

# DOE Bioenergy Technologies Office (BETO) 2021 Project Peer Review **Upgrading of C2 Intermediates – ORNL**

Catalytic Upgrading

March 10, 2021

**Zhenglong Li** Oak Ridge National Laboratory



## Integrated and collaborative portfolio of catalytic technologies and enabling capabilities

		• •
Catalytic Technologies	Enabling Capabilities	Industry Partnerships
<b>Catalytic Upgrading of Biochemical</b>	Advanced Catalyst Synthesis and	(Phase II Directed
Intermediates	Characterization	Funding)
(NREL, PNNL, ORNL, LANL)	(NREL, ANL, ORNL)	Opus12 (NREL)
Upgrading of C1 Building Blocks	Consortium for Computational	Visolis (PNNL)
(NREL)	Physics and Chemistry	Sironix (LANL)
Upgrading of C2 Intermediates	(ORNL, NREL, PNNL, ANL, NETL)	
(PNNL, ORNL)	<b>Catalyst Deactivation Mitigation</b>	
Catalytic Fast Pyrolysis	for Biomass Conversion	
(NREL, PNNL)	(PNNL)	
Electrocatalytic CO <sub>2</sub> Utilization		
(NREL)		
	Cross-Cutting Sunnort	

Cross-Cutting Support

ChemCatBio Lead Team Support (NREL)

**ChemCatBio DataHUB** (NREL)

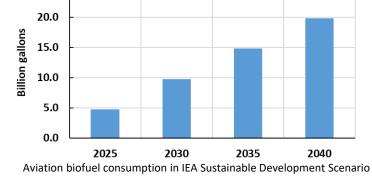
**ChemCatBio** 

# Ethanol is an attractive feedstock for middle distillate fuels and chemical products:

- Growing market of aviation biofuels
- Established **ethanol market** (~29.0 billion gallons globally in 2019)
- Lower ethanol prices over the last five years creates opportunities for jet fuel production from ethanol
  - Chemical coproduct from ethanol can further enhance the cost competitiveness
- Factors expected to expand ethanol availability:
  - Ethanol "blend wall"

ChemCatBio

- Advancement in ethanol production efficiency
- Feedstock diversification

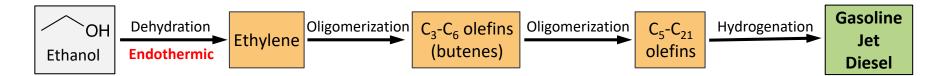


25.0



US historical ethanol selling price (source: Iowa State University)

https://www.iea.org/commentaries/are-aviation-biofuels-ready-for-take-off Renewable Fuels Association, ethanolrfa.org/statistics/annual-ethanol-production



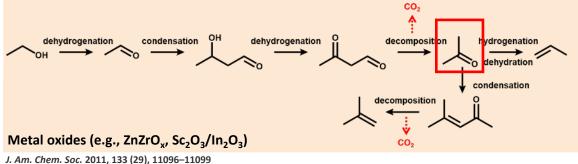
#### Limitations for conventional ETJ:

- Two-step process operation for C<sub>3+</sub> olefin production
- Ethanol dehydration is endothermic (requires significant energy input)
- Large unit operation temperature differences → increase heat management duty
  - Dehydration, 1<sup>st</sup>-stage and 2<sup>nd</sup>-stage oligomerization: e.g., 200-350, ~85 and 200-300°C
- Higher CapEx and OpEx

#### **Opportunities for developing one-step ethanol to C\_{3+} olefin (ETO) technology:**

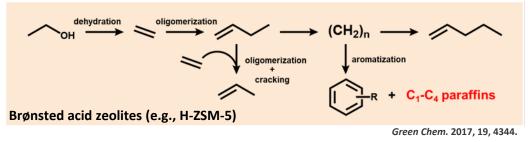
- Reduce the number of unit operations
- Minimize external energy inputs and reduce the demand of heat management
- Reduce CapEx and OpEx

ETO over oxide-based catalysts via acetone:



Chem. Lett. 2012, 41 (9), 892-894.

