Wax Behavior in Crude Oils by Pour Point Analyses

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Crude oils containing large quantities of waxes, and in situations where sudden temperature drops occur, a crystalline network can be formed, generating flow difficulties in production, transfer and offloading lines. Despite the scientific and economic importance of this phenomenon, correlations between the behavior of petroleum in relation to its pour point are scarce in the literature. In this work, nine crude oil samples were characterized regarding density, water content, wax fraction (one- and two-dimensional chromatography), pour point, yield stress and wax appearance temperature. The results showed that the storage conditions and oil characteristics had a strong influence on the pour point. Yield stress was influenced by initial temperature of the sample, cooling rate and shear rate during cooling. Finally, the content of aromatic compounds in oil with low concentration of \( n \)-alkanes can contribute to increase the pour point.

Keywords: crude oils, pour point, yield stress, gas chromatography, waxes

Introduction

In the reservoir, crude oil behaves as a Newtonian fluid, since the waxes with high molar mass are in solution, causing the oil to act as a monophasic liquid, i.e., in laminar flow, in which the viscosity is only a function of temperature, independent of the shear rate.\(^1\)\(^2\) As soon as the oil leaves the reservoir and flows through the line at a lower temperature, the temperature gradient between the cold wall of the pipe and the hot oil causes the waxes to precipitate. When the temperature of a crude oil containing high quantities of waxes drops suddenly, a crystalline network forms, impeding its flow in production, transfer or offloading lines. The precipitated waxes, when aggregating, can alter the flow, resulting in non-Newtonian behavior of the oil. The presence of wax crystals can also lead to an increase in viscosity, requiring application of a much higher pressure to assure continued flow.\(^3\)\(^-\)\(^9\) Other components present in petroleum, such as polar compounds, resins and asphaltenes, can co-precipitate with the wax crystals, generating sludges that are hard to remove.\(^10\)\(^-\)\(^11\) In general, organic compounds can be used to prevent wax deposition.\(^12\)\(^,\)\(^13\)

The determination of several parameters, such as the wax appearance temperature (WAT), crystallization enthalpy (using microcalorimetry), pour point and yield stress, can help the evaluation of the crystallization and deposition of waxes in crude oil.\(^12\)\(^,\)\(^14\)\(^-\)\(^16\) The WAT is the temperature at which the first wax crystals start to form under cooling. It can be measured by differential scanning calorimetry (DSC), a technique also applicable to determine the variation of enthalpy of the crystallization process. Microcalorimetry is significantly more sensitive,\(^17\) revealing the transition more clearly,
besides allowing observation of other more discrete transitions that cannot be detected by calorimetry. It can also be used for tests under pressure, thus more faithfully reproducing real conditions.\textsuperscript{12,14,18}

Another procedure used to assess wax deposition involves determining the pour point, which is the lowest temperature at which the oil is able to flow only under the action of gravity. A pour point above the ambient temperature is an indication of the presence of large amounts of paraffinic hydrocarbons with long chains, known as waxes. The exception is heavy oils, which have high pour points due to the large quantity of asphaltenes and resins in their composition.\textsuperscript{19-22} Knowledge of the pour point is important because the conditions for pumping, storage and transfer of oil in pipelines and tanks are defined based on this parameter and the viscosity.

The yield stress is one of the properties that helps to describe and to understand the complex behavior of the gelled structure formed when an oil with high paraffin content is cooled under a static condition. The term yield stress denotes the minimum stress that must be applied to a determined material to produce a shear flow. It is a rheological property of petroleum that represents the transition between the elastic solid behavior of a gelled oil and its behavior as a viscous liquid when a production shutdown occurs.\textsuperscript{23-25}

Gas chromatography (GC) has been used to determine petroleum compounds.\textsuperscript{26} However, two-dimensional gas chromatography (GC×GC) has been used to better identify many kind of organic compounds.\textsuperscript{27,28}

In this work, we characterized and quantified the waxes present in crude oil samples using traditional methods, such as one-dimensional gas chromatography, and recently developed chromatographic methods, such as comprehensive two-dimensional gas chromatography, seeking to correlate the distribution of the wax chains with the pour point of the oil samples.

\section*{Experimental}

\subsection*{Materials}

Nine samples of crude oil, called here AP01, AP02, AP03, AP04, AP05, AP06, AP07, AP08 and AP09, were used for the main study. Four crude oil samples, called here SAMG, SAMM, SAMS and SAMU, were used to evaluate two pour point methods (ASTM D97\textsuperscript{29} and ASTM D5853\textsuperscript{30} standards). Nine crude oil samples, named here Petro1, Petro2, Petro3, Petro4, Petro5, Petro6, Petro7, Petro8 and Petro9 (all collected at three different points of primary processing), were used to evaluate the influence of thermal history on the pour point results following ASTM D97.\textsuperscript{29}

