

To: Barbara Miller **Date:** January 6, 2022

(Ameren Missouri)

From: XDD (DRAFT) cc: Michael Marley (XDD)

RE: Rush Island Treatability Memo

Rush Island Energy Center

100 Big Hollow Road, Festus, MO

1.0 INTRODUCTION

XDD Environmental (XDD) was retained by Ameren Missouri (Ameren) to perform a metals treatability study for the remediation of metals of concern (MOC) from ash pond groundwater at the Rush Island Energy Center (RIEC) in Festus, MO. Groundwater is currently monitored as required by the United States Environmental Protection Agency (USEPA) in 40 CFR Part 257 "Hazardous and Solid Waste Management System; Disposal of Coal Combustion Residuals from Electric Utilities; Final Rule" (the CCR Rule), the facility's NPDES permit (No MO-000043), and the facilities UIC permit (UI-0000043). Groundwater is analyzed for metals via EPA Methods 200.7 and 200.8, alkalinity via SM Method 2320B, total dissolved solids via SM Method 2450C, ferric and ferrous iron via SM Method 3500, anions via EPA Method 300, and phosphorous via EPA Method 365.4. Based on statistical analysis, elevated levels of arsenic and molybdenum exceed site-specific groundwater protection standards (GWPS) established under the CCR Rule.

While metals cannot be destroyed, they can be susceptible to treatment and undergo changes in form to become either (a) less soluble; or (b) more sorbent and bind to particle surfaces. Both methods involve the physical removal of metals from the dissolved state (a very mobile state) to either a solid state or an adsorbed state. The three primary approaches for metals removal from groundwater are:

- <u>Precipitation:</u> Transformation of a dissolved species to a solid form, which can then settle out of suspension.
- <u>Co-precipitation with other minerals</u>: Transformation of a dissolved species to a solid form that combines with another material (such as iron), which can then settle out of suspension.
- <u>Adsorption</u>: Introduction or production of a solid that will absorb the MOC from the groundwater.

Where multiple metals are present, there is a potential that one metal can inadvertently affect



other metal(s) either positively (reduced dissolved concentration) or negatively (increased dissolved concentration). Therefore, a treatment chain consisting of a sequence of multiple technologies is often needed to address all metals of concern at a site. The focus of this study is on MOC which have regulatory action levels exceeded at RIEC downgradient of the ash pond.

Since the treated water will be injected into the extracted aquifer for hydraulic control, the treated water also will be required to meet groundwater permit levels for compounds such as sulfate and boron, along with the primary MOC (arsenic and molybdenum), which currently exceed permit discharge levels. The primary MOC at RIEC include arsenic and molybdenum.

This memo will address the treatability work performed for RIEC water to remove the two primary MOC along with an evaluation of sulfate and boron removal to meet the discharge permit levels. The treatability lab testing for RIEC was done in parallel with the treatability studies performed for Labadie Energy Center (LEC) and Sioux Energy Center (SEC). All three sites have overlap on treatment objectives and similar water chemistry; therefore, promising results at one site were used to guide treatment at the other two sites.

Preliminary evaluations focused on in-situ treatment options; however, due to concerns of precipitation clogging pore space thereby affecting subsurface flow conditions, along with the lack of available in-situ options for removing or stabilizing boron, the focus transitioned to ex-situ treatment options. This memo will focus on the ex-situ treatment options only. The findings presented will focus on RIEC but will include relevant results obtained from treatability tests from LEC and SEC.

2.0 TREATABILITY OBJECTIVE

The treatment objective for RIEC is to create hydraulic controls where groundwater is extracted along the flow path at the site to capture and contain the groundwater MOC plume. The water will then be treated above ground and reinjected between the extraction wells to create a hydraulic control that will minimize the extraction rate, maintain natural hydraulic flow, and prevent further migration of MOC from the ash pond groundwater. The focus of the treatability study is to remove the site MOC so that the water will meet the groundwater permit discharge levels. This will allow safe injection of the treated groundwater back into the aquifer from which it was extracted. For RIEC, the proposed treatment system must address arsenic, molybdenum, sulfate, and boron. The target discharge permit levels for these compounds are:

- arsenic = 10 micrograms per liter (μg/L)
- molybdenum = 100 μg/L
- sulfate = 250,000 μg/L
- boron = 2,000 μg/L

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This memo will summarize the tests performed to address each compound, conditions under which each method was tested, how successful each remedial option was shown to be, and recommendations for full scale implementation.

