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Differences in the catalytic properties of Fe isotopes

Songtao Xiao,^{†*} Yubing Xue,^{ID} Jing Zhao, Xiechun Liu, Haifeng Cong, Tian Lan, Xiaojuan Liu, Dashu Pan,^{ID} Lingyu Wang, Guoan Ye^{*} and Yinggen Ouyang ^{ID*}

The significant differences in the catalytic properties caused by different 'isotopic catalysts' were discovered for the first time. The commonly purchased Fe_2O_3 is a 'mixture' of different Fe isotopic oxides which means the catalytic effect of Fe_2O_3 is theoretically a synthetical result of all isotopic compounds. In this work, the differences in catalytic properties of $\alpha\text{-Fe}_2\text{O}_3$ with natural abundance ratio and separated isotopic $\alpha\text{-Fe}_2\text{O}_3$ ($\alpha\text{-}^{54}\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{56}\text{Fe}_2\text{O}_3$, and $\alpha\text{-}^{58}\text{Fe}_2\text{O}_3$) catalyzing thermal decomposition of ammonium perchlorate (AP) were investigated, and are mainly attributed to the difference in the charge distribution of the nuclei of different iron isotopes. The result suggests that isotope effects in different isotopes when utilized as catalysts are caused by nuclear morphology and the nuclear charge distribution. This study will serve as a base as well as an initiation for future studies of the isotopic catalyst.

Introduction

Isotopes are nuclides of the same element with the same charge number (Z) and electron shell structure but different neutron numbers. Since isotopes have a different mass number (A), isotopes can be classified into three groups *viz.* light isotopes ($A \leq 50$), medium-weight isotopes ($50 < A < 100$), and heavy isotopes ($A \geq 100$). Over 270 stable isotopes and over 2000 radioactive isotopes of 118 known elements have been found since the term 'isotope' was proposed by F. Soddy¹ in 1910, in which only the elements with $Z \leq 83$ have been found to have stable isotopes.

Generally determined by the nuclear charge, the physical and chemical properties of isotopes are very similar, but not identical due to the distinct neutron number. Therefore, the differences in the properties of isotopes of an element or molecules substituted with sister isotopes (*i.e.*, isotopomers) are called isotope effects (IE).² According to the source of the isotope effect, it can be divided into the mass-dependent isotope effect (MDE) caused by the different masses and mass-independent isotope effect (MIE, also the anomalous isotope effect) caused by other nuclei properties like spin and volume.³

Isotope effect can also be categorized into equilibrium isotope effect (EIE) and kinetic isotope effect (KIE) according to the effect on the chemical behavior.⁴ EIE is the distribution difference of isotopes presented in different phases or chemical states at the equilibrium state and also the basis of chemical

isotope separation. In addition, KIE is the kinetic difference when any atom of the reactant molecules is substituted with its sister isotopes, which is generally accepted as a consequence of the different minimum vibrational energy (*i.e.*, zero-point energy) affected by different masses of isotopic isomers.

As a ubiquitous effect in chemical reactions, KIE is broadly utilized for studying mechanisms,^{4,5} preparing isotope reagents,⁶ controlling the reaction products,^{7,8} and so on. Most researchers in the field of KIE have focused on primary kinetic isotope effects (PKIE) and secondary kinetic isotope effects (SKIE) commonly expressed by the ratio of rate constants of light to heavy isotopomers, k_L/k_H .⁹ The value of PKIE is usually between 2 to 9, but in some special systems, it can be way more than 9 or less than 1 (inverse isotope effect). To date, two theories have been employed to explain the different values of KIE: one is the zero-point energy (ZPE), and the other is the quantum tunneling effect. Neither of them can explain all of the experiment results but only a part which means they all need to be further improved.⁹

Several studies have examined the difference of chemical properties for isotopes *via* the KIE method: (1) a large KIE effect ($k_H/k_D = 40.4$) was measured by U. Weisflog *et al.* at 157 °C with the compound without additional alkyl groups, $\text{C}_6\text{H}_5\text{-C}^2\text{H}_2\text{-O-C}^2\text{H}_2\text{-C}_6\text{H}_5$, as oxidation takes place also on the alkyl groups but not deuterated.¹⁰ (2) Huang Gang *et al.*¹¹ from China Academy of Engineering Physics investigated the p - t curves of deuterium and tritium absorption by uranium at 150–300 °C and constant volume system through reaction rate analysis. They obtained the activation energy value of deuterium and tritium absorption as (-42.8 ± 0.3) and (-43.2 ± 1.2) kJ mol⁻¹, respectively. (3) Eric M. Simmons and John F. Hartwig¹² have reviewed the studies of deuterium KIE in C–H bond functionalization by

