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ALD-Modified USY Zeolite Characterization Using Single-Event MicroKinetics

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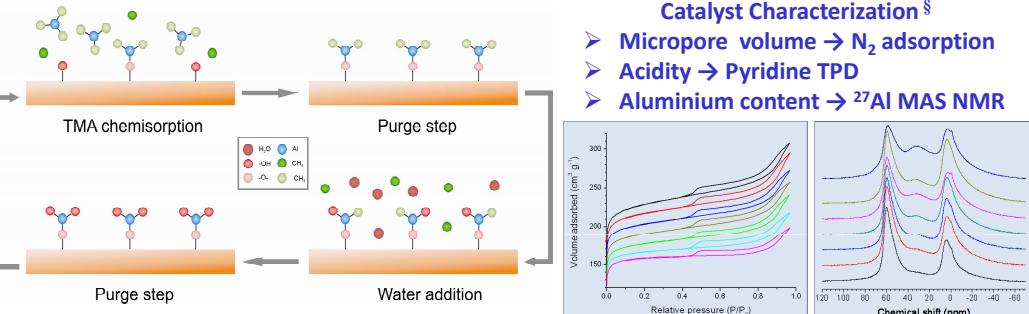
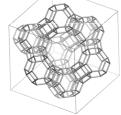
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CATALYST MODIFICATION

Modification of a commercial Pt/H-USY zeolite by Atomic Layer Deposition (ALD) making use of the $\text{Al}(\text{CH}_3)_3/\text{H}_2\text{O}$ process

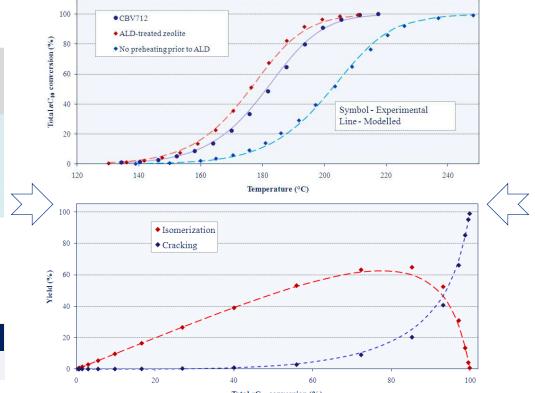
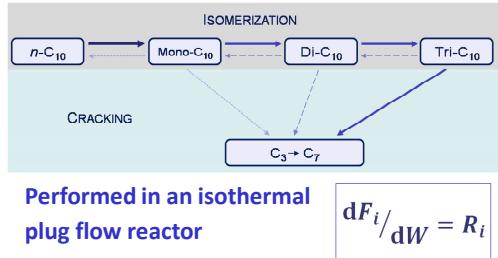
1. $\text{||-OH} + \text{Al}(\text{CH}_3)_3(\text{g}) \rightarrow \text{||-O-Al}(\text{CH}_3)_2 + \text{CH}_4(\text{g})$
2. $2\text{||-OH} + \text{Al}(\text{CH}_3)_3(\text{g}) \rightarrow (\text{||-O})_2\text{AlCH}_3 + 2\text{CH}_4(\text{g})$
3. $\text{||-CH}_3 + \text{H}_2\text{O}(\text{g}) \rightarrow \text{||-OH} + \text{CH}_4(\text{g})$

Parent material: CBV712
 $\text{Si}/\text{Al} = 5.8$



SINGLE-EVENT MICROKINETIC (SEMK) MODELING

Hydrocracking experiments using *n*-decane



$$r_{\text{iso/cra}}(m_1; m_2) = \frac{k_{\text{iso/cra}}(m_1; m_2) C_{\text{sat}} C_{\text{acid}} K_{\text{prot}} K_{\text{deh}} K_{\text{L}} p_{\text{pp}} p_{\text{H}_2}^{-1}}{(1 + \sum K_{\text{L}} p_{\text{pp}}) \left(1 + \frac{\sum C_{\text{sat}} K_{\text{prot}} K_{\text{deh}} K_{\text{L}} p_{\text{pp}} p_{\text{H}_2}^{-1}}{1 + \sum K_{\text{L}} p_{\text{pp}}} \right)}$$