All crude oil samples were provided by the Petrobras Research Center (CENPES), Rio de Janeiro, Brazil. 99.5% toluene, supplied by Vetec Química Fina Ltda, Duque de Caxias, Brazil, was distilled at 110 °C,\textsuperscript{31} while 99.9% cyclohexane, 99.9% dichloromethane and 95% \textit{n}-hexane were supplied by Tedia Brasil Ltda., Rio de Janeiro, Brazil. Activated aluminum oxide (150 mesh) and silica gel pore size 30 Å (75-150 µm; 100-200 mesh) were supplied by Sigma-Aldrich, São Paulo, Brazil. Silica gel pore size 60 Å (40-63 µm; 230-400 mesh) and silica gel pore size 60 Å (63-200 µm; 70-230 mesh) were supplied by SiliCycle Inc, Quebec, Canada. Helium (99.9999% purity) was purchased from Linde Gas Ltda., Rio de Janeiro, Brazil. Cholestane-D\textsubscript{6}, dibenzothiophene-D\textsubscript{8}, hexadecane-D\textsubscript{34}, pyrene-D\textsubscript{10}, pristane-D\textsubscript{40} and tetracosane-D\textsubscript{50}, > 97% purity, were purchased from CDN Isotopes, Quebec, Canada.

\subsection*{Density}

In an Anton Paar DMA 4500M densimeter, a small quantity of oil was inserted in a cell (U-shaped tube) with controlled temperature. The oscillation frequency was recorded and the density of the sample was calculated by using cell constants, previously determined by measuring the oscillation frequency of calibration fluids with known densities. In the petroleum industry, the density of crude oils is usually given in °API, since this permits better evaluation of the difference of density between oils. The °API is given by \((141.5/dR) – 131.5\), where dR is the ratio between the specific mass of the oil and the specific mass of water, both measured at 15 °C.\textsuperscript{32}

\subsection*{Water content (Karl Fischer method)}

The water content of the crude oil samples was determined based on the ASTM D4377-00 standard.\textsuperscript{33} A known quantity of sample was added to the titration jar, already containing a solution of chloroform and methanol in proportion of 3:1. Using a specific Karl Fischer reagent for titrant, the quantity of water present in the sample was determined and reported as %m/m. The device used was a Metrohm 841 Titrrando, with double platinum wire electrode, which identifies the presence of water in the medium by measuring conductivity.

\subsection*{Crude oil fractionation}

Oil samples (ca. 200 mg) were fractionated and the saturated fraction was ascertained through liquid chromatography under vacuum, using as stationary phase:
Wax Behavior in Crude Oils by Pour Point Analyses

J. Braz. Chem. Soc.

(i) silica gel pore size 60 Å (40-63 µm; 230-400 mesh);
(ii) silica gel pore size 60 Å (63-200 µm; 70-230 mesh);
(iii) silica gel pore size 30 Å (75-150 µm; 100-200 mesh);
and (iv) activated aluminum oxide (150 mesh). The saturated hydrocarbon fraction was eluted using n-hexane (250 mL). The solvent was evaporated under nitrogen gas flux.34

One-dimensional gas chromatography

The whole crude oil samples were diluted with CS2 at 2% m/m. The reference n-alkane sample was a mixture of n-alkanes in the range C3 to C120, and it was also prepared with CS2. The chromatographic system used was an Agilent Technologies 6890N Network GC system. The injector temperature program was from 50 to 430 °C at 15 °C min−1 and held at 430 °C for 26 min. The GC oven temperature program was: 20 to 430 °C at 10 °C min−1 and held at 430 °C for 5 min. The detector temperature was set at 430 °C. Helium was used as carrier gas at a constant flow rate of 22.5 mL min−1 and the injected volume was 1.0 µL.35

The saturated fractions were analyzed by gas chromatography with flame ionization detection (GC-FID). The injector temperature program was from 100 to 330 °C at 15 °C min−1 and held at 330 °C for 5 min. The GC oven temperature program was 80 to 340 °C at 8 °C min−1 and held at 340 °C for 7 min. The detector was at 340 °C. Helium was used as carrier gas at a constant flow rate of 5 mL min−1, and the injected volume was 1.0 µL. Linear, iso and cycloparaffins were calculated by ASTM D5442.36

Comprehensive two-dimensional chromatography time-of-flight mass spectrometry (GC×GC-TOFMS)

Approximately 10.0 mg of each crude oil sample was weighed using an analytical balance and dissolved with a mixture containing internal standards and dichloromethane. The GC×GC-TOFMS system used was a Pegasus 4D (Leco, St. Joseph, MI, USA), which is an Agilent Technologies 7890 GC (Palo Alto, CA, USA) equipped with a Pegasus H11 time-of-flight mass spectrometer. The GC columns consisted of a DB-17 (Agilent Technologies, Palo Alto, CA, USA) with 50% phenyl-50% methylsiloxane (30 m × 0.25 mm i.d., 0.25 µm df) as the first dimension column (1D) and a DB-5 (Agilent Technologies, Palo Alto, CA, USA) with 5% phenyl-95% methylsiloxane (1.5 m × 0.18 mm i.d., 0.18 µm df) as the second dimension column (2D). The injections were performed in a splitless mode of 0.5 µL at 310 °C using a purge time of 60 s and a purge flow of 5 mL min−1. Helium (99.9999% purity) was used as carrier gas at a constant flow rate of 1.0 mL min−1. The oven temperature program was 40 °C for 5 min and ramped up to 330 °C at 3 °C min−1. The temperature of the secondary oven was 5 °C higher than that of the primary oven. Modulation period was 9 s, with 4.5 s hot-pulse duration and modulator temperature 30 °C higher than the primary oven temperature. The MS transfer line was maintained at 280 °C, and the TOFMS was operated in the electron ionization mode with a collected mass range of m/z 40-650. The ion source temperature was 230 °C, the detector was operated at −1400 V, with electron energy 70 eV, and an acquisition rate of 100 spectra s−1.37 GC×GC-TOFMS data acquisition and processing were performed using the ChromaTOF® software version 4.51 (Leco, St. Joseph, MI, USA). After data acquisition, samples were submitted to a data-processing method in which individual peaks were automatically detected based on a 50:1 signal-to-noise ratio.

