POLYCYCLOALCANE HYDROCARBONS IN TARIBANI OIL

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ABSTRACT
For the purpose of extraction and identification of cyclic alkane hydrocarbons in oil from the Tarihani field, Georgia, concentrates isolated from 200-350°C fractions have been studied. The concentrate preparation included the following steps: (a) distillation of a crude oil, (b) dearomatization of distillates by silica gel, (c) three-stage thermal diffusion separation of isoalkanes and cycloalkanes, and (d) thiourea adduction. Ten thermal diffusion fractions were obtained. A mixture of the IX and X fractions and separately the VIII fraction were studied. The extracts from these fractions with thiourea were studied as well. The composition of concentrates has been determined by the gas chromatography–mass spectrometry (GC-MS) method. The main components of the VIII thermodiffusion fraction were C11-C22 isoprenoid alkanes. Among them the predominant were pristane C19 and the phytane C20, the well-known biomarkers of oil. In the VIII thermodiffusion fraction including dimethyl-, trimethyl-, tetramethyl-, ethyl-, methyl-ethyl-, diethyl-, propyl, and methyl-propyl adamantanes, were identified.

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Introduction. Naphthenic hydrocarbons make up the main part of petroleum hydrocarbons (40-60%). They are the most interesting components, the presence of which determines the valuable chemical and technical properties of petroleum products. Consequently, their study is an integral part of the problems of petrochemistry and oil refining. Lower naphthenes C3-C9, boiling up to 200 °C, have been studied quite well. In the middle fractions, the complexity of the composition and structure of bi-, tri-, tetra- and pentacyclic naphthenes represented in them with different type of bonding of five- and six-membered rings (joined, condensed, bridged type, etc.) significantly increases. These hydrocarbons, due to the variety of structural and spatial isomers, are the least studied hydrocarbons, with the
exception of adamantane compounds. On the GLC chromatograms of naphthenic concentrates a continuous undivided background in the form of a "hump" is obtained, indicating the exceptional complexity of these fractions. Interest towards the study of polycyclic naphthenes present in the middle fractions of petroleum was caused by the scarcity of information about these fractions and about the tri-, tetra- and pentacyclic alkanes present in them in the form of compact structure of C_{11}-C_{16} composition. Studies of these hydrocarbons in Georgian oils were carried out for the first time at the molecular level, with the aim of determining individual chemical compounds in oils with definite deciphering of their structure and spatial structure.

On the territory of Georgia oil deposits are known from the ancient times, and there are more than 1500 manifestations of oil and gas. The Taribani field is a large under-developed oil field covering an area of approximately 80 km² located in the southeast of Georgia, on the Shiraki Valley (Kakheti, the 12th licensed block). Productive zones are situated in the Miocene and Pliocene aged sandstone reservoirs that have been deposited in fluvo- To study the individual hydrocarbon composition of the polycyclic naphthenes of the middle fractions of the investigated oil, modern methods of separation and research were selected to achieve greater differentiation by the type of the structure of molecules of complex hydrocarbon mixtures, since the higher the degree of differentiation of the investigated objects, the higher and more reliable is the effectiveness of methods for their analysis. The crude oil sample was taken from the well # 23 of Taribani oils at a depth between -2345 and -2374m. The Taribani oil is paraffinic (7%), although like all other Georgian oils, at the same time it contains a considerable amount of isoprenoid structures; the content of light fractions (55-350°C) makes up to 60%, sulfur – 0.2%, tar– 8.5%, asphaltenes– 6.2% [1-3].

The main goal of the present study was to develop the feasible sequential procedure of oil distillation, dearomatization, separation of isoalkanes from cycloalkanes, and extraction of cycloalkanes for the subsequent analysis by the gas chromatography-mass spectrometry (GC-MS) method.

**Purpose of the study.** Qualitative determination of polycyclic saturated hydrocarbons in the high-boiling fraction (200-350°C) of the Taribani oil

**Materials and methods.** All used chemicals were analytical grade. The certified analytical standards of adamantane, methyladamantanes, n-prolyladamantane, 2-n-buthyladamantane, and diamantane were supplied by Sigma-Aldrich (Germany). The HPLC/GC grade methanol, n-pentane, benzene, and petroleum ester were purchased from Sigma-Aldrich and Merck (Germany).Activated fine silica gel(0.07–0.15 and 0.2–0.3 mm) was supplied by Labstatus (Ukraine ) and Chem Reactive Snab (Almaty, Kazakhstan), coarse silica gel (0.4–2.5 mm) by Salavat catalyst plant (Russia), the thin layer chromatography silica gel plates were from Sanpont (Czech Republic) and from Merck (Germany); sodium sulfate purchased from Salavat catalyst plant (Russia), thiourea – from Biochem (France). Carrier gas - Hydrogen (CB Index: 87; Product Catalog: 9628, Customer Evaluation: 6, CAS: 1333-74-0; Remarks: Brand: Sigma-Aldrich | Product Number: 295396 | Purity: ≥99.99%/.