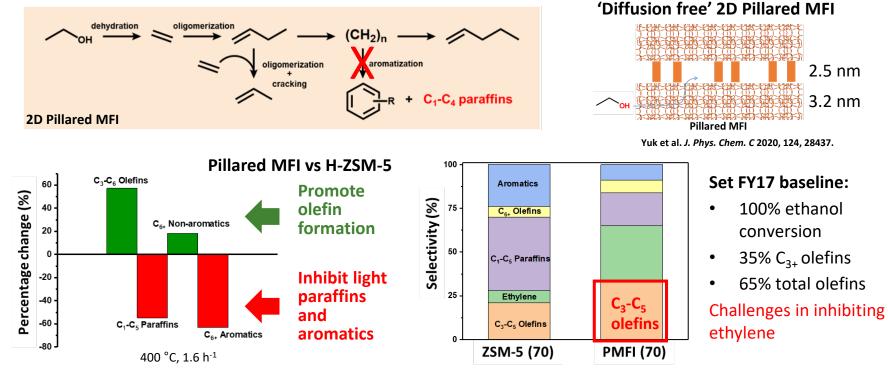
Brønsted acid-catalyzed reactions to C<sub>3+</sub> olefins:



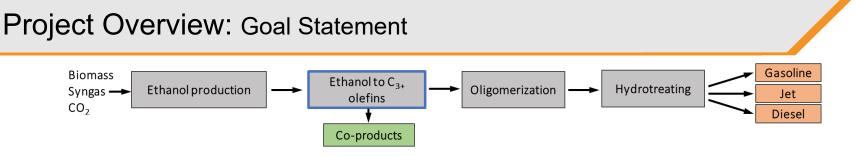
• Common challenge: significant side product formation (e.g., light paraffins, CO<sub>2</sub>)

Project Overview: 2D Pillared MFI to Optimize C<sub>3+</sub> olefins (ORNL, FY17)

Minimize downstream aromatization and light paraffin formation to promote C<sub>3+</sub> olefin formation by significantly reducing the diffusion length:



• Strong R&D need in new catalysis and reactions for one-step ETO to selectively produce C<sub>3+</sub> olefins



**Goal:** Develop efficient **one-step ETO technology** to enable high carbon efficiency to liquid hydrocarbon fuels at lower cost, and enable the development of a **market-responsive biorefinery** concept through C<sub>2</sub> platform providing control over gasoline, diesel, and jet fuel, as well as high value co-products

**Outcome:** Advance C<sub>2</sub> upgrading technology and reduce commercialization risk by addressing catalysis and process challenges and demonstrating liquid fuel production at a modeled minimum fuel selling price (MFSP) of \$3.0/GGE

### Differentiator

#### **Conventional ETJ**

- Two-step ethanol to C<sub>3+</sub> olefins
- **Higher energy demand** due to endothermic dehydration and temperature differences between unit operations
- **Higher CapEx and OpEx** (e.g., ethanol upgrading cost is \$1.19/GGE based on two-step Ziegler reaction\*)

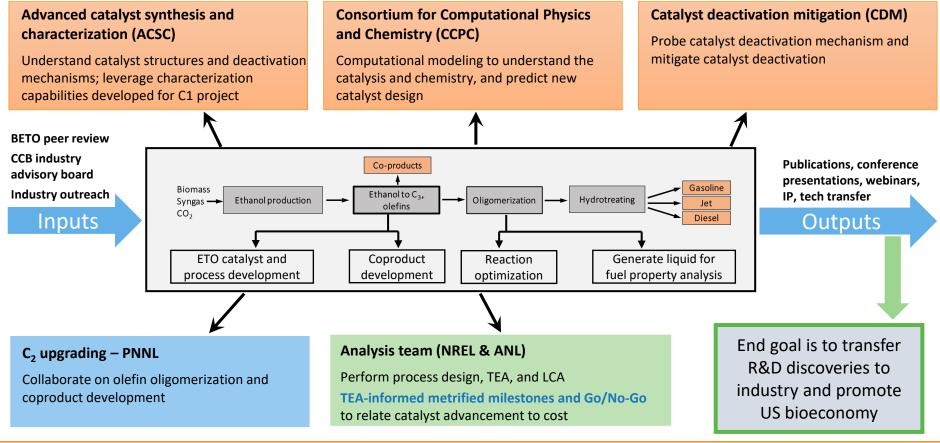
\*Tao et al. Green Chem. 2017, 19, 1082-1101.

