

## Reference

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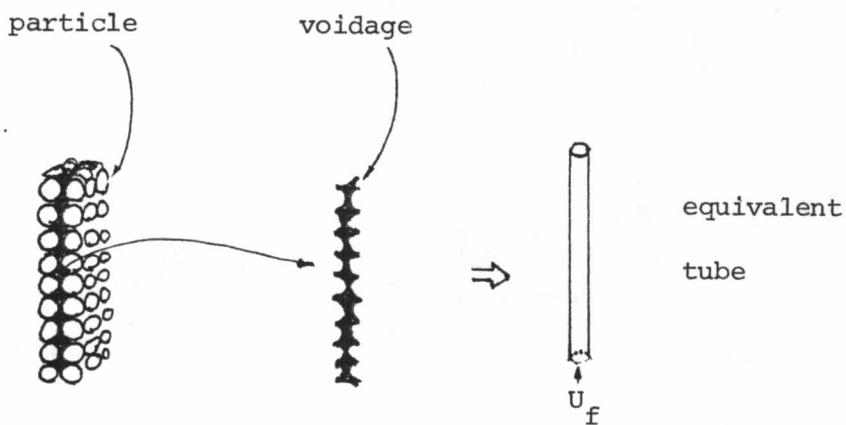
APPENDICES

## Appendix A

Minimum Fluidization Velocity ( $U_{mf}$ )

The Ergun equation derivation for predicting minimum fluidization velocity ( $U_{mf}$ ) is shown followingly.

In fixed bed, the particles of the same size are assumed. The small tubes are assumed to be the equivalent tubes for the voidages between particles. The surface area of the equivalent tube is assumed to be equal to the surface area of the particles.



Let the equivalent tube diameter be  $d_c$  and  $U_f$  is the air velocity passing through the tube.

From the Bernoulli equation

$$\frac{\Delta P}{\rho_g} + \Delta z \frac{g}{g_c} + \frac{\Delta (\frac{U_f^2}{2g_c})}{2g_c} + \sum F = w_f \dots \dots \quad (1)$$

where  $\Delta P$  = pressure drop across the tube

$\rho_g$  = density of gas

F = force

$w_f$  = work

$$\frac{\Delta P}{\rho_g} + 0 + 0 + F = 0$$

$$\frac{\Delta P}{\rho_g} = F_{friction} + F_{kinetic} \dots \dots \quad (2)$$

where  $F_{friction}$  = friction force

$F_{kinetic}$  = kinetic force

$$F_{friction} = f \frac{L}{d_c} \frac{\bar{U}_f^2}{2g_c} \dots \dots \quad (3)$$

$$F_{kinetic} = \frac{k \bar{U}_f^2}{2g_c} \dots \dots \quad (4)$$

where  $f, k$  = constant

$$\frac{\Delta P}{\rho_g} = f \frac{L}{d_c} \cdot \frac{\bar{U}_f^2}{2g_c} + \frac{k \bar{U}_f^2}{2g_c} \dots \dots \quad (5)$$

$$d_c = 2r = 2r \cdot \frac{2\pi r}{2\pi r} = \frac{4\pi r^2}{2\pi r} \cdot \frac{L}{L} \cdot \frac{N_c}{N_c} \dots \dots \quad (6)$$

where  $r$  = radius of the tube

$d_c$  = tube diameter

$N_c$  = numbers of tube

$L$  = length of tube

$$\text{hence } d_c = \frac{4 \cdot \pi r^3 L N_c}{\frac{2 \pi r L N_c}{d_c}} = 4 \frac{\text{total void volume}}{\text{total surface of particles}}$$

$$\begin{aligned} d_c &= \frac{4 \cdot N \cdot \frac{4}{3} \frac{\pi d_p^3}{8} \cdot \frac{\epsilon_m}{1-\epsilon_m}}{N \cdot \frac{4 \pi d_p^2}{4}} \\ &= \frac{2}{3} \cdot \frac{\epsilon_m}{1-\epsilon_m} d_p \quad \dots \dots \quad (7) \end{aligned}$$

where  $N$  = numbers of particles

$d_p$  = particle diameter

$\epsilon_m$  = void fraction in fixed bed

substitute Eq. (7) in Eq. (5)

$$-\frac{\Delta P}{\rho_g} = f \cdot \frac{3}{2} \frac{(1-\epsilon_m)}{\epsilon_m \frac{d_p}{d_c}} \cdot \frac{L \bar{U}_f^2}{2g_c} + \frac{k \bar{U}_f^2}{2g_c} \quad \dots \dots \quad (8)$$

$$\text{from } U_o A_t = \bar{U}_f \epsilon_m A_t \quad \dots \dots \quad (9)$$

$A_t$  = bed cross section area

$U_o$  = superficial velocity

$$K \propto \frac{L}{d_c} \quad \dots \dots \quad (10)$$

$$f = \frac{64}{Re} = \frac{64 \mu}{\rho_g U_f d_c} \quad \dots \dots \quad (11)$$

where  $Re$  = Reynolds number

substitute Eq. (9), (10) and (11) into Eq. (8)

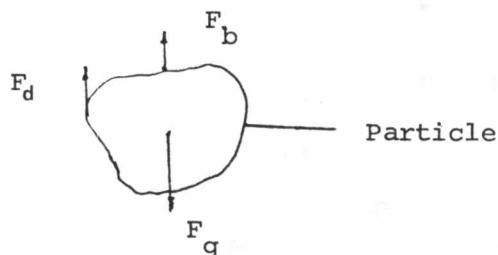
$$\left(-\frac{\Delta P}{L}\right) g_c = \frac{K_1 (1-\epsilon_m)^2 \mu U_o}{\epsilon_m^3 d_p^2} + K_2 \frac{(1-\epsilon_m) \rho_g U_o^2}{d_p \epsilon_m^3} \quad \dots \dots \quad (12)$$

Ergun found that  $K_1 = 150$ ,  $K_3 = 1.75$

$$\therefore \left(-\frac{\Delta P}{L}\right) g_c = 150 \frac{(1-\epsilon_m)^2}{\epsilon_m^3} \cdot \frac{\mu U_o}{d_p^2} + \frac{1.75 (1-\epsilon_m) \rho_g U_o^2}{d_p \epsilon_m^3} \quad \dots \dots \quad (13)$$

In a fluidized bed, a particle is floating and follows the equation

$$\sum F = 0 \quad \dots \dots \quad (14)$$



$$F_b + F_d = F_g \dots \dots \dots \quad (15)$$

where  $F_b$  = bouyancy force

$F_d$  = drag force

$F_g$  = gravitational force

$$F_g = (1 - \epsilon_{mf}) A_t L \rho_s \frac{g}{g_c} \dots \dots \dots \quad (16)$$

$$F_b = \rho_g (1 - \epsilon_{mf}) A_t L \frac{g}{g_c} \dots \dots \dots \quad (17)$$

$$F_d = \Delta P A_t \dots \dots \dots \quad (18)$$

where  $\rho_s$  = solid density

$\epsilon_{mf}$  = minimum fluidization void fraction

substitute Eq. (16), (17) and (18) in Eq. (15)

$$(1 - \epsilon_{mf}) A_t L \rho_s \frac{g}{g_c} = \rho_g (1 - \epsilon_{mf}) A_t L \frac{g}{g_c} + (-\Delta P A_t) \\ \left( \frac{-\Delta P}{L} \right) g_c = (1 - \epsilon_{mf}) (\rho_s - \rho_g) g \dots \dots \dots \quad (19)$$

From Eq. (13) and Eq. (19) and letting  $\epsilon_m = \epsilon_{mf}$ ,  $U_o = U_{mf}$

$$\text{hence } 150 \frac{(1-\epsilon_{mf})^2 \mu u_{mf}}{\epsilon_{mf}^3 (\varphi d_p)^2} + 1.75 \frac{(1-\epsilon_{mf}) \rho_g u_{mf}^2}{(\varphi d_p) \epsilon_{mf}^3} \dots \quad (20)$$

$$= (1-\epsilon_{mf}) (\rho_s - \rho_g) g$$

$$\text{or } G_a = 150 \frac{1-\epsilon_{mf}}{\varphi^2 \epsilon_{mf}^3} \frac{Re_{mf}}{d_p} + 1.75 \frac{Re_{mf}^2}{\epsilon_{mf}^3} \dots \quad (21)$$

where  $Re_{mf} = \frac{u_{mf} d_p \rho_g}{\mu}$ ; particle Reynolds number at fluidization velocity

$$Ga = g d_p^3 \rho_g \cdot (\rho_s - \rho_g) / \mu^2; \text{ Galilei number}$$

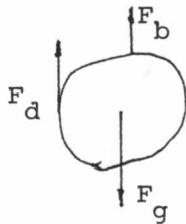
$\varphi$  = sphericity of particle

## Appendix B

Terminal Velocity of Particles ( $U_t$ )

The gas flow rate through a fluidized bed is limited on one hand by  $U_{mf}$  and on the other by entrainment of solids by the gas. This upper limit to gas flow rate is approximated by the terminal or free-fall velocity of the particles, ( $U_t$ ).

The derivation of  $U_t$  is as follow.



$F_d$  = drag force

$F_b$  = buoyancy force

$F_g$  = gravitational force

$$F = ma$$

$$\therefore a = 0$$

$$F = 0 \quad \dots\dots\dots \quad (1)$$

$$F_d = F_g - F_b \quad \dots\dots\dots \quad (2)$$

$$F_d = A_t K C_d$$

where  $A_t$  = cross sectional area of bed

$C_d$  = drag coefficient

$K$  = characteristic kinetic energy

$$K = \frac{1}{2} \rho_g u_t^2 \dots \dots \dots \quad (3)$$

$$F_g = \frac{4}{3} \frac{\pi d_p^3}{8} \rho_s \frac{g}{g_c} \dots \dots \dots \quad (4)$$

$$= \frac{\pi d_p^3}{6} \rho_s \frac{g}{g_c} \dots \dots \dots \quad (4)$$

$$F_b = \frac{\pi d_p^3}{6} \rho_g \frac{g}{g_c} \dots \dots \dots \quad (5)$$

substitute Eq. (3), (4) and (5) in Eq. (2)

$$\frac{1}{2} \rho_g u_t^2 \frac{\pi d_p^2}{4} C_d = \frac{\pi d_p^3}{6} (\rho_s - \rho_g) \frac{g}{g_c}$$

$$u_t = \frac{\frac{4}{3} d_p (\rho_s - \rho_g) \frac{g}{g_c} \frac{1}{2}}{\rho_g C_d} \dots \dots \dots \quad (6)$$

Eq. (6) is used for spherical particles

An alternate way of finding  $u_t$  for spherical particles uses analytic expressions for the drag coefficient  $C_d$ . Unfortunately,

no single simple expression can represent the experimental findings in the flow regime of interest, so for the particle Reynolds number ranges shown we have

$$C_d, \text{ spherical} = \frac{24}{Re_p} \quad \text{for } Re_p < 0.4 \quad \dots \dots \quad (7)$$

$$C_d, \text{ spherical} = \frac{10}{\frac{1}{2} Re_p} \quad \text{for } 0.4 < Re_p < 500 \quad \dots \dots \quad (8)$$

$$C_d, \text{ spherical} = 0.43 \quad \text{for } 500 < Re_p < 200,000 \quad \dots \dots \quad (9)$$

substituting these values of  $C_d$  in Eq. (6) gives analytic expressions for  $U_t$ , or

$$U_t, \text{ spherical} = \frac{g(\rho_s - \rho_g) d_p^3}{18} \quad \text{for } Re_p < 0.4 \quad \dots \dots \quad (10)$$

$$U_t, \text{ spherical} = \frac{\frac{4}{225}}{} \frac{(\rho_s - \rho_g)^2 g^2}{\rho_g} \frac{1}{3} d_p \quad \text{for } 0.4 < Re_p < 500 \\ \dots \dots \quad (11)$$

$$U_t, \text{ spherical} = \frac{3.1g (\rho_s - \rho_g) d_p^{\frac{1}{2}}}{\rho_g} \quad \text{for } 500 < Re_p < 200,000 \\ \dots \dots \quad (12)$$

$$\text{where } Re_p = \frac{d_p \rho_g U_t}{\mu}$$

For non-spherical particles, Pettyjohn and Christiansen (1948) have proposed that the terminal velocity can be estimated by multiplying the above  $U_t$  values by a correction factor.

$$\gamma = 0.843 \log \left( \frac{\varphi}{0.065} \right) \dots \dots \quad (13)$$

where  $\varphi$  = sphericity of particle.

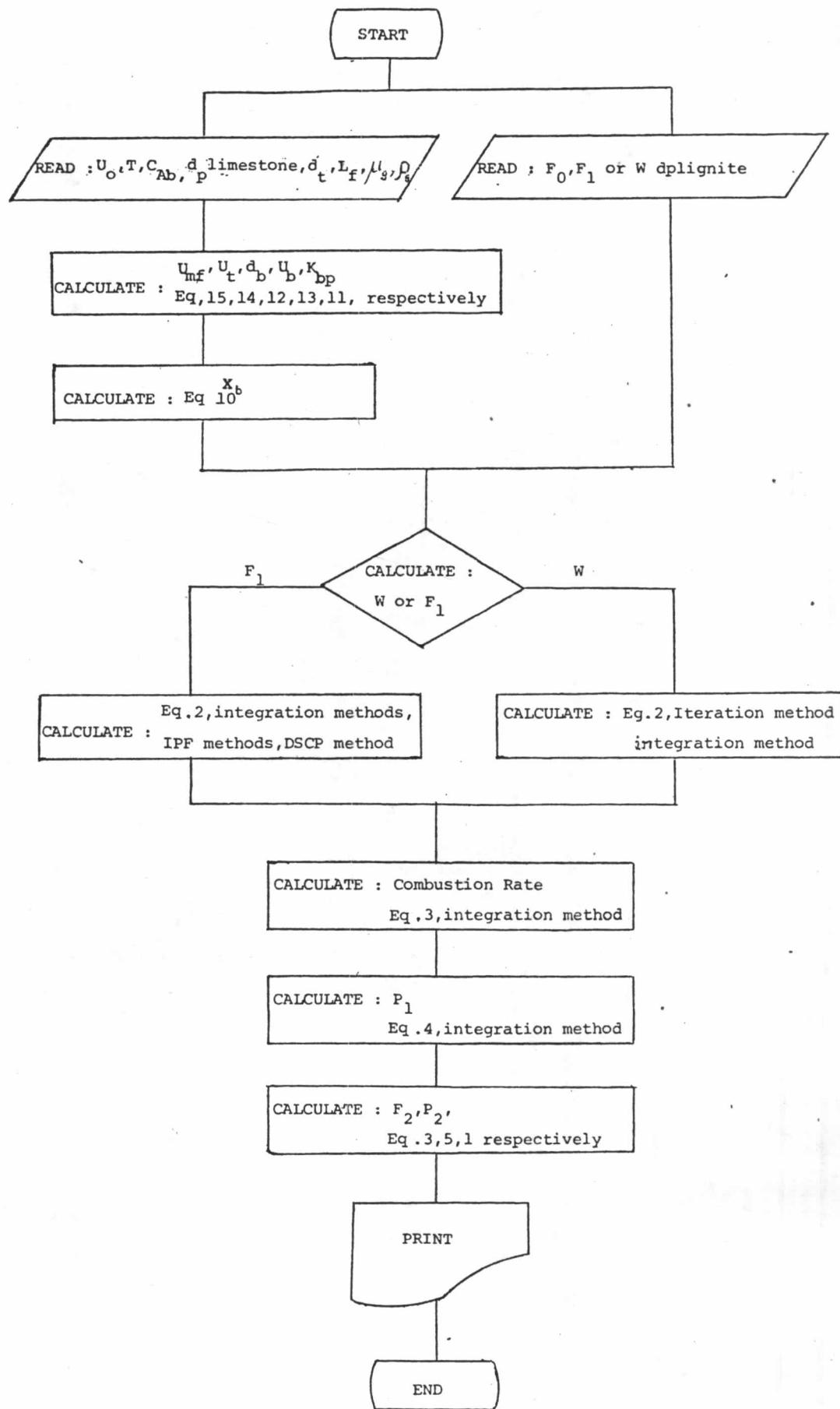
## Appendix C

### System Model Computation

The system model is computed by a numerical digital computation method.

