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## Produced Formation Water Sample Results from Shale Plays

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Chesapeake Energy Corporation (Chesapeake) and Environmental Consultants reviewed non-potable produced formation water sample results available from the literature and other sources. The data review was specifically directed at the chemical characteristics of source water used for hydraulic fracturing fluids, hydraulic fracturing fluids, produced formation water from natural gas wells following hydraulic fracturing activities (i.e., less than 30 or 90 days following hydraulic fracturing), and produced formation water from operating natural gas wells.. The largest amount of data represents sampling conducted of produced formation water from the Marcellus Shale, including the GTI study (Hayes, 2009) and the USGS Produced Water database (USGS, 2010). Additional data was available from private sources. This data set included limited data for three other shale formations, the Barnett, Fayetteville and Haynesville. Some of the data evaluated is subject to attorney-client privilege (herein after "privileged data").

The zones in the four shale plays from which natural gas is being produced vary in average depth below the surface from 4,500 feet in the Fayetteville to 11,500 feet in the Haynesville. The average depth from the surface to the natural gas producing zones in each of the four shale plays are thousands of feet below the geologic zones which produce potable groundwater. Furthermore, temperatures and pressures found at these depths vary from 130°F and 3,000 pounds per square inch (psi), respectively, in the Fayetteville to 320°F and 10,000 psi, respectively, in the Haynesville. These temperatures and pressures have marked effects on the sampling and field analysis of produced formation waters and create potential safety issues for sampling personnel.

The primary purpose of the produced formation water data review was to support Chesapeake's baseline water sampling program for potable water. Of major interest was the identification of potential sentinel chemicals and/or radiochemicals that could be included in the baseline potable water sampling program. Further, the review was intended to assist in decision making regarding disposal or treatment for reuse or recycling of produced formation waters.

Data available from the GTI study included general chemistry parameters and metals as well as volatile organic compounds and semi-volatile organic compounds from 70 wells in the Marcellus. The data evaluation included a review of chemicals currently being used in Chesapeake wells during hydraulic fracturing. Initially, Chesapeake had chosen the parameters for analyses based on their potential mobility in groundwater water systems, toxicity, and the availability of analytical methods. Eventually, the complete list found at 40 CFR Part 264, Appendix IX, and commonly associated with groundwater monitoring, was selected. The complete list was specified because many of the chemicals in the treatment fluids are proprietary or not disclosed in information available to Chesapeake. All of the analyses were conducted using EPA analytical methods and analyses were performed by certified laboratories. The majority of analyses were conducted by one laboratory.

Some additional data was available in the privileged data for special volatile organic compounds, primarily glycols, and radiochemistry data. The glycols had been included because of the frequency of use in hydraulic fracturing fluids and in order to evaluate their usefulness as sentinel species.

Both the authors of the GTI study and the sources for the privileged data indicated that there are significant issues regarding sampling of produced formation water. Natural gas is contained under high pressure in specialized equipment that is not designed for producing high quality environmental samples. The sample matrix itself presents challenges such as foaming and changes in surface tension.

Analytical techniques are also impacted by the presence of elevated concentrations of total dissolved solids and chloride. The inorganic and wet chemistry methods were most affected by the presence of high total dissolved solids. Radiochemistry methods are particularly affected by the elevated concentrations of barium and total dissolved solids.

As anticipated total dissolved solids and the divalent cations, including barium and strontium, are elevated in produced formation water samples. The concentrations noted in the data evaluated are consistent with that gleaned from the literature (USGS, 2010). Concentrations of these parameters appear to increase with time following hydraulic fracture activities and remain at high concentrations in the produced formation water from operating wells in each shale formation. The concentrations of barium and strontium appear to correlate to the concentrations of total dissolved solids. Chloride represented the most abundant anion. These data are consistent with the generalized data available in the USGS database.

The literature supports the interaction and release of arsenic from host rock in the presence of refined hydrocarbons products, therefore, a specific review of the arsenic data was conducted. In general, arsenic concentrations were not found in many of the samples above method detection limits. The GTI study found some detectable concentrations of arsenic with produced formation water sampled on day five following hydraulic fracturing having the highest concentration, 124 µg/L. Data available from analysis of produced formation water from the Marcellus prior to disposal confirmed the general absence of arsenic. There were four detected concentrations from 87 fluids samples. The highest detected concentration was 4.2 µg/L. The limited data available for the Barnett, Fayetteville and Haynesville formations showed not detected concentrations of arsenic; detection limits ranged from 1 to 10 µg/L.

