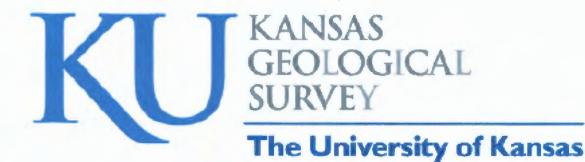
ANALYSIS OF CHEROKEE GROUP CUTTINGS SAMPLES FOR GAS CONTENT -- DART CHEROKEE BASIN C. BONNER #D3-7; sec. 7-T.30S.-R.14E., WILSON COUNTY, KS

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SUMMARY

Cuttings samples from the Pennsylvanian Cherokee Group were collected from the Dart Cherokee Basin C. Bonner et al. #D3-7; NW SW SE sec. 7-T.30S.-R.14E., Wilson County, KS. The samples calculate as having the following gas contents:

•	Dawson coal at 789' depth ¹	(scf/ton)
٠	Unidentified coal at 939' depth ²	(129 scf/ton)
•	Mulberry coal at 998' depth ²	(94 scf/ton)
•	Shale associated with "Lexington" coal at 1026' depth ³	(35 scf/ton)
	Mulky coal at 1096' depth ¹	(scf/ton)
	Bevier coal at 1152' depth ²	(63 scf/ton)
	Mineral coal at 1222' depth ¹	(scf/ton)
	Weir-Pittsburg coal at 1297' depth ²	(209 scf/ton)
	Riverton coal at 939' depth ⁴	(scf/ton)
	1	

¹no significant coal or gas-bearing shale in sample, no significant gas evolved off sample ²assuming accompanying dark shales in sample desorb 3 scf/ton

³no significant coal in sample

⁴canister defective

BACKGROUND

The Dart Cherokee Basin C. Bonner et al. #D3-7; NW SW SE sec. 7-T.30S.-R.14E., Wilson County, KS, was selected for cuttings desorption tests in association with an ongoing coalbed gas research project at the Kansas Geological Survey. The samples were gathered April 9, 2005, by personnel from Dart Cherokee Basin L.L.C., and turned over to LeaAnn Davidson of the Kansas Geological Survey. Samples were obtained during normal drilling of the well, with no cessation of drilling before zones of interest (i.e., coals and dark shales in the Cherokee Group) were penetrated.

The samples were canistered, with surface time and canistering times noted. These samples were collected in canisters that were supplied by Dart Cherokee Basin L.L.C. and the Kansas Geological Survey. Lag times for samples to reach the surface (important for lost-gas determinations) were determined by the wellsite geologist and driller.

The cuttings samples from the Pennsylvanian Cherokee Group were:

•	Unidentified coal at 939' depth	(136 grams)
٠	Mulberry coal at 998' depth	(195 grams)
٠	Shale associated with "Lexington" coal at 1026' depth	(403 grams)
٠	Bevier coal at 1152' depth	(492 grams)
•	Weir-Pittsburg coal at 1297' depth	(356 grams)

If correct wellsite procedures were followed, the cuttings were caught in kitchen strainers as they exited the air-stream pipe emptying to the mud pit. The samples were then washed in water while in the kitchen strainers to rid them of as much drilling mud as possible before the cuttings were placed in desorption canisters. Water with zephyrn chloride biocide was then added to the canisters, with a headspace of 1 to 2 inches being preserved at the top of the canister.

Desorption measurements at the Kansas Geological Survey in Lawrence, KS were continued at approximately 70 °F. Desorption measurements were periodically made until the canisters produced negligible gas with daily testing for at least two successive days.

DESORPTION MEASUREMENTS

The equipment and method for measuring desorption gas is that prescribed by McLennan and others (1995). The volumetric displacement apparatus is a set of connected dispensing burettes, one of which measures the gas evolved from the desorption canister. The other burette compensates for the compression that occurs when the desorbed gas displaces the water in the measuring burette. This compensation is performed by adjusting the cylinders so that their water levels are identical, then figuring the amount of gas that evolved by reading the difference in water level using the volumetric scale on the side of the burette.

The desorption canisters were obtained from SSD, Inc., in Grand Junction, CO. These canisters are 12.5 inches high (32 cm), 3 1/2 inches (9 cm) in diameter, and enclose a volume of approximately 150 cubic inches (2450 cm³). The desorbed gas that collected in the desorption canisters was periodically released into the volumetric displacement apparatus and measured as a function of time, temperature, and atmospheric pressure.

The time and atmospheric pressure were measured in the field using a portable weather station (model BA928) marketed by Oregon Scientific (Tualatin, OR). The atmospheric pressure was displayed in millibars on this instrument, however, this measurement was not the actual barometric pressure, but rather an altitude-compensated barometric pressure automatically converted to a sea-level-equivalent pressure. To translate this measurement to actual atmospheric pressure, a regression correlation was determined over several weeks by comparing readings from the Oregon Scientific instrument to that from a pressure transducer in the Petrophysics Laboratory in the Kansas Geological Survey in Lawrence, KS (Figure 1). The regression equation shown graphically in Figure 1 was entered into a spreadsheet and was used to automatically convert the millibar measurement to barometric pressure in pounds per square inch (psi).

A spreadsheet program written by K.D. Newell (Kansas Geological Survey) was used to convert all gas volumes at standard temperature and pressure. Conversion of gas volumes to standard temperature and pressure was by application of the perfect-gas equation, obtainable from basic college chemistry texts:

n = PV/RT

where n is moles of gas, T is degrees Kelvin (i.e., absolute temperature), V is in liters, and R is the universal gas constant, which has a numerical value depending on the units in which it is measured (for example, in the metric system R = 0.0820 liter atmosphere per degree mole). The number of moles of gas (i.e., the value n) is constant in a volumetric conversion, therefore the conversion equation, derived from the ideal gas equation, is:

 $(P_{stp}V_{stp})/(RT_{stp}) = (P_{rig}V_{rig})/(RT_{rig})$

Customarily, standard temperature and pressure for gas volumetric measurements in the oil industry are 60 °F and 14.7 psi (see Dake, 1978, p. 13), therefore P_{stp} , V_{stp} , and T_{stp} , respectively, are pressure, volume, and temperature at standard temperature and pressure, where standard temperature is degrees Rankine (°R = 460 + °F). P_{rig} , V_{rig} , and T_{rig} , respectively, are ambient pressure, volume, and temperature measurements taken at the rig site or in the desorption laboratory.

The universal gas constant R drops out as this equation is simplified and the determination of Vstp becomes:

 $V_{stp} = (T_{stp}/T_{rig}) (P_{rig}/P_{stp}) V_{rig}$

The conversion calculations in the spreadsheet were carried out in the English metric system, as this is the customary measure system used in American coal and oil industry. V is therefore converted to cubic feet; P is psia; T is °R.

The desorbed gas was summed over the time period for which the coal samples evolved all of their gas.