**Preparation of cycloalkane concentrates**

The concentrate preparation included the following steps:

- Distillation collecting two consecutive fractions 200-250°C and 250-350°C;
- Deearomatization of distillates by silica gel adsorption chromatography;
- Three-stage thermal diffusion separation [4-6] of isoalkanes and cycloalkanes on the Mel polder’s fractionating column [7] obtaining ten thermodiffusion fractions; fractions IX and X had almost identical content of polycyclic saturated hydrocarbons and have been united, the fraction VIII was studied separately;
- Extraction of cycloalkanes by thiourea adduction [8].

One of the most difficult problems to solve in oil studies is separation of isoparaffins from cycloparaffins and the latter in terms of their cyclicity. Thermal diffusion is almost the only method that allows solving this problem successfully. The most widely distributed were Mel polder’s vertical columns of "pipe in a pipe" type with a hollow gap and a coil on the outer surface of the inner tube. The coil in the working space significantly increases an efficiency of separation and much shortens the analysis time. By selecting the optimal conditions (temperature gradient, separation time) and column efficiency, it becomes possible to successfully extract the concentrates of different hydrocarbons from the petroleum fractions.

In this study large and micro - TDF columns were used. The height of the large columns was 1500 cm, the volume of the annular space – 50 ml, the clearance – 0.3 mm with a coil in the working space. Stainless steel microcolumns of the original design had a volume of 3, 4 and 4.5 ml. These columns are of the Melpolder’s columns type, their efficiency factor in dividing the model mixture of
cis-trans decalines (1: 1) is maximal $S = 99\%$ in 8-10 hours. Their height is 110 mm. These columns were designed and manufactured at the Petre Melikishvili Institute of Physical and Organic Chemistry, Laboratory of Petroleum Chemistry. The columns have 10 drain cocks that provide separation of mixtures into ten equal-volume fractions.

For extraction of naphthenic concentrates, saturated fractions of 200-250 °C and 250-350 °C were subjected to three-fold thermal diffusion. At each stage mixtures of the two lower fractions (IX + X) were placed into a TDF column for the next separation step, at the end of the process, a mixture of fractions (IX + X) - naphthenic concentrates was taken. The degree of TDF separation was high, estimation was made by change in the refractive index: fraction 200-250 °C – $n_D^{20}$ was changed from 1.4486 to 1.4768 and fraction 250-350 °C – $n_D^{20}$ was changed from 1.4495 to 1.4972.

Gas chromatography-mass spectrometry analysis
The chromatographic separation of mixture in the thiourea extract of IX+X thermal diffusion fractions have been carried out on capillary columns 60m × 0.25mm (200-250°C distillation fraction) and 40m × 0.25mm (250-350°C distillation fraction), linear programming of temperature – from 100°C with a speed of 2°C/min (200-250°C fraction) and from 130°C with a speed of 3°C/min (250-350°C fraction). Compounds in the VIII thermo diffusion fraction and corresponding extract have been separated on the dimethylpolysiloxane GC capillary column 200 m × 0.25 mm, temperature rose from 40°C to 280°C at a speed of 2°C/min during 70 min. Mass spectra have been measured on GC-MS systems with quadruple and magnetic sector analyzers at the department of spectrometric investigations of the National Institute of Standards and Technology (NIST, Gaithersburg, USA), the automated mass deconvolution and identification system (AMDIS) was used for data analysis.

Research results. Chromatograms of the thiourea extracts obtained from the naphthenic concentrate (mixture of the 3rd stage of thermal diffusion fractions IX + X) of the saturated distillation fractions of Taribani oil are shown in Figures 1 and 2.

