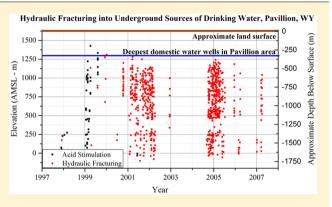
Impact to Underground Sources of Drinking Water and Domestic Wells from Production Well Stimulation and Completion Practices in the Pavillion, Wyoming, Field

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Supporting Information

ABSTRACT: A comprehensive analysis of all publicly available data and reports was conducted to evaluate impact to Underground Sources of Drinking Water (USDWs) as a result of acid stimulation and hydraulic fracturing in the Pavillion, WY, Field. Although injection of stimulation fluids into USDWs in the Pavillion Field was documented by EPA, potential impact to USDWs at the depths of stimulation as a result of this activity was not previously evaluated. Concentrations of major ions in produced water samples outside expected levels in the Wind River Formation, leakoff of stimulation fluids into formation media, and likely loss of zonal isolation during stimulation at several production wells, indicates that impact to USDWs has occurred. Detection of organic compounds used for well stimulation in samples from two monitoring wells installed by



EPA, plus anomalies in major ion concentrations in water from one of these monitoring wells, provide additional evidence of impact to USDWs and indicate upward solute migration to depths of current groundwater use. Detections of diesel range organics and other organic compounds in domestic wells <600 m from unlined pits used prior to the mid-1990s to dispose diesel-fuel based drilling mud and production fluids suggest impact to domestic wells as a result of legacy pit disposal practices.

INTRODUCTION

Between 2005 and 2013, natural gas production in the U.S. increased by 35% largely due to unconventional gas production in shale and tight gas formations.¹ Between 2013 and 2040, natural gas production is expected to increase another 45% with production from tight gas formations in particular increasing from 4.4 to 7.0 trillion cubic feet (59%) primarily in the Gulf Coast and Dakotas/Rocky Mountain regions.¹ Tight gas formations already account for 26% of total natural gas production in the United States today.²

In the U.S. Code of Federal Regulations (CFR), there are two federal regulations for protecting groundwater resources for present and future use relevant to oil and gas extraction – "Underground Source of Drinking Water" (USDW) and "usable water." A USDW is defined in 40 CFR 144.3 in requirements for the Underground Injection Control program promulgated under Part C of the Safe Drinking Water Act (SDWA) as "an aquifer or its portion: (a)(1) Which supplies any public water system; or (2) Which contains a sufficient quantity of ground water to supply a public water system; and (i) Currently supplies drinking water for human consumption; or (ii) Contains fewer than 10 000 mg/L total dissolved solids; and (b) Which is not an exempted aquifer." With the exception of use of diesel fuels, the Energy Policy Act of 2005 ("EPAct") exempted hydraulic fracturing from the SDWA, thereby allowing injection of stimulation fluids into USDWs. However, under Section 1431 of the SDWA, the Administrator of EPA may take action if impact to a USDW "may present an imminent and substantial endangerment to the health of persons."

The term "usable water" applies to lands containing federal or tribal mineral rights regulated by the Bureau of Land Management (BLM). This term is applicable to the Pavillion Field because tribal mineral rights are associated with more than half of production wells there. In the BLM Onshore Oil and Gas Order No. 2, usable water is defined as water containing $\leq 10\,000$ mg/L total dissolved solids (TDS) – a definition maintained in the March 2015 BLM rule on hydraulic fracturing (43 CFR 3160). In 43 CFR 3160, BLM retained a threshold for groundwater protection at 10 000 mg/L stating, "Given the increasing scarcity and technological improvements in water treatment, it is not unreasonable to assume aquifers with TDS levels above 5000 ppm are usable now or will be usable in the future." However, on September 30, 2015, the U.S. District Court for Wyoming granted a

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preliminary injunction filed by the States of Wyoming and Colorado to stop implementation of the BLM rule based on the assertion that the EPAct precludes BLM rulemaking.³

In 2004, EPA⁴ documented the widespread use of hydraulic fracturing in USDWs colocated in formations used for coal bed methane (CBM) recovery. EPA⁴ acknowledged likely groundwater contamination as a result of this activity but stated that the attenuation factors of dilution, adsorption, and biodegradation would reduce contaminant concentrations to safe levels prior to reaching domestic wells that are generally shallower than production wells. Thus, EPA⁴ distinguished impact to USDWs from impact to domestic wells. In 2014, while defining the chemical abstract numbers of fluids designated as diesel fuels, EPA revised its position and stated that injecting stimulation fluids directly into USDWs "presents an immediate risk to public health because it can directly degrade groundwater, especially if the injected fluids do not benefit from any natural attenuation from contact with soil, as they might during movement through an aquifer or separating stratum."

The Pavillion Field (Figure 1) is located east of the Town of Pavillion in Fremont County, WY, in the west-central portion

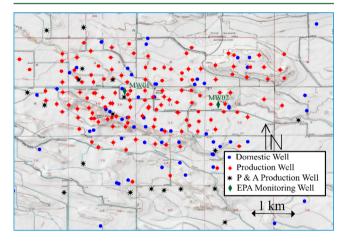


Figure 1. Central portion of the Pavillion Field illustrating locations of domestic water wells, production wells, plugged and abandoned (P&A) wells, and EPA monitoring wells (labeled). The entire Field, with labels for production and domestic wells and approximate locations of unlined pits, is illustrated in Figure SI A5. The geographic area in which the Field is located is illustrated in Figure SI A1.

of the Wind River Basin (WRB) (Figure SI A1). The field consists of 181 production wells including plugged and abandoned wells. Conventional and unconventional (tight gas) hydrocarbon production in the Pavillion Field is primarily natural gas from sandstone units in the Paleocene Fort Union and overlying Early Eocene Wind River Formations. However, oil has also been produced from production wells in these formations, primarily in the western portion of the field close to the suspected location of a fault (SI Sections A.1 and A.2).

In response to complaints regarding foul taste and odor in water from domestic wells within the Pavillion Field, EPA initiated a groundwater investigation in September 2008 under the Comprehensive Environmental Response and Liability Act (CERCLA).⁶ This investigation remains the only one in which CERCLA has been invoked to investigate potential groundwater contamination due to hydraulic fracturing.⁷ Under CERCLA, impact to both groundwater resources and domestic wells is evaluated, in contrast to limiting evaluation to impact to

domestic wells as is common in oil- and gas-field-based investigations.

EPA conducted two domestic well sampling events in March 2009 (Phase I)⁶ and January 2010 (Phase II).⁸ Between June and September 2010, EPA installed two monitoring wells, MW01 and MW02, using mud rotary drilling with screened intervals at 233–239 m and 296–302 m below ground surface (bgs), respectively. These monitoring wells were installed to evaluate potential upward solute transport of compounds associated with well stimulation to maximum depths of current groundwater use (~322 m).⁹ EPA sampled MW01 and MW02 during the Phase III (October 2010) and Phase IV (April 2011) sampling events.

In December 2011, EPA⁹ released a draft report summarizing results of the Phase I–IV sampling events. EPA documented groundwater contamination in surficial Quaternary unconsolidated alluvium attributable to numerous unlined pits used for disposal of diesel-oil-based (invert) drilling mud and production fluids including flowback, condensate, and produced water prior to the mid-1990s. EPA⁹ also documented injection of stimulation fluids into USDWs and concluded that inorganic and organic geochemical anomalies at MW01 and MW02 appeared to be attributable to production well stimulation. EPA received numerous comments both challenging and supporting its findings in the draft EPA report.^{10–37} We reviewed and considered these comments when preparing this manuscript.

A substantial amount of data has been collected since publication of the 2011 draft EPA report, adding to an already extensive data set. In April 2012 (Phase V) the EPA^{38,39} split samples with the U.S. Geological Survey at MW01^{40,41} and MW02.⁴² In 2014, the Wyoming Oil and Gas Conservation Commission (WOGCC) released a report on production well integrity⁴³ and in 2015 released a report on surface pits.⁴⁴ In December 2015, the Wyoming Department of Environmental Quality (WDEQ) released a report on sample results of a subset of domestic wells previously sampled by EPA.⁴⁵

We conducted a comprehensive analysis of all publicly available online data and reports, to evaluate impact to USDWs and usable water as a result of acid stimulation and hydraulic fracturing. Although injection of stimulation fluids into USDWs in the Pavillion Field was previously documented by EPA,⁹ the potential impact to USDWs at depths of stimulation was not assessed. We evaluate potential upward migration of contaminants to depths of current groundwater use using data from MW01 and MW02. We also evaluate potential impact to domestic wells as a result of legacy disposal of production and drilling fluids in unlined pits.

MATERIALS AND METHODS

Sources of EPA reports, versions of the Quality Assurance Project Plan (QAPP), and Audits of Data Quality (ADQs) are provided in Table SI H1. Sources of analytical data and associated information on quality assurance and control are summarized in Table SI H2. ADQs were conducted by EPA for Phase I–IV investigations to verify the quality of analytical data and consistency with requirements specified in the QAPP.

