

Thermochemical conversion of Biomass

Gasification

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IPESE

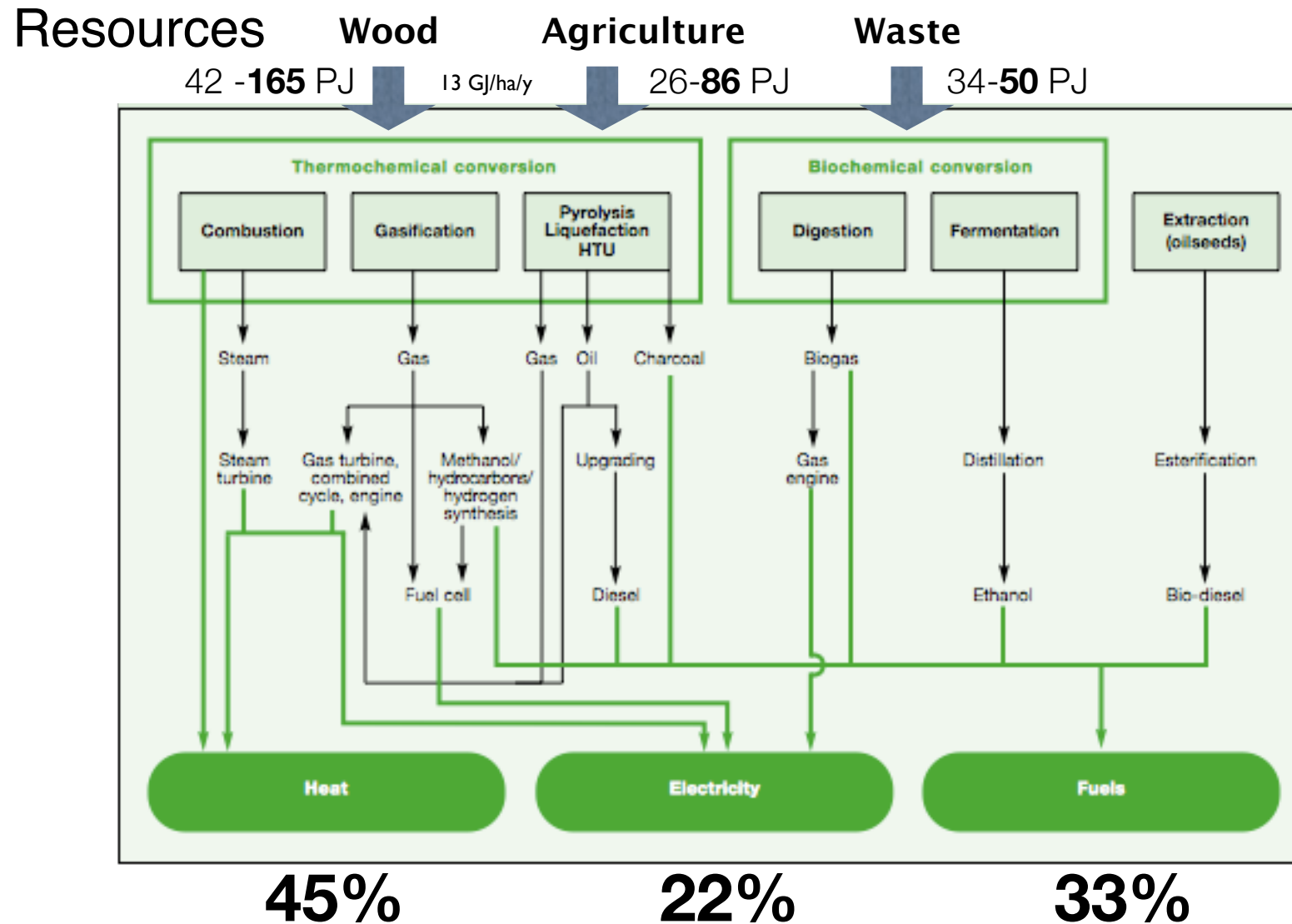
Industrial Process and Energy Systems Engineering

EPFL Valais-Wallis

CH - 1950 Sion

Biomass conversion routes and production potential in Switzerland

sustainable - **Technical** in PJ/year



Figures for Switzerland (sustainable potential : total = 82-301 PJ)

Source : world energy assessment : UNDP 2000

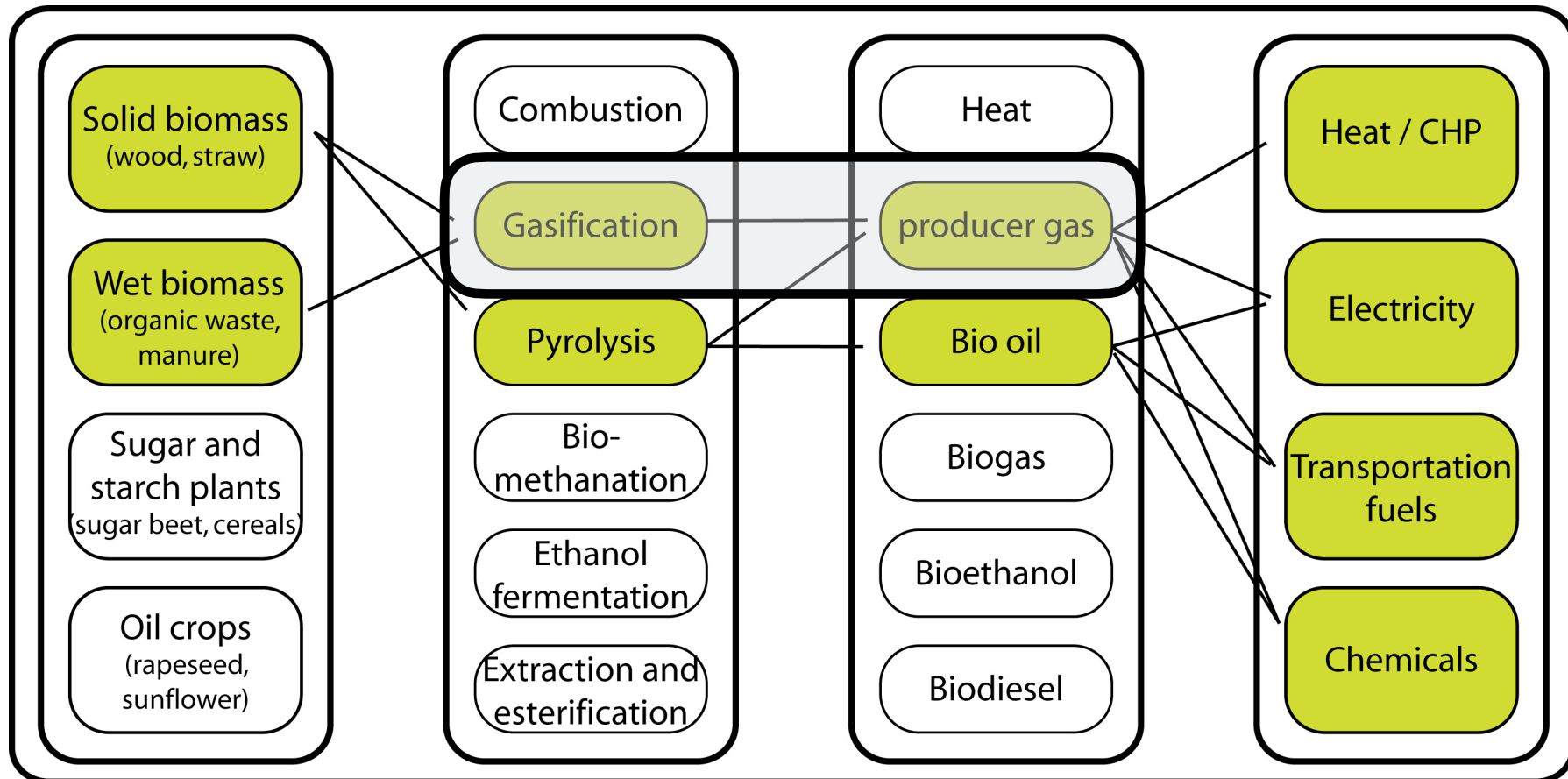
Bioenergy in Switzerland: [assessing the domestic sustainable biomass potential](#)

