ORIGINAL RESEARCH



Study on coal gasification with soot formation in two-stage entrained-flow gasifier

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Abstract Concerning the global warming due to large CO₂ emission, the efficient use of coal becomes important for getting sustainable energy production. Coal gasification under CO₂-rich condition is expected to be an effective way to produce a concentrated and pressurized carbon dioxide stream, resulting in reduction in separation energy of CO₂ for CCS. Moreover, the soot formation, which is of significant environmental concern, is still being neglected in the past studies of coal gasification. A one-step soot formation reaction mechanism is proposed in this study and implemented in numerical simulations of coal gasification with the aim of describing the gasification behaviors in a two-stage entrained-flow gasifier. In addition, the effects of O₂ ratio and CO₂ concentration on soot concentration, syngas heating value and carbon conversion are numerically studied in an effort to increase the syngas production. The Eulerian-Lagrangian approach is applied to solve the Navier-Stokes equation and the particle dynamics. Finite rate/eddy dissipation model is used to calculate the rate of nine homogeneous gas-to-gas phase reactions including soot formation and soot oxidation. While only finite rate is used for the heterogeneous solid-to-gas phase reactions. It is found that formation of

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soot enhances the H_2 production in the gasifier. Carbon conversion gradually increases with an increase in O_2 ratio, while producing a low heating value syngas beyond a certain limit of O_2 ratio. In contrast, an increase in CO_2 concentration in the gasifier increases heating value of product syngas.

Keywords Coal gasification \cdot Soot formation \cdot Two-stage gasifier \cdot Syngas \cdot CO₂

List of symbols

a	Absorption co-efficient (m^{-1})
a_p	Equivalent absorption co-efficient (m^{-1})
À	Surface area (m ²)
A_{f}	Pre-exponential factor (kg/m ² s Pa), (s ^{-1})
$A_{\mathbf{R}}$	Magnussen constant for reactants (-)
B_P	Magnussen constant for products (-)
c_p	Specific heat of gas (J/kg K)
$C_{\rm P}$	Specific heat of coal particle (J/kg K)
d	Diameter (m)
$D_{\rm k}$	Diffusion co-efficient in <i>k</i> th reaction (m^2/s)
Ε	Energy (J)
$E_{\rm p}$	Equivalent emission of coal particles (W/m ³)
$f_{\rm p}$	Particle scattering factor (-)
$f_{\rm w}$	Fraction of water present in coal particles (-)
$f_{\rm h}$	Fraction of heat absorbed by coal particles (-)
g	Gravitational acceleration (m/s ²)
h	Heat transfer co-efficient (W/m ² K)
H	Enthalpy (J/kg)
$H_{\rm comb}$	Height of combustor (m)
Ι	Number of species (-)
<i>I</i> _{rad}	Radiation intensity (W/m ²)
$I_{\rm t}$	Turbulent intensity (-)
J_i	Mass flux of species i (kg/m ² s)
k	Turbulent kinetic energy (m ² /s ²)



k.	Reaction rate constant (unit vary)			
K K	Number of reactions (_)			
I	Latent heat of water present in coal (I/kg-coal)			
	Mass (kg)			
111 122	Mass of coal particle (kg)			
$m_{\rm p}$	Molecular weight of species <i>i</i> (kg/kmol)			
NI i	Notecular weight of species i (kg/killor)			
IV D	Drassura (Da)			
p	Pressure (Fa)			
r D	Position vector (iii) Universal ass constant (8.214 \times 10 ³) (Ultral K)			
К D	Universal gas constant $(8.314 \times 10^{\circ})$ (J/kmol K)			
<i>K</i> _i	Source of chemical species t due to reaction (kg/m ³ s)			
$\hat{R}^{(A)}_{i,k}$	Rate of production (Arrhenius) of species <i>i</i> in <i>k</i> th reaction ($kmol/m^3$ s)			
$\hat{\boldsymbol{\rho}}(R)$	Rate of production (Eddy dissipation) of reactant			
$\mathbf{K}_{i,k}$	<i>i</i> in <i>k</i> th reaction (kmol/m ³ s)			
$\hat{\boldsymbol{\rho}}^{(P)}$	Rate of production (Eddy dissipation) of product			
$\mathbf{K}_{i,k}$	<i>i</i> in <i>k</i> th reaction (kmol/m ³ s)			
\bar{R}_{L}	Rate of particle surface species depletion in kth			
- 1 _K	reaction (kg/s)			
\tilde{R}_{L}	Rate of particle surface species reaction per unit			
ιĸ	area in kth reaction (kg/m ² s)			
Red	Revnolds number based on the particle diameter			
u	(-)			
S	Path length (m)			
\vec{s}	Direction vector (m)			
Sm	Rate of mass added from coal particle $(kg/m^2 s)$			
Sh reac	Source of heat due to reaction $(W/m^2 s)$			
t	Time (s)			
Т	Temperature (K)			
u, v, w	Velocity magnitude (m/s)			
\vec{v}	Velocity vector (m/s)			
$\overline{u_i}$	Mean velocity component			
u':	Fluctuating velocity component			
$\stackrel{'}{V}$	Volume (m ³)			
X_i	Molar concentration of species i (kmol/m ³)			
v^{+}	Distance (–)			
Y_i	Mass fraction of species i (–)			
Z	Height of reactor (m)			
-				
Greek letters				