2.1 Arsenic and Molybdenum

At RIEC, arsenic is the primary MOC that causes groundwater protection exceedances. The highest concentrations are greater than 10x the discharge permit limit (**Table 1**). The primary focus of arsenic removal is on its lower solubility formations at low pH ranges and its ability to coprecipitate with iron. Molybdenum is also present above permit discharge levels at RIEC with varying concentrations. Through testing, it was determined that processes shown to treat arsenic were also successful at removing molybdenum. The two MOC are therefore discussed in unison.

Tests were performed for precipitation at a pH of 5, filtering the water through zero valent iron (ZVI) columns, pH adjustment followed by ferric chloride (FeCl₃) precipitation, and pH adjustment followed by FeCl₃ and oxidative (hydrogen peroxide) precipitation.

2.1.1 pH adjustment

The pH adjustment process involved adjusting the pH of RIEC water to 5 using hydrochloric acid (HCl) followed by flow through a sand filter than a SIR-150 boron resin filter (see Section 2.2 for more information on SIR-150). Since the resin is specially designed for boron removal, it is not desirable to have the resin capacity exacerbated with high concentrations of other groundwater constituents. Samples were therefore collected prior to the resin to determine the effects of the pH adjustment to arsenic and molybdenum treatment (**Table 1, Test A,M-1**).

Groundwater samples collected between the sand filter and the resin columns showed that the pH adjustment followed by sand filtration did not effectively remove the arsenic and molybdenum in the groundwater to the discharge permit levels (**Table 1**, **Test A,M-1**). A further review of the data and the procedures used in this test suggests that, for pH adjustment to be successful for removing arsenic and molybdenum from RIEC groundwater, the groundwater needs to be maintained at a reduced pH for longer than 40 minutes (the residence time in the sand filter tested columns). Hence, pH adjustment alone would not be a viable ex-situ remedial approach.

2.1.2 ZVI Columns

Iron can precipitate molybdenum as a low-solubility iron-arsenic and iron-molybdenum. To test the feasibility of using a ZVI column to remove arsenic and molybdenum ex-situ, test columns were constructed using a mixture of the microscale ZVI and commercial sand. Microscale ZVI was tested



due to its highly reactive surface and increased potential to remove arsenic and molybdenum. The sand was added to allow the required flow through the ZVI column without clogging due to the microscale ZVI particle size. The columns were prepared using a 5:1 and 2:1 ratios of sand to microscale ZVI. The columns were operated for 7 days, with effluent groundwater samples collected from the column after 1, 3, and 7 days of flow.

The Day 7 results are shown on **Table 1 Tests A,M-2** (5:1 sand to ZVI ratio) and **A,M-3** (2:1 sand to ZVI ratio). The results show partial removal of arsenic and molybdenum, though not to action levels. Boron concentrations did not change passing through the ZVI columns though research suggested there may be a reduction. It was concluded that the ZVI removal effectiveness (at the design sand to ZVI dosages) was questionable, and likely not reliable as a sustainable remedial option.

2.1.3 <u>Ferric Chloride Co-Precipitation</u>

The ex-situ remediation method that proved most successful and reliable for arsenic and molybdenum treatment of groundwater is pH adjusted (using HCl) followed by FeCl₃ aided flocculation. Preliminary testing with the ZVI and pH adjustment, discussed above, helped guide the design of the FeCl₃ treatment train.

FeCl₃ testing was performed to determine reaction time needed (**Table 1 Tests A,M-4** to **A,M-12**), optimal pH ranges (**Table 1 Tests A,M-7** to **A,M-12**), optimal FeCl₃ dosage (**Table 1 Tests A,M-4** to **A,M-9**), and if oxidation through hydrogen peroxide addition could perform better (**Table 1 Test A,M-13**). A summary of the finding are:

- Both arsenic and molybdenum can be reduced to concentrations at or below action levels, using FeCl₃ addition.
- An initial pH of 6 (prior to the addition of FeCl₃) caused faster settling of the precipitants than an initial pH of 4.
- Higher FeCl₃ dosage (40 mg/L vs. 20 mg/L) provided greater removal of arsenic and molybdenum.
- The additional of hydrogen peroxide did not improve the arsenic removal efficiency.