B Steubing, R Zah, P Waeger, C Ludwig

Renewable and Sustainable Energy Reviews 14 (8), 2256-2265

Biomass conversion

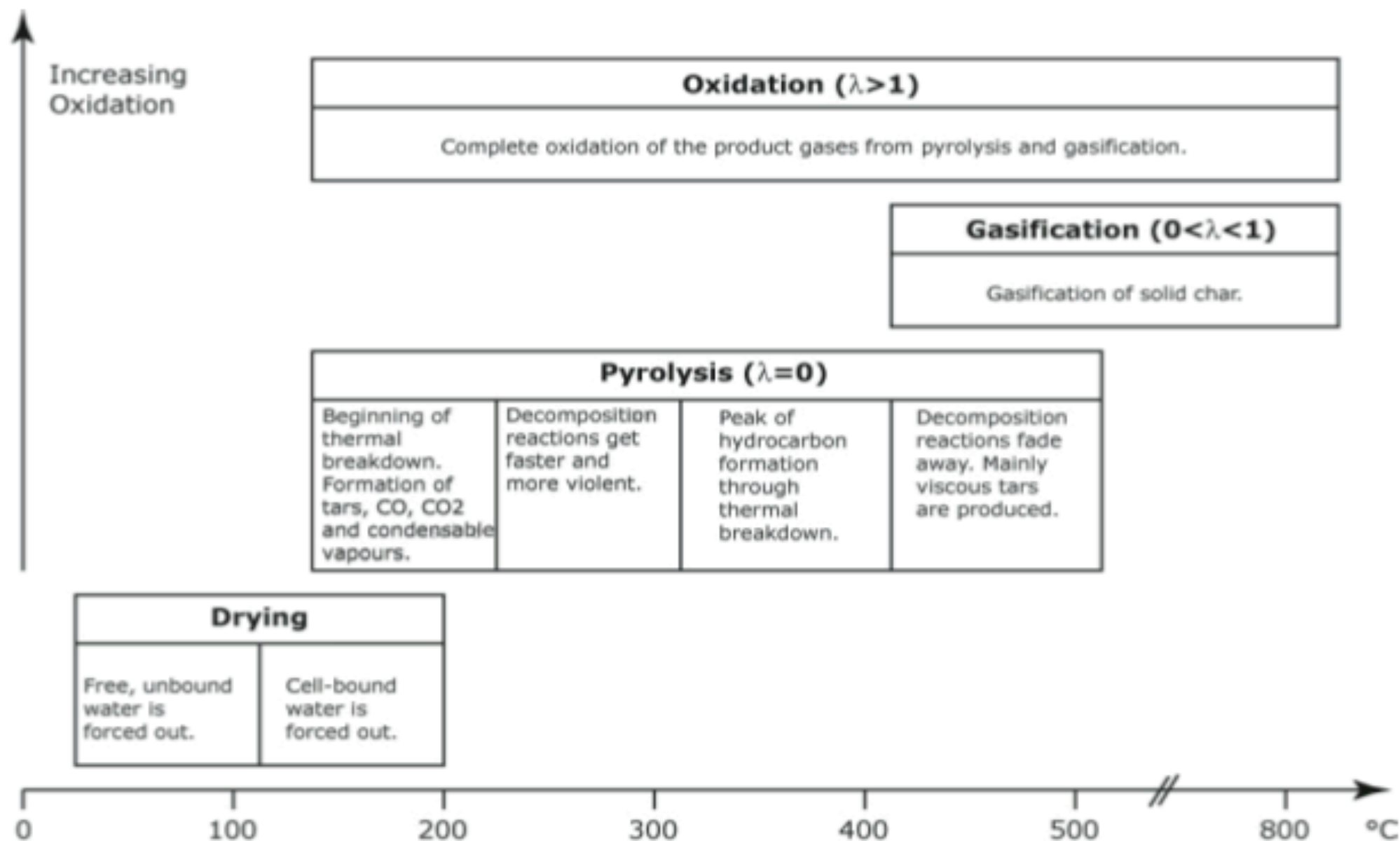
Thermochemical routes



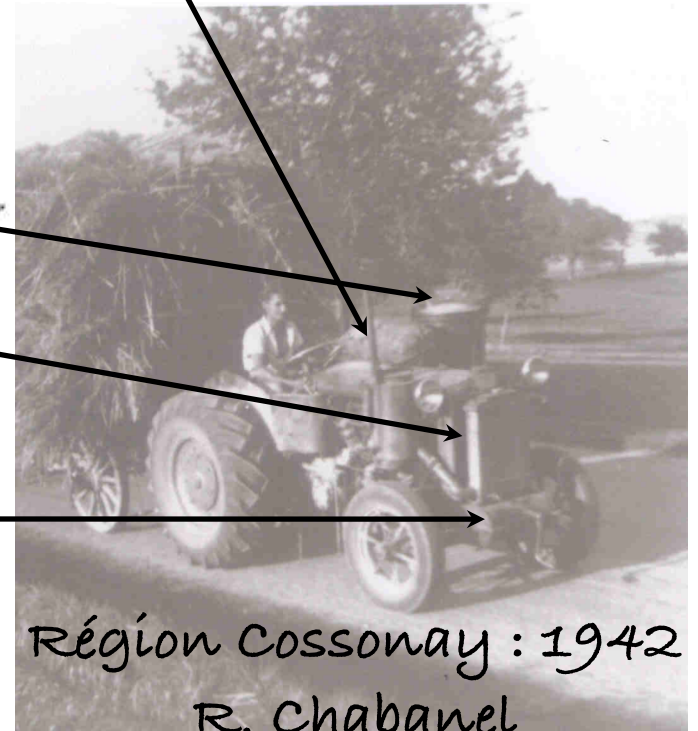
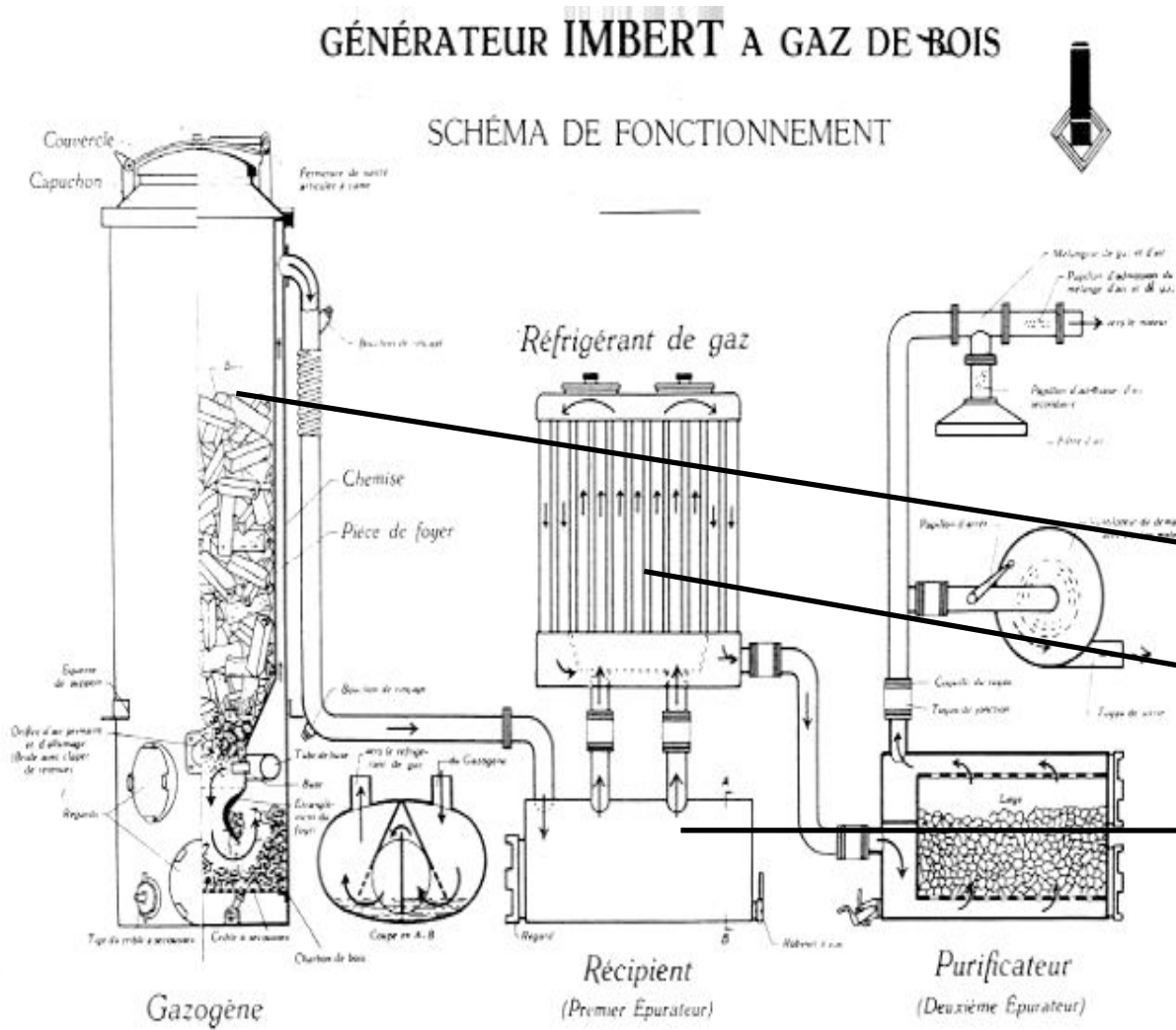
adapted from Chemical Engineering 10 (2006)