Greek letters

- Yield parameter for first step devolatilization (-) α_1 Yield parameter for second step devolatilization (-) α_2 8 Turbulent dissipation rate (m^2/s^3) Emissivity of coal particle (-) εp η', η'' Rate exponent for reactants, products (-) v', v'' Stoichiometric co-efficient for reactants, products (-) $\theta_{\rm R}$ Radiation temperature (K) Dynamic viscosity (Pa s) μ Turbulent viscosity (Pa s) µ, Density (kg/m^3) ρ
- $\sigma \qquad \begin{array}{l} \text{Stefan-Boltzmann constant (5.669 \times 10^{-8})} \\ (\text{W/m}^2 \text{ K}^4) \end{array}$

- σ_k Turbulent Prandtl number for k (–)
- σ_{ε} Turbulent Prandtl number for ε (–)
- $\sigma_{\rm s}$ Scattering co-efficient (m⁻¹)
- $\sigma_{\rm p}$ Equivalent particle scattering factor (m⁻¹)
- Ω Solid angle (°)

Subscripts

- a Ash
- ac Activation
- b Backward
- f Forward i Species
- Species
- h Heat
- m Mass
- P Product species
- p Particles
- R Reactant species
- rad Radiation t Turbulent
- 0 Initial stage

Introduction

Global energy consumption in 2030 is predicted to increase 1.4 times than that in 2007, where about half of the increase will be contributed by Asia. It is also predicted that remaining years of exploitable global energy resources in sequences are: coal (122 years), uranium (100 years), natural gas (60 years) and oil (42 years) [1]. Because of more exploitable coal resource compared to other resources, it is expected that coal will continue to play a significant role in meeting the future energy demand. However, due to use of fossil fuel mainly coal to generate power, large amounts of CO₂ is discharged from conventional coal fired power plant, which is deemed as one of the major causes of global warming. Although technologies for employing renewable energy such as solar, wind, ocean, hydro, and biomass have been developed, the advantage of utilizing fossil fuels (mainly coal) for providing the most affordable electrical energy cannot be replaced overnight by any other technologies today [2]. However, clean coal technologies need to be implemented in the power sector in an effort to meet the environmental targets.

The clean coal technology field is moving in the direction of coal gasification with a second stage so as to produce a concentrated and pressurized carbon dioxide stream followed by carbon sequestration, including the capture and storage of carbon dioxides. However, CO_2 concentration in the conventional coal–air combustion flue gas is too low for carbon sequestration to be considered economically feasible. Recycling CO_2 in coal gasification process with the addition of oxygen will further increase CO_2 concentration in the flue gas. Flue gases with CO_2 concentration higher than 90 % can also be economically used for deep sea CO_2 storage and enhanced oil recovery [3]. This technology has the potential to provide what may be called "zero emissions"—in reality, extremely low emissions of the conventional coal pollutants, and as low-as-engineered carbon dioxide emissions. This has come about as a result of the realization that efficiency improvements, together with the use of natural gas and renewable such as wind will not provide the deep cuts in greenhouse gas emissions necessary to meet future national targets.

There are only few studies on coal gasification in twostage entrained-flow gasifier found in World Wide Web. Moreover, no study on coal gasification under CO2-rich condition in two-stage entrained-flow gasifier is found. Chen et al. [4, 5] performed a series of numerical simulations under various operating conditions for a two-stage air blown entrained-flow gasifier. It was reported that increasing air ratio leads to increased CO₂ and decreased CO and H₂ concentrations, and accordingly, had a strong effect on the heating value of the product gas. The effect of air/coal partitioning to the two stages, and the feed rate of recycle char was found to be limited. Silaen et al. [6] conducted numerical simulation of coal gasification process inside a two-stage entrained-flow coal gasifier. They reported that smaller particles produced more CO and less CO₂ which result in an increased syngas heating value. Luan et al. [2] studied the simulation of the coal combustion and gasification processes in a two-stage entrainedflow gasifier using the finite rate model for heterogeneous reactions. They reported that the increased O₂/coal ratio leads to higher exit temperature and CO₂ concentration, but lower CO concentration, resulting in a decrease of syngas heating value. However, the soot formation, which is of significant environmental concern, is still being neglected in the past studies of coal gasification [2, 4-6]. Soot formation has been observed in many pulverized coal utilization processes, including coal gasification and combustion. The formation of soot during coal gasification causes substantial heat losses due to radiative heat transfer. Therefore, an understanding of soot formation and its mechanism is necessary for the better design of coal gasification systems.

The main objectives of this study are to conduct numerical simulation including one-step soot formation mechanism in coal gasification and to discuss the effect of soot formation on the outcome of the simulation. In addition, a number of numerical simulations under O_2 -rich and CO_2 -rich gasification condition are carried out in an effort to increase the syngas production. The numerical results obtained from this study are considered to be an important step towards better designs of gasifiers.



