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ALTERNATIVE SOURCE DEMONSTRATION REPORT

2024 FIRST SEMIANNUAL EVENT TEXAS STATE CCR RULE

Welsh Power Plant Bottom Ash Storage Pond Registration No. CCR 110 Pittsburg, Texas

Prepared for

American Electric Power 1 Riverside Plaza Columbus, Ohio 43215-2372

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Project CHA8495

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- Attachment B Historical Potentiometric Maps
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LIST OF ACRONYMS AND ABBREVIATIONS

AEP	American Electric Power
ASD	alternative source demonstration
BASP	Bottom Ash Storage Pond
CCR	coal combustion residuals
EPRI	Electric Power Research Institute
HDPE	high-density polyethylene
LPL	lower prediction limit
meq/kg	milliequivalents per kilogram
mg/L	milligrams per liter
PBAP	Primary Bottom Ash Pond
SSI	statistically significant increase
SU	standard units
TAC	Texas Administrative Code
TCEQ	Texas Commission on Environmental Quality
UPL	upper prediction limit



1. INTRODUCTION AND SUMMARY

This alternative source demonstration (ASD) report has been prepared to address statistically significant increases (SSIs) for boron, calcium, and chloride in the groundwater monitoring network at the Bottom Ash Storage Pond (BASP) located at the Welsh Power Plant (Welsh Plant) in Pittsburg, Texas, following the first semiannual detection monitoring event of 2024. The Welsh Plant has three coal combustion residuals (CCR) storage units regulated by the Texas Commission on Environmental Quality (TCEQ) under Registration No. CCR 110, including the BASP (**Figure 1**). The BASP was being closed by CCR removal at the time of the detection monitoring event, and the other two CCR units were still active.

Background groundwater values for the BASP were originally calculated in January 2018 and have been updated periodically in accordance with the *Statistical Analysis Plan* prepared for the Welsh Plant (Geosyntec 2021). Under this plan, prediction limits were calculated for each well using intrawell comparisons. Applicable background values for the first semiannual event of 2024 are the revised upper prediction limits (UPLs) calculated in January 2024 for each Appendix III parameter (Geosyntec 2024). Revised lower prediction limits (LPLs) were also calculated for pH. Prediction limits were calculated based on a one-of-two retesting procedure to maintain an appropriate site-wide false positive rate. With this procedure, an SSI is concluded only if both samples in a series of two have reported results above the UPL or, in the case of pH, are below the LPL. In practice, if the initial result was not above the UPL or was not below the LPL, a second sample was not collected or analyzed.

The first semiannual detection monitoring event of 2024 was performed in April 2024 (initial sampling event), and the results were compared to the calculated prediction limits. Where initial values were identified above the UPL or below the LPL, verification resampling was completed in June 2024. Following verification resampling, intrawell comparisons identified SSIs for boron and calcium at monitoring well AD-4C and chloride at monitoring well AD-3. A summary of the detection monitoring analytical results for the downgradient compliance wells and the calculated prediction limits to which they were compared is provided in **Table 1**.

1.1 CCR Rule Requirements

TCEQ regulations regarding detection monitoring programs for CCR landfills and surface impoundments provide owners and operators with the option to make an ASD when an SSI is identified (Title 30 §352.941(c)(2) of the Texas Administrative Code (TAC) [30 TAC §352.941(c)(2)]):

In making a demonstration under this section, the owner or operator must . . . within 90 days of making a determination of an SSI over the background value for any Appendix III constituent adopted by reference in §352.1421 of this title, submit a report prepared and certified in accordance with §352.4 of this title (relating to Engineering and Geoscientific Information), to the executive director, and any local pollution agency with jurisdiction that has requested to be notified, demonstrating that a source other than a coal combustion residuals unit caused the SSI or that the SSI resulted from error in sampling, analysis, statistical evaluation, or natural variation in groundwater quality.



Pursuant to this regulation, Geosyntec Consultants, Inc. (Geosyntec) has prepared this ASD report on behalf of American Electric Power (AEP) to document that the SSIs identified for boron and calcium at well AD-4C and chloride at well AD-3 are from sources other than a release from the BASP at the Welsh Plant.

1.2 Demonstration of Alternative Sources

An evaluation was completed to assess possible alternative sources to which each identified SSI could be attributed. Alternative sources were categorized into the following five types, based on methods provided by the Electric Power Research Institute (EPRI 2017):

- ASD Type I: Sampling Causes
- ASD Type II: Laboratory Causes
- ASD Type III: Statistical Evaluation Causes
- ASD Type IV: Natural Variation
- ASD Type V: Anthropogenic Sources

A demonstration was conducted to show that the identified SSIs at AD-3 and AD-4C were based on Type IV (natural variation) causes and not by a release from the BASP.



2. SUMMARY OF SITE CONDITIONS

The site background summary included in this section was primarily taken from Arcadis (2022), unless otherwise noted.

2.1 BASP Location and Design

The BASP was a 22-acre CCR surface impoundment located in the southern portion of the Welsh Plant, immediately south of the Landfill and Primary Bottom Ash Pond (PBAP) (**Figure 1**). It was designed with approximately 20-foot-high compacted-clay perimeter embankments and a 60-mil-thick high-density polyethylene (HDPE) liner placed over the base of the pond and the interior embankment slopes. The BASP was constructed and placed into operation in 2000 to receive bottom ash and economizer ash dredged and sluiced from the PBAP.

A Closure Plan for the BASP was developed in October 2016 and revised in February 2021 (AEP 2021a). This document details the closure activities which are to take place throughout the closure of the BASP. AEP submitted a certified notification that as of April 6, 2021, the BASP ceased receipt of CCR and non-CCR waste streams and closure activities had been initiated in accordance with the certified Closure Plan (AEP 2021b). Thus, the BASP no longer received CCR material or transport waters and no longer received non-CCR wastewaters such as stormwater runoff from the landfill and surrounding areas. In November 2021, removal of the CCR material from the BASP began with the CCR material stockpiled in the northern portion of the BASP (AEP 2022a). Dewatering activities began in early 2022 and included installation of dewatering pumps and trenches. As a result of the closure activities, the BASP no longer contained impounded water as of November 3, 2022 (AEP 2022b). The removal of all CCR materials from the BASP as part of closure activities was completed in September 2024.

2.2 Regional Geology and Site Hydrogeology

The Welsh Plant is located within the West Gulf Coastal Plain. The BASP is immediately underlain by the Eocene-age Recklaw Formation, which consists of very-fine- to fine-grained sand and clay (Flawn 1966). The Recklaw Formation ranges in thickness from approximately 10 to 110 feet in Titus County, where the Welsh Plant is located. This formation is underlain by the Eocene-age Carrizo Sand, consisting of fine to coarse sand, silt, and clay.

The uppermost aquifer in the vicinity of the BASP consists of an interval of the Recklaw Formation that is approximately 12-feet thick and composed of very-fine- to fine-grained silty sand and sandy silt. This aquifer is first encountered approximately 8 feet below the base of the BASP (Arcadis 2022). It is recharged primarily through infiltration of regional precipitation. Groundwater flow velocities in the uppermost aquifer in the vicinity of the BASP have been reported as approximately 1–20 feet per year (AEP 2022a).

Monitoring well AD-3 is screened from 7-17 feet below ground surface and monitoring well AD-4C is screened from 5-15 feet below ground surface. Both monitoring wells AD-3 and AD-4C are screened within the Recklaw Formation. Subsurface lithology at and near monitoring wells AD-3 and AD-4C are shown on geologic cross sections from Arcadis (2022) (Attachment A).



2.3 BASP Monitoring Well Network and Flow Conditions

The BASP monitoring well sampling network consists of background monitoring wells AD-1, AD-5, and AD-17 and downgradient compliance monitoring wells AD-3, AD-4C, and AD-16R (Figure 1). The groundwater flow direction near the BASP is generally to the southeast (Figure 2). Potentiometric groundwater flow maps from sampling events completed within the past year are provided as Attachment B. Seasonal variability in groundwater flow direction has not been observed in the immediate vicinity of the BASP.



3. ALTERNATIVE SOURCE DEMONSTRATION

The ASD evaluation method and proposed alternative source of boron and calcium at well AD-4C and chloride at well AD-3 are described below.

3.1 Proposed Alternative Sources

An initial review of groundwater sampling field forms did not identify alternative sources due to a Type I (sampling) issue. A review of the laboratory quality assurance and quality control data and the statistical analyses did not identify any Type II (laboratory) or Type III (statistical evaluation) issues. Groundwater sampling, laboratory analysis, and statistical evaluations were generally completed in accordance with 30 TAC §352.941(a) and draft TCEQ guidance for groundwater monitoring (TCEQ 2020).

As described below, the SSIs for boron, calcium, and chloride have been attributed to natural variation, which is a Type IV issue.

3.1.1 Comparison to Background Concentrations - Boron

Recent fluctuations of aqueous boron concentrations at downgradient well AD-4C are attributed to natural variability of boron in the uppermost aquifer. The maximum boron concentration observed at AD-4C of 0.217 milligrams per liter (mg/L) falls within the range of boron concentrations observed in the background wells also screened within the Recklaw Formation and located upgradient or cross-gradient of the BASP (Figure 3; Attachment A). Background monitoring well AD-1 has consistently contained boron concentrations greater than the AD-4C maximum reported value of 0.217 mg/L. Background monitoring well AD-17 has reported boron concentrations greater than or comparable to the AD-4C maximum as well. Groundwater boron concentrations at all locations discussed are lower than the Texas Risk Reduction Program (TRRP) Class I residential ingestion pathway limit (^{GW}GW_{Ing}) of 4.9 mg/L (TCEQ 2009; updated TCEQ 2023), and groundwater boron concentrations at AD-4C are more than 50 times less than the TRPP limit. Historical data indicate that the recent boron concentrations observed at AD-4C are still within the expected range associated with site background conditions. TCEQ established a Texasspecific soil background concentration of 30 milligrams per kilogram of boron in 30 TAC §350.51(m). Given the abundance of boron in Texas soils, some contribution of boron to groundwater from the aquifer is anticipated. Variable boron concentrations observed in recently collected samples are greater than historical tendencies, but concentrations in these samples fluctuate between increasing and decreasing over the previous five events and do not display a definitive increasing trend.