In response to a comprehensive information request by EPA regarding oil and gas production and disposal activities in the Pavillion Field, the field operator, Encana Oil & Gas (U.S.) Inc., provided Material Safety and Data Sheets (MSDSs) of products used for well stimulation to EPA⁴⁶ (Table SI C3). During the Phase V sampling event, EPA developed a gas chromatography-

Table 1. Summary of Major Ion Concentrations of Domestic Wells in the Wind River Indian Reservation (WRIR), Fremont County, WY, and within and around the Pavillion Field

	WRIR ^a			Fremont County ^b			within and around Pavillion $Field^c$		
parameter (mg/L)	п	median	range	п	median	range	n	median	range
TDS	154	490	211-5110	77	1030	248-5100	65	925 [†]	229 ⁺ -4901 ⁺
Ca	149	10	1-486	77	45	1.7-380	48	50.8	3.32-452
Mg	128	2.2	0.1-195	77	8.2	0.095-99	45	5.32	0.024-147
Na	153	150	5-1500	77	285	4.5-1500	72	260	38.0-1290
K	149	2.0	0.2-30	77	2.45	0.1-30	43	1.36	0.179-10.5
SO_4	154	201	2-3250	77	510	12-3300	88	590	29.0-3640
Cl	154	14	2-466	77	20	3-420	48	21.1	2.60-77.6
F	154	0.7	0.1-8.8	76	0.9	0.2-4.9	46	0.88	0.20-4.1

^{*a*}With the exception of potassium, from Daddow.⁴⁸ Information on potassium extracted from Daddow.⁵³ ^{*b*}From Plafcan et al.⁵¹ There is overlap of 19 sample results with Daddow.^{48,53} ^{*c*}Major ion concentrations in domestic wells^{6,8,9,39,45,52} summarized in Table SI B2. Mean values used for domestic wells sampled more than once. ^{*n*}Number of sample results. ^{*†*}TDS for EPA data estimated using linear regression equation from Daddow⁴⁸ TDS (mg/L) = 0.785 × specific conductance (μ s/cm) – 130 (n = 151, $r^2 = 0.979$)

flame ionization-based approach to obtain a lower reporting limit (50 μ g/L) for methanol compared to commercial laboratory analysis (5000 μ g/L). We obtained this data set as the result of a Freedom of Information Act request to EPA.⁴⁷

We reviewed over 1000 publicly available well completion reports, sundry notices, drilling reports, and cement bond and variable density logs accessed from the WOGCC Internet site using API search numbers to determine dates of well completion, depths of surface casing, top of original or primary cement, and numbers and depths of cement squeeze jobs (injection of cement through perforated production casing to remediate or extend existing primary cement). Similarly, we reviewed online information to document well stimulation practices summarized in Tables SI C1 and SI C2.

The field operator analyzed major ions in produced water samples at 42 production wells in 2007 (Table SI D1). EPA collected produced water samples at four production wells in 2010 and analyzed them for organic compounds (Table SI D3).8 The field operator also conducted mechanical integrity and bradenhead (annular space between production and surface casing) testing between November 2011 and December 2012. In addition to sustained casing pressure at many production wells during that period (Table SI D2), water flowed through the bradenhead valve to the surface at four production wells (SI Section D.3). Aqueous analysis of bradenhead water samples by the field operator was limited to major ions (Table SI D1). Production well string and brandenhead gas samples were collected for benzene, toluene, ethylbenzene, xylenes (BTEX) and light hydrocarbons (Table SI D2).

To evaluate the effect of purging volume on water quality, EPA collected ten samples through time (Table SI 3a) during the Phase V sampling event at MW01. Based on EPA's purging procedure, we developed a model incorporating plug flow in casing and mixing in the screened interval (SI Section E.3, Figure SI E4). Our simulations indicated that virtually all (99.997%) of water entering the sampling train at the surface at the time of the first sample collection at MW01 originated directly from the surrounding formation (i.e., no stagnant casing water) (Figure SI E6). MW02 was a low flow monitoring well. The cause of low flow is unknown but could be due to several factors, including low relative aqueous permeability due to gas flow or insufficient removal of drilling mud during well development. During the Phase V sampling event, MW02 was repeatedly purged over a 6-day period to ensure that sampled

water originated from the surrounding formation (SI Section E.2, Figure SI E5). A discussion of monitoring well construction, including schematics for MW01 (Figure SI E1) and MW02 (Figure SI E2), is provided in SI Section E.1.

RESULTS AND DISCUSSION

Groundwater Resources in the Pavillion Area. The Wind River and Fort Union Formations are variably saturated fluvial depositional systems characterized by shale and fine-, medium-, and coarse-grained sandstone sequences. Lithology is highly variable and difficult to correlate from borehole data. No laterally continuous confining layers of shale exist below the maximum depth of groundwater use to retard upward solute migration. A comprehensive review of regional and local geology, including a lithologic cross-section in the vicinity of MW01 and MW02 (Figure SI A4), is provided in SI Sections A.1–A.6.

Domestic wells in the Pavillion area draw water from the Wind River Formation—a major aquifer system in the WRB.^{48,49} From the surface to approximately 30 m bgs, groundwater exists under unconfined conditions.⁵⁰ Below this depth, groundwater is present in lenticular, discontinuous, confined sandstone units with water levels above hydrostatic pressure, and in some instances flowing to the surface,^{48,50,51} indicating the presence of strong localized upward gradients. The majority of documented domestic well completions in Fremont County⁵¹ and five municipal wells in the Town of Pavillion⁵² west of the Field are completed in the Wind River Formation.

Flow to the surface was observed in a domestic well during the Phase II sampling event,⁶ and as mentioned, at four production wells during bradenhead testing in 2012. While the overall vertical groundwater gradient in the Pavillion Field is downward, these observations indicate that localized upward hydraulic gradients exist in the Field, which is relevant to potential upward solute migration from depths of production well stimulation. The deepest domestic wells in the Pavillion Field and immediate surrounding area are 229 and 322 m bgs, respectively (Table SI B1). Two municipal wells were proposed, but not drilled, in the Pavillion Field as replacement water for domestic wells at depths of 305 m bgs,⁵² similar to the depth of MW02 installed by EPA.

Major ion concentrations of domestic wells in the Pavillion field (summarized in Table SI B2) are typical of the Wind River Indian Reservation (WRIR),⁴⁸ west of the Pavillion Field, and

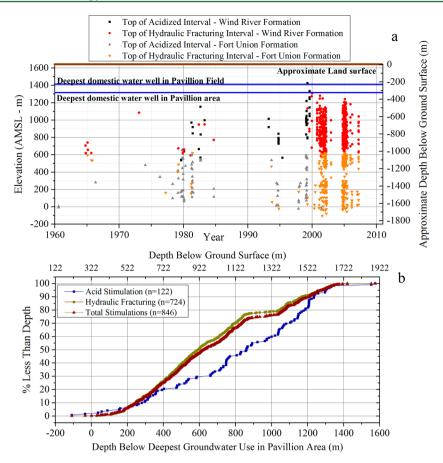


Figure 2. (a) Elevation in absolute mean seal level (AMSL) and approximate depth below ground surface of documented acid and hydraulic fracturing stimulation stages. (b) Cumulative distribution of stimulation stages as a function of depth below deepest groundwater use in the Pavillion Field. Documentation of stimulation stages is absent at a number of production wells so that numbers presented here are a lower bound.

in Fremont County,⁵¹ where the Pavillion Field lies, (Table 1) with TDS levels <5000 mg/L. TDS concentrations in the Wind River Formation appear to vary with lithology rather than depth (white coarse sandstone associated with lower TDS values).⁵² There are no apparent trends in TDS levels with depth from data sets from the WRIR,⁵³ Fremont County,⁵¹ and domestic wells in and around the Pavillion Field.

The Fort Union Formation is not used for water supply in the Pavillion area. However, the formation is highly productive and permeable where fractured⁴⁹ with TDS values from 1000 to 5000 mg/L.⁵⁴ An aquifer exemption was obtained to enable disposal of produced water in a disposal well perforated in the Fort Union Formation⁵⁵ at a location 5.6 km northwest of the Pavillion Field. Use of this well was suspended due to failure of well casing. Thus, the Wind River and Fort Union Formations in the Pavillion Field meet the regulatory definition of USDWs, as explicitly stated by EPA,^{9,55} and of usable water as defined by the BLM.

Well Stimulation Depths, Treatments, and Chemical Additives. Exploration of oil and gas in the Pavillion Field commenced in August 1953 with increasingly shallow stimulations through time (Figure 2). The first acid stimulation and hydraulic fracturing stages (injection over one or more discrete intervals) occurred in June 1960 and October 1964, respectively. Acid stimulation ceased in 2001. To date, the last stimulation stage (hydraulic fracturing) occurred in April 2007. Most production wells were completed and stimulated during several periods of increased activity, especially after 1997

(Figure 2a). Acid stimulation and hydraulic fracturing occurred as shallowly as 213 and 322 m bgs, respectively, at depths comparable to deepest domestic groundwater use in the area (Figure 2a). Approximately 10% of stimulation stages were <250 m of deepest domestic groundwater use whereas approximately 50% of stimulation stages were <600 m and 80% were <1 km of deepest domestic groundwater use (Figure 2b).

Surface casing of production wells—the primary line of defense to protect groundwater during conventional and unconventional oil and gas extraction—is relatively shallow in the Pavillion field with a median depth of 185 m bgs (i.e., shallower than the deepest groundwater use) and range of 100–706 m bgs (Figure SI C1). There is no primary cement below surface casing, often for hundreds of meters, for 55 of 106 (~52%) production wells for which cement bond logs are available (Table SI C1, Figure SI C1). There is currently no requirement in Wyoming for placement of primary cement to surface casing or to ground surface.⁴⁵

Instantaneous shut in pressures (ISIP) (wellhead gauge pressure immediately following fracture treatment) were similar for acid stimulation and hydraulic fracturing (Figure SI C2) suggesting that both matrix acidizing and acid fracturing (no proppants used⁵⁶) occurred in the Pavillion Field. Acidizing solutions used in the Pavillion Field typically consisted of a 7.5% or 15% hydrochloric acid solution plus additives described in well completion reports as inhibitors, surfactants, diverters, iron sequestration agents, mutual solvents, and clay stabilizers.

