INSTITUTE OF GAS TECHNOLOGY IIT CENTER CHICAGO, ILLINOIS 60616

PREPARATION OF A COAL CONVERSION SYSTEMS TECHNICAL DATA BOOK

ERDA Contract No. E (49-18)-1730 Report No. 7 May 1975

Project 8964 Status Report

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ENERGY RESEARCH AND DEVELOPMENT ADMINISTRATION



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Project Status Report for

ENERGY RESEARCH AND DEVELOPMENT ADMINISTRATION

Report for

May 1975

Project Title: Preparation of a Coal Conversion Systems Technical Data Book

ERDA Contract No. E(49-18)-1730

I. Project Objective

The objective of this work is to provide a single, comprehensive source of data on coal conversion systems. This compilation shall be entitled <u>The Coal Conversion Systems Technical Data Book</u> and shall provide up-to-date data and information for the research, development, design, engineering, and construction of coal conversion processes and/or plants. Other concurrent objectives are to identify those areas where data are required and to suggest research programs that will provide the required data.

II. Summary

Liquefaction

More experimental coal liquefaction data were examined using the method described in the Project 8964 March 1975 Status Report. The "cut end point" for the liquefied residue (vacuum bottoms) of catalytic operations was determined to be 1200° to 1300°F. For noncatalytic operations, it was 2900° to 3600°F. A procedure is given to estimate yields of all intermediate cuts, given the total oil yield and hydrocarbon gas yield.

It was also found that coal liquefaction operations, in which less than 50 weight % of MAF coal fed is converted to oil, do not exhibit normal product distribution.

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Gasification

Conversion versus solids residence time charts are presented for a steam-char gasification system. In this system, the heat is supplied by some external source, such as electricity, instead of by char-oxygen reaction as in the case of steam-oxygen gasification, which was described in earlier reports.

All other assumptions for this system are the same as those for the steam-oxygen-char system.

Fluidization

A new correlation for estimating the bed-expansion ratio is proposed, which predicts about 95% of the evaluated data within $\pm 12\%$. Further work is in progress to express the correlation in pertinent dimensionless groups to facilitate reliable extrapolation beyond the operating range of the data considered.

Published correlations to predict transport disengagement height (TDH), entrainment, and elutriation are compiled.

Combustion

Low-Temperature (Fluidized-Bed)

The data-correlation approach described in last month's report was applied to an earlier set of data developed by Argonne National Laboratory (Illinois coal and limestone No. 1359 system).

Coal, Char, and Oil Shale Properties

Analytical and other related data on 50 coal deposits, which were received from Penn State are presented in a tabular compilation form deemed suitable for the Data Book.

Notice to Readers of Open File

Any comments about the material presented in this report or suggestions about the format and the content of the Data Book as well as the priorities of the needed data are most welcome. Please direct any communications to Mr. Bipin Almaula of ERDA (202/634-6643) or to Dr. Al Talwalkar of the Institute of Gas Technology (312/225-9600, Ext. 869).

III. Work Accomplished

A. LIQUEFACTION

1. Correlation of Coal Liquefaction Yield Structure

We have examined more experimental coal liquefaction data using the method described in the Project 8964 March 1975 Status Report. The "cut end point" for the liquefied residue (vacuum bottoms) of catalytic operations was determined to be 1200° to 1300°F. For noncatalytic operations, it was 2900° to 3600°F. By assigning a nominal cut end point temperature for vacuum bottoms of 1250°F for catalytic operations at 3250°F for noncatalytic operations, the point for total oil yield, which is often the only figure reported, can be plotted on a graph similar to Figure 1 or 2. If, in addition, one other yield temperature point is available, an estimate is then available of the complete yield structure for the operation.

We also found that coal liquefaction operations in which less than 50 wt % of MAF coal fed was converted to oil do not exhibit normal product distributions.

Experimental data for the H-COAL operation,^{2,3} the Synthoil Process,¹ and the SRC Process⁴ show normal product distribution. Figure 1 shows experimental data for several H-COAL operations and for the Synthoil operation using various types of catalysts. Figure 2 shows autoclave data for the SRC operation and for a noncatalytic run made in the H-COAL reactor. The data were plotted as follows:

First, the cumulative yields of products up to but not including vacuum bottoms were plotted. Then, a line was drawn through the points and the added yield of vacuum bottoms was plotted on that line. Thus, the "cut end point" for the residue (vacuum bottoms) was determined.

It should be noted that the vacuum bottoms cut end point is not an actual temperature corrected to atmospheric pressure at which all the vacuum bottoms would be distilled off. It is merely a way of adding the residue yield data to the product distribution curve determined from yields of lighter fractions. Residue oils for catalytic operations all show nominal cut end points of 1200° to 1300°F. Considering experimental difficulties, this agreement is quite good. Residue oil for noncatalytic operations generally has a nominal end point of 2900° to 3600°F, a considerably wider spread. This is probably due in part to the considerable experimental difficulties associated with quanti-





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Figure 1. CATALYTIC COAL LIQUEFACTION PROCESSES



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Figure 2. NONCATALYTIC COAL LIQUEFACTION PROCESSES

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tative separation of the solvent in batch tests. Data for batch autoclave tests in which total oil yield (including vacuum bottoms) were less than 50 wt % of MAF coal fed are not shown because operations with less than 50 wt % total oil yield do not exhibit normally distributed product slates.

Example*

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Correlation of experimental data has shown that a particular catalytic process operating on a specific coal gives a yield structure with mean temperature $\mu = 800^{\circ}$ F and a spread temperature $c = 700^{\circ}$ F. Determine the yield of 1000°F + residue that can be expected.

The nominal cut end point temperature for the residue cut for a catalytic operation is 1250°F. Therefore,

$$X_{1} = \frac{1000 - 800}{700} = 0.2857$$

$$X_{2} = \frac{1250 - 800}{700} = 0.6429$$

$$N(X_{1}) = N(0.2857) = 0.6124^{\dagger}$$

$$N(X_{2}) = N(0.6429) = 0.7399^{\dagger}$$

$$N(X_{2}) - N(X_{1}) = 0.1275$$

The yield of residual oil (vacuum bottoms) boiling above 1000°F would be approximately 12.75% by weight of the MAF coal fed.

Note that the total oil yield, about 74% for this case, is <u>not</u> equal to 100% minus the percentage of unconverted MAF coal. Some of the coal reacts to form H₂O and carbon oxides, which are not included in the oil yield.

- 2. References Cited
- Akhtar, S. et al., "The Synthoil Process Material Balance and Thermal Efficiency." <u>Paper No. 35B</u> presented at 67th Annual A.I.Ch.E. Meeting, December 1-5, 1974, Washington, D.C.
- Hydrocarbon Research, "Liquefaction of Kaiporowits Coal," <u>EPRI</u> <u>123-2</u>, Palo Alto, Calif.: Electric Power Research Institute, October 1974.

Nomenclature is as given in the Project 8964 March 1975 Status Report.

