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Influence of support on the catalytic properties of Pt–Sn–K/ θ -Al₂O₃ for propane dehydrogenation†

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A study on the performance of Pt(0.3 wt%)Sn(0.2 wt%)K(0.5 wt%) catalysts supported on four different θ -Al₂O₃ for propane dehydrogenation is reported in this study. The θ -Al₂O₃ used as supports were prepared by four different methods as (a) calcination of the commercial γ -Al₂O₃ at 1223 K for 12 h, (b) synthesis by hydrochloric acid reflux method, (c) precipitation of Al(NO₃)₃·9H₂O with ammonia solution and (d) extrusion use pseudo-boehmite powder, respectively. They were characterized by using XRD, BET, SEM, H₂-TPR, NH₃-TPD, CO-chemisorption, XPS and TG-DTA methods to study which characteristics of the carrier will affect the performance of the catalyst. The results show that high acidity and strong interactions of the Sn support can improve the propane dehydrogenation activity of the catalyst, large pore volume and large pore diameter can enhance the stability of the catalyst. Al₂O₃ synthesized by method (b) has the largest pore volume, pore diameter, relatively high surface acidity and strong interaction with Sn, which meant the catalyst with the support prepared by method (b) showed the highest propane conversion and superior selectivity. The average conversion is 38.6% and the average selectivity is 95.2% during the reaction time of 25 h.

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Introduction

There is growing interest in the catalytic dehydrogenation of propane due to the high demand for propylene.¹ Propylene is important in the production of various basic compounds, such as propylene oxide, polypropylene, cumene, acrylonitrile, isopropyl alcohol, *etc.*^{2–5} In alkylation and oligomerization reactions, propylene is also used to produce clean fuel with a high octane number, an alkylate-blended, *etc.*⁶ To achieve high yields of propylene, high reaction temperatures (823–883 K) and low pressures are always required since the propane dehydrogenation (PDH) is highly endothermic ($\Delta H_{298}^{\circ} = +124 \text{ kJ mol}^{-1}$) and equilibrium limited. Thus, the deactivation of catalyst caused by coke formation is unavoidable.⁷ To overcome this problem, more efficient catalysts with high-activity, high-selectivity and high-stability are under developed to enhance the propylene yield.

The platinum–tin–potassium-supported catalysts have drawn intense attention in paraffin dehydrogenation reactions.^{8–12} Over the platinum-based catalysts, it is benefit to improve the stability and selectivity while suppressing hydrolysis reactions, cracking and coke formation process. The addition of alkali metal K can generally neutralize some acid

sites of the support. The interaction between metal Pt and Sn, the valence of Sn, the properties of the support and the interactions between metal and support strongly influence the catalytic properties of PtSnK trimetallic catalysts. Sn⁰ may be a poison, while it could also acts as a promoter as Sn⁴⁺ or Sn²⁺.^{13,14} Moreover, suitable textural properties of the support and strong interactions between loaded metals and support are beneficial for the improvement of catalyst performance for propane dehydrogenation.^{15,16} Zhang *et al.*¹⁷ found uniform pore size distribution and a relatively large surface area of the support were favorable for the improvement of the dispersion of active metallic particles. Supports which with large pore size is beneficial for coke to cover the external surface of the support first instead of the metallic surface. There are many papers studied the influence of different supports on the alkane dehydrogenation process,^{17–22} the supports they studied include ZSM-5, SBA-15, SiO₂, spinels (ZnAl₂O₄, MgAl₂O₄), γ -Al₂O₃, *etc.* But the effects of the preparation methods of θ -Al₂O₃ supports on the catalytic propane dehydrogenation performances have not yet been reported so far. In addition, θ -Al₂O₃ was selected as support by UOP Oleflex process in the industry. So, the study about the influence of different θ -Al₂O₃ support on the catalytic propane dehydrogenation is necessary.

In this paper, four different synthesis ways of θ -Al₂O₃ supports have been employed, and were applied as Pt based catalyst supports for propane dehydrogenation. The performance of these catalysts on propane dehydrogenation was systematically compared. Many physicochemical characterization methods were applied to demonstrate the influence of

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synthesize routes on the properties of θ - Al_2O_3 supports and further clarify the influence on PtSnK propane dehydrogenation catalysts, then the most suitable support was selected.

Experimental

Preparation of different alumina supports

Four kinds of θ - Al_2O_3 supports (20–40 mesh) were used in this work. Among them, Al_2O_3 -A support was prepared by calcining the commercial γ - Al_2O_3 at 1223 K for 12 h. Al_2O_3 -B support is synthesized by HCl reflux method. Aluminum foil (99.999%, 40 g) and HCl (11%, 307 g) were mixed under a rotating rate of 50 rpm and slowly heated to 95 °C. When the aluminum foil began to dissolve, the rotating rate was adjusted to 300 rpm. The alumina sol was obtained until the aluminum foil was completely dissolved. Hexamethylenetetramine (HMT) solution (40%, 24 g) was then added to the above alumina sol (70 g). After being well mixed, the mixture was dropped into the oil column. The alumina spheres were then aged at 140 °C for 17 h in an autoclave, which were further treated by washing, drying and calcining at 1223 K for 12 h. Al_2O_3 -C support is synthesized by ammonia precipitation method. Ammonia (8%) was dropped into aluminum nitrate solution (1 mol L⁻¹) at 40 °C and 300 rpm until the pH value reached 8. The precipitation reaction lasted for 1 h and the precipitate was then subjected to aging 4 h, washing with de-ionized water, drying at 120 °C for 12 h and calcining at 1223 K for 12 h. Al_2O_3 -D support is synthesized by extrusion with pseudo-boehmite powder bought from Korea, then dried and calcined at 1223 K for 12 h. The complete conversion to θ - Al_2O_3 of the four supports were proved by XRD measurements.