The reaction time determined for the FeCl₃ coagulation and flocculation/precipitation and associated removal of arsenic and molybdenum from groundwater in the preliminary testing was adequate for the conceptual ex-situ remedial approach (an hour or less).



2.2 Boron

Resins and reverse osmosis are the primary methods used to remove boron from water sources. There is a low efficiency of treated water to wastewater using reverse osmosis, so the boron treatment has focused on resins. Three commercially available boron-removal resins were identified and tested; SIR-150, IRA-743, and PWA-10.

As discussed in Section 2.1.1, a pH of 5 adjusted water followed by a column of the SIR-150 was tested over 7 days. After 7 days of passing through the column, boron concentrations were still below the detection limit of 10 ug/L (**Table 1 Test B-1**). As shown in the sample collected between pH adjustment to 5 and the resin column (**Table 1 Test A,M-1**), arsenic and molybdenum were not treated by the pH adjustment showing that all the arsenic and molybdenum removed in **Table 1 Test B-1** is from attachment to the resin. By Day 7, arsenic began to break through and the resin showed visual discoloration.

While the resin beds in this treatability test were designed for a 15 minute residence time, when scaling the lab test to field application flow rates, the mass of resin was much larger than would be used in a site application. The breakthrough of arsenic after 7 days suggests that, given field-sized vessels, the resin utilization would be too high without pre-treatment for arsenic and molybdenum prior to resin filtration.

Subsequent testing used the procedure discussed in Section 2.1.3 prior to filtration through the resin beds. The process involved RIEC water adjusted to a pH of 6 with HCl followed by FeCl₃ (40 mg/L), settling, and sand filtration. The water was then passed through one of three columns containing SIR-150 (**Table 1 Test B-2**), IRA-743 (**Table 1 Test B-3**), or PWA-10 (**Table 1 Test B-4**). While all resins were capable of removing the boron and polishing the arsenic and molybdenum, SIR-150 showed the highest performance for sulfate removal. Additional temporal testing was performed on the SIR-150 and PWA-10 resins for sulfate removal along with surfactant coated zeolite (**Table 1 Tests B,S-1** to **B,S-24**). These tests are discussed further in Section 2.3.2.

2.3 Sulfate

2.3.1 Sorption and Precipitation

Sulfate can be precipitated out at high concentrations (thousands of milligrams per liter (mg/L) concentration ranges) but is difficult to reduce at lower concentrations (hundreds of mg/L concentration ranges). The primary approaches used for the lower level concentration of sulfate are reverse osmosis and resin removal. As mentioned for boron, there is a low efficiency of treated water to wastewater using reverse osmosis. The cost, maintenance, and waste stream of resins are such that it would be preferable to not rely on a second resin bed for sulfate removal which would result in a second resin waste stream. Other options discussed in literature were:



- sorption using
 - zeolite (Table 1 Tests S-1, S-5, S-22);
 - o surfactant coated zeolite (Table 1 Tests S-2, S-6, S-11, S-12, S-19 to S-21);
 - sodium chloride coated zeolite (Table 1 Tests S-3, S-7, S-9, S-10).
- precipitation using
 - limestone (Table 1 Tests S-4, S-8);
 - cement (Table 1 Tests S-13 to S-15);
 - o calcium hydroxide (Table 1 Tests S-16 to S-18);
 - o chitosan (Table 1 Tests S-23 to S-25);
 - sodium aluminate (Table 1 Test S-26);
 - calcium aluminate (Table 1 Test S-27);
 - calcium aluminate cement (Table 1 Test S-28).

Since the success of these processes are sensitive to water quality and water chemistry (alkalinity, pH, total dissolved solids, etc.), several dosages, treatment train applications, product formulations, and pH ranges were tested to identify if there was a method to make them successful.

Of the methods and conditions tested, none were shown to greatly reduce the sulfate concentration relative to the baseline value.

2.3.2 <u>Resin</u>

Based on the results of RIEC pilot study which was conducted in parallel with the sulfate portion of the lab treatability testing, approximately 20% of the influent sulfate can be removed through the FeCl₃ coagulation/flocculation process and is removed after the sand filtration. This reduction brings the sulfate concentrations at RIEC below the discharge permit levels. In addition, the pilot study showed an average of 58% reduction in sulfate between the influent and post resin treatment (**Table 1 Tests P1** to **P20**).