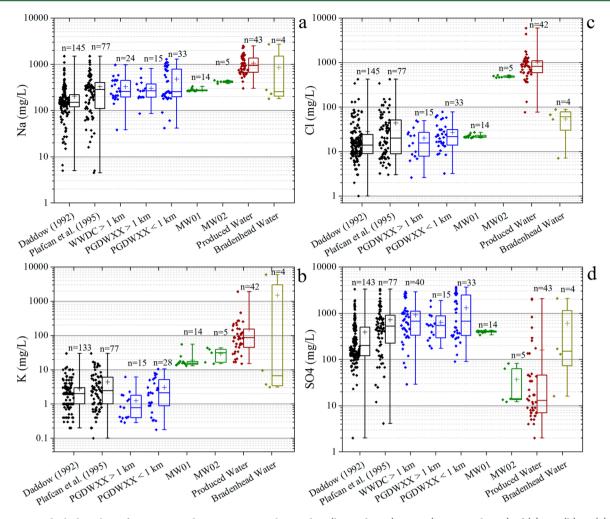


Figure 3. Box and whisker plots of minimum and maximum, quartiles, median (line in boxes), mean (crosses in boxes) of (a) Na, (b) K, (c) Cl, (d) SO_4 for domestic wells inventoried by Daddow^{48,53} and Plafcan⁵¹ in the Wind River Indian Reservation and Fremont County, respectively, sampled by EPA^{68,9,39} and WDEQ⁴⁵ (PGDWXX series) greater than and less than 1 km from a production well, Wyoming Water Development Commission⁵² (WWDC series) greater than 1 km from a production well, EPA monitoring wells^{9,39} (Tables SI E2b, SI E3b), and produced water and bradenhead water samples (Table SI D1). Domestic wells sampled more than once, including data from Daddow,⁵³ are represented with a mean value. Fourteen measurements in Daddow⁵³ < 1 mg/L for potassium are not illustrated. Data points at MW01 and MW02 are samples collected during Phase III, IV, and V sample events.

Acidizing solutions were often flushed with a 2, 4, or 6% potassium chloride (KCl) solution. Pad acid, to initiate fractures, contained 10–50% heavy aromatic petroleum naptha. Corrosion inhibitors contained isopropanol and propargyl alcohol. Clay stabilizers contained methanol. Musol solvents used for acid stimulation consisted of 60-100% 2-butoxyethanol and 10-30% oxylated alcohol (Table SI C3).

Prior to 1999, "salt solutions" were commonly used for hydraulic fracturing. After 1999, a 6% KCl solution was used extensively for hydraulic fracturing often combined with CO_2 foam, with subsequent flushing using a 6% KCl solution. There were reported losses of KCl solutions during stimulation (e.g., at Tribal Pavillion 12–13 "lost thousands of bbls KCl"). Undiluted diesel fuel was used for hydraulic fracturing at three production wells before 1985. From the mid-1970s through 2007, there was widespread use of gelled fracture fluids (gelled water, linear gel, and cross-linked gel). Diesel fuel #2 was used for liquid gel concentrates (Table SI C3). Ammonium chloride, potassium hydroxide, potassium metaborate, and a zirconium complex were used as cross-linkers.

Gelled fracture fluids were used extensively with CO₂ foam (Table SI C4). Between 2001 and 2005, "WF-125" was used with CO₂ foam (often with a 6% KCl solution) for hydraulic fracturing (Table SI C5). A stimulation report (one of only three publicly available throughout the operating history of the Field) and MSDSs indicate that WF-125 contained diesel fuel #2, 2-butoxyethanol, isopropanol, ethoxylated linear alcohols, ethanol, and methanol. During 2001, WF-125 and unidentified product mixtures were used with a 6% KCl and a 10% methanol solution and CO₂ foam for hydraulic fracturing followed with a 6% KCl and 10% methanol solution flush. Other WF-series compound mixtures of unknown composition were also used with CO₂ foam and in some cases with N₂ gas. Methanol, isopropanol, glycols, and 2-butoxyethanol were used in foaming agents (Table SI C3). Ethoxylated linear alcohols, isopropanol, methanol. 2-butoxyethanol, heavy aromatic petroleum naptha. naphthalene, and 1,2,4-trimethylbenzene were used in surfactants (Table SI C3). Slickwater (commonly with a 6% KCl solution) was used for hydraulic fracturing with and without CO₂ foam in 2004 and 2005, respectively (Table SI C6).

At least 41.5 million liters (or ~11 million gallons) of fluid was used for well stimulation in the Pavillion Field (calculated from Table SI C2). Given lack of information at numerous production wells, this is an underestimate of actual cumulative stimulation volume. The cumulative volume of well stimulation in closely spaced vertical wells in the Pavillion Field is characteristic of high volume hydraulic fracturing in shale units.⁵⁷ In evaluating solute attenuation in USDWs, EPA⁴ did not consider cumulative volumes of injection of well stimulation fluids in closely spaced vertical production wells common to CBM and tight gas production.

Evaluation of Impact to USDWs and Usable Water. In the Pavillion Field, impact to USDWs and usable waters depends upon the advective-dispersive solute transport of compounds (or their degradation products) used for well stimulation to water-bearing units (sandstone units at or near water saturation). Water-bearing units exist throughout the Wind River and Fort Union Formations in the Pavillion Field. For instance, production well Unit 41X-10 was recommended for plugging and abandonment in 1980 because of "problems with water production and casing failure." In 1980, drilling logs at Tribal Pavillion 14-2 stated "Hit water flow while drilling at 4105-4109 ft" bgs. The magnitude of produced water production in the Pavillion Field is variable with some wells having high produced water production (e.g., 17.9 million liters ~4.7 million gallons at Tribal Pavillion 23-10 from July 2000 to present) (Table SI C2). In some cases, stimulation fluids were injected directly into water bearing units. For instance, at Tribal Pavillion 14–1, a cast iron bridge plug was used to stop water production in 1993 from an interval where hydraulic fracturing occurred using undiluted diesel fuel in 1964 (Table SI C2).

The migration of stimulation fluid to water-bearing sandstone units in the Pavillion Field also likely occurred during fracture propagation and subsequent leakoff (loss of fluid into a formation in or near the target stratum). Leakoff increases in complex fracture networks as a result of lithologic variation over short distances and contact of stimulation fluid with permeable strata⁵⁸⁻⁶¹ expected during hydraulic fracturing in fluvial depositional environments of the Wind River and Fort Union Formations. Leakoff can remove much or most of the fracturing fluid even for moderate sized induced fractures.^{58,59} Maximum ISIP values for acid stimulation and hydraulic fracturing were 19.5 and 40.1 MPa (Figure SI C2), respectively, equivalent to ~2000 and ~4100 m of hydraulic head. Pressure buildup during hydraulic fracturing far in excess of drawdown expected during produced water extraction makes full recovery of stimulation fluids unlikely.^{4,62}

The migration of stimulation fluids to water-bearing units also likely occurred as a result of loss of zonal isolation during well stimulation (SI Section D.1). Casing failure occurred at five production wells following well stimulation. Cement squeezes were performed above primary cement often days after hydraulic fracturing without explanation⁶³ at six production wells, potentially because of migration of stimulation fluid above primary cement. At one production well, stimulation fluid was injected just 4 m below an interval lacking cement outside of the production casing with a stimulation pressure of only 1.3 MPa indicating potential entry into the annular space.

Major ion concentrations in produced water sampled after stimulation (Table SI D1) were distinct from values expected in the Wind River Formation as evidenced by sample data from the WRIR,^{48,53} Fremont County,⁵¹ and domestic wells in and around the Pavillion Field which were representative of the Wind River Formation regardless of distance from production wells (Table 1, Figure 3). Using combined data sets in and around the Pavillion Field, and the nonparametric Mann-Whitney test (null hypothesis that two sample sets come from the same population), sodium, potassium, and chloride concentrations were higher and sulfate concentrations lower in produced water compared to concentrations expected in the Wind River Formation $(p = 6.6 \times 10^{-19}, 2.1 \times 10^{-15}, 2.6 \times 10^{-15})$ 10^{-16} , and 4.4×10^{-19} , respectively), providing direct evidence of impact to USDWs at depths of stimulation. Also, potassium increased with calcium concentrations and sulfate increased with TDS concentrations, respectively, in domestic wells but not in production wells (Figures SI D1). Chloride is a major component of TDS concentrations in production wells. Potassium/calcium and chloride/sulfate concentration ratios were higher in production wells than in domestic wells (Figures SI D2), further indicating anomalous potassium, chloride, and sulfate concentrations in production wells.

Produced water samples were collected from gas-water separators at four production wells and analyzed for organic compounds (Table SI D3, Figure SI D3) during the Phase II sampling event.⁶ Samples from one production well appeared to be from both an aqueous and an apparent nonaqueous phase liquid with the latter exhibiting thousands of mg/L of benzene, toluene, ethylbenzene, xylenes (BTEX). Synthetic organic compounds methylene chloride and triethylene glycol (TEG) were detected in produced water samples at 0.51 and 17.8 mg/L, respectively indicating anthropogenic origin. Methylene chloride has been detected in flowback water in other systems,⁶⁴ including 122 domestic wells above the Barnett Shale TX,⁶⁵ and in air sampled near well sites.⁶⁶

Sample Results at MW01 and MW02. Concentrations of potassium in MW01 and MW02 were higher than expected values in the Wind River Formation (Figure 3) at p-values of 2.6×10^{-13} and 1.2×10^{-06} , respectively. High pH values (>11 standard units) were observed during purging at both monitoring wells (Tables SI E3b, SI E4b, Figures SI E5, SI E6, SI E7), indicating that elevated potassium concentrations may have been attributable to release of potassium from potassium oxides and sulfates during curing of cement^{0/-} used for monitoring well construction. However, a number of observations were inconsistent with cement interaction as a causative factor for elevated pH, and there was extensive use of compounds containing potassium including potassium hydroxide during stimulation (Table SI C3). Water in contact with hydrating cement is saturated or oversaturated to portlandite $(Ca(OH_2))^{72-74}$ and remains oversaturated prior to degradation or carbonation.⁷⁵⁻⁷⁸ In contrast, water from monitoring wells was highly undersaturated to portlandite. Elevated pH in monitoring wells was not observed during monitoring well development until natural gas intrusion occurred in the wells, suggesting degassing as a possible cause of elevated pH (SI Section E.5). Also, potassium was detected at a concentration of 6000 mg/L in a bradenhead water sample having a pH of 10.86 standard units from Tribal Pavillion 13–1 (Table SI D1). This may indicate either high potassium concentration at depths below EPA monitoring wells due to well stimulation (water from bradenhead samples originated at some unknown distance above cement outside production casing at each production well) or interaction of bradenhead water with wellbore cement.

