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## APPENDICES

## APPENDIX A

### SAMPLES OF CALCULATION

#### Preparation of 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> Catalysts with the Dry Impregnation Method

Reagent: Chloroplatinic acid (H<sub>2</sub>PtCl<sub>6</sub>·6H<sub>2</sub>O)

Molecular weight = 517.92 g

(Atomic weight of Platinum = 195.08)

Support : Alumina (γ-Al<sub>2</sub>O<sub>3</sub>); type KNH-3

Pore volume = 1.0 ml/g

From Sumitomo Aluminium Smelting Co., Ltd., Japan.

#### Calculation for prepared 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> (%by weight)

Based on: 0.3%Pt/Al<sub>2</sub>O<sub>3</sub>                      Catalyst Weight = 100 g

Assume : alumina support used is X grams.

So that, the catalyst 100 grams would composed of

Platinum            0.3            g

Alumina             X             g

Then            0.3 + X = 100    g

Support (X) = 99.7    g

The alumina support weight used for preparation is 2 grams and H<sub>2</sub>PtCl<sub>6</sub>·6H<sub>2</sub>O used as precursor salt.

Platinum required            =    2 × 0.3 / 99.7            g

    =    6.018 × 10<sup>-3</sup>            g

Platinum (Pt) 6.018 × 10<sup>-3</sup> g was prepared from 25 ml of the stock solution of chloroplatinic acid, which prepared by dissolving 1 g of H<sub>2</sub>PtCl<sub>6</sub> in de-ionized water.

$$\begin{aligned} \text{Pt content in stock solution} &= 1 \times 195.08 / 518.1 && \text{g} \\ &= 0.377 && \text{g} \end{aligned}$$

Therefore:

$$\begin{aligned} \text{The required-solution} &= 6.018 \times 10^{-3} \times 25 / 0.377 && \text{ml} \\ &= 0.3993 && \text{ml} \end{aligned}$$

### Preparation of 0.3%Pt/TiO<sub>2</sub> Catalysts with the Wet Impregnation Method

Reagent: Chloroplatinic acid (H<sub>2</sub>PtCl<sub>6</sub> · 6H<sub>2</sub>O)

Molecular weight = 517.92 g.

(Atomic weight of Platinum = 195.08)

Support: Titanium dioxide (TiO<sub>2</sub>); Anatase phase

From Fluka., A Sigma-Aldrich Company, Switzerland.

### Calculation for prepared 0.3%Pt/TiO<sub>2</sub> (%by weight)

Based on: 0.3%Pt/TiO<sub>2</sub> Catalyst Weight = 100 g

Assume : Titanium oxide support used is X grams.

So that, the catalyst 100 grams would composed of

$$\begin{array}{rcl} \text{Platinum} & 0.3 & \text{g} \\ \text{Titanium oxide} & X & \text{g} \\ \text{Then} & 0.3 + X = 100 & \text{g} \\ \text{Support (X)} & = 99.7 & \text{g} \end{array}$$

The titanium oxide support weight used for preparation is 3 grams and H<sub>2</sub>PtCl<sub>6</sub> · 6H<sub>2</sub>O used as precursor salt.

$$\begin{aligned} \text{Platinum required} &= 3 \times 0.3 / 99.7 && \text{g} \\ &= 9.027 \times 10^{-3} && \text{g} \end{aligned}$$

Platinum (Pt)  $9.027 \times 10^{-3}$  g was prepared from 25 ml of the stock solution of chloroplatinic acid which prepared by dissolving 1 g of  $\text{H}_2\text{PtCl}_6$  in de-ionized water.

$$\begin{aligned}\text{Pt content in stock solution} &= 1 \times 195.08 / 518.1 \quad \text{g} \\ &= 0.377 \quad \text{g}\end{aligned}$$

Therefore:

$$\begin{aligned}\text{The required-solution} &= 9.027 \times 10^{-3} \times 25 / 0.377 \quad \text{ml} \\ &= 0.599 \quad \text{ml}\end{aligned}$$

## APPENDIX B

### CALCULATION OF METAL ACTIVE SITE

#### Calculation of metal active site on catalyst

The weight of catalyst used	=	w	g
Area of CO peak after adsorption	=	A	unit
Average area of 50 $\mu$ l standard CO peak	=	B	unit
Amounts of CO adsorbed on catalyst	=	B – A	unit
Volume of CO adsorbed on catalyst	=	$[(B - A)/B] \times 50$	$\mu$ l
Volume of gas 1 mole at 30°C	=	$24.86 \times 10^6$	$\mu$ l
Mole of CO adsorbed on catalyst (mole)	=	$[(B - A)/B] \times [50/24.86 \times 10^6]$	

1 mole is  $6.02 \times 10^{23}$  molecules

Then, Molecule of CO adsorbed on catalyst

$$= 2.01 \times 10^{-6} \times [(B - A)/B] \times 6.02 \times 10^{23} \text{ molecules}$$

Metal active site =  $1.21 \times 10^{18} \times [(B - A)/B]/w$  molecules of CO / g catalyst

#### Example of calculated active site of 0.3 % Pt/Al<sub>2</sub>O<sub>3</sub>

0.3 % Pt/Al <sub>2</sub> O <sub>3</sub>	=	0.1	g
Area of CO peak after adsorption	=	1185	unit
Average area of 50 $\mu$ l. standard CO peak	=	2170	unit
Amounts of CO adsorbed on catalyst	=	2170 – 1185	unit
Volume of CO adsorbed on catalyst	=	$[(2170 - 1185)/2170] \times 50$	$\mu$ l
	=	22.70	$\mu$ l

The adsorption CO condition was carried out at 30°C

Then, Mole of CO adsorbed on catalyst =  $(22.70/24.86 \times 10^6)$



Molecule of CO adsorbed on catalyst (0.1 g)

$$= (22.70/24.86 \times 10^6) \times 6.02 \times 10^{23} \quad \text{molecules}$$

$$= 5.50 \times 10^{17} \quad \text{molecules}$$

Metal active site =  $5.50 \times 10^{17} / 0.1$

$$= 5.50 \times 10^{18} \quad \text{molecules of CO / g catalyst}$$

## APPENDIX C

### CALCULATION OF NO, CO AND C<sub>3</sub>H<sub>8</sub> CONVERSIONS

#### Calculation of NO, CO and C<sub>3</sub>H<sub>8</sub> conversion

The effluent gas was analyzed by gas chromatograph, the NO reduction activity was evaluated in terms of the conversion of NO to N<sub>2</sub>



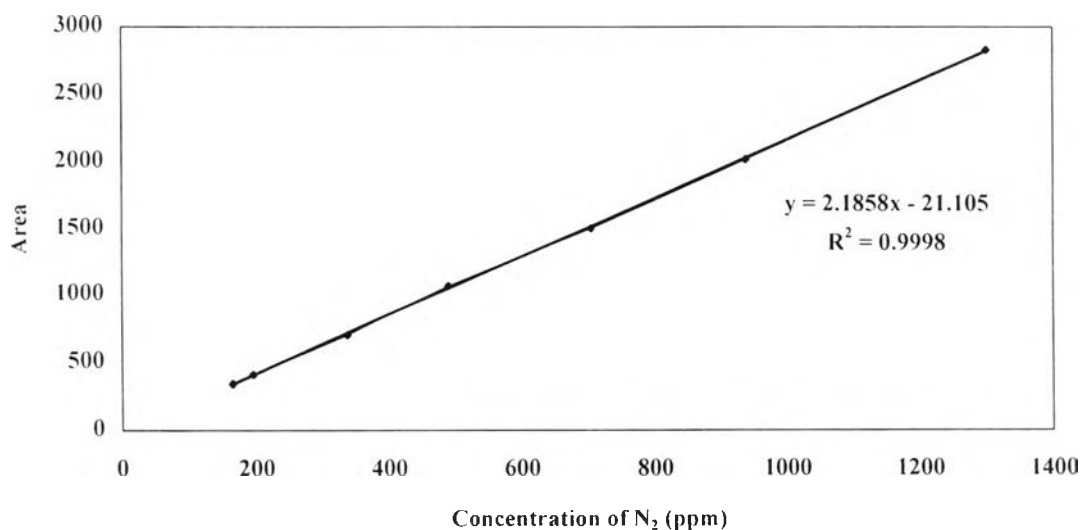
$$\text{NO conversion (\%)} = (2[\text{N}_2]_{\text{out}}/[\text{NO}]_{\text{in}}) \times 100$$

$$\text{Where } [\text{NO}]_{\text{in}} = 500 \text{ ppm}$$

[N<sub>2</sub>] : analyzed by gas chromatograph from calibration curve  
(Figure C-1)