Values of N corresponding to X_1 and X_2 were determined from a table of normal distribution.

- Hydrocarbon Research, "Project H-COAL Report on Process Development," <u>OCR R& D Rep.</u> No. <u>26</u>. Washington, D. C.: Office of Coal Research, n. d.
- Wright, C. H. <u>et al.</u>, "Development of a Process for Producing an Ashless, Low-Sulfur Fuel From Coal," Vol II, Part 1 - Autoclave Experiments, <u>OCR R&D Rep.</u> No. <u>53</u>, Int. Rep. No. <u>6</u>. Washington, D. C.: Office of Coal Research, n. d.

B. GASIFICATION

1. Steam-Char Gasification in Fluidized Bed

The design charts presented up to now have been for the steam-oxygenchar gasification system in which the heat requirement for the steam-char reaction was supplied within the system by combustion of a part of the feed char with oxygen. In this monthly report, we begin presenting information on a different system - the gasification of char with steam only. Since there is no oxygen feed into this system, all the feed carbon converted is by the steam-char reaction. To operate at a constant temperature, such a system requires an external source of heat. The amount of heat required to maintain a given temperature is shown by lines of constant heat inputs in Figures 3 to 6.

Besides the exceptions just note \hat{a} , the assumption and the basis for the present system are the same as those in the steam-oxygen-char system. The charts presented here are also for the base case conditions - fluidized-bed model with both gases and solids in backmixed flow and the base carbon conversion fraction in the feed char, $X_0 = 0$. Additionally, the same correction charts for adjustments for variations from the base case can be applied to the present system.

For a given gasification temperature and 70-atm pressure, Figures 3 to 6 present curves that give the residence time required to achieve a specified feed carbon conversion at different steam feeds to the gasifier. The figures also have the constant heat input lines that show the amount of heat input required by the gasifier in order to maintain the specified operating temperature. In contrast to the charts for the steam-oxygen-char system, the time required to achieve specified feed carbon conversion is relatively higher because none of the feed carbon is being combusted with oxygen.

2. Erratum

In the Project 8964 April Status Report, the following correction is to be noted. In the calculation of C_b (p. 9), the first quantity should be 0.078 lb ash/lb raw coal instead of 0.78.

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C. FLUIDIZATION

1. Bed Expansion on Fluideation

During the past month, more data on the expansion of fluidized beds were collected. A complete list of the published data used to evaluate the fluidized-bed expansion correlations is shown in Table 1.

An interesting correlation in the literature, developed by Furukawa and Ohmae,⁸ assumes the principle of corresponding states and in particular states that the expansion characteristics of a fluidized bed are analogous to the thermal expansion of liquids. The relationship between the fluidized-bed expansion ratio and the superficial gas velocity is given by,

$$U^{0.5} \alpha \left[1 - \left(\frac{L_{mf}}{L_{f}}\right)^{1/3}\right]$$
(1)

for low fluidization gas velocities. Furukawa and Ohmae note that Equation 1 is valid only for the linear part of the bed expansion ratio versus the superficial gas velocity plot. Because the published data used in this investigation cover both the linear and nonlinear parts of the bed expansion plots, the comparison of these data with the Furukawa-Ohmae correlations was not attempted.

The evaluations of the Lewis <u>et al</u>.¹⁵ and the Shen-Johnstone¹⁹ correlations reported in the Project 8964 April 1975 Status Report were repeated to check their comparison with the additional data by Tarman <u>et al</u>.²⁰ and Knowlton.¹⁰ These comparisons are shown in Figure 7 and 8, and it is apparent that more than 90% of the data falls within $\pm 20\%$ of the calculated values of bed expansion ratios.

To explore the possibility of verifying whether a modified version of the Shen-Johnstone correlation will show a greater than 20% deviation, the published data were plotted in terms of L_f/L_{mf} versus $(U - U_{mf})/D_p^{0.5}$, as shown in Figure 9. The data fall into distinctly different groups, permitting the drawing of a straight line through each set of data points. It is, therefore, possible to deduce empirical correlations of the form -

$$\frac{L_{f}}{L_{mf}} = 1 + \alpha \left(\frac{U - U_{mf}}{D_{p}^{0.5}} \right)$$
(2)

for each set of data, by determining the slope of the linear plots.

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•	Investigators	Bed Diameter, inches	Fluidized Solids	Particle Diameter, inch	Particle Density, <u>lb/cu ft</u>	Fluidizing Medium	Operating Pressure, psig	Range of U/Umf	Range of L _f /L _{mf}					
	Curran and Gorin ³	l and 2	Lignite, char, dolomite, periclase	0.0028- 0.0173	51-222	N2, H2, CO2	0	1-20,5	1-2					
	Feldmann <u>et al</u> . ⁵	2	Char	0,0052	23	COz	0	112, 0	1-1,53					
	Tarman <u>et al</u> . ²⁰	2.5 5.5 11.5	Siderite	0.00269- 0.0141	245	Air, freon, N ₂	0-1000	1-10.5	1-1.49					
16	Knowlton ¹⁰	11.5	Lignite, FMC char, Illinois No, 6 siderite	0.0096- 0.0114	73.0- 244	Nz	0-1000	1-2.637	1-1.12					

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Table 1. SOURCES OF FLUIDIZED-BED EXPANSION DATA FOR COALS AND RELATED MATERIALS (at 70° to 80°F)

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Figure 7. COMPARISON OF THE LEWIS et al. CORRELATION WITH MEASURED FLUIDIZED-BED EXPANSION RATIOS



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Figure 9. PLOT OF EXPANSION RATIO VERSUS $(U - U_{mf}) / D_{p}^{0.5}$

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However, a single correlation is desirable that could adequately describe the entire data shown in Table 1. To arrive at such a correlation, a very fundamental approach was followed and the bed expansion ratio was related to some of the pertinent and varied parameters by the following equation:

$$\frac{L_{f}}{L_{mf}} - 1 = \alpha (U - U_{mf})^{a} (U_{mf})^{b} (D_{p})^{c} (D_{T})^{d} (\rho_{s})^{e} (\rho_{g})^{f}$$
(3)

The constant α and the exponents a through f can be estimated by converting Equation 3 to a linear form and into its logarithmic form and by using linear regression analysis. The linear regression analysis was carried out assuming that the relative precision of L_f / L_{mf} is constant. Using the appropriate weighing factor for the linear form of Equation 3 as determined by propagation of variance to be $\left[(L_f - L_{mf}) / L_f \right]^2$, the regression analysis of the data shown in Table 1 resulted in the following empirical correlation:

$$\frac{L_{f}}{L_{mf}} = 1 + \frac{0.748 (U - U_{mf})^{0.571} \rho_{g}^{0.089}}{U_{mf}^{0.033} \rho_{s}^{0.182} D_{p}^{0.030} D_{T}^{0.405}}$$
(4)

A comparison of the calculated fluidized-bed expansion ratios, using Equation 4, with the published data is shown in Figure 10.