Catalyst preparation

The trimetallic catalysts were prepared through a previously developed vacuum complex impregnation method.²⁰ A pre-determined amount of $\text{H}_2\text{PtCl}_6 \cdot 6\text{H}_2\text{O}$ (Aladdin, Pt 37.5% min) and $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ (Aladdin, AR, 98%) were dissolved in de-ionized under N_2 (purity 99.995%) atmosphere in order to form Pt–Sn complex, which indicated by the red color of the mixture.²³ Then KNO_3 aqueous solution was added thereafter. The impregnation solution was impregnated onto the alumina supports which were previously degassed for 30 min under vacuum at room temperature. The catalysts were dried at 60–70 °C in vacuum for 30 min before drying at 120 °C overnight. Finally, catalysts were calcined at 600 °C for 6 h in air atmosphere with a ramp rate of 10 °C min⁻¹. Those four catalysts were denoted as Cat-A, Cat-B, Cat-C and Cat-D, respectively. The nominal loading amounts on each sample were 0.3 wt% for Pt, 0.2 wt% for Sn and 0.5 wt% for K.

Catalyst characterizations

The powder X-ray diffraction (XRD) patterns were obtained on an Empyrean powder X-ray diffractometer (Netherlands) at 0 kV and 40 mA in the scan 2θ range of 10–90°.

The textural properties of the alumina supports were calculated from N_2 adsorption–desorption isotherms collected at

liquid nitrogen temperature by a volumetric adsorption system (Quantachrome Autosorb-1, American). All the samples were previously degassed for 5 h under vacuum hood at 300 °C. The Brunauer–Emmett–Teller (BET) method was applied to calculate the specific surface areas of the samples and the desorption branch of the isotherms was used to calculate the average pore diameter by the Barrett–Joyner–Halenda (BJH) pore size model.

The SEM images were taken using a scanning electron microscope (JSM-7800F) operating at 20 kV.

The acidic of each samples were measured by NH_3 -TPD experiments on a Micromeritics Auto-Chem II 2910 (American) chemisorption analyzer. Samples (0.10 g) which pre-dried at 120 °C overnight were placed in a U-type quartz sample tube. NH_3 was saturated at 100 °C after pretreatment at 600 °C for 1 h under helium stream (30 mL min⁻¹). Subsequently, the desorption of ammonia was determined by a thermal conductivity detector at temperatures from 100 °C to 600 °C at a temperature ramp rate of 10 °C min⁻¹.

Temperature-programmed reduction (TPR) method was used to measure the reducibility of the catalysts by a Micromeritics Auto-Chem II 2910 apparatus (American). Samples (0.20 g) which pre-dried at 120 °C overnight were placed in a U-shaped quartz reactor. The samples were pretreated *in situ* by flowing dry argon (99.99%, 30 mL min⁻¹) at 300 °C for 2 h. After cooled to room temperature, 10% H_2 in Ar was switched and the samples were heated to 700 °C at a ramp rate of 10 °C min⁻¹. The hydrogen consumption was monitored by a TCD detector.

The XPS data of Pt–Sn–K/ θ - Al_2O_3 catalysts were investigated on ESCALAB 250Xi using Al K α radiation. All catalysts were previously reduced in a hydrogen flow at 600 °C for 2 h. The binding energies (BE) were calibrated using the C1s level at 284.6 eV with an uncertainty of ± 0.2 eV.

The Pt dispersion of the catalysts were measured by pulse chemisorption of CO experiments (Micromeritics AutoChem II 2910, American). Samples (0.10 g) were pretreated under a He stream at 500 °C for 1 h to remove the moistures and other impurities. Then the samples were reduced under H_2 atmosphere (99.99%, 30 mL min⁻¹) at 600 °C for 1 h, and purged in He (99.99%, 30 mL min⁻¹) at 600 °C for 1 h, CO was saturated after cooling to 50 °C in flowing He (30 mL min⁻¹). The amount of the chemisorbed CO was determined by a TCD detector. A stoichiometry of CO : Pt = 1 was assumed to estimate the dispersion of Pt.

The coke content was measured by a thermo gravimetric and differential thermo analysis (TG-DTA) apparatus (TA Q600, P R China) with a temperature increasing rate of 10 °C min⁻¹ from room temperature to 800 °C in an air flow of 50 mL min⁻¹.

Catalytic activity test

The propane dehydrogenation reaction was performed with the prepared catalysts in a quartz fixed-bed reactor (inner diameter: 8 mm) heated by an electric furnace. The catalysts were heated to 600 °C at a ramp rate of 5 °C min⁻¹ and reduced *in situ* in flowing H_2 for 2 h prior to the reaction. The reaction was conducted at 600 °C with 0.25 g samples and WHSV was 4 h⁻¹ ($\text{H}_2/\text{C}_3\text{H}_8$ molar ratio = 0.5 : 1). Ten minutes later, the gas



compositions of the reactants and products were collected and analyzed by an on-line gas chromatograph (Agilent 7890A, American) using a flame ionization detector (FID). The gas chromatograph was operated under an inlet temperature of 180 °C, an oven temperature of 105 °C and detector temperature of 200 °C.