China Institute of Atomic Energy, Beijing, 102413, China. E-mail: ouyang_yinggen@163.com; xiao_songtao@126.com; guo_anye_ciae@163.com

[†] Songtao Xiao and Guoan Ye contributed equally.



Table 1 Band gap value (E_g) and the catalytic effects of common metal oxide semiconductor materials on AP^a

Metal oxide semiconductor catalysts		E_g eV ⁻¹	$\Delta T_H/^\circ\text{C}$
P-type	NiO	3.7	65
	La ₂ O ₃	2.86	71
	Cu ₂ O	2.02	90
	Co ₃ O ₄	0.9	117
N-type	MnO ₂	0.25	57
	Fe ₂ O ₃	2.34	67
	V ₂ O ₅	2.49	88
	ZnO	3.35	132

^a ΔT_H is the reduction of the peak temperature at high-temperature decomposition process of AP with addition of catalysts, compared with the pure AP.

transition-metal complexes and concluded factors affecting the KIE like multi-step reactions, the emergence, and structure of the transition state. Zhou Yu-jing *et al.*¹³ from Peking University also reviewed the studies as well as applications of KIE and calculated the difference of ZPE *via* symmetric stretching models obtaining the calculating result of the PKIE of ¹H and D around 6.5, which is very close to the experiment result.

The present work has strongly confirmed the existence of differences in chemical characteristics of isotopes and some of the differences in certain systems are significant, although isotopes 'behave similarly' in most chemical systems due to the limitation of the precision of analysis instrument and the lacking of researches on isotope effects. Here, we first develop isotopes as catalysts and have found the significant differences at the chemical level, which can no longer be ignored.

Thermal decomposition of NH₄ClO₄ (ammonium perchlorate, AP) catalyzed by Fe₂O₃ is a classical catalytic system¹⁴ that has been thoroughly studied and thus this system was selected to investigate the association between isotopes and catalytic ability. The band gap value and catalytic activity described by ΔT_H of different catalytic materials from other public research are listed in Table 1.¹⁵ The data reveals an apparent law that the higher the E_g is, the better the catalytic performance is for Fe₂O₃ and other N-type semiconductor catalysts, while for the P-type the worse the catalytic performance is.

The commonly purchased Fe₂O₃ is actually a mixture of different Fe isotopic oxides consisted of 5.8% ⁵⁴Fe₂O₃, 91.72% ⁵⁶Fe₂O₃, 2.1% ⁵⁷Fe₂O₃, and 0.28% ⁵⁸Fe₂O₃, which means the catalytic performance of Fe₂O₃ on the thermal decomposition of AP is a synthetical result of all isotopic compounds. Here, *n*Fe₂O₃ (abbreviation for Fe₂O₃ in natural abundance ratio), ⁵⁴Fe₂O₃, ⁵⁶Fe₂O₃ and ⁵⁸Fe₂O₃ were transformed into the alpha-crystalline form through a same hydrothermal approach,^{16,17} and the differences in catalytic performance of these Fe-isotopomers were investigated by TG-DSC methods.

Results and discussion

Characterization results

The typical SEM images of the four kinds of α -Fe₂O₃ catalysts obtained *via* the hydrothermal procedure with different

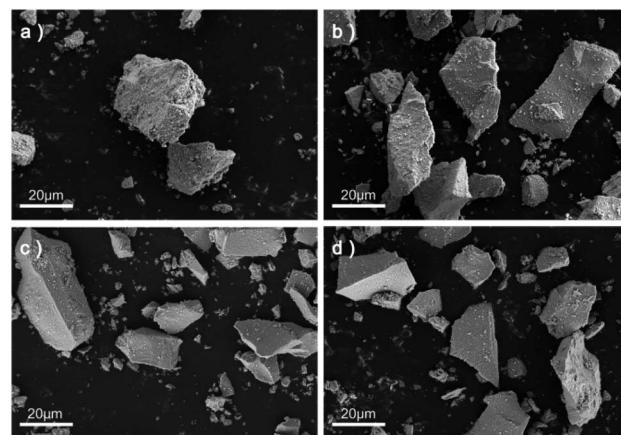


Fig. 1 SEM images of (a) α -*n*Fe₂O₃, (b) α -⁵⁴Fe₂O₃, (c) α -⁵⁶Fe₂O₃ and (d) α -⁵⁸Fe₂O₃.