Fig. 1 Schematic of computational model adopted from CRIEPI [7]

Numerical methods

Computational domain

The coal gasifier (Fig. 1) considered here consists of a combustor stage and a reductor stage. Coal and char are injected into the combustor stage with O2-rich gas mixtures. The gasifier has two levels of injectors that are positioned axisymmetrically at combustor and reductor stage. The combustor injectors are placed similar to a tangential firing system to create swirling flow inside the gasifier. The reductor injectors are directed towards the center of the gasifier. The diameter of the coal/char inlet zone is <10 mm which is very small compared to the height of the gasifier (4.94 m). Thus, we make the mesh with various size ranges, from 2 to 10 mm. Moreover, making a uniform mesh with 2 mm size will significantly increase the computational time. A three-dimensional mesh consisting of 247,818 computational cells is used with the small cell size being around 2 mm and the largest one around 10 mm. The near wall y^+ value is 250, which is appropriate $(30 > y^+ > 300)$ to apply the standard wall functions in the standard $k-\varepsilon$ turbulence model.

Soot formation

Soot formation in coal gasification is a very complicated process. This is due to the fact that the molecules of coal volatiles, particularly polycyclic aromatic hydrocarbons (PAHs), are much larger and more chemically diverse than those of simple hydrocarbon fuels. There are several soot models that have been proposed during the recent decades.



Some of the most important empirical models are Khan and Greeves model [8], Tesner model [9] and Lindstedt model [10]. There are also some detailed models that take complex physical phenomena and detailed chemistry into account. One of the most comprehensive detailed models is that proposed by Frenklach [11]. A detailed kinetic model describing the formation and consumption of PAHs and soot in hydrocarbon combustion has also been developed by Richter et al. [12]. Although the detailed models have undergone remarkable development recently, these models are still computationally demanding and cannot be used for complex geometries. In our previous work [13], we investigated soot formation model in a plug flow reactor (PFR) and reported that the following two reactions are typically considered to be the main reaction path in soot formation: (a) particle nucleation—PAH of increasing size are mainly formed by sequence of chemical reactions between PAH and their radicals, and between PAH radicals. This process is repeated producing large PAHs that form soot particles [12, 13] and (b) soot/PAH oxidation—reaction of soot/PAH with oxygen/hydroxyl radicals that depletes PAH/soot [13]. Corresponding to these concepts, a one-step soot formation mechanism is proposed in the present study to investigate the effect of soot formation on product gas concentration and gas temperature in coal gasification. A schematic of soot formation mechanism is shown in Fig. 2. In the onestep soot formation mechanism, an aromatic hydrocarbon molecule benzene (C_6H_6) or naphthalene $(C_{10}H_8)$ or phenanthrene ($C_{14}H_{10}$) or pyrene ($C_{16}H_{10}$) is considered as a precursor of soot formation. Before introducing the soot mechanism, the calculated results obtained using one-step soot formation mechanism are compared with those obtained using detailed soot formation mechanism under various gasification conditions [13].

Governing equations

For the fluid phase, the steady-state Reynolds Averaged Navier–Stokes (RANS) equations as well as the mass and



Fig. 2 Schematic of soot formation mechanism

energy conservation equations are solved for two-stage entrained-flow coal gasifier shown in Fig. 1. The governing equations for the conservation of mass, momentum, energy and species in 3D Cartesian coordinates are given as follows:

Continuity:
$$\nabla \cdot (\rho \vec{v}) = S_{\rm m}$$
 (1)

Momentum: $\nabla \cdot (\rho \vec{v} \vec{v}) = -\nabla p + \nabla \cdot (\overline{\overline{\tau}}) + \rho \vec{g} + \vec{F}$ (2)

Energy:
$$\nabla \cdot (\vec{v}(\rho E + p)) = -\nabla \cdot \left(\sum_{i} H_{i}J_{i}\right) + I_{\text{rad}} + S_{\text{h,reac}}$$
 (3)

Species:
$$\nabla \cdot (\rho \vec{v} Y_i) = -\nabla \cdot \vec{J}_i + R_i + S_i$$
 (4)

Turbulent flows are characterized by fluctuating velocity fields. Turbulence models seek to solve a modified set of transport equations by introducing averaged and fluctuating components. In Reynolds averaging, the solution variables in the instantaneous Navier–Stokes equations are decomposed into the mean (time-averaged) and fluctuating components. For the velocity component:

$$u_i = \overline{u_i} + u_i',\tag{5}$$

where $\overline{u_i}$ and u'_i are the mean and fluctuating velocity components (i = 1, 2, 3). A standard $k-\varepsilon$ model [14–16] is used to solve the turbulence. The turbulence kinetic energy, k, and its rate of dissipation, ε , are obtained from the following transport equations:

$$\frac{\partial}{\partial x_i}(\rho k u_i) = \frac{\partial}{\partial x_j} \left[\left(\mu + \frac{\mu_i}{\sigma_k} \right) \frac{\partial k}{\partial x_j} \right] + G_k - \rho \varepsilon, \tag{6}$$

$$\frac{\partial}{\partial x_i}(\rho \varepsilon u_i) = \frac{\partial}{\partial x_j} \left[\left(\mu + \frac{\mu_t}{\sigma_{\varepsilon}} \right) \frac{\partial \varepsilon}{\partial x_j} \right] + C_{1\varepsilon} G_k \frac{\varepsilon}{k} - C_{2\varepsilon} \rho \frac{\varepsilon^2}{k}, \quad (7)$$

where G_k represents the generation of turbulence kinetic energy related to the mean velocity gradient, and μ_t is the turbulent viscosity. The turbulent model constants are $C_{1\varepsilon} = 1.44$, $C_{2\varepsilon} = 1.92$, $C_{\mu} = 0.09$, $\sigma_k = 1.0$, and $\sigma_{\varepsilon} = 1.3$ [15, 16].