Lost gas for samples (i.e., the gas lost from the sample from the time it was drilled, brought to the surface, to the time it was canistered) are normally determined using the direct method (Kissel and others, 1975; also see McLennan and others, 1995, p. 6.1-6.14) in which the cumulative gas evolved is plotted against the square root of elapsed time. Time zero is assumed to be the moment that the rock is cut and its cuttings circulated off bottom. Lost gas, however, had to be inferred for the samples collected from this well because no desorption apparatus was on site when those samples were collected. The procedure used to infer lost gas for these samples is outlined in the section below on Lost Gas.

LITHOLOGIC ANALYSIS

Upon removal from the canisters, the cuttings were washed of drilling mud and dried in air for at least a week. After drying, the cuttings were weighed and then dry sieved into 5 size fractions: >0.0930", >0.0661", >0.0460", >0.0331", and <0.0331". For large sample sizes, the cuttings were run through a sample splitter and a lesser portion (approximately

75 grams) was sieved and weighed, and the derived size-fraction ratios were applied to the entire sample.

The size fractions were then inspected and sorted by hand under a dissecting microscope. Three major lithologic categories were differentiated: coal, dark shales (generally Munsell rock colors N3 [dark gray], N2 [grayish black], and N1 [black] on dry surface), and lighter-colored lithologies and/or dark and light-colored carbonates. The lightercolored lithologies are considered to be incapable of generating significant amounts of gas. After sorting, and for every size class, each of these three lithologic categories was weighed and the proportion of coal dark shale and light-colored lithologies were determined for the entire cuttings sample based on the weight percentages.

DATA PRESENTATION

Data and analyses accompanying this report are presented in the following order: 1) data tables for the desorption analyses, 2) lost-gas graphs, 3) "lithologic component sensitivity analyses" showing the interdependence of gas evolved from dark shale versus coal in each sample, and 4) a summary component analysis for all samples showing relative reliability of the data from all the samples.

Data Tables of the Desorption Analyses (Table 1)

These are the basic data used for lost-gas analysis and determination of total gas desorbed from the cuttings samples. Basic temperature, volume, and barometric measurements are listed at left. Farther to the right, these are converted to standard temperature, pressure and volumes. The volumes are cumulatively summed, and converted to scf/ton based on the total weight of coal and dark shale in the sample. At the right of the table, the time of the measurements are listed and converted to hours (and square root of hours) since the sample was drilled.

Lost-Gas Graphs (Figure 2, 3)

Surface and canistering times for each sample were noted by the wellsite geologist. Bottom hole time was inferred by subtracting lag time from surface time, with lag time determined from a depth-lag-time plot for a nearby well (Figure 2). To infer an approximate lost-gas value for each sample, a correlation of the total gas desorbed from a sample after it had been canistered to its rate of lost gas was developed using desorption data accumulated for 42 cuttings samples obtained from air-drilled wells in the Cherokee basin in southeastern Kansas (Figure 3). The rate of lost gas used in this correlation was that amount of gas lost by the square root of 0.6 square-root hours (the square root of 0.36 hours). By knowing the total gas given up by the sample after canistering (i.e., the total gas desorbed) a hypothetical rate of lost-gas could be calculated using the a regression line:

lost gas rate per square root of 0.36 hours = 0.1241 X (total gas desorbed in ccs) + 48.14

Once the hypothetical lost-gas rate was calculated, the lost gas could be calculated by taking the square root of the bottom-hole to canister time (derived from subtracting the lag time from the surface time), and multiplying it times the hypothetical lost-gas rate. Analysis of the lithology of the cuttings used in this correlation revealed no consistent relationship (see Figure 2), therefore further refinement of the relationship of the rate of lost gas to the total gas desorbed after canistering is not possible..

"Lithologic Component Sensitivity Analyses" (Figures 4-8)

The rapid penetration of an air-drilled well through relatively thin-bedded strata makes it difficult to collect lithologies that are unmixed. Mixed lithologies are more the norm rather than the exception. Some of this mixing is due to cavings from strata farther up hole. The mixing may also be due to collection of two or more successively drilled lithologies in the kitchen sieve at the exit line, or differential lifting of relatively less-dense coal compared to other lithologies, all of which are more dense than coal.

The total gas evolved from the sample is due to gas being desorbed from both the coal and dark shale. Both lithologies are capable of generating gas, albeit the coal will be richer in gas than the dark-colored shale. Even though dark-colored shale is less rich in sorbed gas than coal, if a sample has a large proportion of dark, organic-rich shale and only a minor amount of coal, the total volume of gas evolved from the dark-shale component may be considerable. The lighter-colored lithologies are considered to be incapable of generating significant amounts of gas.

The total amount of gas evolved from a cuttings sample can be expressed by the following equation:

Total gas $(cm^3) = [weight_{coal} (grams) X gas content_{coal} (cm^3/gram)] + [weight_{dark shale} (grams) X gas content_{dark shale} (cm^3/gram)]$

A unique solution for gas $content_{coal}$ in this equation is not possible because gas $content_{dark shale}$ is not known exactly. An answer can only be expressed as a linear solution to the above equation. The richer in gas the dark shales are, the poorer in gas the admixed coal has to be, and vice versa. If there is little dark shale in a sample, a relatively well constrained answer for gas $content_{coal}$ can be obtained. Conversely, if considerable dark shale is in a sample, the gas content of a coal will be hard to precisely determine.

The lithologic-component-sensitivity-analysis diagram therefore expresses the bivariant nature inherent in the determination of gas content in mixed cuttings. The gas content of dark shales in Kansas can vary greatly. Proprietary desorption analyses of dark shales in cores from southeastern Kansas have registered as much as 50 scf/ton, but can be as low as 2-4 scf/ton.

A value of 3 scf/ton for average dark shale is based on the assay of the gas content of cores of dark shales in nearby wells. However, high-gamma-ray shales (such as the Excello Shale), also colloquially known as "hot shales", typically have more organic

matter and associated gas content than dark shales with no excessive gamma-ray level. Determination of gas content for a coal associated with a "hot" shale therefore carries more uncertainty than if the coal were associated with a shale without a high gamma-ray value.

In general, shale gas content does not have to be very much greater that 10 scf/ton before the associated coal starts to have a gas content less than that of the dark shale. In all the lithologic-component-sensitivity-analysis diagrams, a "break-even" point is therefore noted where the gas content of the coal is equal to that of the dark shale. This "breakeven" point corresponds to the minimum gas content assignable to the coal and maximum gas content assignable to the dark shale. It can also be thought of the scf/ton gas content of the cuttings sample minus the weight of any of the lighter-colored lithologies, which are assumed to have no inherent gas content. Conversely, to assume that all the gas evolved from a cuttings sample is derived solely from the coal would result in an erroneously high gas content for the coal.