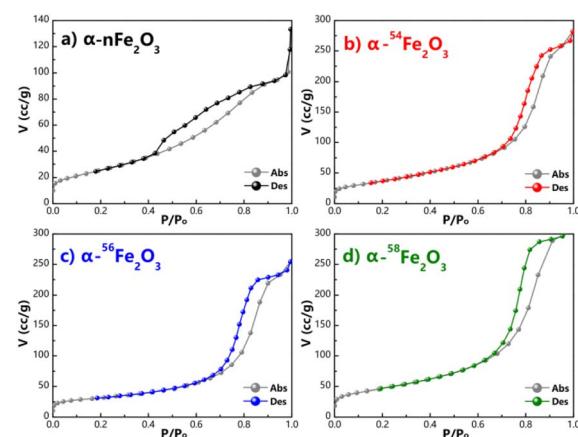


Fig. 2 Adsorption and desorption isotherms of N₂ of (a) α -*n*Fe₂O₃, (b) α -⁵⁴Fe₂O₃, (c) α -⁵⁶Fe₂O₃, and (d) α -⁵⁸Fe₂O₃.

contents of Fe isotopes are shown in Fig. 1. It can be clearly seen from Fig. 1 that the particle shape of the catalysts are all irregular lump and the surface of the α -*n*Fe₂O₃ is rougher with more pores and gullies on the surface compared with separated isotopic α -Fe₂O₃.

The adsorption and desorption isotherms of nitrogen at -196°C obtained from four α -Fe₂O₃ catalysts are presented in Fig. 2. The four isotherms are identified as Type IV isotherms according to IUPAC (International Union of Pure and Applied Chemistry) classification because of the inflection point and the hysteresis loop, which indicating the existence of abundant mesopores in the catalysts. The hysteresis loops could be further divided into H2(b) as the α -*n*Fe₂O₃ and H2(a) as other three separated isotopic α -Fe₂O₃ indicating that the α -*n*Fe₂O₃ has much larger size distribution of neck widths of pore-blocking structure. The BET surface area and pore size data are listed in Table 2. The smaller average pore diameter and pore volume of α -*n*Fe₂O₃ and the rougher surface support this result, which also explain why the surface area of α -*n*Fe₂O₃ is about 14% smaller than that of separated isotopic α -Fe₂O₃.



Table 2 BET surface area and pore size of four α -Fe₂O₃ catalysts

Catalysts	Surface area/(m ² g ⁻¹)	Average pore diameter/nm	Pore volume/(cm ³ g ⁻¹)
α -nFe ₂ O ₃	120.29	3.91	0.211
α - ⁵⁴ Fe ₂ O ₃	144.64	4.80	0.478
α - ⁵⁶ Fe ₂ O ₃	144.01	5.16	0.432
α - ⁵⁸ Fe ₂ O ₃	142.82	4.79	0.516

Based on the electron microscope images and the results of N₂ adsorption and desorption isotherms, the morphology of four α -Fe₂O₃ catalysts is relatively similar that they are all bulk particle with abundant mesopores. The surface area of α -nFe₂O₃ (120.29 m² g⁻¹) is around 14% smaller than those of α -⁵⁴Fe₂O₃ (144.64 m² g⁻¹), α -⁵⁶Fe₂O₃ (144.01 m² g⁻¹), and α -⁵⁸Fe₂O₃ (142.82 m² g⁻¹) because pore-blocking of α -nFe₂O₃ causing smaller pore diameter and volume than those of the separated isotopic α -Fe₂O₃ (around 5 nm and cm³ g⁻¹).