It is recognized that the regression analysis used above is constrained because all the parameters in Equation 3 were forcibly fitted to the data used in this investigation. To circumvent this problem, the linear regression analysis was expanded to use the t-test with 95% confidence limit to determine the relative statistical significance of the chosen parameters; as a result the following correlation was obtained:

$$\frac{L_{f}}{L_{mf}} = 1 + \frac{0.855 (U - U_{mf})^{0.569} \rho_{g}}{U_{mf}^{0.049} \rho_{s}^{0.173} D_{T}^{0.403}}$$
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A comparison of the calculated expansion ratios using this correlation with the published data related to coal gasification is shown in Figure 11. Figures 10 and 11 appear to predict about 95% of the data within $\pm 12\%$ of the measured values. This is a considerable improvement over the correlations shown in Figures 7 and 8. Most of the data with the higher expansion ratios and greater than 12% deviation are seen to be light char materials of particle sizes less than 0.0043 inch (about 150 mesh) fluidized at U/U_{mf} values ranging from 16 to 20. Even though Curran and Gorin³ did not report any difficulty in measuring bed heights, it is conceivable that possible slugging or the inability to measure bed heights with accuracy at high superficial velocities could account for the deviations of these data. The difficulty of measuring precisely the expansion ratios of dense materials at low bed expansions is the reason for the calculated values being higher than the experimental data reported by Tarman et al., Curran and Gorin, and Knowlton.

By comparing Equations 4 and 5, a strong correlation appears to exist between U_{mf} and D_p for the data considered. Further work is in progress to modify the correlations given by Equations 4 and 5 to express the fluidizedbed expansion ratio in terms of physically pertinent dimensionless groups to facilitate reliable extrapolation beyond the range of operating conditions of the data shown in Table 1. Following this effort, the rationale will be developed for recommending specific correlations to estimate the fluidized-bed expansion characteristics.

2. Transport Disengagement Height

Fluidized beds are usually operated with a wide size distribution of solids, containing a substantial portion of fines. More fines are also created by attrition and by virtue of reacting solids, and the exiting gases conceivably exceed the terminal velocity of many of these fine particles, thereby carrying them out of the reactor. In addition to the entrainment of fines, solids are carried into the freeboard by erupting bubbles. However, as the particles lose their kinetic energy while ascending the freeboard space, some particles will return to the bed depending on the particle size and particle density. As a result, the particle loading in the escaping freeboard gas drops rapidly to a certain point beyond which it attains a constant value at which the terminal velocity of the accompanying particles are equal to or less than the velocity of the exiting gas stream.



Figure 11. COMPARISON OF THE EMPIRICAL CORRELATION GIVEN BY EQUATION 5 WITH MEASURED FLUIDIZED-BED EXPANSION RATIOS

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The freeboard height corresponding to the constant entrainment rate is known as the transport disengagement height (TDH), which determines the optimum location for gas exit ports in a fluidized bed.

The published correlations and empirical procedures to estimate TDH are summarized in Table 2. Attempts to evaluate the suitability of these correlations with the published data will be initiated during the subsequent months.

3. Entrainment and Elutriation

By definition, "entrainment" refers to the carry-over of solids in the exiting fluid stream from a fluidized bed. The particle-size distribution in the entrained-gas stream changes with freeboard height and remains constant beyond the TDH. The phenomenon of separation or classification of particles, either based on their size or particle density, during entrainment irrespective of the freeboard height is known as "elutriation."

The estimation of entrainment and the extent of accompanied elutriation are vital to the satisfactory operation of a fluidized-bed reactor. These quantities will determine the solids inventory in the fluidized bed and the extent of off-gas cleanup, and will be the basis for the design of dust removal equipment.

The published correlations to estimate entrainment are listed in Table 3. The suitability of these correlations to the entrainment and elutriation data of materials related to coal gasification will be tested and published in the future status reports.

4. Nomenclature

- A₊ = cross-sectional area of fluidized bed, sq ft
- d_b = bed-particle diameter, ft
- d_r = particle diameter of fines, ft

 d_{fm} = maximum diameter of entrained fine particles, ft d_{p} = particle diameter, ft

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Table 2. SOME PUBLISHED CORRELATIONS TO PREDICT TRANSPORT DISENGAGEMENT HEIGHT (TDH)

Investigators	Bed Dlameter, inches	Fluidized Solide	Fluidizing Medium	Proposed Correlation	itemarke
Zens and Wall ²³	2, 3, 8, 12, 24, and 192			$TIDH = \frac{\rho_{a} d_{p}^{2}}{2R_{1} \mu_{p}} \left[V_{0} + (U_{1} - U) \cdot \ln (I - \frac{V_{0}}{U_{1} - U}) \right]^{2}$	Theoretical model
				A plot of TDH versus superficial velocity, U, with tube diameter, $\mathbf{D}_{\mathbf{r}}$, as a parameter,	Emplrical correlation
Amitin et al.	•5,67	Catalyst	Air	TDH = 19,016 $\sqrt{D_T} / \{U_1 - U\}$	
				where U _l is the terminal velocity based on the average diameter of coarser particles,	
Lova and Wen ¹⁴		••		$TDH = \frac{\rho_n \cdot d_{\mu}^2}{18\mu} \left[V_0 - (U_1 - U) \cdot \ln(1 + \frac{V_0}{U_1 - U}) \right]$	Theoretical model
Do <u>et al</u> ."	22 (t × 0, 4 In.	Ciass heads	Air	 	Theoretical model; requires the numerical integration of a differential equation.
Franiz and Juhl ⁷	1, 2, 6, 12	Cataly st, soud	H ₂ , N ₂ , Natural gas, atr. argon.	TDH « 8.4 X 10 ⁴ D _p ρ ₈ ^{0,33} μ ^{0,71} D ₁ ^{2,14} U ^m	•-
			CO	$n_1 = D_2^{(0,1)}$	
Fournol <u>et al</u> . ⁶	24	FCC catalyst	Air	Трн - <u>1000 U⁴</u> 8	
					B 75061 31 8
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Table 3. SOME PUBLISHED CORRELATIONS TO PREDICT ENTRAINMENT RATES AND ELUTRIATION RATES

