

Editorial

# Catalysts for Syngas Production

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Synthesis gas (or syngas) is a mixture of hydrogen and carbon monoxide, that may be obtained from alternative sources to oil, such as natural gas, coal, biomass, organic wastes, etc. [1–3] Biomass is a promising raw material for syngas production, due to its renewable character and potentially zero CO<sub>2</sub> emissions [4]. Syngas is an excellent intermediate for the production of high value compounds at the industrial scale, such as hydrogen, methanol, liquid fuels, and a wide range of chemicals.

This Special Issue on “Catalysts for Syngas Production” shows new research about the development of catalysts and catalytic routes for syngas production, and the optimization of the reaction conditions for the process.

This issue includes ten articles. Yu et al. analyze the performance of Ni-Co bi-metallic catalysts in *n*-decane steam reforming [5]. The addition of Co to the catalyst improves the hydrogen selectivity and anti-coking ability compared with the mono-Ni/Ce-Al<sub>2</sub>O<sub>3</sub> catalyst. A synergistic effect between Ni and Co is observed, with 12% Co showing the best catalytic activity in the series Co-Ni/Ce-Al<sub>2</sub>O<sub>3</sub> catalysts. In situ regeneration of a spent alumina-supported cobalt-iron catalyst for catalytic methane decomposition is reported by Fakeeha et al. [6] The main factors responsible for the catalyst deactivation are coke deposition and weak sintering of the metallic active phase (Co-Fe), which occur during the catalytic methane decomposition reaction and regeneration process. A facile fabrication of supported Ni/SiO<sub>2</sub> catalysts for dry reforming of methane is developed by Xu et al. [7] Due to the formation of much smaller Ni nanoparticles, this Ni/SiO<sub>2</sub> catalyst exhibits excellent coke-resistance performance and effectively suppresses the side reaction toward RWGS compared to that prepared with the conventional wetness impregnation method. The dry reforming of methane over combined magnesia, ceria and nickel catalysts, supported on  $\gamma$ -Al<sub>2</sub>O<sub>3</sub> and doped with TiO<sub>2</sub>, is investigated by Al-Fatesh et al. [8] The addition of CeO<sub>2</sub> and MgO to the catalyst enhances the interaction between the Ni and the support, and improves the activity of the solid. Liu et al. describe a novel one-step conversion of CO<sub>2</sub> and H<sub>2</sub>S to syngas induced by non-thermal plasma, with the aid of Ni-Mo sulfide/Al<sub>2</sub>O<sub>3</sub> catalyst under ambient conditions [9]. The optical and structural properties of the synthesized catalysts are significantly influenced by the Ni/Mo molar ratio. Moreover, the Ni-Mo sulfide/Al<sub>2</sub>O<sub>3</sub> catalysts possess excellent catalytic activities for CO<sub>2</sub> and H<sub>2</sub>S conversion, compared to the single-component NiS<sub>2</sub>/Al<sub>2</sub>O<sub>3</sub> and MoS<sub>2</sub>/Al<sub>2</sub>O<sub>3</sub> catalysts. The paper by Park et al. describes the effect that reaction parameters have on hydrogen production via steam reforming of methane, using lab- and bench-scale reactors to identify critical factors for the design of large-scale processes [10]. The temperature at the reactor bottom is crucial for determining the methane conversion and hydrogen production rates when a sufficiently high reaction temperature is maintained (above 800 °C). However, if the temperature of one or more of the furnaces decreases below 700 °C, the reaction is not equilibrated at the given space velocity. Liu et al. study a novel sulfur tolerant water gas shift catalyst (SWGS) developed for the applications under lean (low) steam/gas ratio conditions [11]. The adoption of the lean steam/gas SWGS catalyst significantly improves the plant efficiency and safety, and remarkably reduces the actual steam consumption for H<sub>2</sub> production, decreasing CO<sub>2</sub> emission. The paper by Fasolini et al. summarizes the synthesis, characterization and catalytic behavior of Rh-based catalysts, obtained by using the

Rh<sub>4</sub> (CO)<sub>12</sub> neutral cluster as the active-phase precursor [12]. The preparation method allows the deposition of the cluster on the surface of Ce<sub>0.5</sub>Zr<sub>0.5</sub>O<sub>2</sub> and ZrO<sub>2</sub> supports, which are synthesized by the microemulsion technique, being the catalysts active in the low-temperature steam reforming process for syngas production. Methane and ethane steam reforming over MgAl<sub>2</sub>O<sub>4</sub>-supported Rh and Ir catalysts is analyzed in the paper by Lopez et al. [13] The Rh- and Ir-supported catalysts exhibit higher activity than Ni catalysts for steam methane reforming. Catalyst durability studies reveal the Rh catalyst to be stable under steam methane reforming conditions. The results of this study conclude that a Rh-supported catalyst enables very high activity and excellent stability, for both the steam reforming of methane and other higher hydrocarbons contained in natural gas, and under conditions of operation that are amendable to solar thermochemical operations. In the paper by Azara et al., iron-rich mining residue is used as a support to prepare a new Ni-based catalyst for C<sub>2</sub>H<sub>4</sub> dry reforming and catalytic cracking [14]. The deposited carbon is found to be filamentous and of various sizes (i.e., diameters and lengths). The analyses of the results show that iron is responsible for the growth of carbon nanofilaments and nickel is responsible for the split of C-C bonds.

In summary, these ten papers clearly show the relevance of obtaining syngas for further applications, such as the production of hydrogen, methanol, liquid fuels, and a wide range of chemicals. Nowadays, efforts are being made on the co-feeding of CO<sub>2</sub> with syngas, as an alternative for reducing greenhouse gas emissions. I would like to thank all the authors of this Special Issue.

I am honored to be the Guest Editor of this Special Issue. I would like to thank the reviewers for improving the quality of the papers with their comments. I am also grateful to all the staff of the Catalysts Editorial Office.

**Conflicts of Interest:** The authors declare no conflict of interest.

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