Effect of impurity on thermally self-sustained double reactor coupling hydrogen production from glycerol reforming and methanol production from carbon dioxide and hydrogen

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Abstract. Thermally self-sustained double reactor (TSSDR) operating without external heat source consists of dual channels for endothermic and exothermic reactions. Hydrogen (H2) is produced from wasted glycerol by aqueous-phase glycerol reforming (APGR) at 200-250 °C and 20-25 bar while carbon dioxide (CO₂) is a by-product. Produced H₂ and CO₂ are used as raw materials for methanol synthesis (MS) at 200-250 °C and 50-80 bar. Methanol synthesis and glycerol reforming occur at inner and outer channels of TSSDR, respectively. The TSSDR is fully packed with catalyst. Generated heat of exothermic reaction is sufficient for endothermic reaction. Main products of glycerol reforming in gas phase are H₂ and CO₂ while CO and CH₄ are by-products. All products in gas phase are totally recycled as a feed stream for exothermic channel. CO and CH₄ in feed reduce CO₂ conversion and methanol yield in MS. The effect of impurities in glycerol feed stream also influences with hydrogen production in APGR. Especially, methanol, which is an impurity in glycerol feed obtained from biodiesel production, significantly reduces glycerol conversion in TSSDR.

1 Introduction

Methanol is an essential chemical and primary feedstock for paraffins, olefins and various organic compounds such as acetic anhydride, acetic acid and formaldehyde [1]. MS is generally carried out in gas phase at high pressure and temperature of 50-80 bar and 200-300 °C with Cu/ZnO/Al₂O₃ as a catalyst. As this reaction is exothermic, researchers have devised schemes to use heat released from MS in other reactions. Rahmanifard et al. [2] studied MS in thermally coupled membrane where MS is a heat source for cyclohexane dehydrogenation and Nimcar et al. [3] utilized the exothermic heat from MS in benzene production. Generally, methanol is produced from reformed natural gas (syngas), resulting in CO₂ emission which has negative impact to environment. Methanol can be produced from CO₂ conversion which can be a promising method to reduce CO₂ emission while producing higher value products. However, hydrogen (H₂) source and availability are still a major barrier for CO₂ conversion.

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Glycerol is waste from biodiesel production and can be available source for H_2 production. Normally, glycerol reforming is endothermic reaction is generated in gas phase at high temperature (400-700 °C) [4]. Glycerol reforming can be carried out at different operating condition-autothermal glycerol reforming, photo-reforming and aqueous phase glycerol reforming (APGR). The features of APGR are that no vaporization of the feedstock is required, which could decrease the input energy compared to steam reforming. This reaction can operate at low temperature (200 – 250 °C) [5, 6]. Commercial catalysts for APGR is noble metal on alumina or carbon. Pt/Al₂O₃ can be used as a catalyst and was reported to provide glycerol conversion around 50-60 % [7] with a feed glycerol to water ratio of 1:9. However, production of H₂ from glycerol produces CO₂ as by-product. Direct utilization of CO₂ with H₂ obtained from APGR can be another promising pathway for glycerol conversion with mitigating CO₂ emission. APGR's products in gas phase consist of H₂, CO₂, CH₄ and CO [8]. Methanation and reverse water gas shift reaction (RWGS) is likely occurred in MS at low temperature [9].

In this study, a novel thermally self-sustained double reactor (TSSDR) coupling endothermic APGR with exothermic MS is proposed. Heat transfer between the two reactions is enabled in this reactor and lead to more favourable conditions for higher reactant conversion. H_2 and CO_2 produced from APGR can be totally utilized in MS for methanol production. A computational fluid dynamic (CFD) model was established to study this TSSDR. Effects of impurities in APGR and MS were investigated.

2 Model description

The TSSDR in our study consists of two concentric channels: 1) an inner tube and 2) an outer annular channel. Fig. 1 shows a schematic diagram of this TSSDR. The diameter of the inner channel is 16 mm and the thickness of the annulus channel is 8 mm. The wall between the two channels is 2 mm thick. The reactor height is 150 mm. The inner channel is intended for MS and the outer channel is designed for APGR. Cu/ZnO/Al₂O₃ catalyst for MS is packed in the inner channel, through which CO₂ and H₂ are supplied. A Pt/Al₂O₃ catalyst for APGR is packed in the outer channel. Generated heat from the exothermic reaction in the MS channel is expected to transfer to the endothermic APGR channel. Both feed streams are fed vertically from the bottom to the top of the reactor as a base case configuration.