Based on the lack of success of the options in Section 2.3.1 and the ability of the current proposed treatment train to treat RIEC sulfate concentrations to below discharge permit levels (250,000 μ g/L), the best option would be to use the treatment train developed for arsenic and molybdenum to decrease the sulfate concentrations and then polish the water with the boron resin.

As mentioned in Section 2.2, there were three resins identified that could remove boron and preliminary results suggested that sulfate coated zeolite could reduce sulfate. Temporal testing was performed to evaluate the longevity of two of the resins (SIR-150 and PWA-10) and the surfactant



coated zeolite. Columns were constructed that were scaled to the proposed full scale vessel sizes for the lab defined flow rate and were run for 6 days. While the residence time and vessel sizing was scaled appropriately, due to water volume restraints, the surface area of vessel media to flow is an underestimate of the full scale system. Regardless, the design allowed for a comparison of the performance of the two resins and the surfactant coated zeolite. Based on the results, the SIR-150 resin (Table 1 Tests B,S-19 to B,S-24) outperformed the PWA-10 resin (Table 1 Tests B,S-1 to B,S-6 and B,S-13 to B,S-18) with lower magnitudes of breakthrough. The surfactant coated zeolite (Table 1 Tests B,S-7 to B,S-12) was shown to be ineffective at treatment under the design residence times.

2.4 Final Design

Final temporal testing was performed with the entire treatment train operating for 7 days (pH adjustment to 6 using HCl, FeCl₃ addition at 40 mg/L, settling, sand filtration, and SIR-150 resin filtration). Based on additional literature research, aeration of the groundwater prior to FeCl₃ addition was added as a remedial step to assist in the formation of flocs. Results of these additional tests are presented in **Table 1 Test B-5**. The final design proposed in this memo was implemented for RIEC pilot study with results shown in **Table 1 Tests P-2**, **P-5**, **P-8**, **P-11**, **P-13**, **P-15**, **P-17**, and **P-20**. Key observations and conclusions from the treatability testing, pilot study, and additional FeCl₃ testing are:

- Aeration of the groundwater prior to the addition of FeCl₃ accelerates the formation of precipitants.
- The initial adjustment pH should be close to pH of 6 at RIEC for optimal arsenic and molybdenum removal and precipitant settling times.
- Higher FeCl₃ concentrations provided larger precipitant particles that settle faster.
- The sand filter was effective as a polishing step to remove unsettled flocs and reduce total arsenic and molybdenum concentrations to below action levels, while also decreasing total iron concentrations.
- The resin filter is needed to remove boron from the groundwater to action levels. The SIR-150 resin performed best of the resins tested. The SIR-150 resin operates optimally between a pH of 4 and 10.
- While removing boron from the groundwater, the resin also acts as a polishing tool for removal of residual arsenic, molybdenum, and additional reduction of sulfate.

The final proposed treatment design is shown in **Figure 1** below:



Figure 1: Proposed Treatment Process

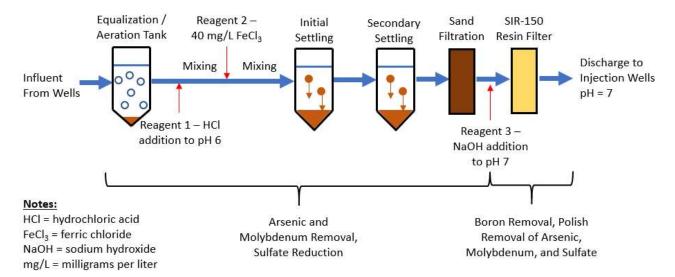


Table 1 Summary of Rush Island Energy Center Treatability Testing

Rush Island Energy Center, Missouri

		Arsenic	Molybdenum	Sulfate	Boron			
			ug/					
Test	Permit Limits	10	100	250,000	2,000			
Condition	Baseline	220	160	230,000	3,850			
Reference	Average Pilot Influent	160	981	254,727	8,550			
	Focus on Arsenic and Molybdenum Remova							
A,M-1	Adjust water to pH of 5 - Day 7	198	142	NM	3,270			
A,M-2 A,M-3	Zero Valent Iron Column - 5 parts sand, 1 Part ZVI - Day 7 Zero Valent Iron Column - 2 parts sand, 1 Part ZVI - Day 7	113 89	151 143	NM	3,900			
	Adjust water to pH 6, FeCl ₃ (20 mg/L) - 1 hour reaction	7.2	20.5	NM NM	3,510 NM			
A,M-4	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	10.9	24.1	NM	NM			
A,M-5	Adjust water to pH 6, FeCl ₃ (20 mg/L) - 3 hour reaction	12.6	28.3	NM	NM			
A,M-6	Adjust water to pH 6, FeCl ₃ (20 mg/L) - 6 hour reaction							
A,M-7	Adjust water to pH 6, FeCl ₃ (40 mg/L) - 1 hour reaction	4.9	<10	NM	NM			
A,M-8	Adjust water to pH 6, FeCl ₃ (40 mg/L) - 3 hour reaction	5.9	<10	NM	NM			
A,M-9	Adjust water to pH 6, FeCl ₃ (40 mg/L) - 6 hour reaction	6.2	<10	NM	NM			
A,M-10	Adjust water to pH 4, FeCl ₃ (40 mg/L) - 1 hour reaction	5.6	<10	NM	NM			
A,M-11	Adjust water to pH 4, FeCl ₃ (40 mg/L) - 3 hour reaction	8.5	<10	NM	NM			
A,M-12	Adjust water to pH 4, FeCl ₃ (40 mg/L) - 6 hour reaction	10.5	<10	NM	NM			
A,M-13	Adjust water to pH 6, FeCl $_3$ (40 mg/L) and 3% H $_2$ O $_2$ - 1 hour reaction	26.9	188	NM	NM			
	·		Focu	s on Boron Re	moval - RIEC			
B-1	Adjust water to pH of 5, SIR 150 Resin Column - Day 7	50.8	<20	NM	<10			
B-2	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin	<5	<10	600	<10			
B-3	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, IRA-743 Boron Resin	<5	<10	17,000	<10			
B-4	Adjust water to pH 6, FeCl₃ (40 mg/L), Sand Filter, PWA-10 Resin	<5	<10	4,400	<10			
B-5	Adjusting water to pH of 6, FeCl ₃ (40 mg/L), Sand Filter, SIR 150 Resin Column - System Operated 7 Days	1.6	<10	NM	<20			
6-0	Operated 7 Days		Focu	s on Sulfate R	emoval - SEC			
S-1	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Zeolite Filter	NM	180	380,000	14,000			
S-2	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Surfactant Coated Zeolite Filter	NM	160	360,000	15,000			
S-3	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, NaCl treated Zeolite Filter	NM	200	370,000	14,000			
S-4		NM	200	380,000	14,000			
	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Limestone Filter	NM	220	390,000	14,000			
S-5	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, adjust to pH 10, Zeolite Filter	INIVI	220	330,000	14,000			
S-6	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, adjust to pH 10, Surfactant Coated Zeolite Filter	NM	220	390,000	15,000			
S-7	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, adjust to pH 10, NaCl treated Zeolite Filter	NM	200	360,000	13,000			
S-8	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, adjust to pH 10, Limestone Filter	NM	220	390,000	14,000			
S-9	Adjust water to pir 5, recig (40 mg/z), saint riter, adjust to pir 15, Emestone riter Adjust water to pH6, NaCl Treated Zeolite Filter	NM	NM	430,000	1,500			
S-10	NaCl Treated Zeolite Filter Only	NM	NM	420,000	1,600			
S-11	Adjust water to pH 6, Surfactant Coated Zeolite Filter	NM	NM	270,000	540			
S-12	Surfactant Coated Zeolite Filter Only	NM	NM	230,000	370			
S-13	Adjust water to pH 6, FeCl ₃ (80 mg/L - Bentonite to help settling), Sand Filter, 1:1 cement:sulfate molar ration (20 min mixing)	NM	NM	460,000	NM			
	Adjust water to pH 6, FeCl ₃ (80 mg/L - Bentonite to help settling), Sand Filter, 1.5:1	NIN 4	NIA 4	470.000	NIA 4			
S-14	cement:sulfate molar ration (20 min mixing)	NM	NM	470,000	NM			
	Adjust water to pH 6, FeCl ₃ (80 mg/L - Bentonite to help settling), Sand Filter, 2:1	NM	NM	490,000	NM			
S-15	cement:sulfate molar ration (20 min mixing)							
S-16	Adjust water to pH 6, FeCl ₃ (80 mg/L - Bentonite to help settling), Sand Filter, 1:1 Ca(OH) ₂ :sulfate molar ration (60 min mixing)	NM	NM	430,000	NM			
3 10	Adjust water to pH 6, FeCl ₃ (80 mg/L - Bentonite to help settling), Sand Filter, 3:1							
S-17	Ca(OH) ₂ :sulfate molar ration (60 min mixing)	NM	NM	420,000	NM			
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	Adjust water to pH 6, FeCl ₃ (80 mg/L - Bentonite to help settling), Sand Filter, 5:1	NM	NM	420,000	NM			