Identification was performed by comparing the mass spectra obtained with the NIST Mass Spectral Library software (NIST 08, software version: 2.0) for correct matching, in addition to the retention times. The hydrocarbon classes were analyzed in extracted ion chromatogram (EIC) mode using m/z 85 (n-alkanes), 57 (branched alkanes), 83 (alkyl-cyclohexanes), 69 (alkyl-cyclopentanes), 137 (alkyl-decalins), 191 (tri-, tetra-, and pentacyclic terpanes), 91 (alkylbenzenes), 128 (alkynaphthalenes), 178 (alkyl-phenanthrenes) and 166 (alkyl-9H-fluorenes). The semi-quantification was performed by classifications for each sample based on the delimitation of regions for the main hydrocarbon classes. Total ions chromatogram (TIC) was used to obtain the total area and the classes are divided into n-alkanes, branched alkanes, monocyclic, bicyclic and polycyclic hydrocarbons, alkylbenzenes, alkynaphthalenes, alkylphenanthrenes and alkyl-9H-fluorenes. For each class, a specific internal standard was applied, e.g., n-hexadecane-D14 for the saturated hydrocarbon classes and pyrene-D10 for the aromatic hydrocarbons classes.38

Analyses of pour point

The tests were performed according to ASTM D97,29 with some adaptations. The determination of pour point is basically a manual method involving the use of a thermometer to read the system’s temperature and check the fluidity of the system after every temperature reduction of 3 °C. The no-flow temperature is recorded when the system ceases flowing when being observed in the horizontal position for 5 s. The flow point was reported as the no-flow temperature plus 3 °C.
For comparison of the results, the ASTM D5853 standard was also used. In this standard, the pour point is represented by a temperature range. Two measures are obtained, the maximum pour point and the minimum pour point. The maximum pour point (determined in a procedure where crystallization of the waxes is encouraged by applying lateral friction of the test tube with a glass rod before starting the test) was measured as described above in ASTM D97, while the minimum pour point (with a procedure that discourages crystallization) was determined by submitting the sample to initial thermal treatment, consisting of heating in a pressure cell at 105 ± 2 °C for 1 h followed by cooling to a temperature above the pour point before analysis. The test procedure was the same as that previously described for ASTM D97. The readings were performed at temperature intervals of 3 °C and the value obtained was increased by 3 °C.

Determination of wax appearance temperature (WAT)

The microcalorimetry tests were conducted in a Setaram µDSCIII instrument, using a Hastelloy C276 cell. The tests were performed with approximately 400 mg of sample and undecane as reference. The initial crystallization temperature was obtained with tests having the following program: heating of the samples from room temperature to 80 °C at a heating rate of 1 °C min⁻¹; holding the temperature at 80 °C for 15 min; cooling to −10 °C at a cooling rate of 0.48 °C min⁻¹; holding the temperature at −10 °C for 15 min; and heating to 30 °C at a heating rate of 1 °C min⁻¹. Based on the curve of heat flow as a function of temperature, the WAT was determined as the onset temperature of the first exothermic peak.

Yield stress

The yield stress was measured with a Haake Mars III rheometer coupled to a microcomputer and a Thermo Haake bath, with a set of parallel plate sensors with diameter of 35 mm, using oscillatory rheology. This set of parallel plates (PP35) was positioned with spacing of 1.2 mm. A 2.0 mL aliquot of previously heated crude oil was injected between the sensors using a syringe with steel needle at the initial testing temperature. After positioning the sample, the set of sensors was covered with a split cover to reduce evaporation of the light fractions during the test. The sample, previously heated for 1 h at 80 °C, was kept heated under stirring at a temperature of 60 °C and submitted to the standard test in the rheometer, as follows: pre-shear of the sample at 45 °C for 15 min at a rate of 10 s⁻¹; cooling of the sample from 45 to 4 °C, under static conditions, at a cooling rate of 0.8 °C min⁻¹; positioning the sensor at the standard distance for analysis (1 mm) with an approach velocity of 0.04 mm min⁻¹; stabilization of the sample for 15 min at 4 °C; application of rising stress steps in oscillatory mode (logarithmic increment from 10⁴ to 10⁶ Pa with duration of 1 h, 120 steps per decade, and frequency of 1 Hz). Since the paraffinic gel formed can present properties similar to those of a solid, i.e., elastically deform when submitted to low shear stress, it will only start flowing when the plastic deformation caused by mechanical stress is greater than the yield stress. By considering a balance of forces, it can be demonstrated that the minimum pressure (ΔPₘᵡᵣ) able to start the flow of a gelled petroleum sample in the tube is given by equation 1:

\[ ΔPₘᵡᵣ = \frac{4L}{D} τ₀ \]  

where L is the length and D the diameter of the tube, and τ₀ is the yield stress.

