Geochemical Characterization of Two Niger Delta Crude Oils and Their Mixtures I: Bulk Properties and Aliphatic Hydrocarbon Distributions

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Abstract: Two crude oils from Rivers and Delta States, Niger Delta Nigeria, samples A and F respectively, and their binary mixtures at different compositions, samples B, C, D, and E, were geochemically characterized using bulk properties and aliphatic hydrocarbon distributions. Results of density and API gravity, which classified samples A, B, C, D as light oils and samples E, F as medium oils, were observed to increase and decrease with increase in the composition of oil sample F respectively. The composition of the saturates, from 68.00 to 71.10%, suggest high maturity of the oil samples which did not show a relationship to the compositional mix of the two Niger Delta crude oils. Bimodal distribution of \(n\)-alkanes in samples A and B with a slight prominence of the short chain suggest marine source with significant terrestrial input, while the prominence of the long chain \(n\)-alkanes in sample F, which suggest a terrestrial source, progressively increased in samples C, D, and E. Pr/Ph ratios from 2.37 to 2.70 suggest source rocks deposited in an oxic environment. Ratios of Pr/\(n\)C\(_{17}\), from 0.94 to 1.20, and Ph/\(n\)C\(_{18}\), from 0.39 to 0.53, suggest shale source rocks with terrestrial higher plant input. From these results, the geochemical characteristics of the individual Niger Delta crude oils were exhibited by their binary mixtures. However, density, API gravity, \(n\)-alkane distribution, Pr/\(n\)C\(_{17}\) and Ph/\(n\)C\(_{18}\) ratios were expressed relative to the compositional mix of the two Niger Delta crude oils.

Keywords: \(n\)-alkane, Composition, Isoprenoid, Crude Oil Mixture, Diagnostic Ratio

1. Introduction

Correlation studies of crude oils require parameters that discriminate a homologous suit of oils (oils from the same source) from another [1]. This is primarily accomplished using bulk properties such as American petroleum institute (API) gravity, compound class fractions - saturates, aromatics, resins and asphaltenes (SARA) and distribution of aliphatic hydrocarbons (AHCs) from C8 to C40 normal alkanes (\(n\)-alkanes) including the isoprenoids pristane and phytane [2, 3, 4]. Additional data from biological marker (biomarker) and alkylated polycyclic aromatic hydrocarbon analysis are used to support these parameters [5, 6, 7]. The qualitative and quantitative comparisons of these parameters are important to geochemists for establishing relationship between crude oils [8].

Crude oils are usually characterized by their API gravity range as heavy (< 25° API), medium (25° - 35°), light (35° - 45°) and condensates (> 45°). Data about the composition of the SARA fraction of crude oils is valuable for oil correlation as well as refining processes. The stability of asphaltenes in organic deposition during petroleum production, processing, and refining is related to the SARA composition of crude oil and cannot be associated with only one fraction [9]. Nwadinigwe et al. (2017) showed that despite API gravities of three Nigerian Niger Delta crude oils, Kokori 34.19°, Afiesere 37.76° and Nembe 39.39°, indicated light oils, the aromatics/saturate ratios ascertained Kokori with the lowest ratio of 0.092 will cause more problem by asphaltene deposition in the refinery than the other oils [10].

Aliphatic hydrocarbon distribution and composition (fingerprint) of crude oils vary for different sources [11]. \(n\)-alkane fingerprints obtained from gas chromatography (GC) analysis and characterized by the prominence of \(n\)C13 to \(n\)C19 reflect marine organic sources such as algae, plankton, and bacteria, while the prominence of the \(n\)C25 to \(n\)C35 are usually...
associated with terrestrial higher plant waxes [12]. The relative abundance of pristane and phytane also provide valuable information on source organic matter input, redox potential of the organic matter depositional environment, lithology of the source rock, maturation and biodegradation [13, 14]. In the characterization of crude oil mixtures, Oyekunle and Famakin (2004) noticed that oils characterized by the same specific gravity do not necessarily exhibit the same viscosity. They showed that only binary mixtures of light crude oils produced linear viscosity - compositional relationship [15]. Thus, this study is aimed at assessing the relationship between the geochemical characteristics of two crude oils from different areas of Nigeria’s Niger Delta and their binary mixtures.