The propane conversion ($X\%$) and the propylene selectivity (S) were calculated according to the following formulate:

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

$$\text{Propylene selectivity } (S\%) = \frac{n(\text{C}_3\text{H}_6)_{\text{out}}}{n(\text{C}_3\text{H}_8)_{\text{in}} - n(\text{C}_3\text{H}_8)_{\text{out}}} \times 100\%$$

where $n(\text{C}_3\text{H}_8)_{\text{in}}$ represents the molar content of propane in feed, $n(\text{C}_3\text{H}_8)_{\text{out}}$ and $n(\text{C}_3\text{H}_6)_{\text{out}}$ is the molar content of propane and propylene in the product.

Results and discussion

Catalytic performance

The performance of the four Pt-Sn-K catalysts supported on different types of θ - Al_2O_3 are compared in Fig. 1, and significant differences among the four catalysts as a function of the alumina supports could be seen obviously. The conversion of

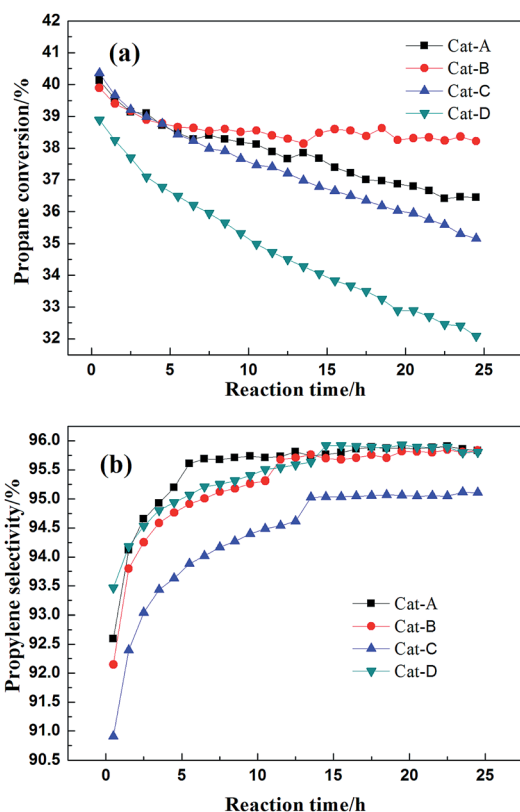


Fig. 1 (a) Propane conversion and (b) propylene selectivity over the four different catalysts (reaction conditions: $T = 600$ °C, $\text{H}_2 : \text{C}_3\text{H}_8 = 0.5 : 1$, $\text{WHSV} = 4 \text{ h}^{-1}$, $m_{\text{cat}} = 0.25 \text{ g}$).

propane catalyzed by those four catalysts were 40.1, 39.9, 40.4 and 38.9%, respectively, initially, and gradually decreased to 36.4%, 38.2%, 35.2% and 32.1%, respectively, after 25 h. A deactivation parameter D (defined as $D = [X_0 - X_f] \times 100\%/X_0$, where X_0 and X_f represent the initial and final propane conversion, respectively) is used to characterize the catalytic stability. The values of D for the four catalysts are 9.4, 4.2, 12.9 and 17.5%, respectively. It is obvious that Cat-B exhibited the highest reaction activity and the lowest deactivation value during the 25 h tests, suggesting that Al_2O_3 -B is more favorable for propane dehydrogenation than the other three supports. Besides, the selectivity of all four catalysts increased gradually with the prolongation of reaction time, which could be explained by that the coke can gradually cover the active metal centers which for coking and cracking reactions with the prolongation of reaction time.^{15,20,24} The average selectivity to propene over the four catalysts during the 25 h tests were 95.5%, 95.2%, 94.3% and 95.4%. The results above shows that the selectivity for all the catalysts is high, but Cat-D prepared by extrusion use pseudo-boehmite powder as support was poorer in propane dehydrogenation activity and stability when compared with others. The by-products (methane, ethane and ethylene) produced in the process were also analyzed (see Fig. S1 in the ESI[†]). It should be noted that under the experimental conditions, Cat-B showed the most satisfactory propane dehydrogenation performance, which attributed to the superior support Al_2O_3 -B.

Properties of alumina supports

XRD patterns of the four types of alumina supports synthesized by different methods are shown in Fig. 2. It can be observed that all samples display the similar XRD patterns. The peaks at $2\theta = 19.54^\circ, 32.78^\circ, 39.86^\circ, 44.83^\circ, 50.70^\circ$ and 67.42° , with d values of 0.454, 0.273, 0.226, 0.202, 0.180 and 0.139 nm, respectively, are assigned to monoclinic θ - Al_2O_3 phase (ICDD file no. 00-035-0121), which should be ascribed to the high calcination temperature of 1223 K.^{25,26}

The isotherms and the pore size distributions of four alumina supports are presented in Fig. 3. According to the

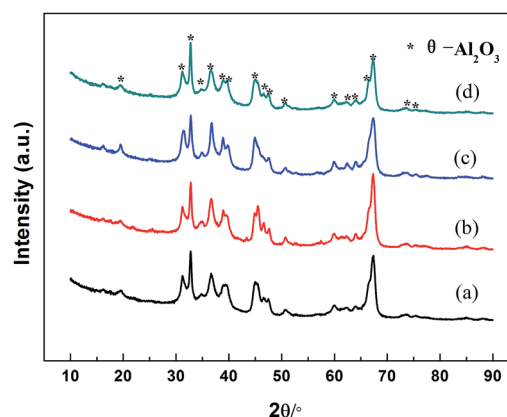


Fig. 2 XRD patterns of (a) Al_2O_3 -A, (b) Al_2O_3 -B, (c) Al_2O_3 -C and (d) Al_2O_3 -D.