Results and Discussion

Characterization and evaluation of the crude oil samples

The nine oil samples were characterized regarding density, water content, pour point, wax appearance temperature (WAT), yield stress, mass of saturates, content of n-alkanes, isoalkanes + cycloalkanes (UCM) by one-dimensional gas chromatography (GC) in the C₅–C₄₀ range; contents of branched alkanes, cycloalkanes and some aromatic compounds in relevant abundance, by two-dimensional gas chromatography (GC×GC-TOFMS) in the C₅–C₄₀ range. The samples were also classified according to the Kᵤₒᵖ characterization factor. This is a factor proposed by Universal Oil Products (UOP) to classify petroleum according to its nature, as paraffinic (Kᵤₒᵖ ≥ 12.5), naphthenic (Kᵤₒᵖ < 11.5), intermediate (11.5 ≤ Kᵤₒᵖ ≤ 12.1) or aromatic (Kᵤₒᵖ ≤ 10). Kᵤₒᵖ = (BT)³/d, where BT is the mean molar boiling temperature in degrees Rankine [(°C + 273.15) × 95] and d is the 15/15 °C density. The data are compiled in Table 1. The results show that the oil samples are relatively distinct, as indicated by the different density values (°API) found.

With respect to yield stress, this represents the solid-liquid transition of the structured material and is determined by the intersection of the G’ (elastic modulus) and G” (viscous modulus) moduli in the graph of these parameters in function of the shear stress applied. In oscillatory rheology, when a variable stress is applied to the sample, gelled fluids have values of G’ higher than G” at low stresses, showing a predominantly solid
### Table 1. Characterization data of the crude oil samples

| Sample | AP01 | AP02 | AP03 | AP04 | AP05 | AP06 | AP07 | AP08 | AP09 |
|--------|------|------|------|------|------|------|------|------|------|
| °API   | 27   | 30   | 27.4 | 31   | 17.3 | 14.8 | 36   | 28.9 | 23   |
| Karl Fischer / %m/m | 0.05 | 0.03 | 0.50 | 0.03 | 0.60 | 0.09 | 0.45 | 0.04 | 0.01 |
| Characterization factor K_{crp} | 11.8 | 12.0 | 11.8 | 12.0 | 11.6 | 11.7 | > 12.6 | 11.9 | 11.9 |
| Yield stress at 4 °C / Pa | 7.2  | 36.6 | 0.7  | 1.7  | 0    | 0    | > 10,000 | 0   | 4.2  |
| ASTM D9738 pour point / °C | -6   | 12   | -24  | 6    | -6   | 3    | 36   | -15  | 24   |
| solidification / °C | -9   | 9    | -24  | 3    | -9   | 0    | 33   | -18  | 21   |
| μDSC: 1st WAT / °C | 39.6 | 35.7 | 16.1 | 33.6 | 19.9 | 19.7 | 55.2 | 40.5 | 40.5 |
| 2nd WAT / °C | 20.1 | 22.0 | -    | 21.6 | -    | -    | 41.2 | 18.6 | 18.6 |
| Mass of saturated fraction | 87.23 | 101.12 | 74.33 | 85.71 | 103.06 | 76.17 | 102.88 | 69.91 | 77.24 |
| WAT / °C | 2162 |
| n-C_{15} | 0.08 | 0.19 | 0.05 | 0.08 | 0.00 | 0.00 | 0.19 | 0.05 | 0.09 |
| n-C_{16} | 0.19 | 0.28 | 0.17 | 0.19 | 0.00 | 0.00 | 0.48 | 0.14 | 0.17 |
| n-C_{17} | 0.24 | 0.34 | 0.23 | 0.30 | 0.00 | 0.00 | 0.69 | 0.20 | 0.28 |
| n-C_{18} | 0.23 | 0.28 | 0.24 | 0.28 | 0.00 | 0.00 | 0.79 | 0.23 | 0.26 |
| n-C_{19} | 0.32 | 0.34 | 0.30 | 0.34 | 0.00 | 0.00 | 0.98 | 0.28 | 0.33 |
| n-C_{20} | 0.25 | 0.23 | 0.23 | 0.24 | 0.00 | 0.00 | 0.90 | 0.23 | 0.25 |
| n-C_{21} | 0.20 | 0.22 | 0.17 | 0.22 | 0.00 | 0.00 | 0.94 | 0.17 | 0.23 |
| n-C_{22} | 0.19 | 0.21 | 0.15 | 0.22 | 0.00 | 0.00 | 0.91 | 0.17 | 0.25 |
| n-C_{23} | 0.20 | 0.20 | 0.15 | 0.21 | 0.00 | 0.00 | 0.94 | 0.17 | 0.23 |
| n-C_{24} | 0.18 | 0.20 | 0.13 | 0.21 | 0.00 | 0.00 | 0.87 | 0.15 | 0.23 |
| n-C_{25} | 0.19 | 0.20 | 0.14 | 0.20 | 0.00 | 0.00 | 0.95 | 0.17 | 0.24 |
| n-C_{26} | 0.17 | 0.20 | 0.12 | 0.21 | 0.00 | 0.00 | 0.83 | 0.16 | 0.25 |
| n-C_{27} | 0.15 | 0.18 | 0.11 | 0.19 | 0.00 | 0.00 | 0.81 | 0.14 | 0.23 |
| n-C_{28} | 0.17 | 0.18 | 0.13 | 0.18 | 0.00 | 0.00 | 0.71 | 0.18 | 0.23 |
| n-C_{29} | 0.12 | 0.16 | 0.09 | 0.16 | 0.00 | 0.00 | 0.67 | 0.11 | 0.18 |
| n-C_{30} | 0.08 | 0.12 | 0.06 | 0.13 | 0.00 | 0.00 | 0.53 | 0.07 | 0.14 |
| n-C_{31} | 0.05 | 0.11 | 0.04 | 0.11 | 0.00 | 0.00 | 0.45 | 0.03 | 0.12 |
| n-C_{32} | 0.04 | 0.08 | 0.03 | 0.08 | 0.00 | 0.00 | 0.34 | 0.03 | 0.09 |
| n-C_{33} | 0.03 | 0.06 | 0.03 | 0.06 | 0.00 | 0.00 | 0.23 | 0.03 | 0.06 |
| n-C_{34} | 0.02 | 0.05 | 0.02 | 0.05 | 0.00 | 0.00 | 0.19 | 0.02 | 0.05 |
| n-C_{35} | 0.02 | 0.03 | 0.02 | 0.03 | 0.00 | 0.00 | 0.14 | 0.02 | 0.03 |
| n-C_{36} | 0.01 | 0.02 | 0.01 | 0.02 | 0.00 | 0.00 | 0.11 | 0.01 | 0.02 |
| n-C_{37} | 0.01 | 0.01 | 0.01 | 0.02 | 0.00 | 0.00 | 0.06 | 0.01 | 0.02 |
| n-C_{38} | 0.01 | 0.01 | 0.01 | 0.01 | 0.00 | 0.00 | 0.06 | 0.01 | 0.01 |
| n-C_{39} | 0.00 | 0.01 | 0.00 | 0.00 | 0.00 | 0.00 | 0.04 | 0.00 | 0.01 |
| n-C_{40} | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.03 | 0.00 | 0.00 |
| n-C_{41} | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.04 | 0.00 | 0.00 |
| n-C_{42} | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.02 | 0.00 | 0.01 |
| n-C_{43} | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.02 | 0.00 | 0.00 |
| n-C_{44} | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.00 | 0.01 | 0.00 | 0.00 |
| total n-alkanes | 3.15 | 3.93 | 2.64 | 3.75 | 0.00 | 0.00 | 13.95 | 2.80 | 4.03 |
| UCM (iso + cycloalkanes) | 34.20 | 30.26 | 29.38 | 29.05 | 23.41 | 24.62 | 26.11 | 30.08 | 23.60 |