#### ETJ via One-step ETO

- One-step ethanol to C<sub>3+</sub> olefins
- Reduced energy management burdens:
  - Shift to slightly exothermic reaction
  - o ETO and oligomerization are operated at similar T
- Lower CapEx and OpEx (ethanol upgrading cost reduced to \$0.5-0.6/GGE, based on our process)

## 1-Management Approach

Joint milestones to tackle challenges associated with catalyst development and deactivation mitigation



#### **ChemCatBio**

## 1-Management Approach: Risks and Mitigation

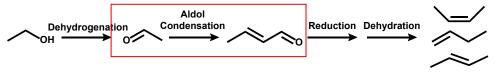
Risks	Mitigation
New ETO catalyst being developed requires investigation of <b>long-term catalyst durability</b> and <b>regeneration.</b>	Catalyst characterization, deactivation and regeneration studies are planned as part of new catalyst testing throughout the course of the project. Accelerated catalyst deactivation may be considered.
Achieving \$3/GGE distillate production may be difficult with the current fermentation-derived ethanol feedstock and process model.	TEA scenario analysis will be conducted for new concepts with <b>coproducts from either ethanol or lignin</b> .
The <b>compositions of C<sub>3+</sub> olefins</b> are not ideal for making high-quality aviation fuels.	Oligomerization of the $C_{3+}$ olefins derived from ETO will be performed and optimized to generate liquid hydrocarbons. Fuel property analysis will be carried out to guide the optimization of $C_{3+}$ olefin composition.

#### Milestones Associated with Risk Mitigations

- FY21 Q2 (C2-CDM): Determine impact of steam on Y/BEA stability and possible deactivation mechanism
- **FY21 Q3 (C2-ACSC-CCPC-CDM):** Develop a regeneration procedure and/or new catalyst formulation to optimize regenerability by characterizing deactivation mechanism(s) of Cu-Zn-Y/Beta for ETO
- **FY21 Go/No-Go:** Evaluation of pathway(s) to meet FY22 carbon efficiency and cost targets

## 2-Technical Approach: Research Challenges and Success Factors

#### Target one-step ETO via acetaldehyde & crotonaldehyde as intermediates



#### **Research Challenges**

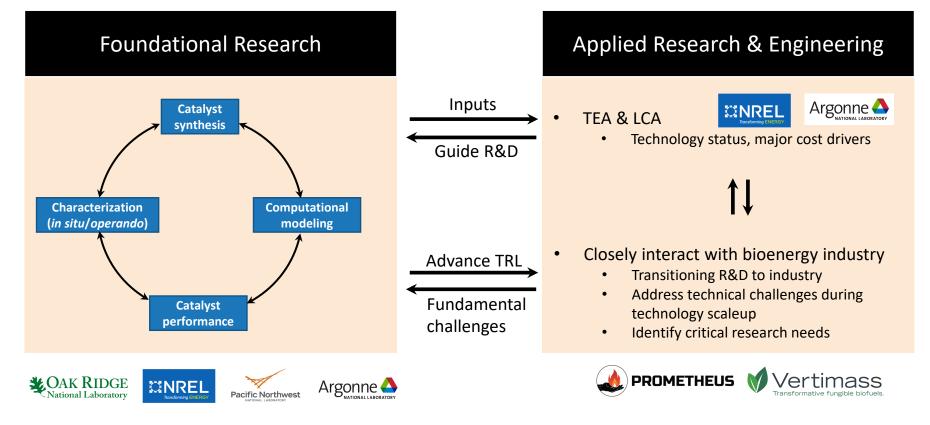
- Design and balance multiple active sites to catalyze cascade reactions to selectively produce targeted C<sub>3+</sub> olefins
- *Minimize side reactions*: ethanol dehydration, C-C cleavage to CO<sub>2</sub>, olefin over hydrogenation
- Characterize and understand the *catalyst structures, reaction mechanisms* for the purpose of designing next-generation catalyst
- Maximize catalyst lifetime

#### **Critical Success Factors**

- *Maximize carbon efficiency to liquid hydrocarbons* via active and selective multifunctional catalyst design
- Understand deactivation and develop regeneration/mitigation approaches though collaborating with ChemCatBio enabling projects (CDM, ACSC, CCPC)
- *Advance technologies* with bioenergy industry partnerships

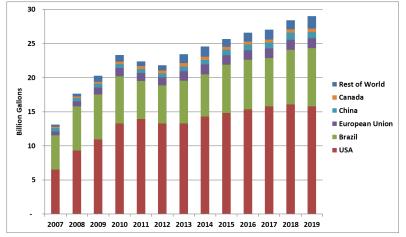
## 2-Technical Approach: Integrated Foundational and Applied Research