In computation, the Simpson rule and the Trapezoidal rule are used for integration, an Iteration method is used for finding the weight of Char in bed W, Davies-Swann-Campey and Powell's method (DSCP) for optimization in order to find the value of the overflow rate  $F_1$ . Interpolation uses Newton Forward, Newton Backward, Everett method, IPF.

The flow chart and computer programs for system model calculation are shown followingly. The numbers of equations in the flow chart are from section 4.4.8.



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C THESE PROGRAM WRITTEN BY MR.NUTTAKUT WACHARAKUNDILOKE
C IT IS SYSTEM MODELLING PROGRAM OF COMBUSTION EFFICIENCY
C DEPARTMENT OF CHEMICAL ENGINEERING, FACULTY OF ENGINEERING
C CHULALONGKORN UNIVERSITY
C H=FINITE DIFFERENCE OF PARTICLE DIAMETER
C HZR=FINITE DIFFERENCE OF PARTICLE .LT.DPCMINT
CHF1TR=FINITE DIFFERENCE OF TRIAL FEED RATE
C THE VALUE OF F1STRT IS .LT. FIEND, THE VALUE OF F1X IS FROM LARGE TO SMALL
C TK=TEMPERATURE OF AIR IN BED IN KELVIN UNIT
C MC=MOLECULAR WEIGHT OF CARBON, MS=MOLECULAR WEIGHT OF SULPHUR
C P=ATMOSPHERIC PRESSURE
C R=GAS CONSTANT
C EXC=EXCESS AIR
C PCTCCL=PERCENT OF FIXED CARBON IN COAL
C FCCL=PERCENT OF ALL CARBON IN COAL
C PSCL=PERCENT OF SULPHUR IN COAL
C THE PERCENT ABOVE IS NOT THE REAL PERCENT MEANING, OTHERWISE HAVE TO *100.
C DG=DENSITY OF AIR AT 35 C
C N=NUMBERS OF PARTICLE DIAMETER VALUES OF FEED
C EMEU=VISCOUSITY OF AIR
C
C NN=NUMBERS OF SUPERFICIAL VELOCITY(LO)
C LET PCTCCL=FCCL
C
1      DIMENSION Y1(999),Y11(999),Y1ZF(999),Y2(999),Y112R(999)
2      DIMENSION DPC(999),DPC12(999),PC(999),P1(999),P2(999)
3      DIMENSION UT(999),II(999),SF(999),RR(999),KK(999)
4      DIMENSION EFFCR(999),F1XX(999),YSUM2(999),ERUPDN(999)
5      DIMENSION LC(50),DAI(50),CAFI(50),DGI(50),EMEU(50),F1I(50)
6      DIMENSION TKI(50)
7      DIMENSION UMF1(50)
8      DIMENSION FOI(50)
9      DIMENSION WI(50)
10     INTEGER TRY,KNOWF1
11     REAL II,KK,IIX,KKX
12     REAL LOWLMT
13     READ(5,10) DPCMINT,DPCMINT,H,DCL,G,DT,DCP,N
14     10 FORMAT(7F10.4,I10)
C F1STRT IS STARTING POINT OF TRIAL F1, AND IS UPPER LIMIT
C FIEND IS END POINT OF TRIAL F1, AND IS LOWER LIMIT
15     READ(5,11) F1STRT,FIEND,HF1TR,EFS,HZR
16     11 FORMAT(5F10.4)
C SPH=Sphericity OF COAL PARTICLE
17     READ(5,20) PCTCCL,SPH,NN
18     20 FORMAT(2F10.4,I10)
19     READ(5,21) (U0(I),I=1,NN)
20     READ(5,21) (CAI(I),I=1,NN)
21     READ(5,21) (FCI(I),I=1,NN)
22     READ(5,21) (WI(I),I=1,NN)
23     READ(5,21) (FII(I),I=1,NN)
24     21 FORMAT(8F10.4)

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25      READ(5,22)  (CAPI(I),I=1,NN)
26      READ(5,22)  (DGI(I),I=1,NN)
27      READ(5,22)  (EMEUI(I),I=1,NN)
28      22 FORMAT(EE10.4)
C FOR BED HEIGHT MORE THAN TDH,A=C.
29      READ(5,23) A
30      23 FORMAT(F10.4)
C*******
C IF WE KNOW F1, THEN HAVE TO TRY FOR W, LET DATA BELOW IN A READ STATEMENT BE
C W=0.0,WINIT=.....,F1=.....,KNOWF1=1,MAX=.....
C IF WE DON'T KNOW F1,BJT WE KNOW THE VALUE OF W, LET DATA BE
C W=... .,WINIT=0.,F1=0.,KNOWF1=0,MAX=0
C*******
C IF F1 KNOWN,W TO BE FOUND, AND F1 .EQ.0.,W FOUND IS THE MAX. CNE
C THAT MAKES THE CARBON IN COAL COMBUSTED ALL
C*******
31      READ(5,25) W,WINIT,KNOWF1,MAX
32      25 FORMAT(2F10.4,2I1)
C IF DESIRE TO USE OPTIMIZATION FOR FINDING F1, LET KYDSCP=1
C KYDSCP=KEY TO OPEN TO DSCP METHOD, KYDSCP=1 MEANS 'OPEN', =0 MEANS 'CLOSE'
C IMAX=ALLOWABLE NUMBERS OF ITERATION IN DSCP
C HDSCP=FINITE DIFFERENCE IN DSCP
C XODSCP=THE INITIAL VALUE OF F1 TRIAL FROM WHICH THE SEARCH START
33      READ(5,26) KYDSCP,IMAX,HDSCP,XODSCP
34      26 FORMAT(2I10,2F10.4)
35      READ(5,30) (PC(I),I=1,N)
36      30 FORMAT(EE10.4)
37      READ(5,32) P,R,PCCL,MC
38      32 FORMAT(3F10.4,I10)
39      READ(5,33) (TKI(I),I=1,NN)
40      READ(5,33) (UMFI(I),I=1,NN)
41      33 FORMAT(8F10.4)
42      DO 100 I=1,N
43      DPC(I)=DFCMIN+FLOAT(I-1)*H
44      100 CONTINUE
45      DPMX=DPCMIX-HZR
46      DPZR=HZF
47      NL=(DPMX-DPZR)/HZR+1.
C NL NOT BE EQUAL TO ZERO
48      NZR=N+NL
C*****
C IF KNOWING F1, INSERT THESE PROGRAM(WITHOUT C) , BUT ALREADY KNOWN W
C   F1=.....
C   KEY=1.
C   IF(KEY.EQ.1.) GOTO 460
C
49      DO 1000 ITR=1,NN
50      W=WI(ITR)
51      F00=FOI(ITR)
52      UMF=UMFI(ITR)
53      TK=TKI(ITR)
54      U00=LO(ITR)
55      DA=DAI(ITR)
56      CAP=CAPI(ITR)
57      DG=DGI(ITR)
58      EMEI=EMEUI(ITR)

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59      F1=F1I(17R)
60      IF(KNOWF1.EQ.1) GO TO 190
61      L=(F1STRT-F1END)/HF1TR+1.1
62      WRITE(6,189) L
63      189 FORMAT(//T1C,'L=',I10)
64      GO TO 191
65      190 CONTINUE
66      W=WINIT
67      W0=W
68      L=MAX
69      191 CONTINUE
70      DO 400 TRY=1,L
71      IF(KNOWF1.EQ.1) GO TO 192
72      Z=TRY-1
73      FIX=F1STRT-Z*HF1TR
74      GO TO 193
75      192 CONTINUE
76      FIX=F1
77      193 CONTINUE
78      DO 200 I=1,N
79      N1=N-I+1
80      DPCX=CPCMAY-FLCAT(I-1)*H
81      CALL IICF(CPCMAY,H,FIX,IIX,DCL,G,DT,DCP,DA,PCTCCL,SPH,
*     U00,W,CAP,CG,EMEU,DPCX,JMF,A,LTX,SHX,RRX,KKX)
82      Y2(I)=PC(N1)/((DPCX**3)*IIX)
83      CALL INTGT(H,I,Y2,SUM2)
84      Y11(N1)=(CPCX**3)*IIX/ABS(RFX)
85      Y11(N1)=Y11(N1)*SU12
86      200 CONTINUE
87      DO 300 I=1,NL
88      NL1=NL-I+1
89      DPX=CPCMAY-FLDAT(I)*HZR
90      CALL IIDP(CPCMAY,HZR,FIX,IIX,DCL,G,DT,DCP,DA,PCTCCL,SPH,
*     U00,W,CAP,CG,EMEU,DPX,UHF,A,UTX,SHX,RRX,KKX)
91      Y11ZR(NL1)=(CPCX**3)*IIX/ABS(RFX)
92      Y11ZR(NL1)=Y11ZR(NL1)*SUM2
93      300 CONTINUE
94      CALL INTGT(H,N,Y11,SUM11)
95      CALL INTGT(HZR,NL,Y11ZR,SUM12)
96      SUM1=SUM11+SUM12
97      IF(KNOWF1.EQ.1) GO TO 331
98      DIF=(W/FC0)-SUM1
99      YDIF=APS(DIF)
100     IF(YDIF.LE.EFS) GO TO 440
101     EFRCF(TFY)=YDIF
102     FIXX(TRY)=FIX
103     GO TO 400
104     331 W1=FCC*SLM1
105     DIF=APS(W0-W1)
106     IF(DIF.LE.EPS1) GO TO 459
107     W0=W1
108     W=WC
109     400 CONTINUE
110     IF(KNOWF1.EQ.1) GO TO 451
111     DIV=1
112     WRITE(6,410)

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```

113    410 FORMAT('1',T1C,'TRYING FOR F1 IS DIVERGE'/  

114      * T10,'TRY AGAIN BY HAND CHECKING AND THEN GO TO NEXT STEP')/  

115      WRITE(6,42C)  

116    420 FORMAT(T10,7('-----')/125,'F1 TRIAL',T45,'|',  

117      * T60,'ERR CF'/T10 ,7('-----'))/  

118      WRITE(6,43C)(F1XX(I),ERROR(I),I=1,L)  

119    430 FORMAT(T25,F1C.4,T45,'|',T55,F13.8)  

120      WRITE(6,434)  

121    434 FORMAT(T1C,7('-----'))  

122      IF(KYDSCP.NE.1) GO TO 437  

123      C FIND FIXX THAT ITS ERROR IS EQUAL TO ZERO BY OPTIMIZATION  

124      C USING DAVIES-SWANN-CAMPEY AND POWELL'S METHOD(DSCP)  

125      LOWLMT=F1XX(L)  

126      UPLMT=F1XX(1)  

127      DO 435 I=1,L  

128      L I1=L-(I-1)  

129      ERUPDN(L11)=ERROR(I)  

130      435 CONTINUE  

131      CALL DSCF(HDSCP,XDDSCP,F1,EFF1,EPS,IMAX,LOWLMT,HFLTR,EFLFCN,L)  

132      WRITE(6,438) F1,ERF1  

133      438 FORMAT(1HC/2CX,'MINIMIZATION BY DSC-POWELL METHCD'//25X,6HX = ,  

134      * F12.5,25X,6HPM = ,F12.5)  

135      WRITE(6,436) F1,ERF1  

136      436 FORMAT(//,T10,'BUT AFTER HAND CHECKING AND DESIRING TO LSE OPTIMIZ  

137      * ATION NUMERICAL CALCULATION',//,T1C,'OVERFLOW,F1=',F10.4,  

138      * T45,'BUT ERROR FROM CALCULATION =',F10.4)  

139      GO TO 460  

140      437 STOP  

141      440 DIV=C  

142      F1=F1X  

143      WRITE(6,45C) F1  

144      450 FORMAT('1',T1C,'TRYING FOR F1 IS CONVERGE',T40,'F1=',F1C.4)  

145      GO TO 46C  

146      451 CONTINUE  

147      WRITE(6,452) W  

148      452 FORMAT('1',T1C,'FINDING W IS DIVERGE, LAST TRIAL OF W IS',F1C.4)  