- **GCxGC-TOFMS**: two-dimensional gas chromatography time-of-flight mass spectrometry.
- **DSC**: differential scanning calorimetry; **WAT**: wax appearance temperature; **UCM**: unresolved complex mixture; **GCxGC-TOFMS**: two-dimensional gas chromatography.

Other characterization data: Mass of saturated fraction (range from 254 to 550 °C, n-C_{15} to n-C_{25}) / %m/m.
characteristic. However, this order is altered beyond a critical stress value, and the fluid starts to behave more like a liquid.\textsuperscript{39}

The rheogram of sample AP01 in the oscillatory rheometry test, shown in Figure 1, depicts the behavior of the \( G' \) and \( G'' \) curves in function of the oscillatory shear stress. When \( G' \) is higher than \( G'' \) at low stress, the sample behaves as a solid. Then, the value of \( G' \) declines (in this case, \( G'' \) only starts to decline after the two moduli cross) and, finally, \( G' \) and \( G'' \) cross at 7.2 Pa, which represents the solid-liquid transition of the materials submitted to oscillatory forces. When \( G' \) is below \( G'' \), it means that viscous behavior predominates in relation to elastic behavior and the sample does not present yield stress. The yield stress values of all the crude oil samples are presented in Table 1.

With the exception of sample AP07, all the other crude oils had low yield stress values. The yield stress (\( \tau_y \)) value obtained for sample AP07 is considered to be high because according to equation 1, depending on the ratio between the length (L) and diameter (D) of the tube, it can require high pressure values (\( \Delta P_{\text{min}} \)) to assure restart of production after prolonged shutdowns, when the oil typically reaches temperature of 15 °C. In general, oils with high pour point values tend to have higher yield stress values. This was observed for sample AP07, but not for any of the other samples. This can be explained by the fact that both the pour point and yield stress measures are influenced by the initial sample temperature, thermal gradient and cooling rate, among other factors, and the difference in procedure likely caused this small difference in behavior.