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Rush Island Energy Center, Missouri

		Arsenic	Molybdenum	Sulfate	Boron				
			ug/						
Test	Permit Limits	10	100	250,000	2,000				
Condition	Baseline	220	160	230,000	3,850				
Reference	Average Pilot Influent	160	981	254,727	8,550				
C 10	Focus on Sulfate Removal - RIE								
S-19 S-20	DS-200 Zeolite Only OC-300 Zeolite Only	160 150	1,000 1,100	300,000 360,000	8,300 9,100				
S-21	HS-200 Zeolite Only	160	890	330,000	8,400				
S-22	Clinoptilolite Zeolite Only	170	1,100	230,000	8,400				
S-23	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Chitosan High MW (50 mg/L), adjust pH		NM	230,000	8,300				
	to 3.5, Mix for 1 hour	NM							
S-24	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Chitosan Medium MW (50 mg/L), adjust	NM	NM	230,000	9,100				
	pH to 3.5, Mix for 1 hour	INIVI							
	Adjust water to pH 6, $FeCl_3$ (40 mg/L), Sand Filter, Chitosan Low MW (50 mg/L), adjust pH to	NM	NM	230,000	9,400				
S-25	3.5, Mix for 1 hour	IVIVI	IVIVI	230,000	3,400				
	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Sodium Aluminate at 2:1 molar ration	NM	NM	230,000	8,100				
S-26	with sulfate, adjust pH to 11.3, Mix for 1 hour				0,100				
	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Calcium Aluminate at 2:1 molar ration	NM	NM	230,000	8,200				
S-27	with sulfate, adjust pH to 11.3, Mix for 1 hour				-,				
	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Calcium Aluminate Cement at 2:1 molar	NIVI	NM	230,000	8,100				
S-28	ration with sulfate, adjust pH to 11.3, Mix for 1 hour		on Culfata and D	lavan Dasin D	amayal DIFC				
D.C. 1	Adjust water to all C. FoCl. (40 mov/L). Cond. Filter, DMA 10 Boxon Bosin, BIFC. 1 Box	<5	on Sulfate and B		<10				
B,S-1	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - RIEC - 1 Day			210,000					
B,S-2	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - RIEC - 2 Day	<5	20	210,000	<10				
B,S-3	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - RIEC - 3 Day	<5	<10	220,000	180				
B,S-4	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - RIEC - 4 Day	<5	<10	220,000	3,400				
B,S-5	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - RIEC - 5 Day	<5	<10	220,000	7,300				
B,S-6	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - RIEC - 6 Day	<5	<10	220,000	9,100				
B,S-7	Adjust water to pH 6, FeCl₃ (40 mg/L), Sand Filter, Surfactant Coated Zeolite - RIEC - 1 Day	<5	200	220,000	8,600				
B,S-8	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Surfactant Coated Zeolite - RIEC - 2 Day	<5	200	220,000	9,100				
B,S-9	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Surfactant Coated Zeolite - RIEC - 3 Day	<5	220	220,000	9,900				
B,S-10	Adjust water to pH 6, FeCl₃ (40 mg/L), Sand Filter, Surfactant Coated Zeolite - RIEC - 4 Day	<5	220	220,000	9,400				
B,S-11	Adjust water to pH 6, $FeCl_3$ (40 mg/L), Sand Filter, Surfactant Coated Zeolite - RIEC - 5 Day	<5	240	220,000	9,300				
B,S-12	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, Surfactant Coated Zeolite - RIEC - 6 Day	<5	240	220,000	9,500				
		Focu	s on Sulfate and	Boron Resin F	Removal - SEC				
B,S-13	Adjust water to pH 6, FeCl $_{ m 3}$ (40 mg/L), Sand Filter, PWA-10 Boron Resin - SEC - 1 Day	NM	<10	360,000	<10				
B,S-14	Adjust water to pH 6, FeCl $_3$ (40 mg/L), Sand Filter, PWA-10 Boron Resin - SEC - 2 Day	NM	<10	380,000	750				
B,S-15	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - SEC - 3 Day	NM	<10	370,000	8,600				
B,S-16	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - SEC - 4 Day	NM	<10	370,000	16,000				
B,S-17	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - SEC - 5 Day	NM	<10	380,000	19,000				
B,S-18	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, PWA-10 Boron Resin - SEC - 6 Day	NM	<10	360,000	17,000				
B,S-19	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin - SEC - 1 Day	NM	<10	330,000	<20				
B,S-20	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin - SEC - 2 Day	NM	<10	370,000	30				
B,S-21	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin - SEC - 3 Day	NM	<10	370,000	4,300				
B,S-22	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin - SEC - 4 Day	NM	<10	380,000	13,000				
B,S-23	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin - SEC - 5 Day		<10	360,000	17,000				
B,S-24	Adjust water to pH 6, FeCl ₃ (40 mg/L), Sand Filter, SIR-150 Boron Resin - SEC - 6 Day		<10	360,000	18,000				
0,3-24	Aujust water to pri o, reciz (40 mg/L), Sand Filter, Sin-150 boron nesin - SEC - 6 Day	14141	110	300,000	10,000				