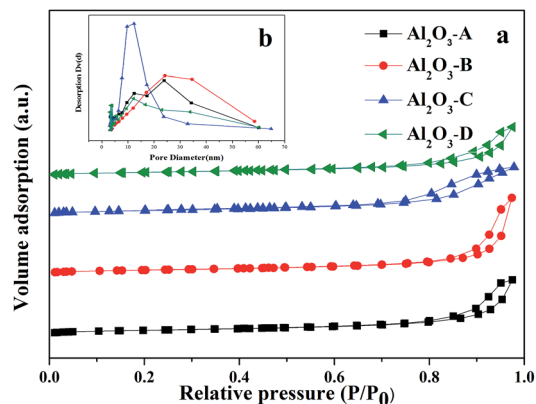


Fig. 3 Nitrogen adsorption-desorption isotherms and pore size distributions of different alumina supports.

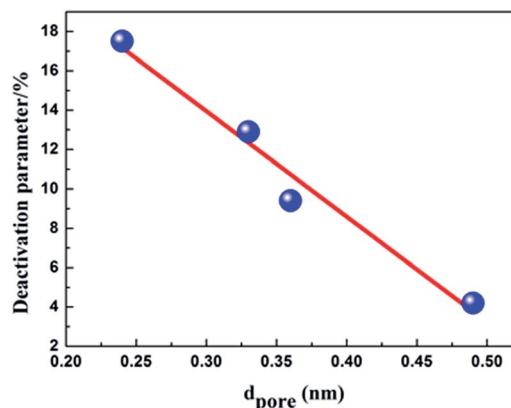


Fig. 4 Deactivation parameters versus average pore diameter of the four alumina supports.

IUPAC classification,²⁷ all samples exhibit type IV N_2 adsorption isotherm with H1-type hysteresis loop, which suggested the presence of meso-pores or macro-pores.^{28,29} When the size of meso-pores in the alumina support decrease (Fig. 3b), the capillary condensation steps will slightly shift to lower relative pressures (Fig. 3a), which is in accord with the literature.¹⁵

The textural properties such as surface area, pore size and pore volume of the samples were calculated from the nitrogen adsorption-desorption study and are listed in Table 1. It is obvious that the four supports are quite different in surface area, pore size distributions and total pore volume. Al_2O_3 -C presented the largest surface area of $104.6 \text{ m}^2 \text{ g}^{-1}$ with abundant small pores,³⁰ while Al_2O_3 -D with less pores only shows the smallest surface area as $56.6 \text{ m}^2 \text{ g}^{-1}$. On the other hand, Al_2O_3 -B shows the largest pore volume ($0.49 \text{ cm}^3 \text{ g}^{-1}$), pore diameter (24.2 nm) and the widest pore size distribution.

It is well known, propane dehydrogenation catalysts lost the activity mainly due to the accumulation of coke.³¹⁻³³ From Fig. 4, we can see that the deactivation parameter of the four catalysts is nearly negatively linear correlated with the average pore diameter, that is to say, the stability of the catalyst increased with the increase of the pore diameter. The deactivation parameter decreased with the increase of the total pore volume too (see Fig. S2 in the ESI†). Those results proved that the catalyst support which have large pore size and pore volume can more effectively overcome diffusion and mass transfer limitations, and could help the coke deposits migrate from the active sites to supports, improve the stability of the catalysts. For Al_2O_3 -B synthesized by hydrochloric acid reflux method has the

largest pore volume ($0.49 \text{ cm}^3 \text{ g}^{-1}$), pore diameter (24.2 nm), the widest pore size distribution, we can speculate that Cat-B ought to be more resistant to coking, show best stability during the reaction, reasonably, and it fits the experimental results well.

The surface structure of alumina synthesized by different methods was revealed by a SEM technique. Fig. 5 showed the representative SEM images of the four alumina samples. It can be seen that Al_2O_3 -A and Al_2O_3 -B show wider pore size distribution and larger pore size, which is in accordance with the results of nitrogen adsorption-desorption study. The pores with diameter of more than 20 nm could also be found in the images. For Al_2O_3 -C and Al_2O_3 -D, most pores of less than 20 nm in diameter implied that the mesoporous structure is exist.

Physicochemical properties of Pt-Sn-K/ θ - Al_2O_3 catalysts

As determined from CO pulse chemisorption, the Pt dispersion and particle sizes are listed in Table 2. Cat-B exhibits the highest Pt dispersion and the smallest Pt particle size which may be ascribed to its wide pore size distribution and relatively large surface area. According to the related studies,³⁴ the Pt particle

Table 1 Textural properties of different alumina supports

Supports	$S_{\text{BET}}(N_2)$ ($\text{m}^2 \text{ g}^{-1}$)	V_{total}^a ($\text{cm}^3 \text{ g}^{-1}$)	d_{pore}^b (nm)
Al_2O_3 -A	77.5	0.36	23.8
Al_2O_3 -B	81.0	0.49	24.2
Al_2O_3 -C	104.6	0.33	12.3
Al_2O_3 -D	56.6	0.24	12.2

^a Total pore volume. ^b Average pore diameter.