Fig. 1. Chromatogram of the thiourea extract obtained from the naphthenic concentrate
According to this experimental data, the concentrate of the 200-250°C fraction contains about 92% of bi-, tri- and tetracycloalkanes, and the concentrate of 250-350°C fraction contains more than 96% of the tri-, tetra- and pentacycloalkanes. More than 50 individual polycyclic alkanes were identified: adamantane and its nine homologues of C\textsubscript{10}–C\textsubscript{13} composition; twelve tricyclic C\textsubscript{11}–C\textsubscript{13} hydrocarbons (tricycloundecanes and tricyclododecanes – bridge tricycloalkanes, the predecessors of adamantanes in oil [9,10]; ten tetracyclic C\textsubscript{11}–C\textsubscript{16} hydrocarbons containing adamantane structure in polycyclic system; four pentacyclotetradecanes – diamantane (C\textsubscript{14}H\textsubscript{20}) and its three derivatives of C\textsubscript{15}H\textsubscript{22} composition; six bicyclic hydrocarbons with sesquiterpene type structure, polymethyl-substituted decalines of C\textsubscript{14}–C\textsubscript{16} composition, structure of which have relict character[11].

It should be noted that when studying Taribani oil [1] ten compounds have been detected in oil for the first time: tricyclo[5.2.2.0\textsuperscript{3,7}]undecan (C\textsubscript{11}H\textsubscript{18}, peak 10), 1-n-propyladamantane (C\textsubscript{13}H\textsubscript{22}, peak 33), 1-ethyl-2-methyl-adamantane (C\textsubscript{13}H\textsubscript{22}, peak 21, Figure 1), methyltetradecylo[6,3,1,1\textsuperscript{6.0}\textsuperscript{10}]tridecan (C\textsubscript{14}H\textsubscript{22}, peak 18); 1-methyl- and 3-methyl-diamantanes (C\textsubscript{15}H\textsubscript{22}, peaks 26 and 29, respectively), tetracyclo[7.3.1.0\textsuperscript{2,7},1\textsuperscript{7,11}]tetradecan (C\textsubscript{14}H\textsubscript{22}, peak 19) and its derivatives up to hexamethyl-substituted analogs (C\textsubscript{15}H\textsubscript{24}–C\textsubscript{16}H\textsubscript{34}, peaks 24,27,28,30,32 and 33 Figure 2).

Chromatogram of the thermodiffusion fraction VIII is shown in Figure 3. Complex study of MS data and GC retention indices made it possible to divide isomers, which in itself is a great difficulty in the process of compounds identification.

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**Fig. 2. Chromatogram of the thiourea extract obtained from the naphthenic concentrate of the saturated 250-350°C fraction.**

**Fig. 3. GC of concentrate VIII obtained after thermal diffusion; Isoprenoides: 1-Undecane 2,6-dimethyl-(C\textsubscript{13}); 2-Dodecane 2,6-dimethyl-(C\textsubscript{14}); 3-Dodecane, 2,6,10-trimethyl-(C\textsubscript{15}); 4-Tridecane 2,6,9trimethyl-(C\textsubscript{16}); 5-Tetradecane 2,6,10-trimethyl-(C\textsubscript{17}); 6-Pentadecane 2,6,10-trimethyl-(C\textsubscript{18}); 7-Pristan-(C\textsubscript{19}); 8-Pristan-(C\textsubscript{20}); 9-Heptadecane 2,6,10,15-tetramethyl-(C\textsubscript{21}); 10-Octadecane 2,6,10,15-tetramethyl-(C\textsubscript{22}); 11-Nonadecane 2,6,10,15-tetramethyl-(C\textsubscript{23}).**
The main components of the VIII thermodynamic fraction, as it was expected, are C_{11}–C_{22} isoprenoid alkanes. Among them the predominant are 2,6,10,14-tetramethylpentadecane (pristane, C_{30}H_{62}) and 2,6,10,14-tetramethylhexadecane (phytane, C_{30}H_{62}), the well-known biomarkers of oil. Cyclopentanes, cyclohexanes and decalines having retention times from 89 to 95 minutes (Figure 4) were characterized by low intensity. The list of compounds detected in the extract is presented in Table 1.

![Figure 4. A part of chromatogram (RT=89-95min) of the VIII fraction:](image)

**Table 1. List of hydrocarbons identified in this segment (89 – 95 min)**

| A1  | 3-methylhexyl-methylcyclopentane | B1, B3, B5, B8, B10 | cyclohexanes with C_9 substituents |
|-----|----------------------------------|----------------------|-----------------------------------|
| A2  | 1,5-dimethylhexyl-methylcyclopentane | B2, B4, B5, B7, B9 | cyclohexanes with C_8 substituents |
| A3  | 3-methylhexyl-dimethylcyclopentane | C1–C9 | Decalines with 4 C-atoms |
| A4  | 3-methylheptyl-methylcyclopentane | C10 – C11, C13 | decalines with C_4 substituents |
| B11 | n-octylcyclohexane | C12 | 1-n-butyldecalin. |

Separation of isoalkanes and cycloalkanes present in the thermal diffusion fraction was successfully carried out by formation of inclusion complex (adduct) of thiourea with cycloalkanes and their extraction. Chromatogram of extract obtained after the thiourea adduct formation in the thermal diffusion fraction VIII is shown in Figure 4; extract contains C_{13} – C_{21} bicyclic and C_{19} tricyclic alkanes separated on long capillary column, mass spectra of relict-type bicyclic compounds (decaline and perhydroindane derivatives) are shown in the Figure 5.