High-resolution transmission electron microscopy (HRTEM) images presented in Fig. 3a-d show the typical crystal plane

spacing of the crystal plane like (110), (104) and (012). The particle distribution obtained from TEM images shown in Fig. 3e-h reveals the lumps in Fig. 1 are agglomeration of particles with around 5 nm average diameter.

It can be seen from Fig. 4a that the diffraction peaks in the XRD pattern of all four α -Fe₂O₃ catalysts are fit for the ICDD (International Center for Diffraction Data) No. 33-0664 of α -Fe₂O₃ with the $a = b = 0.50356$ nm and $c = 1.37489$ nm. The diffraction peaks indexed to the other phases are not observed in Fig. 4a. The diffraction peaks located in $2\theta = 24.1^\circ$, 33.2° , 35.6° , 40.9° , 49.5° , 54.1° , 57.6° , 62.4° , and 64° are corresponding to the (012), (104), (110), (113), (024), (116), (018), (214), and (300) facets. HRTEM images also show the crystal plane spacing of the above crystal plane.

The FTIR spectra of prepared four α -Fe₂O₃ catalysts basically shows no difference in the peaks as presented in Fig. 4b. The peaks around 560 and 430 cm⁻¹ are distinct for the vibrations of Fe-O from α -Fe₂O₃ which means the samples are all in the high-purity of α -Fe₂O₃ phase. The peaks around 3000 and 1530 cm⁻¹ originate from the absorbed water molecules caused by the

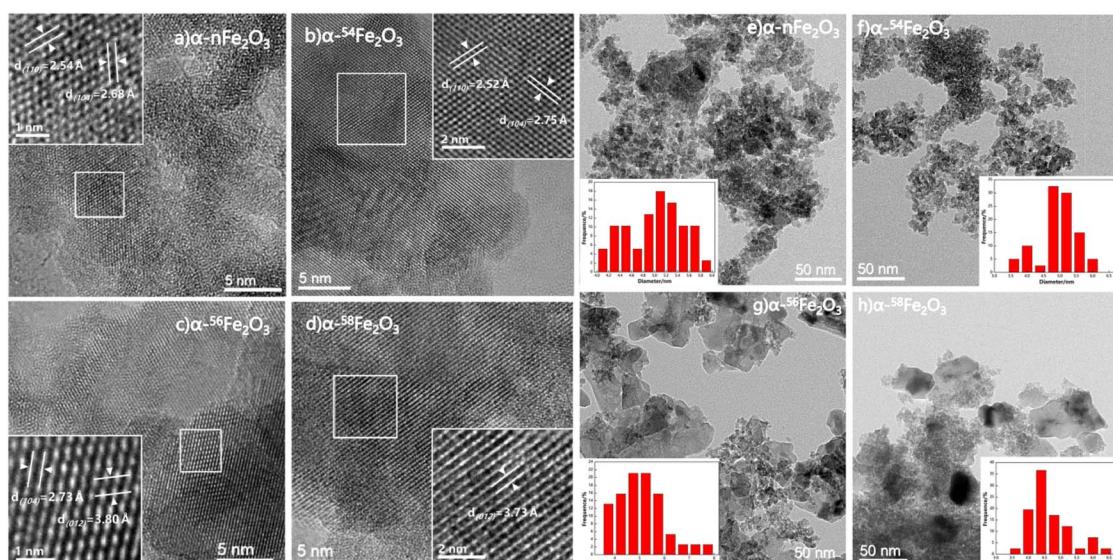


Fig. 3 HRTEM images of synthesized catalysts (a) α -nFe₂O₃, (b) α -⁵⁴Fe₂O₃, (c) α -⁵⁶Fe₂O₃ and (d) α -⁵⁸Fe₂O₃, and particle size distributions of synthesized catalysts (e) α -nFe₂O₃, (f) α -⁵⁴Fe₂O₃, (g) α -⁵⁶Fe₂O₃ and (h) α -⁵⁸Fe₂O₃.