**Figure C-1** Calibration curve of N<sub>2</sub>

Calibrate N<sub>2</sub>, Inj T = 100, Col T = 70, Slope test = 100, Current = 80  
(Molecular sieve 5A)



$$\begin{aligned}\text{Area} &= \text{area of N}_2 \text{ peak on GC 8 ATP} \\ \text{Area} &= 2.1858 \times \text{conc. of N}_2 \text{ (ppm)} - 21.105 \\ \text{Thus, [N}_2\text{]} &= (\text{Area} + 21.105)/2.1858\end{aligned}$$

The CO oxidation activity was evaluated in terms of the conversion of CO into CO<sub>2</sub>

$$\text{CO conversion (\%)} = \frac{([\text{CO}]_{\text{in}} - [\text{CO}]_{\text{out}}) \times 100}{[\text{CO}]_{\text{in}}}$$

The C<sub>3</sub>H<sub>8</sub> oxidation activity was evaluated in terms of the conversion of C<sub>3</sub>H<sub>8</sub> into CO<sub>2</sub> and H<sub>2</sub>O

$$\text{C}_3\text{H}_8 \text{ conversion (\%)} = \frac{([\text{C}_3\text{H}_8]_{\text{in}} - [\text{C}_3\text{H}_8]_{\text{out}}) \times 100}{[\text{C}_3\text{H}_8]_{\text{in}}}$$

**APPENDIX D**  
**CHEMICAL AND PHYSICAL PROPERTIES OF SUPPORTS**

**Table D-1** Specification of Alumina Support (Al<sub>2</sub>O<sub>3</sub>) Type KNH-3

<b>Chemical component</b>	<b>weight percent (%)</b>
Al <sub>2</sub> O <sub>3</sub>	60-70
SiO <sub>2</sub>	30-35
Fe <sub>2</sub> O <sub>3</sub>	0.3-0.5
TiO <sub>2</sub>	0.5-0.7
CaO	0.1-0.2
MgO	0.2-0.4
Na <sub>2</sub> O	0.3-0.4
K <sub>2</sub> O	0.2-0.3
ZrO <sub>2</sub> + HfO <sub>2</sub>	0.03-0.04

<b>Physical properties</b>	
Bulk density (g/ml)	1.3-1.5
Apparent Specific Gravity	3.1-3.3
Packing Density (lb/ft <sup>3</sup> )	20-25
Pore Volume (ml/g)	1.0-1.3
Surface Area (m <sup>2</sup> /g)	340-350

**Table D-2** Chemical component of TiO<sub>2</sub> support

<b>Chemical component</b>	<b>weight percent (%)</b>
TiO <sub>2</sub>	99
PO <sub>4</sub>	0.1
SO <sub>4</sub>	0.1
Cl	0.01
Pb	0.001
As	0.002
Fe	0.005
Zn	0.005
Cu	0.0005

## APPENDIX E

### OPERATING CONDITIONS AND SAMPLES OF CHROMATOGRAM

1. A thermal conductivity detector gas chromatography (model 8ATP) was used to analyze the concentrations of oxygen, nitrogen and carbon monoxide.

Operating conditions are as follows :

GC	:	SHIMADZU-GC-8ATP
Detector	:	TCD
Packed column	:	MS-5A
Carrier gas	:	Ultra high purity helium (99.999%)
Flow rate of carries gas	:	45 ml/min
Column temperature	:	70°C
Detector temperature	:	100°C
Injector temperature	:	100°C
Detector current	:	80 mA

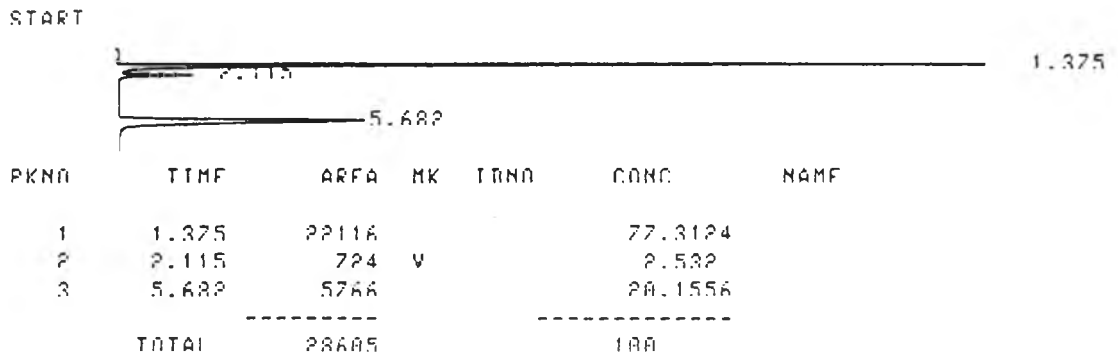
2. Gas chromatography model 8AIT was used to analyze the concentration of H<sub>2</sub>O, propane, carbon dioxide.

Operating conditions were similar to model 8ATP except:

Packed column	:	Porapak-Q
Flow rate of carries gas	:	30 ml/min
Column temperature	:	90°C
Detector temperature	:	110°C
Detector current	:	90 mA

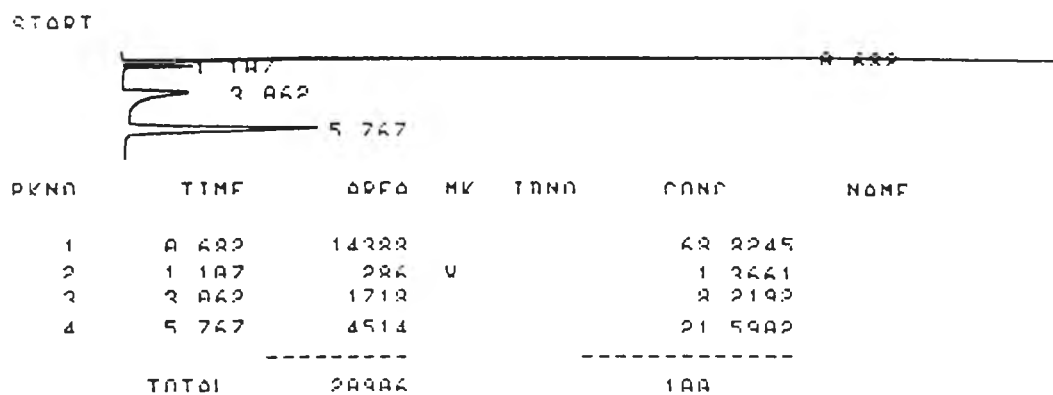
The samples of chromatogram from gas analysis are shown in Figures E-1, E-2.