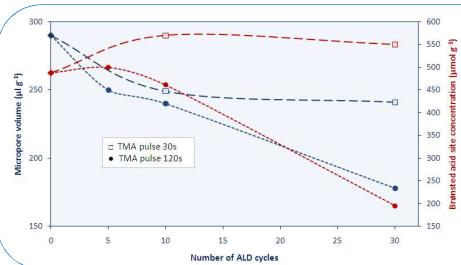
$$k_{\text{iso/cra}}(m_1; m_2) = n_e \bar{k}_{\text{iso/cra}}(m_1; m_2)^\dagger$$

\tilde{k} - unique rate coefficient of reaction family
 n_e - number of geometrically independent ways in which the transition state can be formed → 'number of single events'
 m_1, m_2 - type of reactant and product carbenium ion

Protonation enthalpy for ion formation estimated; $\Delta H_p(t) \approx \Delta H_p(s) - 30 \text{ kJ mol}^{-1}$

EFFECT OF ALD ON CATALYST PROPERTIES

Catalyst dried at 473 K for 6 h prior to ALD
TMA/ H_2O pulse and purge times 30 or 120 s
Total number of ALD cycles 5, 10 or 30
ALD reaction temperature 473 K



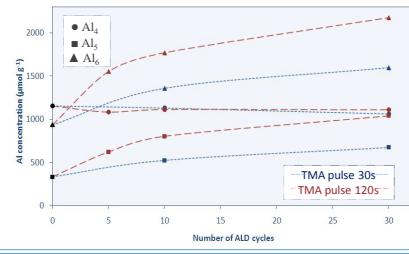
TMA deposition in micropores ⇒ MV ↗
Creation of new acid sites, covered by extra-framework Al
 $\Rightarrow C_{\text{acid}}$

| TMA pulse (s) | # ALD cycli | $-\Delta H_p(s)$ kJ mol^{-1} | $-\Delta H_p(t)$ kJ mol^{-1} |
|---------------|-------------|--|--|
| - | - | $70.8 (\pm 0.1)^*$ | $101.6 (\pm 0.2)$ |
| 30 | 10 | $72.8 (\pm 0.3)$ | $102.4 (\pm 0.5)$ |
| 30 | 30 | $72.0 (\pm 0.2)$ | $100.4 (\pm 0.4)$ |
| 120 | 5 | $72.8 (\pm 0.3)$ | $101.1 (\pm 0.5)$ |
| 120 | 10 | $72.9 (\pm 0.2)$ | $101.2 (\pm 0.5)$ |
| 120 | 30 | $78.5 (\pm 0.3)$ | $110.6 (\pm 0.3)$ |

* 95% confidence region

Formation of new and possibly stronger sites
Inductive effect of extra-framework $\text{Al}_2\text{O}_3(s)$ ⇒ $-\Delta H_p$ ↗
Improvement of hydrocracking activity explained through an increase in average acid site strength

Framework remains unharmed ⇒ $\text{Al}_4 \approx$
Reaction TMA and surface $-\text{OH}$, formation of $\text{Al}_2\text{O}_3(s)$ through chemical vapour deposition
 $\Rightarrow \text{Al}_5 \nearrow$ and $\text{Al}_6 \nearrow$



ALD reaction temperature 573 K ⇒ steaming of zeolite ⇒ $\text{Al}_4 \searrow$
No pretreatment catalyst ⇒ formation of weaker sites ⇒ $-\Delta H_p \nearrow$
High purge times ⇒ longer reaction times $\text{H}_2\text{O} \Rightarrow -\Delta H_p \searrow, C_{\text{acid}} \nearrow$

CONCLUSIONS

- The single-event methodology has proven to be a useful tool in the assessment of catalytic modifications
- Each ALD parameter has a specific effect on the hydrocracking behavior of the catalyst through changes in micropore volume, Brønsted acid site concentration and average acid site strength
- The creation of new acid sites through ALD opens up the route towards the production of new active materials tailored to the requirements of a target reaction

ACKNOWLEDGEMENTS

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