Resolution Test Chart

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PROBE MOLECULE STUDIES:
ACTIVE SPECIES IN ALCOHOL SYNTHESIS

11th Quarterly Report
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TABLE OF CONTENT

1. OBJECTIVE AND SCOPE OF WORK	1
2. SUMMARY OF PROGRESS	1
3. DETAILED DESCRIPTION OF TECHNICAL PROGRESS	2
4. PLANS FOR NEXT REPORTING PERIOD	4
5. REFERENCES	5
 Table 1.	 6
Table 2.	7
Table 3.	8
 Figure 1(a)	 9
Figure 1(b)	10
Figure 2(a)	11
Figure 2(b)	12

1. OBJECTIVE AND SCOPE OF WORK

The goal of this research is to develop a better understanding of the mechanisms of formation of alcohols and other oxygenates from syngas over supported catalysts. Probe molecules are added in situ during the reaction to help delineate reaction pathways and identify reaction intermediate species. The key of our study is to investigate how the species generated by these probe molecules interact with surface species present during oxygenate formation.

The catalysts chosen for this investigation is Co/Cu/ZnO/Al₂O₃. Detailed motivations for studying this system as well as using CH₃NO₂ as the probe molecule were given in a previous report^[1].

2. SUMMARY OF PROGRESS

(A) CO hydrogenation reactions were carried out over a series of Cu/Co/Cr catalysts prepared by coprecipitation method by Murty's group^[2]. The results were compared with the data obtained with ZC45 catalyst^[3] as well as the Co(0, 5, 10%)/Cu/ZnO/Al₂O₃ catalysts^[4].

(B) High pressure reduction found to enhance both activity and selectivity of the Co-Cu catalysts for C₁-C₆ alcohol synthesis was used to retest some of the catalysts which had already been studied.

(C) Experiments of nitromethane addition to the CO hydrogenation over ZC45 with both the conventional and the new reduction treatment and over the Co(10%)/Cu/ZnO/Al₂O₃ catalyst reduced at high pressure have been carried out to study the reaction mechanism.

3. DETAILED DESCRIPTION OF TECHNICAL PROGRESS

(A) CO hydrogenation over the Cu/Co/Cr catalysts.

Four catalyst samples from Murty's group have been tested with CO hydrogenation reaction at 500 psig, 290°C, GHSV $\approx$ 7000 hr⁻¹, H₂/CO=2 for 2~4 hours. The composition of the catalysts are listed in Table 1^[2]. The steady state data obtained over these four catalysts around 3 hours after the reactions were started are given in Table 2 and compared with the data obtained with ZC45 and the Co(0, 5, 10%)/Cu/ZnO/Al₂O₃ catalysts also 2~4 hours after the reactions were started. It is shown in Table 2 that:

- (a) #1, #3 and #4 of the Co/Cu/Cr samples had much lower activities and higher selectivities towards hydrocarbons than the Co(5, 10%)/Cu/ZnO/Al₂O₃ samples;
- (b) only sample #2 of the Co/Cu/Cr catalysts had similar activity and selectivity to those of the Co(5, 10%)/Cu/ZnO/Al₂O₃ catalysts;
- (c) none of the catalysts prepared by coprecipitation, including ZC45, demonstrated significantly higher activities for C₂⁺ alcohols than those obtained with the catalysts prepared by impregnation of CuO/ZnO/Al₂O₃ with cobalt nitrate (Co(5,10%)/Cu/ZnO/Al₂O₃ catalysts), at least when they were reduced at atmospheric pressure;
- (d) all the catalysts containing Co had much lower activities for methanol formation;
- (e) the Co containing catalysts, except for #1, #3 and #4 of Co/Cu/Cr, had higher activities of C₃⁺ alcohols than the Co(0%)/Cu/ZnO/Al₂O₃ catalyst.

(B) Enhancement of both activity and selectivity of the Co-Cu catalysts for C₁-C₆ alcohol synthesis using a new reduction procedure.

It has been found that reducing the catalyst at the reaction pressure, 500

psig, rather than at atmospheric pressure enhanced both the activity and selectivity of ZC45 and Co(10%)/Cu/ZnO/Al₂O₃ catalysts for C₁-C₆ alcohol formation. The steady state data of CO hydrogenation over these two catalysts which received the new reduction treatment are listed in Table 3 and compared with the data over the same samples reduced at atmospheric pressure. It is shown that:

- (a) the activities and selectivities for C₁-C₆ alcohol formation over both ZC45 and Co(10%)/Cu/ZnO/Al₂O₃ catalysts were enhanced by high pressure reduction;
- (b) on ZC45 the selectivity was shifted to higher alcohols;
- (c) the activities for hydrocarbons over ZC45 were enhanced by high pressure reduction, while they were hardly affected on Co(10%)/Cu/ZnO/Al₂O₃;

(C) CH₃NO₂ addition to CO hydrogenation over catalyst samples with the two different reduction treatments.