Fig. 1. TSSDR schematic design

2.1 Kinetic rate

2.1.1 Methanol synthesis (MS)

To calculate TSSDR model, methanol is synthesized by CO_2 hydrogenation. Busshe et al. [10] explains the kinetic rate of MS with Cu/ZnO/Al₂O₃ at 200-250 °C. In methanol production, CH₄ and CO are undesired products in process. Possible reactions in MS are shown in Table 1.

Table 1. Possible reactions and kinetic rates in MS [10,11]

Possible reaction	Rate expression
CO ₂ hydrogenation:	$R_{H_2} = \frac{\left(k_1 P_{H_2} P_{H_2 O}\right) - \left(k_6 P_{CO_2} P_{CO} / P_{H_2 O}^2\right)}{\left(k_1 P_{H_2 O} + \frac{1}{2}\right)}$
$CO_2 + 3H_2 \leftrightarrow CH_3OH + 2H_2O$	$(1 + (k_2 P_{CO_2} P_{H_2O}) + (k_3 P_{H_2O}^{0.5}) + (k_4 P_{CO_2}))^3$
CO hydrogenation:	$P_{L} = k \left(P_{L} P_{2}^{2} - \frac{P_{CH_{3}OH}}{P_{CH_{3}OH}} \right)$
$CO + 2H_2 \leftrightarrow CH_3OH$	$K_{MS,2} = K_{1,0}(r_{CO}r_{H_2} - \frac{K_{1,0}}{K_1})$
Methanol dehydration:	$P_{CH_3OH}P_{H_2O}$
$2CH_3OH \leftrightarrow CH_3OCH_3 + H_2O$	$K_{DME} = K_{2,0}(F_{CH_3OH} - \frac{K_2}{K_2})$
Reverse water gas shift (RWGR):	$(k_5 P_{H_2}) - (k_7 P_{CO_2} P_{CH_3OH} / P_{H_2O})$
$CO_2 + H_2 \leftrightarrow CO + H_2O$	$K_{RWGS} = \frac{1}{(1 + (k_2 P_{CO_2} P_{H_2O}) + (k_3 P_{H_2O}^{0.5}) + (k_4 P_{CO_2}))}$
Methanation:	$P_{CH_4}P_{H_2O_1}$
$CO + 3H_2 \leftrightarrow CH_4 + H_2O$	$\kappa_{methane} - \kappa_{3,0} (r_{CO} r_{H_2} - \frac{K_3}{K_3})$

2.1.2 Aqueous-phase glycerol reforming (APGR)

Glycerol and water were feedstock of APGR for hydrogen production with CO_2 as a byproduct. Iliuta et al. [12] formulated a kinetic model of APGR with a Pt/Al₂O₃ catalyst in a trickle bed reactor. Arely et al. [8] investigated the APGR in a batch slurry reactor. Methane, ethane and propane were detected as gaseous by-products while ethylene glycol and alcohols were detected as liquid products. The rate for side reactions in APGR are show in Table 2 when $k_{i,0}$ is calculated by Arrhenius's equation.

