# Alternative Methods to RSK 175 Using Purge and Trap Concentration and Automated Headspace for the Analysis of Dissolved Gases in Drinking Water By Nate Valentine and Tom Hartlein

#### **Abstract**

With the growing price of oil, many alternative energy sources are being explored. Natural gas prices are actually decreasing, in large part to the use of hydraulic fracturing in areas like the Marcellus Shale in Pennsylvania. Due to increased concern over the hydraulic fracturing process and the release of methane and other chemicals into the local drinking water, a need has developed for fast and accurate analysis of methane in water. This study will evaluate methods developed for the determination of methane, ethane, ethene, and propane in water using a Purge and Trap concentrator and automated headspace analysis with GC/FID. Calibration curves, method detection limits (MDLs), and carryover data will be presented and comparisons between the methods will be made.

## Introduction

Hydraulic fracturing, also known as "fracking", is a drilling process currently used to recover natural gas from sources like coalbeds and shale formations. This involves injecting large amounts of water, mixed with sand and chemicals at high pressures to break up the shale to release the gas. An outline of the hydraulic fracturing process can be found in Figure 1. Natural gas from these hard-to-extract sources is becoming increasingly popular and is projected to grow to nearly 45% of the nation's natural

gas supplies by 2035<sup>3</sup>.