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	Investigators	Bed Diameters inches	Fluidized Solide.	Fluidizing Nedium	Propaged Correlation	
	Leva ¹³	1, 32	Sand, Fischer+ Tropsch catalyst	Air	$X_f = X_{fo} e^{-t_s \log k \theta}$, $k \alpha U^{s, 2}$	R > 1 DH
	Osberg and Charlesworth ¹⁸	J	Scotchilte glass loads	Air	$X_{f} = X_{fo} e^{-2\pi N + k0} k o \frac{(1 - U_{f})^{1/2} d_{b}^{0/2}}{d_{f}^{2} L_{b}^{1/2} d_{b}^{0/2}}$	H > TDH Low files concentration
	Yogi and Auchi ¹²				$X_f = X_{fo} e^{\frac{-K A_t \theta}{W}}$	H > TIM
					Graphical correlation of	
					$\frac{\mathrm{Kd}_{I}}{\mu} \cdot \frac{\mathrm{W}_{I}}{\mathrm{W}_{I}} \operatorname{vernum}_{I} \operatorname{vernum}_{I} \operatorname{vernum}_{I} \mathcal{V}_{I} \mathcal{P}_{K} / \mu$	
	Zong and Weil ⁽³	••			Numerical calculation procedure based on a theoretical model	H < TDH and H > TDH
					Empirical approach by graphical correlation	
26	Wen and Hashinger ⁴¹	4. 2	Scotchite glass beads, bituminuus cual powder	Atr, helium	$X_f = X_{to} e^{\frac{-KA_t\theta}{W}}$	H > T DH for binary and multiparticle system
					$\frac{K}{\rho_{g}(U-U_{1})} = 1.52 \times 10^{-5} \left[\frac{(U-U_{1})^{2}}{8 \sigma_{f}} \right]^{0.5} + \left[\frac{d_{f}U_{1}\rho_{g}}{\mu} \right]^{0.75} + \left[\frac{\rho_{g}}{\rho_{g}} \right]^{0.75} + \left[\rho_$	
					$K_{X_{f} > 0, 25} = K_{X_{f}} = (0, 25) (\frac{N_{f}}{0, 25})$	
	Andrews	1,8, 1,2	Catalyst	Alr, N ₂	$F(d_f) = -\frac{\lambda \Lambda_f \rho_B \beta}{6} (2g)^{1/2} m_p g \int_{H}^{\infty} \sqrt{x} e^{-\beta m_p g x} dx$	Theoretical model
					$F = \Sigma X_{f} F(d_{f})$	
	Fridland and Skoblo "	••	Natural and synthetic micro- spherical alumi- nosilicate catalyst	Air	$\frac{F}{\Lambda_{t} U \rho_{g}} (10^{2}) = \frac{\Lambda U^{1} X_{f}^{0.5} L_{f}^{p} m_{v}^{4}}{d_{xv}^{1.5} \Pi^{0} f}$	li ≤ TDif and Il ≥ TDii
					$d_{au} = \Sigma X_{a} d_{f}$	
					m, is a correction factor and can be determined by graphical correlation: values of A, n, and p are tabulated by the investigators.	
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$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Investigators	Bed Diameter, inches	Fiuldized Sullds	Hadring Median	Proposed Correlation	Remarks
Lewis <u>gl_al</u> . ¹⁴ <u>3</u> /4, 1-1/4, Glass, polystyrene, Air 2, 3, 5-3/4 from, cracking catalyst $ \frac{Y}{A_1 U} \neq C \exp \left[(-b_0 + A \text{ It } U) / U^4 \right], \qquad \qquad$	Kazumov und Sarionova ²⁸	7, 5	Microsphezes of natural clay catalysi	Air	$\frac{F}{A_{t} \cup \rho_{g}} \approx 5.95 \left[F_{g} + \frac{\rho_{g}}{\rho_{g}} + M_{f} \right]^{1.445}$ $F_{g} = U^{f}/g_{g}$	H > TDH
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Lowis <u>et al</u> , ¹⁶	3/4, 1-1/4, 2, 3, 5-3/4	Glass, polystyrene, iron, cracking catalyst	ALT.	$\frac{F}{A_1 U} = C \exp\left[(-b_0 + A H U)/U^4\right]$	H < TOH and H > TOH for single particle size solids
$\begin{array}{cccc} \text{Horeslan and} & 3 & \text{Glass beads} & \text{Air} & \frac{X_{f}}{X_{f_{0}}} & -b e^{-ik_{1}} A + (1-b) e^{-k_{2}} B & \text{Horesland} \\ & & & & & & & & & & & & & & & & & & $					b_ = 1,26 X 10 ⁴ j _a d ² C and A are constants and depend upon the column diameter and fluidized autida,	
Kuni and Levenspiel"Constants k_1, k_2, and is are tabulated for various gas velocities and particle-size distributionsKuni and Levenspiel" $F + F_0 e^{-a+it}$ and $a \pm \frac{K^a}{U_a}$ Theorem It of The It of The It of The to aggiomerate and dispersed phase moving down, attLeve and Wen ¹⁴ $F + F_0 e^{-a+it}$ and $a \pm \frac{K^a}{U_a}$ Theorem U_a Leve and Wen ¹⁴ $F + F_0 e^{-a+it}$ and $a \pm \frac{K^a}{U_a}$ Theorem U_a Leve and Wen ¹⁴ $F + F_0 e^{-a+it}$ and $a \pm \frac{K^a}{U_a}$ Theorem U_a Leve and Wen ¹⁴ $F + F_0 e^{-a+it}$ and $a \pm \frac{K^a}{U_a}$ Theorem U_a Leve and Wen ¹⁴ $F + F_0 e^{-a+it}$ and $a \pm \frac{K^a}{U_a}$ Theorem U_a Leve and Wen ¹⁴ Theorem U_a Leve and Wen ¹⁴ Merrick andMerrick andMerrick andMerrick andMerrick andMerrick andMerrick andMerrick andMerrick and </td <td>Haneslan and Rankell[®]</td> <td>3</td> <td>Glass beads</td> <td>Air</td> <td>$\frac{X_{l}}{X_{fu}} \rightarrow b e^{-k_{1}\theta} + (1-b) e^{-k_{1}\theta}$</td> <td>H > TDH</td>	Haneslan and Rankell [®]	3	Glass beads	Air	$\frac{X_{l}}{X_{fu}} \rightarrow b e^{-k_{1}\theta} + (1-b) e^{-k_{1}\theta}$	H > TDH
Kuni and Levenspiel ¹¹ FF $F_0 e^{-aH}$ and $a = \frac{K^a}{U_a}$ Theorem U_a where $F_0 = rate of subids projected from bed surface, 1b/sq fireHTheoremH > TheH > TheH > TheoremH > Theorem$					Cunstants k., k., and bare tabulated for various gas velocities and particle-size distributions	
$F_{n} = rate of sulide projected from bed surface, 16/sq fire K0 = rate of sulide projected from bed surface, 16/sq fire K0 = rate of sulide projected from bed surface, 16/sq fire K0 = rate of sulide projected from signomerates moving up to aggiomerates and dispersed phase moving down, s-1 U2 = velocity of aggiomerates moving up, fr/s Leva and Wen14 Grightical correlation of F/A1 J versus \left(\frac{U - U_{L_1}}{U_1}\right)^1 U_{mf}^{-4} based on thetheoretical model and data of single-size glass spheros.Merrich andHighley17J6 x 36Coal and ash\frac{K}{U_{Py}} = A + 130 \exp\left[-10.4 \left(\frac{U_{L_1}}{U}\right)^{0.5} \left(\frac{U_{mf}}{U - U_{mf}}\right)^{0.55}\right]A is constantA is constantA = 0,0001 for 36 x 36-inch combustor unit with 13-ft freeboardA = 0,0015 for 48 x 23-inch BCURA with with 6-ft freeboard$	Kunil and Levenspiel ¹¹		`		$F = F_0 e^{-aH}$ and $a = \frac{K^a}{V_1}$.	Theoretical model H < TDH and H > TDH
Leve and Wen ¹¹ Leve and Wen ¹¹ $U_2 = velocity of aggloinerates moving up, (i/s) U_2 = velocity of aggloinerates moving up, (i/s) Grightical correlation of F/A1 J versus \left(\frac{U - U_{L_1}}{U_1}\right)^1 U_{mf}^{-1} based on thetheoretical model and data of single-size glass spheros.Merrich and J6 x 36 Cosl and ash\frac{K}{U\rho_y} = X + 130 \exp\left[-10.4 \left(\frac{U_{L_1}}{U}\right)^{CS} \left(\frac{U_{mf}}{U - U_{mf}}\right)^{0.CS}\right]A is constantA = 0.0001$ for 36 x 36-inch combustor unit with 13-fit freeboard A = 0.0001 for 36 x 23-inch DCURA will with 6-fit freeboard					$F_{\rm p} = rate of sulids projected from bed surface, 1b/sq fi-s K^{\rm B} = rate cuefficient for transfer from agglomerates moving upto reclamerate the surface from agglomerates moving up$	
Leve and Wen ¹⁴ Graphical correlation of F/A ₁ J versus $\left(-\frac{U-U_{L-}}{U_{L-}}\right)^{1} U_{mf}^{-4}$ beacd on the Unit theoretical model and data of single-size glass spheros. Merrick and 36 x 36 Coal and ash Highley ¹⁷ = A + 140 exp $\left[-10.4 \left(-\frac{U_{L-}}{U}\right)^{0.5} \left(-\frac{U_{mf}}{U-U_{mf}}\right)^{0.5}\right]$ A is constant A is constant A = 0,0001 for 36 x 36-inch combinator unit with 13-ft freeboard A = 0,0015 for 48 x 23-inch BCURA with with 6-ft freeboard					U apprometates shifting ersent phase moving down, s" U, = velocity of apploinerates moving up, fi/s	
theoretical model and data of single-size glass spheros. Merrich and 36 x 36 Coal and ash $\frac{K}{U\rho_y} = X + \frac{1}{30} \exp\left[-10.4 \left(\frac{U_L}{U}\right)^{C_3} \left(\frac{U_{mil}}{U-U_{mil}}\right)^{0.65}\right]$ A is constant A = 0.0001 for 36 x 36-inch combustor unit with 13-ft freeboard A = 0.0015 for 48 x 23-inch BCURA with with 6-ft freeboard	Leva and Wen ¹⁴				Graphical correlation of $F/A_{\rm p}$ J. versus $\left(-\frac{U-1}{U_{\rm p}}\right)^{1} U_{\rm mf}^{-1}$ based on the	
Merrich and 36 x 36 Goal and xah Highley 17 $\frac{K}{U_{py}} = A + 130 \exp \left[-10.4 \left(\frac{U_{t-1}}{U} \right)^{0.5} \left(\frac{U_{mif}}{U - U_{mif}} \right)^{0.75} \right]$ V_{py} A is constant A = 0.0001 for 36 x 36-inch combinator unit with 13-fit freeboard A = 0.0001 for 36 x 36-inch DCUKA will with 6-fit freeboard					theoretical model and data of single-size glass spheros,	
A is constant $A = 0,0001 \text{ for } 16 \times 36 \text{-Inch combustor unit with 1 J-ft freeboard}$ $A = 0,0015 \text{ for } 48 \times 24 \text{-inch DCURA with } with 6 - (1 freeboard)$	Merrick and Highley 17	36 x 36	Coal and ash		$\frac{K}{U\rho_{y}} = A + i H \exp\left\{-10.4 \left(\frac{U_{t}}{U}\right)^{1.5} \left(\frac{U_{tinf}}{U-U_{nif}}\right)^{0.15}\right\}$	
$\Lambda = 0,0001 \text{ for 36 x 36-inch combustor unit with 13-ft freeboard}$ $\Lambda = 0,0015 \text{ for 48 x 24-inch DCURA with with 6-ft freeboard}$					A is constant	
A = 0,0015 for 4H x 21-inch BCURA with 6-ft freeboard					$\Lambda = 0.0001$ for 36 x 36-inch combinator unit with 1.3-ft freeboard	
					A = 0,0015 for 48 x 21-inch BCURA will with 6-ft freeboard	