149      STOP  

150      459 CONTINUE  

151      W=W1  

152      460 CONTINUE  

153      ****  

154      C FIND TABLE OF DPC(I),PD(I),SH(I),KK(I),RR(I),II(I)  

155      C FROM NEAR ZERO TO DPCMAX  

156      C INPUT F1 FROM HAND CHECKING IF TRYING FOR F1 IS DIVERGE  

157      DO 480 I=1,NL  

158      DPC12(I)=HZR*FLOAT(I)  

159      480 CONTINUE  

160      DO 490 I=1,N  

161      NLI=NL+I  

162      DPC12(NLI)=DPCMIN+FLOAT(I-1)*H  

163      490 CONTINUE  

164      DO 500 I=1,NZR  

165      DPX=DPC12(I)  

166      IF(I.LE.NL) GO TO 495  

167      HZR=F  

168      GO TO 496

```

```

158   495 CONTINUE
159     HHZ R=HZF
160   496 CONTINUE
161     CALL IICP(DPCMAX,HZF,F1,IIX,DCL,G,CT,DCP,DA,PCTCCL,SPH,
162       * UOO,k,CAP,EG,E4EJ,DPX,UMF,A,UTX,SFX,RRX,KKX)
163     UT(I)=UTX
164     II(I)=IIX
165     SH(I)=SFX
166     RR(I)=RRX
167     KK(I)=KKX
168   500 CONTINUE
169     WRITE(6,510) UOO,F1,W
170   510 FORMAT('1',T1C,*UU =',F10.4,',',T30,'F1 =',F10.4,',',T5C,'W =',
171       * F1C.4/)
172     WRITE(6,520)
173   520 FORMAT(110,8('-----')/115,'DPC',T21,'|',T26,'UT',T33,'|',
174       * T38,'SH',T44,'|',T52,'KK',T58,'|',T65,'RR',T72,'|',T78,'II'/
175       * T1C,E('-----'))
176     WRITE(6,530) (DPC12(I),UT(I),SH(I),KK(I),RF(I),II(I),I=1,NZ)
177   530 FORMAT(110,F1C.4,T21,'|',T22,F10.4,T33,'|',T34,F10.4,T44,'|',
178       * T45,F1C.4,T58,'|',T59,F10.4,T72,'|',T73,F10.4)
179     WRITE(6,540)
180   540 FORMAT(T1C,8('-----'))
C***FIND P1(I)
C***FIND P1(EFC1), SIZE DISTRIBUTION OF COAL PARTICLE IN F1
181   DO 600 I=1,N
182     N1=N-I+1
183     NZR1=NZR-I+1
184     YSUM2(I)=PO(N1)/((DPC(N1)**3)*II(NZR1))
185     CALL INTGT(H,I,YSJ42,SUM2)
186     A=FOO*((DPC(N1)**3)*II(NZR1))/(W*ABS(RR(NZR1)))
187     P1(NZR1)=A*SLM2
188   600 CONTINUE
189   DO 700 I=1,NL
190     NZR1=NZR-I+1
191     DPCX=DPCM1-FLCAT(I)*HZR
192     P1(NZR1)=FCC*(DPCX**3)*II(NZR1)*SUM2/(W*ABS(RR(NZR1)))
193   700 CONTINUE
194   DO 7C1 I=1,NZ
195     NT=NZR-(I-1)
196     P1T=P1(NT)
197     IF(P1T.EQ.C.) GO TO 702
198   701 CONTINUE
199     GO TO 7C4
200   702 NT1=NT-1
201     DO 7C3 I=1,NT1
202     P1(I)=C.
203   703 CONTINUE
204   704 CONTINUE
205     WRITE(6,7C5)
206   705 FORMAT('1',T1C,7('-----'))
207     WRITE(6,706)
208   706 FORMAT(T1C,'PARTICLE DIAMETER OF COAL,CM',T40,'|',T51,
209       * 'SIZE DISTRIBUTION ,CM**(-1)')
210     WRITE(6,7C7)
211   707 FORMAT(T1C,7('-----'))

```

```

206      WRITE(6,70E) (CPC(I),P0(I),I=1,N)
207      70E FORMAT(718,F12.4,T40,'|',T51,F13.4)
208      WRITE(6,707)
209      WRITE(6,710) FCO,I,U00,F1
210      710 FORMAT('1',T1C,'F0 =',F10.4,',',T25,'h=',F10.4,',',T40,'UC =',
211           * F10.4/T1C,'F1 =',F10.4)
212      WRITE(6,720)
213      720 FORMAT(71C,7('-----'))
214           * T25,'CFC1,CN',T45,'|',T60,'P1,CM**-1'/T10,7('-----')
215           WRITE(6,730) (CPC12(I),P1(I),I=1,NZ)
216           730 FORMAT(T25,F10.4,T45,'|',T6C,F10.4)
217           WRITE(6,740)
218           740 FORMAT(71C,7('-----'))
C***  

C**FIND ELUTRIATION RATE, F2  

219      DPC X=HZF  

220      DO 800 I=1,NL  

221           YSUM2(I)=3.*W*P1(I)*RR(I)/DFCX  

222           DPC X=CPCX+HZR  

223           800 CONTINUE  

224           CALL INTGT(HZR,NL,YSUM2,SMF21)  

225           DO 850 I=1,N  

226           NLI=NL+I  

227           DPC X=CFCMIN+FLOAT(I-1)*H  

228           YSUM2(I)=3.*W*P1(NLI)*RP(NLI)/DPCX  

229           850 CONTINUE  

230           CALL INTGT(H,N,YSUM2,SMF22)  

231           SMF2=SMF21+SMF22  

232           WRITE(6,E60) SMF2  

233           860 FORMAT('1',T1C,'RATE OF COAL COMBUSTED =',F10.4)
234           F2=FCO-F1-SMF2  

C**  

C* FIND P2(CPC2), SIZE DISTRIBUTION OF COAL PARTICLE IN F2  

235           DO 900 I=1,NZF  

236           P2(I)=W*KK(I)*P1(I)/F2  

237           900 CONTINUE  

238           WRITE(6,S05) F2  

239           905 FORMAT('1',T1C,'F2 =',F10.4)
240           WRITE(6,S10)
241           910 FORMAT(71C,7('-----'))
242           * T25,'CFC2,CN',T45,'|',T60,'P2,CM**-1'/T10,7('-----')
243           WRITE(6,73C) (CPC12(I),P2(I),I=1,NZ)
244           WRITE(6,74C)  

C*  

C CONCLUSION  

245           EFF=(SMF2/FCC)*100.  

246           IF(F2.GT.C.) GO TO 915
247           F2=C.C  

248           915 CONTINUE  

249           WRITE(6,S20) SMF2  

250           920 FORMAT(71C,'FATE OF COAL COMBUSTED =',F10.4)
251           WRITE(6,S3C) FOO,F1,F2,EFF
252           930 FORMAT(T10,'FC =',F10.4,T25,'F1 =',F10.4,T40,'F2 =',F10.4,
253           * T55,'EFF =',F10.4)
C FIND PERCENT OF EXCESS AIR IN BED  

254           AA=0.21*P*(3.1416*(DT+DDP)*(DT-DDP)/4.)*U00
255           BB=R*TK*FCC*FCCL/4C
256           EXC=AA/BB-1.
257           PCTEXC=EXC*100.0
258           WRITE(6,S4C) TK,PCTEXC
259           940 FORMAT(//T10,'TEMPERATURE OF AIR IN BED =',F10.4,3X,'K'/
260           * T10,'EXCESS AIR =',F10.4)
261           1000 CONTINUE
262           STOP
263           END

```

```

C SUBROUTINE FOR FINDING II,SH,RR,KK
C H= FINITE DIFFERENCE OF COAL DIAMETER
C F1= FEED RATE OF COAL
C DPCM=MAXIMUM PARTICLE DIAMETER
C DCL=DENSITY OF COAL PARTICLE
C DT=DIAMETER OF BED
C DDP=DIAMETER OF DRAIN PIPE IN BED
C DA=CoeffICIENT DIFFUSIVITY
C PCTCCL=PERCENT CARBON IN COAL PARTICLE(IN FRACTION)
C SPH=SUPERFICITY
C W=WEIGHT OF COAL IN BED
C CAP=CONCENTRATION OF OXYGEN IN PARTICULATE PHASE
C
1      SUBROUTINE IIDP(DPCM,DH,F1,II,DCL,G,DT,DDP,DA,PCTCCL,SPH,
2      * UO,W,CAP,DG,EMEU,DPC,UMF,A,UT,SH,RR,KK)
3      DIMENSION Y(1000)
4      REAL II,KK,KKX
5      N=(DPCM-DPC)/DH+1.
6      DO 10 I=1,N
7      DPX=DPC+FLOAT(I-1)*F
8      CALL TMV(DCL,G,DPX,CG,EMEU,SPH,UTX)
9      CALL SRK(DCL,G,DT,DDP,DA,PCTCCL,UO,W,CAP,DG,EMEU,DFX,UMF,A,
* UTX,SHX,RRX,KKX)
10     Y(I)=(F1/W+KKX)/FRX
11     CONTINUE
12     CALL INTGT(H,N,Y,SLM)
13     IF(SUM.GE.70) GO TO 50
14     II=1./EXP(SLM)
15     CALL TMV(DCL,G,DPC,CG,EMEU,SPH,UT)
16     CALL SRK(DCL,G,DT,DDP,DA,PCTCCL,UO,W,CAP,DG,EMEU,DPC,UMF,A,
* UT,SH,RR,KK)
17     RETURN
18     50 WRITE(6,60) SUM,DPC
19     60 FORMAT('1',T10,'SUM IS MORE THAN 70, =',F10.4,'DPC= ',F10.4)
20     STOP
END

```

```
C SUBROUTINE FOR CALCULATING INTEGRATION WITH FINITE DIFFERENCE
C SUBROUTINE INTGT(H,N,Y,SLN)
C H=FINITE DIFFERENCE OF X
C Y=FUNCTION OF X
1      SUBROUTINE INTGT(H,N,Y,SUM)
2      DIMENSION Y(N)
3      RFAL ITRAP,ISIMP
4      IF(N.GE.7) GO TO 5
5      SUM1=0
6      DO 3 I=1,N
7          SUM1=SLN1+Y(I)*H
8      3 CONTINUE
9      SUM=SUM1
10     GO TO 60
11     5 CONTINUE
12     KT1=FLOAT(N)/2.+0.6
13     KT2=N/2
14     IF(KT1-KT2) 1C,4C,1C
C SIMPSON RULE
15     10 N1=N-1
16     YM1=Y(2)
17     DO 20 I=4,N1,2
18     YM1=YM1+Y(I)
19     20 CONTINUE
20     N2=N-2
21     YM2=Y(3)
22     DO 30 I=5,N2,2
23     YM2=YM2+Y(I)
24     30 CONTINUE
25     ISIMP=(I/3.)*(Y(1)+4.*YM1+2.*YM2+Y(N))
26     SUM=ISIMP
27     GO TO 60
C TRAPEZOIDAL RULE
28     40 YMID=Y(2)
29     N1=N-1
30     DO 50 I=3,N1
31     YMID=YMID+Y(I)
32     50 CONTINUE
33     ITRAP=(I/2.)*(Y(1)+2.*YMID+Y(N))
34     SUM=ITRAP
35     60 RETURN
36     END
```

```

C SUBROUTINE FOR CALCULATING SHERWOOD NUMBER(SH),
C SHRINKING RATE OF PARTICLE (FR)
C ELUTRIATION CONSTANT (KK)
C CAP=CONCENTRATION OF OXYGEN IN PARTICULATE PHASE,GMMCLE/CM**3
C PCTCC= PERCENT OF CARBON IN CCAL
C DCL = DENSITY OF COAL, CPC = DIAMETER OF CCAL PARTICLE
C UO USED IN CALCULATING KK MUST BE LESS THAN UMF
1      SUBROUTINE SRK(DCL,E,CT,CDF,DA,FCTCCL,UO,W,CAP,CG,EMEU,CPC,LMF,A,
* UT,SH,FR,KK)
2      REAL KK,K,K1
3      SH = 2.0+0.6*((UO*DCE/EMEU)**0.5)*((EMEL/(CG*DA))**(.1./3.))
* *(CPC**0.5)
4      RR=57.6*CAF*SH*DA/(CPC*FCTCCL*DCL)
5      IF(UT.GT.UO) GO TO 10
6      K1=130.*EXP(-10.4*((LT/UO)**0.5)*((UMF/(UO-UMF))**0.25))+A
7      K=3.1416*(CT-CDF)*(DT+DDP)/4.
8      KK=(CG*LC*K/W)*K1
9      RETURN
10     KK=0
11     RETURN
12     END

```

```

C SUBROUTINE FOR CALCULATING TERMINAL VELOCITY (UT)
C DS= DENSITY OF PARTICLE
C G= GRAVITATIONAL ACCELERATION
C DP=PARTICLE DIAMETER
C DG=DENSITY OF GAS
C EMEU=VISCOUSITY OF GAS
C SPH=SUPERICITY
1      SUBROUTINE TMV(DS,G,DP,DG,EMEU,SPH,LT)
2      ANU=C.843*ALOG10(SPH/C.165)
3      UTS PHR=C.153*(DP**1.14)*(G**0.71)*((DS-DG)**0.711)/((EMEU**0.43)
* *(DG**0.29))
4      UT=ANU*UTSPHR
5      RETURN
6      END

```

```

C DSC POWEL METHOD USING THE COMBINED CAVIES-SWANN-CAMPEY AND POWEL
C ****
C THIS SUBROUTINE USING THE COMBINED CAVIES-SWANN-CAMPEY AND POWELL
C METHOD(DSCP),TC COMPUTE THE MINIMUM OF A CONTINUOUS FUNCTION
C THE FOLLOWING PROVISIONS FOR USERS TO SPECIFY,
C X0-THE INITIAL VALUE OF X FROM WHICH THE SEARCH START
C DX-THE SEARCH STEP SIZE
C EPS-THE PRECISION INDEX
C MAX-THE MAXIMUM ALLOWABLE NUMBER OF ITERATIONS
C XOO,HH,YY,NN ARE ARGUMENTS IN SUBROUTINE SUBPROGRAM IPF,
C LOWER LIMIT ,FINITE DIFFERENCE,VALUES OF Y,NUMBERS OF Y RESPECTIVELY
C **** WRITTEN BY KUTTAWUT WACHARAKUNDILOK****

1  SUBROUTINE DSCP(X0,XM,FM,EPS,MAX,XCC,HH,YY,NN)
2    DIMENSION YY(NN)
3    DOUBLE PRECISION X1,X2,X3
4    DATA ITAF,JTAF/5,6/
5      H=DX
6    2 CONTINUE
C START THE SEARCH THE BOUND MINIMUM
7    ITR=0.
8    M=0
9    M1=0
10   CALL IPF(X00,HH,YY,NN,X0,FA)
11   X=X0+H
12   CALL IPF(X00,HH,YY,NN,X,FB)
13   IF(FB.LE.FA) GO TO 4
C REVERSE THE SEARCH DIRECTION TO OBTAIN DECREASING IN FUNCTION VALUE
C
14   X=X0
15   H=-2.*H
16   GO TO 6
C THE FUNCTION STILL DECREASING ,INCREASE THE STEP SIZE BY DOUBLE
C THE PREVIOUS INCREASE IN STEP SIZE
17   4 H=2.*H
18   6 CONTINUE
19   X=X+H
20   CALL IPF(XCC,HH,YY,NN,X,FC)
21   IF(FC.GT.FB) GO TO 25
22   FB=FC
23   ITR=ITR+1
24   IF(ITR.GT.MAX) GO TO 25
25   GO TO 4
C MINIMUM IS FOUND
26   8 CONTINUE
27   X2=X-H
28   IF(X2.LT.X) GO TO 9
29   X1=X
30   X3=X2-H/2.
31   GO TO 10
32   9 CONTINUE
33   X3=X
34   X1=X2-H/2.
35   10 H=-H/2.

```