Thermochemical biomass conversion

Process principles: gasification vs. pyrolysis



Gasification & Fuel

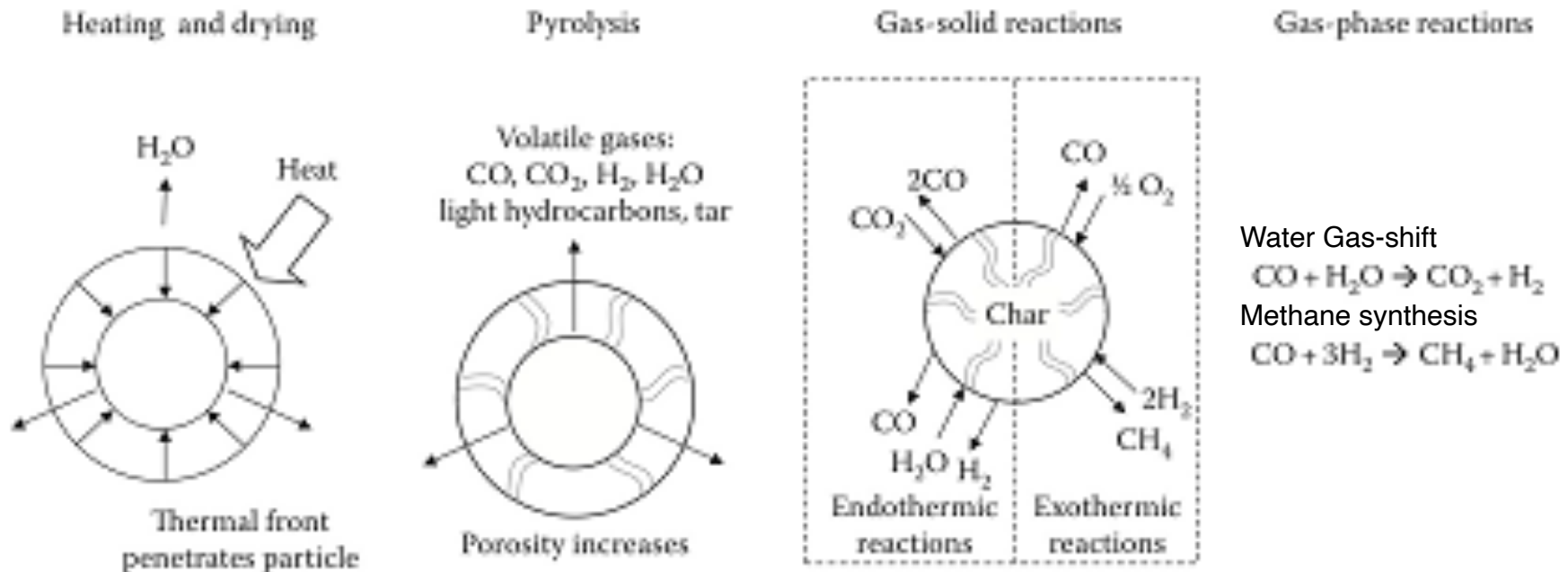


Région Cossonay : 1942
R. Chabanel

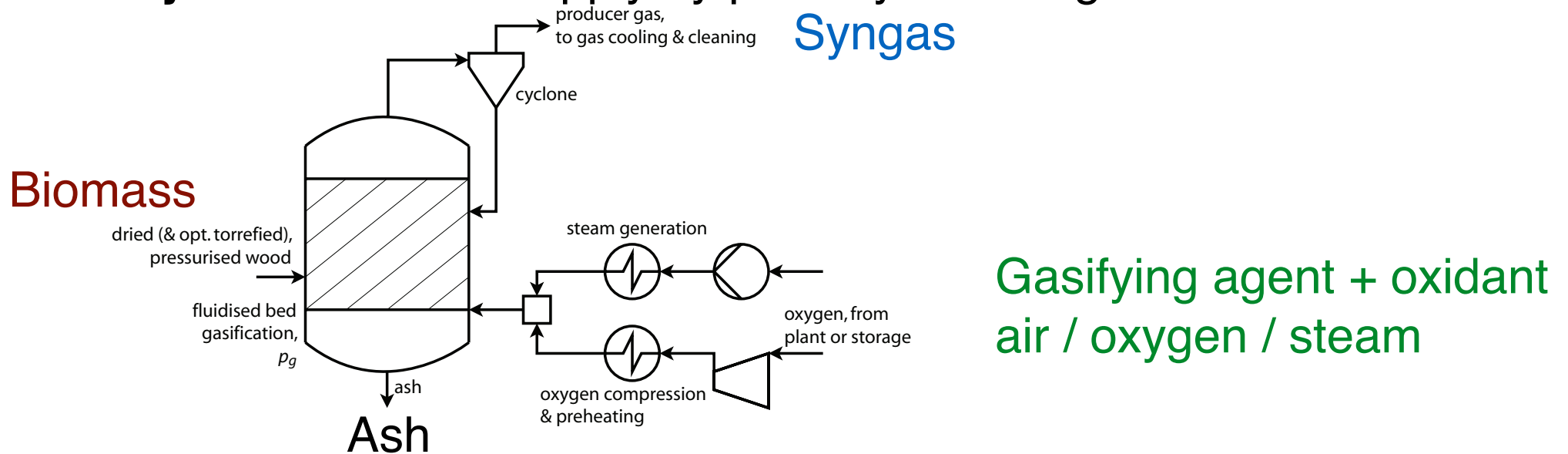
endothermic



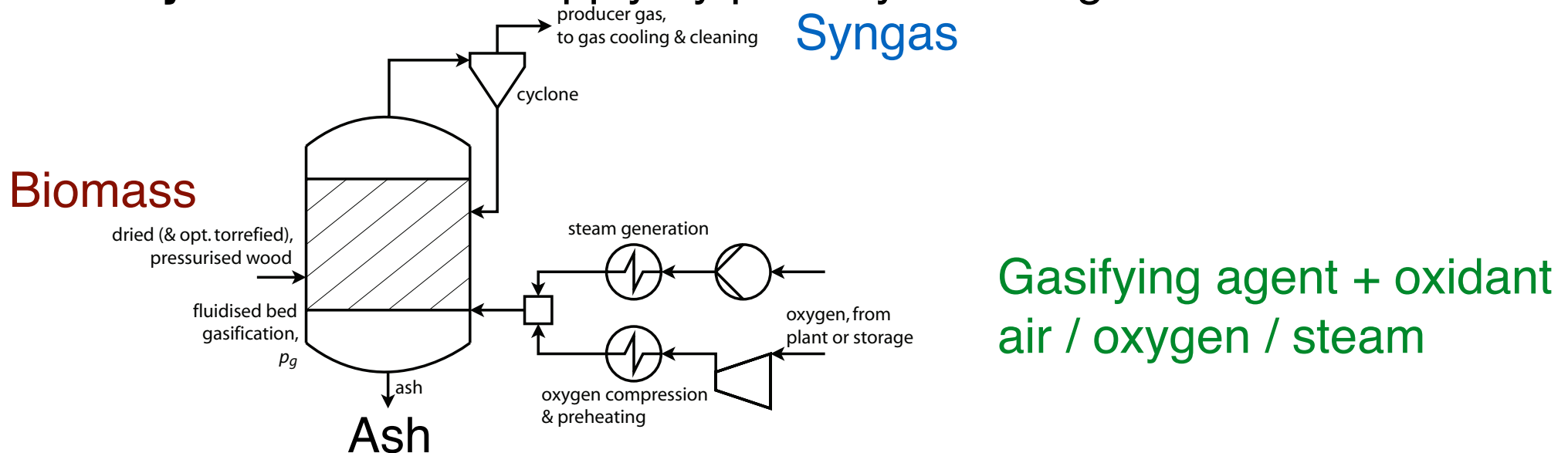
wood steam syngas
(gasifying agent)



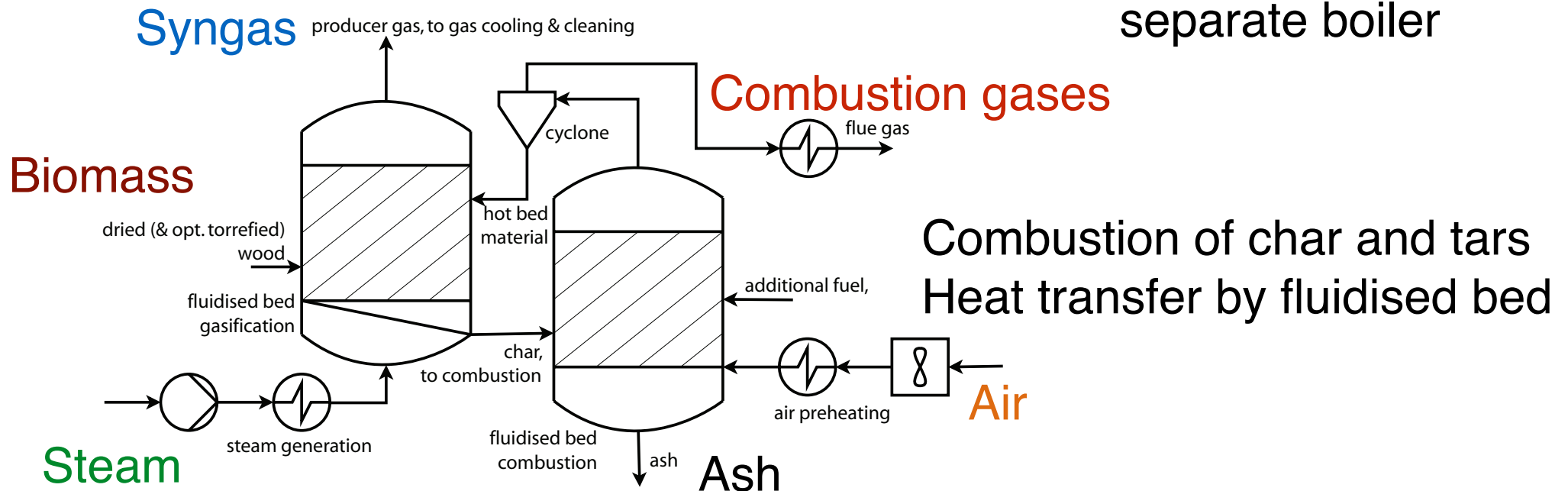
Directly heated : heat supply by partially oxidising the biomass