3.1.2 Comparison to Background Concentrations - Calcium

Calcium concentrations at background wells AD-1, AD-5, and AD-17, which are located upgradient or cross-gradient of the BASP and AD-4C, have historically been much higher than those observed at AD-4C (**Figure 4**). Since background monitoring was initiated in 2016, calcium concentrations at AD-1, which is the closest background well as well as an upgradient well, have been subject to significant variability, with a peak concentration of 147 mg/L in June 2017 and a minimum of 3.88 mg/L in October 2020. In contrast, calcium concentrations at AD-4C have ranged between 0.341 in December 2020 and 1.83 mg/L in April 2024. Calcium concentrations at background cross-gradient wells AD-5 and AD-17 have also been consistently above those

observed at AD-4C. Protective concentration levels have not been established for calcium in groundwater through the TRRP program (TCEQ 2023). Given that the concentrations of calcium at AD-4C have consistently been one to two orders of magnitude lower than those of the background wells, the recently observed higher calcium concentrations at AD-4C may represent the migration of groundwater from upgradient locations such as AD-1 within the uppermost aquifer which contain geogenic calcium at concentrations exceeding the UPL for AD-4C.

Regional scale sampling data from shallow wells located in Titus County (Attachment C; Texas Water Commission 1965) further support the existence of naturally occurring calcium concentrations exceeding the UPL of 1.44 mg/L. At the time of publication in 1965, calcium concentrations were reported for 38 samples from 27 wells within Titus County screened at shallow (60 feet or less) depths. Of these 38 samples, only 1 contained reported calcium concentrations below the UPL of 1.44 mg/L (Figure 5). This dataset contained an average calcium concentration of 46.73 mg/L, a median concentration of 13 mg/L, and a maximum of 308 mg/L. Both the average and the median values exceed the calcium UPL of 1.44 mg/L at well AD-4C. These data demonstrate the common natural calcium concentrations in regional groundwater and contextualize the calcium SSI relative to what would be expected within the surrounding region.

3.1.3 Comparison to Background Concentrations - Chloride

Chloride concentrations at downgradient well AD-3 have remained generally consistent since monitoring began in 2016 and are within the range of values observed at monitoring wells located upgradient of the BASP (**Figure 6**). The maximum chloride concentration observed at AD-3 of 10.3 mg/L falls within the range of chloride concentrations observed in background wells screened within the Recklaw Formation and located cross-gradient or upgradient of the BASP (**Figure 3**; **Attachment A**). Cross-gradient monitoring wells AD-5 and AD-17 have consistently contained chloride concentrations greater than the AD-3 maximum reported value of 10.3 mg/L. Protective concentration levels have not been established for chloride in groundwater through the TRRP program (TCEQ 2023). Chloride concentrations at AD-3, the well of concern, are comparable to and often lower than those observed at the downgradient monitoring well AD-4C. This indicates that the recent chloride concentration exists naturally in the groundwater of the uppermost aquifer at concentrations which exceed the UPL for AD-3.

Historical data from AD-3 indicate that chloride concentrations can range from 7.0 to 10.3 mg/L. Regional scale sampling data from shallow wells located in Titus County (**Attachment C**; Texas Water Commission 1965) further support the existence of naturally occurring chloride concentrations exceeding the UPL of 9.40 mg/L. At the time of publication in 1965, chloride concentrations were reported for 44 samples from 27 wells within Titus County screened at shallow (60 feet or less) depths. Of these 44 samples, only 6 contained reported chloride concentrations below the UPL of 9.40 mg/L (**Figure 7**). This dataset contained an average chloride concentration of 130.4 mg/L, a median concentration of 39.5 mg/L, and a maximum of 450 mg/L. Both the average and the median values exceed the chloride UPL of 9.40 mg/L at well AD-3. These data indicate that chloride concentrations vary within groundwater at comparable depths at the regional scale.

3.1.4 AD-3 and AD-4C Aqueous Geochemical Stability

A CCR unit release would be expected to impact the major ion chemical signature of downgradient groundwater. A Piper diagram was created to visualize major ion chemistry of AD-3 and AD-4C



groundwater (Figure 8). Piper diagrams represent the relative proportions of major cations and anions in water samples in the lower left and right triangles respectively and provide a combined view in the middle diamond which is created by projecting each triangle's axes onto a singular plot. Placement of data on the diamond therefore does not incorporate the full extent of the data plotted in individual triangles (i.e., movement along one axis in each triangle is not reflected in the diamond). The BASP sample included on Figure 8, which was collected in August 2020, is the most recently collected water sample from the unit and represents the final geochemical composition prior to the initiation of closure activities. No additional BASP samples can be collected due to closure activities.

The geochemical signature of AD-4C groundwater has displayed some variability (particularly among anion proportions) but has remained generally similar throughout the monitoring period, as illustrated by the clustering of sample results on the Piper diagram. In the event of a BASP release, AD-4C groundwater chemistry would be expected to shift to reflect the major ion signature of the BASP sample. The cation signature of AD-4C groundwater has remained consistent, and the anion signature has diverged further from the BAP sample since installation. The lack of a temporal shift towards the BASP sample suggests a lack of influence from the BASP on the groundwater chemistry. This conclusion reinforces the determination that recently observed chemical concentrations at AD-4C are associated with natural variability in groundwater composition of the uppermost aquifer.

AD-3 groundwater has also displayed some variability (particularly among anion proportions) throughout the monitoring period but has remained generally consistent. In the event of a BASP release, AD-3 groundwater chemistry would be expected to shift to reflect the major ion signature of the BASP sample. The cation signature of AD-3 has remained consistent except for one sample collected in 2019, and the anion signature does not display a clear temporal trend among the minor variability. This observation reinforces the determination that chloride concentrations at AD-3 are not associated with a BASP release, but rather due to natural variability in groundwater composition of the uppermost aquifer.



4. CONCLUSIONS AND RECOMMENDATIONS

The preceding information serves as the ASD prepared in accordance with 30 TAC 352.941(c)(2) and supports the position that the boron and calcium SSIs at AD-4C and the chloride SSI at AD-3 identified during the first semiannual detection monitoring event of 2024 should be attributed to natural variation and not to a release from the Welsh BASP. Therefore, no further action is warranted. Certification of this ASD by a qualified professional engineer is provided in **Attachment D**.

5. REFERENCES

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- AEP, 2021b. Notification of Intent to Close a CCR Unit. Pirkey Power Plant, East Bottom Ash Pond. April 6, Revised June 1.
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TABLES

Table 1. Detection Monitoring Data SummaryAlternative Source Demonstration Report - 2024 First Semiannual EventWelsh Plant, Bottom Ash Storage Pond

Analyta	Unit	Description	AI)-3	AD)- 4C	AD	-16R	
Analyte	Unit	Description	4/1/2024	6/10/2024	4/1/2024	6/10/2024	4/2/2024	6/10/2024	
Boron	ma/I	Intrawell Background Value (UPL)	0.0	407	0.0	882	0.0	577	
DOIOII	mg/L	Analytical Result	0.027		0.217	0.176	0.033		
Calcium	ma/I	Intrawell Background Value (UPL)	1.	38	1.	44	2.	90	
Calcium	mg/L	Analytical Result	0.65		1.83	1.56	0.54		
Chloride	mg/L	Intrawell Background Value (UPL)	9.	40	18	8.6	8.00		
Chionae	mg/L	Analytical Result	10.3	10.1	14.0		6.68		
Fluoride	mg/L	Intrawell Background Value (UPL)	0.2	263	0.1	180	0.2	296	
Tuonac	mg/L	Analytical Result	0.12		0.12		0.10		
		Intrawell Background Value (UPL)	5	.2	5	.7	4.6		
pН	SU	Intrawell Background Value (LPL)	3	.8	4	.0	2.8		
		Analytical Result	3.7	4.4	4.6	4.2	2.7	3.6	
Sulfate	ma/I	Intrawell Background Value (UPL)	10).6	1	23	73	5.4	
Sullate	mg/L	Analytical Result	2.4		120		66.5		
Total Dissolved Solids		Intrawell Background Value (UPL)	1.	36	3	32	242		
Total Dissolved Solids	mg/L	Analytical Result	100		320		170		

Notes:

1. Bold values exceed the background value.

2. Background values are shaded gray.

LPL: Lower prediction limit

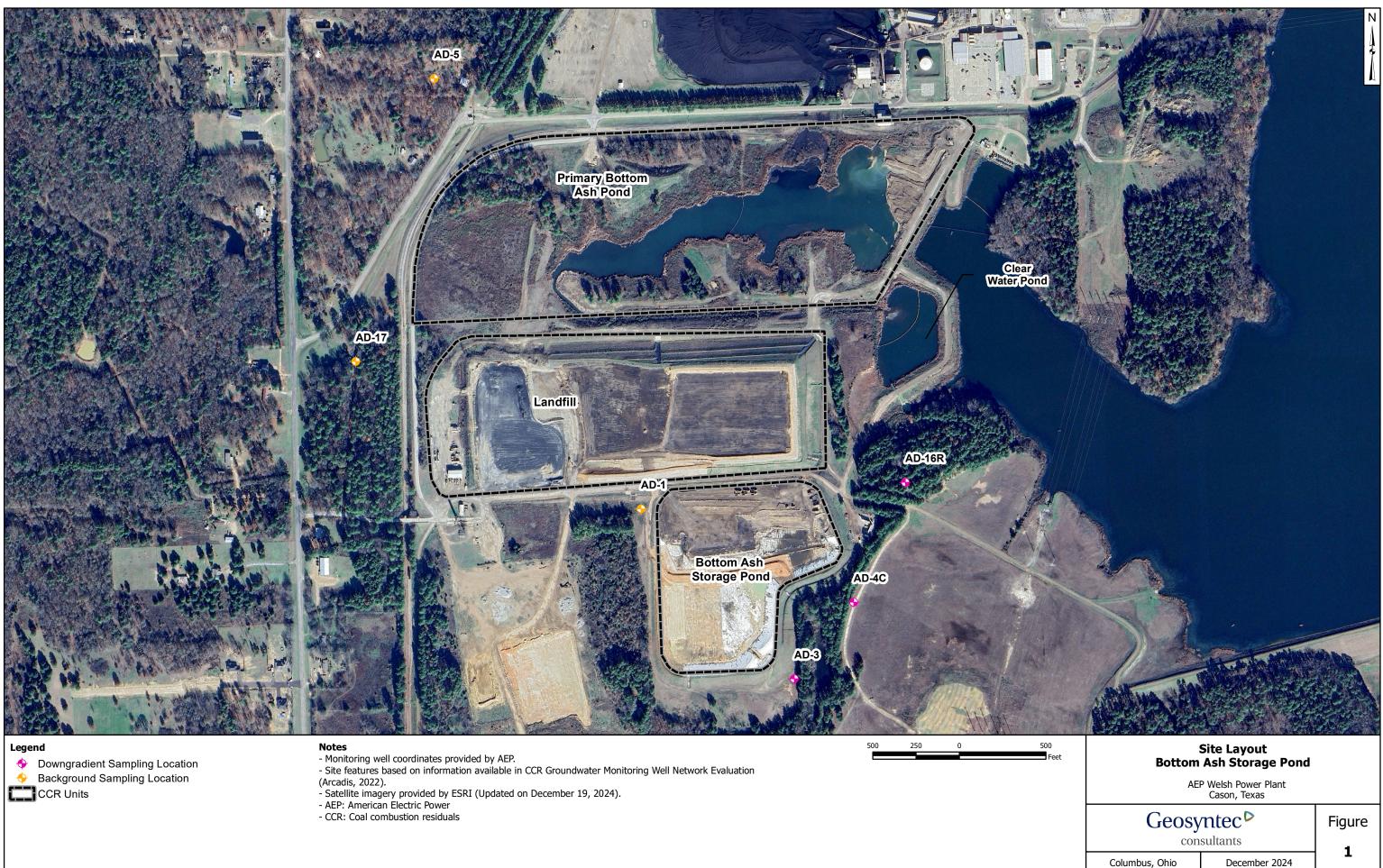
mg/L: milligrams per liter

SU: standard units

UPL: Upper prediction limit



FIGURES





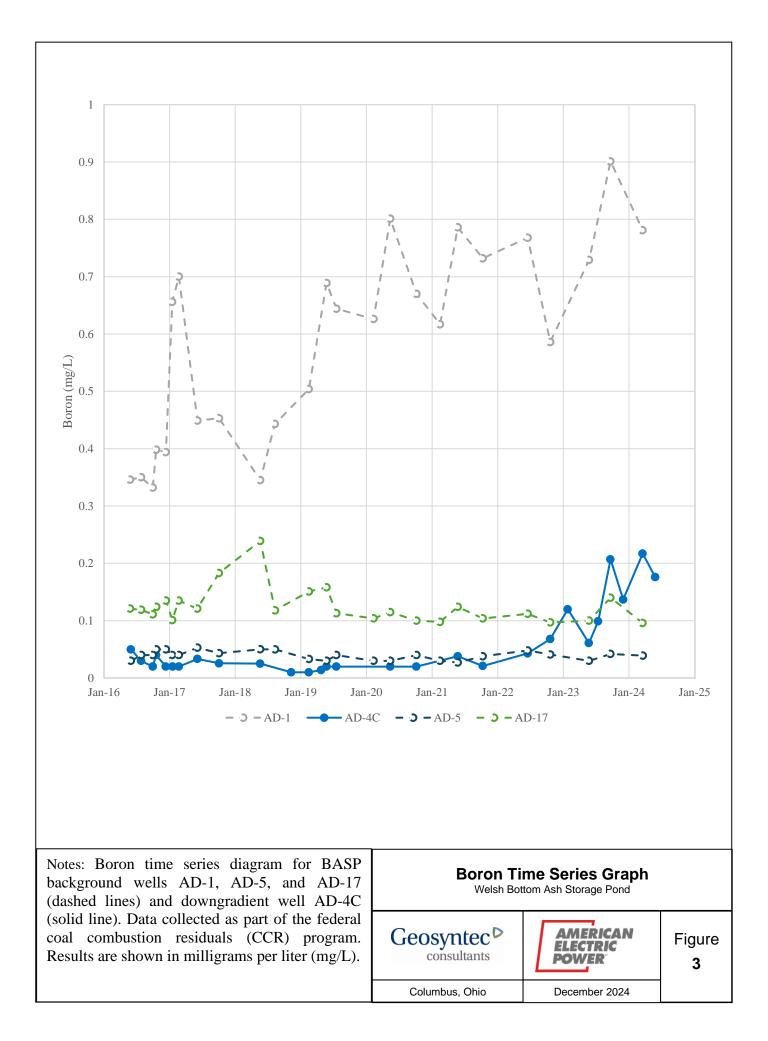
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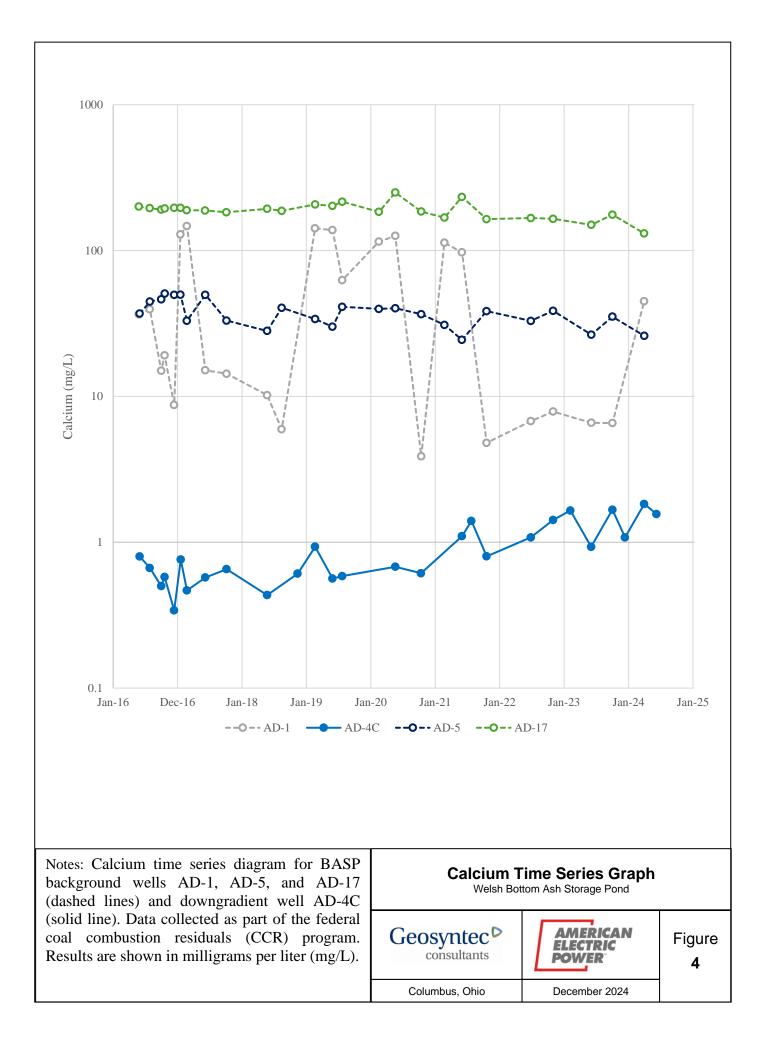