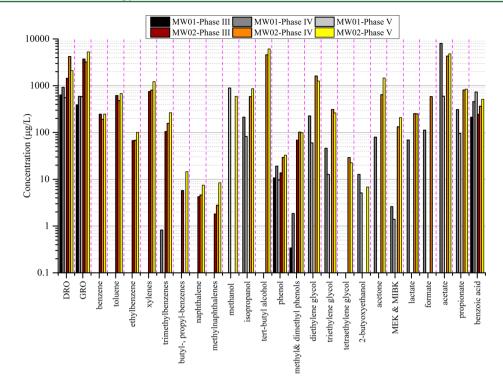


Figure 4. Summary of organic compounds detected by EPA in MW01 and MW02 during Phase III, IV, and V sampling events. Glycols, alcohols, and low molecular weight organic acids were not analyzed in Phase III. Alkylphenols and methanol (GC-FID method) were only analyzed in Phase V. Organic compounds detections for MW01 and MW02 are summarized in Table SI E3a and Table SI E4a, respectively.

The median chloride concentration at MW02 was 469 mg/L (Figure 3), well above expected values in the Wind River Formation ($p = 7.0 \times 10^{-07}$). Compounds containing chlorides (e.g., KCl solutions) were used extensively for stimulation in the Pavillion Field. Sulfate concentrations in MW02 were below expected values in the Wind River Formation ($p = 2.7 \times 10^{-07}$) and not dissimilar (p = 0.40) to produced water concentrations. The Cl/SO₄ concentration ratio was similar to produced water (Figure SI D2) at MW02. Chloride and sulfate concentrations in MW01 were more typical of the Wind River Formation which may be due variation in well stimulation practices both spatially and over time.

Concentrations of organic compounds detected in MW01 and MW02 are summarized in Tables SI E3a, SI E4a and Figure 4. Diesel range organics (DRO) and gasoline range organics (GRO) were detected in MW01 and MW02 with maximum DRO concentrations of 924 and 4200 μ g/L, respectively and GRO concentrations of 760 and 5290 μ g/L, respectively. Benzene, toluene, ethylbenzene, *m*,*p*-xylenes, and *o*-xylene were detected in MW02 at maximum concentrations of 247, 677, 101, 973, and 253 μ g/L, respectively, but were not detected at MW01. The maximum contaminant level (MCL) of benzene is 5 μ g/L, so the observed maximum value was 50 times higher than the MCL. Nondetection of BTEX at MW01 is surprising given that the well was gas-charged (foaming during sampling, Figure SI E9) with similar light hydrocarbon composition to MW02 (Table SI E5). Nondetection of BTEX may be due to increased dispersion and biodegradation of these compounds at the shallower depth of this well. We could find no published information on BTEX compounds in groundwater at concentrations detected in MW02 occurring above a gas field in the absence of well stimulation. However, further testing, such as compound specific isotope analysis of BTEX components present in natural gas from the Pavillion Field

(Table SI D2) and water from MW02, is necessary to attribute detection of BTEX to well stimulation.

1,3,5-, 1,2,4-, and 1,2,3-Trimethylbenzene were detected at maximum concentrations of 71.4, 148, and 45.8 μ g/L, respectively in MW02 and at an order of magnitude lower concentrations in MW01. Naphthalene, methylnaphthalenes, and alkylbenzenes were also detected in MW02 at concentrations up to 7.9, 10.2, and 21.2 μ g/L, respectively. Similar to BTEX compounds, detection of trimethylbenzenes, alkylbenzenes, and naphthalenes could in principle reflect nonanthropogenic origin but natural gas from the Pavillion Field and in EPA monitoring wells is "dry" (ratio of methane to methane through pentane concentration >0.95) (SI Section A.2, Table SI E5). Also, oil production in the vicinity of monitoring wells is very low or zero especially in the vicinity of MW02 (Table SI C2, Figure SI A5). Thus, the detection of higher molecular weight hydrocarbons in groundwater is unexpected. Trimethylbenzenes and naphthalenes were present in mixtures used for well stimulation (Table SI C3).

Other organic compounds used extensively for well stimulation were detected in MW01 and MW02 (Figure 4). Methanol, ethanol, and isopropanol were detected in monitoring wells at up to 863, 28.4, and 862 μ g/L, respectively (Figure 4). *Tert*-butyl alcohol (TBA) was detected at 6120 μ g/L in MW02. Detection of TBA in groundwater has been associated with degradation of *tert*-butyl hydroperoxide used for hydraulic fracturing.⁷⁹ Another potential source of TBA is degradation of methyl *tert*-butyl ether (MTBE) associated with diesel fuel.^{80–84}

Diethylene glycol (DEG) and TEG were detected in both monitoring wells at maximum concentrations of 226 and 12.7 μ g/L, respectively, in MW01, and at 1570 and 310 μ g/L respectively, in MW02 (Figure 4). Tetraethylene glycol was detected only in MW02 at 27.2 μ g/L. MSDSs indicate that

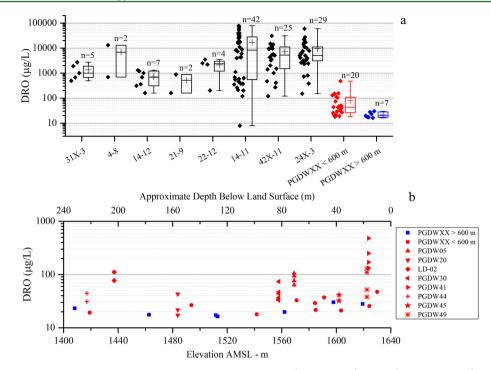


Figure 5. (a) Box and whisker plots of minimum and maximum, quartiles, median (line in boxes), mean (crosses in boxes) of diesel range organics (DRO) in shallow monitoring wells near unlined pits potentially receiving production fluids (abbreviations of production wells in Table SI C1) and domestic wells^{6,8,9,39,45} (LD-20 and PGDWXX series) less than and greater than 600 m from pits. Mean values are used for domestic well sampled more than once. (b) DRO as a function of elevation and approximate depth below surface for domestic wells with results of multiple sample events illustrated.

DEG was used for well stimulation. Use of TEG was not specified. Polar organic compounds, including DEG, are commonly used as cement grinding agents.85-88 DEG and TEG have been detected in leachate from cured cement samples under static (no flow) conditions.⁸⁹ Similar to elevated potassium detection, it is possible that detection of glycols could be attributable to cement used for monitoring well construction. However, mass flux scenario modeling, commonly used to evaluate potential concentrations of exposure of compounds released from materials in contact with drinking water under dynamic (flowing) conditions,⁹⁰ was conducted on MW01 (SI Section E.7) indicating unlikely impact. The relevance of dynamic testing is corroborated by the observation that detection of DEG and TEG was limited to a water sample from a gas production well⁹¹ with nondetection in water samples from 83 domestic wells at five retrospective study sites^{79,91-94} using high performance liquid chromatography with dual mass spectrometry at a reporting limit 5 μ g/L in EPA's national study on hydraulic fracturing. 2-Butoxyethanol, a glycol ether used extensively for well stimulation in the Pavillion Field (Table SI C3), was detected in both monitoring wells at a maximum concentration of 12.7 μ g/L. 2-Butoxyethanol was not detected in leachate from cured cement.⁸⁹.

The low molecular weight organic acids (LMWOAs) lactate, formate, acetate, and propionate were detected in both monitoring wells at maximum concentrations of 253, 584, 8050, and 844 μ g/L, respectively (Figure 4). LMWOAs are anaerobic degradation products associated with hydrocarbon contamination in groundwater.^{95,96} Acetate has been detected in produced water,^{97–99} in impoundments used to hold flowback water from the Marcellus Shale,¹⁰⁰ and in produced water from the Denver-Julesburg Basin, CO.¹⁰¹ Acetate and formate were detected in flowback water from two different fracturing sites in Germany with investigators concluding that these compounds were likely of anthropogenic origin resulting from degradation of polymers used in the fracturing fluid.¹⁰² Formate and acetate are also degradation products of methylene chloride.¹⁰³ Benzoic acid, a degradation product of aromatics, was also detected in both monitoring wells at a maximum concentration of 513 μ g/L.

Phenols were detected in both monitoring wells with maximum concentrations of phenol, 2-methylphenol, 3&4methylphenol, and 2,4-dimethylphenol at MW02 at 32.7, 22.2, 39.8, and 46.3 μ g/L, respectively. Ketones were also detected in both monitoring wells with maximum concentrations of acetone, 2-butanone (MEK), and 4-methyl-2-pentanone (MIBK) at MW02 at 1460, 208, and 12.5 μ g/L, respectively. Acetone, MEK, phenol, 2-methylphenol, 3&4 methylphenol, and 2,4-dimethylphenol were detected in produced water from the Denver-Julesburg Basin.¹⁰¹ MIBK, MEK, and acetone may result from microbial degradation of biopolymers used for hydraulic fracturing.¹⁰¹ Nonylphenol and octylphenol, commonly present in mixtures of ethoxlyated alcohols, were detected in both monitoring wells with maximum concentrations at MW02 at 28 and 2.9 μ g/L, respectively. Ethoxlyated alcohols were used for well stimulation in the Pavillion Field.

Detection of organic compounds, especially those that cannot be attributed to cement, and degradation products of compounds known to have been used for production well stimulation in both MW01 and MW02 provide additional evidence of impact to USDWs and indicate upward solute migration to depths of current groundwater use. Installation of additional monitoring wells at depths similar to MW02, with sample analysis supplemented by state-of-the-art analytical methods better suited to detection of compounds present in stimulation fluids (e.g., liquid chromatography coupled with quadrupole time-of-flight mass spectrometry $^{104-106}$), is necessary to evaluate long-term risk to domestic well users in the Pavillion Field.