To confirm that assumption, we performed another oscillatory rheometry test to measure the yield stress of samples AP02, AP04, AP07 and AP09. In this new test, the initial temperature was equal to that of the pour point test, 45 °C, without previous thermal treatment at 80 °C. The yield stress results, reported in Table 2, showed that when the test was conducted under the same conditions as the pour point test (initial temperature of 45 °C as indicated by ASTM D97),\textsuperscript{29} the results were more coherent, i.e., the greater the pour point, the higher was the yield stress.

| Oil sample | Pour point (±3 °C) / °C | Yield stress at 4 °C (\( T_i = 80 \) °C) / Pa | Yield stress at 4 °C (\( T_i = 45 \) °C) / Pa |
|------------|------------------------|---------------------------------|---------------------------------|
| AP02       | 6                      | 36.6                            | 367.7                           |
| AP04       | 12                     | 1.7                             | 672.8                           |
| AP07       | 36                     | >10,000                         | >10,000                         |
| AP09       | 24                     | 4.2                             | 3,368                           |

To eliminate any suspicion that the difference between the pour points of the samples was related to some difference in the testing conditions, we evaluated the precision of the methods of determining the pour point and the influence of the thermal treatments on the pour point test result. In a first test, we selected at random four of the oil samples used in this work and performed the pour point tests (ASTM D97\textsuperscript{29} and ASTM D5853\textsuperscript{30} standards) with four replicates by two technicians, to assess the precision of the laboratory test. The results obtained by each technician are presented in Table 3 separated by “/”.

| Crude sample | Measure | ASTM D97 | ASTM D5853 max | ASTM D5853 min |
|--------------|---------|----------|-----------------|----------------|
| SAMG         | 1\textsuperscript{st} | -36/–30 | -30/–27 | -45/–48 |
|              | 2\textsuperscript{nd} | -33/–30 | -36/–24 | -48/–48 |
|              | 3\textsuperscript{rd} | -36/–30 | -33/–27 | -48/–48 |
|              | 4\textsuperscript{th} | -36/–30 | -36/–27 | -45/–51 |
| SAML         | 1\textsuperscript{st} | -24/–24 | -24/–24 | -42/–24 |
|              | 2\textsuperscript{nd} | -24/–24 | -24/–24 | -33/–24 |
|              | 3\textsuperscript{rd} | -27/–24 | -24/–24 | -33/–24 |
|              | 4\textsuperscript{th} | -24/–24 | -27/–24 | -30/–24 |
| SAMS         | 1\textsuperscript{st} | 18/15   | 15/15          | -15/–12        |
|              | 2\textsuperscript{nd} | 15/15   | 15/15          | -12/–12        |
|              | 3\textsuperscript{rd} | 15/15   | 15/15          | -15/–12        |
|              | 4\textsuperscript{th} | 18/15   | 15/15          | -15/–12        |
| SAMU         | 1\textsuperscript{st} | 3/0     | -3/-3          | -15/-18        |
|              | 2\textsuperscript{nd} | 3/0     | 0/-3           | -12/-18        |
|              | 3\textsuperscript{rd} | 3/0     | 0/-3           | -15/-18        |
|              | 4\textsuperscript{th} | 3/0     | 0/-3           | -15/-18        |

\textsuperscript{a}Test not performed due to insufficient quantity of sample.
According to results obtained for each sample by each method, the maximum difference between the repetitions was 6 °C, with the majority being 3 °C at most. These values are within the repeatability ranges of the methods used. Between the two test methods, the pour point value obtained by ASTM D97, in most cases, was near the maximum value obtained by ASTM D5853, while the minimum value was always substantially below the others, as is inherent to the method.

The differences in the values found by the two technicians were within the intermediate precision range expected for the method and can be considered not significant. Probably the differences in the pour point results found for the crude oil samples studied are not related to fluctuation of the laboratory readings. To assure the reliability of the data generated, we decided to conduct all the pour point tests of the samples in replicate.

After eliminating the hypothesis of repetitiveness problems of the method, we assessed the influence of the thermal history on the pour point results. For that purpose, we carried out a thermal sensitivity test, in which the nine oil samples used (Petro1, Petro2, Petro3, Petro4, Petro5, Petro6, Petro7, Petro8 and Petro9) had been collected at three different points of primary processing (arrival at the platform, exit from the treatment unit and offloading (transfer of the oil from the production platform/vessel to a relief ship)). The results are shown in Table 4.

The pour point of the samples varied according to the temperature to which they were submitted, both in primary processing and in the laboratory simulation. The samples

**Table 4. Pour point (ASTM D97) for 27 crude oil samples from 9 production systems**

| Crude oil | Collect point | Pour point / °C | ± 3 °C | Cooled at 4 °C for 48 h (± 3 °C) | Cooled at 4 °C for 7 days (± 3 °C) | Heated to 60 °C (± 3 °C) | Heated to 105 °C (± 3 °C) |
|-----------|---------------|-----------------|--------|----------------------------------|-------------------------------|------------------------|----------------------------|
|           | A             |                 |        |                                  |                               |                        |                            |
| Petro1    | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro2    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro3    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro4    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro5    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro6    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro7    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro8    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |
| Petro9    | A             |                 |        |                                  |                               |                        |                            |
|           | B             |                 |        |                                  |                               |                        |                            |
|           | C             |                 |        |                                  |                               |                        |                            |