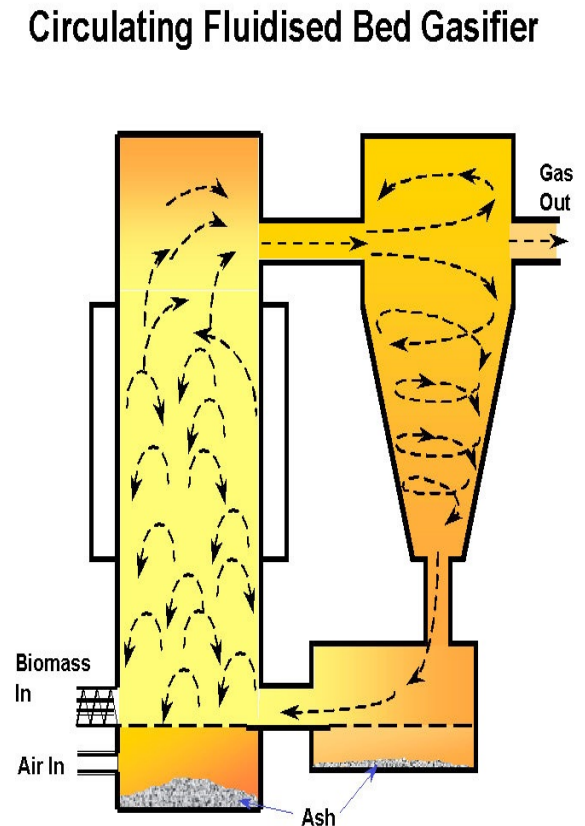
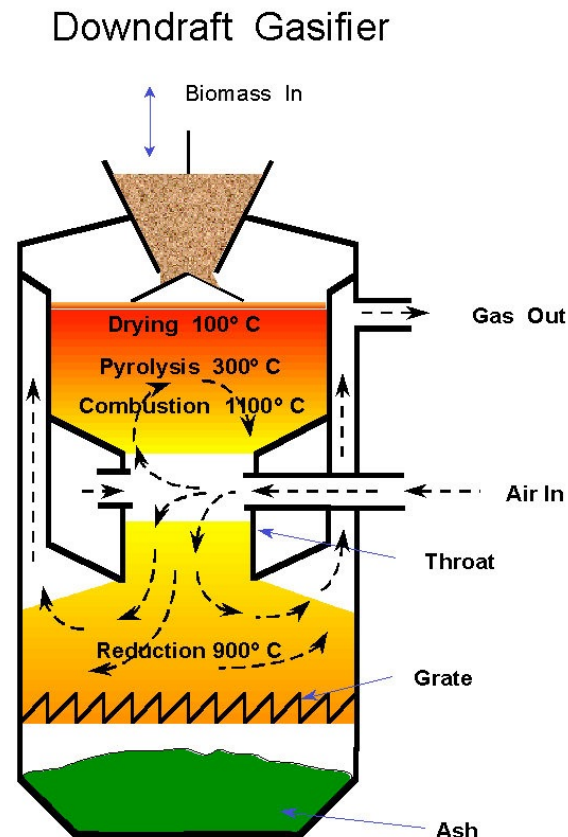
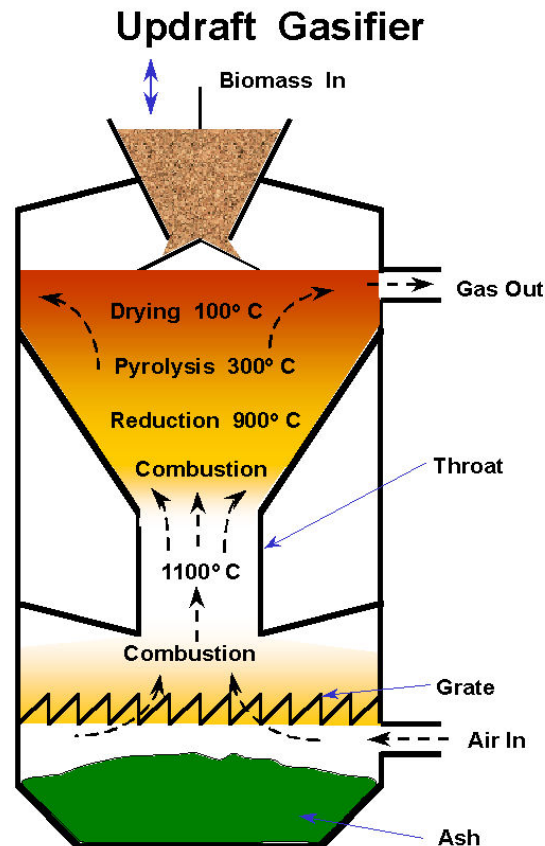
Directly heated : heat supply by partially oxidising the biomass



Indirectly heated : heat supply by burning part of the biomass in a
separate boiler



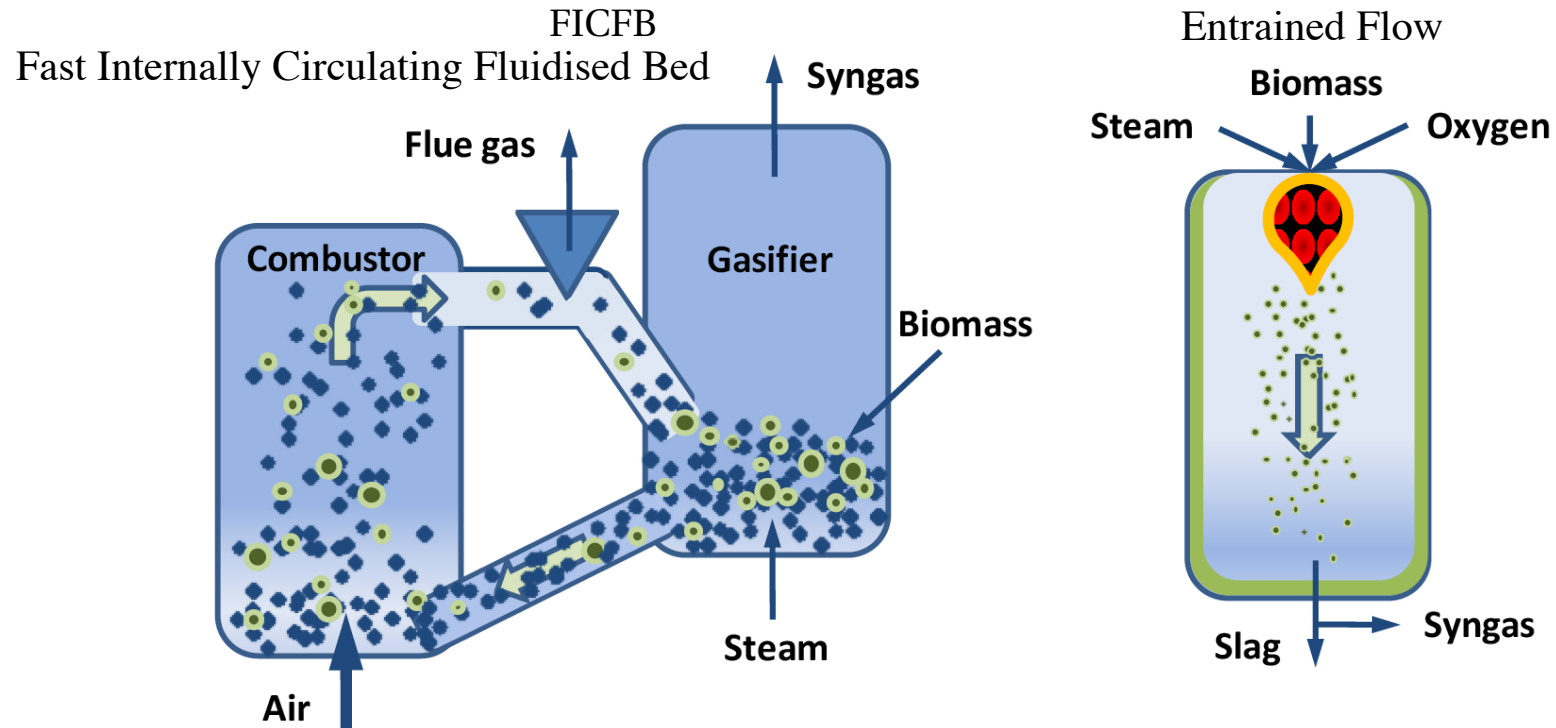
Solid - Fluid dynamics is important



Heat is supplied by partial oxidation of the biomass

Types of gasifiers

Comparing operating conditions



Temperature	800-1000 °C	1200-1500 °C
Pressure	1-4 bar	30-80 bar
Gas residence time	10 s	seconds
Solid residence time	several minutes	seconds
Particle size	50-150 mm ^a	< 0.1 mm
Gasification agent	steam	steam/O ₂