The discrete ordinates (DO) radiation model is used to solve the radiative heat transfer equation. The DO radiation model considers the radiative transfer equation as:

$$\frac{\mathrm{d}I_{\mathrm{rad}}(\vec{r},\vec{s})}{\mathrm{d}s} = -\left(a + a_{\mathrm{p}} + \sigma_{\mathrm{p}}\right)I_{\mathrm{rad}}(\vec{r},\vec{s}) + E_{\mathrm{p}} + a\phi^{2}\frac{\sigma T^{4}}{\pi} + \frac{\sigma_{s}}{4\pi}\int_{0}^{4\pi}I_{\mathrm{rad}}(\vec{r},\vec{s}')\Phi(\vec{s}\cdot\vec{s}')\mathrm{d}\Omega.$$
(8)

In discrete phase modeling, pulverized coal particles are injected into the gasifier and tracked throughout the computational domain using a Lagrangian approach. The continuity and momentum of particles are expressed as follows:

$$\frac{\mathrm{d}m_{\mathrm{p}}}{\mathrm{d}t} = \dot{m},\tag{9}$$

$$\frac{\mathrm{d}u_{\mathrm{p}}}{\mathrm{d}t} = F_{\mathrm{D}}(u - u_{\mathrm{p}}) + \frac{g(\rho_{\mathrm{p}} - \rho)}{\rho_{\mathrm{p}}}.$$
(10)

 $F_{\rm D}(u-u_p)$ is the drag force per unit particle mass and $F_{\rm D}$ is determined from

$$F_{\rm D} = \frac{18\mu}{\rho_{\rm p} d_{\rm p}^2} \frac{C_D \mathrm{Re}_{\rm d}}{24}.$$
(11)

 Re_d is the relative Reynolds number based on the particle diameter and relative velocity.

The change of particle temperature during devolatilization is determined from the energy balance of particles governed by convective, latent heat and radiative heat transfer as follows [17, 18].

$$m_{\rm p}C_{\rm p}\frac{\mathrm{d}T_{\rm p}}{\mathrm{d}t} = hA_{\rm p}(T-T_{\rm p}) + \frac{\mathrm{d}m_{\rm p}}{\mathrm{d}t}L + A_{\rm p}\varepsilon_{\rm p}\sigma\Big(\theta_{\rm R}^4 - T_{\rm p}^4\Big).$$
(12)

After the volatile species of the coal particle has evolved completely, char $-O_2$, char $-CO_2$ and char $-H_2O$ surface reactions begin. During surface reaction, the following heat balance equation is used:

$$m_{\rm p}C_{\rm p}\frac{\mathrm{d}T_{\rm p}}{\mathrm{d}t} = hA_{\rm p}(T-T_{\rm p}) - f_{\rm h}\left(\frac{\mathrm{d}m_{\rm p}}{\mathrm{d}t}\right)\Delta H + A_{\rm p}\varepsilon_{\rm p}\sigma\left(\theta_{\rm R}^4 - T_{\rm p}^4\right).$$
(13)

Reaction models

When the temperature of the coal particles reaches the vaporization temperature (600 K), chemical reactions occur producing various amounts of gases, tar, and coke. The tar and gases are usually referred as volatiles. The volatiles are released according to Kobayashi model [19]. This model assumes two kinetic rates, $k_{kin,1}$ and $k_{kin,2}$, which may control the devolatilization over different temperature ranges, and yields an expression for the devolatilization as:

$$\frac{m_{\rm p}(t)}{(1-f_{\rm w,0})m_{\rm p,0}-m_{\rm ash}} = \int_0^t (\alpha_1 k_{kin,1} + \alpha_2 k_{kin,2}) \\ \times \exp\left(-\int_0^t (k_{kin,1} + k_{kin,2}) dt\right) dt.$$
(14)

For the gas phase reactions including soot formation (R1–R9 shown in Table 1), the smaller of the two reaction rates (finite rate and eddy dissipation) is used as the overall reaction rate ($\hat{R}_{i,k}$). The finite rate and the eddy dissipation models consider the reaction rate as follows:

Finite rate:
$$\hat{R}_{i,k}^{(A)} = (v_{i,k}'' - v_{i,k}')$$

 $\times \left(k_{kin,f,k} \prod_{i=1}^{I} [X_i]^{\eta_{i,k}'} - k_{kin,b,k} \prod_{i=1}^{I} [X_i]^{\eta_{i,k}'} \right)$
(15)