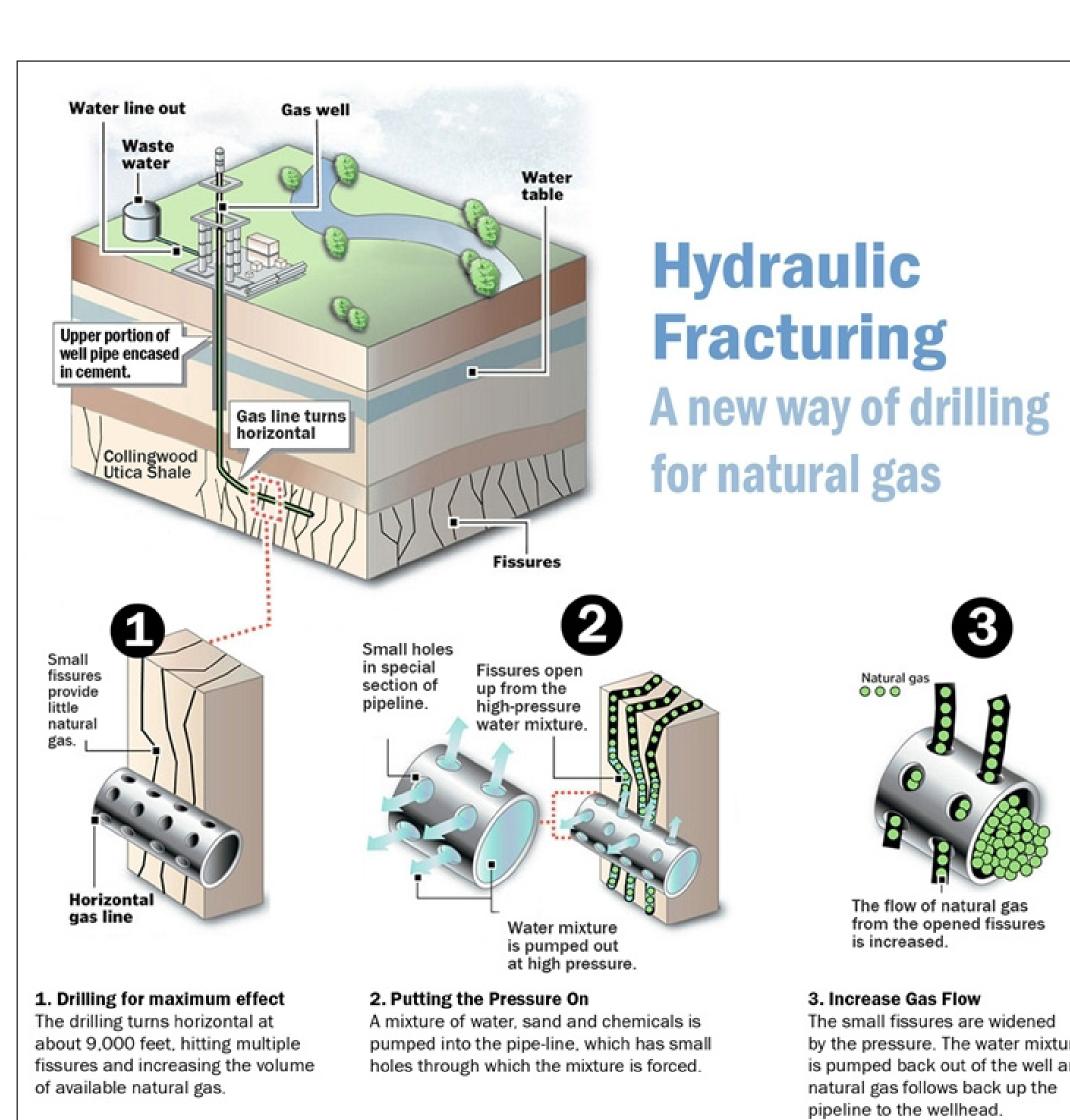


Figure 1: Process of Hydraulic Fracturing to Drill for Natural Gas<sup>4</sup>

Even though hydraulic fracturing is a relatively old practice, first employed over 60 years ago to drill for oil in Oklahoma<sup>5</sup>, there has been little research into the impact of its increasing use as a drilling process for natural gas. With growing concern over the environmental effects of fracking on water quality, the United States EPA has begun studies to monitor the treatment methods and environmental impact with the goal of standardization by 2013/2014<sup>6</sup>. Figure 2 shows a map of shale gas formations in the North America indicating the potential widespread environmental impact the fracking process could have.

The current method for determining natural gas constituents (methane, ethane, and ethene) in water is RSK 175<sup>1</sup>. This method is employed for the analysis of dissolved gases in drinking water using a headspace equilibration technique. Propane has been added to this list in modified methods such as BOL60192, developed by the Pennsylvania Department of Environmental eia Protection (PADEP). This analysis also requires more



modern automated headspace analyzers. A flame ionization detector (FID) will be employed for this study, although RSK 175<sup>1</sup> also allows thermal conductivity (TCD) as well as electron capture detectors (ECD) to be used. This application demonstrates alternative analyses, including purge and trap concentration and automated headspace, to create simple and efficient methods for analyzing dissolved gases.

For this study, a Stratum Purge and Trap Concentrator (PTC) was used in conjunction with an AQUATek 100 Autosampler. This set-up allows for complete automation of sample preparation for the analysis of liquid samples for purge and trap. A recirculating chiller bath was also utilized to maintain a sample temperature of less than 10° C. This technique also requires a 5 mL purge volume. Additionally, the HT3 and Versa Headspace analyzers, utilizing a loop sampling technique, used a 10 mL sample volume for this analysis.

Utilizing a GC/FID, a linear calibration was performed and percent Relative Standard Deviation (%RSD) and Method Detection Limits (MDLs) were determined for the full list of compounds. Similarly to BOL6019<sup>2</sup>, calibrations were performed on aqueous standards rather than the gaseous standards used in RSK 175<sup>1</sup>. Percent carryover was also evaluated for the purge and trap method.

### **Experimental-Instrument Conditions**

The Stratum PTC and AQUATek 100 Autosampler, as well as the HT3 and Versa Headspace analyzers, were coupled to a GC/FID for analysis. Teledyne Tekmar's new proprietary trap (P/N: 15-0885-403) was also utilized for the purge and trap method. The GC was configured with a Restek Rt-U-BOND 15 m x 0.53 mm x 20 μm column. The GC/FID parameters are outlined in Tables 1 and 2. Table 3 outlines the P&T and autosampler conditions. A recirculating chiller bath was also employed to maintain sample temperatures below 10° C. Table 4 outlines the conditions for the headspace analyses.