```

36      X=X+H
37      X2 S=X2
38      CALL IPF(XCC,HH,YY,NN,X2S,FXX)
39      CALL IPF(XCC,HH,YY,NN,X,FXX)
40      IF(FX2.LT.FXX) GO TO 12
41      X1=X2
42      X2=X
43      GO TO 14
44      12 X3=X
45      14 CONTINUE
46      X1S=X1
47      X2S=X2
48      X3S=X3
49      CALL IPF(XOO,HH,YY,NN,X1S,FA)
50      CALL IPF(XOO,HH,YY,NN,X2S,FB)
51      CALL IPF(XCC,HH,YY,NN,X3S,FC)
52      B=2.* (FA-2.*FB+FC)
53      IF(B) 15,24,15
54      15 X=X2-ABS(H)*(FA-FC)/B
55      WRITE(6,111) X
56      111 FORMAT(//,T1C,'X=' ,F1C.4)
57      ITR=C
58      16 CONTINUE
59      XX1=X-X1
60      XX2=X-X2
61      XX3=X-X3
62      IF(ABS(XX1).LE.EPS) GO TO 40
63      IF(ABS(XX2).LE.EPS) GO TO 40
64      IF(ABS(XX3).LE.EPS) GO TO 40
65      WRITE(6,222) X
66      222 FORMAT(T1C,'TEST=' ,F1C.4)
67      ITR=ITR+1
68      IF(ITR.GT.MAX) GO TO 38
69      CALL IPF(XOO,HH,YY,NN,X,FP)
70      XP=X
71      XP2=XP-X2
72      IF(ABS(XP2).LE.EPS) GO TO 40
73      IF(FB.LT.FP) GO TO 18
74      IF(X.LT.X2) GO TO 17
75      X1=X2
76      FA=FB
77      X2=X
78      FB=FP
79      GO TO 20
80      17 X3=X2
81      FC=FB
82      X2=X
83      FB=FP
84      GO TO 20
85      18 CONTINUE
86      IF(X.LT.X2) GO TO 19
87      X3=X
88      FC=FP
89      GO TO 20
90      19 X1=X
91      FA=FP

```



```

92      20 CONTINUE
C NOW PERFORM THE PARABOLIC INTERPOLATION
93      A=(X2-X3)*FA+(X3-X1)*FB+(X1-X2)*FC
94      IF(A) 21,28,21
95      21 X=(1./2.)*((X2**2-X3**2)*FA+(X2**2-X1**2)*FB+(X1**2+X2**2)*FC)/A
96      IF(X.GT.X3) GO TO 22
97      IF(X.LT.X1) GO TO 22
98      GO TO 23
99      22 X31.2=(X3-X1)/2.
100      B=2.**(FA-2.*FB+FC)
101      IF(B.EQ.0.) GO TO 24
102      X=X2-AB$IX312)*(FA-FC)/B
103      23 CONTINUE
104      GO TO 16
105      24 CONTINUE
106      IF(M) 26,26,30
107      26 M=M+1
108      GO TO 8
109      28 CONTINUE
110      IF(M1) 29,29,32
111      29 M1=M1+1
112      GO TO 8
113      30 WRITE(JTAP,281)
114      GO TO 40
115      32 WRITE(JTAP,282)
116      GO TO 40
117      38 CONTINUE
118      WRITE(JTAP,28C) ITR
C
119      40 CONTINUE
C
120      XM=X
121      CALL IFF(XCC,HH,YY,NN,XM,FM)
122      280 FORMAT(1HC,2CX,' SEARCH FAILED WITHIN',I6,' ITERATIONS')
123      281 FORMAT(1HC/2CX,' THE DSC DIVISOR BEING ZERO SECOND TIME')
124      282 FORMAT(1HC/2CX,'THE POWELL DIVISOR BEING ZERO SECOND TIME')
C
25      RETURN
6       END

```

```

C SUBROUTINE IPF(INTERPOLATION) INCLUDES
C NEWTON FORWARD INTERPOLATION FORMULAR
C NEWTON BACKWARD INTERPOLATION FORMULAR
C AND EVERETT INTERPOLATION FORMULAR
C M=DEGREE OF THE INTERPOLATION PCLINCMAL,M=N-1
C N= THE NUMBER OF TABULATED DATA POINTS
C X1=INITIAL VALUE OF X,H=FINITE DIFFERENCE,CY=THE DIFFERENCE OF Y OR ANY DEGRE
C XP=THE POINT OF X WHICH IS USED TO FIND YP OR Y-REQUIRED
C N NOT BE LESS THAN 7
1      SUBROUTINE IPF(X1,H,Y,N,XP,YP)
2      DIMENSION Y(N)
3      DIMENSION CY(200,2CC)
4      DOUBLE PRECISION DY,PF,QQ,A,F,Q
5      M=N-1
6      N1=N-1
7      DO 2 I=1,N1
8      2 DY(I,1)=Y(I+1)-Y(I)
9      DO 6 K=2,M
10     L=N-K
11     DO 4 I=1,L
12     4 DY(I,K)=CY(I+1,K-1)-CY(I,K-1)
13     6 CONTINUE
14     XN=X1+H*FLCAT(N1)
15     SECT=(XN-X1)/3.
16     SECT1=X1+SECT
17     SECT2=X1+2*SECT
18     IF(XP.GT.SECT2) GO TO 30
19     IF(XP.GE.SECT1) GO TO 20
C XP=X1+P*H
C NFIF
20     10 P=(XP-X1)/H
21     PP=1.
22     YP=Y(1)
23     DO 12 K=1,4
24     K1=K-1
25     PP=PF*(P-FLOAT(K1))/FLCAT(K)
26     12 YP=YP+PP*DY(1,K)
27     RETURN
C XN=XP-P*H
C NBIF
28     30 P=(XN-XP)/H
29     PP=1.
30     YP=Y(N)
31     NN=N
32     DO 33 K=1,4
33     K1=K-1
34     PP=-PP*(P-FLCAT(K1))/FLCAT(K)
35     NN=NN-1
36     33 YP=YP+PP*CY(NN,K)
37     RETURN
C EIF
38     20 K=FLOAT(N)/2.+0.6
39     KI=N/2+C.6

```

```
40      X0=X1+(FLOAT(K)-1.)*H
41      P=(XP-XC)/H
42      C=1.-P
43      A=1.
44      PP=P
45      QQ=Q
46      YP=Q*Y(K)+P*Y(K+1)
47      IF(K-KI) 22,21,22
48      21 L=M/2
49      GO TO 25
50      22 L=M/2-1
51      25 CONTINUE
52      DO 23 I=1,3
53      II=2*I
54      A=A*II*(II+1)
55      QQ=(Q+I)*QQ*(Q-I)
56      PP=(P+I)*PP*(P-I)
57      KK=K-I
58      23 YP=YP+(QQ*DY(KK,II)+PP*DY(KK+1,II))/A
59      RETURN
60      END
```

C SUBROUTINE FOR FINDING CAP,X,KBP  
 C CAP= CONCENTRATION OF OXYGEN IN PARTICULATE PHASE  
 C CAB= CONCENTRATION OF OXYGEN IN BUBBLE AT SPECIFIED HEIGHT  
 C CAO= CONCENTRATION OF OXYGEN IN BUBBLE AT INLET OF FBC(GAS INLET)  
 C UC= SUPERFICIAL VELOCITY, UB= BUBBLE VELOCITY, UMF= MINIMUM FLUIDISATION VEL.  
 C DB= BUBBLE DIAMETER , LF= HEIGHT OF BED  
 C G= GRAVITATIONAL ACCELERATION, DA= OXYGEN DIFFUSION COEFFICIENT  
 C KBP= GAS INTERCHANGE COEFFICIENT FROM BUBBLE TO PARTICULATE PHASE  
 C X= INTERPHASE CHANGE  
 C LB NOT EQUAL TO ZERO  
 C THIS SUBROUTINE DOUBLE PRECISION CAB,CAO,B,C

```
SUBROUTINE DXYC(CAB,CAC,UMF,UC,DB,DA,G,LF,UB,KBP,X,CAP)
DOUBLE PRECISION CAB,CAO,B,C
REAL LF,KBP
A=(DA**0.5)*(G**0.25)/(DB**1.25)
KBP=4.5*(UMF/DB)+5.85*A
UB=0.711*((G*DB)**0.5)+(UO-UMF)
X=KBP*LF/UB
IF(X.GT.50.) GO TO 10
B=EXP(-X)
GO TO 20
10 B=0.0
20 CONTINUE
C=1.
CAP=(CAB-CAO*B)/(C-B)
RETURN
END
```

C SUBROUTINE FOR FINDING BUBBLE DIAMETER IN FBC
 C LF=HEIGHT OF FBC BED
 C UO=SUPERFICIAL VELOCITY
 C UMF=MINIMUM FLUIDIZATION VELOCITY
 C DB=BUBBLE DIAMETER