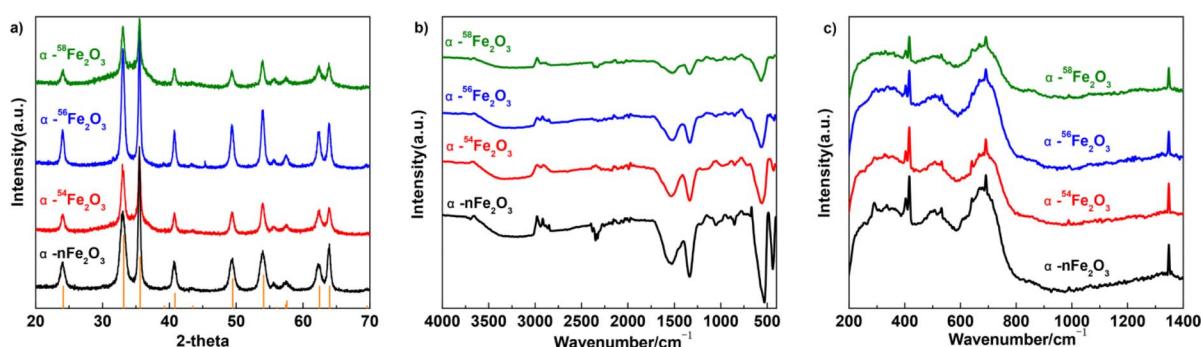


Fig. 4 The (a) XRD patterns, (b) FTIR spectra, and (c) Raman spectra of four α -Fe₂O₃ catalysts.



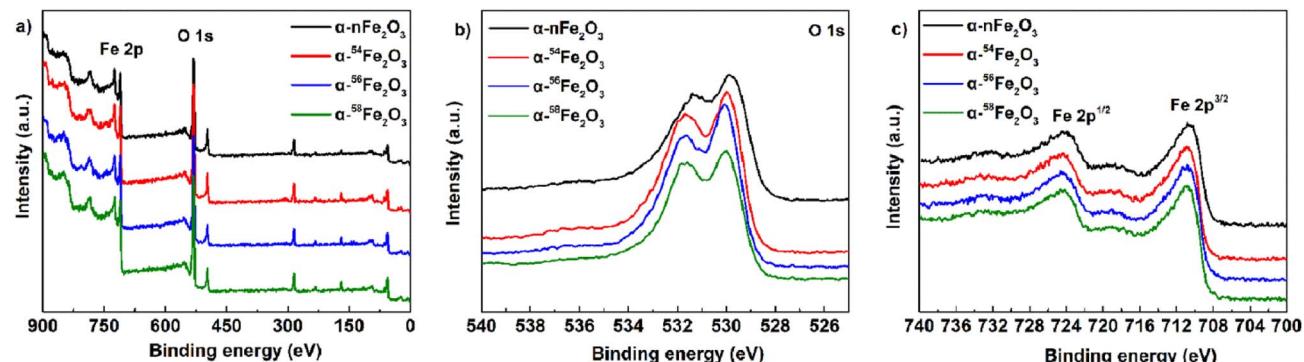


Fig. 5 XPS spectra of four α -Fe₂O₃ catalysts: (a) survey spectrum, (b) O 1s, and (c) Fe 2p.

Table 3 Peak positions of the XPS O 1s peak and Fe 2p peak of four α -Fe₂O₃ catalysts

Compound	Peak position of O 1s/eV		Peak position of Fe 2p/eV		
	O-H	Fe-O	Fe 2p _{1/2}	Satellite	Fe 2p _{3/2}
α -nFe ₂ O ₃	531.4	529.9	724.4	720.0	710.8
α - ⁵⁴ Fe ₂ O ₃	531.7	530.0	724.4	720.4	710.9
α - ⁵⁶ Fe ₂ O ₃	531.6	530.1	724.4	720.0	710.9
α - ⁵⁸ Fe ₂ O ₃	531.6	530.0	724.4	719.9	710.9

potassium bromide pellet technique, and peaks around 1340 cm⁻¹ originate from residual impurities.¹⁸

From Fig. 4c, the four kinds of α -Fe₂O₃ catalysts have similar peaks in the Raman spectra while the peaks located at 403 cm⁻¹, 669 cm⁻¹, and 1348 cm⁻¹ are related to the A_{1g} peak of α -Fe₂O₃, and the peaks at around 416 cm⁻¹, 532 cm⁻¹, 691 cm⁻¹ are related to the E_{1g} of α -Fe₂O₃. There's no obvious difference between the four α -Fe₂O₃ catalysts with different Fe isotopes.¹⁹