GC Parameters					
GC:	GC/FID				
Column:	Restek Rt-U-Bond 15m x 0.53mm x 20µm, 1.0 psi constant pressure				
Oven Program:	35° C for 4 min, 20° C/min to 190° C hold for 2 min				
Inlet:	190° C				
Gas:	Helium				
Split Ratio:	20:1				
Table 1: GC Paran	neters				

FID	Parameters			
Temperature:	190°C			
Hydrogen Flow:	35 mL/min			
Air Flow:	300 mL/min			
Mode:	Constant Makeup Flow			
Makeup Flow:	30 mL/min			
Makeup Gas:	Helium			
Table 2: FID Parameters				

Stratum FTC and AQUATER 100 Parameters				
Variable	Value	Variable	Value	
Pressurize Time	0.35 min	Purge Time	1.5 min	
Sample Transfer Time	0.35 min	Purge Temp	20° C	
Rinse Loop Time	0.30 min	Purge Flow	10 mL/min	
Sweep Needle Time	0.30 min	Dry Purge Time	0.0 min	
Bake Rinse	On	Dry Purge Temp	20° C	
Bake Rinse Cycles	1	Dry Purge Flow	100 mL/min	
Bake Rinse Drain Time	0.35 min	GC Start	Start of Desorb	
Presweep Time	0.25 min	Desorb Preheat Temp	95° C	
Water Temp	90° C	Desorb Drain	On	
Valve Oven Temp	80° C	Desorb Time	2.00 min	
Transfer Line Temp	80° C	Desorb Temp	100° C	
Sample Mount Temp	60° C	Desorb Flow	300 mL/min	
Purge ready Temp	35° C	Bake Time	15.00 min	
Condenser Ready Temp	40° C	Bake Temp	100° C	
Condenser Purge Temp	20° C	Bake Flow	400 mL/min	
Standby Flow	10 mL/min	Condenser Bake Temp	200° C	
Dro Durgo Timo	0.5 min			

Sample Temp	40°C		
Table 3: Stratum PTC and AC	QUATek 100 Pard	imeters (Stratu	m PT(
Parameters are in Blue)			

# **Calibration Data**

re-Purge Flow

Sample Heater

ample Preheat Time

To make the stock solutions, a 500 mL volumetric flask filled with de-ionized water was placed in an ice water bath and purged with a reference gas corresponding to each of the four analytes. Each gas was bubbled through chilled water for two hours to make individual concentrated standards. Unlike in RSK 175<sup>1</sup>, calibrations in BOL6019<sup>2</sup> are performed using aqueous rather than gaseous standards. This study also employs an aqueous calibration, where standards are analyzed under the same conditions as samples.

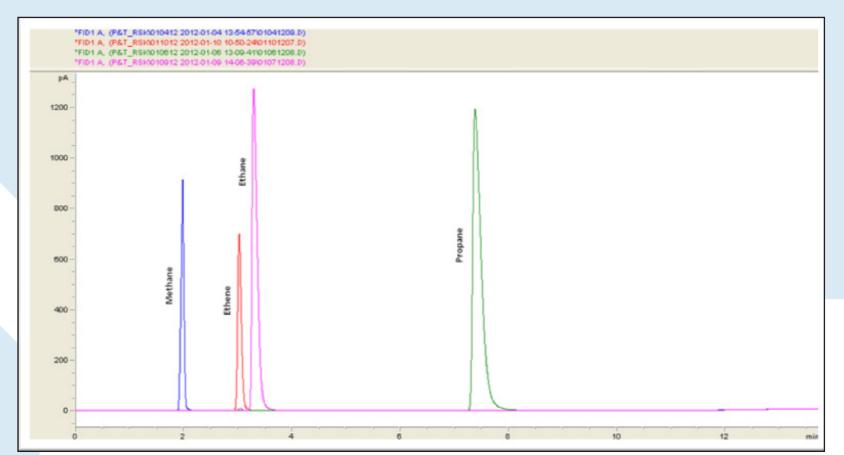
Calibration standards were made from serial dilutions of these stock standards by calculating

Parameters (gray parameters were not used). the concentration of saturated gas solutions in water at 0°C. For example, the saturation point of methane in 0°C water is 39.59 mg/L at atmospheric pressure. Calibration standards were made in 50 mL volumetric flasks filled to volume with chilled de-ionized water over a range of 7.92 ppb to 19.8 ppm. Samples were transferred to headspace free 40 mL vials for analysis. These standards were additionally made for the remaining compounds of interest. Similarly, calibration standards for the headspace analysis were generated from dilutions of this stock standard, but due the fixed volumes used in the method the calibration ranges differed slightly from the purge and trap method.