### Collaborative efforts to enable fundamental catalysis findings and advance technology



# New ETJ technology addresses the needs for both ethanol producers and growing markets of middle-distillate fuels

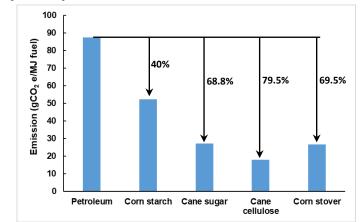
Increasing ethanol production, ethanol 'blend wall' and decreasing ethanol price urge for **new applications** 



Global Ethanol Production by Country or Region

ETJ present a great opportunity to expand the ethanol applications, **diversifying the product portfolios** for ethanol producers

Middle distillate fuels from renewable ethanol can help to address the **decarbonization** challenges for **heavy duty transportation and aviation**.



Well-to-wake analysis for jet production from different feedstocks

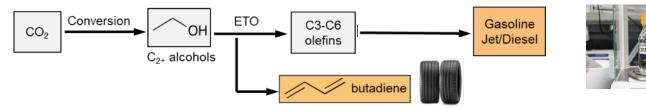
Vertimass ethanol upgrading technology licensed from ORNL shows significant opportunities for GHG emission reduction (joint study with Vertimass\*)

\*Hannon et al. PNAS. 2020, 117 (23), 12576-12583.

afdc.energy.gov/data

• ORNL licensed new ethanol-to-butadiene-jet (ETBJ) technology to Prometheus to expand the applications to CO<sub>2</sub>-derived alcohols



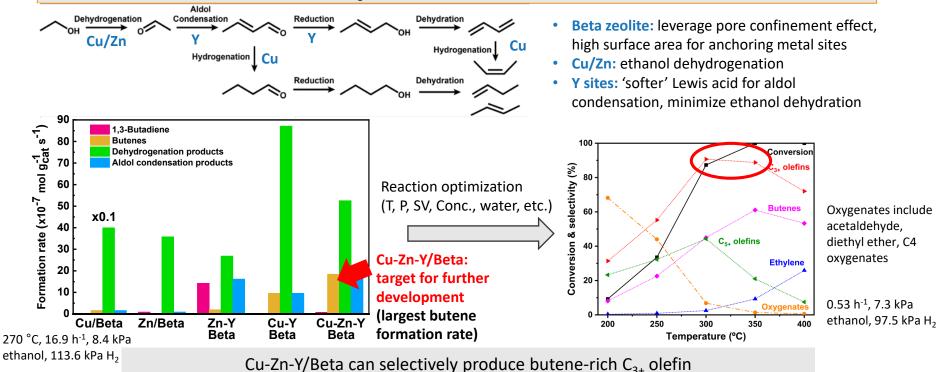




- High yield of liquid hydrocarbon fuels
- Lower CapEx and OpEx
- Flexibility to produce chemical co-product
- Additional IPs developed on ethanol upgrading technologies to expand the collaboration opportunities with industry
  - US Patent 10,696,606, and additional three US patent applications
- Disseminate scientific learnings on catalysis for bioenergy through peer-review publications
  - Hannon et al. PNAS. 2020, 117 (23), 12576-12583; Yuk et al. J. Phys. Chem. C 2020, 124, 28437.
  - Sun et al. Nat Commun 2018, 9, 4454; Cordon et al. ACS Catal. Under revision.

## 4-Progress and Outcomes: One-Step ETO Catalyst Design

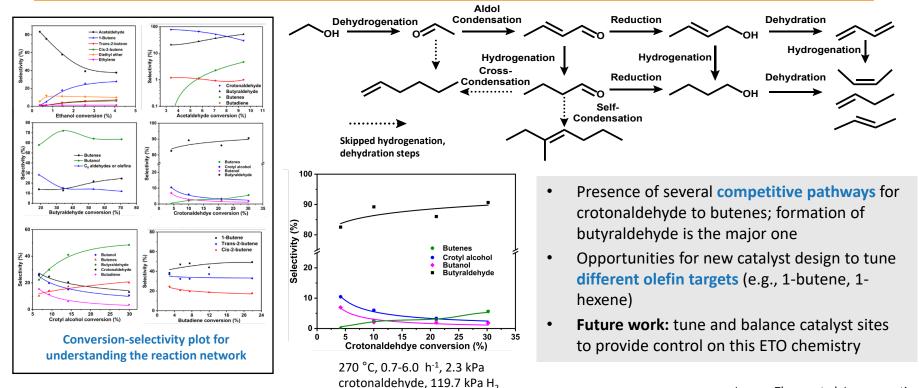
**Objective:** develop multifunctional Lewis acid zeolite-based catalyst(s) to catalyze cascade ethanol conversion to butene-rich  $C_{3+}$  olefins