![Figure 5. Chromatogram of extract obtained after thiourea adduct formation in the VIII fraction](image)

1-Bicycloalkane(C13); 2- Bicycloalkane(C14); 3,4,6,11,13,15- Bicycloalkane(C15); 5,12,14,16,17 - Bicycloalkane(C16); 18 - Bicycloalkane(C17); 19-20 – Bicycloalkane (C20); 23,24,28 – Tricycloalkane (C19); 25-27 - Bicycloalkane(C21).
The initial part of a chromatogram (retention time from 70 to 85 minutes, noted by a dotted rectangle on the Figure 5) is shown on the Figure 7 in the increased scale, and a list of hydrocarbons identified in this segment is presented in Table 2.

![Figure 7: Chromatogram](image)

**Fig. 6. A segment of chromatogram (RT=70 – 85 min) of thiourea extract**

**Table 2. List of hydrocarbons identified in this segment (70 – 85 min)**

|   | Hydrocarbon                          |   | Substituted Hydrocarbon                           |
|---|-------------------------------------|---|--------------------------------------------------|
| a1| 3-Methyladamantane                  | b5| Tricyclo[7.3.0.0\textsuperscript{10,16}]dodecane, |
| a2| 1,4-Dimethyladamantane              | b6| Tricyclo[7.2.1.0\textsuperscript{5,9}]dodecane   |
| a3| 1,2-Dimethyladamantane              | b7| Tricyclo[6.3.1.0\textsuperscript{4,12}]dodecane  |
| a4| 1,3-Dimethyladamantane              | e1–e5| Methyldecalines                                    |
| a5| 2-Ethyladamantane                   | e6–e7| Ethyldecalines                                     |
| d1| 1,3,5,6-Tetramethyladamantane       | c1| 1,3,4-Trimethyladamantane                          |
| d2| 1-Methyl-3-propyladamantane         | c2| 1,3,5-Trimethyladamantane                          |
| d3| 1,3-Diethyladamantane               | c3| 1,3,4-Triethyladamantane                          |
| d4| 2-n-Butyladamantane                 | c4| 1-Ethyl-2-methyladamantane                        |
| d5| 1-n-Butyladamantane                 | c5| Tricyclo[7.3.1.0\textsuperscript{5,13}]tridecane |
| b1| Tricyclo[4.3.1.1\textsuperscript{13,16}]undecane | c6| 1-n-Propyladamantane                              |
| b2| Tricyclo[6.3.0.0\textsuperscript{10,16}]undecane | c7| 1-Methyl-3-ethyladamantane                        |
| b3| Tricyclo[6.2.1.0\textsuperscript{1,16}]undecane | c8| 2-Methyl-1-ethyladamantane                        |
| b4| Tricyclo[5.2.2.0\textsuperscript{10,16}]undecane | c9| 2-n-Propyladamantane                              |

Among bicycloalkanes the content of the relict-type structures – the long-chain derivative of perhydroindan – 1-(2-methyl, hexyl)-perhydroindan and polymethyl derivatives of decalin – were determined. Typical structures and corresponding mass-spectra are presented on Figure 7.
In the same concentrate were determined protoadamantanes (the adamantanes predecessor compounds in petroleum) – tricycloundecanes (C\textsubscript{11}), homoadamantane – tricyclo [4,3,1,1\textsuperscript{5,8}] undecane having a seven-member cycle in its structure, tricyclododecanes (C\textsubscript{12}) and perhydrophenalene – tricyclo [7,3,1,0\textsuperscript{5,13}]tridecane (C\textsubscript{13}). The structures of tricycloalkanes with compact structures – polyedrane found in the extract are of particular interest. An efficient separation of these compounds became possible on the above mentioned 200 m capillary column. Typical structures and corresponding mass-spectra are presented on Figure 8.