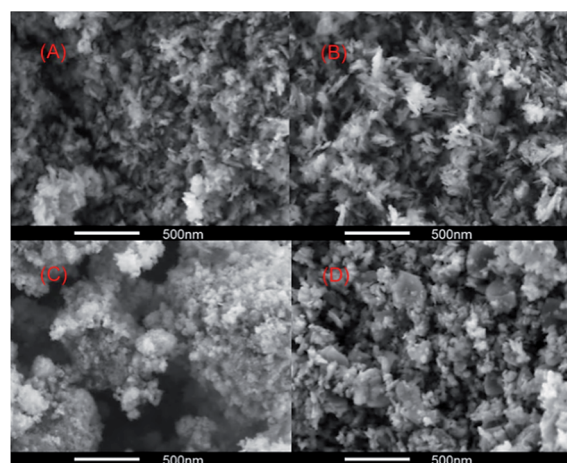


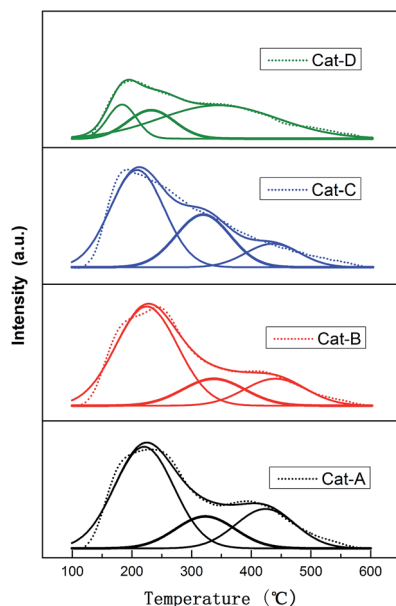
Fig. 5 SEM morphologies of (A) Al_2O_3 -A, (B) Al_2O_3 -B, (C) Al_2O_3 -C and (D) Al_2O_3 -D.



Table 2 Characterization of the four catalysts

Catalysts	Pt dispersion ^a (%)	Average particle size ^a (nm)	Average conversion ^b (%)	Coke contents ^c (%)	Deactivation parameter ^d (%)
Cat-A	51.1	2.2	37.8	2.61	9.4
Cat-B	55.1	2.0	38.6	2.99	4.2
Cat-C	37.5	3.0	37.3	3.66	12.9
Cat-D	47.6	2.4	34.8	0.99	17.5

^a Calculated by CO-pulse chemisorption of fresh PtSnK catalysts. ^b The average conversion of propane during the 25 h reaction. ^c The coke contents of the deactivated catalysts by thermogravimetric experiments. ^d Deactivation parameter for the 25 h reaction.

Fig. 6 NH₃-TPD curves of the catalysts.

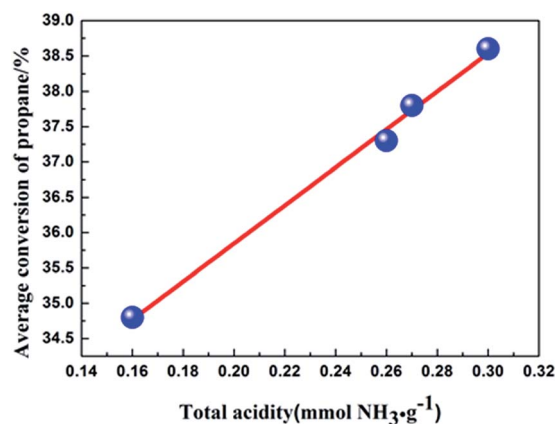
size has a significant influence on the dehydrogenation activity of Pt-Sn-K/Al₂O₃ catalysts since the dehydrogenation reaction is more likely to take place on small Pt clusters on the catalyst surface and the side reactions need larger platinum ensembles. Therefore, the superior activity of Cat-B could be revealed partly by the smaller Pt particles present on its surface.

NH₃-TPD method was employed to determine the acidity of solid catalysts. It is well known that the acid properties of the catalyst markedly affect the dehydrogenation performance of the catalyst. The acidity of the four Pt-Sn-K/Al₂O₃ catalysts was examined by NH₃-TPD profiles as depicted in Fig. 6.

Gaussian deconvolution method was used to the semi-quantitative analysis of the ammonia desorption peaks. The

calculated results are compiled in Table 3. All samples exhibit three peaks. For Cat-A, Cat-B and Cat-C, the peak centered around 210 °C (peak I), 320 °C (peak II) and 430–450 °C (peak III) should be attributed to weak, medium and strong acid sites, respectively. For Cat-D, the first (peak I) and the second (peak II) peak can be ascribed to weak acid sites, and the third peak (peak III) can be assigned to medium and strong acid sites.^{35,36} According to the total desorption peak area, it can be inferred that the order of the total acid content of the four catalysts are as follows: Cat-B > Cat-A ≈ Cat-C > Cat-D. Note that a majority of acidic centers exhibited on Cat-A, Cat-B and Cat-C is weak or medium acid sites. But as for Cat-D, the medium or strong acidic centers are dominant.