```
SUBROUTINE DRCG(UO,UMF,LF,DB)
REAL LF
DB1=0.0326*((UO-UMF)**1.11)*(LF**0.81)
DB=DB1/1.81
RETURN
END
```

## An Example of Output from the Computer Simulation

UJ = 394.2300, F1 = 0.0259, W = 8.6584

CPC	UT	SH	KK	RR	II
0.0020	5.3542	2.3501	122.2946	0.5239	0.0001
0.0040	11.7996	2.4951	83.3734	0.2781	0.0001
0.0060	18.7332	2.6063	61.4450	0.1937	0.0002
0.0080	26.0041	2.7001	47.2101	0.1505	0.0004
0.0100	33.5367	2.7d27	37.2730	0.1241	0.0007
0.0120	41.2845	2.8575	30.0110	0.1062	0.0013
0.0140	49.2160	2.9262	24.5308	0.0932	0.0022
0.0160	57.3083	2.9901	20.2947	0.0833	0.0037
0.0180	65.5437	3.0502	16.9581	0.0756	0.0060
0.0200	73.90E6	3.1070	14.2896	0.0693	0.0092
0.0220	82.3915	3.1610	12.1284	0.0641	0.0137
0.0240	90.9833	3.2126	10.3591	0.0597	0.0196
0.0260	99.6760	3.2621	8.8972	0.0559	0.0274
0.0280	108.4627	3.3098	7.6797	0.0527	0.0371
0.0300	117.3381	3.3558	6.6584	0.0499	0.0492
0.0320	126.2966	3.4002	5.7963	0.0474	0.0633
0.0340	135.3329	3.4433	5.0645	0.0452	0.0803
0.0360	144.4461	3.4852	4.4400	0.0432	0.0993
0.0380	153.6292	3.5259	3.9047	0.0414	0.1212
0.0400	162.8804	3.5655	3.4439	0.0397	0.1449
0.0420	172.1967	3.6042	3.0456	0.0383	0.1718
0.0440	181.5752	3.6419	2.7001	0.0369	0.1994
0.0460	191.0136	3.6788	2.3994	0.0357	0.2304
0.0480	200.5058	3.7149	2.1369	0.0345	0.2612
0.0500	210.0615	3.7503	1.9070	0.0334	0.2953
0.0520	219.6669	3.7849	1.7053	0.0325	0.3283
0.0540	229.3241	3.8189	1.5276	0.0315	0.3647
0.0560	239.0313	3.8523	1.3709	0.0307	0.3988
0.0600	258.5903	3.9173	1.1095	0.0291	0.4990
0.0700	308.2705	4.0710	0.6707	0.0259	0.6631
0.0800	358.9575	4.2139	0.4181	0.0235	0.8507
0.0900	410.5413	4.3482	0.0000	0.0215	0.9040
0.1000	462.9353	4.4753	0.0000	0.0200	0.9053
0.1100	516.0653	4.5961	0.0000	0.0186	0.9067
0.1200	569.8855	4.7115	0.0000	0.0175	0.9082
0.1300	624.3323	4.8222	0.0000	0.0165	0.9060
0.1400	679.3696	4.9288	0.0000	0.0157	0.9077
0.1500	734.9612	5.0316	0.0000	0.0150	0.9095
0.1600	791.0750	5.1310	0.0000	0.0143	0.9113
0.1700	847.6821	5.2273	0.0000	0.0137	0.9171
0.1800	904.7559	5.3209	0.0000	0.0132	0.9191
0.1900	962.2759	5.4119	0.0000	0.0127	0.9174
0.2000	1020.2227	5.5005	0.0000	0.0123	0.9197
0.2100	1078.5762	5.5870	0.0000	0.0119	0.9219
0.2200	1137.3201	5.6714	0.0000	0.0115	0.9281
0.2300	1196.4395	5.7539	0.0000	0.0112	0.9306
0.2400	1255.91E9	5.8347	0.0000	0.0108	0.9293
0.2500	1315.7473	5.9137	0.0000	0.0105	0.9319
0.2600	1375.9124	5.9912	0.0000	0.0103	0.9346
0.2700	1436.4011	6.0673	0.0000	0.0100	0.9377
0.2800	1497.2051	6.1419	0.0000	0.0098	0.9440
0.2900	1558.3132	6.2152	0.0000	0.0096	0.9431
0.3000	1619.7173	6.2873	0.0000	0.0093	0.9461
0.3100	1681.4102	6.35E1	0.0000	0.0091	0.9491
0.3200	1743.3799	6.4279	0.0000	0.0090	0.9523
0.3300	1805.6230	6.49E5	0.0000	0.0083	0.9594
0.3400	1868.1252	6.5642	0.0000	0.0086	0.9528
0.3500	1930.8962	6.6308	0.0000	0.0094	0.9622

CPC	UT	SH	KK	RR	II
0.3600	993.9114	6.6965	0.0000	0.0083	0.9656
0.3700	2057.1733	6.7613	0.0000	0.0081	0.9691
0.3800	2120.6768	6.8252	0.0000	0.0080	0.9767
0.3900	184.4124	6.8862	0.0000	0.0079	0.9766
0.4000	2248.3772	6.9505	0.0000	0.0077	0.9762
0.4100	2312.5674	7.0120	0.0000	0.0076	0.9800
0.4200	2376.9771	7.0728	0.0000	0.0075	0.9838
0.4300	2441.6018	7.1328	0.0000	0.0074	0.9878
0.4400	2506.4380	7.1921	0.0000	0.0073	0.9959
0.4500	2571.4797	7.2508	0.0000	0.0072	0.9958

PARTICLE DIAMETER OF COAL, CM	SIZE DISTRIBUTION , CM** ( 1 )
0.0600	0.0200
0.0700	0.0400
0.0800	0.0500
0.0900	0.1000
0.1000	0.1500
0.1100	0.3750
0.1200	0.7500
0.1300	1.1200
0.1400	1.5000
0.1500	1.8750
0.1600	2.3500
0.1700	2.8200
0.1800	3.2500
0.1900	3.8800
0.2000	4.3000
0.2100	4.5500
0.2200	5.3200
0.2300	5.5000
0.2400	5.6200
0.2500	5.6500
0.2600	5.6800
0.2700	5.6700
0.2800	5.6300
0.2900	5.5500
0.3000	5.3500
0.3100	4.8000
0.3200	3.9500
0.3300	3.0500
0.3400	2.3000
0.3500	1.7500
0.3600	1.4000
0.3700	1.1000
0.3800	0.8200
0.3900	0.4600
0.4000	0.3500
0.4100	0.2500
0.4200	0.1800
0.4300	0.1200
0.4400	0.0700
0.4500	0.0300

$F_0 = 1.6717, W = 8.6584, U_0 = 354.2300$   
 $F_1 = 0.0259$

DPC1, CM		F1, CM** 1
0.0020		0.0000
0.0040		0.0000
0.0060		0.0000
0.0080		0.0000
0.0100		0.0000
0.0120		0.0000
0.0140		0.0000
0.0160		0.0000
0.0180		0.0000
0.0200		0.0000
0.0220		0.0000
0.0240		0.0000
0.0260		0.0000
0.0280		0.0000
0.0300		0.0000
0.0320		0.0000
0.0340		0.0000
0.0360		0.0000
0.0380		0.0000
0.0400		0.0000
0.0420		0.0000
0.0440		0.0000
0.0460		0.0000
0.0480		0.0000
0.0500		0.0000
0.0520		0.0000
0.0540		0.0000
0.0560		0.0576
0.0600		0.0841
0.0700		0.2017
0.0800		0.3911
0.0900		0.6065
0.1000		0.8859
0.1100		1.2372
0.1200		1.6409
0.1300		2.0832
0.1400		2.5667
0.1500		3.0791
0.1600		3.6045
0.1700		4.1363
0.1800		4.6311
0.1900		5.0430
0.2000		5.4172
0.2100		5.6884
0.2200		5.8689
0.2300		5.9194
0.2400		5.8615
0.2500		5.7194
0.2600		5.5003
0.2700		5.1653
0.2800		4.7696
0.2900		4.2186
0.3000		3.6099
0.3100		2.9339
0.3200		2.3286
0.3300		1.7944
0.3400		1.3921
0.3500		1.0553

DPC1, CM	P1, CM**-1
0.3600	0.7936
0.3700	0.5685
0.3800	0.3873
0.3900	0.2572
0.4000	0.2251
0.4100	0.1514
0.4200	0.0966
0.4300	0.0550
0.4400	0.0260
0.4500	0.0081

RATE OF COAL COMBUSTED = 1.5768

F2 = 0.0690

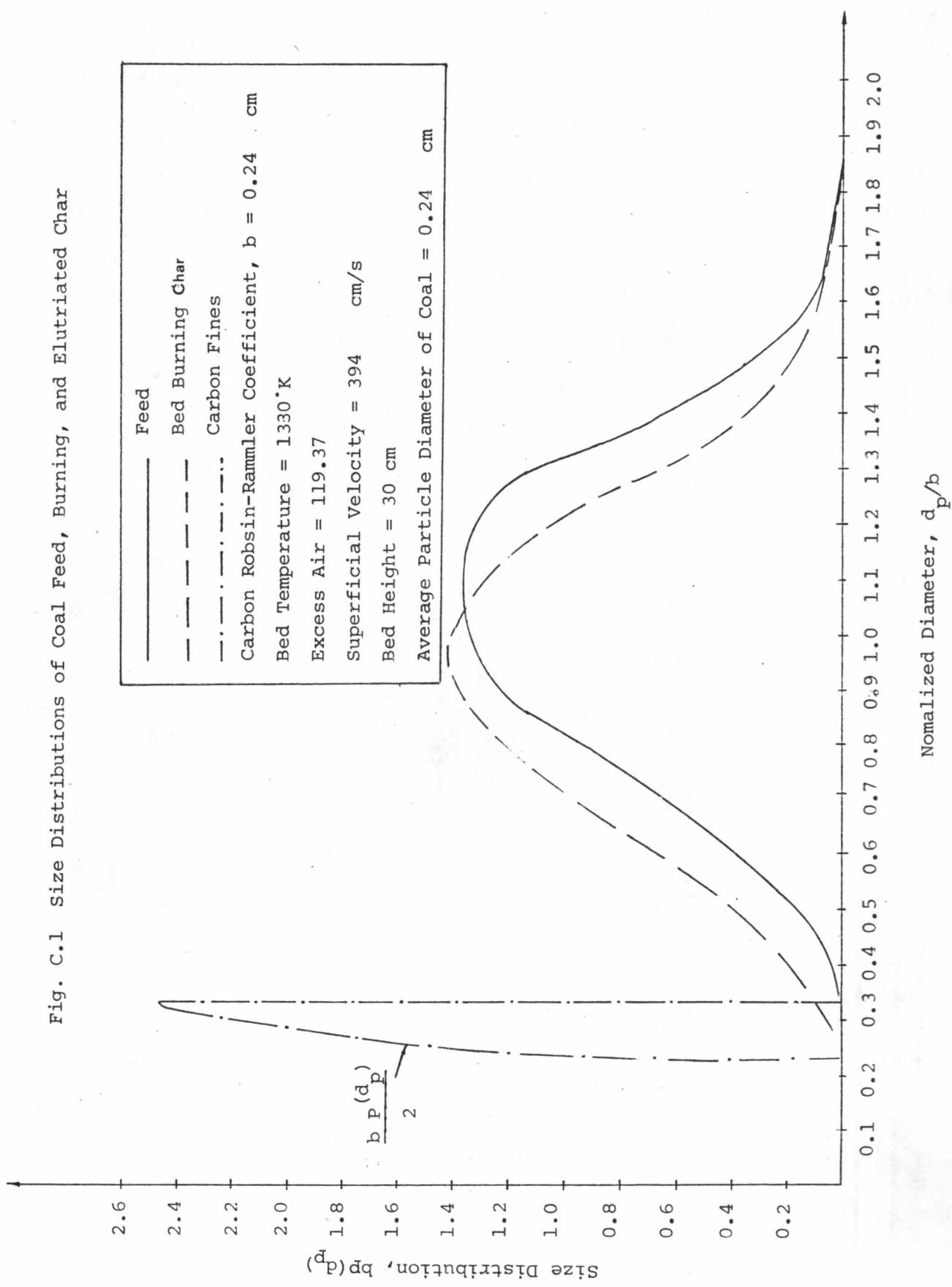
DPC2, CM	F2, CM** 1
0.0020	0.0000
0.0040	0.0000
0.0060	0.0000
C.0080	0.0000
C.0100	0.0000
C.0120	0.0000
C.0140	0.0000
C.0160	0.0000
C.0180	0.0000
C.0200	0.0000
0.0220	0.0000
C.0240	0.0000
0.0260	0.0000
C.0280	0.0000
C.0300	0.0000
0.0320	0.0000
0.0340	0.0000
0.0360	0.0000
C.0380	0.0000
0.0400	0.0000
0.0420	0.0000
0.0440	0.0000
0.0460	0.0000
0.0480	0.0000
C.0500	0.0000
0.0520	0.0000
0.0540	0.0000
0.0560	9.9063
C.0600	11.7023
C.0700	16.9619
C.0800	20.5056
0.0900	0.0000
0.1000	0.0000
C.1100	0.0000
0.1200	0.0000
0.1300	0.0000
0.1400	0.0000
C.1500	0.0000
C.1600	0.0000
C.1700	0.0000
0.1800	0.0000
C.1900	0.0000
0.2000	0.0000
0.2100	0.0000
C.2200	0.0000
0.2300	0.0000
0.2400	0.0000
0.2500	0.0000
C.2600	0.0000
0.2700	0.0000
0.2800	0.0000
0.2900	0.0000
0.3000	0.0000
C.3100	0.0000
C.3200	0.0000
C.3300	0.0000
0.3400	0.0000
0.3500	0.0000
C.3600	0.0000

DPC2, CM	F2, CM**-1
0.3700	0.0000
0.3800	0.0000
0.3900	0.0000
0.4000	0.0000
0.4100	0.0000
0.4200	0.0000
0.4300	0.0000
0.4400	0.0000
0.4500	0.0000

RATE OF COAL COMBUSTED = 1.576E  
F0 = 1.6717 F1 = 0.0259 F2 = 0.0690 EFF = 94.3203

TEMPERATURE OF AIR IN BED = 1330.0000 K  
EXCESS AIR = 119.3747

Fig. C.1 Size Distributions of Coal Feed, Burning, and Elutriated Char



## Appendix D

## Average Particles Diameter