Table 3, Cont. SOME PUBLISHED CORRELATIONS TO PREDICT ENTRAINMENT RATES AND ELUTRIATION RATES

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particle diameter of a sieved fraction, ft D_{f} $\frac{1}{\Sigma(X/D_f)}$ average particle diameter, ft = Dp tube diameter, ft DT ~ E total energy of fluidized system, ft-lb = free area of gas distributing grid (fraction of total), sq ft/sq ft f Ξ F total entrainment rate, lb/s = $F(d_f) =$ entrainment rate of particles of diameter d_f , lb/s acceleration of gravity, ft/s^2 g Ξ H distance between bed surface and gas outlet, ft Ξ rate of solids carried up by wake of bubbles per unit time per J = unit surface area of bed, 1b/sq ft-s elutriation rate constant, s⁻¹ k Ξ ĸ elutriation rate constant, lb/sq ft-s = L_{mî} = height of minimum fluidized bed, ft Ξ height of fluidized bed, ft Lf settled bed height, ft Ls = mp mass of entraining particle, 1b = Ν = total number of particles in fluidized system U = superficial gas velocity, ft/s Umf minimum fluidization velocity, ft/s = Ut terminal velocity, ft/s \simeq vo particle velocity leaving bed surface, ft/s = W weight of solids in fluidized bed, 1b Ξ Х weight fraction of sieved particles = Xf weight fraction of fines in the fluidized bed at any time, θ = X_{fo} weight fraction of fines in the fluidized bed at $\theta = 0$ = weight fraction of ith size fraction with respect to total weight $\mathbf{X}_{\mathbf{i}}$ = of fines density of fluidizing gas, lb/CF ۶g particle density of fluidizing solids, Ib/CF ρ_s =

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- $\rho_{\rm B}$ = fluidized-bed density, 1b/CF
- μ = viscosity of fluidizing gas, lb/ft-s
- β = entrainment rate constant = N/E
- λ = constant coefficient of conductance
- θ = time, s

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D. COMBUSTION

In the Project 8964 April 1975 Status Report, it was proposed that sulfur recovery data in FBC experiments be organized on the basis of a comparatively simple reactor model. The equations presented were applied to Argonne National Laboratory (ANL) data for a Pittsburgh coal-Tymochtee Dolomite system, and a completely backmixed reactor model appeared to

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give a reasonable characterization of the reported results. Therefore, the same approach was applied to an earlier but not quite as extensive set of data developed by ANL.¹ The system was of Illinois coal-limestone No.1359 with the material properties and operating conditions shown in Tables 4 and 5. Unlike the previously analyzed data set, this one does not report directly measured bed compositions in most runs. Therefore, for this analysis, bed compositions have been calculated using material balance considerations as described in the April Report, with the assumption that there was no significant preferential elutriation.

