## Combined steam and CO<sub>2</sub> reforming of methane for syngas production over carbon-resistant boron-promoted Ni/SBA-15 catalysts

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## A R T I C L E I N F O

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## ABSTRACT

The unpromoted and B-promoted 10%Ni/SBA-15 catalysts synthesized via sequential incipient wetness impregnation approach were assessed for combined steam and  $CO_2$  reforming of methane (CSCRM) at various reaction temperatures of 973–1073 K and stoichiometric feed composition. An expected and noteworthy drop in mean NiO crystallite size and BET surface area with boron promotion from 1% to 5%B loading could be due to the agglomeration of  $B_2O_3$  particles and deboration reaction during calcination and hence blocking mesopores of SBA-15 support at elevated B composition. The complete NiO reduction to metallic Ni<sup>0</sup> form was achieved during H<sub>2</sub> activation and the reduction temperature of NiO phase was shifted towards higher temperature with B addition owing to enhancing interaction between the acidic  $B_2O_3$  and basic NiO phases. For all reaction temperature employed, 3%B appeared to be the optimal promoter loading in terms of reactant conversions and 3%B-10%Ni/SBA-15 catalyst revealed the greatest H<sub>2</sub> yield (69.4%) at 1073 K. In addition, CH<sub>4</sub> and CO<sub>2</sub> conversions were enhanced about 23.2% and 32.4%, correspondingly with rising reaction temperature from 973 to 1073 K. Ratio of H<sub>2</sub> to CO varied from 1.26 to 2.71 and the desired H<sub>2</sub>/CO ratio of about 2 favored for Fischer-Tropsch synthesis was achieved on 3%B-10%Ni/SBA-15 sample at 973 K. Boron promoter suppressed graphitic carbon formation and the amount of carbonaceous deposition was reduced about 4 times. Noticeably, 3%B-10%Ni/SBA-15 was also resilient to metallic Ni<sup>0</sup> re-oxidation throughout CSCRM.