```

C FIND THE AVERAGE DIAMETER OF LIGNITE PARTICLE
C DBPMAX=MAXIMUM DIAMETER OF BED PARTICLE
C DBPMIN=THE MINIMUM DIAMETER OF BED PARTICLE
C PBP(I)= SIZE DISTRIBUTION OF PARTICLE ,BETWEEN DP AND DP+D(I)
C NM=(DBPMAX-DBPMIN)/H+1, THE NUMBER OF DATAS TO BE CALCULATE
C H=FINITE DIFFERENCE OF DBP
1      DIMENSION DBP(100),PBP(100),YFB(100)
2      DO 200 J=1,2
3      READ(5,21) DBPMAX,DBPMIN,H,NM
4      21 FORMAT(3F10.4,I10)
5      READ(5,31) (PBP(I),I=1,NM)
6      31 FORMAT(8F10.4)
7      DO 100 I=1,NM
8      DBP(I)=DBPMIN+(I-1)*H
9      YFB(I)=PBP(I)/DBP(I)
10     100 CONTINUE
11     CALL INTGT (N,NM,YFB,SUM)
12     DPA=1/SUM
13     WRITE(5,36)
14     36 FORMAT('1',T11,'FIND THE AVERAGE DIAMETER OF LIGNITE PARTICLE')
15     WRITE (6,37)
16     37 FORMAT('0',T11,7('-----'))
17     WRITE(5,38)
18     38 FORMAT (T11,'DIAMETER OF LIGNITE PARTICLE ,CM',T45,' ',T51,'SIZE
* DISTRIBUTION, CM**(-1)')
19     WRITE(6,39)
20     39 FORMAT(T11,7('-----'))
21     WRITE(5,41) (DBP(I),PBP(I),I=1,NM)
22     41 FORMAT(T18,F13.4,T45,' ',T58,F13.4)
23     WRITE(6,42)
24     42 FORMAT(T11,7('-----'))
25     WRITE(6,43) DBPMAX,DBPMIN,NM,H,DPA
26     43 FORMAT(T11,'MAXIMUM DIAMETER OF BED MATERIAL = ',T51,F13.4/
*           T11,'MINIMUM DIAMETER OF BED MATERIAL = ',T51,F13.4/
*           T11,'RANGE NUMBERS OF PARTICLE DIAMETER = ',T51,I4/
*           T11,'FINITE DIFFERENCE = ',T51,F13.4//'
*           T11,'THE AVERAGE BED PARTICLE DIAMETER = ',T51,F13.4)
27     200 CONTINUE
28     STOP
29     END
C SUBROUTINE FOR CALCULATING INTEGRATION WITH FINITE DIFFERENCE
C SUBROUTINE INTGT(H,N,Y,SUM)
C H=FINITE DIFFERENCE OF X
C Y=FUNCTION OF X
30     SUBROUTINE INTGT(H,N,Y,SUM)
31     DIMENSION Y(')
32     REAL ITRAP,TSTMP
33     KT1=FLOAT(N)/2.+0.6
34     KT2=Y/2
35     IF(KT1-KT2) 10,40,10
C SIMPSON RULE
36     10 N1=N-1
37     YM1=Y(2)
38     DO 20 I=4,N1,2
39     YM1=YM1+Y(I)
40     20 CONTINUE
41     N2=N-2
42     YM2=Y(3)
43     DO 30 I=5,N2,2
44     YM2=YM2+Y(I)
45     30 CONTINUE
46     TSIIMP=(4/3.)*(Y(1)+4.*YM1+2.*YM2+Y(N))
47     SUM=TSIIMP
48     GO TO 60
C TRAPEZOIDAL RULE
49     40 YMID=Y(2)
50     N1=N-1
51     DO 50 I=3,N1
52     YMID=YMID+Y(I)
53     50 CONTINUE
54     ITRAP=(H/2.)*(Y(1)+2.*YMID+Y(N))
55     SUM=ITRAP
56     60 RETURN
57     END

```

## Appendix E

## Estimation of Air-to-fuel Ratio

$$\text{Air-to-fuel Ratio} = \frac{\text{air feed rate (by weight)}}{\text{coal feed rate}}$$

$$= \frac{\frac{PA_t U_o}{RT} M_{\text{air}}}{F_o}$$

$$= \frac{\frac{PA_t U_o}{RT F_o} M_{\text{air}}}{}$$

where  $P$  = air pressure

$A_t$  = bed cross section area

$U_o$  = superficial velocity

$M_{\text{air}}$  = molecular weight of air

$R$  = gas constant

$T$  = air temperature

$F_o$  = coal feed rate

## Appendix F

## Estimation of Excess Air

$$\begin{aligned}
 \text{Excess Air} &= \left( \left( \frac{\text{mole of air used}}{\text{mole of carbon}} \right) / \left( \frac{\text{mole of air}}{\text{mole of carbon}} \right)_{\text{theory}} - 1 \right) \times 100 \\
 &= \frac{\left( \frac{\text{mole of air used}}{\text{mole of Carbon}} \right)_{\text{actual}} - 1}{\left( \frac{\text{mole of oxygen} \times \frac{1}{0.21}}{\text{mole of carbon}} \right)_{\text{theory}}} \times 100 \\
 &= \left( \frac{PV}{RT \times \text{mole of carbon}} \times 0.21 - 1 \right) \times 100 \\
 &= \left( 0.21 \frac{PA_t U_o}{RT} \times \frac{\text{molecular weight of carbon}}{\text{weight of carbon / time}} - 1 \right) \times 100 \\
 &= \left( 0.21 \frac{PA_t U_o M_c}{RTYF_o} - 1 \right) \times 100
 \end{aligned}$$

where P = air pressure

$A_t$  = bed cross section area

$U_o$  = superficial velocity

$M_c$  = carbon molecular weight

R = gas constant

T = air temperature

$F_o$  = coal feed rate

Y = percentage of carbon in coal

## Appendix G

## Fuel Mixture Feed Rate



The fuel mixture in these experiments mentioned before is fed with screw feeder which is controlled by a variable speed motor.

The fuel mixture feed rates and the speed numbers are shown in Table G.1 and Table G.2

Table G.1

## Fuel Mixture Feed Rate

Ratio of limestone and lignite in fuel mixture = 1:1 by weight

$\bar{d}_p$  of coal = 1 mm.

$\bar{d}_p$  of limestone = 0.5 mm.

motor speed No.	fuel mixture feed rate	
	gm/sec	gm/min.
1	1.5	87.3
2	2.0	118.1
3	2.7	163.4
4	3.6	214.0
5	4.6	277.1
6	5.2	311.3

$\bar{d}_p$  = average diameter of particles

Table G.2

## Fuel Mixture Feed Rate

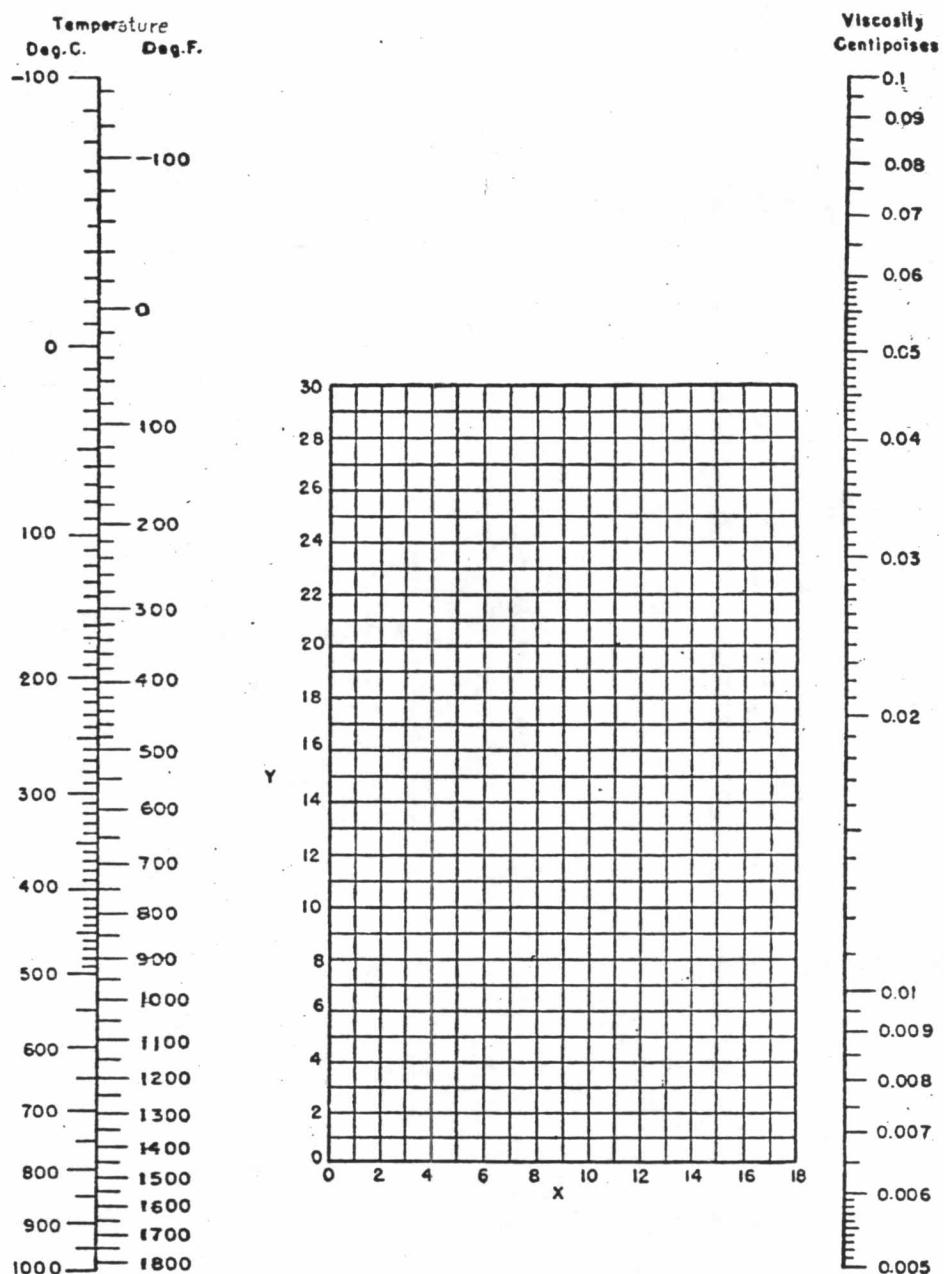
Ratio of limestone and lignite in fuel mixture = 1:1 by weight

 $\bar{d}_p$  of coal = 2.4 mm. $\bar{d}_p$  of limestone = 1 mm.

motor speed No.	fuel mixture feed rate	
	gm/sec	gm/min.
1	1.2	71.6
2	1.6	98.2
3	2.4	144.6
4	3.3	200.6
5	4.3	259.4
6	5.1	303.3
7	5.4	325.6

 $\bar{d}_p$  = the average diameter of particles

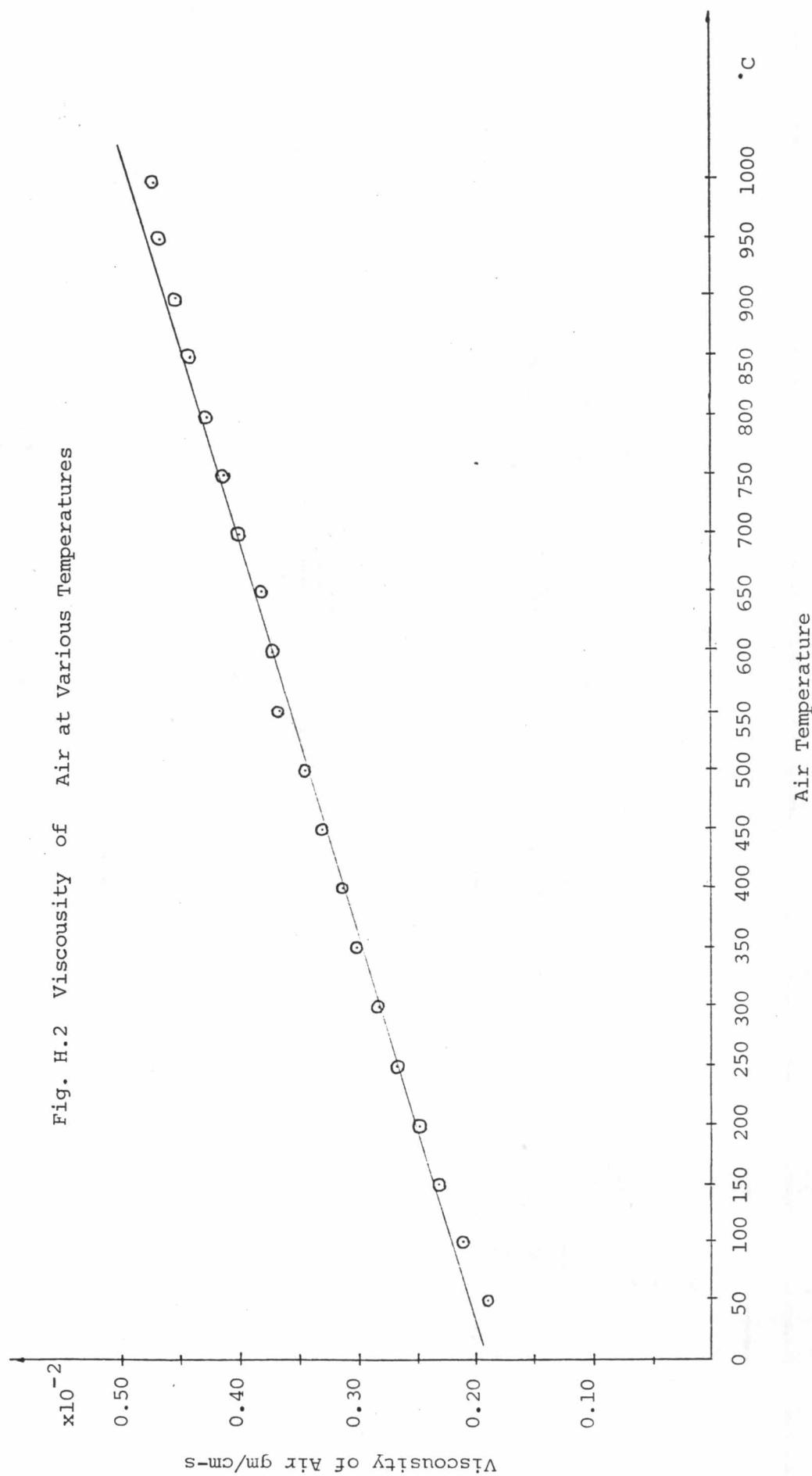
Appendix H  
37  
Viscosity



Viscosities of gases at 1 atm. For coordinates, see

Coordinates for use with Fig.											
No.	Gas	X	Y	No.	Gas	X	Y	No.	Gas	X	Y
1	Acetic acid	7.7	14.3	15	Chloroform	8.9	15.7	29	Freon-113	11.3	14.0
2	Acetone	5.9	13.0	16	Cyanogen	9.2	15.2	30	Helium	10.9	20.5
3	Acetylene	9.5	14.9	17	Cyclohexane	9.2	12.0	31	Hexane	8.6	11.5
4	Air	11.0	20.0	18	Ethane	9.1	14.5	32	Hydrogen	11.2	12.4
5	Ammonia	5.4	16.0	19	Ethyl acetate	8.5	13.2	33	$3H_2 + 1N_2$	11.2	17.2
6	Argon	10.5	22.4	20	Ethyl alcohol	9.2	14.2	34	Hydrogen bromide	6.5	20.9
7	Benzene	5.5	13.2	21	Ethyl chloride	8.5	15.6	35	Hydrogen chloride	5.8	18.7
8	Bromine	5.9	19.2	22	Ethyl ether	8.9	13.0	36	Hydrogen cyanide	9.5	14.9
9	Butene	9.2	13.7	23	Ethylene	9.5	15.1	37	Hydrogen iodide	9.0	21.3
10	Butylene	5.9	13.0	24	Fluorine	7.3	23.8	35	Hydrogen sulfide	8.6	18.0
11	Carbon dioxide	9.5	18.7	25	Freon-11	10.6	15.1	39	Iodine	9.0	18.4
12	Carbon disulfide	5.0	16.0	26	Freon-12	11.1	16.0	40	Mercury	5.3	22.9
13	Carbon monoxide	11.0	20.0	27	Freon-21	10.8	15.3	41	Methane	9.9	15.5
14	Chlorine	9.0	18.4	28	Freon-22	10.1	17.0	42	Methyl alcohol	8.5	15.6
									Xenon	9.3	23.0

Fig H.1 Viscosity of Gases



## Appendix I

Diffusivity<sup>38</sup>

For low pressure of binary gas mixture and high temperature, the diffusivity of oxygen in air is expressed in the following equation 32 , 33

$$D_{AB} = \frac{0.001858 T^{3/2} (1/M_A + s/M_B)^{1/2}}{P \sigma_{AB}^2 \Omega_D} \text{ cm}^3/\text{s} \quad \dots \dots \quad (1)$$

where  $D_{AB}$  = Mutual diffusion coefficient species A in binary of A and B,  $\text{cm}^2/\text{s}$

$P$  = total pressure = 1 atm.

$M_A$  = Molecular weight of A

here, the molecular weight of oxygen = 32

$M_B$  = Molecular weight of B,

here, the molecular weight of air = 28.84

$\sigma$  = Lennard-Jones potential parameter, "A (see Table

Table 1.1 LENNARD-JONES POTENTIAL PARAMETERS

Molecule	Compound	$\sigma, \text{Å}$	$\epsilon/k, ^\circ\text{K}$
A	Argon	3.542	93.3