2. Materials and Methods

2.1. Study Area

The Niger Delta region is situated in Southern Nigeria between longitude 5° - 9° E and latitude 4° - 6° N on the West African continental margin and at the apex of the Gulf of Guinea [16]. The region is an extremely prolific hydrocarbon province with three stratigraphic sequences namely: the marine between longitude 5° - 9° E and latitude 4° - 6° N on the West petroleum resources (DPR) and used for the study. The crude oils samples A and F were mixed at different compositions to give samples B, C, D, and E 4:1, 3:2, 2:3 and 1:4 respectively.

2.2. Sampling

Crude oils were obtained from Nigeria’s Niger Delta region with the assistance of officials from the department of petroleum resources (DPR) and used for the study. The crude oils (one litre each) were collected from Rivers State and Delta State and labelled samples A and F respectively. Mixtures of samples A and F were made at different compositions to give sample B 4:1, sample C 3:2, sample D 2:3 and sample E 1:4 respectively.

2.3. Crude Oil Characterization

The density and API gravity of the crude oil samples were determined using the density meter [18].

<table>
<thead>
<tr>
<th>Samples</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
<th>E</th>
<th>F</th>
</tr>
</thead>
<tbody>
<tr>
<td>Density (mg/l)</td>
<td>0.829</td>
<td>0.834</td>
<td>0.843</td>
<td>0.844</td>
<td>0.858</td>
<td>0.865</td>
</tr>
<tr>
<td>API Gravity (°)</td>
<td>39.19</td>
<td>38.16</td>
<td>37.32</td>
<td>36.15</td>
<td>33.42</td>
<td>32.08</td>
</tr>
<tr>
<td>Saturates (%)</td>
<td>71.10</td>
<td>70.57</td>
<td>70.60</td>
<td>70.25</td>
<td>71.10</td>
<td>68.00</td>
</tr>
<tr>
<td>Aromatics (%)</td>
<td>25.95</td>
<td>25.90</td>
<td>26.00</td>
<td>25.89</td>
<td>26.06</td>
<td>24.00</td>
</tr>
<tr>
<td>Resin (%)</td>
<td>2.75</td>
<td>3.40</td>
<td>3.23</td>
<td>3.62</td>
<td>2.61</td>
<td>7.62</td>
</tr>
<tr>
<td>Alphaphtene (%)</td>
<td>0.18</td>
<td>0.11</td>
<td>0.15</td>
<td>0.17</td>
<td>0.18</td>
<td>0.32</td>
</tr>
</tbody>
</table>

The densities of the crude oil samples range from 0.829 mg/l, for sample A to 0.865 mg/l for sample F (table 1). Crude oils samples A and F were mixed at different compositions of 4:1, 3:2, 2:3 and 1:4 to give samples B, C, D, and E respectively. From table 1, it was observed that the densities of the crude oil samples increased with increase in the composition of oil sample F. API gravity describes the class of the chemical compounds in crude oils, i.e. light or heavy.

2.4. Fractionation of Crude Oil Samples

The separation of the crude oil samples into saturates, aromatics, resins and asphaltenes (SARA) fractions was achieved by the column chromatography according to Manilla and Onyema [19]. 50 mg of each crude oil sample was weighed into labelled centrifuge tubes and excess pentane added to precipitate the asphaltenes. The content concentrated and introduced onto the top of a glass chromatographic column (25 cm x 1 cm) packed with activated silica gel. n-Hexane was poured into the packed column to elute the saturates, dichloromethane poured to elute the aromatics and dichloromethane/methanol (1:1) mixture poured to elute the resins. The eluents were collected into pre-weighed vials and the solvents evaporated to dryness under a gentle stream of nitrogen at 40°C. The weights of saturates, aromatics, resins and asphaltenes were determined and the normalized percent reported.

2.5. Gas Chromatography Analysis

Analysis of the aliphatic hydrocarbon was performed with an Agilent 7890B gas chromatography (GC) system fitted with an HP-5 silica capillary column 30 m x 320 μm ID and 0.25 μm film thickness and equipped with a flame ionization detector (FID). One microlitre (1μL) of the saturate fraction was injected into the GC capillary column with the aid of a G4513A automatic liquid sampler using a split ratio mode of 100:1. Helium was used as the carrier gas. Oven temperature was programmed from 40 to 315°C at 5°C/min with a 5 min hold at 40°C and 30 min hold at 315°C. Peak identification was by their relative retention times in comparison with internal standard and quantification by area integration of each identified peak. These were processed by Chemstation OPEN LAB CDS software.

3. Results and Discussion

3.1. Bulk Properties

The bulk properties of crude oils are useful for initial screening of the quality and class of chemical compounds present in the oil. The result of the bulk properties determined for crude oil samples A, F and their mixtures, samples B, C, D, and E are presented in Table 1.