• 89% C<sub>3+</sub> olefin, 98% total olefins at ~100% conversion (350 °C)

## 4-Progress and Outcomes: Reaction Network Understanding

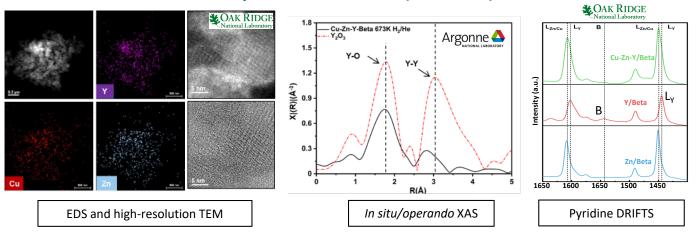
## **Objective:** understand ETO reaction network over Cu-Zn-Y/Beta catalyst by feeding ethanol and various reaction intermediates



Junyan Zhang, et al. In preparation.

## 4-Progress and Outcomes: Catalyst Structures

**Objective:** accelerate understanding of catalyst structures via collaborations with ACSC and CCPC, providing guidance for next-generation catalyst design



#### **Catalyst characterizations (with ACSC)**

#### **Outcomes:**

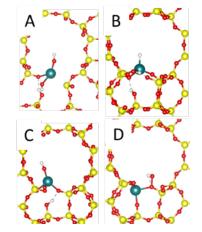
- Characterizations showed atomically dispersed metal sites over Cu-Zn-Y/Beta
  - Isolated Y induced both L acid sites and B acid sites
- Joint efforts with CCPC and ACSC provided structural information for further catalyst design
  - Eliminate B acid sites to inhibit ethanol dehydration; alternative L acid centers for condensation

Future work: further establish structure-function relation for new catalyst design

#### **ChemCatBio**

#### Bioenergy Technologies Office

#### Modeling (with CCPC)



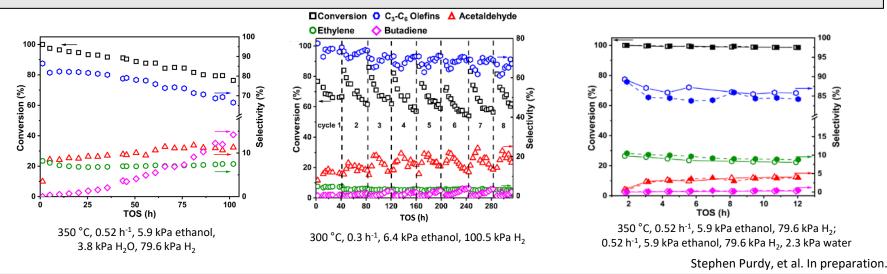
Potential Y/Beta structures identified via modeling (PNNL)

- Catalyst structures
- Reaction mechanism
- Guide catalyst design

Junyan Zhang, et al. In preparation.

## 4-Progress and Outcomes: Catalyst Stability

**Objective:** evaluate Cu-Zn-Y/Beta catalyst stability, regenerability and water tolerance



#### **Outcomes:**

- Cu-Zn-Y/Beta was operated for >100 h TOS in one cycle, regenerable for multiple cycles via oxidative regeneration
- Water does not show significant impact on ethanol conversion and product selectivity for short TOS (Preliminary)
- Offer potential for feeding wet ethanol to cut cost by reducing separation burden

#### **Future work:**

• **Collaboration with CDM, ACSC, CCPC** to further understand catalyst deactivation (i.e., steam stability, coke formation, metal sites stability) in FY21 Q2 and Q3, providing insights for long-term catalyst stability