Assessment of Potential Impact of Unlined Pits to Domestic Wells. EPA⁷ previously reported disposal of diesel fuel-based (invert) drilling mud and production fluids (flowback, condensate, produced water) in unlined pits in the Pavillion Field and resultant groundwater contamination in surficial Quaternary deposits in shallow monitoring wells sampled by EPA in the vicinity of three unlined pits but did not document the extent of these disposal practices. At least 64 unlined pits were used for disposal of drilling fluids of which invert mud was disposed in 57 pits consisting of up to 79% diesel fuel (Tables SI F1, SI F2). As many as 44 of 64 unlined pits were used or likely used for disposal of production fluids. Unlined pits were emptied and closed in 1995.^{107,108}

A summary of information available on disposal of drilling and production fluids in pits is provided in Table SI F2. This summary includes results of soil and groundwater sampling, excavation volumes and associated criteria (1000–8500 mg/kg total petroleum hydrocarbons), proximity and direction of unlined pits to domestic wells, and recommendations by WOGCC⁴⁴ for further investigation (or no investigation).

The field operator has collected groundwater samples in surficial Quaternary deposits at 12 unlined pit locations.⁴⁴ The highest reported concentrations of GRO and DRO were 91 000 and 78 000 μ g/L, respectively (Figure 5, Table SI F2). Benzene, toluene, ethylbenzene, and xylenes were detected at five locations at concentrations up to 1960, 250, 240, and 1200 μ g/l, respectively (Table SI F2). Thus, sample results indicate impact to surficial groundwater in Quaternary deposits.

There may be as many as 48 domestic wells within 600 m of unlined pits of which 22 domestic wells were sampled by $EPA^{6,8,9,39}$ and 11 were resampled by $WDEQ^{45}$ (Table SI F3). DRO concentrations in domestic wells <600 m from unlined pits likely receiving production fluids were elevated (p = 0.003) compared to domestic wells >600 m from unlined pits (Figure 5a). DRO was detected at 752 mg/kg in a reverse osmosis filter sample from a domestic well (PGDW20) during the Phase II sampling event⁸ (Table SI F3). Concentrations of DRO in domestic wells generally decreased with depth (Figure 5b). Another potential source of DRO in some domestic wells (Table SI G1) is invert mud remaining in boreholes. However, differentiation from other source terms (unlined pits and stimulation) is not possible with currently available data (SI Section G.1).

At two domestic wells (PGDW05 and PGDW30), chromatograms for DRO analysis suggest a diesel fuel source (Figure SI F1a, b). Chromatograms of aqueous (Figure SI F2a) and carbon trap samples (Figure SI F2b) for DRO at another domestic well (PGDW20) indicated the presence of heavy hydrocarbons in water. All three domestic wells are located near unlined pits likely used for disposal of production fluids.

Adamantanes were detected at low aqueous concentrations (<5 μ g/L) at four domestic wells (PGDW05, PGDW20, PGDW30, and PGDW32) (Table SI F3). Admantane, 2-methyl adamantane, and 1,3-dimethyladamantane were detected in a reverse osmosis filter sample at PGDW20 at concentrations of 420, 9400, and 2960 μ g/kg, respectively. Adamantanes were detected in produced water up to 74 mg/L (Table SI D3) indicating disposal in unlined pits as a potential source term. The inherent molecular stability of admantanes and other

diamondoid compounds imparts thermal stability resulting in enrichment in manufactured petroleum distillates.¹⁰⁹ Diamondoids are resistant to biodegradation^{110,111} resulting in their use as a fingerprinting tool to characterize petroleum and condensate induced groundwater contamination.¹¹²

2-Butoxyethanol was detected at 3300 μ g/L in a domestic well (PGDW33)⁴⁵ (Table SI F3). The depth of this domestic well is only 9.1 m bgs and is located within 134 m of an unlined pit used for disposal of production fluids. Other compounds, including BTEX, associated with production well stimulation (e.g., isopropanol) were detected at lower concentrations (<10 μ g/L) in other domestic wells (Table SI F3). Sample results at domestic wells suggest impact from unlined pits and the immediate need for further investigation including installation of monitoring wells in the Wind River Formation. Since flood irrigation is common in the vicinity of unlined pit areas, the lateral extent of groundwater contamination is potentially greater in the Wind River Formation than in overlying surficial Quaternary deposits due to "plume diving" (i.e., uncontaminated water overlies portions of a contaminant plume).^{113–115}

Our investigation highlights several important issues related to impact to groundwater from unconventional oil and gas extraction. We have, for the first time, demonstrated impact to USDWs as a result of hydraulic fracturing. Given the high frequency of injection of stimulation fluids into USDWs to support CBM extraction and unknown frequency in tight gas formations, it is unlikely that impact to USDWs is limited to the Pavillion Field requiring investigation elsewhere.

Second, well stimulation in the Pavillion Field occurred many times less than 500 m from ground surface and, in some cases, at or very close to depths of deepest domestic groundwater use in the area. Shallow hydraulic fracturing poses greater risks than deeper fracturing does,^{57,116} especially in the presence of well integrity issues^{117,118} as documented here in the Pavillion Field. Additional investigations elsewhere are needed.

Finally, while disposal of production fluids in unlined pits is a legacy issue in Wyoming, this practice has nevertheless caused enduring groundwater contamination in the Pavillion Field. Impact to groundwater from unlined pits is unlikely to have occurred only in the Pavillion Field, necessitating investigation elsewhere.

ASSOCIATED CONTENT

S Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.est.5b04970.

Supplemental discussion and tables summarizing data sets are provided in the Supporting Information (SI) portion of the paper (PDF)

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REFERENCES

(1) U.S. Energy Information Administration. *Annual Energy Outlook* 2015 with Projections to 2040; Office of Integrated and International Energy Analysis, U.S. Department of Energy: Washington, DC, April 2015; http://www.eia.gov/forecasts/aeo/pdf/0383(2015).pdf.

(2) U.S. Energy Information Administration. *Annual Energy Outlook* 2012 with Projections to 2035; Office of Integrated and International Energy Analysis, U.S. Department of Energy: Washington, DC, June 2012; http://www.eia.gov/forecasts/aeo/pdf/0383(2012).pdf.

(3) U.S. District Court for the District of Wyoming. Order on Motions for Preliminary Injunction, Case No. 2:15-CV-043-SWS. Petitioners: States of Wyoming, State of Colorado and Intervenor-Petitioners: State of North Dakota, State of Utah, and Ute Indian Tribe vs. U.S. Department of the Interior. Respondents, Sierra Club, Earthworks, Western Resource Advocates, Conservation Colorado Education Fund, the Wilderness Society and Southern Utah Wilderness Alliance. Intervenor-Respondents: Independent Petroleum Association of American and Western Energy Alliance. Filed September 30, 2015; http://www.wyd.uscourts.gov/pdfforms/ orders/15-cv-043%20130%20order.pdf (accessed 3/1/2016).

(4) U.S. Environmental Protection Agency. Evaluation of Impacts to Underground Source of Drinking Water by Hydraulic Fracturing of Coalbed Methane Reservoirs, Office of Water, Office of Ground Water and Drinking Water (4606M), EPA 816-R-04-003, June 2004.

(5) U.S. Environmental Protection Agency. Permitting Guidance for Oil and Gas Hydraulic Fracturing Activities Using Diesel Fuels: Underground Injection Control Program Guidance #84, Office of Water, EPA 816-R-14-001, February 2014.

(6) U.S. Environmental Protection Agency. Site Inspection – Analytical Results Report Pavillion Area Groundwater Investigation Site, Pavillion, Fremont County, Wyoming, CERCLIS ID# WYN000802735, URS Operating Services, Inc., START 3, EPA Region 8, Contract No. EP-W-05-050, August 2009. http://www2.epa.gov/sites/production/ files/documents/Pavillion GWInvestigationARRTextAndMaps.pdf.

(7) Folger, P.; Tiemann, M.; Bearden, D. M. *The Draft Report of Groundwater Contamination Near Pavillion*, Congressional Research Service, R42327; Main Findings and Stakeholders Responses, WY, January 26, 2012.

(8) U.S. Environmental Protection Agency. Expanded Site Investigation – Analytical Results Report Pavillion Area Groundwater Investigation, Fremont County, START 3, EPA Region 8, Contract No. EP-W-05-050; Superfund Technical Assessment and Response Team: WY, August 30, 2010. http://www2.epa.gov/sites/production/ files/documents/PavillionAnalyticalResultsReport.pdf.

(9) DiGiulio, D. C., Wilkin, R. T., Miller C., Oberley, G. Investigation of Ground Water Contamination near Pavillion, Wyoming – Draft Report; U.S. Environmental Protection Agency, Office of Research and Development, National Risk Management Research Laboratory, Ada, OK and Region 8, Denver CO, December 2011 http://www2.epa. gov/region8/draft-investigation-ground-water-contamination-nearpavillion-wyoming.

(10) Wyoming Oil and Gas Conservation Commission. *Four page question sheet submitted to EPA at Stakeholders meeting in Riverton, WY*, November 30, 2011. http://wogcc.state.wy.us/pavillionworkinggrp/PWG_Meeting_Part_B_11302011.pdf.

(11) Wyoming Oil and Gas Conservation Commission. Moving Forward in Evaluating Contamination in the Pavillion Gas Field. Presented to EPA at Stakeholders meeting in Riverton, WY, November 30, 2011. http://wogcc.state.wy.us/pavillionworkinggrp/PWG_ Meeting Part B 11302011.pdf.