The calibration data generated during this study was evaluated by linearity (r<sup>2</sup>) and percent Relative Deviation (%RSD). Method detection limits were also established for all compounds by analyzing seven low level replicates. Calibration data and MDLs can be found in Table 5 for the P&T method and Table 6 for the HT3 and Versa. In

addition, an example of an overlay chromatogram showing each standard can be found in Figure 3. Figure 4 presents a chromatogram of a mixed gas standard as well as an overlay showing there is no discernible difference in response between the headspace systems. A blank analyzed after the highest calibration standard was used to calculate the percent carryover for the P&T method which was less than 0.04% for all compounds.

Compound	Calibration Range	Relative Response Factor (RRF)	Linearity (r²)	% Relative Deviation (%RSD)	Method Detection Limit (MDL)	% Carryover
Methane	7.92 ppb to 19.8 ppm	166	1.000	2.0	0.4 ppb	0.04%
Ethene	56.2 ppb to 281 ppm	587	0.9995	4.5	31 ppb	0.03%
Ethane	26.4 ppb to 132 ppm	621	0.9998	13.9	21 ppb	0.04%
Propane	29.4 ppb to 147 ppm	803	0.9999	12.0	18 ppb	0.04%



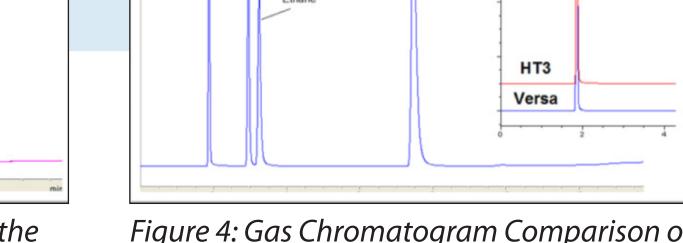


Figure 4: Gas Chromatogram Comparison of Methane and Ethane between the HT3 and the Versa and a Chromatogram of the Four Gases.

Compound	Calibration Range	Linearity (r²)		% Relative Standard Deviation (%RSD)		Method Detection Limit (MDL)	
		HT3	Versa	HT3	Versa	HT3	Versa
Methane	20 ppb to 24 ppm	0.9981	0.9958	3.8	7.3	2 ppb	2 ppb
Ethene	141 ppb to 169 ppm	0.9979	0.9966	6.0	11.9	25 ppb	2 ppb
Ethane	66 ppb to 79 ppm	0.9987	0.9992	6.3	7.8	5 ppb	7 ppb
Propane	74 ppb to 88 ppm	0.9989	0.9992	2.1	12.0	6 ppb	4 ppb

Table 6: Calibration Data for Methane, Ethene, Ethane, and Propane using the HT3 and Versa Headspace Analyzers

**Teledyne Tekmar HT3 Parameters** 

GC Cycle Time

Loop Fill Pressure

On N/A

21.00 min | 21.00 min

30.00 min 30.00 min

Off Off

10 psig 10 psig

0.20 min 0.25 min

7 psig 7 psig

2.00 min | 2.00 mir

0.50 min | 0.50 min

2.00 min

Table 4: HT3 and Versa Automated Headspace

100° C

With increased interest in alternative energy sources, hydraulic fracturing has become a common practice in the extraction of natural gas from coalbeds and shale formations across the United States. Unfortunately, there has not been adequate time to measure the environmental impact of these procedures. Regulatory agencies are looking for easy and reliable testing methods to monitor these effects.

This study demonstrates methods for analyzing these gases using the Teledyne Tekmar's Stratum PTC/AQUATek 100 Autosampler and HT3 and Versa Headspace analyzers coupled with a GC/FID system. These methods met all performance criteria outlined in the current Headspace methods, RSK 175<sup>1</sup> and BOL6019<sup>2</sup>. By completely automating the sample preparation, efficiency and throughput can be greatly increased while saving time and money. Using the purge and trap method, there is no need to manipulate the samples which eliminates the potential for human error and employs instrumentation already familiar to many environmental laboratories.

#### References

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AQUATek 100 Waters-Only Autosampler and Stratum PTC Purge and Trap Concentrator