<table>
<thead>
<tr>
<th>Bulk properties</th>
<th>Samples</th>
<th>A</th>
<th>B</th>
<th>C</th>
<th>D</th>
<th>E</th>
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<td></td>
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<td>0.11</td>
<td>0.15</td>
<td>0.17</td>
<td>0.18</td>
<td>0.32</td>
<td></td>
</tr>
</tbody>
</table>
saturated etc. The API gravities of the samples range from 32.08° to 39.19° (table 1). Samples A, B, C, and D have high API gravities (> 35°) which classify them as light crude oils, while samples E and F are classified as medium crude oils. Table 1 showed that the API gravities of the crude oil samples decreased with increase in the composition of oil sample F.

The SARA compositions of the crude oil samples are presented in Table 1. During thermal maturation of crude oils, the high molecular weight compounds are cracked leading to an increase in hydrogen-rich low molecular weight hydrocarbons and saturates [20]. Saturates were the most abundant fraction in all the samples with composition from 68.00 to 71.10%, while the asphaltenes were the least abundant with composition from 0.11 to 0.32% (table 1). The saturate hydrocarbons were significantly more than the other compound classes and indicate all the oil samples were highly matured. This result indicates that the characteristic high-level maturity of the Niger Delta crude oils, sample A and F, was preserved in their mixtures, samples B, C, D, and E.

3.2. Distribution of n-Alkanes

n-Alkanes were identified by GC analysis of the saturate fractions which were well resolved. The identified n-alkanes range from nC8 to nC38 (figure 1). GC fingerprints of n-alkanes characterized by the dominance of short carbon chain (nC13 - nC20) reflect crude oils derived from marine organism sources, while the dominance of the long carbon chain (nC25 - nC35) are associated with terrestrial higher plant wax sources [21]. Distribution of n-alkanes in the samples consist of both the short and long carbon chain n-alkanes (figure 1). Sample A showed a bimodal distribution of n-alkanes with a slight prominence of the short chain with the maximum peak carbon number (Cmax) of nC14, while sample F showed the prominence of the long chain n-alkanes and a (Cmax) of nC35. This result suggests that sample A crude oil from Rivers State, which exhibited a bimodal distribution, was derived from marine source with significant terrestrial input to source and sample F crude oil from Delta State was derived from terrestrial organic source. Sample B showed a bimodal n-alkane distribution with a slight prominence of the short chain and Cmax of nC14, which is similar to sample A and suggest marine source with significant terrestrial input. The n-alkane distribution of samples C, D, and E showed the prominence of the long chain n-alkanes. This prominence of the long chain n-alkanes, which is similar to sample F, was observed to progressively increase respectively. Cmax for sample C is nC31, sample D nC33, and sample E nC33. These n-alkane distributions suggest terrestrial source for samples B, C and D. These results reveal that the n-alkane features of the two Niger Delta crude oils, samples A and F, were expressed in their mixtures relative to their compositional mix.

3.3. Composition of Isoprenoids

The most common isoprenoids in crude oils are pristane (C19) and phytane (C20). They are generated from the phytyl (C20) side chain of chlorophyll molecule in an oxidizing (oxic) or reducing (anoxic) depositional environment respectively [5]. Pristane (Pr) and phytane (Ph) were identified and well resolved in all the oil samples (figure 1). Common ratios derived from the abundance of these isoprenoids, Pr/Ph, Pr/nC17 and Ph/nC18, are calculated for the samples and presented in Table 2.
Two Niger Delta crude oils from Rivers and Delta States (samples A and F respectively) and their mixtures at different compositions (samples B, C, D, and E) were geochemically characterized using bulk compositional properties and aliphatic hydrocarbon distributions. API gravities classified samples A, B, C, D, and E as light oils, samples E and F as medium oils. The API gravities were relative to the compositional mix of samples A and F (decreased with increase in the composition of oil sample F). The saturate hydrocarbons were significantly more than the other compound classes (aromatics, resins, and asphaltenes), but did not show a relationship to their compositional mix. The prominence of the long chain n-alkanes for samples A, B, C, D and E which progressively decreased suggest terrestrial organic source, while bimodal distribution with a slight prominence of the short chain n-alkanes for samples E and F suggest mixed marine and terrestrial organic sources. This revealed the n-alkane distribution features were expressed relative to the compositional mix of samples A and F. Pr/Ph ratios for all the samples were consistent with deposition in an oxic environment, but with a nonlinear relationship. Pr/nC17 and Ph/nC18 ratios suggest shale source rocks with terrestrial higher plant input, which were expressed relative to the compositional mix of samples A and F.