## 4-Progress and Outcomes: Performance at Varied H<sub>2</sub> Partial Pressures

**Objective:** understand Cu-Zn-Y/Beta performance at different H<sub>2</sub> partial pressures or without H<sub>2</sub>

H <sub>2</sub> Partial Pressure	Conversion	Product Selectivity (%)								
(kPa)	(%)	AA	C <sub>2</sub> =	C <sub>3</sub> =	C <sub>4</sub> =	C <sub>5+</sub> =	C <sub>3+</sub> =	Butadiene	Oxygenates	Paraffins
101	96.4	6.3	17.0	5.0	56.7	10.8	72.6	0.3	9.4	0.6
75	95.8	6.6	17.0	5.0	56.9	11.1	72.9	0.5	9.0	0.5
50	94.5	6.9	16.9	4.9	57.2	8.5	70.6	1.7	10.5	0.3
25	93.0	6.5	16.0	5.0	52.9	7.3	65.2	8.3	10.3	0.2
0	90.6	6.5	16.9	5.1	12.9	6.8	24.8	47.5	10.7	0.1
350 °C, 0.52 h <sup>-1</sup> , 7.4 kPa ethanol AA: acetaldehyde; oxygenates include diethyl ether, AA, C4 oxygenates Junyan Zhang, et al.				nang, et al. In p						

AA: acetaldehyde; oxygenates include diethyl ether, AA, C4 oxygenates

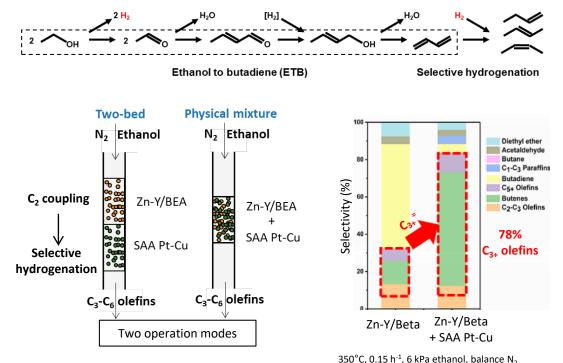
Junyan Zhang, et al. In preparation.

#### **Outcomes:**

- $H_2$  partial pressure (<50 kPa) can be adjusted to tune butadiene/ $C_{3+}$  olefins -
- At low H<sub>2</sub> partial pressure or without H<sub>2</sub>, this process can produce middle distillate and butadiene simultaneously -
- Preliminary TEA indicates coproduction of butadiene can significantly reduce MFSP -
- How to achieve selective ETO without hydrogen cofeeding? (industry interests in this approach)

## 4-Progress and Outcomes: Ethanol to Butenes without External H<sub>2</sub>

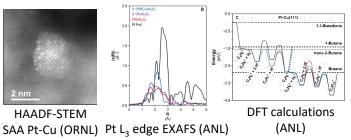
**Objective:** utilize *in situ* generated H<sub>2</sub> to achieve selective ethanol to butene-rich olefins **Challenge:** require active and selective catalyst that can utilize these H<sub>2</sub> at ETO conditions (>300 °C)



Michael Cordon, et al. ACS Catalysis. Under revision.

#### (~94% ethanol conversion)

#### **ACSC and CCPC**



**Outcome:** composite catalyst (Zn-Y/Beta + SAA Pt-Cu) can achieve selective ETO without  $H_2$  cofeeding (**78% C<sub>3+</sub> olefin** selectivity, 94% conversion)

 Characterizations and DFT calculations demonstrate SAA Pt-Cu ensemble sites are critical for BD hydrogenation at ETO conditions

**Impact:** avoid external H<sub>2</sub> supply, separation and recycle; reduce OpEX and CapEx

#### Joint catalysis R&D significantly advanced the state of technology for ETJ via single-step ETO over ORNL catalysts

	FY17	FY18	FY20	Target
ETO Catalyst	Pillared H-MFI	Cu-Zn-Y/Beta	Cu-Zn-Y/Beta	Improved design
ETO Single-pass EtOH Conversion	100%	~99%	100%	100%
C <sub>3+</sub> Olefin Selectivity	33%	87%	89%	95%
Total Olefin Selectivity	65%	92%	98%	98%
Liquid Hydrocarbon Yield (GGE/gal. EtOH)	0.27	0.54	0.58	0.62
Demonstrated Time on Stream		~40 h	>300 h	>300 h*
EtOH Feed	Pure EtOH	Pure EtOH	Aqueous EtOH (up to 60 wt.% water)	Biomass/CO <sub>2</sub> derived EtOH
EtOH Upgrading Cost (\$/GGE)	1.35	0.94	0.60	0.50
Technology Readiness Level	1	1	3	