Possible reaction	Rate expression
$C_3H_8O_3 + 3H_2O \rightarrow 3CO_2 + 7H_2$	$R_{APGR,1} = 2.678 \exp\left(-\frac{60000}{RT}\right) C_{C_3 H_8 O_3}$
$C_3H_8O_3 \rightarrow 3CO \ + \ 4H_2$	$R_{APGR,2} = k_{2,0} \exp\left(\frac{-83990}{RT}\right) C_{C_3 H_8 O_3}$
$CH_3OH + H_2 \rightarrow CH_4 + H_2O$	$R_{APGR,3} = k_{3,0} \exp\left(\frac{-84510}{RT}\right) C_{CH_3OH} \frac{C_{H_{2,g}}}{H_{H_2}}$
$CO + H_2O \rightarrow CO_2 + H_2$	$R_{APGR,4} = k_{4,0} \exp\left(\frac{-130300}{RT}\right) \frac{C_{CO_g}}{H_{CO}}$
$CO + 3H_2 \rightarrow CH_4 + H_2O$	$R_{APGR,5} = k_{5,0} \exp\left(\frac{-133260}{RT}\right) \frac{C_{H_{2,g}}^{1.5}}{H_{H_{4}}}$
$CH_3OH \rightarrow CO + 3H_2$	$R_{APGR,6} = \frac{(1.17 \times 10^{-7} \exp(\frac{-40999}{RT}))C_{CH_3OH}}{(7.10 \times 10^{-7} \exp(\frac{-40999}{RT}))C_{CH_3OH}}$
$3CO_2 + 4H_2 \rightarrow CH_4 + 2H_2O$	$R_{APGR,7} = (1.75 \times 10^{-7}) \exp\left(\frac{-95000}{RT}\right) C_{CO_2}^{0.66} C_{H_2}^{0.21}$

Table 2. Possible reactions and kinetic rates in APGR [8	, 9]
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2.3 Simulation method

MS and APGR in TSSDR were simulated with COMSOL Multiphysics 5.3a. Steady-state flows of the feed gas mixtures was solved by mass, momentum, and energy balance equations of a discretized flow domain of the reactor geometry. The dual reactions were simulated in a 2D-axis-symmetric system in cylindrical coordinate. The flow regime in each channel was assumed to be steady and laminar. The diffusion of materials and the heat conduction in direction flow axis was neglected. The ideal gas and Peng-Robinson model was employed. The physical properties of the chemicals were initially defined with Perry's Chemical Engineers' Handbook [13]. The mathematical equations for the CFD model were divided into three parts. For Part I, MS was calculated at the surface of the solid catalyst. A constant bed void fraction was maintained for the entire length of the reactor. For Part II, APGR was calculated with three phases (gas-phase products, liquid-phase reactants and solid catalyst). For Part III, heat transfer through the solid wall was determined. Numerical solutions were obtained with the finite element method in COMSOL Multiphysics version 5.3a. The mesh resolution was free triangular with a mesh area of 0.0012 m² and the number of elements of 1416. Performance of both reactions are measured by conversion, yield and selectivity, show in Eq. (1-6).

$$CO_2 \ conversion = \frac{mol \ of \ CO_2 \ inlet - mol \ of \ CO_2 \ outlet}{mol \ of \ CO_2 \ inlet} \times 100$$
(1)

$$C_3 H_8 O_3 \text{ conversion} = \frac{\text{mol of } C_3 H_8 O_3 \text{ inlet-mol of } C_3 H_8 O_3 \text{ outlet}}{\text{mol of } C_3 H_8 O_3 \text{ inlet}} \times 100$$
(2)

% yield of
$$H_2 = \frac{mol \ of \ H_2 \ generated}{mol \ of \ C_2 \ H_2 \ O_2 \ feed} \times 100$$
 (3)

% yield of
$$CH_3OH = \frac{mol of CH_3OH generated}{mol of CO_5 ford} \times 100$$
 (4)

% selectivity of APGR product =
$$\frac{\text{mol of desired product}}{\text{mol of C_2H_2O_2 reacted}} \times 100$$
 (5)

% selectivity of MS product =
$$\frac{mol \ of \ desired \ product}{mol \ of \ CO_2 \ reacted} \times 100$$
(6)

3 Result & Discussion

Simulation for the model validation was performed for the MS and APGR channels separately in adiabatic model. The MS model accuracy was assured at an average error of only 0.88 % based on average of all components when compared with Busshe et al [10]. In APGR, the average error is 2.82% when comparing output composition with Iliuta et al. [12]. This research divides TSSDR model into three cases. Case I is the base case condition: MS inlet temperature of 220 °C; MS inlet pressure of 50 bar; MS total flow rate of 0.003 mol s⁻¹; MS external feed H₂ to CO₂ ratio of 82:3; APGR inlet temperature of 220 °C ; APGR inlet pressure of 20 bar; MS total flow rate of 0.005 mol s⁻¹; APGR external feed glycerol to water ratio of 1:9. Case II is the TSSDR model without external feed of H₂ and CO₂ in MS feed stream. MS feed totally relied on APGR's gas products. Case III is the TSSDR model was similar to case II plus methanol as impurity in APGR feed side.