Figure E-1 Sample of Chromatogram from GC-8ATP (column MS-5A)



<u>PKNO</u>	<u>TIME</u>	<u>AREA</u>	<u>CONC</u>	<u>NAME</u>
1.	1.375	22116	77.3124	OXYGEN
2.	2.115	724	2.5320	NITROGEN
3.	5.682	5766	20.1556	CARBON MONOXIDE
		.....	.....	
TOTAL		28605	100	

Figure E-2 Sample of Chromatogram from GC-8AIT (column Porapak-Q)



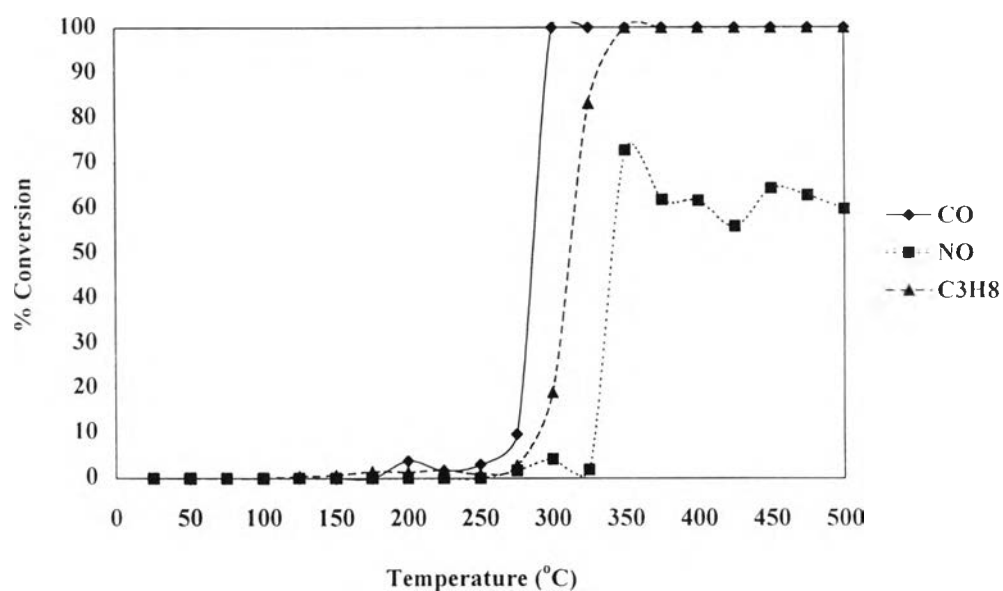
<u>PKNO</u>	<u>TIME</u>	<u>AREA</u>	<u>CONC</u>	<u>NAME</u>
1.	0.682	14388	68.8245	AIR (N <sub>2</sub> +O <sub>2</sub> )
2.	1.107	286	1.3661	CO <sub>2</sub>
3.	3.062	1718	8.2192	H <sub>2</sub> O
4.	5.767	4514	21.5902	PROPANE
TOTAL		28605	100	



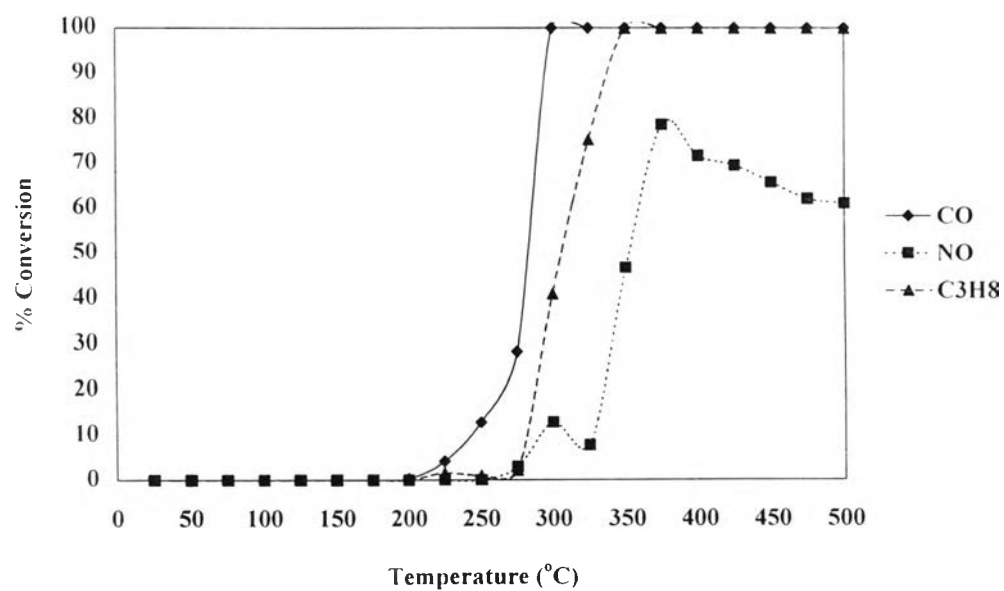
## APPENDIX E

### CO, NO AND C<sub>3</sub>H<sub>8</sub> CONVERSIONS OF PREPARED CATALYSTS

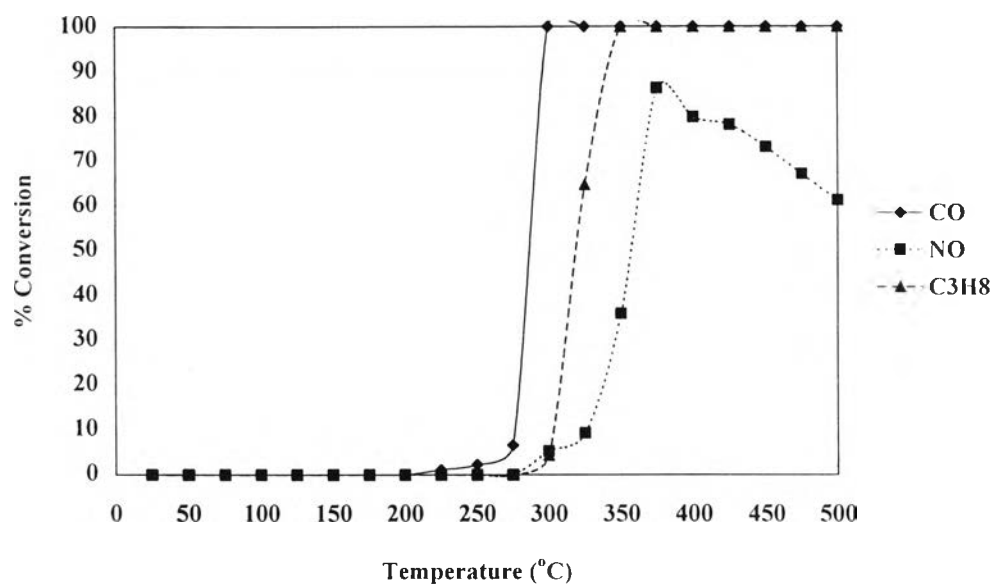
**Figure F-1** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in air at 380°C under stoichiometric condition