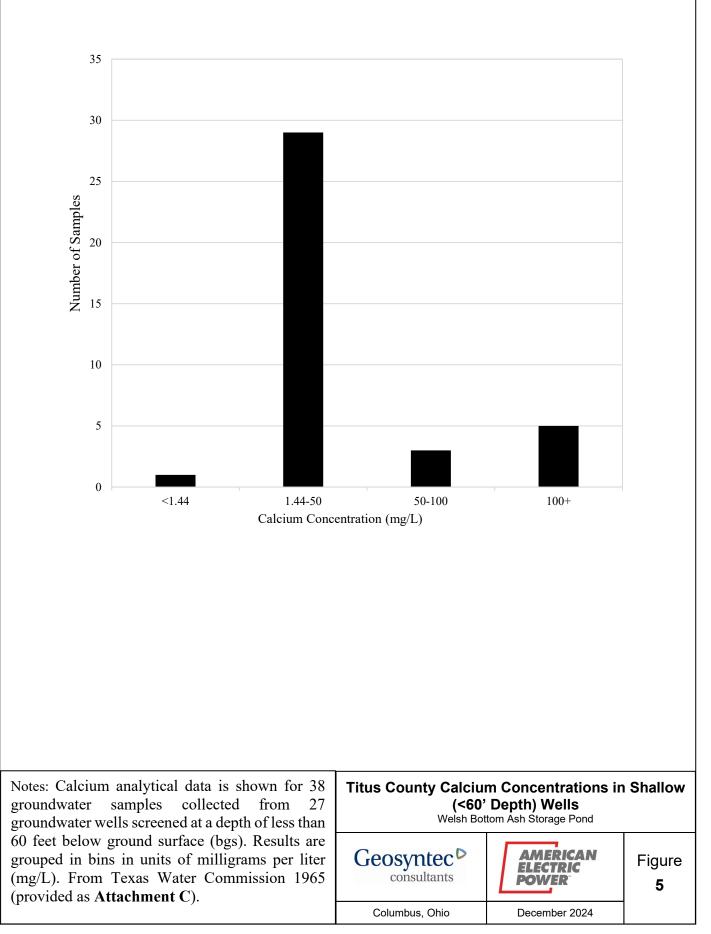
consultants

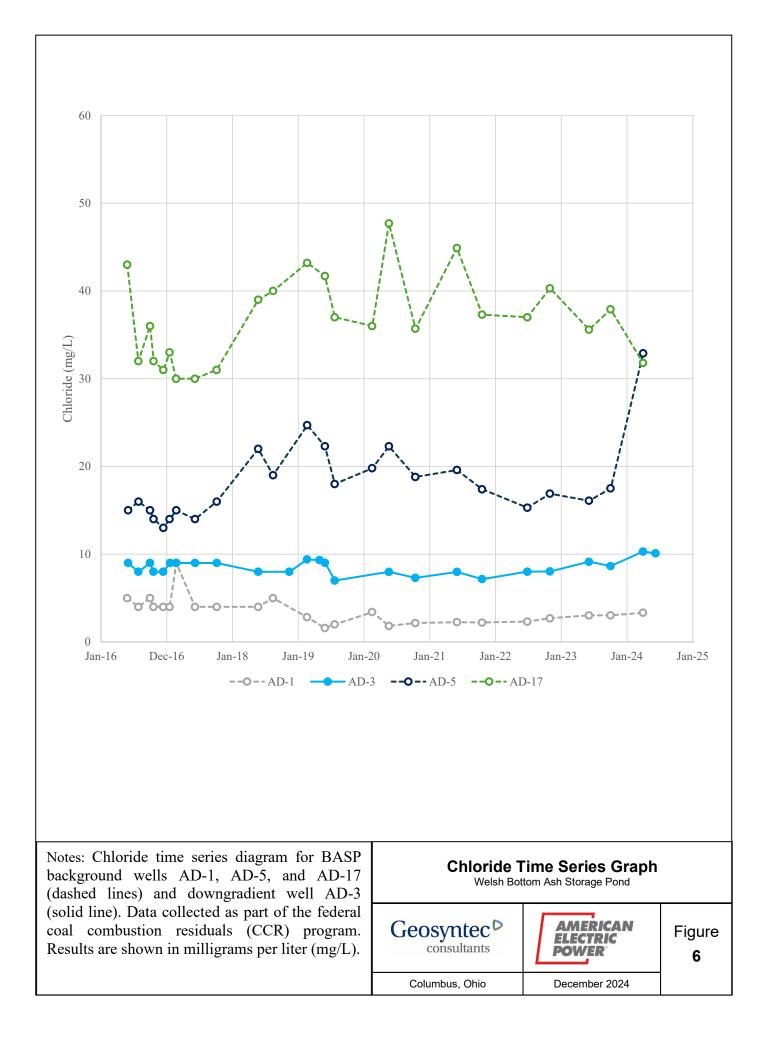
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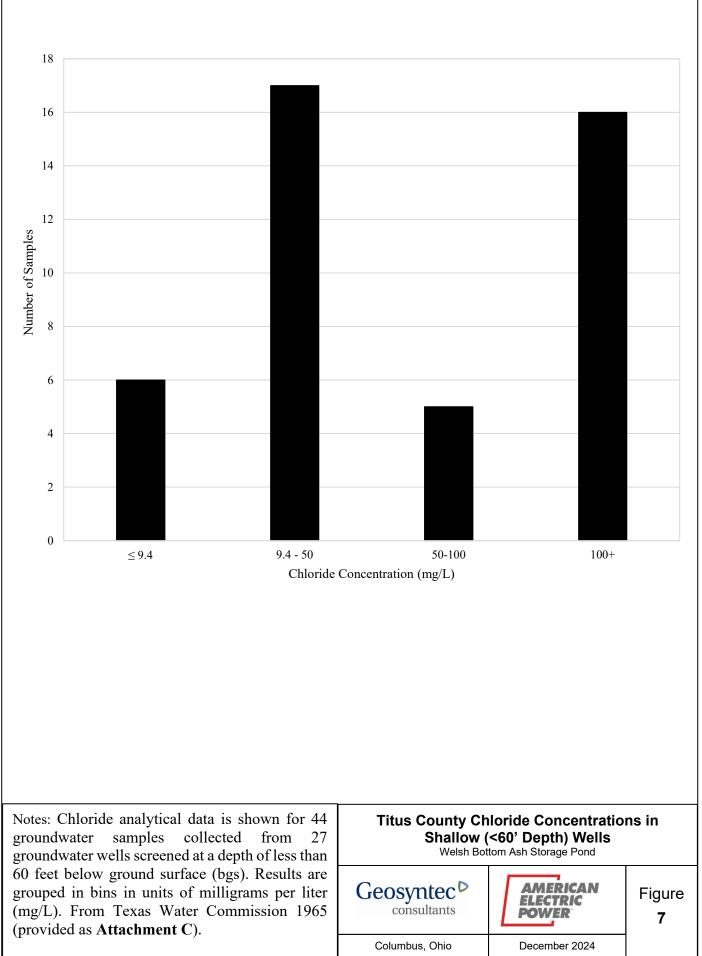
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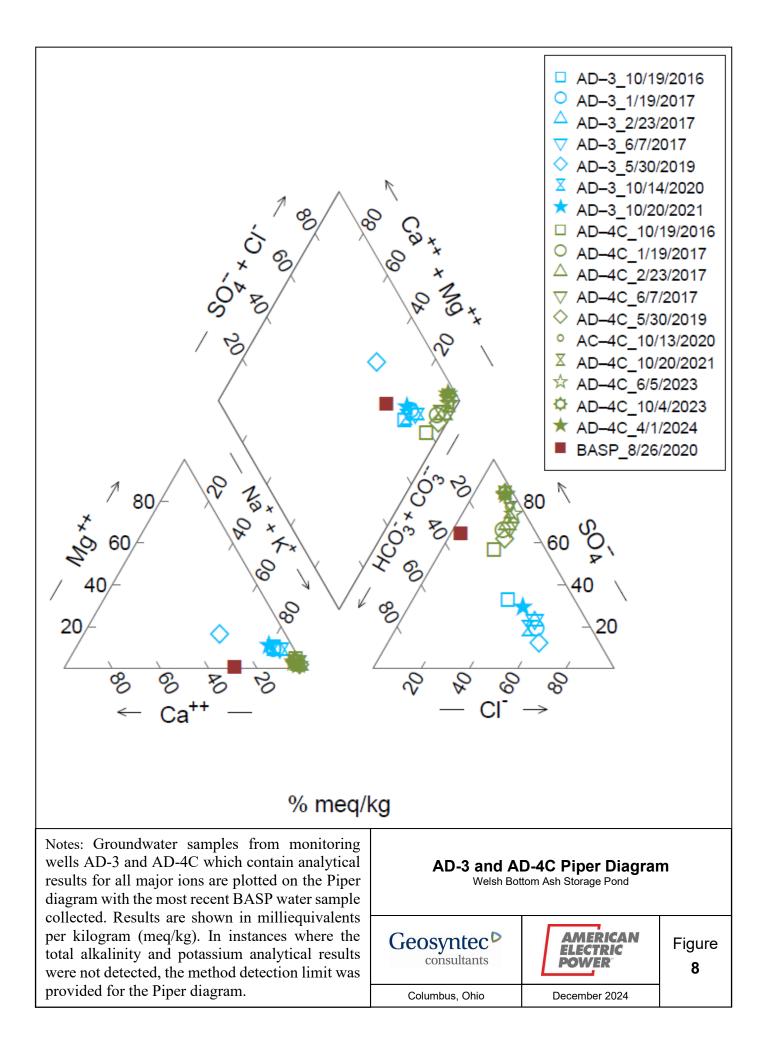