(12) Wyoming Oil and Gas Conservation Commission. Evaluating Potential Sources of Contamination in the Pavillion Gas Field. Presented to EPA at Stakeholders meeting in Riverton, WY, November 30, 2011. http://wogcc.state.wy.us/pavillionworkinggrp/PWG_Meeting_Part_ B 11302011.pdf. (13) Wyoming Water Development Office. Review of U.S. EPA Deep Monitoring Well Laboratory Data (Phase III and Phase IV) and Well Construction and Development Data (2010–2011) Pavillion Area, Fremont County, Wyoming, Cheyenne, Wyoming, November 29, 2011. http://wogcc.state.wy.us/pavillionworkinggrp/PWG_Meeting_ Part B 11302011.pdf.

(14) Encana Oil & Gas (USA) Inc. Letter to James Martin, EPA Region 8 Administrator from John Schoup, re: EPA Pavillion Field Investigation, December 21, 2011. http://wogcc.state.wy.us/ pavillionworkinggrp/

Encana%20ltr%20to%20James%20Martin%20EPA%2012212011.pdf.

(15) Encana Oil & Gas (USA) Inc. Letter to James Martin, EPA Region 8 Administrator from John Schoup, re: EPA Pavillion Field Investigation, December 22, 2011. http://wogcc.state.wy.us/ p a villion w or k in g g r p /

Encana%20ltr%20to%20James%20Martin%20EPA%2012222011.pdf.

(16) Encana Oil & Gas (USA) Inc. Letter to James Martin, EPA Region 8 Administrator from John Schoup, re: Testimony during Hearing before the House Subcommittee on energy and Environment, February 7, 2012.

(17) Encana Oil & Gas (USA) Inc. Initial Comments on U.S. Environmental Protection Agency Draft Report "Investigation of Ground Water Contamination Near Pavillion, Wyoming" (December 2011), EPA Docket ID No. EPA-HQ-ORD-2011-0895, April 18, 2012.

(18) Itasca Denver, Inc. Review of the United States Environmental Protection Agency DRAFT Report Entitled "Investigation of Ground Water Contamination near Pavillion, Wyoming", EPA Docket ID No. EPA-HQ-ORD-2011-0895; Prepared for Encana Oil & Gas (USA) Inc., Denver CO, March 2012.

(19) Itasca Denver, Inc. Evaluation of Groundwater-Quality Data Collected by the USEPA and USGS during Phase V Sampling of Wells – Pavillion Area. Prepared for Encana Oil & Gas (USA) Inc., Denver CO, January 10, 2013.

(20) Newfields Denver, CO. Critical Review of ATSDR's Health Consultation: Evaluation of Contaminants in Private Residential Well Water, Pavillion, Wyoming (August 31, 2010), November 2011.

(21) Stimulation Petrophysics Consulting, LLC. A Review and Evaluation of EPA's Drilling and Cement Jobs for EPA's MW01 and MW02 Monitoring Wells in the Pavillion Field, Wyoming, EPA Docket ID No. EPA-HQ-ORD-2011-0895; Prepared for Encana Oil & Gas (USA) Inc., April 2012.

(22) American Petroleum Institute. API's Review Shows EPA's Monitoring Wells at Pavillion, Wyoming are Improperly Constructed and Unsuitable for Groundwater Quality Assessment. 2012 http://www. api.org/~/media/Files/Policy/Hydraulic_Fracturing/Pavillionreviews/API-Fact-Sheet-on-Pavillion-Monitoring-Wells-2013.pdf.

(23) American Petroleum Institute. API's Review of Recent USGS Pavillion, Wyoming. Reports Show USGS Groundwater Sampling Results Differ from EPA's Results in 2011 Draft Report. October 2012. http:// www.api.org/~/media/Files/News/2012/12-October/Pavilion_ Review_v4.pdf.

(24) American Petroleum Institute. Comments of the American Petroleum Institute on EPA's Notice of Extension of the Public Comment Period of the draft Research Report titled, "Investigations of Ground Water Contamination near Pavillion, Wyoming." (78 Federal Register 2396) – Docket ID No. EPA-HQORD-2011–0895, July 1, 2013. http://www. api.org/~/media/files/policy/hydraulic_fracturing/pavillion-reviews/ 13-july-1-api-letter-to-epa-docket-on-draft-pavillion-report.pdf.

(25) American Petroleum Institute. American Petroleum Institute's Review of EPA's Pavillion December 8, 2011Draft Report with Focus on Monitoring Well Drilling, Completion, Development, And Sampling Activities Related to Deep Monitoring Wells MW-01 and MW-02. Prepared by: The American Petroleum Institute. 1220 L Street, NW, Washington, DC, June 25, 2013; 20005–4070, http://www.api.org/ ~/media/Files/Policy/Hydraulic_Fracturing/Pavillion-reviews/API-Report-on-Pavillion-WY.pdf.

(26) Gradient. Review of US EPA and USGS Groundwater Sampling Data Reports Pavillion, Wyoming, Prepared for Halliburton Energy Services, Inc., P.O. Box 42806, Houston, TX, August 1, 2013; pp

77242–2806, http://yosemite.epa.gov/sab/sabproduct.nsf/ 290F553A676E920885257C2400694519/\$File/11+-+Gradient+Review+of+Pavillion+Sampling+Data+Reports+2013.pdf.

(27) Gradient and Environmental Resources Management. Review of US EPA's "Draft Investigation of Ground Water Contamination near Pavillion, Wyoming"; Prepared for Halliburton Energy Services, Inc.: Houston, TX, June 25, 2012. EPA Docket ID No. EPA-HQ-ORD-2011-0895.

(28) S.S. Papadopulos & Associates Inc. (Cohen, H.A.; Grigsby, F.B; Bessinger, B.D.) in Association with QA/QC Solutions (McAteer Jr, J.J.) and Echelon Applied Geoscience Consulting (Baldassare, F.) Review of U.S. EPA's December 2011 Draft Report "Investigation of Ground Water Contamination near Pavillion, Wyoming. Prepared for The Independent Petroleum Association of America, Washington D.C., April 26, 2012.

(29) Kelly, S. Comments on EPA Draft Research Report: Investigation of Ground Water Contamination near Pavillion, Wyoming, March 8, 2012; EPA Docket ID No. EPA-HQ-ORD-2011-0895.

(30) Kelly, S. Comments on EPA Draft Research Report: Investigation of Ground Water Contamination near Pavillion, Wyoming, Phase V Sampling Event, January 8, 2013. EPA Docket ID No. EPA-HQ-ORD-2011–0895.

(31) Stephens, D. B. Analysis of the groundwater monitoring controversy at the Pavillion, Wyoming natural gas field. *Groundwater* **2015**, 53 (1), 29–37.

(32) U.S. Department of the Interior. Bureau of Land Management. Letter to James B. martin, Regional Administrator, EPA Region 8, Denver, CO from Donald A. Simpson, State Directory, March 1, 2012.

(33) American Water Works Association. Letter to Cynthia Sonich-Mullin, Director of the National Risk Management Research Laboratory, EPA Office of Research and Development from Thomas W. Curtis, Deputy Executive Director, January 27, 2012.

(34) New York State Water Resources Institute. A Perspective on the USEPA Study of Pavillion, Wyoming-Relevance and Lessons for NY, January 16, 2012.

(35) Delaware Riverkeeper Network. Review of EPA Investigation of Ground Water Contamination near Pavillion, Wyoming, March 9, 2012.

(36) Natural Resources Defense Council. Sierra Club, Oil and Gas Accountability Project, Wyoming Outdoor Council, Review of Draft: Investigation of Ground Water Contamination near Pavillion Wyoming Prepared by the Environmental Protection Agency, Ada, OK, May 1, 2012.

(37) New Mexico Citizens for Clean Air & Water. Comments on Draft Investigation of Ground Water Contamination near Pavillion, Wyoming, February 1, 2012.

(38) U.S. Environmental Protection Agency. Investigation of Ground Water Contamination near Pavillion, Wyoming Phase V Sampling Event, Summary of Methods and Results. Office of Research and Development, National Risk Management Research Laboratory and Region 8, Denver, CO. September 2012. ftp://ftp.epa.gov/r8/pavilliondocs/ phase5/PavillionSeptember2012Narrative.pdf.

(39) U.S. Environmental Protection Agency. Groundwater Sampling Results at Locations near Pavillion, WY, Pavillion Phase V (April 2012) Groundwater Quality Results and Quality-Control (QC) Data. Office of Research and Development, National Risk Management Research Laboratory and Region 8, Denver, CO. 2012. ftp://ftp.epa.gov/r8/ pavilliondocs/phase5/PavillionSeptember2012Appendices.pdf.

(40) Wright, P. R., McMahon, P. B. Sampling and Analysis Plan for the Characterization of Groundwater Quality in Two Monitoring Wells near Pavillion, Wyoming: U.S. Geological Survey Open-File Report 2012-1197, 2012. http://pubs.usgs.gov/of/2012/1197/.

(41) Wright, P. R., McMahon, P. B., Mueller, D. K., Clark, M. L. Groundwater-Quality and Quality-Control Data for Two Monitoring Wells near Pavillion, Wyoming, April and May 2012; U.S. Geological Survey Data Series 718, 2012. http://pubs.usgs.gov/ds/718/.

(42) Cottrell, G. L. and Myers, D. N. U.S. Geological Service (USGS). Transmittal of Contract Laboratory Results and Evaluation of Laboratory-Specific Quality Control Measures, U.S. Environmental Protection Agency Monitoring Well MW02, Pavillion Wyoming 2012, Administrative Report Prepared for the U.S. Environmental Protection Agency. Director Approved August 30, 2012. ftp://ftp.epa.gov/r8/ pavilliondocs/phase5/USGS_MW02_AdministrativeReportSep2012. pdf.

(43) Wyoming Oil and Gas Conservation Commission. *Pavillion Field Well Integrity Review*, October 8, 2014. http://wogcc.state.wy.us/pavillionworkinggrp/PAVILLION_REPORT_1082014_Final_Report.pdf.