\*Long-term durability testing may be achieved via other opportunities

#### Future work on catalysis R&D:

- Improve C<sub>3+</sub> olefin selectivity via new catalyst design, leading to increased liquid hydrocarbon yield
- Understand and improve catalyst stability
- Develop **coproduct(s)** from ethanol based on TEA guidance

## Summary

#### **Project Goal**

Develop *efficient one-step ETO* technology to enable *high carbon efficiency to liquid hydrocarbon fuels at lower cost*, and enable the development of a market-responsive biorefinery concept through C<sub>2</sub> platform providing control over gasoline, diesel, and jet fuel, as well as high value co-products

#### Approach

- Developing *multifunctional Lewis acid catalysts* to catalyze cascade reactions to selectively produce C<sub>3+</sub> olefins
- Integrated foundational and applied research to accelerate the catalyst development

#### Impact

- New ETJ technology addresses needs for both ethanol producers and growing markets of middle distillates
- Transitioning R&D discoveries from national labs to industry

#### **Research progress**

- Demonstrate a new Cu-Zn-Y/Beta catalyst for *selective* ETO (~89% C<sub>3+</sub> olefins, 98% total olefins)
- Cu-Zn-Y/Beta is *regenerable* for multiple cycles via oxidative regeneration, and *stable in the presence of water*
- Collaboration among ACSC, CCPC and C<sub>2</sub> upgrading enables the understanding of catalyst structure and chemistry, providing guidance for further catalyst development
- R&D efforts have significantly *advanced the state of the technology*

## **Quad Chart Overview**

## Timeline

- 10/01/2019
- 09/30/2022

	FY20	Total Planned Funding (FY20-Project End Date)
DOE Funding	\$400K	\$1.2M

## Barriers addressed

#### **Ct-F: Increasing the Yield from Catalytic Processes**

• Developing catalysts that enable processes with higher carbon efficiency and yield

#### **Ct-E.** Improving Catalyst Lifetime

• Exploring deactivation via characterization, and developing regeneration/mitigation strategies

## Project Goal

Develop *efficient one-step ETO technology* to enable high carbon efficiency to liquid hydrocarbon fuels at lower cost, and enable the development of a *marketresponsive biorefinery concept* through C2 platform providing control over gasoline, diesel, and jet fuel, as well as high value co-products

## End of Project Milestone

Demonstrate *improvements to the direct ethanol to butene-rich olefins catalyst technology* thereby enabling a new, market-responsive biorefinery pathway through C2+ oxygenates providing control over gasoline, diesel, jet, and co-products, with the ability to *obtain a distillate MFSP of at least \$3/GGE*.

#### Funding Mechanism AOP Project (Direct-Funded Lab Project)

## Acknowledgement

#### **ORNL team**

Zhenglong Li Michael Cordon Kinga Unocic Shiba Adhikari Bruce Adkins Junyan Zhang Stephen Purdy Brian Davison James Parks Todd Toops

Susan Habas

Qiyuan Wu

#### **NREL team**

Ling Tao Dan Ruddy

#### **PNNL team**

Roger Rousseau Vanda Glezakou Robert Dagle Huamin Wang Gregory Collinge Mal-Soon Lee Vanessa Dagle Fan Lin

#### ANL team

Ted Krause Rajeev Assary Evan Wegener Mingxia Zhou

#### **Collaborators**

Jeffrey Miller (Purdue University) James Harris (UA) Dongxia Liu (UMD)



**Bioenergy Technologies Office** 



CCCPC Consortium for Computational Physics and Chemistry

> U.S. DEPARTMENT OF ENERGY BIOENERGY TECHNOLOGIES OFFICE



# DOE Bioenergy Technologies Office (BETO) 2021 Project Peer Review **Upgrading of C2 Intermediates – ORNL**

Catalytic Upgrading

March 10, 2021

**Zhenglong Li** Oak Ridge National Laboratory



## **Additional Slides**



## Responses to Previous Reviewers' Comments

- "In general, the project has very focused tasks to develop a wide range of potential full candidates from low molecular weight oxygenates. The team successfully leverages interactions with many groups within ChemCatBio to enhance the project. The majority of the project has seen activity and selectivity metrics met, though room to improve catalyst reusability remains. In general, future studies appear to build on prior successes and will leverage capabilities in ChemCatBio to attempt to further catalyst performance. The project is productive in both connecting with fundamental science, publications in field-leading journals, technology, and several patents and a successful technology transfer with one of the processes."
  - We agree that catalyst reusability remains an important part of our research. This is the focus of our endof-year goal, where we will explore deactivation and regeneration in all three of our oxygenate conversion pathways.