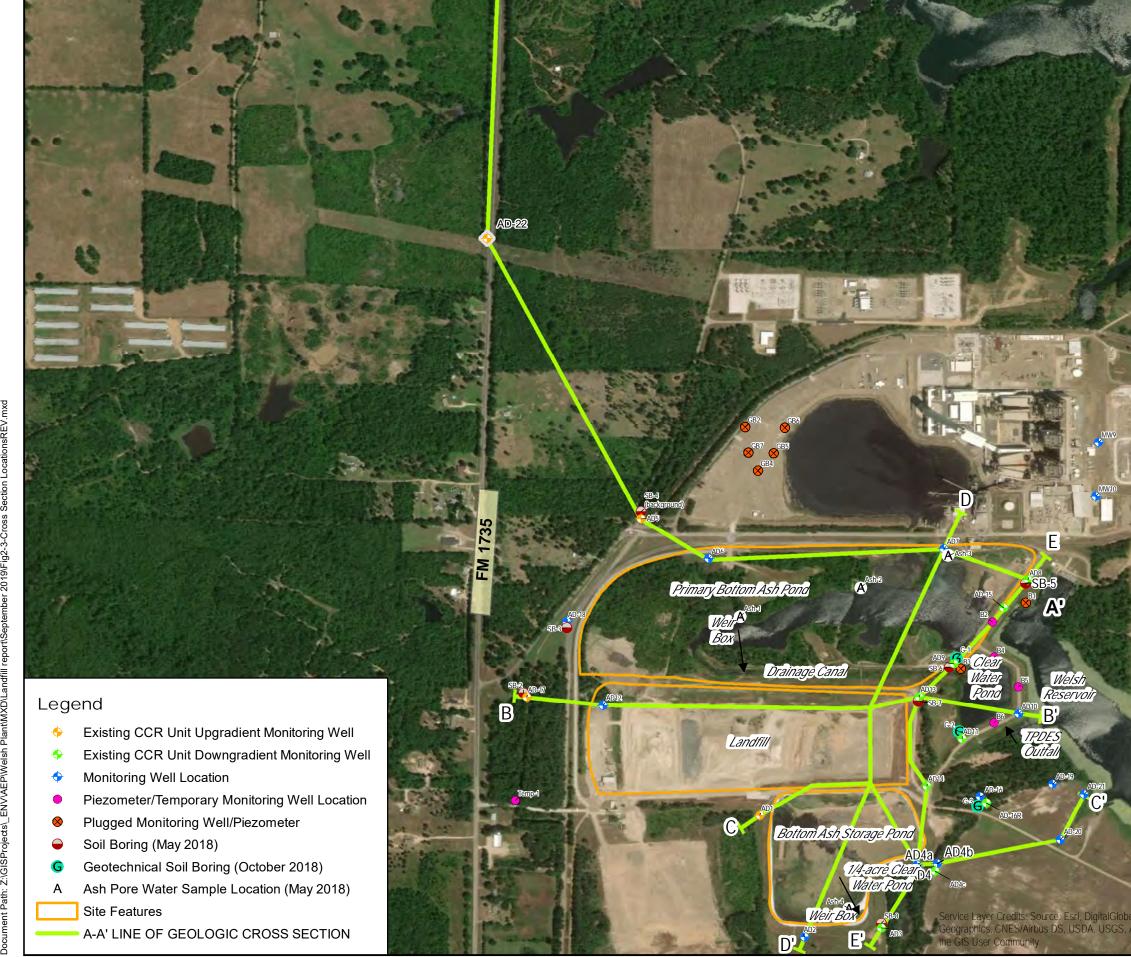






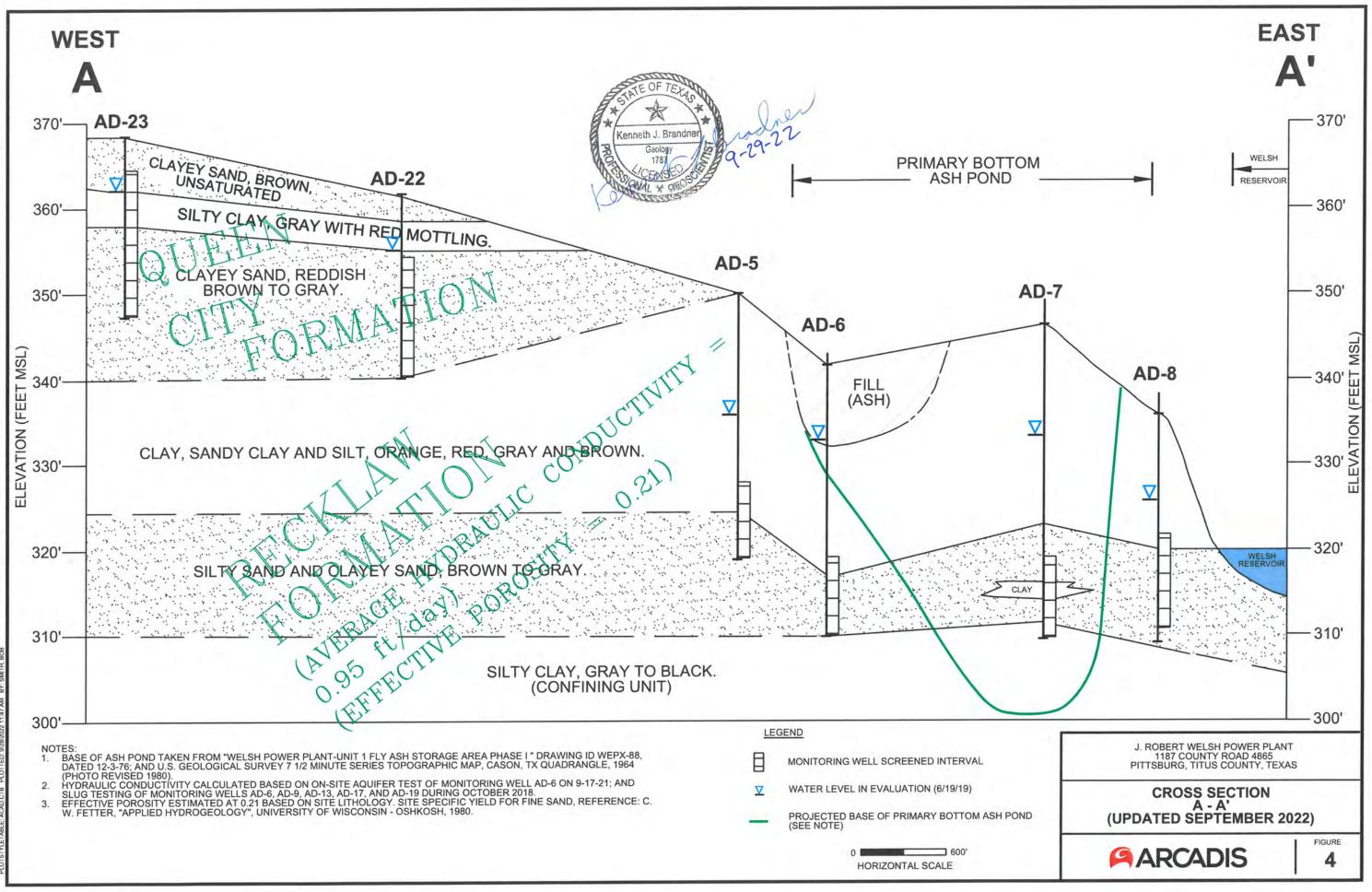


ATTACHMENT A Geologic Cross Sections

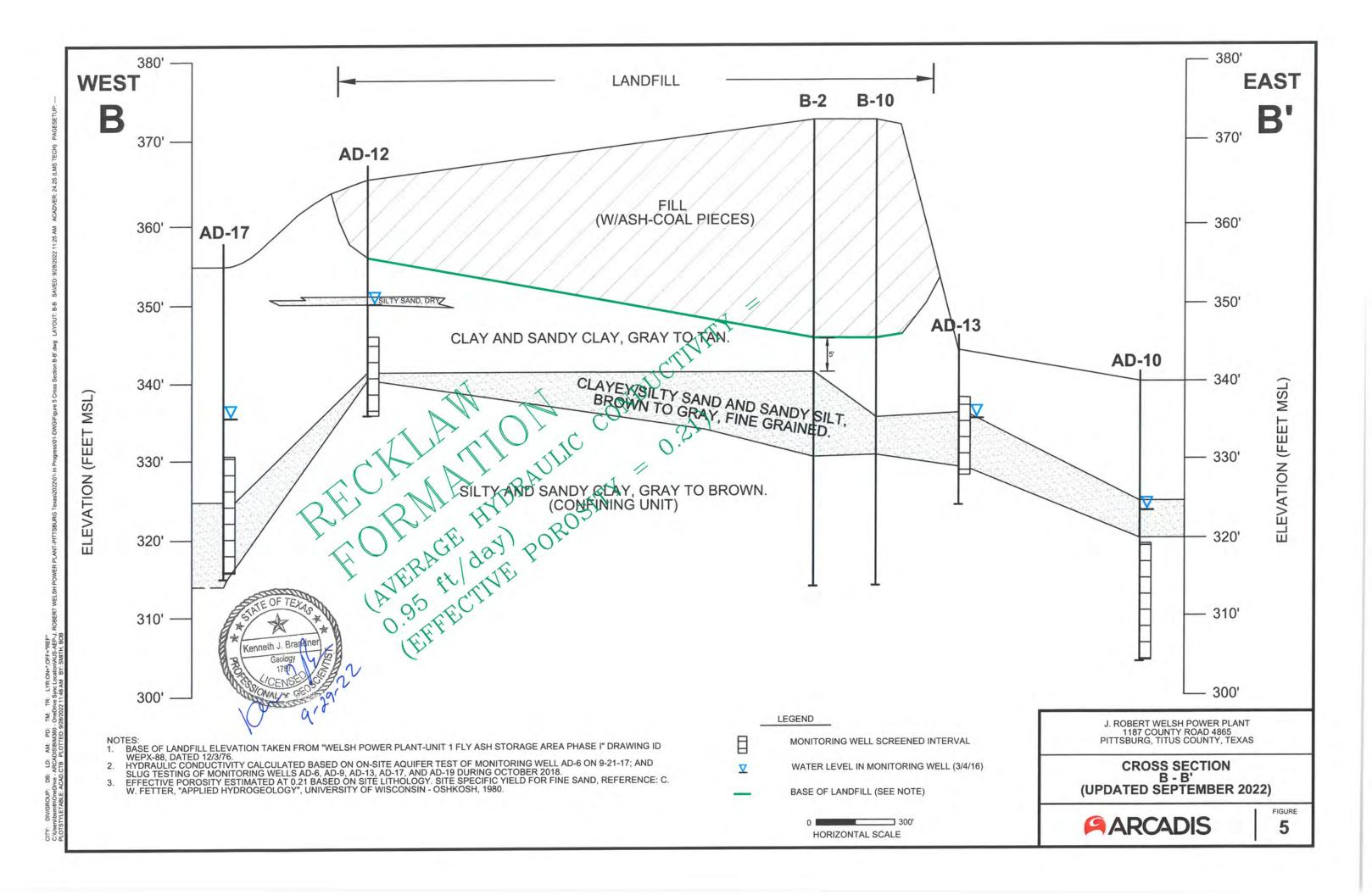


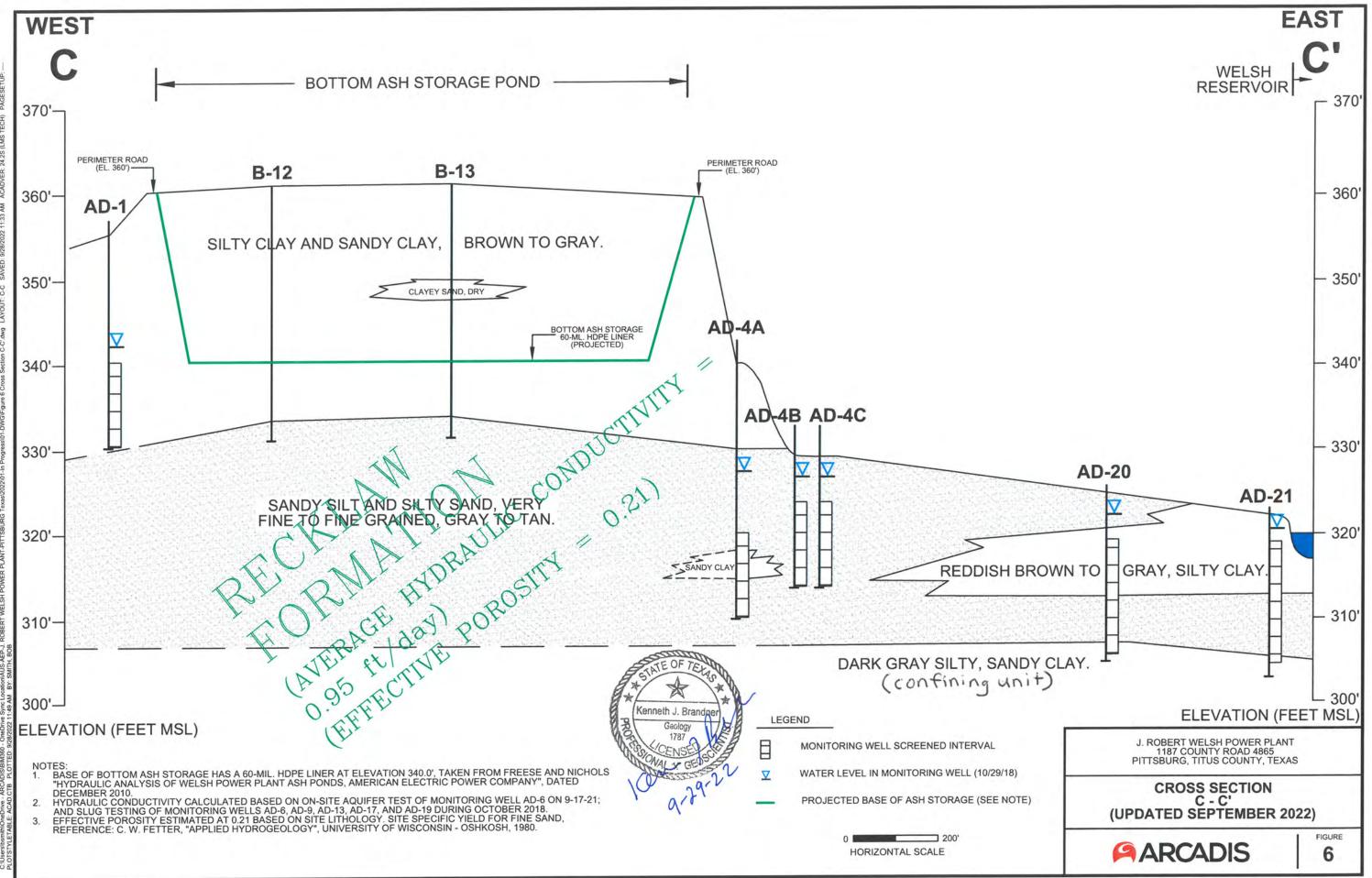
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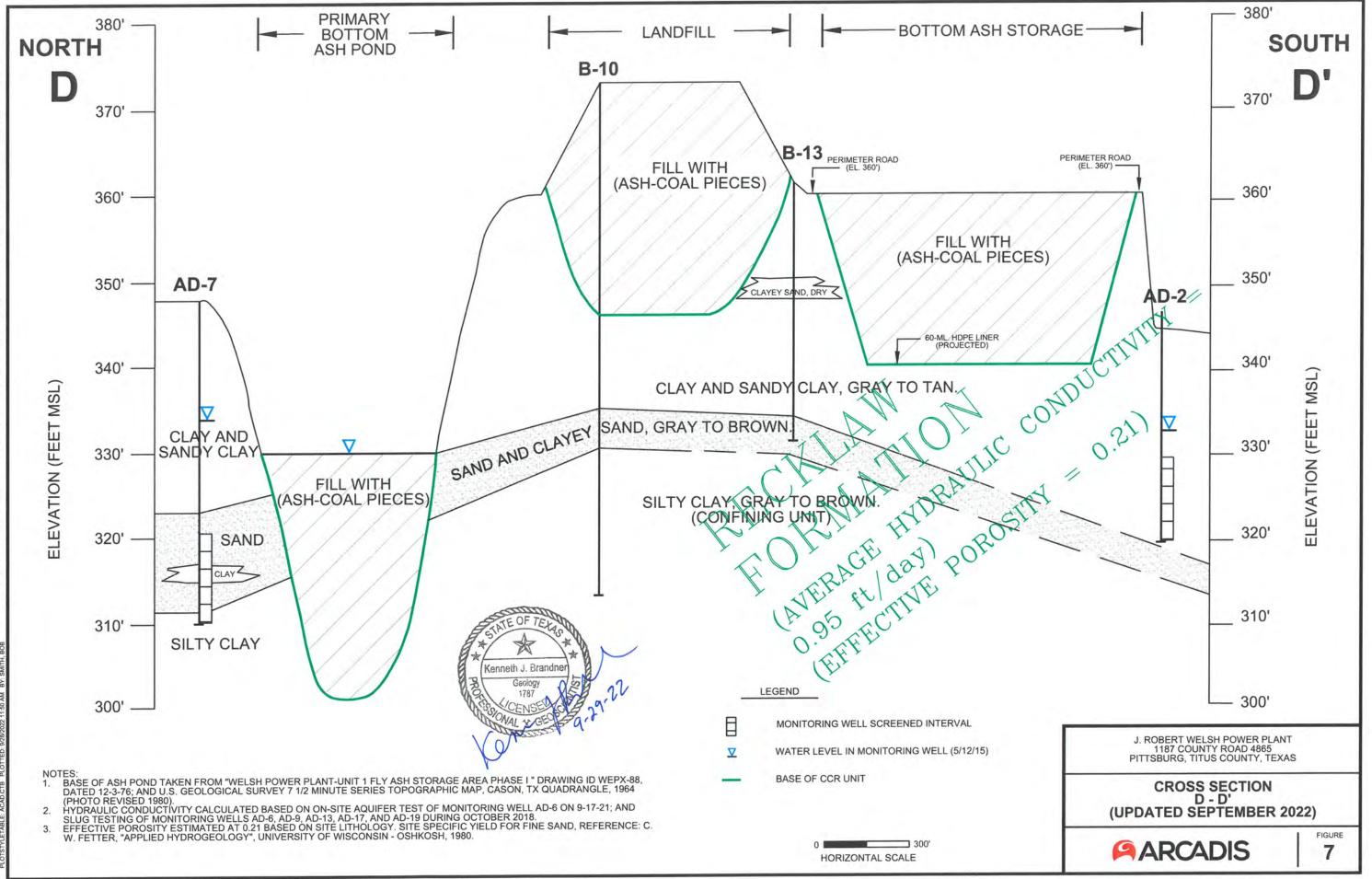




