environmental setting with some marine inputs (Huang & Meinschein 1979). The marine influence is also supported by the occurrence of oleanane (c.f., Murray et al. 1997). Diasterane abundances in comparison to regular steranes are often used to distinguish carbonate facies (low diasteranes) from clastic ones (Waples & Machihara 1991). The noticeable presence of diasteranes compared to steranes of the studied shale samples characteristically supports the dominance of clastic depositional facies. The cross-plot of Pr/nC17 versus Ph/nC18 (Figure 7) based on the analyzed samples implies the source of the organic matter of Bhuban shales was mostly terrestrial-derived. The condition of deposition was an alternation of oxic to anoxic within a deltaic to shallow marine environment. This is supported by the cross-plot of Pr/Ph versus C_{27} $(C_{27}+C_{29})$ sterane of the present study (Figure 8). This also indicates the dominant terrestrial environmental settings with oxic-anoxic depositional condition (Waseda & Nishita 1998). Minor marine (pelagic) influence is also suggested by this diagram. The presence of bicadinanes in the gas chromatograms (m/z 191) of the investigated Bhuban samples also supports this terrestrial depositional setting as previously reported by Pearson and Alam (1993).



FIGURE 6. Relationship between sterane compositions, source input and depositional environment. The analyzed Bhuban shales are dominated by terrestrial organic matter input with minor contribution from marine sources (adopted after Waples & Machihara 1991)



FIGURE 7. A plot of pristane/nC₁₇ versus Phytane/nC₁₈ of the examined samples indicates the condition of deposition for the analyzed samples. The analyzed Bhuban samples suggest a mixture of terrigenous Type III and II organic matter inputs deposited under oxic-anoxic condition

(adopted after Peters et al. 2005)



FIGURE 8. A cross-plot of Pr/Ph and sterane $C_{27}/(C_{27}+C_{29})$ ratios indicates the analyzed Bhuban shales deposited within an oxicanoxic terrestrial environmental settings with minor influence from pelagic source (adopted after Waseda & Nishita 1998)

The oxic condition of any terrestrial depositional environment has extensively been appraised by various authors considering the ratio of Pr/Ph (Peters & Moldowan 1993). The source rocks with Pr/Ph ratio greater than 1 are more likely to have formed in an oxidizing environmental setting. Thus the Pr/Ph ratio of the analyzed shales ranged from 0.99 to 3.41 indicates a predominant terrestrial environment. The condition of deposition was oxic. The terrestrial depositional environment is also supported by petrographic data such as dominating vitrinite macerals and the high abundance of woody fragments. The marine influence is suggested by the presence of liptinite macerals such as liptodetrinite and fluorescing amorphous organic matter.

CONCLUSION

The investigated Bhuban shales consist of a mixture of Type III/II kerogens with Type III dominant. They were determined to be thermally immature to mature for hydrocarbon generation based on the mean vitrinite reflectance and T_{max} values. The production index value and the biomarker parameters of 22S/(22S + 22R) hopane, moretane/hopane ratio and sterane data have also supported this level of thermal maturity. SRA and biomarkers data concludes that the analyzed Bhuban shales of the Bengal Basin (Bangladesh) possess poor to fair quality source potential for hydrocarbon generation. The depositional environment is evaluated as dominantly terrestrial setting with minor marine influences. The condition of deposition was an alternation of oxic-anoxic.

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