To determine the surface chemical states of four α -Fe₂O₃ catalysts, the XPS spectra was measured and displayed in Fig. 5. Fig. 5a is the survey spectrum that confirms the presence of Fe and O elements located at around 720 eV and 530 eV, respectively. For the high-resolution O 1s spectra in Fig. 5b, the peak at 531.6 eV is attributed to the absorbed water molecules, and the peak at 530.0 eV is the lattice oxygen of α -Fe₂O₃. All the catalysts show two main peaks of Fe 2p_{3/2} at 710.9 eV and Fe 2p_{1/2} at

724.4 eV from Fig. 5c. The peak positions of the XPS O 1s and Fe 2p peak are listed in Table 3. The major compound α -Fe₂O₃ in the prepared catalysts was further demonstrated by all the characteristic peaks mentioned above.²⁰

As presented in Fig. 6a the UV-Vis diffuse reflectance spectra of four separated isotopic α -Fe₂O₃ catalysts showing the variation of absorption coefficient A, as a function of photon energy. The A values for all catalysts decreased with the wavelength increasing around 570 nm. The optical band gap was extracted from the UV-Vis diffuse reflectance spectra according to the Tauc plot method,^{21,22}

$$(Ahv)^2 = B(hv - E_g) \quad (1)$$

where A is the absorption coefficient mentioned above, B is a constant relating to the material, E_g is the optical band gap, h is Planck constant and ν is the frequency of the photon. From the Fig. 6b, the band gap values of four α -Fe₂O₃ catalysts are: α -

Table 4 Band gap values (E_g) and catalytic results of α -Fe₂O₃ on the thermal decomposition of AP

Compound	Endothermal peak		Exothermal peak		E _g /eV
	T _{endo} /°C	RSD	T _{exo} /°C	RSD	
AP	242.50	0.28%	439.76	0.21%	—
α -nFe ₂ O ₃ /AP	242.59	0.08%	341.34	0.47%	1.97
α - ⁵⁴ Fe ₂ O ₃ /AP	243.36	0.11%	341.98	0.32%	1.87
α - ⁵⁶ Fe ₂ O ₃ /AP	243.20	0.15%	346.25	0.32%	1.73
α - ⁵⁸ Fe ₂ O ₃ /AP	243.95	0.11%	344.34	0.16%	1.86

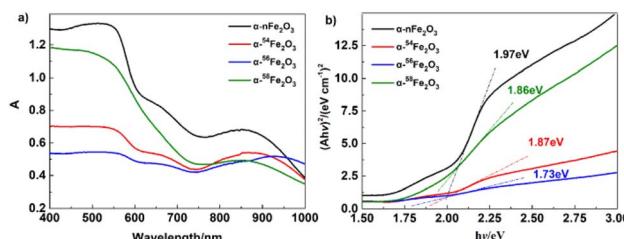


Fig. 6 (a) UV-Vis diffuse reflectance spectra, and (b) (Ahv)² - hν curves of α -Fe₂O₃ catalysts.

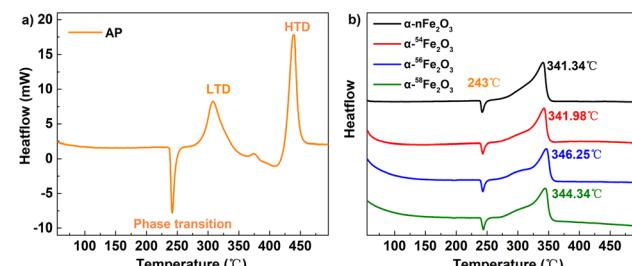


Fig. 7 The DSC curves of (a) pure AP and (b) AP catalyzed by α -Fe₂O₃.



$n\text{Fe}_2\text{O}_3$ (1.97 eV) > $\alpha\text{-}^{54}\text{Fe}_2\text{O}_3$ (1.87 eV) $\gtrsim \alpha\text{-}^{58}\text{Fe}_2\text{O}_3$ (1.86 eV) > $\alpha\text{-}^{56}\text{Fe}_2\text{O}_3$ (1.73 eV) as shown in Table 4.