CH₃NO₂ was added as probe molecule to the CO hydrogenation over the catalyst samples of ZC45 reduced at both atmospheric and high pressure and over the Co(10%)/Cu/ZnO/Al₂O₃ catalyst reduced at high pressure. Figure 1 and 2 show the reaction results obtained with the samples of ZC45 and Co(10%)/Cu/ZnO/Al₂O₃, respectively, reduced at (a) low pressure, and (b) high pressure. The data for the Co(10%)/Cu/ZnO/Al₂O₃ catalyst reduced at low pressure were reported in previous report^[5]. The followings are the observations from the experimental results.

- (a) With all these four catalyst samples:
 - methanol was suppressed and trimethylamine was detected during the addition of nitromethane;
 - the ratios of the formation rates with CH₃NO₂ addition to those without CH₃NO₂ addition, R, for C₂-C₆ alcohols increased with carbon number with the R values for C₄-C₆ alcohols being all greater than 1;

- the effects of nitromethane addition on the formation rates of hydrocarbons, or the R values of hydrocarbons, did not follow the same trend as those of alcohols.
- (b) With the samples of ZC45 reduced at both low and high pressures, nitromethane addition suppressed the formation of C₁-C₆ hydrocarbons, while it enhanced the formation rates of C₁-C₆ hydrocarbons over Co(10%)/Cu/ZnO/Al₂O₃.

4. PLANS FOR THE NEXT REPORTING PERIOD

The following studies are planned for the next reporting period:

- a) Nitromethane addition to the CO hydrogenation over ZC45 samples reduced at high and low pressures with detection of N₂ to assist in the understanding of the reaction pathways.
- b) Nitromethane addition to the CO hydrogenation over Co(0%)/Cu/ZnO/Al₂O₃ and Co(5%)/Al₂O₃ reduced differently to assist in the understanding of both the reaction pathway(s) and the active site(s).
- c) XRD experiments with the catalyst samples which received different reduction treatments to assist in the understanding of the active sites as well as the effect of reduction pressure.
- d) Experiments of temperature programmed reduction at high pressure to generate information on the effect of the reduction pressure.

5. REFERENCES

- [1] DOE Quarterly Report, No. 1&3&5, 1991, under Grant No. DE-FG22-90PC-90305.
- [2] U. A. Donatto, Tess L. Hoard, and A. N. Murty, 1992, (unpublished).
- [3] DOE Quarterly Report, No. 10, 1993, under Grant No. DE-FG22-90PC-90305.
- [4] DOE Quarterly Report, No. 7&8&9, 1992, under Grant No. DE-FG22-90PC-90305.
- [5] DOE Quarterly Report, No. 9, 1992, under Grant No. DE-FG22-90PC-90305.
- [6] DOE Quarterly Report, No. 5&6&7, 1992, under Grant No. DE-FG22-90PC-90305.
- [7] F. A. P. Cavalcanti, R. Oukaci, I. Wneder, D. G. Blackmond, j. Catal. 123, 270-274 (1990).

Table 1. Cu/Co/Cr Catalyst Composition (from Ref. 2).

Sample (#)	Cu/Co (atomic ratio)	Cu/Co/Cr
1	1.3	40/30/30
2	1.7	50/30/20
3	3.3	50/15/35
4	4.6	70/15/15

Table 2. Formation Rates^a of the Products of CO Hydrogenation at 500 psig; 290°C; H₂:CO = 2.

CATALYSTS:	Co/Cu/Cr				Co/Cu/Zn/Al			
	#1	#2	#3	#4	ZC45	10%-Co	5%-Co	0%-Co
ACTIVITY, $\mu\text{mol-C/g-cat/min}$								
OXYs + HCs:	12.9	53.7	3.0	7.6	25.1	57.3	58.7	320.1
OXYGENATES:	2.0	17.5	0.2	0.1	9.6	28.1	25.5	290.4
CH ₃ OH	1.4	7.6	0.2	0.1	2.3	15.2	16.0	248.6
CH ₃ OCH ₃	0.0	0.0	0.0	0.0	0.0	0.4	0.4	18.6
CH ₃ CH ₂ OH	0.4	7.0	0.0	0.0	3.3	7.9	6.4	23.2
CH ₃ CH ₂ CH ₂ OH	0.2	2.9	0.0	0.0	2.0	3.7	2.2	0.0
CH ₃ CH ₂ CH ₂ CH ₂ OH	0.0	0.0	0.0	0.0	2.0	0.9	0.5	0.0
HYDROCARBONS:	10.9	36.2	2.8	7.5	15.5	29.2	33.2	29.7
CH ₄	6.4	9.8	32.2	6.7	6.0	15.7	12.7	5.5
C ₂ H ₄ +C ₂ H ₆	1.3	4.4	0.2	0.4	2.7	6.4	8.0	1.8
C ₃ + HCs	3.2	22.0	0.4	0.4	6.8	7.1	12.5	22.4
CO ₂ :	10.0	34.2	7.4	6.0	28.8	27.7	31.5	18.5
SELECTIVITY ^b , mol-C %								
OXYGENATES:	15.5	32.6	6.7	1.3	38.2	49.0	43.5	90.7
CH ₃ OH	10.9	14.2	6.7	1.3	9.1	26.5	27.3	77.7
CH ₃ OCH ₃	0.0	0.0	0.0	0.0	0.0	0.7	0.7	5.8
CH ₃ CH ₂ OH	3.1	13.0	0.0	0.0	13.1	13.8	10.9	7.2
CH ₃ CH ₂ CH ₂ OH	1.5	5.4	0.0	0.0	8.0	6.4	3.7	0.0
CH ₃ CH ₂ CH ₂ CH ₂ OH	0.0	0.0	0.0	0.0	8.0	1.6	0.9	0.0
HYDROCARBONS:	84.5	67.4	93.3	98.7	61.8	51.0	56.5	9.3
CH ₄	49.6	18.2	73.3	88.2	23.9	27.4	21.6	1.7
C ₂ H ₄ +C ₂ H ₆	10.1	8.2	6.7	5.3	10.8	11.2	13.6	0.6
C ₃ + HCs	24.8	41.0	13.3	5.3	27.1	12.4	21.3	7.0
CO CONVERSION, %								
	0.6	2.2	0.3	0.3	1.4	2.2	2.2	1.3