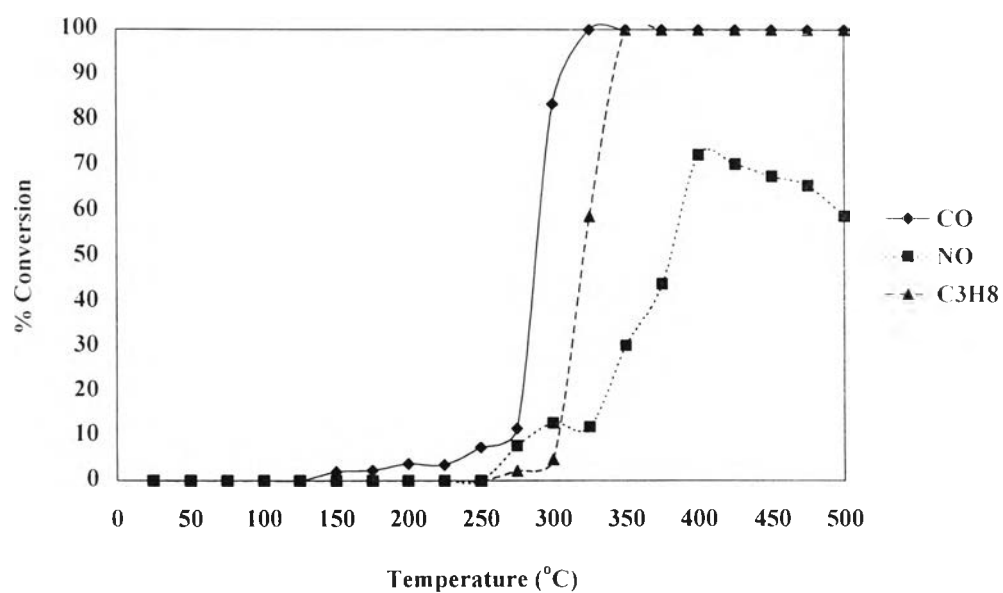
**Figure F-2** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in air at 450°C under stoichiometric condition



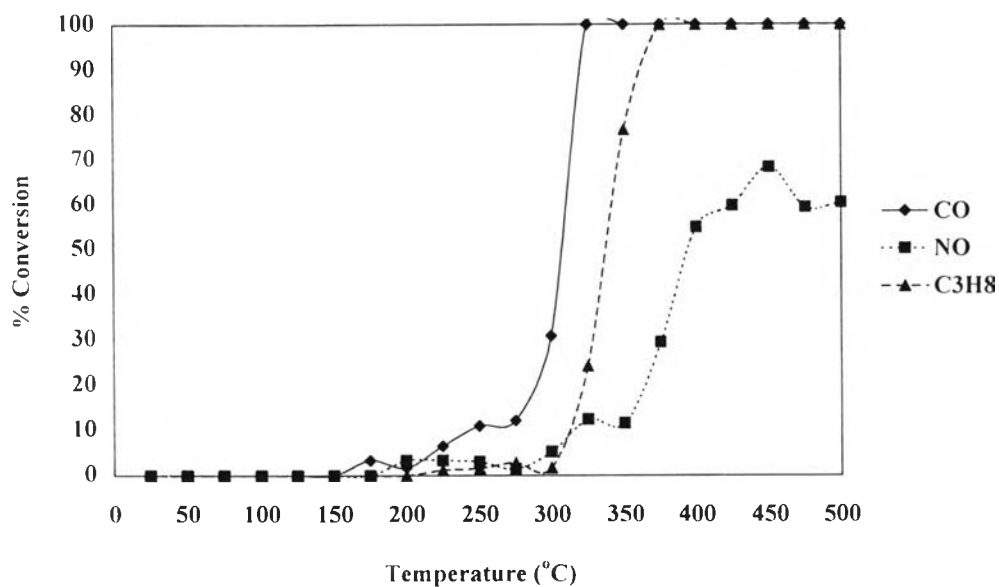
**Figure F-3** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in air at 500°C under stoichiometric condition



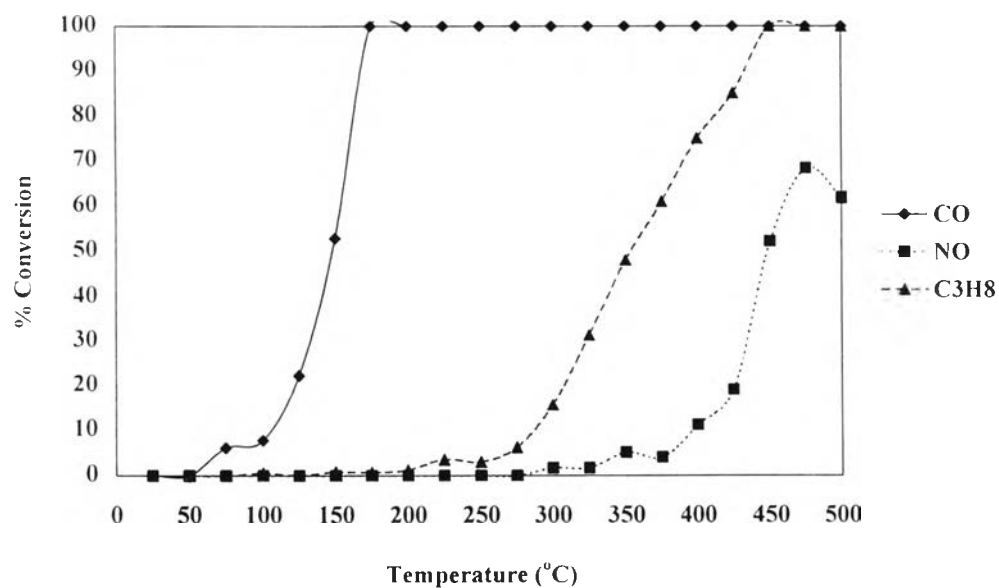
**Figure F-4** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in air at 550°C under stoichiometric condition



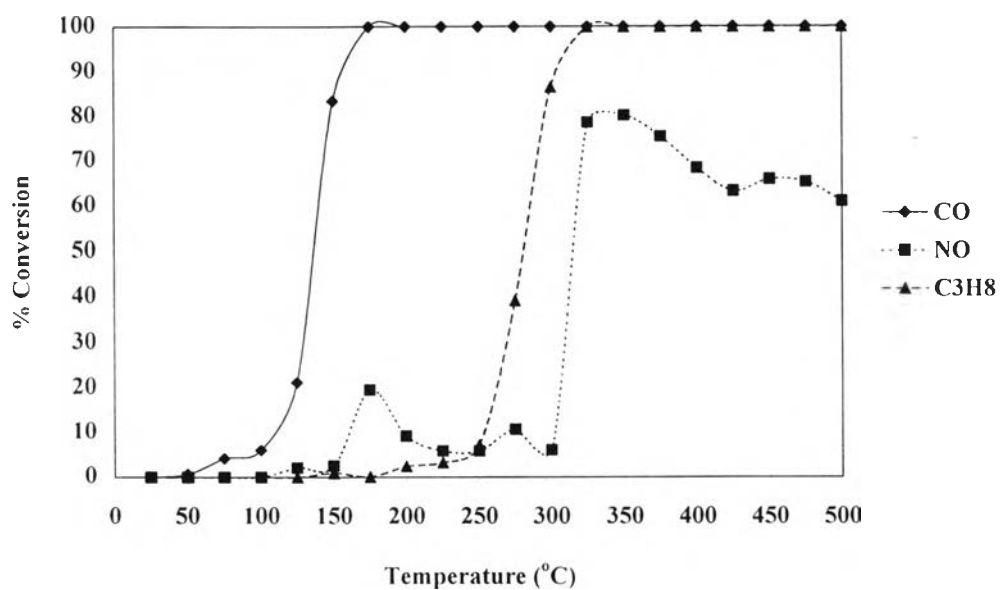
**Figure F-5** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in air at 650°C under stoichiometric condition