As presented in Fig 2, methanol yield and CO_2 conversion in base case were higher than other conditions since H_2 was highly excess. CO_2 hydrogenation is based on theoretical ratio of H_2 to CO_2 at 3. There was a reaction between hydrogen and CO (CO hydrogenation). Therefore, methanol yields in base case was higher than CO_2 conversion. On the other hand, total recycle of APGR's gas product to MS influenced CO_2 conversion decreased to 36.36% because the ration of H_2 to CO_2 in feed was decreased to 3:1. Gas products in APGR consisted of the main product (H_2 , CO_2) and undesired products (CO, CH_4 and methanol) which led to a reduction in methanol production.



Fig. 2. Comparing CO₂ conversion and CH₃OH yield in exothermic side: (1) TSSDR with external feed H₂ and CO₂, (2) TSSDR with total recycle APGR's gas product, (3) Effect of methanol impurity in APGR's feed stream (ratio = 1:80), (4) Effect of methanol composition in APGR's feed stream (ratio = 1:40), (5) Effect of methanol composition in APGR's feed stream (ratio = 1:20), (6) Effect of methanol composition in APGR's feed stream (ratio = 1:10).

As presented in Fig. 3, there were a minor change in H_2 yield and glycerol conversion in all cases because the molar of pure glycerol in water solution was constant although methanol was introduced as impurity in feed.



Fig. 3. Comparing $C_3H_8O_3$ conversion and H_2 yield in endothermic side: (1) TSSDR with external feed H_2 and CO_2 , (2) TSSDR with total recycle APGR's gas product, (3) Effect of methanol composition in APGR's feed stream (ratio = 1:80), (4) Effect of methanol composition in APGR's feed stream (ratio = 1:40), (5) Effect of methanol composition in APGR's feed stream (ratio = 1:20), (6) Effect of methanol composition in APGR's feed stream (ratio = 1:10).

In reality, crude glycerol which is obtained from biodiesel production could contain methanol as impurity. Fig. 4 shows the effect of methanol in glycerol feed in APGR on product selectivity in MS. Selectivity of MS's product insignificantly decreased. There was a separation of liquid product by condensation before recycling to MS. Methanol was removed before entering MS.



Fig. 4. Selectivity of MS's product base on CO₂ reacted: (1) TSSDR with external feed H₂ and CO₂, (2) TSSDR with total recycle APGR's gas product, (3) Effect of methanol composition in APGR's feed stream (ratio = 1:80), (4) Effect of methanol composition in APGR's feed stream (ratio = 1:40), (5) Effect of methanol composition in APGR's feed stream (ratio = 1:20), (6) Effect of methanol composition in APGR's feed stream (ratio = 1:10).

Fig. 5 presents effect of methanol impurity in APGR feed on the selectivity of APGR products. Methanol selectivity dropped to the negative value. Generated methanol is less than molar feed of methanol. Methanol could decompose to H_2 and CO and generated CH₄ following the reaction presented in Table 2. Methanol in glycerol solution decreased when the ratio of impurity increased.



Fig. 5. Selectivity of APGR's product base on theoretical $C_3H_8O_3$ reacted: (1) TSSDR with external feed H₂ and CO₂, (2) TSSDR with total recycle APGR's gas product, (3) Effect of methanol composition in APGR's feed stream (ratio = 1:80), (4) Effect of methanol composition in APGR's feed stream (ratio = 1:40), (5) Effect of methanol composition in APGR's feed stream (ratio = 1:20), (6) Effect of methanol composition in APGR's feed stream (ratio = 1:10).

4 Conclusion

Thermally self-sustained double reactor (TSSDR) consisted of dual channels for APGR and MS. TSSDR could operate without external heat and external feed supplied in MS. H₂ and CO₂ was produced APGR and was recycled to produce methanol in MS. Therefore, byproducts in APGR affected methanol production in MS. The effect of methanol in glycerol feed stream influenced hydrogen production in APGR. Therefore, feed compositions in MS and APGR were important variables and significantly affected the reactor's performance.

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