a: 2~3 hours after the CO hydrogenation reaction was started.

b: CO₂ is not included.

Table 3. Effect of Reduction Pressure on the Activity and Selectivity of CO Hydrogenation^a over ZC45 and Co(10%)/Cu/ZnO/Al₂O₃ at 500 psig; 290°C; H₂:CO = 2.

	ACTIVITY				SELECTIVITY ^b			
	$\mu\text{mol-C/g-cat/min}$				%, mol-C			
	ZC45		10%-Co		ZC45		10%-Co	
Catalyst:	LPR	HPR	LPR	HPR	LPR	HPR	LPR	HPR
Reduction Condition ^c :	LPR	HPR	LPR	HPR	LPR	HPR	LPR	HPR
OXYs + HCs :	29.0	97.5	45.9	58.7				
OXYGENATES:	14.5	60.7	29.6	42.0	50.0	62.2	64.5	71.6
Methanol	4.6	7.9	22.3	28.9	15.9	8.1	48.5	49.2
Dimethylether	0.0	0.0	0.3	0.1	0.0	0.0	0.7	0.2
Ethanol	5.2	12.8	4.3	5.9	17.9	13.1	9.4	10.1
n-Propanol	2.7	7.2	2.3	4.7	9.3	7.4	5.0	8.0
n-Butanol	2.0	9.6	0.4	1.6	6.9	9.8	0.9	2.7
n-Pentanol	0.0	10.0	0.0	0.8	0.0	10.3	0.0	1.4
n-Hexanol	0.0	13.2	0.0	0.0	0.0	13.5	0.0	0.0
HYDROCARBONS:	14.5	36.8	16.3	16.7	50.0	37.8	35.5	28.4
Methane	4.5	11.4	9.6	9.5	15.5	11.7	20.9	16.2
C ₂	2.6	7.2	4.2	4.0	9.0	7.4	9.2	6.8
C ₃ +	7.4	18.2	2.5	3.2	25.5	18.7	5.4	5.4
CO ₂ :	31.9	128.	20.1	64.0				
CO CONVERSION, %								
	1.6	6.5	1.7	2.6				

a: 24 hours after the reaction was started.

b: CO₂ is not included.

c: LPR represents low-pressure-reduction, and HPR represents high-pressure-reduction.

Figure 1(a). Effect of CH_3NO_2 addition on the products of the CO hydrogenation over the sample of ZC45 reduced at low pressure. R is the ratio of the rate with CH_3NO_2 addition to the rate without CH_3NO_2 addition.

0708ALC
ZC45HPR

0708HCS
ZC45HPR

Figure 1(b). Effect of CH₃NO₂ addition on the products of the CO hydrogenation over the sample of ZC45 reduced at high pressure. R is the ratio of the rate with CH₃NO₂ addition to the rate without CH₃NO₂ addition.

Figure 2(a). Effect of CH₃NO₂ addition on the products of the CO hydrogenation over the sample of Co(10%)/Cu/ZnO/Al₂O₃ reduced at low pressure. R is the ratio of the rate with CH₃NO₂ addition to the rate without CH₃NO₂ addition.

0720ALC
CO10%HPR

0720HCS
CO10%HPR

Figure 2(b). Effect of CH₃NO₂ addition on the products of the CO hydrogenation over the sample of Co(10%)/Cu/ZnO/Al₂O₃ reduced at high pressure. R is the ratio of the rate with CH₃NO₂ addition to the rate without CH₃NO₂ addition.

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