

# Organic and Isotope Geochemistry of Hybrid Hydrocarbon Accumulations

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# Summary

Shales are the most common type of sedimentary rock in the Western Canadian Sedimentary Basin (WCSB). The development and production of shale gas in this region, however, do not have a long history. The Lower Triassic Montney Formation hosted the first application of horizontal drilling and multi-stage hydraulic fracturing only in 2004, and its hydrocarbon resources represent 145 years of Canada's 2012 natural gas consumption. The significance of this play relies not only on favorable reservoir properties but also on the high natural gas liquid  $(C_2-C_4)$  and condensate  $(C_5-C_{12})$  content. The condensate contributes with a significant value to well production, which thus increases our interest in understanding its spatial distribution and abundance through the formation.

Lithofacies distribution in the Montney Formation is very complex, and often leads to the occurrence of closed and semi-closed systems where hydrocarbons have been generated or otherwise accommodated after migration, and thermally degraded. Such conditions resulted in isotope anomalies in  $C_1$ - $C_3$  alkanes after thermal cracking of heavier molecules. We have investigated these kinetic isotope effects (sometimes normal distribution, sometimes reversals) in order to define reservoir compartments, methane intraformational leakage, thermal maturation stage of the rock, and fluids mixing. We additionally present a preliminary interpretation on the molecular composition of liquid hydrocarbons from Montney reservoirs.

## **Methods**

To attempt a regional survey, the carbon isotopic compositions of methane, ethane and propane in fluids from the Montney Formation, as reported in different sources (e.g. Desrocher, 1997; Tilley and Muehlenbachs, 2006; Sereda, 2017; this study), has been combined. The obtained δ<sup>13</sup>C values usually correspond to produced gases from wells drilled in Alberta and British Columbia. When produced gas was not available, mud-gas data has been included. In our investigation, 48 gas samples (IsoTubes) from a well drilled in Northeast British Columbia, were also analyzed and results were added to the database. For the molecular analysis, a 5mL aliquot of gas was removed from each IsoTube via a modified IsoTube septum valve and locking gas-tight syringe. Gas compositional analysis (H2, He, N2, O2, Ar, CO2 and alkanes C<sub>1</sub> to C<sub>6</sub>) was completed by injecting the sample into a Scion 450/456 gas chromatograph (GC). The GC utilizes four separate analytical columns connected to three thermal conductivity detectors and a flame-ionization detector for gas separation and quantification. The lower detection limit for hydrocarbons is 1 ppm and for non-hydrocarbon gasses is 50 ppm. Certified gas standards were used to calibrate the GC immediately prior to the analysis. Analytical drift was monitored by injecting the appropriate gas standards after every ten samples analyzed. Analytical precision and accuracy for gas composition analysis is typically better than  $\pm 2.5\%$  of the reported concentrations.



The  $C_1$  to  $C_3$  alkanes compound specific  $^{13}\text{C}/^{12}\text{C}$  ratios of the gas samples were determined using continuous flow technology. The system is comprised of a Thermo Trace GC – GC-IsoLink system interfaced to a Thermo 253 mass spectrometer via a Thermo Conflo IV. Gas sampling was conducted as indicated above for GC compositional analysis. Aliquots of gas were injected into a helium carrier stream through the inlet of the GC using a gas tight syringe. Air,  $CO_2$  and the hydrocarbon compounds of the gas are separated on a GC column before passing through a high temperature, combustion reactor (maintained at 1030 °C). All hydrocarbon gas species are quantitatively converted to  $CO_2$  in passing through the combustion furnace. The separate  $CO_2$  gas pulses are then swept sequentially, by the carrier gas, through a water trap (Nafion®) then into the open split interface which 'leaks' the gas into the mass spectrometer. The  $\delta^{13}$ C values of the unknown species were calculated by the instrument software (ISODAT 3.54). Results are expressed in the usual delta notation in per mil (‰) relative to the international V-PDB standard.