Catalytic performance

The TG-DSC results of pure AP and AP in presence of $\alpha\text{-Fe}_2\text{O}_3$ are listed in Table 4. Fig. 7a shows the three processes of thermal decomposition of pure AP: (1) phase transition corresponding to the negative peak in DSC curves. AP is transforming into the cubic crystal from the orthorhombic crystal without weight loss. (2) The low-temperature decomposition (LTD) corresponding to the first exothermic peak (310 °C) is related to the partial decomposition of AP to form NH_3 and HClO_4 intermediates by dissociation and sublimation. And (3) the high-temperature decomposition (HTD) corresponding to the last peak (440 °C) related to the complete decomposition of the intermediates to N_2O , NO , H_2O , HCl , Cl_2 , etc. products.²³

It's obvious from Fig. 7b that the HTD peaks of the separated isotopic $\alpha\text{-}n\text{Fe}_2\text{O}_3$ shift to the lower temperature direction much more than that of $\alpha\text{-Fe}_2\text{O}_3$ shifts, indicating the existence of remarkable differences in catalytic performance between separated isotopic $\alpha\text{-Fe}_2\text{O}_3$ and $\alpha\text{-}n\text{Fe}_2\text{O}_3$. From Table 4 we can further see that the HTD peak exothermal temperatures of four kinds of $\alpha\text{-Fe}_2\text{O}_3/\text{AP}$ show significant differences which reveals that the catalytic performance is affected by the Fe isotopes in $\alpha\text{-Fe}_2\text{O}_3$. The catalytic performance order from the best to the least of the four $\alpha\text{-Fe}_2\text{O}_3$ catalysts is $\alpha\text{-}n\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{54}\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{58}\text{Fe}_2\text{O}_3$, and $\alpha\text{-}^{56}\text{Fe}_2\text{O}_3$. The catalytic performance positively correlates with the optical band gap, which follows the rules found in experiment results of thermal decomposition of AP catalyzed by N-type metal oxide semiconductor materials.²²

According to the electron-transfer theory,²⁴ the partially filled 3d orbit in $\alpha\text{-Fe}_2\text{O}_3$ catalysts promotes the LTD process by providing more pathways for electrons and intermediate products. Then the gases arising from LTD process would be absorbed on the surface of catalysts enhancing the HTD process. The mechanism of thermal decomposition process of AP/ $\alpha\text{-Fe}_2\text{O}_3$ is illustrated in Fig. 8.

Although the surface area of $\alpha\text{-}n\text{Fe}_2\text{O}_3$ is the smallest, it still shows the best catalytic performance which is not in accordance with the rule that larger surface area leads to better catalytic

performance. We infer that the difference of catalytic activity in $\alpha\text{-}n\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{54}\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{56}\text{Fe}_2\text{O}_3$ and $\alpha\text{-}^{58}\text{Fe}_2\text{O}_3$ is due to the different Fe isotope content.

Experimental

Materials

$^{54}\text{Fe}_2\text{O}_3$ ($\geq 99.88\%$), $^{56}\text{Fe}_2\text{O}_3$ ($\geq 99.7\%$) and $^{58}\text{Fe}_2\text{O}_3$ ($\geq 93.13\%$) were purchased from Trace Sciences International. Fe_2O_3 ($\geq 99.95\%$), NaOH solid, concentrated hydrochloric acid, concentrated nitric acid and NH_4ClO_4 ($\geq 99.5\%$) were purchased from Aladdin Industrial Corporation. All the above reagents are analytically pure and used directly as received without further purification.

Preparation and characterization

Four kinds of $\alpha\text{-Fe}_2\text{O}_3$ catalysts, *i.e.*, $\alpha\text{-}n\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{54}\text{Fe}_2\text{O}_3$, $\alpha\text{-}^{56}\text{Fe}_2\text{O}_3$, and $\alpha\text{-}^{58}\text{Fe}_2\text{O}_3$ were all prepared through the typical hydrothermal procedure from the four kinds of purchased Fe_2O_3 , respectively. The mixture of 0.2 g Fe_2O_3 and 8 mL aqua regia was sealed into a Teflon-lined stainless-steel autoclave maintained at 100 °C for 3 hours and then cooled to room temperature naturally. Adding 8 mL deionized water into the solution and then adding 5 M NaOH solutions drop by drop with magnetic stirring until the pH of the supernatant arrived at 6–7. The obtained precipitate was collected by centrifugation and washed 3 times with deionized water as well as absolute ethanol. The precipitate after being dried through a vacuum dryer for 8 h at 60 °C was roasted at 400 for 2 h in the Muffle furnace to get the $\alpha\text{-Fe}_2\text{O}_3$ catalysts.