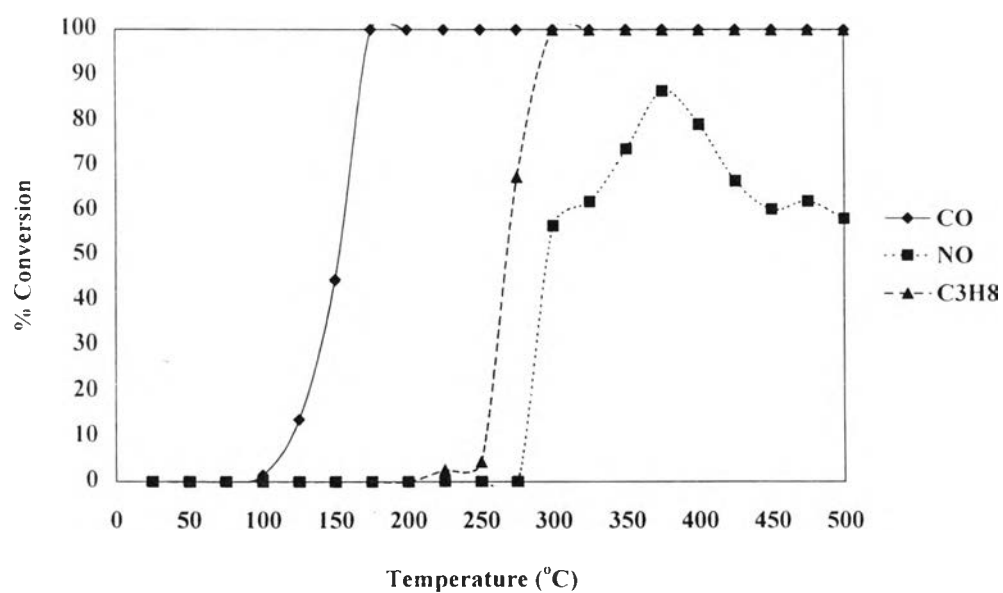
**Figure F-6** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in a reducing atmosphere at 380°C under stoichiometric condition



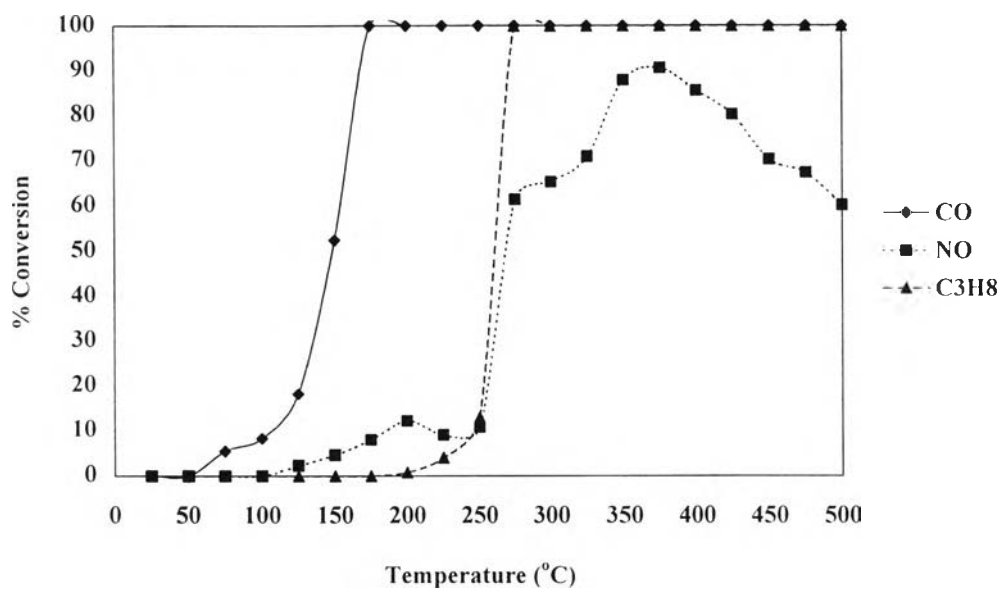
**Figure F-7** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in a reducing atmosphere at 450°C under stoichiometric condition



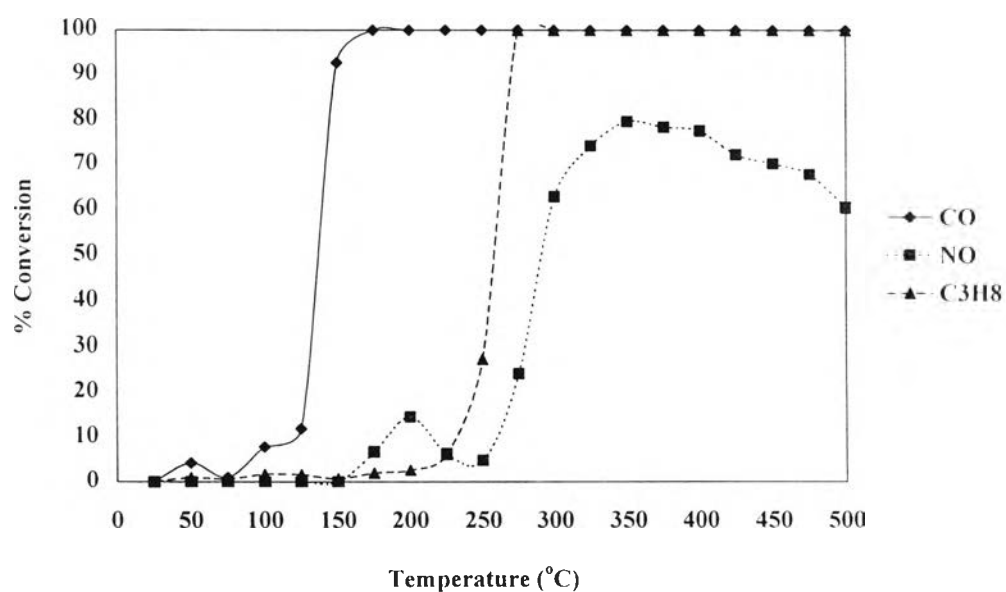
**Figure F-8** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in a reducing atmosphere at 500°C under stoichiometric condition



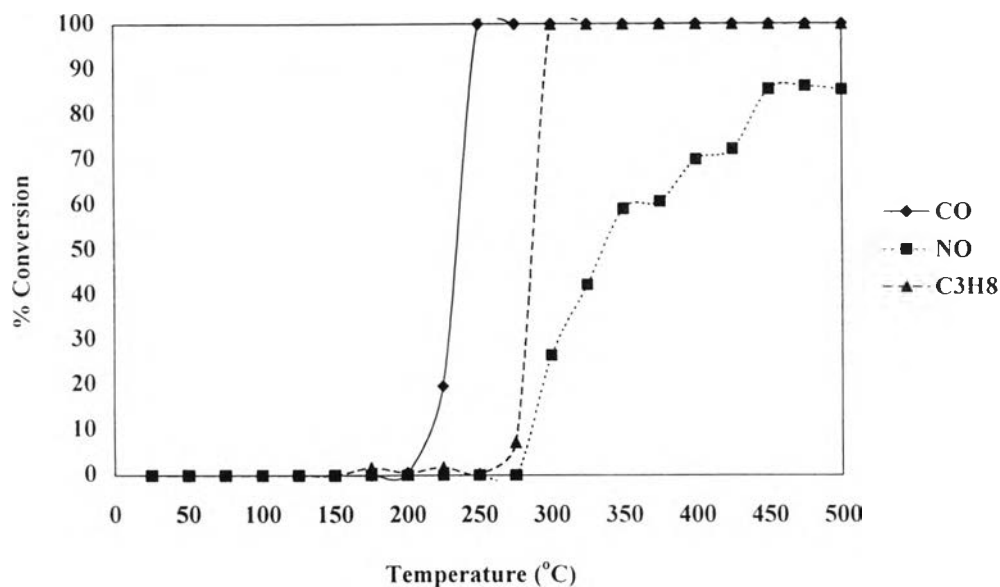
**Figure F-9** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in a reducing atmosphere at 550°C under stoichiometric condition



