Thermochemical Recycling of Municipal Solid Waste

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Cyprus 2016

Institute of Chemical and Environmental Engineering

Reactor Engineering Reaserch Group

- Experimental study and mathematical modeling of fuel thermal processes
- Pyrolysis, gasification and combustion of solid fuels
- Biomass, polymer waste, MSW, and coal thermal and catalytic processing for production energy and materials

National center for research and application of renewable energy sources



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Refuse-Derived Fuel (RDF)



RDF composition

Component	Material	w _i [kg/kg]
Paper	White paper, recycled paper	0,6317
Foil	LDPE, HDPE	0,1578
Plastics	Rigid plastics, polystyrene, polyurethane	0,1910
Textile	Polyamide, polyester, cotton , wool	0,0194

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Proximate and Elemental Composition of RDF

Com.	Mois.	VM*	FC*	ASH*	С	н	N	S	O**
Wt. %	10	75.5	8.9	15.6	51.7	5.9	0.9	0.4	25.5

*moisture free basis
**calculated to 100%



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Behaviour of Thermal decomposition



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Behaviour of Thermal decomposition



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Heating value of RDF

Component	Heating value [kJ/kg]
Paper	13410
Foil	43860
Plastics	33570
Textile	19770
Mixed RDF	20810

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Tar content measurement



Gasification Model

Assumptions:

- Steady state flow is considered inside the gasifier
- No temperature and concentration gradient exist inside the reactor
- The residence time is enough long to reach complete decomposition of RDF and unreacted part of RDF is only carbon.
- Only the major species are considered in the product gases, i.e CO, CO₂, H₂, CH₄ , H₂O, NH₃, H₂S, N₂ and Tar

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Global material balance of RDF gasification

 $CH_{b}O_{c}N_{d}S_{e} + x_{1}O_{2} + x_{2}H_{2}O \rightarrow x_{3}CO + x_{4}CO_{2} + x_{5}H_{2} + x_{6}CH_{4} + x_{7}H_{2}O$ RDF $+x_{8}NH_{3} + x_{9}H_{2}S + x_{10}CH_{b1}O_{c1}N_{d1}S_{e1}$ TAR

Reactions:

$C+0, 5O_2 \longrightarrow CO$	$C + H_2 O \longleftrightarrow H_2 + CO$
$CO+0, 5O_2 \longrightarrow CO_2$	$C + CO_2 \longleftrightarrow 2CO$
$H_2 + 0,5O_2 \longrightarrow H_2O$	$CH_4 + H_2O \longleftrightarrow 3H_2 + CO$
$CH_4 + 2O_2 \longrightarrow CO_2 + 2H_2O$	$C+2H_2 \longrightarrow CH_4$
	$CO + H_2O \longleftrightarrow CO_2 + H_2$

Equilibrium constant:

$$K_{a} = \left(\frac{P}{P^{0}}\right)^{\sum v_{i}} \prod \phi_{i}^{v_{i}} \prod x_{i}^{v_{i}}$$

$$K_{a}^{298} = e \frac{-\Delta_{r} G^{298}}{RT} , \quad \Delta_{r} G^{298} = \Delta_{r} H^{298} - T\Delta_{r} S^{298}$$

$$\Delta_{r} H^{298} = \sum v_{i} \Delta_{f} H_{i}^{298} \qquad \Delta_{r} S^{298} = \sum v_{i} \Delta_{f} S_{i}^{298}$$

$$\Delta_{r} H = \Delta_{r} H^{298} + \sum v_{i} c_{pi} \cdot (T - 298) \qquad \Delta_{r} S = \Delta_{r} S^{298} + \sum v_{i} c_{pi} \cdot \ln \frac{T}{298}$$

Enthalpy balance:

$$H_{RDF} + H_{O2(air)} + H_{steam} + Q_R = H_{gas} + H_{ash} + H_C + Q_{loss}$$

 $Q_{R} = m_{RDF} \sum w_{i} Q_{i} - \sum (-\Delta_{c} H_{i}) n_{i} \qquad \text{IF, } T_{air} = T_{RDF} = T_{ref}, \text{ then } H_{RDF} = 0, H_{O2(air)} = 0$

$$T = T_{ref} + \frac{m_{RDF} \sum w_i Q_i - \left(\sum (-\Delta_c H_i) n_i\right) - Q_{loss}}{\left(\sum n_i c_{pi}\right) + m_C \overline{c}_{pC} + m_{ash} \overline{c}_{pash} - m_{steam} \overline{c}_{steam}}$$

 Q_{R} – heat of reaction [J],

 H_{RDF} – enthalpy of RDF feed [J],

$$H_{O2(air)}$$
 – enthalpy of oxygen and air respectively [J],

 H_{steam} – enthalpy of water steam [J],

 H_{gas} – enthalpy of gas [J],

$$H_{ash}$$
 – enthalpy of ash [J],

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 H_{C} – enthalpy of unreacted carbon [J],

 Q_{loss} – heat losses from the reactor [J]

SLOVENSKÁ TECHNICKÁ UNIVERZITA V BRATISLAV *m*_{RDF}– mass flow of RDF feed [kg]

 n_i – mole flow of component i in the products [kmol]

 w_i – maas fraction of component *i* in the feed (paper, foil, plastics, textil

 Q_i – lower heating value of component *i* in the feed (paper, foil, plastics, textile) [Jkg⁻¹],

 $\Delta_{i}H_{i}$ - heat of combustion of component *i* in the products [Jkmol⁻¹]

m_{ash}– mass flow of ash [kg]

*m*_{ash}– mass flow of remaining carbon [kg]

m_{steam}- mass flow of steam [kg]

 \overline{c}_{pash} – specific heat capacity of ash [Jkg⁻¹K⁻¹]

 \overline{c}_{pc} – specific heat capacity of remaining carbon [Jkg⁻¹K⁻¹]

 \overline{c}_{psteam} – specific heat capacity of steam [Jkg-1K⁻¹]

Results of modelling RDF gasification

Observed parameters:

Conversion of RDF Reactor Temperature Gas composition Content of pollutants (NH3, H2S, TAR)

Variables:

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Oxygen (air) to RDF mass ratio Steam to RDF mass ratio



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Gasification of RDF Using O₂



Effect of RDF composition

Com.	Wt. %
Mois	10
VM	75.5
FC	8.9
ASH	15.6
С	51.7
Н	5.9
Ν	9.9
S	0.4
0	25.5



Com.	Wt. %
Mois	1.2
VM	80.22
FC	5.23
ASH	13.34
С	51.66
Н	8.82
Ν	9.66
S	0.08
0	25.42
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Effect of Steam in RDF Gasification



Conclusion

- For RDF studied in this work,100% of RDF conversion in gasification by air was reached at m_{air}/m_{RDF}=2,2. However, the gas heating value was 4,4 MJ/Nm³
- Gasification of RDF using Oxygen enables production of a gas with heating value around 10 MJ/Nm³ at m_{O2}/m_{RDF}=0,45
- Elemental Composition of RDF has a crutial effect on riquired $m_{air}\!/m_{RDF}$
- Raw untreated gas tar content was 3.3 mass %; tar fraction content a solid phase insoluble in isopropanol
- By increasing the m_{steam}/m_{RDF} the content of H2 and CO₂ increased, However, the content of CO, reactor temperature and gas heating vale decreased



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Thank you for attention