Summary Component Analysis for all Samples (Figure 9)

This diagram is a summary of the individual "lithologic component sensitivity analyses" for each sample, all set at a common scale. The steeper the angle of the line for a sample, the more uncertainty is attached to the results (i.e., *gas content_{coal}*) for that sample. If the coal content is miniscule (i.e., < approximately 5%), the results are a better reflection of the *gas content_{dark shale}*.

RESULTS and DISCUSSION

The least constrained results are those from the Bevier coal at 1152'. The unidentified coal at 939" has the best constrained results. Weir-Pitt at 1297' and Mulberry at 998' have roughly comparable reliabilities (see Figure 9).

REFERENCES

- Dake, L.P., 1978, Fundamentals of Reservoir Engineering, Elsevier Scientific Publishing, New York, NY, 443 p.
- Kissel, F.N., McCulloch, C.M., and Elder, C.H., 1975, The direct method of determining methane content of coals for ventilation design: U.S. Bureau of Mines, Report of Investigations, RI7767.
- McLennan, J.D., Schafer, P.S., and Pratt, T.J., 1995, A guide to determining coalbed gas content: Gas Research Institute, Chicago, IL, Reference No. GRI-94/0396, 180 p.

FIGURES and TABLES

FIGURE 1. Correlation of field barometer to Petrophysics Lab pressure transducer.

TABLE 1. Desorption measurements for samples.

FIGURE 2. Lag time determination for samples

FIGURE 3. Correlation of the rate of lost gas to the total gas desorbed after canistering.

FIGURE 4. Sensitivity analysis for unidentified coal at 939' depth.

FIGURE 5. Sensitivity analysis for Mulberry coal at 998' depth.

FIGURE 6. Sensitivity analysis for shale associated with "Lexington" coal at 1026' depth.

FIGURE 7. Sensitivity analysis for Bevier coal at 1152' depth.

FIGURE 8. Sensitivity analysis for Weir-Pittsburg coal at 1297' depth.

FIGURE 9. Lithologic component sensitivity analyses for all samples.



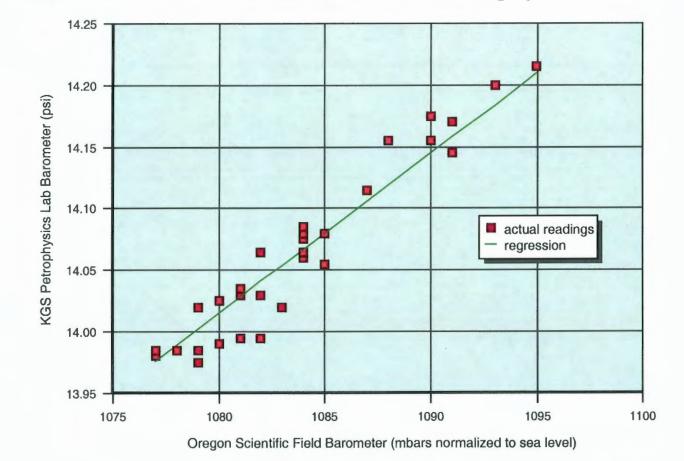


FIGURE 1.

TABLE 1 -- Desorption data for DART CHEROKEE BASIN C. BONNER et al. #D3-7; NW SW SE 7-T.30S.-R.14E.

SAMPLE:																				
		lbs.		grams										est. lost gas () = (TIME OF:				elapsed time (off bottom to canistering)
dry sample we	eight:		0.3894	176.62	2										0	off bottom		at surface	in canister	5.4 minutes
																4/5/05	12:42	4/5/05 12:43	4/5/05 12:4	7 0.090 hours
RIG/LAB MEASL	JREMENTS			CONVER	ISION OF R	IGA A	B MEASL	IREMENTS TO	STP (060 deg	F; 14.7 psl)	CUMULATIVE VC	DLUMES	SCF/TON	SCF/TON				TIME SINCE		0.299536679 SQRT (hrs)
measured cc	measured T	(F) mea	asured P	cubic ft	absolute 1	T (R)	psia	cubic ft (OS	FP) cc (OSTF	9	cubic ft (@STP)) cc (@STP)	without lost gas	with lost gas		TIME OF MEAS	URE	off bottom	in canister	SQRT hrs. (since off bottom)
-11		69	1072	-0.0004		529	13.914	-0.000361	439	-10.23	-0.00036143	9 -10.23	-1.86		1.86	4/9/05	14:24	97:42:00	97:36:3	7 9.884331035
-1		70	1074	-4E-05		530	13.940	-3.28573E	-05	-0.93	-3.28573E-0	5 -0.93	-0.17		0.17	4/10/05	10:31	117:49:00	117:43:3	7 10.85433861
3		71	1066	0.0001		531	13.836	9.76533E	-05	2.77	9.76533E-0	5 2.77	0.50		0.50	4/11/05	17:12	148:30:00	148:24:3	7 12.18605761
-2		73	1075	-7E-05		533	13.953	-6.54055E	-05	-1.85	-6.54055E-0	5 -1.85	-0.34		0.34	4/12/05	14:19	169:37:00	169:31:3	7 13.02369635
-7		69	1091	-0.0002		529	14.161	-0.000234	083	-6.63	-0.00023408	3 -6.63	-1.20		1.20	4/15/05	11:08	238:26:00	238:20:3	7 15.44128665
DESORPTION	TERMINATE	ED 4/15/2	005 DUE	TONOG	GAS BEING	EVOL	VED; no	coal in sample	; sample not sa	wed										

SAMPLE: 939' to 941' (? coal) cuttings in Dart SSD canister NOTE: lost gas is estimated by time interval between at surface and canister times, and total gas evolved lbs. grams est. lost gas (cc) = TIME OF: elapsed time (off bottom to canistering) 0.0315 14.29 4.7 minutes dry sample weight: 26 off bottom at surface in canister 4/5/05 14:10 4/5/05 14:11 4/5/05 14:15 0.078 hours CONVERSION OF RIG/LAB MEASUREMENTS TO STP (@60 deg F; 14.7 ps) CUMULATIVE VOLUMES RIGILAB MEASUREMENTS SCF/TON TIME SINCE 0.279880927 SQRT (hrs) SCF/TON measured cc measured T (F) measured P cubic ft absolute T (R) psia cubic ft (@STP) cc (@STP) cubic ft (OSTP) cc (OSTP) without lost gas with lost gas TIME OF MEASURE off bottom in canister SQRT hrs. (since off bottom) 1072 0.0004 529 13.914 0.000361439 11 69 10.23 0.000361439 10.23 22.95 81.24 4/9/05 14:24 96:13:19 96:08:37 9.809278487 1074 0 530 13.940 0 70 0 0.00 0.000361439 10.23 22.95 81.24 4/10/05 10:31 116:20:19 116:15:37 10.78603779 531 13.836 8.13778E-05 2.5 71 1066 9E-05 2.30 0.000442817 12.54 28.11 86.40 4/11/05 17:12 147:01:19 146:56:37 12.12526059 1075 -7E-05 533 13.953 -6.54055E-05 -1.85 0.000377411 -2 73 10.69 23.96 82.25 4/12/05 14:19 168:08:19 168:03:37 12.96682733 -9 69 1091 -0.0003 529 14.161 -0.000300964 -8.52 7.64471E-05 2.16 4.85 63.14 4/15/05 11:08 236:57:19 236:52:37 15.39335174 70 1089 -7E-05 530 14.135 -6.66324E-05 -1.89 9.81473E-06 0.28 0.62 58.91 4/16/05 15:38 -2 265:27:19 285:22:37 16.29279834