(44) Wyoming Oil and Gas Conservation Commission. *Pavillion Field Pit Review*, November 24, 2014. http://wogcc.state.wy.us/pavillionworkinggrp/Draft%20Pavillion%20Field%20Pits%20Review_11242014.pdf.

(45) Wyoming Department of Environmental Quality. *Pavillion, Wyoming Domestic Water Wells Draft Final Report and Palatability Study*; Prepared by Acton Michelson Environmental, Inc: El Dorado Hills, CA, December 14, 2015; http://deq.wyoming.gov/wqd/ pavillion-investigation/ (accessed 3/1/2016).

(46) EnCana Oil & Gas (USA) Inc.. Submittal of Material Safety and Data Sheets and Letter to U.S. EPA, Region 8, Denver, CO, October 19, 2009.

(47) Natural Resources Defense Council. Freedom of Information Act (FOIA) Online. Accessed on 8/25/2015. https://foiaonline. regulations.gov/foia/action/public/view/request?objectId= 090004d2806a7021.

(48) Daddow, R. L. Water Resources of the Wind River Indian Reservation, Wyoming, U.S, Geological Survey Report 95-4223, 1996. (49) Wyoming Water Development Office. Wyoming State Water

Plan, Wind/Bighorn River Basin Plan, Jan. 14, 2003. http://waterplan. state.wy.us/plan/bighorn/techmemos/grnddet.html.

(50) Morris, D. A.; Hackett, O. M.; Vanlier, K. E.; Moulder, E. A.;Durum, W. H. *Ground-Water Resources of Riverton Irrigation Project Area*, Paper 1375; Geological Survey Water-Supply: WY, 1959.

(51) Plafcan, M., Eddy-Miller, C. A., Ritz, G. F., and Holland, J. P. R. *Water resources of Fremont County*, Water-Resources Investigations Report 95-4095; U.S. Geological Survey: WY, 1995.

(52) Gores and Associates. Pavillion Area Water Supply I Study, Final Report for the Wyoming Water Development Commission, October 2011 http://www.jamesgoresandassociates.com/DocFiles/ Pavillion Area Water Supply Level I Study Final Report.pdf.

(53) Daddow. R. L. Ground-Water and Water Quality Data Through 1991 for Selected Wells and Springs on the Wind River Indian Reservation, Wyoming; U.S. Geological Survey Open-file Report 92-455, 1992.

(54) McGreevy, L. J.; Hodson, W. G.; Rucker IV, S. J. Ground-Water Resources of the Wind River Indian Reservation Wyoming, Geological Survey Water-Supply Paper 1576-1, 1969.

(55) U.S. Environmental Protection Agency, Region 8, Denver, CO. Pavillion Gas Well Integrity Evaluation, July 25, 2013 ftp://ftp.epa.gov/ r 8 / p a v illion d o c s / O ther D o c u m e n t s /

W e l l A n d F i e l d P i t s E v a l u a t i o n J u l y 2 0 1 3 / GasWellIntegrityEvaluation25July2013Final.pdf.

(56) California Council on Science and Technology. Advanced Well Stimulation Technologies in California; Lawrence Berkeley National Laboratory, Pacific Institute, 2014 http://www.ccst.us/publications/ 2014/2014wst.pdf.

(57) Jackson, R. B.; Lowry, E. R.; Pickle, A.; Kang, M.; DiGiulio, D.; Zhao, K. The depths of hydraulic fracturing and accompanying water use across the United States. *Environ. Sci. Technol.* **2015**, *49* (15), 8969–8976.

(58) Adachi, J.; Siebrits, E.; Peirce, A.; Desroches, J. Computer simulation of hydraulic fractures. *International Journal of Rock Mechanics & Mining Science* 2007, 44, 739–757.

(59) Fisher, K.; Warpinski, N. Hydraulic fracture – height growth: Real data. *SPE 145949* **2011**, DOI: 10.2118/145949-MS.

(60) Valkó, P. P.; Economides, M. J. Fluid-leakoff delineation in high permeability fracturing. SPE Prod. & Facilities 1999, 14 (2), 117–130.
(61) Yarushina, V. M.; Bercovici, D.; Oristaglio, M. L. Rock deformation models and fluid leak-off in hydraulic fracturing. Geophys. J. Int. 2013, 194, 1514–152.

Article

(62) Myers, T. Potential contaminant pathways from hydraulically fractured shale to aquifers. *Groundwater* **2012**, *50* (6), 872–882.

(63) Ingraffea, A. R. An Analysis of the "Pavillion Field Well Integrity Review," September 5, 2014. http://wogcc.state.wy.us/ pavillionworkinggrp/publiccomments11202014/PRBRC09052014. pdf.

(64) Maguire-Boyle, S. J.; Barron, A. R. Organic compounds in produced waters from shale gas wells. *Environmental Science Processes & Impacts* 2014, *16* (10), 2237–48.

(65) Hildenbrand, Z. L.; Carlton, D. D.; Fontenot, B. E.; Meik, J. M.; Walton, J. L.; Taylor, J. T.; Thacker, J. B.; Korlie, S.; Shelor, P.; Henderson, D.; Kadjo, A. F.; Roelke, C. E.; Hudak, P. F.; Burton, T.; Rifai, H. S.; Schug, K. A. A Comprehensive Analysis of Groundwater Quality in the Barnett Shale Region. *Environ. Sci. Technol.* **2015**, *49*, 8254–8262.

(66) Colborn, T.; Schultz, K.; Herrick, L.; Kwiatkowski, C. An Exploratory Study of Air Quality near Natural Gas Operations. *Hum. Ecol. Risk Assess.* **2014**, *20* (1), 86–105.

(67) Lothenbach, B.; Le Saout, G.; Gallucci, E.; Scrivener, K. Influence of limestone on the hydration of Portland cements. *Cem. Concr. Res.* **2008**, 38, 848–860.

(68) Lothenbach, B.; Matschei, T.; Möschner, G.; Glasser, F. P. Thermodynamic modeling of the effect of temperature on the hydration and porosity of Portland cement. *Cem. Concr. Res.* **2008**, *28*, 1–18.

(69) Lothenbach, B.; Winnefeld, F. Thermodynamic modeling of the hydration of Portland cement. *Cem. Concr. Res.* **2006**, 36 (2), 209–226.

(70) Brouwers, H. J. H.; van Eijk. Alkali concentrations of pore solution in hydrating OPC. *Cem. Concr. Res.* 2003, 33, 191–196.

(71) Leemann, A.; Lothenbach, B. The influence of potassiumsodium ratio in cement and concrete expansion due to alkai-aggregate reaction. *Cem. Concr. Res.* **2008**, *38*, 1162–1168.

(72) Andersson, K.; Allard, B.; Bengtsson, M.; Magnusson, B. Chemical composition of cement pore solutions. *Cem. Concr. Res.* **1979**, *19*, 327–332.

(73) Thomas, J. J.; Rothstein, D.; Jennings, H. M.; Christensen, B. J. Effect of hydration temperature on the solubility behavior of Ca-, S-, Al-, and Si-bearing solid phases in Portland cement pastes. *Cem. Concr. Res.* **2003**, *33*, 2037–2047.

(74) van Eijk, R. J.; Brouwers, H. J. J. Prediction of hydroxyl concentrations in cement pore water using a numerical cement hydration model. *Cem. Concr. Res.* **2000**, *20*, 1801–1806.

(75) Lothenbach, B.; Le Saout, G.; Gallucci, E. and Scrivener, K. Influence of limestone on the hydration of Portland cements. *Cem. Concr. Res.* **2008**, 38, 848–860.

(76) Lothenbach, B.; Matschei, T.; Möschner, G.; Glasser, F. P. Thermodynamic modeling of the effect of temperature on the hydration and porosity of Portland cement. *Cem. Concr. Res.* **2008**, *28*, 1–18.

(77) Lothenbach, B.; Winnefeld, F. Thermodynamic modeling of the hydration of Portland cement. *Cem. Concr. Res.* **2006**, *36* (2), 209–226.

(78) Rothstein, D.; Thomas, J. J.; Christensen, B. J.; Jennings, H. M. Solubility behavior of Ca-, S-, Al-, and Si-bearing solid phases in Portland cement pore solutions as a function of hydration time. *Cem. Concr. Res.* **2002**, *32*, 1663–1671.

(79) Beak, D. G.; Oberley, G. G.; Ruybal, C. J.; Acree, S. D.; Ross, R. R. Retrospective Case Study in Killdeer, North Dakota: Study of the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources, EPA/600/R-14/103; U.S. Environmental Protection Agency, Office of Research and Development, Washington, DC, May 2015.

(80) Cummins, T. M.; Robbins, G. A.; Henebry, B. J.; Goad, C. R.; Gilbert, E. J.; Miller, M. E.; Stuart, J. D. A Water Extraction, Static Headspace Sampling, Gas Chromatographic Method to Determine MTBE in Heating Oil and Diesel Fuel. *Environ. Sci. Technol.* 2001, 35, 1202–1208.

(81) Deeb, R. A.; Chu, K.-H.; Shih, T.; Linder, S.; Suffer, I.; Kavanaugh, M. C. Alvarez-Cohen, L. MTBE and other oxygenates: Environmental sources, analysis, occurrence, and treatment. *Environ. Eng. Sci.* **2003**, 20 (5), 433–447.

(82) U.S. Environmental Protection Agency. *Monitoring and reporting* of *MTBE and other oxygenates at UST release sites*, memorandum from Director Office of Underground Storage Tanks to Regional UST Program Mangers and State UST Program Managers. Dated January 18, 2000. http://www.epa.gov/oust/mtbe/jan1800.pdf.

(83) Robbins, G. A.; Henebry, B. J.; Schmitt, B. M.; Bartolomeo, F. B.; Green, A.; Zack, P. Evidence of MTBE in heating oil. Groundwater. *Groundwater Monit. Rem.* **1999**, *19* (2), 65–69.