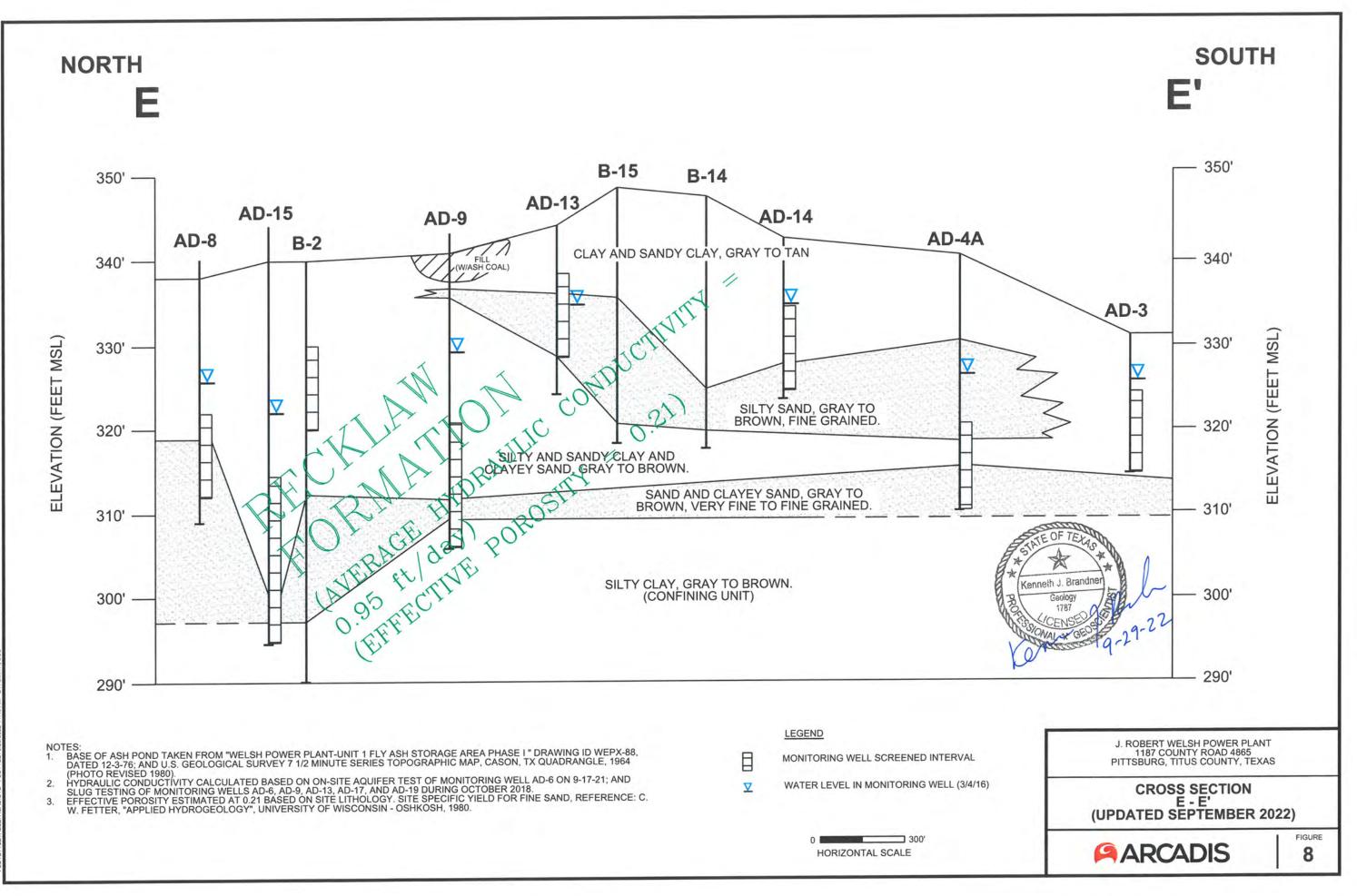
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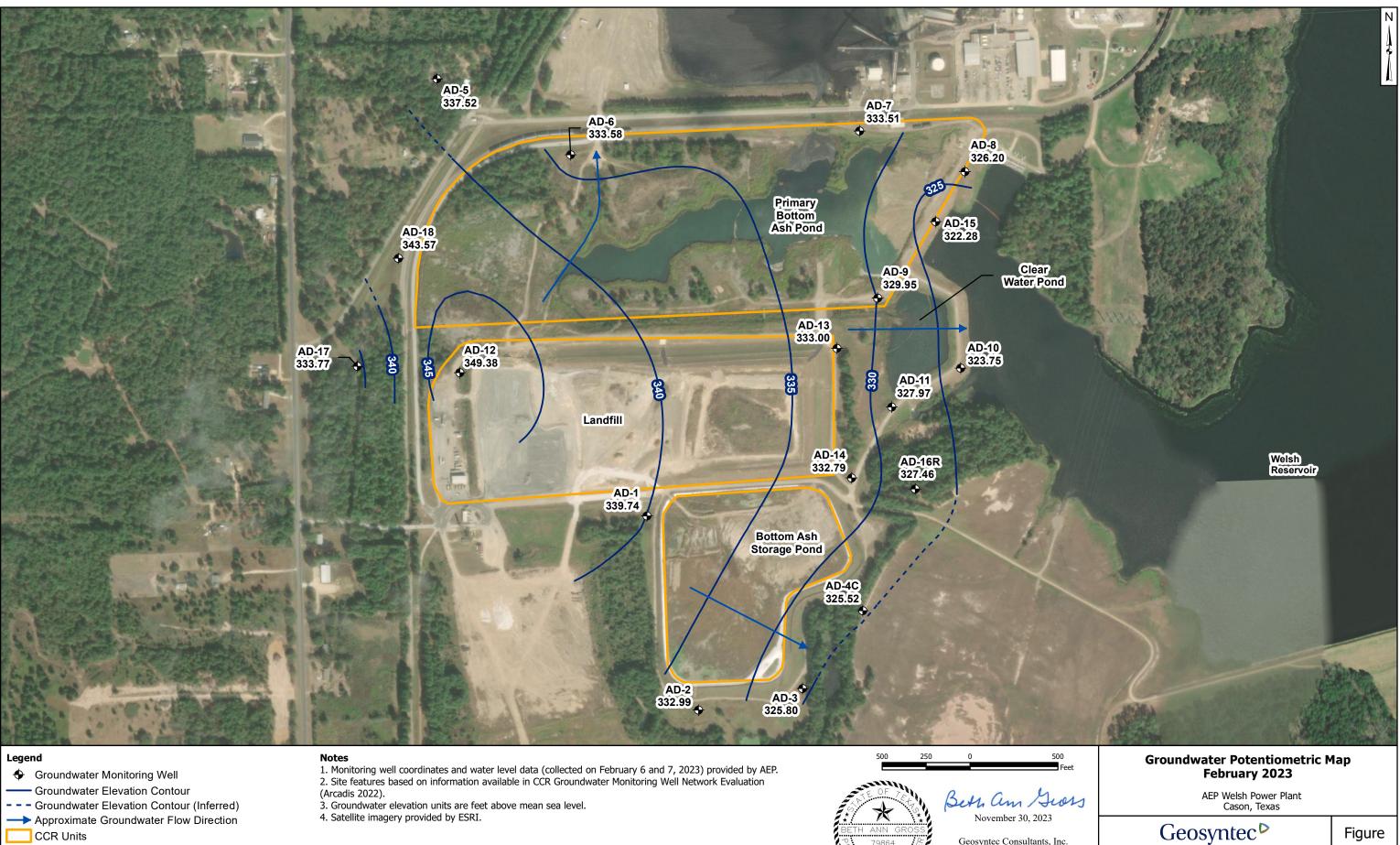