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	Scam 6, Mine 10 Peabody Coal Co. Christian Co., Ill.	Lime 	estone 1359			
Analysis		-				
Moisture	10.12	Ca	37.9			
Ash	10.85	COz	42.2			
Sulfur	3.72	Ash	56.2			
Carbon	61.54					
Hydrogen	4.47					
Mean Particle Size, mm	490	550-1000				

Table 4. MATERIAL PROPERTIES

As has been reported in many FBC studies at atmospheric pressure, the reaction rate appears to go through a maximum at about 1500°F, an effect not observed at higher (8-atm) pressures. Also, for this limestone, at these conditions, only a fraction of the CaO appears to be effective. Simultaneous allowance for these peculiarities makes data correlation difficult. Koppel¹ in his analysis of these (or similar) data implies that only 50% of the CaO is effective. With this value, the reaction rate constants, k_0 , are plotted as a function of temperature. They are calculated from the ANL data and defined by -

Jonke, A. A. et al., <u>Reduction of Atmospheric Pollution by the Application</u> of Fluidized-Bed Combustion, Annual Report July 1970-June 1971, <u>ANL ES-CEN-1004</u>. Argonne National Laboratory, U.S. Environmental Protection Agency, 1971.

Table 5. OPERATING RANGES*

Temperature, °F	1400-1600
Pressure, atm	1
Dry Flue Gas/Coal, SCF/lb	105-130
Bed Diameter, ft	0.5
Bed Weight, 1b	14-45
Ca/S Mole Ratio	1.0-5.5
Ca Conversion (Estimated)	0.16-0.38
Ca Bed Weight Fraction (Estimated)	0.22-0.45
Sulfur Recovery Fraction	0.38-0.96

Not all the ANL tabulated results have been used in this analysis in view of the caveat concerning tabulation errors.

Reaction rate (moles of S/min) = $k_0 X_{SO_2} (X_{Ca, m} - X_{Ca}) Ca_{bed} W_{bed}$ where

 X_{SO_2} = mole fraction of SO₂ in actual flue gas

 $C_{abed} W_{bed}$ = weight of calcium in the bed, lb

 X_{C_2} = fraction of calcium sulfated

 $(X_{Ca})_m$ = maximum possible fraction of calcium sulfated.

Figure 12 shows the results. Each point is shown as a range corresponding (quite arbitrarily) to $\pm 5\%$ of the fraction of sulfur recovered as the sulfate. It is seen that the maximum rate near 1500° F is not very reproducible. There are, however, other systems that have indicated this maximum so its existence cannot be rejected. But one can entertain the thought that operation at this peak condition is unstable and not always achievable. Because of the uncertainty in the peak value, the curve shown was drawn on the basis of engineering conservatism. The observed value at 1500° F is 3.7 times the value on the curve, but, if one substitutes the lower value (at the conditions of the run), the fraction of sulfur recovered would be estimated as 0.81 rather than the 0.92 observed. Other data in this set imply such discrepancies are encountered. A similar calculation for the high value at 1550° would result in an estimation of 0.85 rather than the 0.92 sulfur recovery observed.

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Figure 12. RATE CONSTANTS FOR THE REACTION OF SULFUR IN ILLINOIS COAL WITH LIMESTONE 1359

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E. COAL, CHAR, AND OIL SHALE PROPERTIES

1. Specific Heat of Coal, Char, and Ash

Work on the specific heat of coals, chars, and related materials was continued. Comparisons of Kirov's correlation (the Project 8964 March 1975 Status Report) with experimental data at elevated temperatures (300° and 600° C) were prepared in graphical form. Then we learned that apparent specific heat data on 20 American coals have just become available from P. L. Walker at Pennsylvania State University, via a paper to be presented at the ACS Fuels Division Meeting at Chicago in August. Next month, if the paper is received in time, we will present correlations incorporating these data as well as previous data.

2. Penn State Data

Data on 50 coal samples from Penn State have been tabulated (Table 6). About one-third of the samples are from deposits too small to be included in our table of large-size deposits (Table 4, Project 8964 April 1975 Status Report). However, they are being retained in the data table for the time being, at least.

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Table 6. PROPERTIES OF SEL

		MINE (OR OTHER ICANTIFICATION)	AANX	PROXIMATE ANALYSIS, WI -												
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AZ	Navajo. Rec	(Blue Var				1.4. 4 7	47.47	72.75	3. 64	1.20	و ن ر ن	2.71	· • • • •	4. 25	37.7	4
co	Weld, Laramie	E. de	NUSA L.L.L	10.00	÷	50.24	v		5. 20	2د ـ ډ	0.04	0. s7	* . * .	7	a* ,*	÷
CC	Roult. Wacke	Esta	1-125 B		· · · · ·	×	÷. ÷0	72.00	÷. 75	1.0.	v. 62	0. *~	÷. 40	: *. *	To	•
co	Patkin, Coal Barin B	Duton Commis	arca		;	-1.1:	10.14	s∿.≟0	4.4,	1.75	6.61	0. 25	:6		01. C	÷.
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11_	Callatin, Illinuis Nu. 5	Earle No.		1				5 P. 44	÷.0:		6.9	9.05	104		10.3	.
щ	Williamson, Davis	Will Summer	N. 21.		••••	44,62	· · ·		÷. ÷	. 5 4	0+	4.7-	. 4. 8-	n. :	D.L.	
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8¥	Letcher, Elkhorn No. 5						•••	c7. hi	4 . •	•• •	0		·•. ·-	4. 44		4.
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КУ	Muhlenburg, Ky. Na. 11	Surfair Strip	71.85	1 4.11		54. 4 2	r0	74,44	5.00	2.20	6,21	4.14	8,40	7.27	54. T	-
мс	Benry, Tebe	Stati ir Strip	h 55	1.7	·	41, - 1	16.64	e4. (4	4, 57	1.67	6,01	÷. ~~	10,04	r.41	et, t	
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-4	21.1	442 -	ند و ه	. v. + 4	Q. 1	e. or	v. + a	+ 2. 24		17. 25	13.4.4					1
. ± r	·. 4	J. +	é, 1	44.5		U. 1-	11	57 7		11.477	1				••	internet.
1 . T	4	4. 4	U. *	1. - 1	4. 24	6, 14	2.00	57.7-		7 .	11.755	57.4				1-24 - 44 - 4
÷	÷. 7	4.1	6.1	- v, 45	4. 45	0.04	1.00	44. 52		1 1. 714	16 1.54					Pol Cal
4 , 0	16.2	1. *	Ú. 1	5. 74	4. 71	C_ 05	1.04	56.27		1.5.100	11. 444		•			1-51 (-,
:2,7	• . •	4,4	4.1	1. 11	0	0.04	1. A.	47.44		10,177		50 a		1	~~	2.31 C-1-
3.5	· r.	1.0	0	6. 44	Ю. V.S	6.6-	0.50	4. 6		7 292	th hut					Pacca.
نحي ہ	4. 4	1. U	n , n	22.49	0.65	n e.	1. <u>-</u> 11				1. 000	61 a				2750 0+24+
4.4	.	÷. 1	C. 1	ಲ್ಲಿಂದ	0.55	a (B	6 2 3			15 051		21. 7 21. 2	ų, -			2861-45
11.0	۰. ۲	1.4	0,4	C. 48	0.6	0.00	0.10	-4.04			10 4-1		••			P36.6 - 2-1
4.4	. -	5. 54	0.7	0.77	1.41	0.20	6 7.	44 21		141.541	1		0 			1500
4		54	0. +	U, 64	4.25	0.02	1.17	30.27		1-1-214		37 4 Án Ú				2°SC C - 27
3.4	۰ <u>،</u> ۲	1. 6	V, ó	4.35	2. 53	0.0.	2.05	54.71		14.144	11.074	55,2	-, U			3154 (
				1			1	1			1			1	{	

