

## The Influence of Different Mn-Co interaction Modes in Mn/Co/TiO<sub>2</sub> Fischer-Tropsch Catalysts

Jingjing Liu<sup>1</sup>, Theresa E. Feltes<sup>1</sup>, Yuan Zhao<sup>2</sup>, Juan-Carlos Idrobo<sup>3</sup>, Robert F. Klie<sup>2</sup>, Jeffrey T. Miller<sup>4</sup>, Christopher L. Marshall<sup>4</sup>, John R. Regalbuto<sup>1,5</sup>, and Randall J. Meyer<sup>1</sup>

<sup>1</sup> Dept. of Chemical Engineering, University of Illinois at Chicago

<sup>2</sup> Dept. of Physics, University of Illinois at Chicago

<sup>3</sup> Materials Science and Technology Division, Oak Ridge National Laboratory,

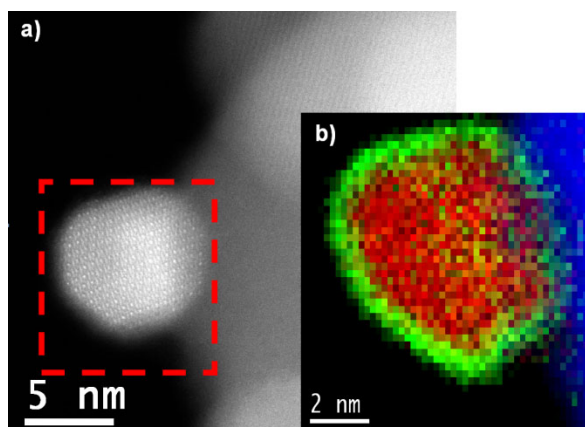
<sup>4</sup> Chemical Science and Engineering Division, Argonne National Laboratory

<sup>5</sup> Dept. of Chemical Engineering, University of South Carolina

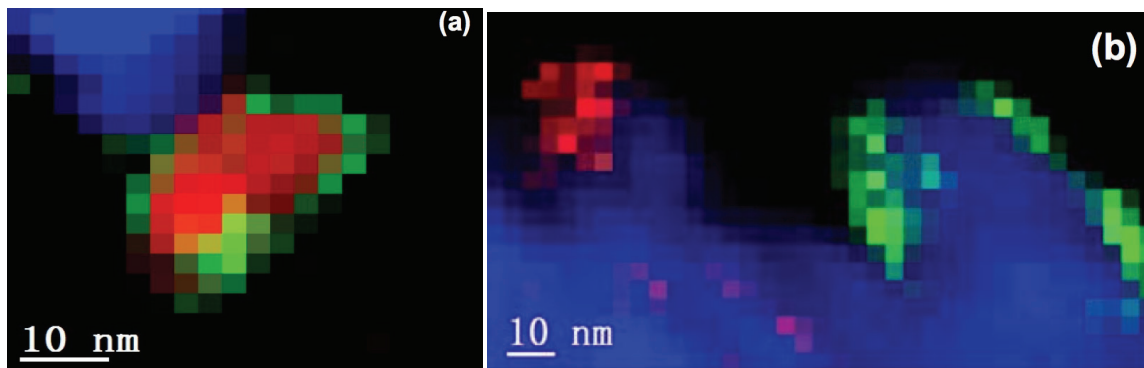
The unpredictable turbulence of crude oil prices has largely defined the prominence of research towards alternative fuel processes, most notably the Fischer-Tropsch (FT) process [1]. The Cobalt based FT catalysts are well known for their activity and stability [2]. In general, cobalt catalysts exhibit both higher activity and selectivity than their iron counterparts. For FT synthesis, Mn is often used as a promoter for both supported and unsupported Co catalysts [3].

The addition of promoter onto supported metal catalysts is a prevalent way in making catalysts to enhance the reactivity results, but it is also a poorly understood phenomenon in heterogeneous catalysis. Being a local effect, intimate interaction of the promoter and the metal is highly desired; therefore, a key design objective is to increase the promoter-metal interactions to maximize their effectiveness. We achieved this goal of creating a monolayer of Mn promoter on Co metal (confirmed by STEM-EELS) by using the Strong Electrostatic Adsorption (SEA [4]), which utilizes pH control to steer the promoter precursor (in this case MnO<sub>4</sub><sup>-</sup>) onto Co oxide supported on TiO<sub>2</sub>. And this type of catalyst showed promising Fischer-Tropsch (FT) reactivity results in the catalytic testing by comparing with our previous studied FT catalysts, which are Mn promoted on Co/TiO<sub>2</sub> by SEA (partial but not core-shell interaction) and Mn promoted on Co/TiO<sub>2</sub> by Dry Impregnation (most common way of catalysts synthesis in industry but least promoter-metal interaction).

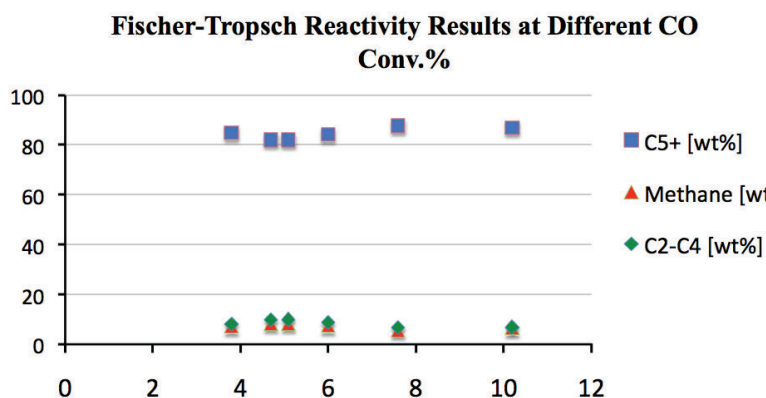
In this presentation, the microscopy results (imaging and EELS mapping, Figure 1 and 2) is linked with the CO hydrogenation reactivity results (Figure 3) to show how will the different promoter-metal interactions affect the FT reaction of producing long chain hydrocarbons.



**Figure 1:** . a) Atomic resolution Z contrast imaging of monolayer Mn promoted Co catalyst supported on TiO<sub>2</sub>. b) EELS mapping showed distinct elemental composition, in the graph, green is Mn, red is Co and blue is TiO<sub>2</sub>. Imaging and spectra acquired from Oak Ridge National Laboratory by using a Nion UltraSTEM.



**Figure 2** a) EELS mapping of  $\text{Mn}^{\text{SEA\_partial}}/\text{Co}/\text{TiO}_2$  catalys, b) EELS mapping of  $\text{Mn}^{\text{DI}}/\text{Co}/\text{TiO}_2$  catalyst, in the graph, green is Mn, red is Co and blue is  $\text{TiO}_2$ . Imaging and spectra acquired from UIC RRC center by using JEOL-2010 STEM.



**Figure 3:** Fischer-Tropsch reactivity results at various CO conversion point by using the  $\text{Mn}^{\text{SEA\_monolayer}}/\text{Co}/\text{TiO}_2$  catalyst. The FT catalytic testing was performed at Argonne National Laboratory.

#### References:

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- [5] This work is supported by a grant from the National Science Foundation (CBET-1067020).