Table 1 Summary of Rush Island Energy Center Treatability Testing

Rush Island Energy Center, Missouri

		Arsenic	Molybdenum	Sulfate	Boron
		ug/L			
Test	Permit Limits	10	100	250,000	2,000
Condition	Baseline	220	160	230,000	3,850
Reference	Average Pilot Influent	160	981	254,727	8,550
			1	Pilot Study	Results - RIEC
P-1	2/10/21 RIEC Pilot Influent	169	1,070	263,000	7,510
P-2	2/10/21 RIEC Pilot Post Resin Filter	6	14	8,000	<250
P-3	2/12/21 RIEC Pilot Influent	145	852	261,000	8,110
P-4	2/12/21 RIEC Pilot Post Sand Filter	65	349	253,000	2,230
P-5	2/12/21 RIEC Pilot Post Resin Filter	0.8	0.7	7,000	20
P-6	2/15/21 RIEC Pilot Influent	167	871	280,000	1,620
P-7	2/15/21 RIEC Pilot Post Sand Filter	16	93	228,000	7,530
P-8	2/15/21 RIEC Pilot Post Resin Filter	1	<5	63,000	<10
P-9	2/25/21 RIEC Pilot Influent	163	880	265,000	7,940
P-10	2/25/21 RIEC Pilot Post Sand Filter	33	134	237,000	NM
P-11	2/25/21 RIEC Pilot Post Resin Filter	2	10	208,000	<10
P-12	3/3/21 RIEC Pilot Influent	166	1,030	255,000	8,550
P-13	3/3/21 RIEC Pilot Post Resin Filter	8	6	212,000	<10
P-14	4/9/21 RIEC Pilot Influent	188	1,060	278,000	9,940
P-15	4/9/21 RIEC Pilot Post Resin Filter	8	33	128,000	62
P-16	5/7/21 RIEC Pilot Influent	167	946	228,000	8,710
P-17	5/7/21 RIEC Pilot Post Resin Filter	47	248	100,000	2,330
P-18	5/11/21 RIEC Pilot Influent	180	1,020	235,000	9,480
P-19	5/11/21 RIEC Pilot Post Sand Filter	<10	4	186,000	6,910
P-20	5/11/21 RIEC Pilot Post Resin Filter	98	8	136,000	NM

Notes and Abreviations:

RIEC = Rush Island Energy Center SEC = Sioux Energy Center

NM = not measured

< = concentration is less than value

Red values exceed discharge permit limits

H₂O₂ = hydrogen peroxide

mg/L = milligrams per liter ug/L = micrograms per liter

Ca(OH)₂ = calcium hydroxide

ZVI = zero valent iron

NaCl = sodium chloride

FeCl₃ = ferric chloride

A = arsenic treatment approach

M = molybdenum treatment approach

B = boron treatment approach

S = sulfate treatment approach

MW = molecular weight