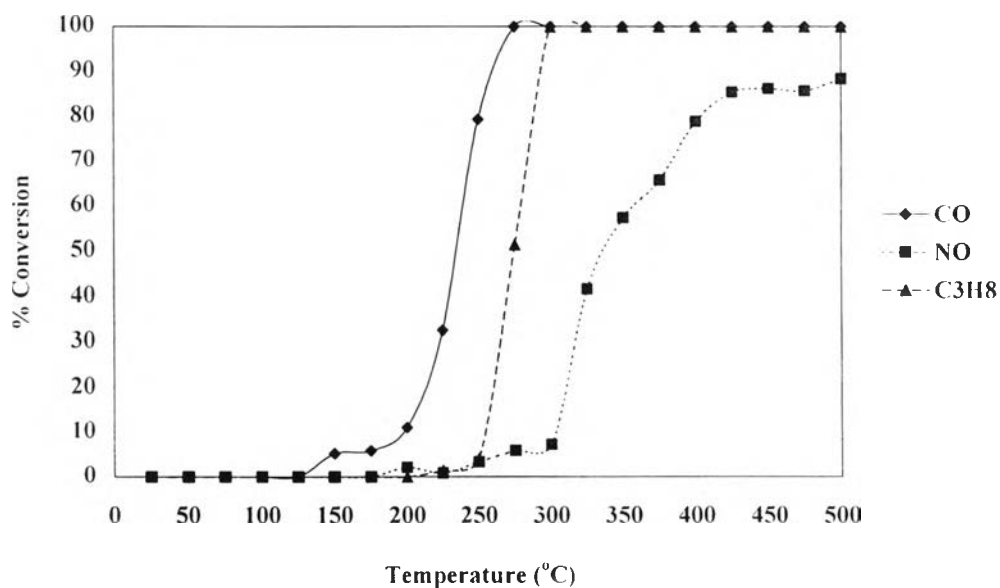
**Figure F-10** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/Al<sub>2</sub>O<sub>3</sub> catalyst calcined in a reducing atmosphere at 650°C under stoichiometric condition



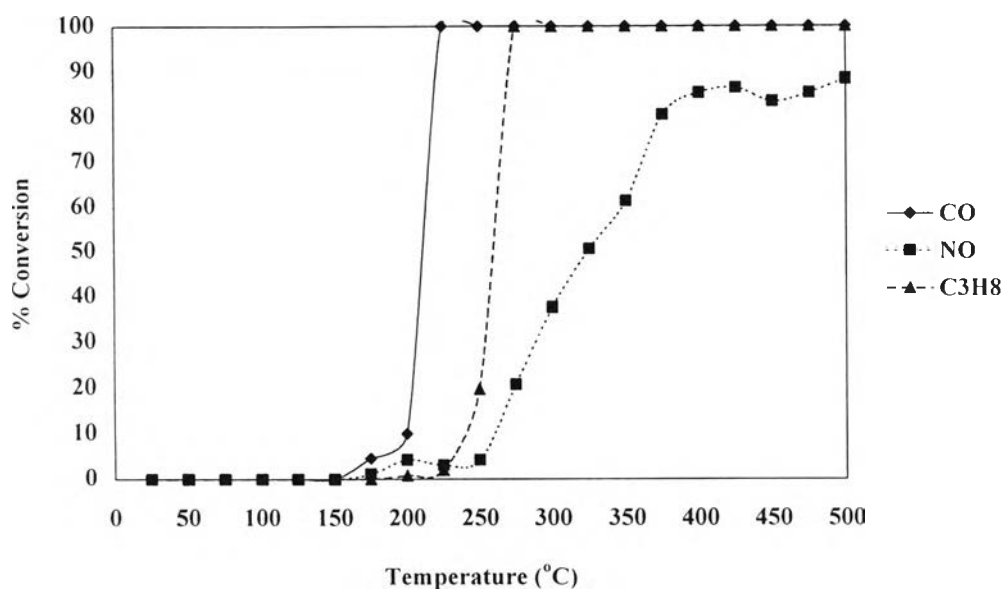
**Figure F-11** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in air at 380°C under stoichiometric condition



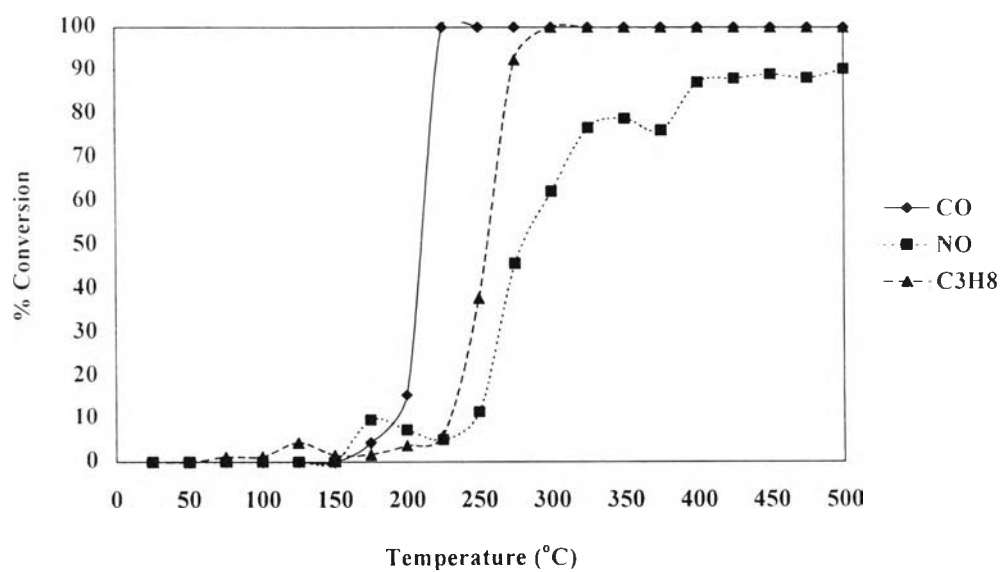
**Figure F-12** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in air at 450°C under stoichiometric condition



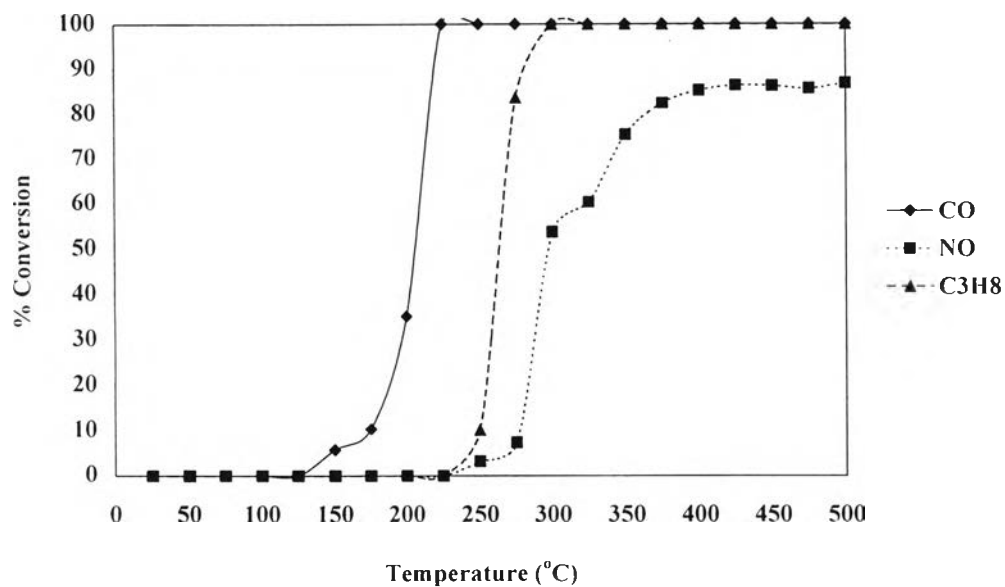
**Figure F-13** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in air at 500°C under stoichiometric condition