A: just arriving to the platform; B: close to electrostatic treatment; C: at the offloading.
that underwent electrostatic treatment were subjected to a temperature that varied from 40 to 90 °C, and in the majority of the cases the samples before treatment presented lower pour points than those that arrived on the platform, as expected. Samples Petro4, Petro6 and Petro7 presented anomalous behavior when comparing the pour point results of each of these systems. The samples from offloading of each system were subjected to prior heating and cooling processes to check the influence of temperature on the pour point. To better visualize the behaviors, the results of the offloading samples are organized in a bar graph (Figure 2). When the samples were heated before the pour point test, the pour point declined in the majority of cases. We can speculate that the rearrangement of the systems due to dissolution of the waxes present in them caused a reduction of the pour point. The majorities of the samples that were submitted to cooling before the test showed higher pour point. When the samples were submitted to cooling before the test, in general, the obtained pour point was higher, and the longer the sample remained at low temperature, the stronger this effect was. We observed, therefore, that leaving the samples at rest for 24 h to erase the thermal history, as specified in ASTM D9729 and ASTM D5853,30 was not sufficient for these types of samples.

Thus, the pour points of the crude oils studied were significantly influenced by the thermal variations to which they were submitted, explaining the different pour point values found for samples from the same stream (system). We can suppose that this behavior was related to the composition and structure of the wax crystals formed due to cooling of the samples. The samples obtained directly from the reservoir formation, without any thermal treatment, when having waxes in the solid state, were amorphous, while those that had undergone temperature variations presented crystalline structures. These crystalline structures, once formed, if not totally melted served as nucleation points for the growth of new crystals, favoring loss of mobility of the sample. Because of this fact, all the samples analyzed were stored at room temperature, taking care not to heat them before the test or to add them under refrigeration, so that the pour point measured was not influenced by this factor.

The content of paraffins (Table 1) and the distribution profile of the saturated compounds (n-alkanes, isoalkanes and cycloalkanes) were determined by one-dimensional gas chromatography. Figure 3 shows the chromatographic profiles of samples AP02, AP05 and AP07, revealing a difference in the distribution of n-paraffins and a more pronounced unresolved complex mixture (UCM) in some samples than others.

Sample AP07 was classified as a paraffinic crude oil (K_UOP factor > 12.6), containing a quantity of n-paraffins in the C_{15}-C_{44} range greater than the other samples. In this sample it was also possible to identify the presence of n-paraffins with larger chain length than C_{37}, which were not identified or were present only in very small quantities in the other samples.

Samples AP02, AP04 and AP09, considered to be intermediate (K_UOP factors of 12.0, 12.0 and 11.9, respectively), presented a certain similarity in the distribution of n-alkanes. For n-alkanes greater than C_{20}, sample AP09 presented slightly higher values than AP02 and AP04. The UCM of AP07 was slightly lower than that of AP01, AP02, AP03, AP04 and AP08, but higher than that of AP05, AP06 and AP09. This UCM was found to be composed predominantly of branched and cyclic saturated compounds. Samples AP05 and AP06 were considered biodegraded, with virtually insignificant presence of.

![Figure 2](image2.png)

**Figure 2.** Pour points (ASTM D97) of different crude oil samples submitted to different thermal histories.

![Figure 3](image3.png)

**Figure 3.** Wax distribution of the crude oils (a) AP02; (b) AP05 and (c) AP07. Black: linear alkanes; gray: iso and cycloalkanes.
$n$-paraffins in the $C_{15}-C_{44}$ range. The chromatographic profiles of samples AP01, AP02, AP03, AP04, AP08 and AP09 were relatively similar to each other.

Samples AP02, AP04 and AP09 were more strongly influenced by the prior temperature in the pour point and yield stress results than AP07. We believe this behavior can be attributed to the fact that the ratio of $n$-alkanes to UCM (iso- and cycloalkanes) was approximately three times higher in crude oil AP07 ($n$-alkane/UCM = 0.534) than in oils AP02, AP04 and AP09, where the $n$-alkane/UCM ratios were 0.130, 0.129 and 0.171, respectively (Table 1). Furthermore, AP07 had a much higher content of linear paraffins than the others. Since the structure of linear molecules, such as $n$-paraffins, can favor packing, the formation of a crystalline network is favored any time the temperature declines. With linear molecules, this behavior can be repeated more easily, irrespective of the number of times the temperature is increased and then decreased, causing the flow to stop. In contrast, the structure of nonlinear molecules (iso- and cycloalkanes) can interfere in this packing and hamper the formation of a crystalline network, which can explain the variation of the pour point ($3 \, ^\circ C$ above the no-flow temperature), depending on how these molecules are arranged during the temperature drop.

Since only four of the nine crude oil samples studied presented relatively high pour points (Table 1), and also considering the complexity of the GC×GC-TOFMS technique, we only characterized samples AP02, AP04, AP07 and AP09 by this method. The samples were analyzed through their total ion chromatograms (TIC) and extracted ion chromatograms (EIC). The classes of saturated compounds that cannot be identified by one-dimensional chromatography (since they elute together) as well as some aromatic compounds could be identified. It is important to mention that by this method, the ion utilized to obtain the EIC is not always the base peak in the mass spectrum, but rather is a characteristic ion for a given family. The ions for determining the EIC were selected to obtain a better visualization of the classes.