In consultation with health physicists from SAIC and IEM, Chesapeake has determined that radium 226 and 228 represent the most useful parameters to provide meaningful information regarding radionuclides. This decision is based on the fact that radium 226 and 228 are more soluble in water, are more potentially mobile, and represent 80 percent of the potential ingested dose of radiation in drinking water. Based on experience with radionuclide analyses in produced formation water, it appears that gamma spectroscopy (EPA Method 901.1) and isotopic analyses of the select decay series provide the most accurate and usable information. It is important for the activity results to be presented accompanied by the standard deviation and minimum detected concentrations in order to put the data in context, particularly for non-detected activity levels.

Certain volatile organic compounds, namely glycols, were also evaluated. Glycols were selected because of their frequency of use in treatment fluids used in natural gas production. EPA Method 8015 has been found to be insufficiently robust to overcome the matrix issues which are attendant to the analysis of produced formation water. The range of detection limits seen in the available data sets ranged from 10,000 to 50,000 µg/L, which does not provide meaningful results. EPA Method 8270 has some utility for larger glycols, such as glycol ethers; however for the smaller, more soluble, ethylene and diethylene glycols, the extraction methods are not useful. Sample results for produced formation

water prior to disposal had similar issues with elevated detection limits even when total dissolved solids levels were lower. In these samples, ethylene glycol was found in 14 of 87 samples. Glycols may prove to be problematic for use as sentinel compounds because of their other uses in natural gas production, such as winterization of equipment similar to their use in automobiles, recreational vehicles, etc.

Measured concentrations of volatile organic compounds were available for the Marcellus from the GTI study and more limited data had been provided for the Barnett, Fayetteville and Haynesville formations. The most consistently detected volatile organic compounds were benzene, toluene, ethyl benzene and xylenes (BTEX). This finding is consistent with data available in the literature and would be anticipated as these compounds are closely associated with hydrocarbon producing geologies from which the produced formation water comes. There were no trends evident for the BTEX data. The appearance of BTEX was somewhat inconsistent and when present was internally inconsistent. That is, the highest concentrations of benzene were not in the same samples as the highest concentrations of the other three. In the iterative sampling, there was no consistent timing for occurrence of the highest concentration. This may well be due to the differences in water solubility in the presence of elevated total dissolved solids for the individual BTEX components. The data is suggestive that there are differences in BTEX levels between and within shale formations. There were few other volatile organic compounds found in produced formation water; however, they were found on an inconsistent basis.

The semi-volatile organic compounds (SVOCs) were generally not present in detectable concentrations. The most frequently detected SVOC was pyridine in both the GTI Study and in the other data available. In the GTI study data set, six of 70 samples had detectable levels of pyridine in the hydraulic fracture fluids. The author speculated that presence of pyridine was due to its use as a precursor in the manufacture of one of the hydraulic fracturing additives. Certain service companies have alkyl pyridines available as corrosion inhibitors (Weatherford, 2011). In the additional data evaluated, only two of eleven sampled hydraulic fracturing fluids had detectable concentrations of pyridine. One of these the concentration was related to the reuse of produced formation water as makeup water for the hydraulic fracturing fluid. Pyridine has been reported as naturally occurring in oil shales (ATSDR, 1992). The additional data sets may indicate support mobilization of the naturally-occurring pyridine since it is soluble in water or may represent a degradation product. There appear to be differences between the shale plays; however, the data set is small and therefore, does not support the development of generalizations.

The iterative (time series) sample results presented by GTI and in other available data does not reveal a consistent trend of increasing or decreasing concentration of volatile organic compounds or SVOCs with increasing times after hydraulic fracturing.

The major conclusions from the review of produced formation water data are:

- Produced formation water appears to be highly variable within and between shale plays.
- Analytical techniques used for chemical and radiochemical characterization of the produced formation water must be robust to the matrix interferences presented by total dissolved solids.
- Few volatile organic compounds and SVOCs are consistently found in produced formation waters. Benzene, ethyl benzene, toluene and xylenes are expected to be present in varying concentrations in natural gas. The concentrations of volatile organic compounds and SVOCs are generally low and tend to preclude their use as sentinel chemicals.
- The most reliable sentinel compounds appear to be total dissolved solids, chloride and divalent cations.

References:

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