He	Helium†	2.551	10.22
Kr	Krypton	3.655	178.9
Ne	Neon	2.820	32.8
Xe	Xenon	4.082	206.9
Air	Air	3.711	78.6
Br <sub>2</sub>	Bromine	4.296	507.9
CCl <sub>4</sub>	Carbon tetrachloride	5.947	322.7
CF <sub>4</sub>	Carbon tetrafluoride	4.662	134.0
CHCl <sub>3</sub>	Chloroform	5.389	340.2
CH <sub>2</sub> Cl <sub>2</sub>	Methylene chloride	4.898	356.3
CH <sub>3</sub> Br	Methyl bromide	4.118	449.2
CH <sub>3</sub> Cl	Methyl chloride	4.182	350.0
CH <sub>3</sub> OH	Methanol	3.626	481.8
CH <sub>4</sub>	Methane	3.758	148.6
CO	Carbon monoxide	3.690	91.7
COS	Carbonyl sulfide	4.130	336.0
CO <sub>2</sub>	Carbon dioxide	3.941	195.2
CS <sub>2</sub>	Carbon disulfide	4.483	467.0
C <sub>2</sub> H <sub>2</sub>	Acetylene	4.033	231.8
C <sub>2</sub> H <sub>4</sub>	Ethylene	4.163	224.7
C <sub>2</sub> H <sub>6</sub>	Ethane	4.443	215.7
C <sub>2</sub> H <sub>5</sub> Cl	Ethyl chloride	4.898	300.0
C <sub>2</sub> H <sub>5</sub> OH	Ethanol	4.530	362.6
C <sub>2</sub> N <sub>2</sub>	Cyanogen	4.361	348.6
CH <sub>3</sub> OCH <sub>3</sub>	Methyl ether	4.307	395.0
CH <sub>2</sub> CHCH <sub>3</sub>	Propylene	4.678	298.9
CH <sub>3</sub> CCH	Methylacetylene	4.761	251.8
C <sub>3</sub> H <sub>6</sub>	Cyclopropane	4.807	248.9
C <sub>3</sub> H <sub>8</sub>	Propane	5.118	237.1
n-C <sub>3</sub> H <sub>7</sub> OH	n-Propyl alcohol	4.549	576.7
CH <sub>3</sub> COCH <sub>3</sub>	Acetone	4.600	560.2
CH <sub>3</sub> COOCH <sub>3</sub>	Methyl acetate	4.936	469.8
n-C <sub>4</sub> H <sub>10</sub>	n-Butane	4.687	531.4
iso-C <sub>4</sub> H <sub>10</sub>	Isobutane	5.278	330.1
C <sub>2</sub> H <sub>5</sub> OC <sub>2</sub> H <sub>5</sub>	Ethyl ether	5.678	313.8
CH <sub>3</sub> COOC <sub>2</sub> H <sub>5</sub>	Ethyl acetate	5.205	521.3
n-C <sub>5</sub> H <sub>12</sub>	n-Pentane	5.784	341.1
C(CH <sub>3</sub> ) <sub>4</sub>	2,2-Dimethylpropane	6.464	193.4
C <sub>6</sub> H <sub>6</sub>	Benzene	5.349	412.3
C <sub>6</sub> H <sub>12</sub>	Cyclohexane	6.182	297.1
n-C <sub>6</sub> H <sub>14</sub>	n-Hexane	5.949	399.3
Cl <sub>2</sub>	Chlorine	4.217	316.0
F <sub>2</sub>	Fluorine	3.357	112.6
HBr	Hydrogen bromide	3.353	449.0
HCN	Hydrogen cyanide	3.630	569.1
HCl	Hydrogen chloride	3.339	344.7
HF	Hydrogen fluoride	3.148	330.0
HI	Hydrogen iodide	4.211	288.7
H <sub>2</sub>	Hydrogen	2.827	59.7
H <sub>2</sub> O	Water	2.641	809.1
H <sub>2</sub> O <sub>2</sub>	Hydrogen peroxide	4.196	289.3
H <sub>2</sub> S	Hydrogen sulfide	3.623	301.1
Hg	Mercury	2.969	750.0
I <sub>2</sub>	Iodine	5.160	474.2
NH <sub>3</sub>	Ammonia	2.900	558.3
NO	Nitric oxide	3.492	116.7
NOCl	Nitrosyl chloride	4.112	395.3
N <sub>2</sub>	Nitrogen	3.798	71.4
N <sub>2</sub> O	Nitrous oxide	3.828	232.4
O <sub>2</sub>	Oxygen	3.467	106.7
PH <sub>3</sub>	Phosphine	3.981	251.5
SF <sub>6</sub>	Sulfur hexafluoride	5.128	222.1
SO <sub>2</sub>	Sulfur dioxide	4.112	335.4
SnBr <sub>4</sub>	Stannic bromide	6.388	563.7
UF <sub>6</sub>	Uranium hexafluoride	5.967	236.8

† Calculated from quantum-mechanical formulas.

$\sigma_{AB}$  = Lennard-Jones potential parameter of the mixture  
of A and B

$\Omega_D$  = collision integral (see Table 1.2 and Fig 1.1.)

T = gas temperature

The value of  $\sigma_{AB}$  can be calculated from

$$\sigma_{AB} = \frac{1}{2} (\sigma_A + \sigma_B) \dots \dots \quad (2)$$

A = oxygen

B = air

hence,  $\sigma_{AB} = \frac{1}{2} (3.711 + 3.467)$   
 $= 3.589$  for binary of oxygen and air  $\dots \dots \quad (3)$

Using the Table 1.1, the value of  $\epsilon/k$  can be found, and  
the value of  $\epsilon/k$  of the binary gas mixture can be calculated by

$$\frac{\epsilon_{AB}}{K} = \left( \frac{\epsilon_A}{K} \cdot \frac{\epsilon_B}{K} \right)^{\frac{1}{2}} \dots \dots \quad (4)$$

Knowing the value of  $\epsilon_{AB}/K$ , the collision integral can be  
found from Table 1.2

Table I.2 VALUES OF THE COLLISION INTEGRAL  $\Omega_b$  BASED ON THE LENNARD-JONES POTENTIAL†

$kT/\epsilon^*$	$\Omega_{b\ddagger}$	$kT/\epsilon$	$\Omega_b$	$kT/\epsilon$	$\Omega_b$
0.30	2.662	1.65	1.153	4.0	0.8836
0.35	2.476	1.70	1.140	4.1	0.8788
0.40	2.318	1.75	1.128	4.2	0.8740
0.45	2.184	1.80	1.116	4.3	0.8694
0.50	2.066	1.85	1.105	4.4	0.8652
0.55	1.966	1.90	1.094	4.5	0.8610
0.60	1.877	1.95	1.084	4.6	0.8568
0.65	1.798	2.00	1.075	4.7	0.8530
0.70	1.729	2.1	1.057	4.8	0.8492
0.75	1.667	2.2	1.041	4.9	0.8456
0.80	1.612	2.3	1.026	5.0	0.8422
0.85	1.562	2.4	1.012	6	0.8124
0.90	1.517	2.5	0.9996	7	0.7896
0.95	1.476	2.6	0.9878	8	0.7712
1.00	1.439	2.7	0.9770	9	0.7556
1.05	1.406	2.8	0.9672	10	0.7424
1.10	1.375	2.9	0.9576	20	0.6640
1.15	1.346	3.0	0.9490	30	0.6232
1.20	1.320	3.1	0.9406	40	0.5960
1.25	1.296	3.2	0.9328	50	0.5756
1.30	1.273	3.3	0.9256	60	0.5596
1.35	1.253	3.4	0.9186	70	0.5464
1.40	1.233	3.5	0.9120	80	0.5352
1.45	1.215	3.6	0.9058	90	0.5256
1.50	1.198	3.7	0.8998	100	0.5130
1.55	1.182	3.8	0.8942	200	0.4644
1.60	1.167	3.9	0.8888	400	0.4170

† From J. O. Hirschfelder, C. F. Curtiss, and R. B. Bird, "Molecular Theory of Gases and Liquids," John Wiley & Sons, Inc., New York, 1954.

‡ Hirschfelder uses the symbols  $T^*$  for  $kT/\epsilon$  and  $\Omega^{(1,1)*}$  in place of  $\Omega_b$ .

Now the value of  $\varepsilon_{AB}/K$  of oxygen in air =  $\varepsilon_{AB}/K$

$$= \frac{1}{2} \left( \frac{\varepsilon_A}{k} \cdot \frac{\varepsilon_B}{k} \right)^{\frac{1}{2}}$$

$$= \frac{1}{2} (78.6 + 106.7)^{\frac{1}{2}}$$

$$= 91.58$$

..... (5)

The collision integral is also the function of temperature.

The collision integral of oxygen in air as the function of temperature is shown in Fig. 1.1

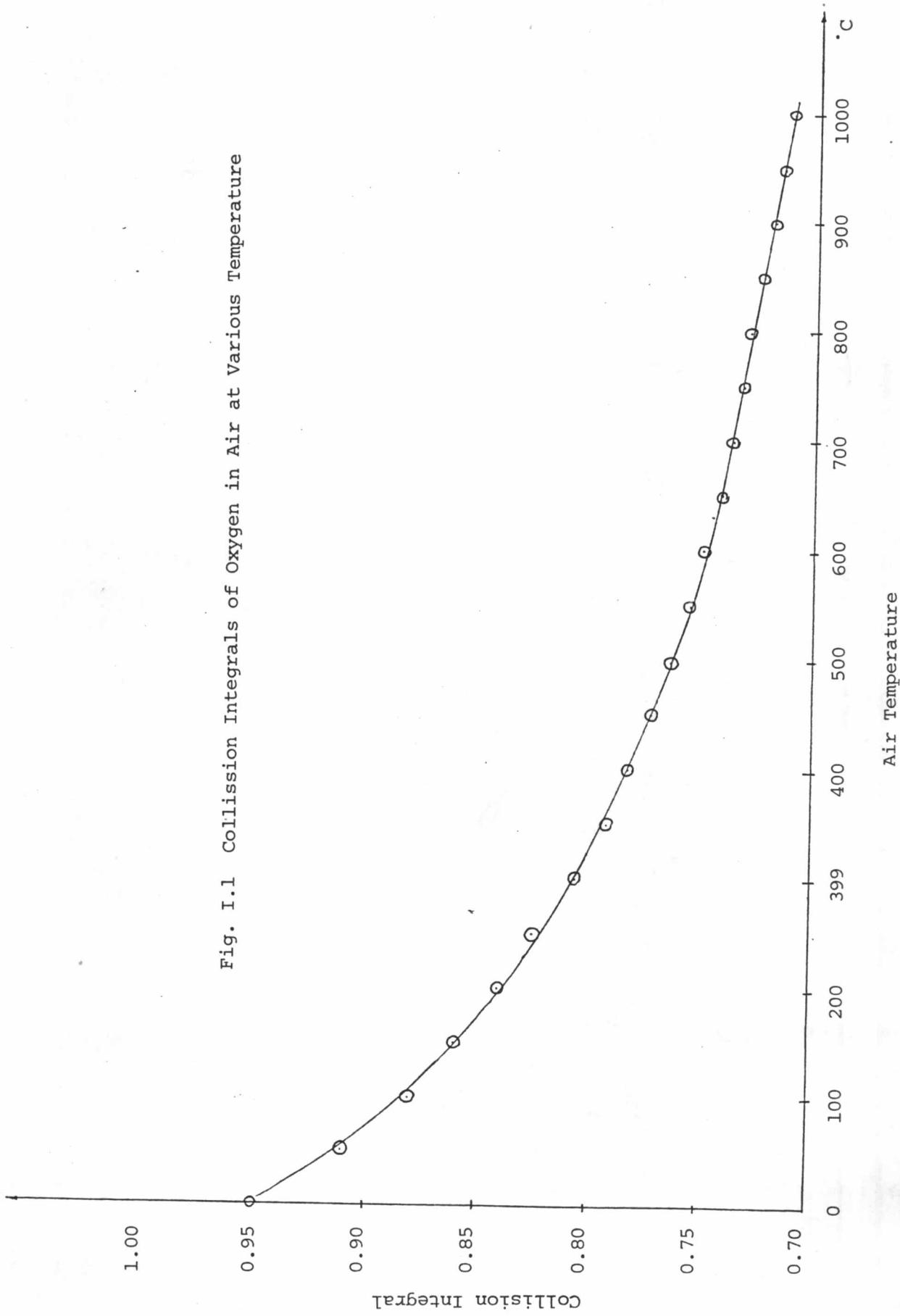
Hence, the diffusion of oxygen in air can be expressed in the following equation

$$D_{AB} = 3.73 \times 10^{-5} \frac{T^{3/2}}{\Omega_{AB}} \quad \dots \dots \dots \quad (6)$$

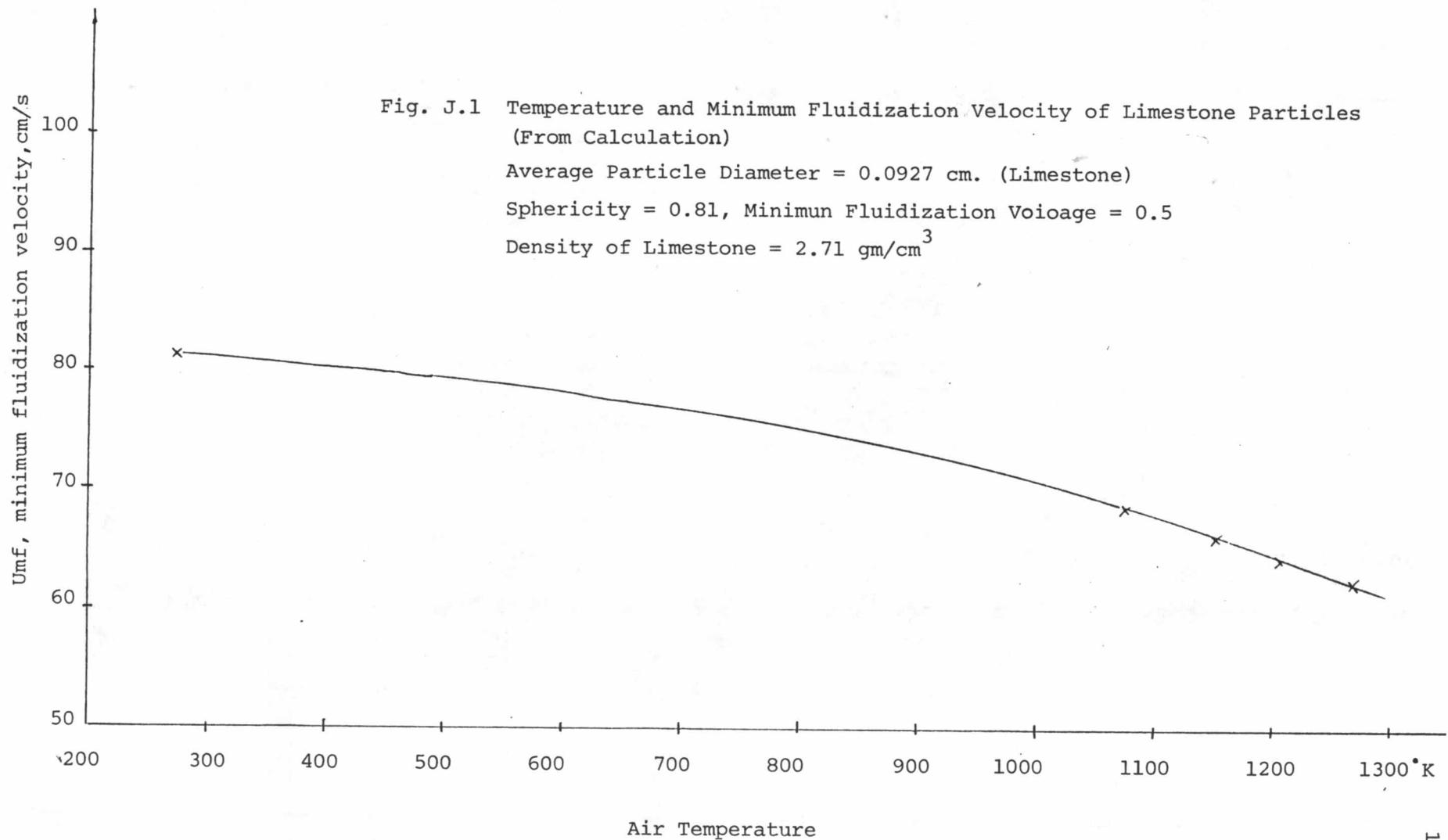
where A = oxygen

B = air

Fig. I.1 Collision Integrals of Oxygen in Air at Various Temperature

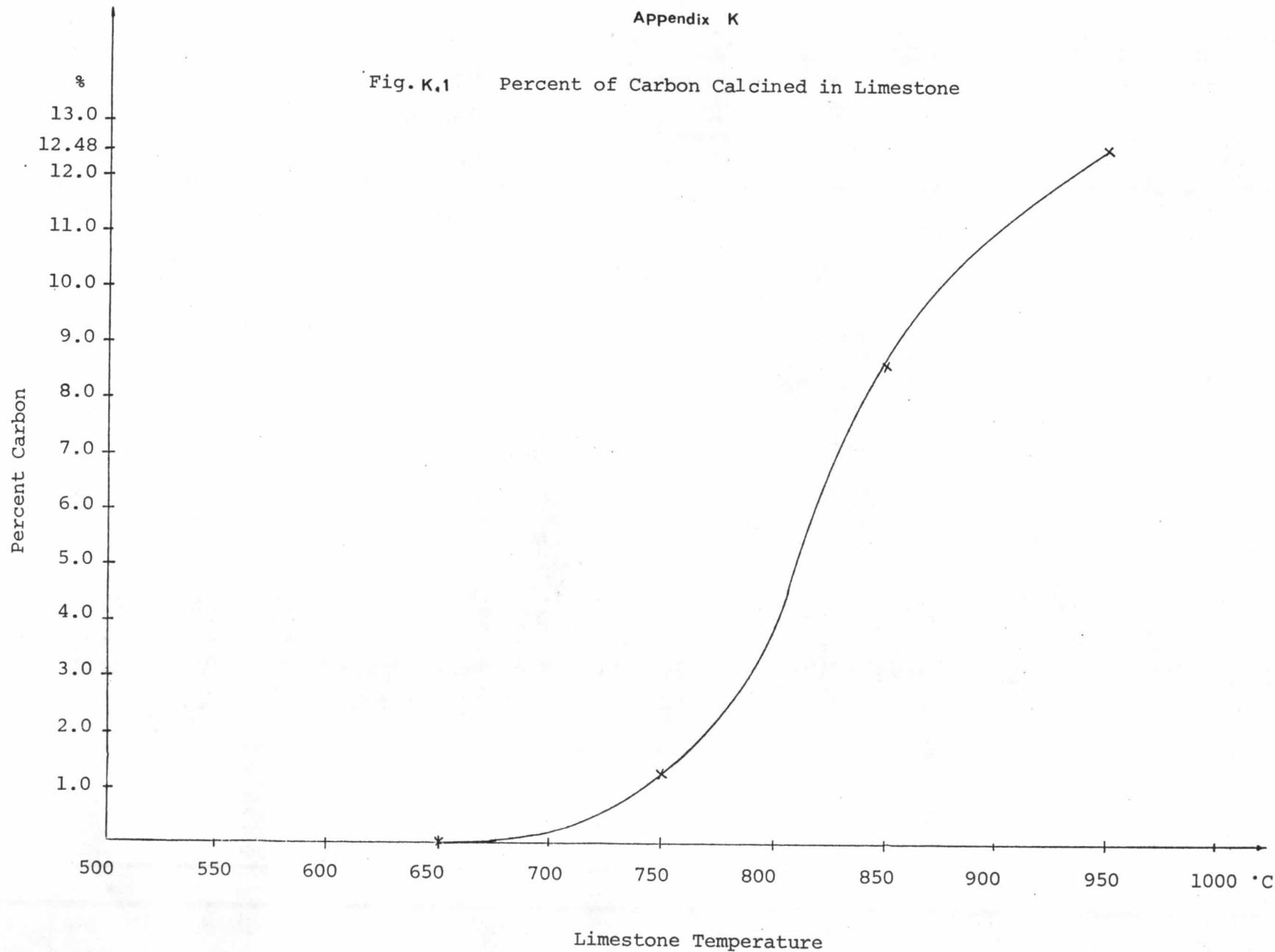


Appendix J



Appendix K

Fig. K.1 Percent of Carbon Calcined in Limestone



## Appendix L

### Approximate Heat Balance in the Combustor

The heat balance in the combustor can be estimated followingly.

Now an example of Calculation is shown

From data No. 3 of Table 6.1

$$\text{average bed temperature} = \bar{T}_B = 995^\circ C$$

$$\text{average gas temperature} = \bar{T}_G = 418^\circ C$$

$$\text{inlet limestone temperature} = 36^\circ C$$

$$\text{inlet lignite temperature} = 30^\circ C$$

$$\text{inlet air temperature} = 35^\circ C$$

$$\text{lignite feed rate} = F_{ocl} = 1.13 \text{ gm/s}$$

$$\text{limestone feed rate} = F_{ol} = 1.13 \text{ gm/s}$$

$$\text{specific heat of coal ash} = 0.31 \text{ cal/gm}^\circ C$$

$$\text{specific heat of limestone} = 0.217 \text{ cal/gm}^\circ C$$

$$\text{specific heat of air at } \bar{T}_G \text{ of } 418^\circ C = 0.26 \text{ cal/gm}^\circ C$$