DESORPTION TERMINATED 4/16/2005 DUE TO NO MORE GAS BEING EVOLVED; sample air dried for 35 days

SAMPLE:	998' to 1000)' (Mult	berry coa	i) cuttings	in Dart SSD	caniste	r					NOTE: lost gas is	estimated by	time int	erval between at s	urface and caniste	er times, and total gas evolved
		lbs		grams								est. lost gas (cc) =	TIME OF:				elapsed time (off bottom to canistering)
dry sample w	veight:		0.2467	111.91								1	7 off bottom		at surface	in canister	5.4 minutes
													4/5/05	14:46	4/5/05 14:48	4/5/05 14:51	0.090 hours
RIG/LAB MEAS	SUREMENTS			CONVER	SION OF RIGA	AB ME/	SUREMENTS TO STI	P (060 deg F; 14.7 psi)	CUMULATIVE VO	LUMES	SCF/TON	SCF/TON			TIME SINCE		0.299536679 SQRT (hrs)
measured cc	measured T	(F) me	asured P	cubic ft	absolute T (I) psia	cubic ft (@STP)	oc (@STP)	cubic ft (@STP)	cc (@STP)	without lost gas	with lost gas	TIME OF ME	SURE	off bottom	in canister	SQRT hrs. (since off bottom)
56	6	69	1072	0.002	52	9 13.9	14 0.001840053	52.10	0.001840053	52.10	14.92	19.7	8 4/9/05	14:33	95:46:56	95:41:33	9.786839236
5	5	70	1074	0.0002	53	0 13.9	40 0.000164286	4.65	0.002004339	56.76	16.25	21.1	4/10/05	10:32	115:45:56	115:40:33	10.7594403
7	7	71	1066	0.0002	53	1 13.8	36 0.000227858	6.45	0.002232197	63.21	18.10	22.9	6 4/11/05	17:13	146:26:56	146:21:33	12.10160687
1	I	73	1075	4E-05	53	3 13.9	53 3.27028E-05	0.93	0.0022649	64.13	18.38	23.2	3 4/12/05	14:20	167:33:56	167:28:33	12.94471149
-2	2	69	1091	-7E-05	52	9 14.1	61 -6.68809E-05	-1.89	0.002198019	62.24	17.82	22.6	4/15/05	11:09	236:22:58	236:17:33	15.37472674
1	l I	70	1089	4E-05	53	0 14.1	35 3.33162E-05	0.94	0.002231335	63.18	18.09	22.9	5 4/16/05	15:37	264:50:56	264:45:33	16.27417859
1	1	71	1089	4E-05	53	1 14.1	35 3.32534E-05	0.94	0.002264589	64.13	18.36	23.2	2 4/17/05	11:03	284:16:56	284:11:33	16.86067087
4	1	72	1082	0.0001	53	2 14.0	44 0.00013191	3.74	0.002396499	67.86	19.43	24.2	4/18/05	11:25	308:38:56	308:33:33	17.56840599
1	l i i i i i i i i i i i i i i i i i i i	70	1076	4E-05	53	0 13.9	66 3.29185E-05	0.93	0.002429418	68.79	19.69	24.5	6 4/20/05	22:46	367:59:56	367:54:33	19.18329713
2	2	70	1072	7E-05	53	0 13.9	14 6.55922E-05	1.86	0.00249501	70.65	20.23	25.0	4/21/05	17:12	386:25:56	386:20:33	19.65787939
-3	3	67	1081	-0.0001	52	7 14.0	31 -9.97791E-05	-2.83	0.002395231	67.83	19.42	24.2	4/22/05	16:11	409:24:56	409:19:33	20.23401976
-5	5	63	1083	-0.0002	52	3 14.0	57 -0.00016788	-4.75	0.00222735	63.07	18.06	22.9	2 4/24/05	13:45	454:58:56	454:53:33	21.33031229
DESORPTION	N TERMINATE	D 4/28/2	2005 DUE	TO NO M	ORE GAS BEI	NG EVO	LVFD: sample air drie	d for 33 days									

DESORPTION TERMINATED 4/28/2005 DUE TO NO MORE GAS BEING EVOLVED; sample air dried for 33 days

SAMPLE: 1026																		
		lbs.	grams									est. lost gas (c	c) = T	IME OF:				elapsed time (off bottom to canistering)
dry sample weight:		0.8738	396.36										31 0	ff bottom		at surface	in canister	6.1 minutes
														4/5/05	15:17	4/5/05 15:19	4/5/05 15:	23 0.102 hours
RIGALAB MEASUREME						IREMENTS TO ST	P (@60 deg F; 1	14.7 pei)	CUMULATIVE VOL	UMES	SCF/TON	SCF/TON				TIME SINCE		0.319722102 SQRT (hrs)
measured cc measured	ured T (F)) measured P	cubic ft	absolute T (R)	psia	cubic ft (@STP)	cc (OSTP)		cubic ft (@STP)	cc (@STP)	without lost gas	with lost gas	Т	ME OF MEAS	SURE	off bottom	in canister	SQRT hrs. (since off bottom)
306	69	1072	0.0108	529	13.914	0.010054576		284.71	0.010054576	284.71	23.01	2	5.52	4/9/05	14:46	95:28:12	95:22:0	9.770875089
43	70	1074	0.0015	530	13.940	0.001412863	1	40.01	0.011467439	324.72	26.25	j 2	8.75	4/10/05	10:33	115:15:12	115:09:0	10.73561052
39	71	1066	0.0014	531	13.836	0.001269494		35.95	0.012736932	360.67	29.15	3	1.66	4/11/05	17:13	145:55:12	145:49:0	12.0797351
11	73	1075	0.0004	533	13.953	0.00035973		10.19	0.013096663	370.85	29.98	3.	2.48	4/12/05	14:21	167:03:12	166:57:0	04 12.92491135
20	69	1091	0.0007	529	14.181	0.000668809		18.94	0.013765472	389.79	31.51	3	4.01	4/15/05	11:10	235:52:12	235:46:	15.35805977
2	70	1089	7E-05	530	14.135	6.86324E-05		1.89	0.013832104	391.68	31.68	3	4.16	4/16/05	15:36	264:18:12	264:12:0	16.25740857