### Table 2. Calculated isoprenoid ratios for the studied Niger Delta crude oils and their mixtures.

<table>
<thead>
<tr>
<th>Samples</th>
<th>Pr/Ph</th>
<th>Pr/nC17</th>
<th>Ph/nC18</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample A</td>
<td>2.40</td>
<td>1.22</td>
<td>0.53</td>
</tr>
<tr>
<td>Sample B</td>
<td>2.70</td>
<td>1.29</td>
<td>0.51</td>
</tr>
<tr>
<td>Sample C</td>
<td>2.54</td>
<td>1.12</td>
<td>0.51</td>
</tr>
<tr>
<td>Sample D</td>
<td>2.37</td>
<td>1.04</td>
<td>0.49</td>
</tr>
<tr>
<td>Sample E</td>
<td>2.42</td>
<td>0.94</td>
<td>0.43</td>
</tr>
<tr>
<td>Sample F</td>
<td>2.44</td>
<td>1.03</td>
<td>0.39</td>
</tr>
</tbody>
</table>

From table 2, Pr/Ph ratios which range from 2.37 - 2.70 indicate pristane was dominant over phytane (Ph) in all the oil samples. Ratios of Pr/Ph less than 1 indicate anoxic deposition, while ratios above 1 reflect oxic depositional environment [22]. Pr/Ph ratios of sample A (2.40) and sample F (2.44) indicate the Niger Delta crude oils were generated from source rocks deposited in an oxic environment. The ratios obtained for the mixtures of samples A and F are sample B 2.70, sample C 2.54, sample D 2.37 and sample E 2.42. These ratios are consistent with crude oils generated from source rocks deposited in an oxic environment. From the results, it was observed that diagnostic feature of Pr/Ph for the two Niger Delta crude oils, samples A and F, was expressed in their mixtures, but did not show a relationship to their compositional mix. Calculated ratios of Pr/nC17 and Ph/nC18 for sample A is 1.22 and 0.53 while for sample F is 1.03 and 0.39 respectively (table 2). Pr/nC17 ratios less than 0.5 indicate marine organic matter input and values greater than 0.6 indicate terrestrial higher plant input to source. Ratios of Ph/nC18 less than or equal to 0.3 are typical of crude oil generated from carbonate source rocks and ratio values greater than 0.3 typify shale source rocks [23]. The Pr/nC17 and Ph/nC18 ratios of samples A and F suggest the studied Niger Delta crude oils were generated from shale source rocks with terrestrial higher plant input. Samples B, C, D and E ratio values for Pr/nC17 are 1.29, 1.12, 1.04 and 0.94, while for Ph/nC18 are 0.51, 0.51, 0.49 and 0.43 respectively (table 2). These ratios suggest crude oils generated from shale source rocks with terrestrial higher plant input. Samples B, C, D and E are mixtures of samples A and F at different compositions of 4:1, 3:2, 2:3 and 1:4 respectively. From the results, it was observed that diagnostic features of Pr/nC17 and Ph/nC18 ratios for the two Niger Delta crude oils, samples A and F, were preserved in their mixtures and relative to their compositional mix.

### 4. Conclusion

Two Niger Delta crude oils from Rivers and Delta States (samples A and F respectively) and their mixtures at different compositions (samples B, C, D, and E) were geochemically characterized using bulk compositional properties and aliphatic hydrocarbon distributions. API gravities classified samples A, B, C, D and E as light oils, samples E and F as medium oils. The API gravities were relative to the compositional mix of samples A and F (decreased with increase in the composition of oil sample F). The saturate hydrocarbons were significantly more than the other compound classes (aromatics, resins, and asphaltenes), but did not show a relationship to their compositional mix. The prominence of the long chain n-alkanes for samples A, B, C and D which progressively decreased suggest terrestrial organic source, while bimodal distribution with a slight prominence of the short chain n-alkanes for samples E and F suggest mixed marine and terrestrial organic sources. This revealed the n-alkane distribution features were expressed relative to the compositional mix of samples A and F. Pr/Ph ratios for all the samples were consistent with deposition in an oxic environment, but with a nonlinear relationship. Pr/nC17 and Ph/nC18 ratios suggest shale source rocks with terrestrial higher plant input, which were expressed relative to the compositional mix of samples A and F.

### References