Eddy dissipation:
$$\hat{R}_{i,k}^{(R)} = v'_{i,k} M_i A_R \rho\left(\frac{\varepsilon}{k}\right) \min_{R} \left(\frac{Y_R}{v'_{R,k} M_R}\right)$$
(16)

Eddy dissipation:
$$\hat{R}_{i,k}^{(P)} = v'_{i,k} M_i A_{\rm R} B_{\rm P} \rho\left(\frac{\varepsilon}{k}\right) \left(\frac{\sum_{\rm P} Y_{\rm P}}{\sum_{j}^{N} v''_{j,k} M_j}\right).$$
(17)

The burning rate of the carbon in coal particle is calculated using the finite rate model proposed by Smith [6, 23, 24]. The rates of depletion of carbon due to surface reactions (R10–R12 shown in Table 1) are given as:

$$\bar{R}_k = A_p \eta_k Y_{\text{carbon}} \tilde{R}_k, \tag{18}$$

$$\tilde{R}_k = k_{kin,k} \left(p_{i,k} - \frac{\tilde{R}_k}{D_k} \right)^{N_k} .$$
(19)

The kinetic reaction rate k_{kin} follows an Arrhenius expression as:

$$k_{kin} = A_{\rm f} \exp^{-(E_{\rm ac}/RT_{\rm p})},\tag{20}$$

and the values of the kinetic parameters that are used to determine k_{kin} for all reactions are also shown in Table 1.

Boundary conditions

Uniform distributions of inlet mass flow rate and temperature are given for all inlet boundary surfaces. The walls are assumed as stationary and smooth with no slip condition. A constant wall heat flux is assigned for wall boundary surfaces. The boundary condition of the discrete phase at walls is assigned as "reflect", which means the discrete phase elastically rebound off once reaching the wall. At the outlet, the discrete phase exits the computational domain.

Numerical solutions procedure

Numerical methods

ANSYS FLUENT 12.0 is used to solve the set of equations discussed earlier. **FLUENT** uses a control volume-based technique to convert a general scalar transport equation to an algebraic equation that can be solved numerically. This control volume technique consists of integrating the transport equation about each control volume, yielding a discrete equation that expresses the conservation law on a control volume basis. General form of the discretized equation for an arbitrary control volume is as follows [16]:

$$\int_{V} \frac{\partial}{\partial t} (\rho \phi) \mathrm{d}V + \oint \rho \phi \vec{v} \cdot \mathrm{d}\vec{A} = \oint \Gamma_{\phi} \nabla \phi \cdot \mathrm{d}\vec{A} + \int_{V} S_{\phi} \mathrm{d}V,$$
(21)



 Table 1
 Kinetic parameters for gas and surface phase reactions

	A_f (consistent unit)	E_{ac} (J/kmol)	Reference(s)	
Devolatilization Ste	p 1: Coal \rightarrow Volatile ₁ + Char ₁			
	2.00×10^5	1.05×10^{8}	[19]	
Devolatilization Step	p 2: Coal \rightarrow Volatile ₂ + Char ₂			
	1.30×10^{7}	1.67×10^{8}	[19]	
(R1) $C_{\alpha 1}H_{\alpha 2}O_{\alpha 3}N_{\alpha 4}$	$\rightarrow \beta_1 CO + \beta_2 CO_2 + \beta_3 H_2 + \beta_4 CO_2$	$CH_4 + \beta_5 H_2 O + \beta_6 C_6 H_6 +$	$\beta_7 N_2$	
K _{kin, I}	3.09×10^{8}	1.67×10^{8}	[20]	
(R2) CO + $\frac{1}{2}O_2 \rightarrow$	CO ₂			
$K_{kin,2}$	2.20×10^{12}	1.67×10^{8}	[6, 21]	
$(\text{R3}) \text{ CO} + \text{H}_2\text{O} \leftrightarrow$	$\mathrm{CO}_2 + \mathrm{H}_2$			
$K_{kin,3f}$	2.75×10^2	8.38×10^{7}	[6, 21]	
$K_{kin,3b}$	2.65×10^{-2}	3.96×10^{3}	[6, 21]	
$(R4) CH_4 + H_2O \leftarrow$	\rightarrow CO + 3H ₂			
$K_{kin,4f}$	4.40×10^{11}	1.68×10^{8}	[6, 21]	
$K_{kin,4b}$	5.12×10^{-14}	2.73×10^4	[6, 21]	
(R5) CH ₄ + $\frac{1}{2}O_2$ –	$\rightarrow CO + 2H_2$			
$K_{kin,5}$	3.00×10^{8}	1.26×10^{8}	[6, 21]	
(R6) H ₂ + $\frac{1}{2}O_2 \rightarrow$	H ₂ O			
$K_{kin,6}$	6.80×10^{15}	1.68×10^{8}	[6, 21]	
$(R7) \ 4C_6H_6 \rightarrow C_{24}H_6$	$H_{12} + 6H_2$	· ·		
$K_{kin,7}$	1.50×10^{10}	4.70×10^{5}	[13, 22]	
(R8) $C_6H_6 + 4.5O_2 \rightarrow 6CO + 3H_2O$				
$K_{kin,8}$	2.00×10^9	3.10×10^{7}	[23]	
(R9) $C_{24}H_{12} + 15O_2 \rightarrow 24CO + 6H_2O$				
$K_{kin,9}$	2.00×10^9	3.10×10^{7}	[23]	
(R10) C + $\frac{1}{2}O_2 \rightarrow$	СО			
$K_{kin, 10}$	0.0520	1.30×10^{8}	[4, 6, 21]	
(R11) C + CO ₂ \rightarrow 2CO				
<i>K</i> _{kin,11}	0.0732	1.62×10^{8}	[4, 6, 21]	
$(R12) C + H_2O \rightarrow CO + H_2$				
<i>K</i> _{kin,12}	0.0782	1.47×10^{8}	[4, 6, 21]	

where ϕ is scalar variable, Γ is diffusion co-efficient, and \vec{A} is surface area vector.