1.5	71	1089	5E-05	531	14.135	4.98802E-05	1.41	0.013881984	393.09	31.77	34.28	4/17/05	11:03	283:45:12	283:39:04	16.84497947
5	72	1082	0.0002	532	14.044	0.000164888	4.87	0.014046872	397.76	32.15	34.68	4/18/05	11:26	308:08:12	308:02:04	17.553822
0	70	1076	0	530	13.986	0	0.00	0.014046872	397.76	32.15	34.68	4/20/05	21:11	365:53:12	365:47:04	19.12816423
1	70	1072	4E-05	530	13.914	3.27961E-05	0.93	0.014079668	398.69	32.23	34.73	4/21/05	17:13	385:55:12	385:49:04	19.64484665
-4	67	1081	-0.0001	527	14.031	-0.000133039	-3.77	0.01394663	394.92	31.92	34.43	4/22/05	16:11	408:53:12	408:47:04	20.22094624
-7	63	1083	-0.0002	523	14.057	-0.000235033	-6.66	0.013711597	388.27	31.38	33.89	4/24/05	13:45	454:27:12	454:21:04	21.31791109
		DAAR DUIT	TO 110 110	DE GAO DENI												

DESORPTION TERMINATED 4/28/2005 DUE TO NO MORE GAS BEING EVOLVED; sample air dried for 33 days

SAMPLE: 1	SAMPLE: 1096' to 1097' (Mulky coal) cuttings in Dart SSD canister																
		lbs.	grams									est. lost gas (co			on a sectoon at o		elapsed time (off bottom to canistering)
dry sample weig	ght:	0.00	00	0									0 off bottom		at surface	in canister	8.9 minutes
													4/5/05	16:39	4/5/05 16:41	4/5/05 16:48	0.148 hours
RIG/LAB MEASUF	REMENTS		CONVE	RSION OF RIG	SALAB MEA	SUREMENTS TO	STP (@60 deg F; 1	4.7 psi)	CUMULATIVE VOI	LUMES	SCF/TON	SCF/TON			TIMESINCE		0.384418753 SQRT (hrs)
measured cc m	easured T ((F) measure	P cubic f	absolute T	(R) psia	cubic ft (@ST	P) cc (OSTP)		cubic ft (@STP)	cc (@STP)	without lost ga	as with lost gas	TIME OF MEA	SURE	off bottom	in canister	SORT hrs. (since off bottom)
1	e	5 <mark>9</mark> 10	72 4E-0	5 f	29 13.9	14 3.28581E	-05	0.93	3.28581E-05	0.93	#DIV/0!	#DIV/0!	4/9/05	14:36	93:56:41	93:47:49	9.692508562
-0.5	7	70 10	74 -2E-0	5 !	530 13.9	40 -1.64286E	-05	-0.47	1.64295E-05	0.47	#DIV/0!	#DIV/0!	4/10/05	10:33	113:53:41	113:44:49	10.67214703
3	7	71 10	66 0.000	1 5	31 13.8	36 9.76533E	-05	2.77	0.000114083	3.23	#DIV/0!	#DIV/0!	4/11/05	17:14	144:34:41	144:25:49	12.02406152
-2	7	73 10	75 -7E-0	5 5	33 13.9	53 -6.54055E	-05	-1.85	4.86773E-05	1.38	#DIV/01	#DIV/0!	4/12/05	14:21	165:41:41	165:32:49	12.8722462
-9	6	59 10	91 -0.000	3 5	29 14.1	-0.000300	964	-8.52	-0.000252287	-7.14	#DIV/0!	#DIV/0!	4/15/05	11:11	234:31:41	234:22:49	15.31430885
-1	7	70 10	89 -4E-0	5 5	30 14.1	35 -3.33162E	-05	-0.94	-0.000285603	-8.09	#DIV/0!	#DIV/0!	4/16/05	15:35	262:55:41	262:46:49	16.21505645
DESORPTION T	COMINIATED	1/16/2005	UE TO NO	GAS BEING P	VOLVED.	olomes of leop on	howes too olomes										

DESORPTION TERMINATED 4/16/2005 DUE TO NO GAS BEING EVOLVED; no coal in sample; sample not saved

SAMPLE: 115																	
		lbs.	grams								est. lost gas	(CC) =	TIME OF:				elapsed time (off bottom to canistering)
dry sample weight	t:	0.7814	354.42									15	off bottom		at surface	in canister	4.4 minutes
													4/6/05	8:08	4/6/05 8:10	4/6/05 8:	12 0.073 hours
RIGALAB MEASURE	MENTS		CONVER	SION OF RIG/LA	B MEASU	REMENTS TO ST	P (@60 deg F; 14.7 psi)	CUMULATIVE VO	LUMES	SCF/TON	SCF/TON				TIME SINCE		0.270287912 SQRT (hrs)
measured oc mea	asured T (F)	measured P	cubic ft	absolute T (R)	psia	cubic ft (@STP)	cc (@STP)	cubic ft (@STP)	cc (@STP)	without lost gas	with lost gas		TIME OF MEAS	URE	off bottom	in canister	SQRT hrs. (since off bottom)
53	69	1072	0.0019	529	13.914	0.001741479	49.31	0.001741479	49,31	4.46		5.81	4/9/05	14:41	78:32:51	78:28:	8.862702748
1	70	1074	4E-05	530	13.940	3.28573E-05	0.93	0.001774336	50.24	4.54		5.90	4/10/05	10:34	98:25:51	98:21:	28 9.921231442
3	71	1066	0.0001	531	13.836	9.76533E-05	2.77	0.001871989	53.01	4.79		6.15	4/11/05	17:15	129:06:51	129:02:	28 11.36284149
-3	73	1075	-0.0001	533	13.953	-9.81083E-05	-2.78	0.001773881	50.23	4.54		5.90	4/12/05	14:22	150:13:51	150:09:	28 12.25686882
-10	69	1091	-0.0004	529	14.181	-0.000334405	-9.47	0.001439476	40.76	3.68		5.04	4/15/05	11:12	219:03:51	218:59:	28 14.80081642
-2	70	1089	-7E-05	530	14.135	-6.66324E-05	-1.89	0.001372844	38.87	3.51		4.87	4/16/05	15:33	247:24:51	247:20:	28 15.72940452
DESORPTION TEF	MINATED 4/1	6/2005 DUE	TO NO M	ORE GAS BEING	GEVOLVE	ED; sample air drie	d for 20 days										