(84) Robbins, G. A.; Henebry, B. J.; Cummins, T. M.; Goad, C. R.; Gilbert, E. J. Occurrence of MTBE in heating oil and diesel fuel in Connecticut. *Groundwater Monit. Rem.* **2000**, *20* (4), 82–86.

(85) Engelsen, C. J. Quality Improvers in Cement Making – State of the Art. COIN Project P1 Advanced Cementing Materials. SP 1.1F Reduced CO₂ missions. COIN Project report 2 – 2008. http://www.sintef.no/upload/Byggforsk/Publikasjoner/coin-no2.pdf.

(86) Ervanne, H., and Hakanen, M. Analysis of Cement Superplasticizers and Grinding Aids, A Literature Survey, Working Report, Posiva, 2007.

(87) Jeknavorian, A. A.; Barry, E. F.; Serafin, F. Determination of grinding aids in Portland cement by pyrolysis gas chromatographymass spectrometry. *Cem. Concr. Res.* **1998**, *28* (9), 1335–1345.

(88) Teoreanu, I.; Guslicov, G. Mechanisms and effects of additives from the dihydroxy-compound class on Portland cement grinding. *Cem. Concr. Res.* **1999**, *29*, 9–15.

(89) Smith, B.; Siegel, D.; Neslund, C.; Carter, C. Organic contaminants in Portland cement used in monitoring well construction. *Groundwater Monit. Rem.* **2014**, *34* (4), 102–111.

(90) U.S. Environmental Protection Agency. *Background Information* for the Leaching Environmental Assessment Framework (LEAF) Test Methods, Appendix C - Preliminary Version of Method 1315, EPA/ 600/R-10/170, November 2010, Office of Research and Development, National Risk Management Research Laboratory, Research Triangle Park, NC.

(91) Wolfe, A. L.; Wilkin, R. T.; Lee, T. R.; Ruybal, C. J.; Oberley, G. G. Retrospective Case Study in the Raton Basin, Colorado Study of the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources; U.S. Environmental Protection Agency Office of Research and Development Washington, DC, EPA/600/R-14/091, May 2015.

(92) Beak, D. G.; Overbay, M. D.; Mravik, S. C. Retrospective Case Study in Wise County, Texas Study of the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources, EPA/600/R-14/090; U.S. Environmental Protection Agency, Office of Research and Development: Washington, DC, May 2015b.

(93) Ludwig, R. D.; Beak, D. G.; Wilkin, R. T.; Ruybal, C. J.; Rectenwald, D. J. *Retrospective Case Study in Northeastern Pennsylvania Study of the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources*, EPA/600/R-14/088; U.S. Environmental Protection Agency, Office of Research and Development: Washington, DC, May 2015.

(94) Wilkin, R. T.; Lee, T. R.; Ruybal, C. J.; Rectenwald, D. J. Retrospective Case Study in Southwestern Pennsylvania Study of the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources, EPA/600/R-14/084; U.S. Environmental Protection Agency, Office of Research and Development: Washington, DC, May 2015.

(95) Cozzarelli, I. M.; Baedecker, M. J.; Eganhouse, R. P.; Goerlitz, D. F. The geochemical evolution of low-molecular-weight organic acids derived from the degradation of petroleum contaminants in ground-water. *Geochim. Cosmochim. Acta* **1994**, *58* (2), 863–877.

(96) Kharaka, Y. K.; Thordsen, J. J.; Kakouros, E.; Herkelrath, W. N. Impacts of petroleum production on ground and surface waters: Results from the Osage-Skiatook petroleum environmental research a site, Osage County, Oklahoma. *Environ. Geosci.* **2005**, *12* (2), 127–138.

(97) Akob, D. M.; Cozzarelli, I. M.; Dunlap, D. S.; Rowan, E. L.; Lorah, M. M. Organic and inorganic composition and microbiology of produced waters from Pennsylvania shale gas wells. *Appl. Geochem.* **2015**, *60*, 116–125.

(98) Cluff, M. A.; Hartsock, A.; MacRae, J. D.; Carter, K.; Mouser, P. J. Temporal changes in microbial ecology and geochemistry in produced water from hydraulically fractured Marcellus Shale gas wells. *Environ. Sci. Technol.* **2014**, *48* (11), 6508–6517.

(99) Orem, W.; Tatu, C.; Varonka, M.; Lerch, H.; Bates, A.; Engle, M.; Crosby, L.; McIntosh, J. Organic substances in produced and formation water from unconventional natural gas extraction in coal and shale. *Int. J. Coal Geol.* **2014**, *126*, 20–31.

(100) Murali Mohan, A.; Hartsock, A.; Hammack, R. W.; Vidic, R. D.; Gregory, K. B. Microbial communities in flowback water impoundments from hydraulic fracturing for recovery of shale gas. *FEMS Microbiol. Ecol.* **2013**, *86* (3), 567–580.

(101) Lester, Y.; Ferrer, I.; Thurman, E. M.; Sitterley, K. A.; Korak, J. A.; Aiken, G.; Linden, K. G. Characterization of hydraulic fracturing flowback water in Colorado: implications for water treatment. *Sci. Total Environ.* **2015**, *512–513*, 637–44.

(102) Olsson, O.; Weichgrebe, D.; Rosenwinkel, K. H. Hydraulic fracturing wastewater in Germany: composition, treatment, concerns. *Environ. Earth Sci.* **2013**, *70*, 3895–3906.

(103) Mägli, A.; Messmer, M.; Leisinger, T. Metabolism of Dichloromethane by the Strict Anaerobe Dehalobacterium formicoaceticum. *Appl. Environ. Microbiol.* **1988**, *64* (2), 646–650.

(104) Ferrer, I.; Thurman, E. M. Analysis of hydraulic fracturing additives by LC/Q-TOF-MS. *Anal. Bioanal. Chem.* **2015**, 407 (21), 6417–6428.

(105) Thurman, E. M.; Ferrer, I.; Blotevogel, J.; Borch, T. Analysis of hydraulic fracturing flowback and produced waters using accurate mass: Identification of ethoxylated surfactants. *Anal. Chem.* **2014**, *86* (19), 9653–9661.

(106) Thurman, E. M.; Ferrer, I.; Rosenblum, J.; Linden, K.; Ryan, J. M. Identification of polypropylene glycols and polyethylene glycol carboxylates in flowback and produced water from hydraulic fracturing. *J. Hazard. Mater.* **2016**, http://dx.doi.org/10.10116/j.jhazmat.2016.02. 041.10.1016/j.jhazmat.2016.02.041

(107) Encana Oil & Gas (USA) Inc. Comments of Encana Oil & Gas (USA) Inc. on the WOGCC Report: Pavillion Field Pit Review (Nov 24, 2014) submitted January 16, 2015 http://wogcc.state.wy.us/pavillionworkinggrp/PitReviewPubComments/Encana20150116CommentstoWOGCCPitRvw.pdf.

(108) Encana Oil & Gas (USA) Inc. Document 2 Summary of Historic Pit Evaluation and Remediation Activities Pavillion Field, Wyoming January 2015 http://wogcc.state.wy.us/ pavillionworkinggrp/PitReviewPubComments/ Encana20150116Doc2CommentsHistPitEval.pdf.

(109) Wingert, W.S. G c-m.s analysis of diamondoid hydrocarbons in smackover petroleums. *Fuel* **1992**, *71*, 37–43.

(110) Grice, K.; Alexander, R.; Kagi, R. I. Diamondoid hydrocarbon ratios as indicators of biodegradation in Australian crude oils. *Org. Geochem.* **2000**, *31*, 67–73.

(111) Williams, J. A.; Bjorøy, M.; Dolcater, D. L.; Winters, J. C. Biodegradation in South Texas Eocene oils – Effects on aromatics and biomarkers. *Org. Geochem.* **1986**, *10*, 451–461.

(112) Stout, S. A.; Douglas, G. S. Diamondoid hydrocarbons - Application in the chemical fingerprinting of natural gas condensate and gasoline. *Environ. Forensics* **2004**, *5*, 225–235.

(113) American Petroleum Institute. Downward Solute Plume Migration: Assessment, Significance, and Implications for Characterization and Monitoring of "Diving Plumes", Regulatory Analysis and Scientific Affairs API Soil and Groundwater Technical Task Force Bulletin 24, April 2006.

(114) United States Environmental Protection Agency (EPA 2001). *Plume Dive Calculator*. Prepared by Jim Weaver for the EPA Office of Research and Development. 2001 http://www.epa.gov/athens/ learn2model/part-two/onsite/diving.htm.

(115) Weaver, J. W.; Wilson, J. T. Diving plumes and vertical migration at petroleum hydrocarbon release sites. *LUSTLine* **2000**, Bulletin *36*, 12–15.

(116) Llewellyn, G. T.; Dorman, F.; Westland, J. L.; Yoxtheimerc, D.; Grievec, P.; Sowersc, T.; Humston-Fulmerd, E.; Brantley, S. L. Evaluating a groundwater supply contamination incident attributed to Marcellus Shale gas development. *Proc. Natl. Acad. Sci. U. S. A.* **2015**, *112* (20), 6325–6330.

(117) Darrah, T. H.; Vengosh, A.; Jackson, R. B.; Warner, N. R.; Poreda, R. J. Noble gases identify the mechanisms of fugitive gas contamination in drinking-water wells overlying the Marcellus and Barnett Shales. *Proc. Natl. Acad. Sci. U. S. A.* **2014**, *111* (39), 14076– 14081.

(118) Jackson, R. B.; Vengosh, A.; Darrah, T. H.; Warner, N. R.; Down, A.; Poreda, R. J.; Osborne, S. G.; Zhao, K.; Karr, J. D. Increased stray gas abundance in a subset of drinking water wells near Marcellus shale gas extraction. *Proc. Natl. Acad. Sci. U. S. A.* **2013**, *110* (28), 11250–11255.