18 derivatives of adamantane have been identified, among them not only methyl- and ethyl-substituted, but also propyl- and butyl-adamantane were found. Formation of alkyladamantanes in oil is explained by process of isomerization of condensed tricyclic hydrocarbons at contact with aluminosilicate rocks [9]. Formation of adamantane derivatives with long chains (>C\textsubscript{2}) hasn’t been
confirmed with model experiments and their existence in oil raised doubts. Nevertheless, in Taribani oil 1-n- and 2-n-propyladamantanes (peaks c6 and c9, Figure 9), 1-methyl-3-propyl-adamantane (peak d2), 1-n- and 2-n-butyladamantanes (peaks d5 and d4, respectively) were detected that is confirmed with the corresponding mass-spectra for all derivatives the adamantly cation C_{10}H_{15}^+ (m/z 135) is observed, as well as molecular ions C_{12}H_{20}^+ (m/z 164) for ethyl-, C_{13}H_{22}^+ (m/z 178) for propyl-, and C_{14}H_{22}^+ (m/z 192) for butyl-derivatives are visible in spectra compared with the NIST database.

Fig. 9. Mass spectra of (b) 2-Ethyl- (peak a5), 2-Propyl- (peak c9) and 2-Butyladamantanes (peak d4)

Alkyladamantanes with substitute larger than ethyl radical were not found in petroleum prior to our investigation. This could be explained by their formation from tricyclic condensed hydrocarbons by isomerization when entering into contact with aluminosilicate rocks. It was considered that as in such conditions the higher derivatives of adamantane were not formed they could not be present in oils. But 1-n-, 2-n-propyl-, 1-methyl-3-propyl- and 2-n-buthyladamantanes were detected in Taribani oil. The presence of these hydrocarbons is difficult to explain, because their probable predecessors were not detected in petroleum, or it can be proposed that they are products of destruction of higher molecular weight petroleum compounds containing adamantane nucleus.

Conclusions.

On the basis of the applied methodology for investigation of individual composition of C_{11}-C_{16} polycycloalkanes from middle 200-250°C and 250-350°C fractions of Taribani oil naphthenic concentrates with high degree of homogeneity were obtained. Subsequent extraction of the concentrates with thiourea made it possible to obtain mixtures of tri-, tetra- and pentacyclic hydrocarbons with compact bridge structure – carcass compounds of C_{11}-C_{16} composition. In the thiourea extracts more than 50 individual polycyclic alkanes were identified by methods of GC, MS and GC-MS: adamantane and its nine homologues of C_{10}-C_{13} composition; twelve tricyclic C_{11}-C_{13} hydrocarbons (tricyclocdecanes and tricyclododecanes – bridge tricycloalkanes, the predecessors of adamantanes in oil); ten tetracyclic C_{12}-C_{16} hydrocarbons containing adamantane structure in polycyclic system; four pentacyclootetradecanes – diamantane (C_{13}H_{30}) and its three derivatives of C_{13}H_{22} composition; six bicyclic hydrocarbons with sesquiterpene type structure, polymethyl-substituted decalines of C_{14}-C_{16} composition, structure of which have relict character.

It should be noted that at a research of the Taribani oil ten compounds have been detected in oil for the first time: tricyclo [5.2.2.0^{1,5}]undecan (C_{11}H_{18}), 1-n-propyladamantane (C_{13}H_{23}), 1-ethyl-2-methyl-adamantane (C_{13}H_{22}), metyltetracyclo[6.3.1.0^{3,7}]tridecan (C_{14}H_{22}); 1-methyl- and 3-methyl-diamantanes (C_{13}H_{23}), tetracyclo[7.3.1.0^{2,6},1^{10}]tetradecan (C_{14}H_{22}) and its homologous range of tetracyclo[7.3.1.0^{2,6},1^{10}]tetradecan up to hexamethyl-substituted (C_{15}H_{31} – C_{20}H_{35}).

The main components of the VIII thermodiffusion faction, as it was expected, were C_{11}-C_{22} isoprenoid alkanes. Among them the predominant were pristane C_{19} and the phytane C_{20}, the well-
known biomarkers of oil. In the thyourea extract of the VIII thermodiffusion fraction 18 hydrocarbons containing adamantan skeleton, including dimethyl-, trimethyl-, tetramethyl-, ethyl-, methyl-ethyl-, diethyl-, propyl, methyl-ethyl adamantanes were identified.

In Taribani oil the presence of following compounds was established for the first time: 2-n-propyladamantane; 1-methyl,3-propyladamantane and 2-n-butyladamantane.

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