The average conversion of propane *versus* the total acidity of the four catalysts are depicted in Fig. 7. Interestingly, the relationship between the average conversion of propane and the total acidity of the four catalysts is linear. The results indicate that the acidity of the catalyst not only affect the distribution of

Fig. 7 Average conversion of propane *versus* total acidity of the four catalysts.Table 3 The semi-quantitative analysis data of NH₃-TPD profiles

Catalysts	T_M (°C)			Total acidity (mmol NH ₃ per g)	Peak fraction (%)			Fitted parameters (R^2)
	I	II	III		I	II	III	
Cat-A	216.7	314.7	432.4	0.27	0.47	0.39	0.14	0.9823
Cat-B	221.3	330.1	451.3	0.30	0.51	0.41	0.08	0.9878
Cat-C	208.8	319.4	436.0	0.26	0.56	0.30	0.14	0.9819
Cat-D	183.5	232.2	345.1	0.16	0.16	0.21	0.63	0.9923



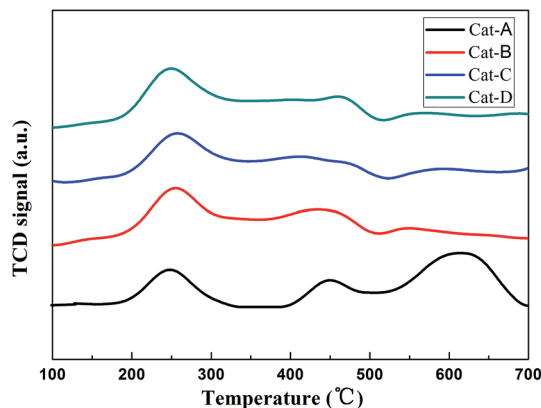


Fig. 8 H₂-TPR curves of the four catalysts.

the products, but also influence the propane dehydrogenation activity of the catalyst in our experiment.

The influences of the four alumina supports on the reduction properties were characterized by H₂-TPR as shown in Fig. 8. It is evident that all the catalysts show at least three reduction peaks at about 250 °C (peak I), 440 °C (peak II) and 600 °C (peak III), respectively. For the signal at 250 °C is belong to the reduction of Pt oxides,³⁷ whereas the peaks at high-temperature can be ascribed to the partial reduction of Sn⁴⁺ to Sn²⁺ and Sn²⁺ to Sn⁰, respectively.^{38,39} Cat-A presents an evident reduction peak higher than 600 °C which corresponding to the reduction of Sn²⁺ to Sn⁰. For Cat-B, the reduction peak corresponding to Sn²⁺ to Sn⁰ is the smallest, and most Sn can keep in the oxidation state, which can act as a promoter, this implying strong interactions of the Sn support.

The XPS spectra and the detailed data of Sn3d_{5/2} of Pt-Sn-K/θ-Al₂O₃ catalysts are shown in Fig. 9 and Table 4, respectively. The Sn3d_{5/2} XPS spectra of the four catalysts were deconvoluted into three peaks at ~485.3, ~486.5 and ~487.5 eV in Fig. 9, corresponding to different chemical states of Sn. Generally speaking, the component at low binding energy (~485.3 eV) is assigned to the reduced tin phase, either in the metallic (Sn⁰) or in the alloyed (SnPt_x) state; whereas the two others (~486.5 and ~487.5 eV) are ascribed to oxides tin with different types (Sn²⁺ or Sn⁴⁺).^{33,40} However, Sn²⁺ and Sn⁴⁺ can't be discriminate

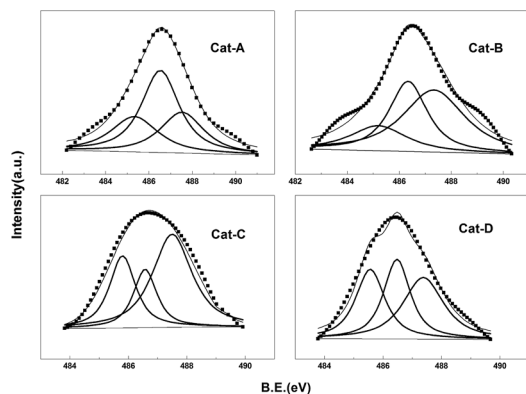


Fig. 9 Sn3d_{5/2} XPS spectra of the four catalysts.

Table 4 Sn3d_{5/2} XPS peak spectra analysis results of the catalysts

Catalysts	Binding energy (eV)		
	Sn3d _{5/2}		
Cat-A	485.3(25.4%)	486.5(47.4%)	487.5(27.2%)
Cat-B	485.2(17.7%)	486.3(33.7%)	487.3(48.6%)
Cat-C	485.8(27.2%)	486.6(19.9%)	487.5(52.9%)
Cat-D	485.5(30.6%)	486.5(31.9%)	487.4(37.5%)

according to XPS spectra alone because their binding energy is very close.^{41–43}

According to the peak percentage given in parentheses in Table 4, the zero valent Sn percentage of the four catalysts are 25.4%, 17.7%, 27.2% and 30.6%, respectively. Tin species in oxidized form is dominant. Sn in zero valent is harmful for catalyst performance and Sn in the oxidation state is benefit for propane dehydrogenation. The presence of SnO_x can increase the catalytic stability for they can keep the Pt sites clean from coke deposits.⁴⁴ This implies that different synthesis routes of alumina can influence the properties of the support, and an appropriate alumina synthesis route could strengthen the Sn–Al₂O₃ interaction, stabilizing the oxidized tin species. For the zerovalent Sn percentage of Cat-B is slightly lower than that of the other three catalysts, which is in accordance with the TPR experiment, indicates that the alumina synthesized by oil-dropped method can interact stronger with Sn species and restrain the reduction of tin species. Cat-B which have the lowest Sn⁰ percentage shows the highest average propane conversion and the lowest deactivation parameter.