- Two stages gasification

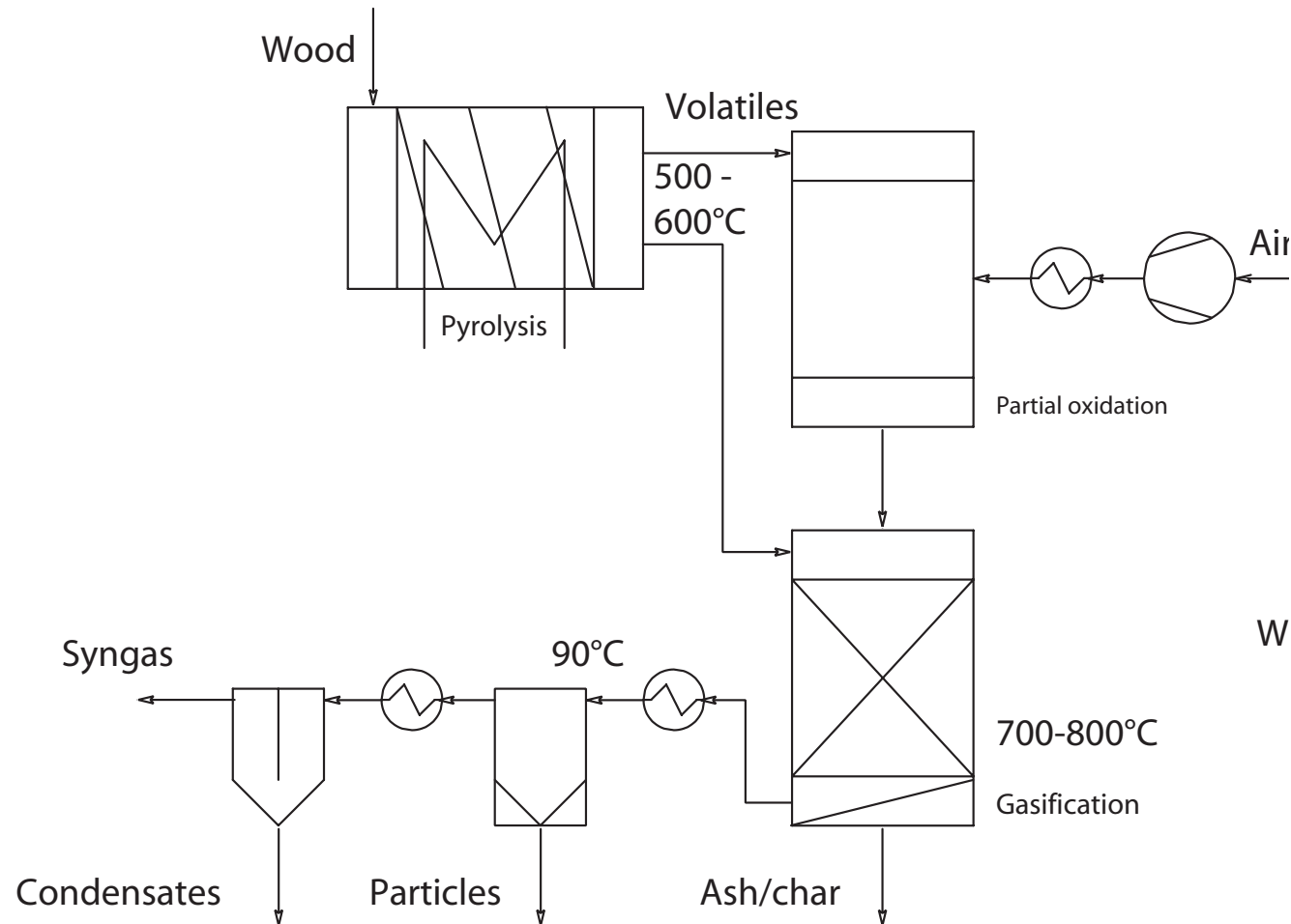


Table 2 – Gasification and methanation reactions.

Interaction	Name	Reaction	$\Delta \tilde{h}_r^0$
Solid–gas	Hydrogenating gasification	$\text{C(s)} + 2\text{H}_2 \rightleftharpoons \text{CH}_4$ (3)	–75 kJ/mol
	Boudouard equilibrium	$\text{C(s)} + \text{CO}_2 \rightleftharpoons 2\text{CO}$ (4)	173 kJ/mol
Gas–gas	Methane synthesis	$\text{CO} + 3\text{H}_2 \rightleftharpoons \text{CH}_4 + \text{H}_2\text{O}$ (5)	–206 kJ/mol
	Ethene reforming	$\text{C}_2\text{H}_4 + 2\text{H}_2\text{O} \rightleftharpoons 2\text{CO} + 4\text{H}_2$ (6)	210 kJ/mol
	Water–gas shift equilibrium	$\text{CO} + \text{H}_2\text{O} \rightleftharpoons \text{CO}_2 + \text{H}_2$ (7)	–41 kJ/mol

Equilibrium model

$$\hat{K}_p = K_p(T_g + \Delta T_{eq}) \quad (14)$$

$$K_p(T) = \exp \left(\frac{\sum_j n_j \tilde{g}_j^0(T) - \sum_i n_i \tilde{g}_i^0(T)}{RT} \right) \quad (15)$$

$$\hat{K}_p = \frac{\prod_j p_j^{n_j}}{\prod_i p_i^{n_i}} \quad (16)$$

Table 5 – Reconciled gasification model parameters.

Gasifier	Indirectly heated	Directly heated
$\Delta T_{eq,(3)}$	–280 °C	–280 °C
$\Delta T_{eq,(7)}$	–112 °C	–112 °C
$k_{\text{C}_2/\text{C}_1}$	0.205	0.476
ε_{cc}	90.3%	93.0%

Table 4 – Reconciliation of the producer gas compositions at nominal T_g and $p_g = 1$ bar (Data/Calculation [%vol]).