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Table ó, Cont. PROPERTIES OF

		MANE (OR OTHER SDENTIFICATION)	RANK	PR0	LIMATE A	NALYSIS. DRY BASK	m 1	ULTIMATE ANALYSIS, #1 "- DR": BASIS							
STATE	COUNTY AND SEAM			MOISTURE	VOLATILE	FIXED CARBON	ASH	1 5 7 7	WYONODEM	NITROCEN	CHLORINE	SULFUR	4SH	OLYGEN I DIFFERENCE I	311M12114
он	Nahoning, No. 5. Lower Bittanning	Five Points	3445	5.18	36, 21	15.50	10.19	67.10		2.14	0.13	0.90	10.15		4.0
сн	Beimont, Chio No. 12	Ne. 10	hvab	3.57	\$1.97	38, 77		56.42	4 21	1.24	0.0é	2.05		6.74	
ск	Le Flore, Hart Shorne	Howe	anvb	0. ++	20.05	00 97	10.1	70.04		0 1 5	0.01	1.05	10.0	- 44	66.5
PA	Schuyladi, Buck Mountain	Renninger	27	4.51	5.77	00.29	13.94	7a. 1a	2.25	0.00	00	0. 55	10.54	4.44	4n 1
PA	Somerret, Lower Kattanning	Bire No. 5	ivb	0, 51	10, 10	71.34	10. e.	78.92	1.17	1.05	0	3.10	16	2.03	46.4
PA	Cumbria, Lower Freeport	Pennsylvania No. 5	1vb	1.14	204	co. 40	14,94	74.27	4. 13	1.11	C. 11	4.04	12.92	1.22	05.0
PA	Cambria, Lower Kutanning	Cambria No. 33	1vb	0.04	18.64	74, 37	7.01	-2 -00	4.40	0.00	0.04	04	7.01	, ó-∔	54.0
PA	Washingiyu, Pittsburgh	Math ex	hváb	1.62	·r. 76		. 8, 70	70.00	4, 71	2.00	0.04		a. 70	4.44	aC. 2
TE	Campbell, Peewee	Long Pit Mining Co.	hvAb	2.77	30. CO	20. 1 l	7.23	TT. 24	5, 24	1. **	0.14	4. 29	7.4 3	0. 04	T3. 60
XL	Harrison, Lignite .	Darco		24.52	42,25	47.44	10.24	60.00	4.45	0. 30	0.0>	6.07	10. <i>2e</i>	:7, 55	
۷۸.	Wise, Jawbone or Shannon	Dal Strip Job	hvab	1.10	94. ND	52.70	18, 54	71.69	4.64	2.03	0.02	6. ĴB	15, 24	ا فرق	
VA	Wite, Imboden	Prescott No. 2	hvAb	1.10	53.00	54.1:	7, 23	79.95	5.05	د تا ، ت	C. 01	0. 54	7, 21	5. 01	
VA	Dickenson, Lower Banner	Camp Branch No. 1	hvAb	0.45	32. re	40.0J	÷. 14	00,1C	4.97	1,75	0.04	C, 71	0.51	3. 93	57.2
UT	Emery, Lower Sunnyside	Geneva	(hvBb	÷. 20	19. 55	55, 30	5, 15	77,15	5. +1	1.34	0.02	0.85	5. 1 S	10.0	8J. 3
UT	Carbon, Scam A	King		5.00	41.75	48.0l	9.46	71, 99	5. 55	1.51	0. D-i	1.70	9.41	e. 77	65.2
UT	Carbon, Hiswatha	Plateau	hvBb	0. 12	·• • •	÷4.60	16.07	00.55	5, 15	1.51	U. D5	C. 01	10.07	10.60	79.9
A.Y	Lewis, Big D	Centralia	*ubB	16.21	19,12	18. 30	22,48	50,27	4.49	1.04	0.02	1.01	22,45	14, 84	70.9
W.C.	Nicholas, Tinga	Tiega No. 1	hudb	1.94	30,71	زە د	15, 30	71 13	4.26	0.07	C. 0o	1, 19	j\$, jö	7.14	00.1
WV	Mcdowell, Pocahontas 3	No. 14		1.82	21, 54	72.07	₽ز_0	04. b	4, 02	1.05	C. C1	ü. 52	0. 34	44.5	70.5
WY	Sheridan, Monarch	Big Hors	-ubB	1 7. 60	40,91	52. 0 0	c. 04	67.99	4,93	1.33	0.04	0.04	6. 9¢	17.00	85,7
WY	Lincoln. Adamile 1	Eikol	sub.A	10.17	43. 3 0	53, 57	2.03	72,96	5. 0 0	1.00	0.04	0.64	وه.د	10.75	76.3
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TTED COAL SAMPLES