As mentioned before, the main reason for the catalyst deactivation in such studies is coking.^{45,46} After the reaction of 25 h, the coke amount deposited over the four catalysts was analyzed by TG measurements as depicted in Fig. 10. The weight losses above 300 °C are attributed to the combustion of coke deposited on the catalysts.⁴⁷ Fig. 10 illustrates that the contents of coke deposited on the four catalysts are 2.61%, 2.99%, 3.66% and 0.99%, respectively, there is no obvious correlation between the amount of coke on the spent catalysts and the final catalytic activity. The highest amount of coke is

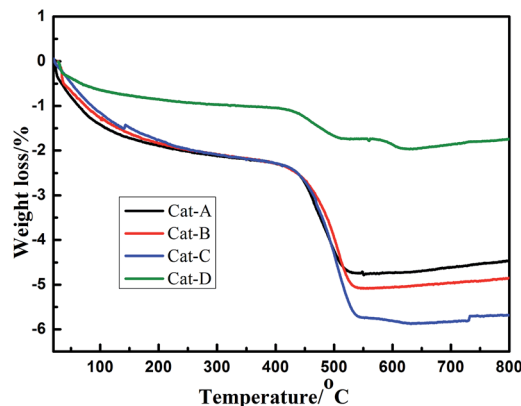


Fig. 10 The TG profiles of deactivated catalysts.



observed over Cat-C may due to its lowest propene selectivity, and the coke on Cat-D with the lowest propane conversion is the least. Although the amount of coke over Cat-B is more than Cat-A, the activity of Cat-B in propane dehydrogenation experiment is better. As mentioned before that large pore size and large pore volume is benefit to overcome mass transfer and diffusion limitations, and could help the coke deposit migration from active sites to supports. With the addition of HMT, the pore-forming mechanism of Al₂O₃-B is very different from the other three alumina support, the decomposition of HMT during the calcination process can produce relatively large pores for Al₂O₃-B. Therefore, we can conclude that the coke in Cat-B is mainly located on the alumina support and does not block active sites, and therefore, though the coke content of Cat-B is not the least, it keeps the best final catalytic activity and the best stability.

Conclusions

In this work, the influences of four different θ -Al₂O₃ supports (synthesized by different methods) on catalytic structure and reaction performances of PtSnK catalysts for propane dehydrogenation were investigated. The correlation between the physicochemical parameters of the catalysts and the dehydrogenation performance of propane was established, and these could afford some guidance for the selection of a θ -Al₂O₃ support for propane dehydrogenation process. High propane dehydrogenation activity was achieved on the catalyst with high acidity and strong interactions of the Sn support. Good stability was achieved on the catalyst with large pore volume and pore diameter. Cat-B with the support prepared by hydrochloric acid reflux method has the largest pore volume (0.49 cm³ g⁻¹), pore diameter (24.2 nm), the widest pore size distribution, relatively large surface area (81.0 m² g⁻¹), and it also has relatively high surface acidity and stronger interactions with Sn species compared to the other three catalysts. All those characteristics make Cat-B an optimal catalytic performance, higher propane conversion and catalytic stability, the average conversion is 38.6%, the average selectivity is 95.2% and the deactivation parameter is only 4.2% during the reaction time of 25 h.

Conflict of interest

We declare that we do not have any conflict of interest.

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References

- 1 F. T. Zangeneh, A. Taeb, K. Gholivand and S. Sahebdehfar, *Appl. Surf. Sci.*, 2015, **357**, 172.
- 2 A. Corma, F. V. Melo, L. Sauvanaud and F. Ortega, *Catal. Today*, 2005, **107**, 699.
- 3 Q. Li, Z. Sui, X. Zhou and D. Chen, *Appl. Catal., A*, 2011, **398**, 18.
- 4 J. S. Plotkin, *Catal. Today*, 2005, **106**, 10.
- 5 R. Watanabe, Y. Hondo, K. Mukawa, C. Fukuhara, E. Kikuchi and Y. Sekine, *J. Mol. Catal. A: Chem.*, 2013, **377**, 74.
- 6 M.-H. Lee, B. M. Nagaraja, P. Natarajan, T. Ngoc Thanh, K. Y. Lee, S. Yoon and K.-D. Jung, *Res. Chem. Intermediat.*, 2016, **42**, 123.
- 7 K. H. Kang, T. H. Kim, W. C. Choi, Y.-K. Park, U. G. Hong, D. S. Park, C.-J. Kim and I. K. Song, *Catal. Commun.*, 2015, **72**, 68.
- 8 D. Rodriguez, J. Sanchez and G. Arteaga, *J. Mol. Catal. A: Chem.*, 2005, **228**, 309.
- 9 Y. Zhang, Y. Zhou, X. Sheng, L. Wan, Y. Li, Y. Xiao, B. Yu and Z. Zeng, *Fuel Process. Technol.*, 2012, **104**, 23.
- 10 Z. Ma, Y. Mo, J. Li, C. An and X. Liu, *J. Nat. Gas Sci. Eng.*, 2015, **27**, 1035.
- 11 Y. Zhang, Y. Zhou, S. Zhang, S. Zhou, X. Sheng, Q. Wang and C. Zhang, *J. Mater. Sci.*, 2015, **50**, 6457.
- 12 B. M. Nagaraja, H. Jung, D. R. Yang and K.-D. Jung, *Catal. Today*, 2014, **232**, 40.
- 13 W. S. Yang, L. W. Lin, Y. N. Fan and J. L. Zang, *Catal. Lett.*, 1992, **12**, 267.
- 14 R. Burch and L. C. Garla, *J. Catal.*, 1981, **71**, 360.
- 15 S. Zhou, Y. Zhou, J. Shi, Y. Zhang, X. Sheng and Z. Zhang, *J. Mater. Sci.*, 2015, **50**, 3984.
- 16 L. Bai, Y. Zhou, Y. Zhang, H. Liu and X. Sheng, *Ind. Eng. Chem. Res.*, 2009, **48**, 9885.
- 17 Y. W. Zhang, Y. M. Zhou, J. J. Shi, S. J. Zhou, X. L. Sheng, Z. W. Zhang and S. M. Xiang, *J. Mol. Catal. A: Chem.*, 2014, **381**, 138.
- 18 S. J. Zhou, Y. M. Zhou, Y. W. Zhang, X. L. Sheng and Z. W. Zhang, *J. Chem. Technol. Biotechnol.*, 2016, **91**, 1072.
- 19 C. T. Shao, W. Z. Lang, X. Yan and Y. J. Guo, *RSC Adv.*, 2017, **7**, 4710.
- 20 S. He, C. Sun, Z. Bai, X. Dai and B. Wang, *Appl. Catal., A*, 2009, **356**, 88.
- 21 S. Bocanegra, A. Ballarini, O. Scelza and S. de Miguel, *Mater. Chem. Phys.*, 2008, **111**, 534.
- 22 S. Bocanegra, A. Ballarini, P. Zgolicz, O. Scelza and S. de Miguel, *Catal. Today*, 2009, **143**, 334.
- 23 L. Y. Jin, *Appl. Catal.*, 1991, **72**, 33.
- 24 C. Yu, H. Xu, Q. Ge and W. Li, *J. Mol. Catal. A: Chem.*, 2007, **266**, 80.
- 25 I. Levin and D. Brandon, *J. Am. Ceram. Soc.*, 1998, **81**, 1995.
- 26 J. M. Hao, *Polyhedron*, 1996, **15**, 2421.
- 27 K. S. W. Sing, D. H. Everett, R. A. W. Haul, L. Moscou, R. A. Pierotti, J. Rouquerol and T. Siemieniowska, *Pure Appl. Chem.*, 1985, **57**, 603.
- 28 C. Larese, J. M. Campos-Martin and J. L. G. Fierro, *Langmuir*, 2000, **16**, 10294.
- 29 M. A. Wahab and C. S. Ha, *J. Mater. Chem.*, 2005, **15**, 508.
- 30 L. D. Sharma, M. Kumar, A. K. Saxena, M. Chand and J. K. Gupta, *J. Mol. Catal. A: Chem.*, 2002, **185**, 135.