SAMPLE: 1222' to 1223' (Mineral coal) cuttings in Dart SSD canister times, and total gas evolved																	
		lbs.	grams								est. lost gas (cc)	= TIME OF:				6	elapsed time (off bottom to canistering)
dry sample weight		0.0000	0.00									0 off bottom	а	t surface	in canister		7.1 minutes
												4/6/05	3:49	4/6/05 8:5	1 4/6/05 8	8:56	0.119 hours
RIG/LAB MEASURE!	AIGLAB MEASUREMENTS CONVERSION OF RIGUAB MEASUREMENTS TO STP (0 60 deg F; 14.7 per) CUMULATIVE VOLUMES SCF/TON SCF/TON TIME SINCE 0.344399639 SQRT (hrs)																
measured cc mea	asured T (F)	measured P	cubic ft	absolute T (R)	psia	cubic ft (@STP)	cc (@STP)	cubic ft (@STP)	cc (OSTP)	without lost gas	with lost gas	TIME OF MEASU	RE o	ff bottom	in canister		SQRT hrs. (since off bottom)
10	69	1072	0.0004	529	13.914	0.000328581	9.30	0.000328581	9.30	#DIV/0!	#DIV/0!	4/9/05 14	1:53	78:03:3	6 77:56	5:29	8.835157044
-1.5	70	1074	-5E-05	530	13.940	-4.92859E-05	-1.40	0.000279295	5 7.91	#DIV/01	#DIV/0!	4/10/05 10	0:34	97:44:3	6 97:37	7:29	9.886522813
1	71	1066	4E-05	531	13.836	3.25511E-05	0.92	0.000311846	8.83	#DIV/01	#DIV/0!	4/11/05 17	7:18	128:26:3	6 128:19	9:29	11.33328431
-2	73	1075	-7E-05	533	13.953	-6.54055E-05	-1.85	0.000246441	6.98	#DIV/01	#DIV/0!	4/12/05 14	:22	149:32:3	6 149:25	5:29	12.22879118
-9	69	1091	-0.0003	529	14.161	-0.000300964	-8.52	-5.45236E-05	-1.54	#DIV/0!	#DIV/0!	4/15/05 11	1:13	218:23:3	6 218:16	5:29	14.778137
-2	70	1089	-7E-05	530	14.135	-6.86324E-05	-1.89	-0.000121156	-3.43	#D1V/01	#DIV/0!	4/16/05 15	5:31	246:41:3	6 246:34	1:29	15.70647425
DECODETION TEE	ALLATTO 44	10/0005 0115	TONOG	AC DEINIC EVC	VED. as	and in complex services	have ten alon										

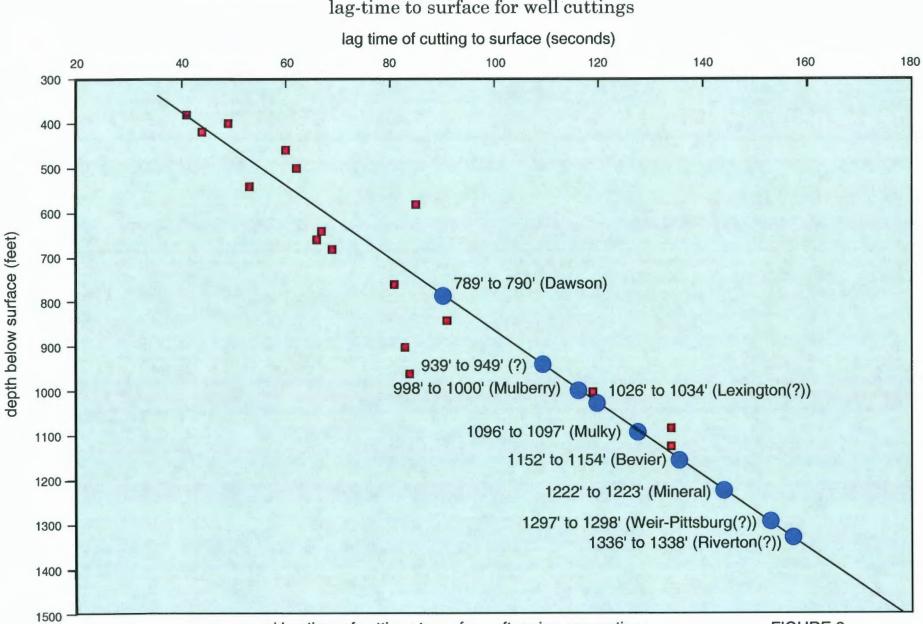
DESORPTION TERMINATED 4/16/2005 DUE TO NO GAS BEING EVOLVED; no coal in sample; sample not saved

SAMPLE:	1297' to 12				cuttings in Dar	t SSD ca	nister								ime inte	rval between at s		er times, and total gas evolved
		lb	98. (grams								est. lost ga	s (CC) =	TIME OF:				elapsed time (off bottom to canistering)
dry sample we	ight:		0.0887	40.24									31	off bottom		at surface	in canister	6.9 minutes
														4/6/05	9:37	4/6/05 9:40	4/6/05 9:44	0.115 hours
RIGALAB MEASU							JAEMENTS TO ST	P (@60 deg F; 14.7 psi)	CUMULATIVE	OLUMES	SCF/TON	SCF/TON				TIME SINCE		0.339118499 SQRT (hrs)
measured cc	measured T	(F) m	easured P	cubic ft	absolute T (F	i) psia	cubic ft (@STP)	cc (OSTP)	cubic ft (@ST	P) cc (@STP)	without lost	gas with lost ga	15	TIME OF MEA	SURE	off bottom	in canister	SQRT hrs. (since off bottom)
44		69	1072	0.0018	52	9 13.914	0.001445756	40.94	0.0014457	56 40.94	32	.59	57.28	4/9/2005 2	:573:00	#VALUE!	#VALUE!	#VALUE!
1		70	1074	4E-05	53	0 13.940	3.28573E-05	0.93	0.0014786	13 41.87	33	.34	58.02	4/10/05	10:35	96:57:24	96:50:30	9.846657639
5		71	1066	0.0002	53	1 13.836	0.000162756	4.81	0.0016413	69 46.48	37	.00	61.69	4/11/05	17:16	127:38:24	127:31:30	11.29778739
-1		73	1075	-4E-05	53	3 13.953	-3.27028E-05	-0.93	0.0016086	66 45.55	36	.27	60.95	4/12/05	14:23	148:45:24	148:38:30	12.19658422

-8	69	1091	-0.0003	529 14.161	-0.000267524	-7.58	0.001341142	37.98	30.24	54.92	4/15/05	11:13	217:35:24	217:28:30	14.75093217
-1	70	1089	-4E-05	530 14.135	-3.33162E-05	-0.94	0.001307826	37.03	29.48	54.17	4/16/05	15:30	245:52:24	245:45:30	15.68034864
DESORPTION TERMI	NATED 4/16/2	005 DUE 1	TO NO GAS BE	ING EVOLVED											
044015 1000	4- 1000L (D)	instan(2)	and) authings	in Dart CCD annister					NOTE	lost non in out	implad by t	ime intenu	I hohuson at sude	and appleter th	here and total and maked