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ATTACHMENT B Historical Potentiometric Maps





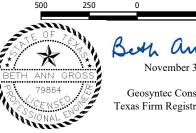
Geosyntec Consultants, Inc. Texas Firm Registration No. 1182

2023/11/30

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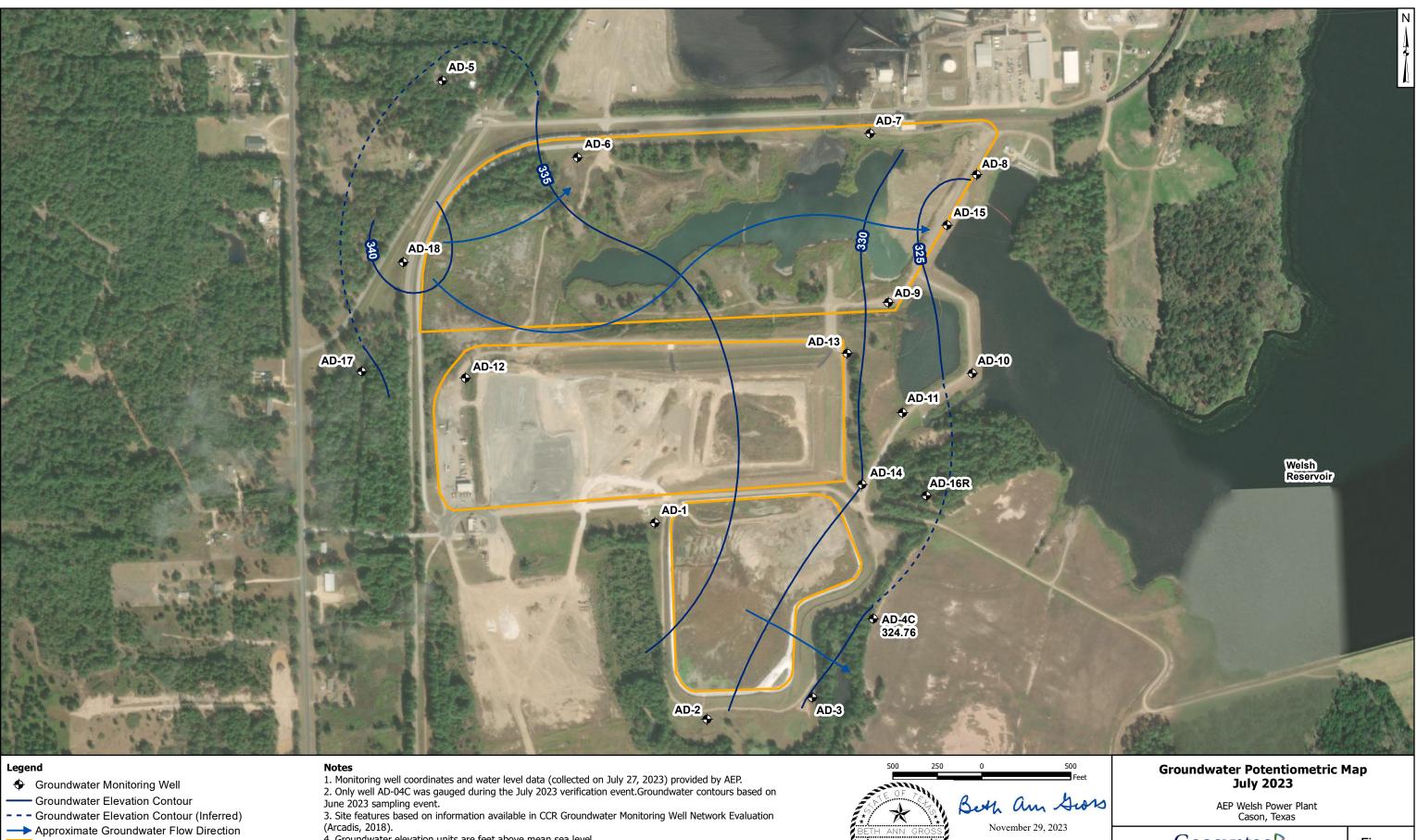
Columbus, Ohio





Columbus, Ohio

2023/11/30



- CCR Units

- Groundwater elevation units are feet above mean sea level.
 Satellite imagery provided by ESRI.



Nannarbor-01\Data\Projects\AEP\Groundwater Statistical Evaluation - CHA8423\Groundwater Mapping\GIS Files\MXD\Welsh\2023\AEP-Welsh_GW_2023-07July_ver.mxd. ASoltero. 11/29/2023. CHA8423/14/08.