Spatial discretization

Solution of Eq. (21) results in values of scalar at each computational node. To calculate convection terms in Eq. (21), scalar values are required at cell surfaces which must be interpolated from cell-centroid values (nodes). First-order upwind scheme is used for spatial discretization of the convective terms. First-order upwind assumes the value of the variable throughout the cell and at the face to be the same as the centroid value.

Pressure-velocity coupling

The discretization of the equations governing the gas phase is solved by the SIMPLE algorithm for pressure-velocity coupling. The algorithm starts with an initial guess for variables in the system. Momentum equations are solved and pressure is corrected using a pressure correction equation. In the next step, all the other transport equations are solved and residuals are checked. If the solution is not converged, the current results would be used as an initial guess for the next iteration. This loop will continue until a converged solution is obtained.

Under-relaxation factor

The following equation is used during iteration to calculate new value of the variable in each cell based on its old value.

$$\phi = \phi_{\text{old}} + \alpha \Delta \phi. \tag{22}$$

 α is the under-relaxation factor and its value controls the change of variables in each iteration. Due to the



nonlinearity of the equations, it is essential to reduce the change of variables in each time step; otherwise the solution becomes unstable and diverges.

Convergence criteria

For any transport equation, the discretized from of the equation has the following form:

$$Z_{\rm p}\phi_{\rm p} = \sum Z_{\rm nb}\phi_{\rm nb} + b_{\rm p} \tag{23}$$

where Z_p and Z_{nb} are central and neighboring co-efficients, respectively. Imbalance of this equation is called residual and can be expressed as:

$$R_{\rm p} = \sum Z_{\rm nb}\phi_{\rm nb} + b_{\rm p} - Z_{\rm p}\phi_{\rm p} \tag{24}$$

This equation will be scaled based on summation of residual in all computational cells. Usually, when the scaled residuals drop by three orders of magnitude, a qualitative convergence has been obtained. In this study, tolerances of pressure and velocity components are set to 1E-3, while tolerance of gas and solid components are set to 1E-5, and energy equation to 1E-6.

Calculation conditions

A bituminous-type CV coal (Coal Valley, Canada) is used to conduct the simulation of coal gasification. The proximate and ultimate analyses of coal are given in Table 2. The initial particle size distributions with a mean diameter of 60 μ m are used in the calculation. The total mass inlet for experiment and calculation are kept same. The coal flow rates for combustor and reductor are set to 40 and 60 kg/h, respectively. The gas flow rates are adjusted in such a way that the inlet O₂ ratio and O₂ concentration

 Table 2 Analyses of coal [25]

Parameters	CV coal (Canada)
Proximate analyses (air dried)	
Moisture (wt%)	6.22
Fixed carbon (wt%)	49.00
Volatile matter (wt%)	34.50
Ash (wt%)	10.28
Ultimate analyses (dry base)	
C (wt%)	69.90
H (wt%)	4.30
O (wt%)	13.70
N (wt%)	1.07
High heating value (MJ/kg)	26.40
Low heating value (MJ/kg)	26.02

become 0.528 and 23 wt%, respectively. The O₂ ratio is defined here as the ratio of the amount of O₂ fed into the gasifier to the amount of O₂ required for complete combustion of carbon present in coal. During devolatilization, all hydrogen, oxygen and nitrogen are assumed to be released as volatiles. Volatiles are considered as a single hypothetical component, $C_{\alpha 1}H_{\alpha 2}O_{\alpha 3}N_{\alpha 4}$. The values of $\alpha 1$, $\alpha 2$, $\alpha 3$ and $\alpha 4$ are calculated from the coal's ultimate and proximate analyses. Once the volatile component is released, it is converted into CO, CO₂, H₂O, H₂, CH₄, C₆H₆ and N₂ according to reaction R1. The pyrolysis data obtained from previous experimental work [26] are used to calculate the β values. In the calculation, all aliphatic and aromatic compounds are lumped into CH₄ and C₆H₆, respectively (Table 3). Chen et al. [27] explained the gas evolution from rapid pyrolysis of a bituminous coal at various pyrolysis temperatures (500-900 °C). It was found that the ratio of CO to CO₂ yield does not change with increasing the pyrolysis temperature. They also showed that the yield of higher hydrocarbon is approximately three times higher than that of CH₄, which is very near to our previous works. Therefore, these two ratios $(Y_{d,CO}/$ $Y_{d,CO2} = 1.27$ and $Y_{d,C6H6}/Y_{d,CH4} = 3.10$) together with three elemental (C, H, O) mass balance equations are used to calculate the stoichiometric co-efficient (β) of product species for reaction R1 (see Table 3). Here Y_d represents the mass yield for the corresponding species.