SAMPLE:																		
		lbs.	grams								est. lost gas ((CC) =	TIME OF:					elapsed time (off bottom to canistering)
dry sample w	eight:	0.6578	3 298.36									0	off bottom		at surface	in caniste	э г	4.4 minutes
													4/6/05	10:05	4/6/05 10:0	8 4/6/05	10:10	0.073 hours
RIG/LAB MEAS	UREMENTS		CONVER	SION OF RIG/LA	B MEASU	REMENTS TO ST	TP (@60 deg F; 14.7 psi)	CUMULATIV	E VOLUMES	SCF/TON	SCF/TON				TIME SINCE			0.270801280 SQRT (hrs)
measured cc	measured T (F	F) measured F	cubic It	absolute T (R)	psia	cubic ft (@STP)	cc (@STP)	cubic ft (@	STP) cc (@STP)	without lost gas	with lost gas		TIME OF MEAN	SURE	off bottom	in caniste	31	SQRT hrs. (since off bottom)
0	6	9 1072	2 0	529	13.914		0.00)	0 0.00	0.00)	0.00	4/9/05	14:53	76:47:2	0 76	6:42:56	8.762926959
1	7	0 1074	4E-05	530	13.940	3.28573E-0	5 0.93	3.28573	E-05 0.93	0.10)	0.10	4/10/05	10:34	96:28:2	0 96	6:23:56	9.822027399
4	7	1 1066	0.0001	531	13.836	0.00013020	4 3.69	0.00016	3062 4.62	0.50)	0.50	4/11/05	17:16	127:10:2	0 127	7:05:56	11.27706621
-8	7:	3 1075	-0.0003	533	13.953	-0.00026162	2 -7.41	-9.85604	E-05 -2.79	-0.30)	-0.30	4/12/05	14:22	148:16:2	0 148	8:11:56	12.17670818
-1	6	9 1091	-4E-05	529	14.181	-3.34405E-0	5 -0.95	-0.00013	2001 -3.74	-0.40)	-0.40	4/15/05	11:13	217:07:2	0 217	7:02:56	14.73506777
SAMPLE DEC	ANISTERED 4/	15/2005 DUE	TO BAD G	ASKET ON CAN	ISTER; no	valid data obtain	ned; sample air dried for	10 days										





(based on lag times from Dart Cherokee Basin #CH-1 Holder; sec. 1-T.30S.-R.14E., Wilson County, KS lag-time to surface for well cuttings

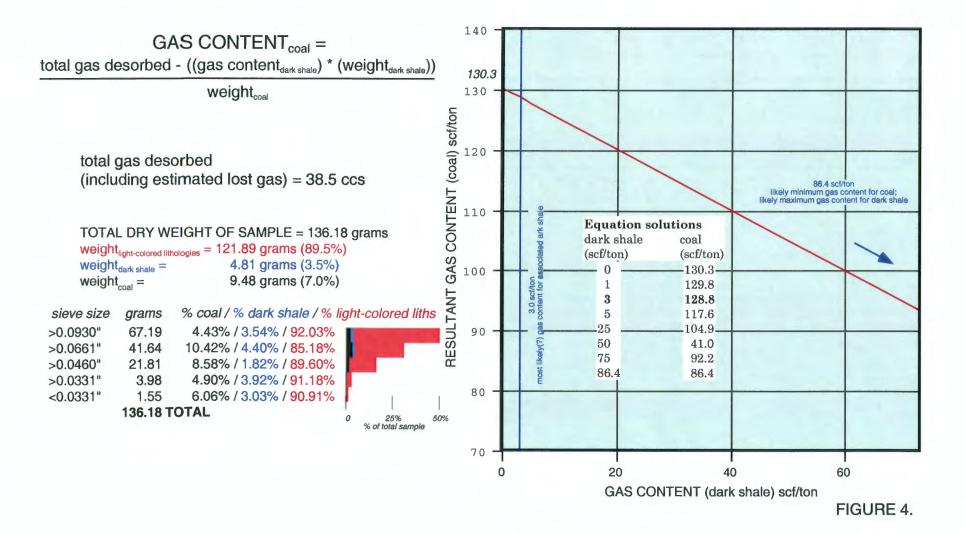
Dart Cherokee Basin #D3-7 C. Bonner, NW SW SE 7-T.30S.-R.14E., Wilson County, KS

measured lag time of cuttings to surface after pipe connections

FIGURE 2.

RELATIONSHIP of TOTAL GAS EVOLVED FROM a CUTTINGS SAMPLE to RATE of LOST-GAS (from 42 cuttings samples from air-drilled wells, Cherokee basin, southeastern Kansas) 600 500 lost gas rate (ccs/SQRT(0.36 hrs)) 400 **REGRESSION LINE** y = 0.1241 (x) + 48.14r squared = 0.81 300 200 2 100 0 0 500 1000 1500 2000 2500 3000 total gas evolved since canistering (ccs at standard temperature and pressure) number of samples (n = 42)100 LOST-GAS ALGORITHM (n=1)10 (n=5)ccs lost gas = \sqrt{X} (Y) 1 coal-to-dark-shale (n=15) 0.1 ratio in sample where X = bottom-hole to canister time (in hours) (n=9) 0.01 where $Y = \cos \log t$ gas at 0.36 hours (n=7) (i.e., value Y from regression equation) coal:dark shale ratio not yet determined (n=5) FIGURE 3.

LITHOLOGIC COMPONENT SENSITIVITY ANALYSIS for calculation of gas content of ? coal from 939' to 941'



LITHOLOGIC COMPONENT SENSITIVITY ANALYSIS for calculation of gas content of Mulberry coal from 998' to 1000'

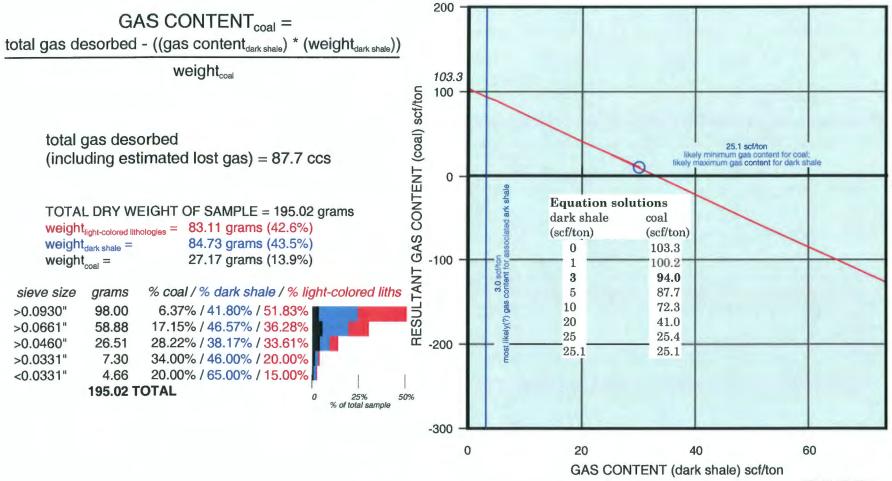


FIGURE 5.