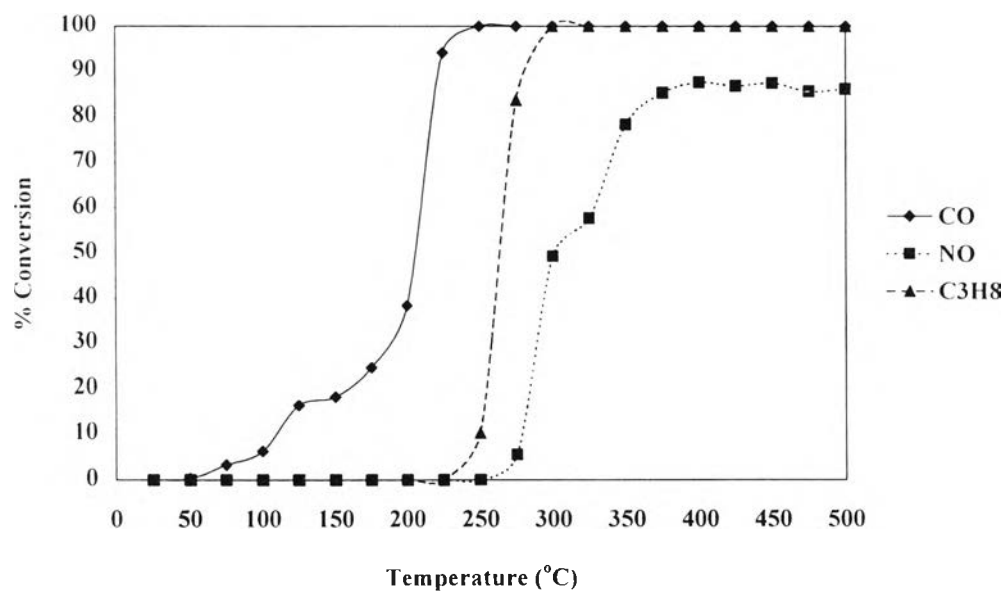
**Figure F-14** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in air at 550°C under stoichiometric condition



**Figure F-15** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in air at 600°C under stoichiometric condition

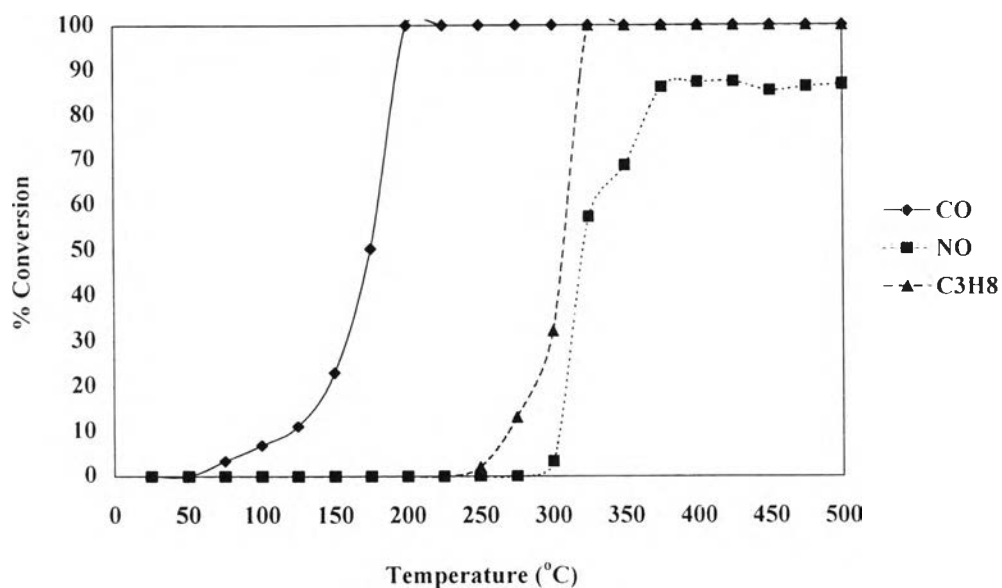


**Figure F-16** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in air at 650°C under stoichiometric condition

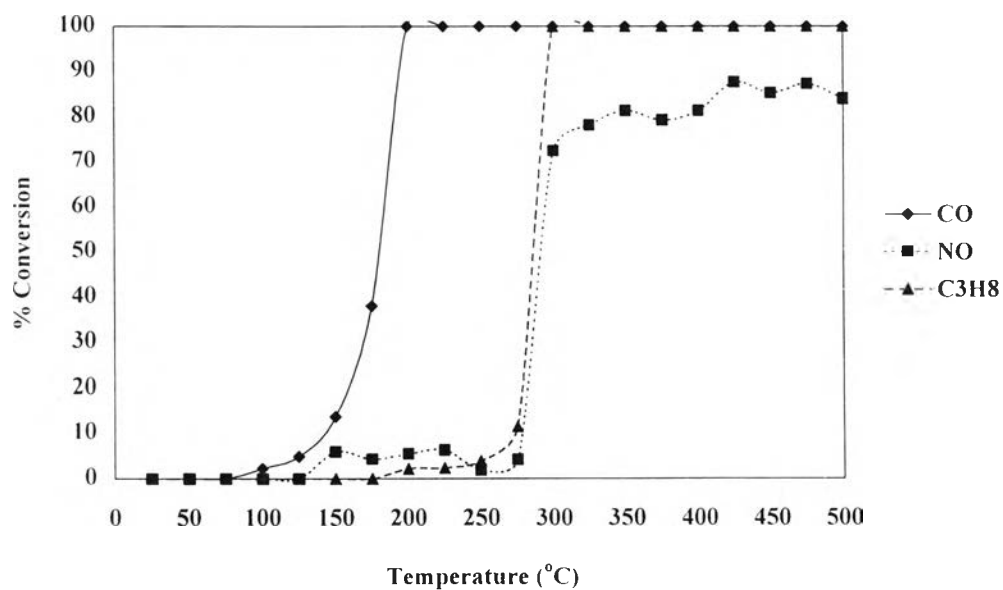




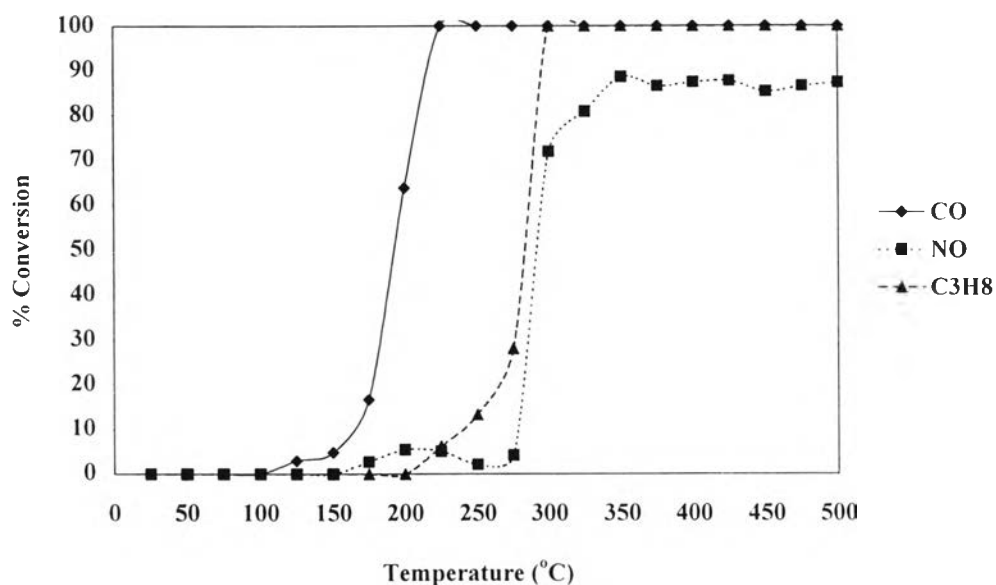
**Figure F-17** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in a reducing atmosphere at 380°C under stoichiometric condition