Results and discussion

Validation of one-step soot model

To validate the one-step soot model, a tubular-type reactor of 0.28 m diameter and 48 m length with the inlet gas velocity of 26.4 m/s is used to conduct the simulation. Eight overall gas phase reactions (R2-R9 shown in Table 1) are considered in the calculation. Benzene (C_6H_6) , naphthalene ($C_{10}H_8$), phenanthrene ($C_{14}H_{10}$) and pyrene (C₁₆H₁₀) are independently considered as a precursor of soot formation. The calculated outlet species concentration of soot and syngas using the proposed soot model are compared with those reported in our previous paper [13] under various gasification conditions. The comparisons are shown in Fig. 3. The trends in outlet species concentration with increasing temperature are found to be similar, in both: detailed mechanism and overall gas phase reactions with proposed one-step soot mechanism. Soot concentration decreases with increasing the temperature while syngas concentration gradually increases. Soot formation as well as soot oxidation tends to increase at higher temperatures, resulting in an increase in CO and H₂ concentration. A detailed explanation of the effect of temperature



Table 3 Volatiles species concentration produced from coal pyrolysis and β values for R1

Species	wt% [13, 26]	β Value (present study)
CO	0.2849	0.110
CO ₂	0.2239	0.057
H ₂	0.0502	0.342
H ₂ O	0.9643	0.470
(CH ₄) aliphatic compounds	0.4621	0.251
(C ₆ H ₆) aromatic compounds	1.4329	0.159
N ₂	96.5817	Balanced with total N_2 present in coal



Fig. 3 Comparison of calculated outlet soot and syngas concentration between detailed reaction mechanism [13] and overall gas phase reactions with one-step soot mechanism calculated at 2.0 MPa [*open diamond* is for benzene (C_6H_6), *open circle* is for naphthalene ($C_{10}H_8$), *open triangle* is for phenanthrene ($C_{14}H_{10}$) and *multiply symbol* is for pyrene ($C_{16}H_{10}$)]

on soot and syngas concentration can also be found in Wijayanta et al. [13]. In addition, very similar results are obtained for various soot precursors (benzene, naphthalene, phenanthrene and pyrene) considered in the calculation. As increasing the molecular weight of species significantly increases the computational time, benzene is chosen as a soot precursor in the simulation of coal gasification in the two-stage entrained-flow gasifier shown in Fig. 1.

Comparisons of species concentration and temperature profile

The comparison for two conditions of without soot and with soot shown in Fig. 4a indicates that there is a small change in outlet species concentration. An increase in H_2 concentration under soot formation condition suggests that the concentration of H_2 will be increased if the soot formation advances in the gasifier. This means that the formation of soot can increase the syngas heating value in this regard, despite having diverse effect of soot. Figure 4a also shows a comparison of outlet species concentration between experiment and calculation. Details of the experimental procedure and condition are described by Kidoguchi et al. [25]. A quite good agreement is obtained for main species CO, CO₂ and H₂. The agreement for the species H₂O is not good due to the lack of information of



Fig. 4 Comparison between experiment [from CRIEPI, 25] and calculation: ${\bf a}$ outlet species concentrations and ${\bf b}$ gas temperature profiles at centerline

experiment. A large deviation for H_2O concentration between experiment and calculation is obtained due to the presence of some moisture in air during experiment which is ignored in the calculation.

The gas temperature profiles at centerline for experiment and calculations are shown in Fig. 4b. In both, trends of gas temperature are found to be similar for experiment and calculations. However, calculation without soot formation overestimates the experimental gas temperature. In contrast, calculation with soot formation provides better agreement with the experiment. In case of soot formation, the reaction R1 includes aromatic species C_6H_6 which is considered as a soot precursor. C_6H_6 is then accumulated to produce a larger species, Coronene ($C_{24}H_{12}$), which is referred here as soot. The gas temperature for calculation with soot decreases significantly because of reducing the heat of reaction (R1). The gas temperature also decreases due to the large heat capacity of aromatic species considered in the soot formation reaction mechanism.

Effect of O₂ ratio

The effect of O_2 ratio on soot concentration, gas temperature, carbon conversion, etc., is numerically investigated under CO_2 -rich gasification condition (CO_2 concentration = 14 wt%). The contours of soot concentration and gas temperature under conditions of two different O_2 ratios at 0.528 and 0.7 are shown in Fig. 5a. A slight decrease in soot concentration at outlet from 1.79 to 1.73 wt% is observed if the O_2 ratio increases from 0.528 to 0.7 in the gasifier. On the other hand, the gas temperature at outlet significantly increases from 1352 to 1588 K with increasing the O_2 ratio. This is because under