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METT		FORMS OF SULFUR, %			RANK DATA			SISCELLANEOUS			ASH-SOFTENING TEMP.ºF					
EBAL ANALYSIS, VAL					DRY BASES			.:							1	
SEMPENSINIE	MCRINITE.	EXIMITE	RESIMITE	NEFLECTANCE OF VITNINITE, S	PYRITK.	SULFATE	ORDANIC	FIXEO CARBON, S. My, Min free	EQUILIBRIUM MOISTURE, *	CALORIFIC VALUE Biu/Ib, maiol, mm fr	CALORIFIC VALUE BIV / Ib, dy buils	HARDOROVE Gaingasility Ihoek	FREE-SMELLING INDEX	DXIDIZIHG &TMOS	REDUCING ATMOS	CODE NO.
÷								1			1					
8	16, 8	21.2	2,2	0, 74	0141	0.0	0.07	56.95		1-2. 38-2	12,157	40,7	1.5			PSCC-296
b .o	9.0	7, 3	0. 2	0,7	1.21	0.01	9,83	57.17		14, 346	10.243	50.4	1.5			PSCC-306
1.0	3. 4	0. C	0.0	1, 43	0,42	C, 00	0.63	77.81			13, 541	115_6				PSOC-142
1.7	p, p	0,0	0. C	4, 17	0.02	6.00	0.53	94. 76			12.603	28.9				FSCC-60
3. 9	o. 4	0. O	0.0	1, 50	2,03	0,16	0,91	61.56			13. 745			1		PSCC-113
3.5	د, ۲	ο, ο	ə, o	1.47	J. 65	0,20	1.16	75.60			13,233					PSCC-116
1.8	હું ચ	0, ŭ	e, o	1,65	0.23	0.03	0, 44	80.81			14,605	122, 5				PSCC-126
	7,9	÷. 2	0, 1	0,81	1, 29	0.01	0,65	60.57		14,577	13. 743	54.4	8.5			1-5CC-245
1.9	9,Ŭ	¥, 3	0, e	Q, 66	0.53	2.06	0,70	61.12		14,057	13, 801	49.0	3.5		`	PSCC-300
- :	11.5	.	4,77	0, 34	0.15	0.05	0.46	53.46			11.467	65.4	0,5			PSCC-135
-	14.5	e, 5	۰. ۲	0, 90	0.D4	0, JO	0.56	P5.59		15.132	12, 754	53.9	7.5			PSCC-265
	14.4 1	3. 3	0.4	1.03	0.02	0.00	e. 54	64.27		15,250	14.239	03. ú	8. C			PSCC-269
9.0	12, 2	ъ. ≟	2, 2		0.16	0.01	0, 54	65,16		15.626	14,265	67.1	5.5			PSCC-302
-4, 2	2,4	<u>і</u> . в	0.6	V. 59	C. 27	0,06	0.52	50.71		13.930	13, 757	46.5				PSCC-068
• 5. 5	9, X	3, 4	0, 5	Ų, 58	Q, 74	0.01	1.03	54.66			13.061	53.2				PSCC-236
2.1	۵,۵	9,0	1, 8	0.57	0.15	0,00	0, 48	54.15		13.326	11,895	49.0	1.0			PSOC-313
10. ¢	4,7*	1. 5	2.5	0,46	0.33	0,00	0,69	50. 59		10.163	9,952	45.6	•-			PSOC-240
÷4. 9	14, 2	5. B	ē, 0	6.89	0, 51	0.00	C. 59	64.93		14,761	12, 567	56.7				PSCC-118
4 , 6	12.0	v. s	0.0	1.42	Q, 04	0.00	0,48	77. 57			14.533	114, 2				PSCC-130
. L. E	3.2	4. 4	û, 4	0, ≼ 8	G_46	0.61	0.37	56, 49		10,357	11,762	45.6				PSCC-241
31. E	¥. 5	1. 4	0,4	0,47	0.03	0, CÚ	10.01	55.42		10,507	12,661	49.7				PSOC-248
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Advisory Committee

Letters were sent out inviting eminent people in various areas of coal conversion technology to serve on an Advisory Committee for the Data Book Project. The purpose of an advisory committee is to obtain reviews and comments on -

- Organization of the total work (Data Book Index)
- Selection of subjects to be covered in priority schedule
- Scope of work to be covered for selected subjects
- General plan for implementing the work
- Progress reports
- Final data presentations.

The list of the people to whom the letters were sent is given in Table 7. We have received eight responses so far; seven of them accepting our invitation. One declined because of lack of time to spare for serving on the committee, although he too was in favor of the idea.

IV. Patent Status

The work performed during May is not considered patentable.

V. Future Work

The data collection and evaluation work will be continued in the selected areas of coal conversion technology.

Approved W. U. Bodle

W. W. Bodle, Director Process Analysis

Signed

A, Talwalkar, Coordinator Process Data

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INSTITUTE

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TECHNOLOGY

Table 7. PERSONS INVITED TO SERVE ON THE DATA BOOK ADVISORY COMMITTEE

Dr. Richard Rosa Avco Corporation Everett Research Laboratories 2385 Revere Beach Parkway Everett, Mass. 02149

Dr. Martin Sherwin Chem Systems, Inc. 275 Hudson St. Hackensack, N. J. 07601

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Mr. Shelton Ehrlich Pope Evans & Robbins, Inc. 320 King St., Suite 503 Alexandria, Va. 22314

Dr. David Archer Westinghouse Electric Corp. Research & Development Center Beulah Road, Churchill Borough Pittsburgh, Pa. 15235

Dr. Jack Jones FMC Corporation P. O. Box 8 Princeton, N.J. 08540

Mr. Bruce SchmidSynthetic FuelPittsburg & Midway Coal Mining Co.Dravo Corp. ≤P. O. Box 199Chemical PlayDu Pont, Wash. 98327One Oliver Pl

Mr. Henry McGrath Procon Inc. 30 UOP Flaza Des Plaines, Ill. 60016

J. D. Stubbs Bechtel Incorporated Fifty Beale St. P. C. Box 3965 San Francisco, Calif. 94119

Mr. J. B. O'Hara, Manager Energy Department Ralph M. Parsons Co. 100 West Walnut St. Pasadena, Calif. 91124

Mr. C. A. Bolez Gilbert Associates, Inc. P. O. Box 1498 525 Lancaster Ave. Keading, Pa. 19603

Mr. E. J. Flavin Fluor Engineers & Constructors, Inc. 5559 Ferguson Drive Los Angeles, Calif. 90022 Mr. Roger Broeker Foster Wheeler Energy Corporation 110 S. Grange Ave. Livingston, N. J. 07039

Mr. Pete Gilman Electric Power Research Institute 3412 Hillview Ave. P. O. Box 10412 Palo Alto, Calif. 94304

Dr. George Skaperdas M. W. Kellogg Co. P. O. Box 696 Piscataway, N. J. 08854

Mr. George Curran Conoco Coal Development Co. Library, Pa. 15129

Mr. Herman Feldman Battelle Columbus Laboratories 505 King Ave. Columbus, Ohio 43201

Mr. Stan Kasper, Chief Process Engineer Synthetic Fuels Department Dravo Corp.⁴ Chemical Plans Div. One Oliver Plaza Pittsburgh, Pa. 15222

Mr. Joseph J. Williams Stone & Webster Engineering Corp. P. O. Box 2325 60 Battery March Boston, Mass. 02107

Dr. Clarence Johnson Vice President, R&D Hydrocarbon Research Incorporated 2233 Wisconsin Ave. N. W. Washington, D. C. 20007

Mr. John Igoe, President Bituminous Coal Research, Inc. 350 Hochberg Road Monroevikle, Pa. 15146

Dr. Irving Wender
U.S. Energy Research & Development Admin.
Pittsburgh Energy Research Center ⁻
4800 Forbes Ave.
Pittsburgh, Pa. 15213

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3424 SOUTH STATE STREET IIT CENTER CHICAGO, ILLINOIS 60616 AFFILIATED WITH ILLINDIS INSTITUTE OF TECHNOLOGY