Geosyntec▷ consultants

Figure 3

Columbus, Ohio 2023/11/29



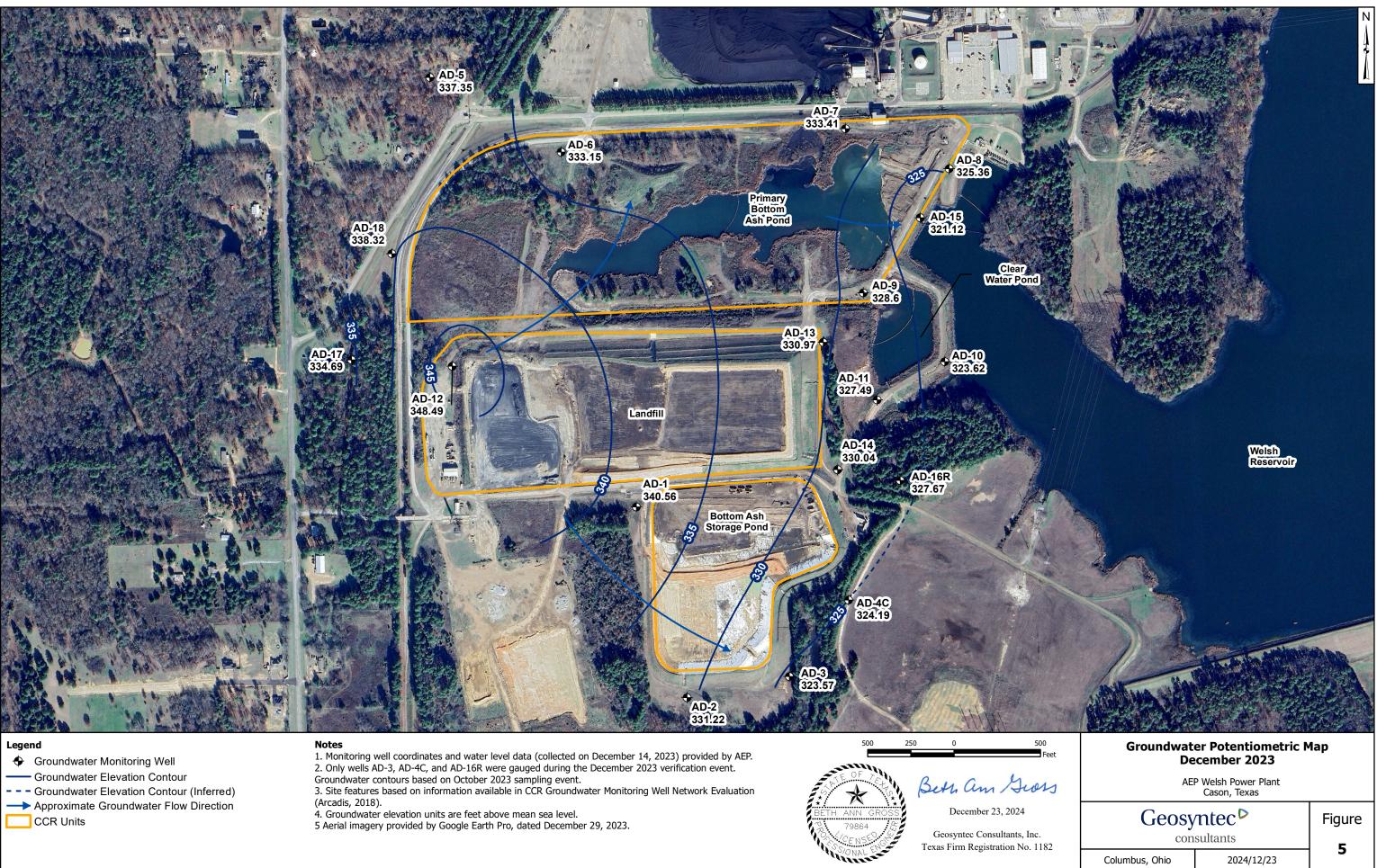


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2023/11/30

4

Columbus, Ohio





ATTACHMENT C Chemical Analysis of Wells in Titus County

Well	Depth of well (ft)	Date of collection	Silica (SiO ₂)	Iron (Fe) (total)	Manga- nese (Mn)	Cal- cium (Ca)	Magne- sium (Mg)	Sodium (Na)	Potas- sium (K)	Bicar- bonate (HCO ₃)	Sul- fate (S0 ₄)	Chlo- ride (Cl)	Fluo- ride (F)		Phos- phate (P0 ₄)	Boron (B)	Dis- solved solids	Hard- ness as CaCO3	Per- cent so-	Sodium adsorp- tion ratio (SAR)	Residual sodium carbonate (RSC)	Specific conduct- ance (micromhos at 25°C)	рН
YA-16-41-101	22	Aug. 20, 1963						*1	.7	7	0.0	18		40				27	57	1.4	0.00	149	5.1
102	34	do								34		358						324				1,250	5.8
201	30	May 26, 1942				12	1.2	*3	2	55	15	24		15			1 2 7	36					
301	60	do				179	126	*12	6	12	1,025	115	0	3.0			1,580	968					
302	60	do				134	39	*15	2	116	211	361		1.0			955	494					
801	200	Feb. 25, 1963	13	0.13		6.8	2.0	*14	7	308	.0	65	.4	.0			385	25	93	13	4.55	703	7.3
802	31	do							-	22	4.0	238		44				181			.00	955	5.2
902	470	July 30, 1963	11	.13		3.2	.7	*32	6	406	.0	272	.9	1.5			815	11	98	43	6.43	1,410	7.5
903	27	May 26, 1942				82	63	*15	5	268	296	191	0	3.0			922	464					
42-401	48	June 3, 1942				226	63	*42	1	549	33	890		0			1,903	824]				
702	22	do				12	5.8	*	58	55	18	77		10			208	54					
49-103	20	May 22, 1942				.8	1.0	*	2	18	11	3.0		1.5			38	6					
202	315	Feb. 20, 1963	51	22		9.8	3.2	*:	4	91	.0	11	.1	.0			144	38	58	1.7	. 74	2 15	5.8
203	30	do						*2	75	64	1,420	700						1 ,92 0	24	2.7	.00	4,090	5.5
206	485	Feb. 25, 1963	14	. 68		20	.7	*	58	122	13	13	.2	1.0			162	8	94	8.9	1.84	2 84	7.3
301	24	May 26, 1942				2.4	1.2	*:	28	31	7	22		10			86	11					
401	24	May 22, 1942				21	3.6	*	3	43	26	20		7.0			112	67					
402	395	Mar. 12, 1963	50	11		9.0	3.9	16	2.6	64	3.4	14	.2	0		0.00	130	38	45	1.1	.28	156	5.9
503	360	Feb. 20, 1963	54	12		9	3.2	*2	26	78	4.6	16	.1	.0			151	36	61	1.9	.57	218	5.8
601	22	May 25, 1942				49	19	*10	59	171	74	138	.2	33			506	2 02					
603	350	July 30, 1963	11	.12		1.5	.1	*{	36	204	.0	16	.2	1.8			217	4	98	19	3.26	353	7.4
701	437	May 27, 1942	20	.07		3.7	1.2	*2	31	370	2	149	.2	2.0			594	14					8.2
701	437	June 22, 1949	15	.14		1.6	.7	196	1.6	<u>a</u> β37	1.6	109	.1	2.2		.79	509					869	8.5
701	437	Feb. 19, 1963	12	2.8		1.5	.5	170	1.1	322	3.2	74	.2	.0		.20	421	6	98	30	5.17	758	7.4
702	597	May 27, 1942	20	.05		3.8	1.0	224		380	2	132	0	.0			567	14					8.4
702	597	Feb. 19, 1963	12	1.4		2.5	.7	218	1.2	368	.0	126	.3	.0		.27	542	9	98	32	5.85	991	7.7
706	430	May 14, 1942	39	5.6		14	6.6	30		126	2	15	.1	.5			176	62					<u> </u>

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See footnotes at end of table.

Well	Depth of well (ft)	Date of collection	Silica (SiO ₂)	Iron (Fe) (total)	Manga- nese (Mn)	Cal- cium (Ca)	Magne- sium (Mg)	Sodium (Na)	Potas- sium (K)	Bicar- bonate (HCO ₃)	fate		Fluo- ride (F)	Ni- trate (NO ₃)	Phos- phate (P0 ₄)	Boron (B)	Dis- solved solids	ness	Per- cent so- dium	Sodium adsorp- tion ratio (SAR)	Residual sodium carbonate (RSC)	Specific conduct- ance (micromhos at 25°C)	рН
YA-16-49-708	460	Mar. 14, 1963	13	0.13		1.5	0.2	152	.8	298	3.2	62	0.2	0.0		0.10	380	5	98	30	4.78	651	8.0
804	18	May 14, 1942				10	1.2	,	1 7.4	24	18	6.0		0			55	31					
924	300	Apr. 25, 1963	33	2.7		20	4.0	*!	 55 	156	13	31	.5	.0			234	66	64	2.9	1.23	352	7.1
50-101	35	June 3, 1942				13	2.4	*/	46 1	49	12	14		86			197	42					
102	31	do				79	35	*/	44	110	11	141		169			533	342					
202	48	do				47	12	*	38	49	2	1 2 5		39			287	168					
403	310	July 30, 1963	18	.09		27	6.9	*1	07	284	57	26	.2	.0			382	96	71	4.8	2.73	601	7.2
404	10	May 25, 1942				31	28	-	*6.4	18	74	78		5.0			231	192					
409	300	July 30, 1963	12	. 05		4.2	.9	*1	16	270	23	13	.2	2.0			304	14	95	13	4.15	485	7.3
501	37	May 25, 1942				308	97	*	76 1	488	274	460	0	2.0			1,457	1,170					
703	18	June 3, 1942				1.6	3.2	,	*2.3	12	4	4.0	.3	2.0			23	17					
57-102	246	Aug. 22, 1963	34	2.9		45	9.7	*	34	130	75	29	.2	.2			291	152	33	1.2	.00	452	6.6
110	700	June 3, 1963	12	.09		4.5	1.2	*4	20	396	0	425	.5	.7			1,060	16	98	46	6.17	1,890	7.6
114	475	Aug. 22, 1963	13	.06		6.0	.7	*	56	155	.0	8.5	.2	.5			161	18	87	5.7	2.18	272	7.2
301	20	May 13, 1942				4.8	3.6	*	10	12	26	7.0	.2	2.0			60	27					
302	420	July 31, 1963	13	.22		3.5	.5	*1	 04 	266	.2	15	.2	2.2		.06	271	11	95	14	4.14	440	7.3
401	300	Aug. 22, 1963	13			4.8	1.0	*1	 57 	286	.0	86	.6	.0			403	16	96	17	4.37	688	7.7
402	300	May 1, 1963	13	1.1	0.00	5.0	1.1	157	1.6	296	.2	88	.5	.0	0.94	.28	414	17	95	17	4.51	708	7.5
601	18	May 13, 1942				8.8	2.4	*	11	18	5	18	.2	12			67	32					
58-101	9	May 14, 1942				12	6.1	*	31	12	63	29	.2	6.0			153	54					
103	24	do				8.8	3.6	*	13	6	12	20		25			85	37					
203	21	do				13	2.4	*:	1 29 1	61	5	28		12			119	42					
401	13	May 13, 1942				4.8	2.4		 *8.1 	18	2	10	.1	10			47	22					
701	25	May 14, 1942				13	12	*	 38 	12	2	35		130			236	83					
17-48-102	26	Aug. 21, 1963							L	53		38						53			.00	299	5.6
202	18	Mar. 22, 1942				11	1.0	*1	1 04 1	12	30	84		120			356	31					
202	18	Aug. 21, 1963							[22		89						86			.00	696	5.6

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See footnotes at end of table.

Well	Depth of well (ft)		Silica (SiO ₂)			Cal- cium (Ca)	Magne- sium (Mg)	Sodium (Na)	Potas- sium (K)	Bicar- bonate (HCO ₃)	fate	Chlo- ride (Cl)	Fluo- ride (F)	Ni- trate (NO ₃)	phate		Dis- solved solids	Hard- ness as CaCO ₃	cent so-	Sodium adsorp- tion ratio (SAR)	Residual sodium carbonate (RSC)	Specific conduct- ance (micromhos at 25°C)	рН
YA-17-48-801	13	May 22, 1942				11	4.9	*37	,	43	12	40		29			155	48					
802	18	May 20, 1942				5.6	6.1	*17	,	31	7	12		33			96	39					
901	25	May 22, 1942				13	4.9	*37	,	12	22	39		55			177	53					
56-201	40	May 27, 1942				2 2	15	*17		12	2	18		141			221	114					
303	20	May 20, 1942				26	4.6	*34	-	55	11	65		10			178	83					
304	310	Aug. 13, 1963	18	1.8		7.5	2.6	*117	7	190	74	35	0.2	.0			347	29	90	9.4	2.53	560	7.2
401	11	May 27, 1942				16	7.3	*134		31	122	102		82			478	70					
402	30	do				4.4	1.2	*19		18	30	4.0	.2	6.0			74	16					
415	225	Jan. 17, 1963	12	.37		3.5	1.2	*132		248	71	15	.1	2.8			360	14	96	15	3.79	526	7.5
601	28	May 20, 1942				98	55	*67	7	171	185	199	.1	1.5			690	469					
701	38	May 15, 1942				6.0	0	*5	.1	18	4	5.0	0	0			29	15					
+ 707	260	Oct. 15, 1962	22	.1		6.7	2.5	*191	.1	201.3	198	50.0						27				892	8.02
707	260	July 27, 1963	7.8	1.7	0.00	8.2	2.3	18 2	2.4	184	202	50	.1	2.8	0.24	0.09	548	30	92	14	2.42	866	7.0
801	Spring	May 15, 1942				8.8	2.4	*]	.2	37	2	1.0		1.5			35	32					
901	502	May 29, 1942				5.2	4.9	*297	7	323	2	288	.2	7.0			764	33					
64-101	380	July 31, 1963	14	.09		3.8	.9	*82		187	1.8	24	.2	1.2			220	13	93	9.9	2.80	356	7.5
102	17	May 15, 1942				4.8	2.4	*22		49	3	15	.1	6.0			77	22					
201	48	do				48	22	*124		43	30	254		66			565	208					
301	40	do				205	126	*239)	580	418	450	0	9.0			1,732	1,033					
401	32	do				24	18	*127	, 	98	30	195		32			474	136					

Titus County

* Sodium and potassium calculated as sodium (Na).

† Analyses by Curtis Laboratories.

a Includes the equivalent of 5 ppm as carbonate (CO_3) .

ATTACHMENT D Certification by a Qualified Professional Engineer

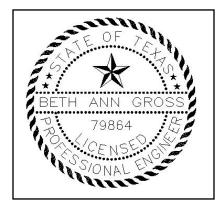
CERTIFICATION BY A QUALIFIED PROFESSIONAL ENGINEER

I certify that the selected and above described alternative source demonstration is appropriate for evaluating the groundwater monitoring data for the Welsh Bottom Ash Storage Pond CCR management area and that the requirements of 30 TAC §352.941(c) have been met.

Beth Ann Gross Printed Name of Licensed Professional Engineer

Beth am Geos

Signature



Geosyntec Consultants 2039 Centre Pointe Blvd, Suite 103 Tallahassee, Florida 32308

Texas Registered Engineering Firm No. F-1182

79864 License Number

Texas Licensing State December 23, 2024 Date