Fig. 5 Effect of O_2 ratio on **a** contours of soot concentration and gas temperature and **b** heating value, carbon conversions and soot concentration at outlet (calculated under conditions at constant CO_2 concentration of 14 wt%)

higher O₂ ratio, exothermic char-O₂ reaction tends to increase. The high gas temperature then advances the endothermic char-CO₂ and char-H₂O gasification reactions. This means increased O₂ ratio significantly enhances char-O₂ reaction as well char-CO₂ and char-H₂O reaction. Although, char-CO2 and char-H2O reactions are endothermic, the gas temperature increases due to significant rise in char-O₂ oxidation reaction under higher O₂ ratios. An increase in char-O₂, char-CO₂ and char-H₂O reactions under a high O₂ ratio at 0.7 results in an increase in carbon conversion and syngas concentration. Figure 5b shows that the carbon conversion gradually increases with increasing the O₂ ratio and reaches a complete (100 wt%) conversion at a O_2 ratio of 0.8. The soot concentration is also found to decrease at higher O2 ratios. In contrast, syngas heating value initially increases with increasing the O₂ ratio, and reaches a maximum value of 3800 kJ/kg with 94 wt% carbon conversion at a O₂ ratio of 0.7 (Fig. 5b). Beyond this value (O_2 ratio = 0.7) heating value decreases with increasing O2 ratio because of shifting the environment from gasification towards combustion. Therefore, if the target is to get a complete conversion of carbon, a lower heating value gas will be produced from the coal gasification. Considering the carbon conversion in real gasification process where unconverted carbon is recycled as char and the use of more O₂ where low heating value gas is produced, the O₂ ratio exceeding 0.7 is not recommended for getting efficient coal gasification. Therefore, to improve the gasification efficiency, the concentrations of other gasification agents (CO2 and/or H2O) need to be increased in coal gasification process keeping O₂ ratio under 0.7. With a target of reducing CO₂ release into the atmosphere from coal gasification, the effect of CO₂ concentration on soot concentration, syngas heating value and carbon conversion is numerically investigated in this study.

Effect of CO₂ concentration

The effect CO_2 concentration on soot concentration, gas temperature, carbon conversion, etc., is numerically investigated by changing the inlet concentration of CO_2 at a constant O_2 ratio (=0.528). The contours of soot concentration and gas temperature under conditions of two different CO_2 concentrations at 14 and 50 wt% are shown in Fig. 6a. No significant difference in soot concentration is found for the two cases, although the gas temperature decreases with increasing the overall concentration of CO_2 . The gas temperature decreases with increasing the CO_2 concentration due to increased char– CO_2 reaction rate. Under CO_2 -rich concentration, endothermic reaction (backward reaction of R3) also increases, resulting in a decrease in gas temperature. The backward tendency of R3 on the other hand increases CO concentration and





Fig. 6 Effect of CO_2 concentration on **a** contours of soot concentration and gas temperature and **b** heating value, carbon conversions and soot concentration at outlet (calculated under conditions at constant O_2 ratio of 0.528)

decreases H₂ concentration. However, the syngas heating value gradually increases with increase of the CO₂ concentration (shown in Fig. 6b) although H₂ concentration decreases at higher CO₂ concentrations. Soot concentration and carbon conversions for various cases are also shown in Fig. 6b. It is found that the soot concentration is nearly independent to CO₂ concentration. On the other hand, the reductor carbon conversion gradually increases with increasing the CO₂ concentration. A 15 % increase in carbon conversion of reductor coal is obtained if the inlet concentration of CO₂ is increased from 14 to 50 wt%. However, the same change in CO₂ concentration gives only a 1 % increase in overall carbon conversion. This is due to low carbon conversion of combustor coal at a high CO₂ concentration (50 wt%). Interestingly, the syngas heating value increases from 2717 to 3501 kJ/kg which corresponds to a 28 % increase in syngas heating value. This indicates that the carbon conversion is not directly related to the syngas heating value. Under higher CO₂ concentrations, char-CO₂ (C + CO₂ \rightarrow 2CO) reaction dominates over char– O_2 (C + 1/2 $O_2 \rightarrow$ CO) and char-H₂O



 $(C + H_2O \rightarrow CO + H_2)$ reactions. This results in an increase in CO concentration with a small increase in carbon conversion. Therefore, it can be concluded that the production of syngas heating value per unit weight of carbon conversion will be higher if CO₂ concentration increases in the gasifier.

Conclusions

A one-step soot formation mechanism is proposed and numerically validated with the detailed reaction mechanism. The proposed mechanism is used to conduct a series of 3D numerical simulation with the aim of describing the gasification process in two-stage entrained-flow gasifier. The calculated results with one-step soot formation reaction mechanism show a good agreement with the experimental results. It is found that formation of soot enhances the H₂ production while predicting a low gas temperature in the gasifier. As the O₂ ratio increases, soot concentration decreases while the gasifier gas temperature and carbon conversion increase. Beyond a certain limit of O_2 ratio at 0.7, soot concentration and syngas heating value sharply decrease. In contrast, syngas heating value gradually increases as the CO₂ concentration increases without affecting the soot concentration and with a small increase in overall carbon conversion. This means that the syngas heating value per unit weight of carbon conversion produced from CO₂-rich gasification condition will be higher than that from the condition with lower CO₂ concentrations and, therefore, coal gasification under CO2-rich condition can be efficiently implemented in IGCC system.

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