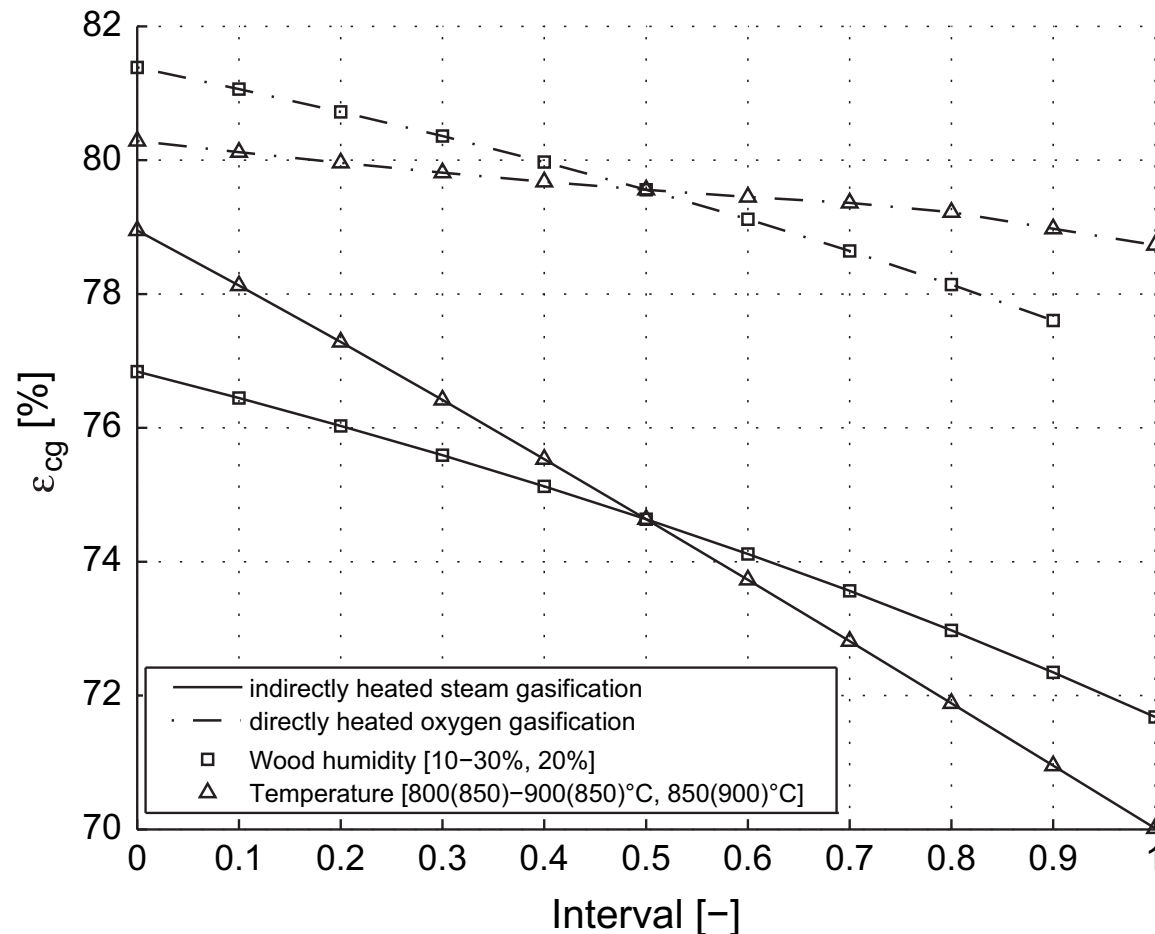
Gasifier	Indirectly heated [66]	Directly heated [29]	Directly heated (O ₂)
T_g	850 °C	800 °C	800 °C
C_2H_4	1.8/1.9	2.0/1.9	–/4.5
CH_4	8.8/9.6	4.2/4.0	–/9.5
H_2	37.3/38.5	14.8/14.7	–/25.8
CO	29.4/27.4	15.4/16.0	–/32.3
CO_2	16.2/15.8	15.0/14.7	–/24.0
N_2	^a –/2.9	39.6/40.3	–/0.1
H_2O	3.6/3.9	–/8.4	–/3.8
Δh^0	12.0/12.2	6.2/6.2	–/12.9
[MJ·Nm ^{–3}]			
SN [–]	0.24/0.26	0.13/0.13	–/0.13

a Although no nitrogen is introduced by the gasification agent, some N₂ is used for inertisation of the raw material, which prohibits to attain the criterion on the Wobbe Index at the process outlet. In the remainder of this work, a cut-down to 0.5%vol of the dry gas by inertisation with CO₂ is assumed feasible.

- Composition depends on gasifying agent

Reactor (<i>n</i> iterations)	Units	O ₂ gasification (15,000)		Air gasification (15,000)		H ₂ O gasification (15,000)	
Tar	mg/N m ³	5189	33,891	15	382	5354	86,034
N ₂	%	0.2	0.3	47.7	59.6	0.2	0.2
H ₂	%	11.5	13.3	10.7	7.5	20.1	14.2
CO ₂	%	49.3	59.5	18.9	21.2	31.2	38.2
CO	%	14.6	9.8	13.8	5.8	16.8	9.4
CH ₄	%	11.5	10.4	6.1	4.4	17.3	17.1
Other HC	%	12.8	5.6	2.7	1.0	14.3	18.3
NH ₃	%	5.44E−03	3.18E−02	5.91E−02	4.63E−01	4.38E−02	4.56E−02
Furfural	%	2.58E−02	7.19E−01	8.41E−05	6.52E−03	6.46E−03	9.03E−02
Phenol	%	1.21E−01	3.99E−01	2.41E−04	1.55E−03	1.40E−01	1.81E+00
Naphthalene	%	1.28E−02	1.62E−02	1.69E−07	2.14E−06	1.34E−02	6.80E−01
Pyridine	%	4.17E−03	1.83E−02	1.06E−04	2.94E−03	6.54E−03	6.58E−02
Char	%	5	23	6	24	10	10

- Efficiency depends on the type of gasification



$$\varepsilon_{cg} = \frac{\Delta h_{gas}^0 \dot{m}_{gas}^-}{\Delta h_{wood}^0 \dot{m}_{wood}^+}$$

Fig. 4 – Cold gas efficiencies in parameter domain.

- Composition depends on gasifying agent
- Composition depends on gasifier technology
- Composition depends on operating composition