The qualitative and semi-quantitative data obtained for samples AP02, AP04, AP07 and AP09, for the classes of compounds that could be identified (isoalkanes, cycloalkanes and some aromatic compounds), were to a certain extent coherent with the data obtained by one-dimensional chromatography regarding pour point and yield stress values. Figure 4 graphically compares the contents of $n$-alkanes identified by the GC-FID technique in the $C_{15}-C_{44}$ range and the contents of isoalkanes, cycloalkanes and aromatic compounds that could be identified by GC×GC-TOFMS in the $C_{6}-C_{40}$ range.

The results obtained by one-dimensional chromatography along with the data obtained by two-dimensional chromatography (Figure 4) revealed that crude oil AP07 had the highest content of $n$-alkanes present in the $C_{15}-C_{44}$ range (GC-FID) (13.95% m/m), a level much higher than the other crude oils analyzed (4.03, 3.93 and 3.75% m/m for samples AP09, AP02 and AP04, respectively), in line with its very high pour point (36 °C) and greater yield stress (> 10,000 Pa). Sample AP09 presented a slightly higher content of $n$-alkanes than AP02 and AP04, agreeing with the pour point and yield stress results, which were higher for AP09 than for AP02 and AP04.

Sample AP07 also contained the largest quantity of branched alkanes (isoalkanes) in the $C_{4}-C_{40}$ range (GC×GC-TOFMS), with 22.95% m/m, although this was not so different than AP09 (20.99% m/m) and AP04 (20.15% m/m). Sample AP02 had the lowest content of isoalkanes (16.40% m/m). Comparison of the UCM (iso- and cycloparaffins) determined by one-dimensional chromatography with the sum of the isoalkanes and cycloalkanes obtained by two-dimensional chromatography revealed results that were relatively close: for AP02, 30.26 (1D) and 31.20% m/m (2D); for AP04, 29.05 (1D) and 35.02% m/m (2D); for AP07, 26.01 (1D) and 31.29% m/m (2D); and for AP09, 23.60 (1D) and 29.23% m/m (2D). Although one-dimensional chromatography cannot distinguish cyclical from branched compounds, it still produces values having a certain correspondence with those detected by two-dimensional chromatography. The second technique presented slightly higher contents, which can be explained by its greater detection range ($C_{2}-C_{40}$) than that of the one-dimensional method ($C_{4}-C_{40}$). In turn, regarding the levels of naphthenic compounds (cycloalkanes), while oils AP02 and AP04 presented total cycloalkane contents that were close to each other (14.80 and 14.87% m/m), with
predominance of compounds with only one ring (Figure 4), samples AP07 and AP09 contained lower, but also similar, contents (8.34 and 8.24% m/m, respectively), as well as similar distribution by number of rings present (Figure 5). These findings suggest that the higher pour points of these two samples (36 and 24 °C, respectively) are also related to the lower levels of naphthenic compounds present.

Samples AP02 and AP09 were found to have a greater content of aromatic compounds in the C₆-C₄₀ range (alkylbenzenes, alkyl-naphthalenes, alkyl-phenanthrenes and alkyl-9H-fluorenes) than the other two crude oils, with a significant contribution from the alkylbenzene compounds (9.61 and 14.78% m/m for AP02 and AP09, respectively) (Figure 6). Samples AP04 and AP07 had content of aromatics in a narrower range of values and also with predominance of alkylbenzenes (3.58 and 4.49% m/m, respectively). The level of aromatic compounds in AP09 was substantially higher than in the others, suggesting that the high pour point of this oil, despite the lower level of n-alkanes than in AP07, is related to the presence of aromatic rings. These rings, due to their intermolecular forces, favor gel formation, diminishing the mobility of the system and consequently increasing the pour point. Similar reasoning can be applied to AP02, also with a higher content of aromatics than the other two crude oils (AP04 and AP07), but lower than AP09, for this reason having a higher pour point, but one lower than AP09, even with the lower content of n-alkanes than in AP07 (Table 1).

Analysis of the distributions of compounds by families based on the combined data generated by the GC-FID and GC×GC-TOFMS techniques, as shown in Figures 4, 5 and 6, reveals that samples AP02, AP04, AP07 and AP09 contain distinct contents of saturated compounds and aromatic compounds, elucidating the pour point behavior of these crude oils.

Conclusions

With respect to the pour point measurement, the results show that when the crude oil samples were submitted to higher temperatures before the tests, the pour point tended to diminish, while the opposite occurred when the samples were submitted to lower temperatures before testing. This indicates the need for careful standardization of the sampling and storage conditions.

The yield stress and pour point were both influenced by the initial sample temperature, thermal gradient, cooling rate and shear rate during cooling. In general, oils with high pour point can be expected to have higher yield stress values, but this relationship is only indicative, not direct. When the initial temperature of the oscillatory rheometry and pour point tests were similar, i.e., without prior treatment at 80 °C, according to ASTM D97, the yield stress and pour point results were coherent, namely that the higher the pour point was, the greater was the yield stress.

The understanding of the behavior of the crude oils’ pour point in response to variations in processing, transport and storage conditions was improved by characterization of the waxes contained in the samples by chromatographic techniques. The high level of aromatic compounds in the crude oil samples with low content of n-alkanes can explain their higher pour point: the presence of aromatic rings, because of the intermolecular forces they exert, can favor gel formation, reducing the system’s mobility and hence increasing the pour point.

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