- 31 G. J. Siri, G. R. Bertolini, M. L. Casella and O. A. Ferretti, *Mater. Lett.*, 2005, **59**, 2319.
- 32 M. L. Casella, G. J. Siri, G. F. Santori, O. A. Ferretti and M. M. Ramirez-Corredores, *Langmuir*, 2000, **16**, 5639.
- 33 S. A. Bocanegra, A. Guerrero-Ruiz, S. R. de Miguel and O. A. Scelza, *Appl. Catal., A*, 2004, **277**, 11.
- 34 H. S. B. Dalian, *Studies on the catalysts and the coke deposition behavior of the dehydrogenation of long chain paraffins(C₁₀-C₁₉)*, Dalian Institute of Chemical Physics, Chinese Academy of Sciences, 2009.
- 35 P. Iengo, M. Di Serio, V. Solinas, D. Gazzoli, G. Salvio and E. Santacesaria, *Appl. Catal., A*, 1998, **170**, 225.
- 36 P. Iengo, M. Di Serio, A. Sorrentino, V. Solinas and E. Santacesaria, *Appl. Catal., A*, 1998, **167**, 85.
- 37 L. S. Carvalho, C. L. Pieck, M. C. Rangel, N. S. Figoli, J. M. Grau, P. Reyes and J. M. Parera, *Appl. Catal., A*, 2004, **269**, 91.
- 38 Y. Zhang, Y. Zhou, H. Liu, Y. Wang, Y. Xu and P. Wu, *Appl. Catal., A*, 2007, **333**, 202.
- 39 J. Shi, Y. Zhou, Y. Zhang, S. Zhou, Z. Zhang, J. Kong and M. Guo, *J. Mater. Sci.*, 2014, **49**, 5772.
- 40 A. D. Ballarini, C. G. Ricci, S. R. de Miguel and O. A. Scelza, *Catal. Today*, 2008, **133**, 28–34.
- 41 J. C. Serrano-Ruiz, G. W. Huber, M. A. Sanchez-Castillo, J. A. Dumesic, F. Rodriguez-Reinoso and A. Sepulveda-Escribano, *J. Catal.*, 2006, **241**, 378–388.
- 42 F. Coloma, A. Sepulveda-Escribano, J. L. G. Fierro and F. Rodriguez-Reinoso, *Appl. Catal., A*, 2009, **148**, 63–80.
- 43 F. Coloma, A. Sepulveda-Escribano, J. L. G. Fierro and F. Rodriguez-Reinoso, *Appl. Catal., A*, 1996, **136**, 231–248.
- 44 A. Iglesias-Juez, A. M. Beale, K. Maaijen, T. C. Weng, P. Glatzel and B. M. Weckhuysen, *J. Catal.*, 2010, **276**, 268–279.
- 45 Q. Li, Z. Sui, X. Zhou, Y. Zhu, J. Zhou and D. Chen, *Top. Catal.*, 2011, **54**, 888.
- 46 Z. Nawaz, F. Baksh, J. Zhu and F. Wei, *J. Ind. Eng. Chem.*, 2013, **19**, 540.
- 47 N. Martin, M. Viniegra, R. Zarate, G. Espinosa and N. Batina, *Catal. Today*, 2005, **107**, 719.