#### **Publications**

- 1. S.F. Yuk, M. Lee, G. Collinge, J. Zhang, A. B. Padmaperuma, Z. Li\*, F. Polo-Garzon, Z. Wu, V. Glezakou, R. Rousseau\*. Mechanistic Understanding of Catalytic Conversion of Ethanol to 1-Butene over 2D-Pillared MFI Zeolite. Journal of Physical Chemistry C, 2020, 124, 28437.
- 2. J.R. Hannon, L.R. Lynd\*, O. Andrade, P.T. Benavides, et al. Technoeconomic and life cycle analysis of single-step catalytic conversion of wet ethanol into fungible fuel blendstocks. PNAS, 2020, 117, 12576-12583.
- 3. M.J. Cordon, J. Zhang, S.C. Purdy, E.C. Wegener, K.A. Unocic, L.F. Allard, M. Zhou, R.S. Assary, J.T. Miller, T.R. Krause, J. Kropf, C. Yang, D. Liu, Z. Li<sup>\*</sup>. Selective butene formation in direct ethanol to C3+ olefin valorization over Zn-Y/Beta and single-atom alloy composite catalysts using in situ generated hydrogen. *ACS Catal*. Under revision.
- 4. J. Zhang, L. Tao\*, B.H. Davison, Z. Li\*. Decarbonize Aviation via Cost-Competitive Middle Distillate Fuels within a Market-Flexible C2 Platform-based Biorefinery Concept. In prep.
- 5. J. Zhang, E.C. Wegener, S.C. Purdy, K.A. Unocic, L.F. Allard, M. Zhou, R.S. Assary, J.T. Miller, T.R. Krause, J. Kropf, C. Yang, D. Liu, Z. Li\*. Isolated metal sites over Cu-Zn-Y/Beta for selective butene-rich C3+ olefins production from ethanol. In prep.

#### **Presentations**

- 1. R. Dagle, V. Dagle, Z. Li. "Single-step catalytic conversion of ethanol to n-butene-rich olefins and 1,3-butadiene chemical product", DOE BETO ChemCatBio Webinar, July 31, 2019.
- J. Zhang, S. Adhikari, C. Yang, N.J. Libretto, J. Miller, T. Krause, K. Unocic, D. Liu, Z. Li. "Multifunctional Zeolite Catalyst for Cascade Conversion of Ethanol to C3+ Olefins: A Pathway to Renewable Jet Fuel", NAM26 2019 North American Catalysis Society Meeting.
- 3. Z. Li, J. Zhang, S. Adhikari, C. Yang, J. Miller, T. Krause, K. Unocic. "Multifunctional Zeolite Catalyst for Tandem Ethanol Conversion to C3-C6 Olefins: A New Pathway to Renewable Jet", ACS Spring 2019 National Meeting.
- 4. Z. Li, J. Zhang, S. Adhikari, C. Yang, E.C. Wegener, J. Miller, T. Krause, K. Unocic, L. Allard, D. Liu, B. Davison, Z. Wu. "Multifunctional zeolite catalyst for cascade ethanol conversion to C3-C6 Olefins: a pathway to renewable aviation biofuel". ACS Fall 2019 National Meeting.

#### Patents

- 1. Narula, C. K., B.H. Davison, Z. Li, "Zeolitic catalytic conversion of alcohols to hydrocarbon fractions with reduced gaseous hydrocarbon content," United States Patent 10,696,606, issued June 30, 2020.
- 2. Zhenglong Li, "Zeolitic catalytic conversion of alcohols to olefins" United States Patent Application No. 16/530,397 filed on August 2, 2019.
- 3. Zhenglong Li, "Method for alcohol upgrading to jet, diesel, gasoline and valuable co-products." United States Patent Application No. 17/083,437 filed on October 29, 2020.
- 4. Zhenglong Li, Junyan Zhang, Brian Davison. "New catalyst materials for ethanol conversion to high olefins." United States Provisional Application Serial No. 63/141,996.

#### Technology Transfer

1. ORNL licensed new ethanol-to-butadiene-jet (ETBJ) technology to Prometheus Fuels: https://www.ornl.gov/news/prometheus-fuels-licenses-energy-saving-ornl-ethanol-jet-fuel-process