$$\text{heating value of lignite} = 4166 \text{ cal/gm}$$

$$\text{air density at STP} = \rho_g = 1.293 \times 10^{-3} \text{ gm/cm}^3$$

$$\text{inlet air velocity} = 376 \text{ cm/s}$$

$$\text{ash content in lignite} = 10 \%$$

$$\text{bed diameter} = .15 \text{ cm}$$

#### Inlet

If carbon is all combusted, heat will

$$\text{be generated} = 4166 \times 1.13 \text{ cal/s}$$

$$= 4708 \text{ cal/s}$$

## Outlet

Due to the specific heat of CO, CO<sub>2</sub>, O<sub>2</sub> N<sub>2</sub> and air at the same temperature of 418°C is approximately the same.

Let the specific heat of outlet gas = 0.26  $\frac{\text{cal}}{\text{gn} \cdot \text{C}}$

heat accepted by outlet gas and limestone

$$= (418-35) \times 1.293 \times 10^{-3} \times \frac{273}{(273+418)} \times 376 \times \frac{4}{15 \times 15} \\ \times 0.26 + (995-30) \times 1.13 \times 0.217 \\ = 3617 \quad \text{cal/s}$$

heat accepted by coal ash

$$= 1.13 \times 0.10 \times (995-30) \times 0.31 \\ = 34$$

hence heat accepted by outlet solids and gas

$$= 365 \quad \text{cal/s}$$

the calculated outlet heat is less than the inlet potential  
heat = 4708 - 3651 = 1057 cal/s

This uncalculated heat can be lost by other way such as the conduction of heat through the refractory and stainless steel of combustor, heat consumed by the endothermic reaction of



## Appendix M

## Some Constant Values

density of limestone  $\pm 2.75 \text{ gm/cm}^3$

density of lignite  $= 1.50 \text{ gm/cm}^3$

density of air at STP  $= 1.293 \times 10^{-3} \text{ gm/cm}^3$

universal gas constant  $= 82.06 \text{ atm-cm}^3/\text{gmole}\cdot\text{K}$

$= 1.987 \text{ cal/gmole}\cdot\text{K}$

gravitational acceleration  $= 980 \text{ cm/s}^2$

sphericity of lime  $= 0.77$

void fraction of limestone of 0.5 mm. in average particles diameter

$= 0.513$

void fraction of limestone of 1.0 mm in average particles diameter

$= 0.45$

inlet oxygen concentration  $= 1.28 \times 10^{-3} \text{ kmole/m}^3$  at  $1173^\circ\text{K}$  and 1 atm.

$= 2.18 \times 10^{-6} \text{ gmole/cm}^3$  at  $1173^\circ\text{K}$  and 1 atm.

## Appendix N

### Output from Computer Simulation

From chapter 7 the graphs of effects of operating variables on carbon combustibles loss are plotted from the following data:

## N.1 The Effect of Superficial Velocity on Carbon Combustibles Loss

Let  $\frac{W}{W_B} = 0.1 \%$ ,  $C_{Ab} = 91.3 \text{ & } C_{Ao}$ ,  $\bar{T}_B = \text{constant} = 1173^\circ\text{K}$ ,  $F_o = \text{constant} = 1.13 \text{ gm/s}$

$\bar{d}_{p \text{ limestone}} = 0.1 \text{ cm}$ ,  $\bar{d}_{p \text{ lignite}} = 0.24 \text{ cm}$ ,  $\rho_g = 0.28 \times 10^{-3} \text{ gm/cm}^3$ ,  $\mu_g = 4.6 \times 10^{-4} \text{ gm/cm-s}$

$D_A = 2.08 \text{ cm}^3/\text{s}$ ,  $C_{Ao} = 2.18 \times 10^{-6} \text{ gmole/cm}^3$ ,  $U_{mf} = 65 \text{ cm/s}$  at  $1173^\circ\text{K}$ ,  $L_f = 30 \text{ cm}$

$d_t = 15 \text{ cm}$ , limestone : lignite = 1:1 by weight

NO	$U_o$ cm/s	X	$C_{Ap}$ gmole/cm <sup>3</sup>	$\bar{t}$ s	W gm	$F_1$ gm/s	$F_2$ gm/s	$\eta$	EXC
1	200	3.7	$1.98 \times 10^{-6}$	6.2	7.1	0.17	0.03	82.7	91.5
2	250	2.9	$1.97 \times 10^{-6}$	6.1	6.9	0.13	0.03	86.2	139.4
3	300	1.8	$1.95 \times 10^{-6}$	6.0	6.8	0.10	0.03	89.0	187.3
4	350	1.7	$1.948 \times 10^{-6}$	5.9	6.7	0.05	0.04	92.2	235.1
5	400	1.5	$1.94 \times 10^{-6}$	5.8	6.7	0.02	0.05	94.2	283.0
6	450	1.4	$1.92 \times 10^{-6}$	5.7	6.4	0.00	0.07	93.7	330.9

$W$  = weight of burning char in the bed

$\bar{t}$  = average residence time of burning char particles in bed

$W_B$  = weight of bed material

$\eta$  = combustion efficiency

$C_{Ab}$  = concentration of oxygen in bubble

EXC = excess air

$C_{Ao}$  = concentration of oxygen in inlet air

$\bar{d}_{p \text{ limestone}}$  = average particles diameter of limestone

$C_{Ap}$  = concentration of oxygen in particulate phase  $\bar{d}_{p \text{ coal}}$  = average particles diameter of coal

$\bar{T}_B$  = average bed temperature

$\rho_g$  = density of air

$F_o$  = coal feed rate

$\mu_g$  = viscosity of air

$F_1$  = overflow rate

$D_A$  = diffusivity of oxygen in air

$F_2$  = elutriation rate

$U_{mf}$  = minimum fluidizing velocity

$U_o$  = superficial velocity

$L_f$  = expanded bed height

X = number of transfer unit

$d_t$  = bed diameter

N. 2 The Effect of Bed Temperature on Carbon Combustibles Loss

Let  $\frac{W}{W_B} = 0.1 \%$ ,  $C_{Ab} = 91.3 \% C_{Ao}$ ,  $U_o = \text{constant} = 350 \text{ cm/s}$ ,  $F_o = 1.13 \text{ gm/s}$

$L_f = 30 \text{ cm}$ ,  $d_t = 15 \text{ cm}$ ,  $\bar{d}_p$  limestone =  $0.1 \text{ cm}$ ,  $\bar{d}_p$  lignite =  $0.74 \text{ cm}$

limestone: lignite = 1:1 by weight

NO.	$\bar{T}_B$ K	$D_A$ $\text{cm}^3/\text{s}$	$\mu_g$ $\text{gm/cm-s}$	$\rho_g$ $\text{gm/cm}^3$	$U_{mf}$ $\text{cm/s}$	X	$C_{Ao}$ $\text{gmols/cm}^3$	$C_{Ab}$ $\text{gmols/cm}^3$	$C_{Ap}$ $\text{gmols/cm}^3$	W gm	$F_1$ $\text{gm/s}$	$F_2$ $\text{gm/s}$	$\eta$ %	EXC %
1	1273	2.39	$4.7 \times 10^{-4}$	$2.6 \times 10^{-4}$	62	1.63	$2.0 \times 10^{-6}$	$1.8 \times 10^{-6}$	$1.8 \times 10^{-6}$	6.6	0.04	0.04	93.3	209
2	1223	2.24	$4.7 \times 10^{-4}$	$2.7 \times 10^{-4}$	64	1.67	$2.1 \times 10^{-6}$	$1.9 \times 10^{-6}$	$1.9 \times 10^{-6}$	6.6	0.05	0.04	92.0	221
3.	1173	2.09	$4.5 \times 10^{-4}$	$2.8 \times 10^{-4}$	65	1.71	$2.2 \times 10^{-6}$	$2.0 \times 10^{-6}$	$2.0 \times 10^{-6}$	6.5	0.06	0.04	91.0	235
4	1123	1.95	$4.4 \times 10^{-4}$	$2.9 \times 10^{-4}$	67	1.76	$2.3 \times 10^{-6}$	$2.1 \times 10^{-6}$	$2.0 \times 10^{-6}$	6.5	0.07	0.04	90.5	250
5	1073	1.81	$4.3 \times 10^{-4}$	$3.1 \times 10^{-4}$	68	1.78	$2.4 \times 10^{-6}$	$2.2 \times 10^{-6}$	$2.1 \times 10^{-6}$	6.5	0.09	0.04	89.1	266
6	1023	1.67	$4.1 \times 10^{-4}$	$3.2 \times 10^{-4}$	70	1.83	$2.5 \times 10^{-6}$	$2.3 \times 10^{-6}$	$2.2 \times 10^{-6}$	6.4	0.10	0.04	88.4	274

N.3 The Effect of Coal Feed Rate on Carbon Combustibles Loss

Let  $\frac{W}{W_B} = 0.1$ ,  $C_{Ab} = 91.3\% C_{Ao}$ ,  $\bar{T}_B = \text{constant} = 1173^\circ\text{K}$ ,  $U_o = \text{constant} = 400 \text{ cm/s}$

$\bar{d}_p$  limestons = 0.1 cm,  $\bar{d}_p$  lignite = 0.24 cm,  $\rho_g = 0.28 \times 10^{-3}$ ,  $\mu_g = 4.6 \times 10^{-4} \text{ gm/cm-s}$ ,  $d_t = 15 \text{ cm}$

$D_A = 2.08 \text{ cm}^3/\text{s}$ ,  $C_{Ao} = 2.18 \times 10^{-6} \text{ gmole/cm}^3$ ,  $U_{mf} = 6.5 \text{ cm/s at } 1173^\circ\text{K}$ ,  $L_f = 30 \text{ cm}$

limestone : lignite = 1:1 by weight

NO.	$F_o$ gm/s	X	$C_{Ap}$ gmole/cm <sup>3</sup>	W gm	$\bar{t}$ s	$F_1$ gm/s	$F_2$ gm/s	$\eta$ %	EXC
1	0.5	1.5	$1.94 \times 10^{-6}$	2.87	13.0	0.00	0.03	94.8	767
2	1.0	"	"	5.77	6.6	0.00	0.05	95.4	333
3	1.5	"	"	6.56	4.4	0.43	0.03	69.3	188
4	2.0	"	"	"	3.3	0.99	0.01	50.4	117
5	2.5	"	"	"	2.6	1.53	0.00	40.0	73
6	3.0	"	"	"	2.2	2.04	0.00	33.5	44

### Bibliography

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