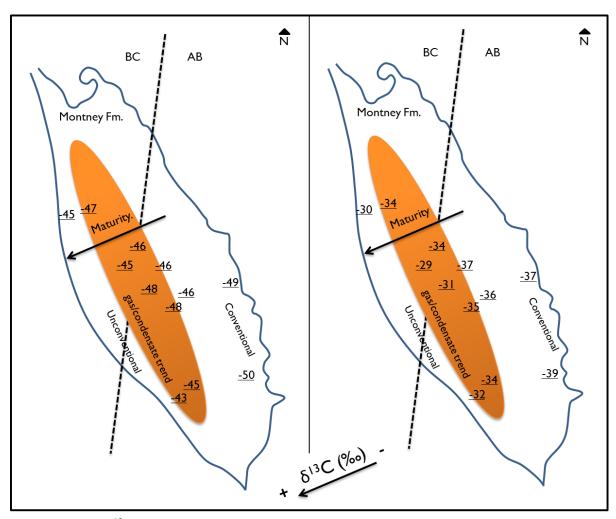
The same instrument was used for  $\delta^2 H$  measurements but using a high temperature reactor (maintained at 1420 °C) instead of a combustion reactor. All hydrocarbon gas species are quantitatively converted to  $H_2$  in passing through the high-temperature-conversion (HTC) furnace. The separate  $H_2$  gas pulses are then swept sequentially, by the carrier gas, through a water trap (Nafion®) then into the open split interface which 'leaks' the gas into the mass spectrometer. The  $\delta^2 H$  values of the unknown species were calculated by the instrument software (ISODAT 3.54), and the results are expressed in the usual delta notation in per mil (‰) relative to the international V-SMOW standard.

# Results

The conducted survey revealed that the  $\delta^{13}C$  values of  $C_1$ - $C_3$  natural gas from the Montney Formation follow, in general, its thermal maturity trend (trend as established by BC Oil and Gas Commission, 2012; Rokosh *et al.*, 2012). This evidence is more effectively expressed in the  $\delta^{13}C$  of ethane (and also propane) compared to methane (Figure 1). A probable explanation for this scenario may rely on the possibility of methane migration into different reservoirs. An excess of methane had already been estimated by Woods and Sanei (2016) to explain anomalous wetness values and suggest migration pathways.

Our preliminary results from a well drilled in Northeast British Columbia, suggest a low maturity hybrid system where little methane migration is expected. Key compositional and isotopic proxies (e.g. iso-alkane ratios, gas wetness,  $\delta^{13}C$  and  $\delta^{2}H$ ) are consistent with depth-increasing maturity.

The  $C_1$ - $C_3$  natural gas from the Montney Formation is isotopically normal although very low carbon isotope ratios for ethane ( $\delta^{13}C \approx 42\%$ ) have been detected in certain wells. The low  $\delta^{13}C$  values of ethane probably occur in localized closed systems caused by facies variations within lithologies.



**Figure 1**. The  $\delta^{13}$ C trend (in ‰) for methane (left) and ethane (right) gases from the Montney Formation (Desrocher, 1997; Tilley and Muehlenbachs, 2006; Sereda, 2017; this study)

## **Novel Information**

For the first time, a preliminary organic geochemistry screening of liquid hydrocarbons (condensate) is reported. Produced condensate from the Montney formation in Northeast British Columbia shows a normal distribution with molar discontinuity at  $C_8$  and  $C_{13}$  alkanes. We currently investigate the source and reservoir conditions for the formation of these fluids, which can potentially address condensate distribution and abundance in the Lower Triassic Montney reservoirs.

## Conclusion

Our understanding of fluid dynamics in the Montney Formation is a fundamental tool for the evaluation of similar hybrid systems such as the Duvernay Formation, also in the WCSB. The isotopic evidence of closed systems in Upper Triassic and Jurassic strata also increases the



probability of fluids yet to be located within those sequences. Ultimately, characterizing the isotope fingerprint of hydrocarbons in hybrid plays from the WCSB is also an essential prerequisite to address environmental concerns about potential thermogenic gas migrating into the shallow hydrosphere.

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