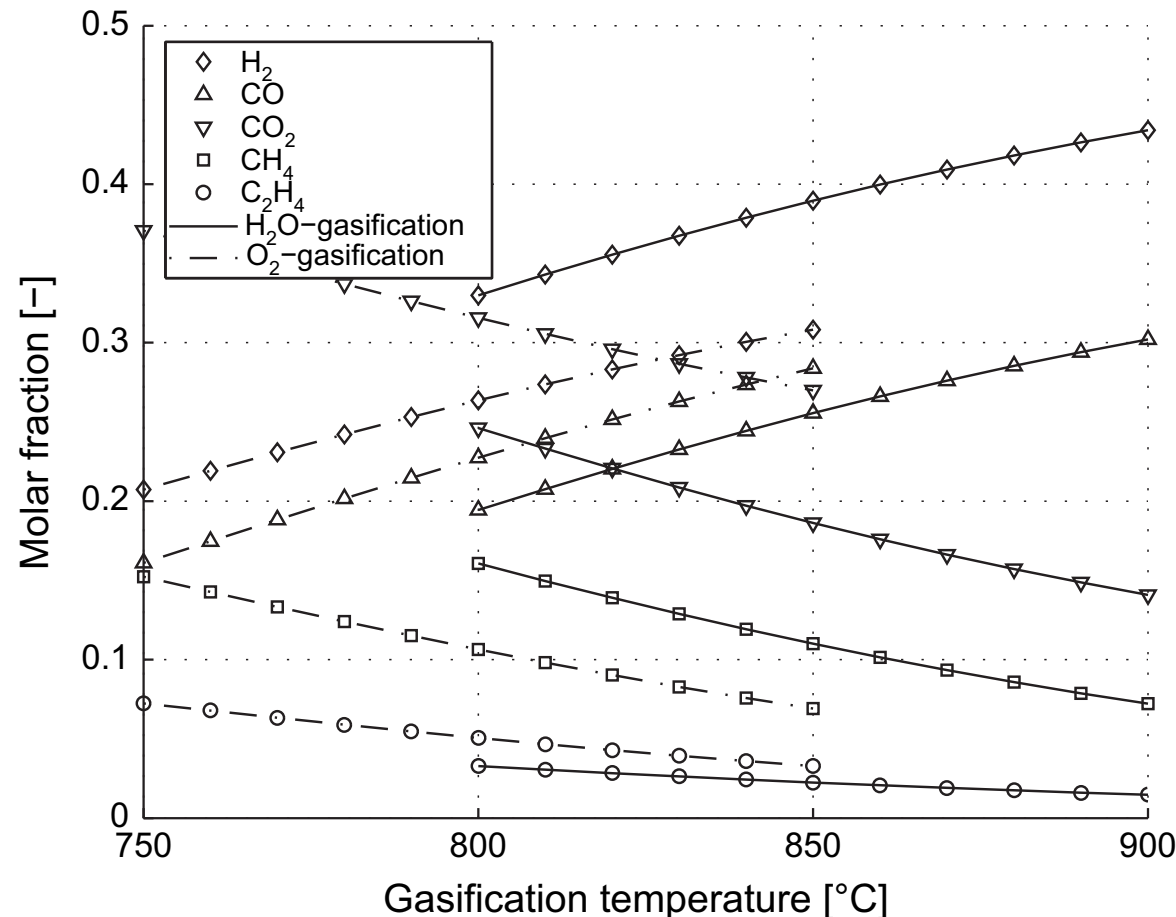
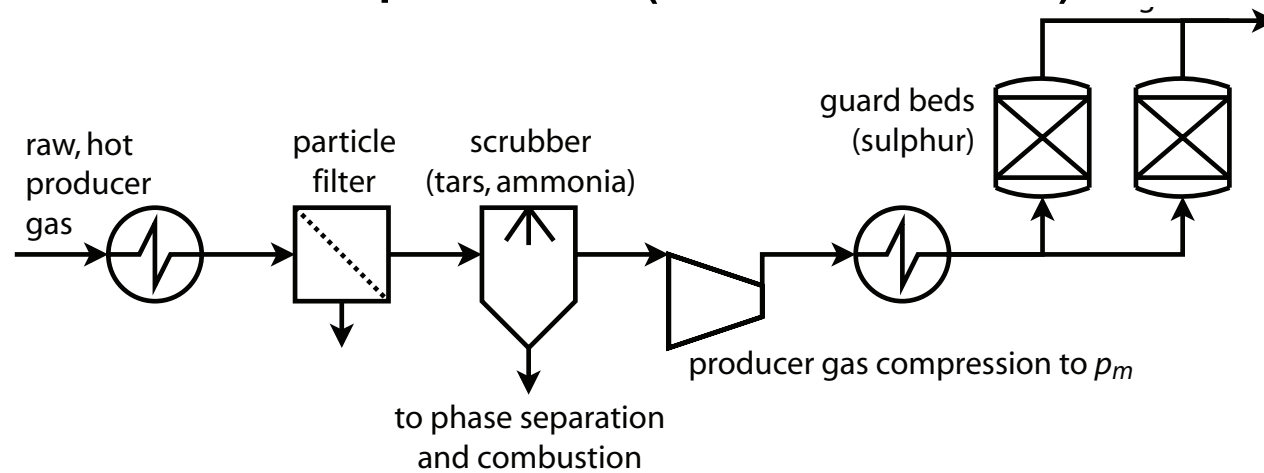
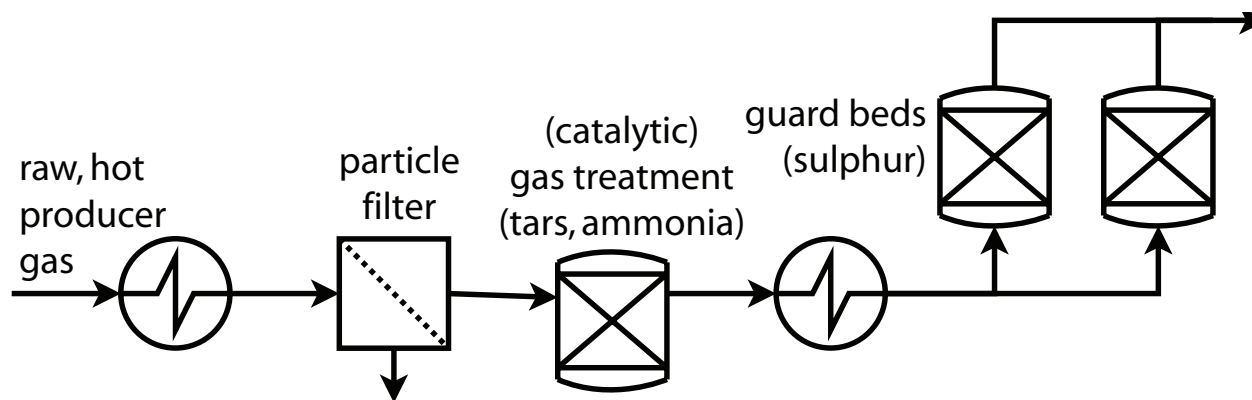


Fig. 3 – Molar compositions of the dry gas from gasification.

- Remove the impurities (inc. H_2S , S_2)

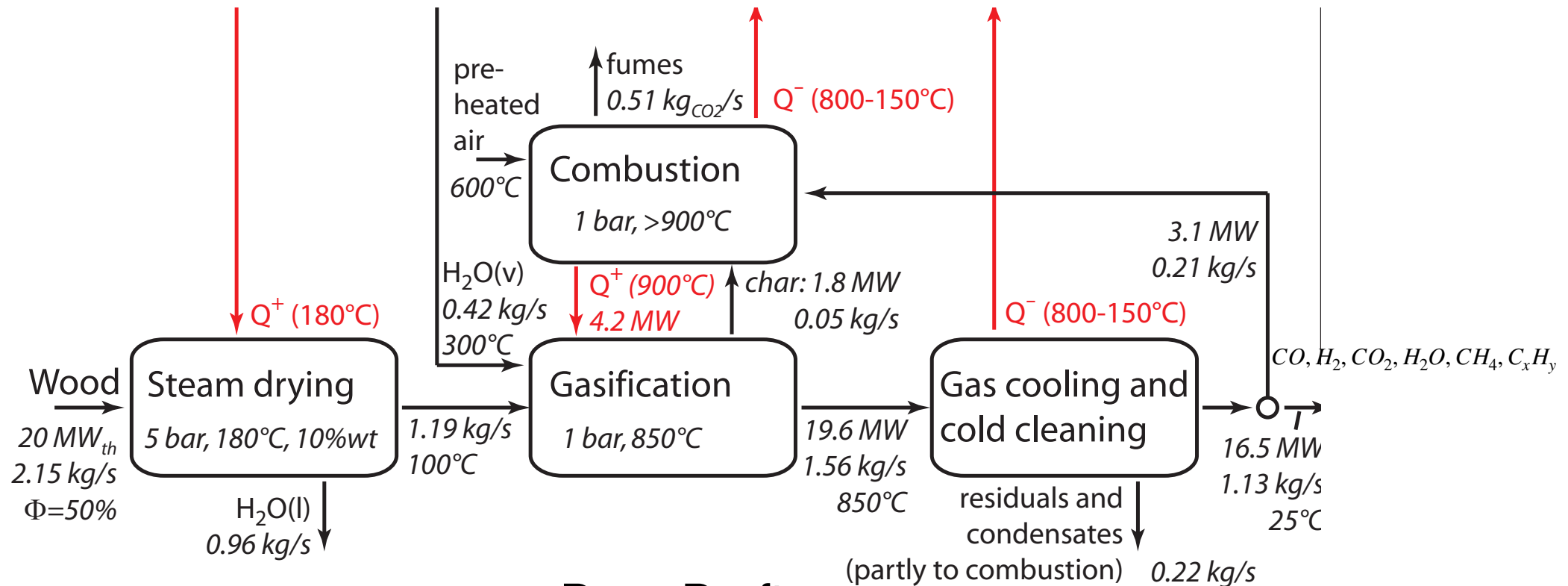


(a) Cold gas cleaning.



(b) Hot gas cleaning.

SOLID -> SYNGAS



DownDraft
Entrained flow
Indirectly heated
Multi-stage

- CO and H₂ are produced (carry the LHV : eff : 80%)
- Gasifying Agent
 - O₂
 - Air
 - Steam
- Heat by oxidation of the part of the biomass
 - Heating : direct or indirect
 - Fluidised bed material (can be catalytic)
- Design, operating conditions and residence time are critical
 - Solid-Gas : drying - pyrolysis - gas release
 - Gas phase : reforming reactions
- Ash separation
- Gas cleaning
- Heat recovery