LITHOLOGIC COMPONENT SENSITIVITY ANALYSIS for calculation of Lexington "coal" from 1026' to 1034'

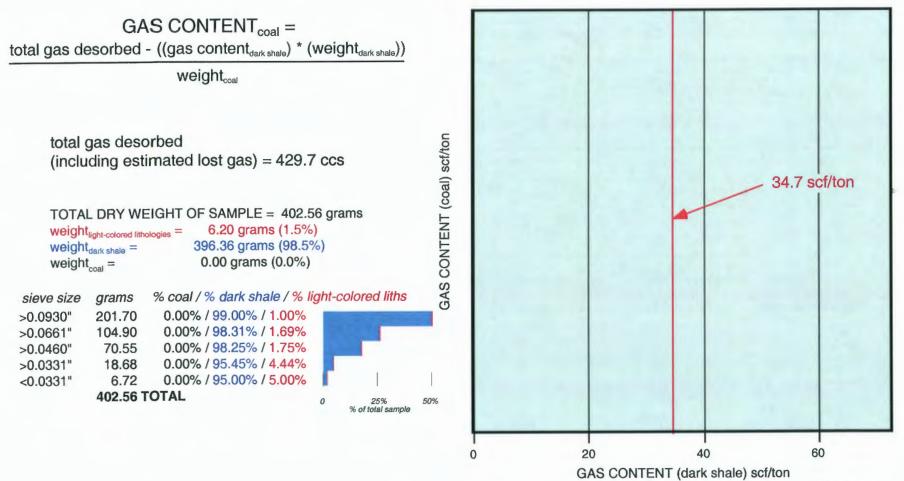
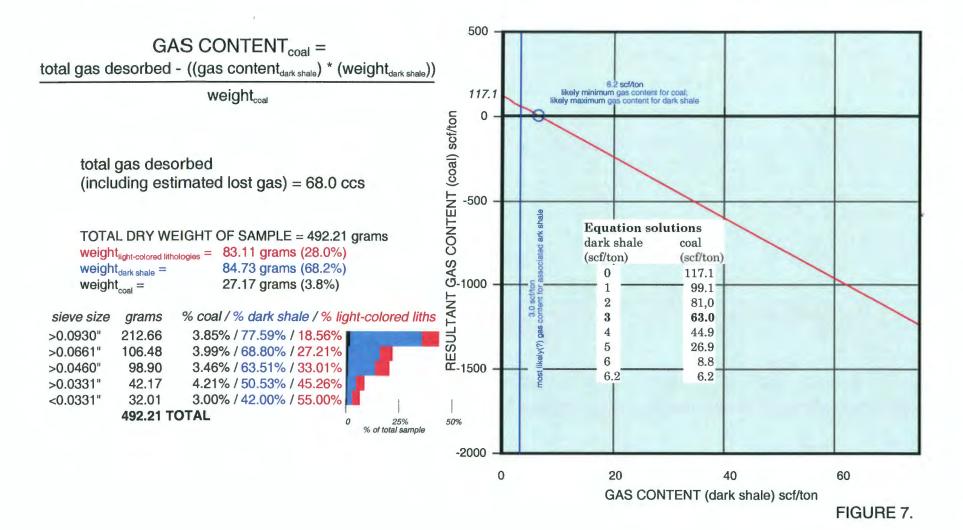
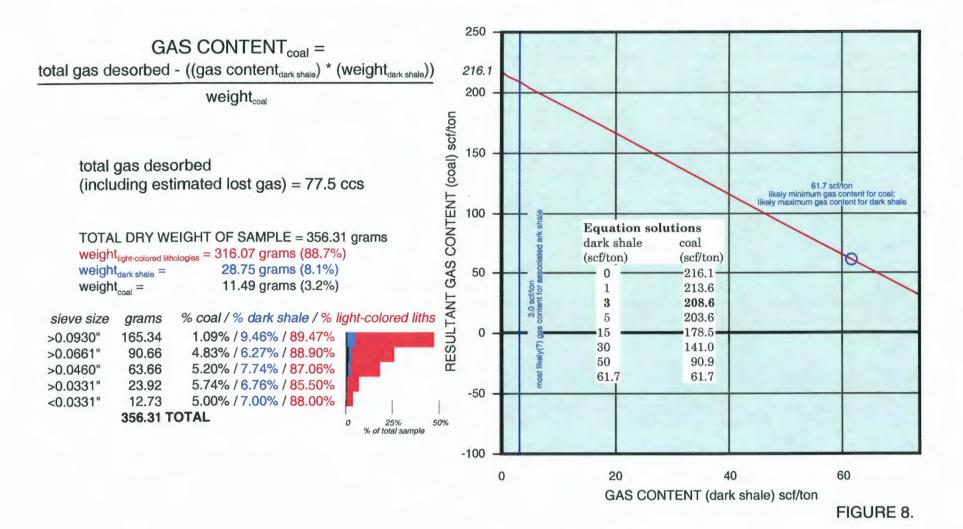


FIGURE 6.

LITHOLOGIC COMPONENT SENSITIVITY ANALYSIS for calculation of gas content of Bevier coal from 1152' to 1154'

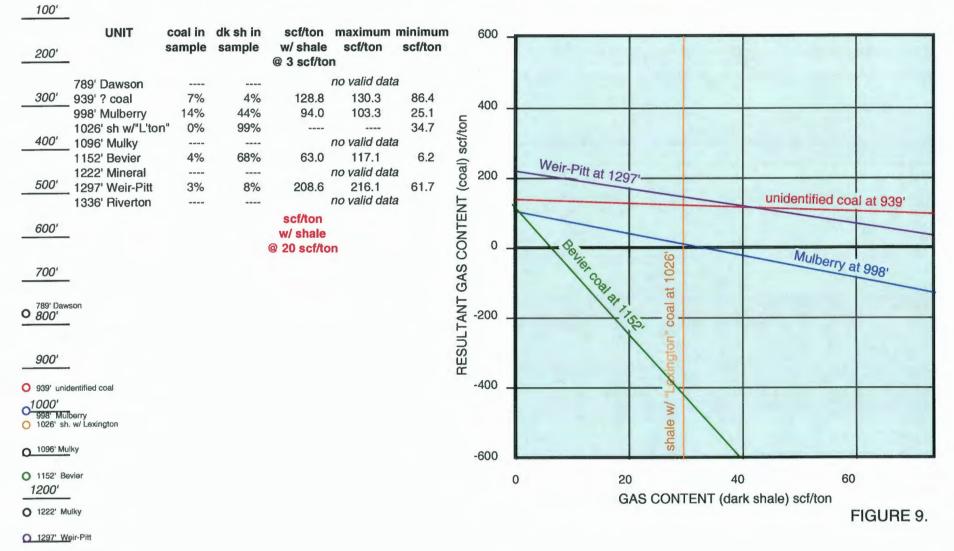


LITHOLOGIC COMPONENT SENSITIVITY ANALYSIS for calculation of gas content of Weir-Pittsburg coal from 1297' to 1298'



surface

LITHOLOGIC COMPONENT SENSITIVITY ANALYSIS for all samples



O 1336' Riverton