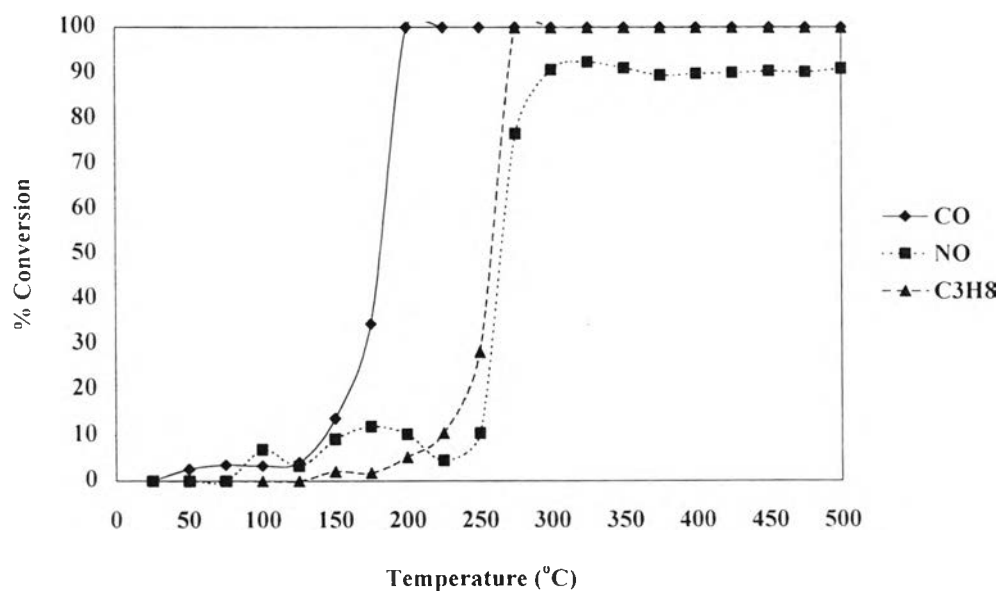
**Figure F-18** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in a reducing atmosphere at 450°C under stoichiometric condition



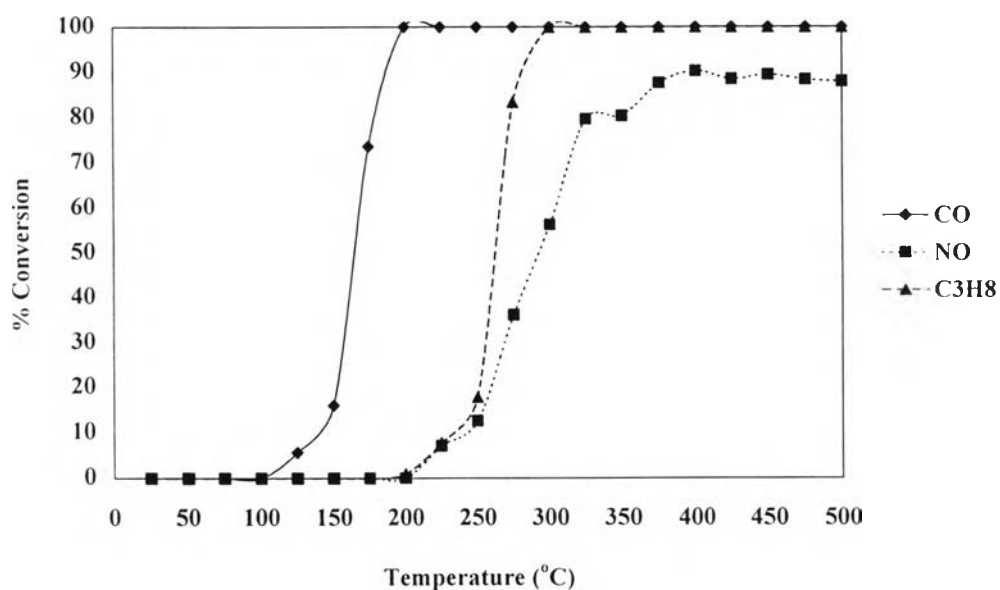
**Figure F-19** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in a reducing atmosphere at 500°C under stoichiometric condition



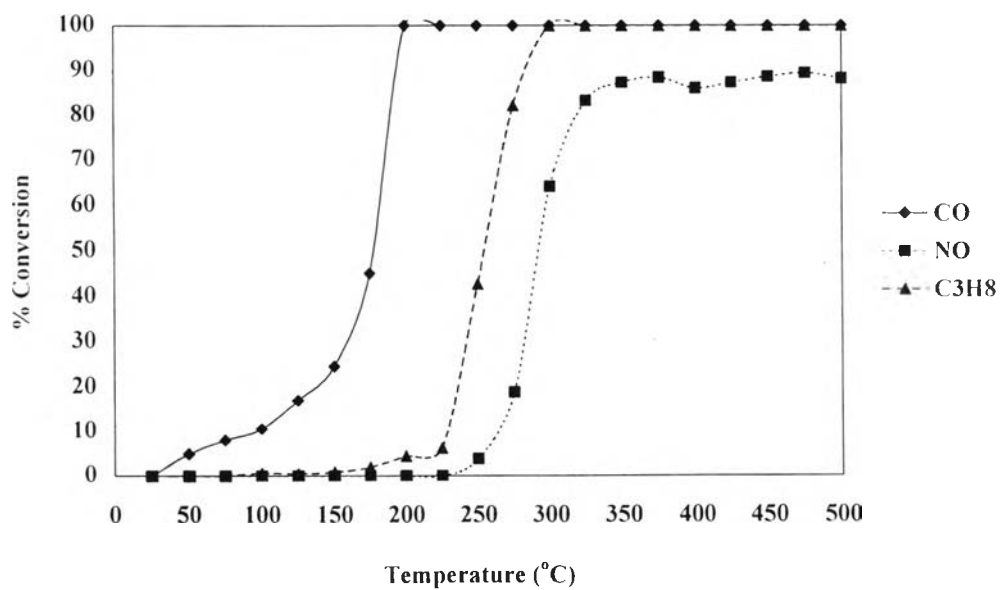
**Figure F-20** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in a reducing atmosphere at 550°C under stoichiometric condition



**Figure F-21** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in a reducing atmosphere at 600°C under stoichiometric condition



**Figure F-22** Conversion of CO, NO and C<sub>3</sub>H<sub>8</sub> over 0.3%Pt/TiO<sub>2</sub> catalyst calcined in a reducing atmosphere at 650°C under stoichiometric condition





## VITA

Mr. Suwat limtrakul was born on October 29, 1976 in Suphanburi, Thailand. He received the Bachelor Degree of Chemical Engineering from Faculty of Engineer, Rangsit University in 1998. He continued his Master's Study at Chulalongkron University in June, 1998.