The form of the catalysts was investigated by scanning electron microscopy (SEM) using a GeminiSEM300. The specific surface area was measured by the Brunner–Emmett–Teller (BET) measurements on an SSA-7000 analyzer. Powder X-ray diffraction (XRD) analyses were characterized by Bruker D8 Advance equipped with $\text{Cu K}\alpha$ radiation at 14° min^{-1} , and the high-resolution transmission electron microscopy (HRTEM) measurements were obtained by FEI Tecnai G2 F30 for a comprehensive understanding of the crystal structure. Fourier transform infrared spectra (FTIR, Nicolet Is 10) were measured from 4000 to 400 cm^{-1} using the potassium bromide pellet method. X-ray photoelectron spectroscopy (XPS) was carried out on Escalab 250Xi.

Catalytic activity

AP and four kinds of $\alpha\text{-Fe}_2\text{O}_3$ catalysts were grounded for about 20 min in an agate mortar with absolute ethanol and further dispersed by ultrasonic wave for 10 min. The mixtures were dried through a vacuum dryer for 12 h at 60 °C and then ground to get the evenly dispersed $\alpha\text{-Fe}_2\text{O}_3/\text{AP}$ sample with the stable 5% mole fraction of the catalyst.

Under the condition of an N_2 gas flow rate of 10 mL min^{-1} , thermal decomposition processes were investigated by the TG-DSC method (TGA/DSC 3+ Simultaneous Thermal Analyzer from METTLER TOLEDO) at a heating rate of 10 °C min^{-1} in the temperature range from 50–500 °C.

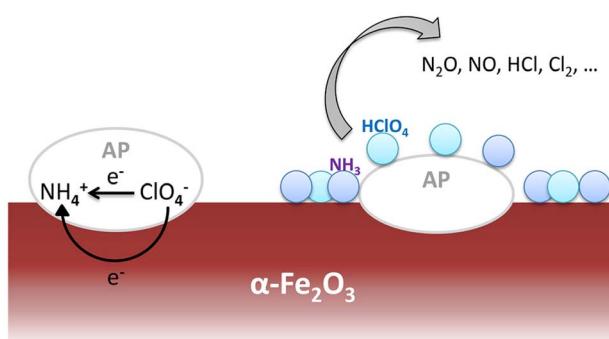


Fig. 8 Schematic of the thermal decomposition process of AP on the $\alpha\text{-Fe}_2\text{O}_3$ catalytic surface.



Conclusions

Four kinds of α -Fe₂O₃ catalysts (α -nFe₂O₃, α -⁵⁴Fe₂O₃, α -⁵⁶Fe₂O₃, and α -⁵⁸Fe₂O₃) were prepared through same hydrothermal procedure to investigate the existence of the difference of catalytic performances of the isotopic catalysts. The characterization results of α -Fe₂O₃ present the uniformity of physical morphology among the α -Fe₂O₃ in natural abundance ratio and the separated isotopic α -Fe₂O₃ with the similar SEM and HRTEM images, adsorption and desorption isotherms of N₂, XRD patterns, FTIR and Raman spectra. While the surface area of α -nFe₂O₃ is about 14% smaller than those of other three separated isotopic α -Fe₂O₃ because of much larger size distribution of neck widths of pore-blocking for the α -nFe₂O₃.

The TG-DSC thermal analysis results of thermal decomposition of AP in presence of α -Fe₂O₃ confirm the existence of differences in the catalytic performance of the isotopic catalysts. The catalytic performance is affected by the contents of Fe isotopes in α -Fe₂O₃ and the order of it is: α -nFe₂O₃ < α -⁵⁴Fe₂O₃ < α -⁵⁸Fe₂O₃ < α -⁵⁶Fe₂O₃ in line with the order of band gap obtained by UV-Vis DRS, which suggests that the catalytic performance of α -Fe₂O₃ catalysts with different contents of Fe isotopes positively correlated with the band gap value but they are not strictly proportional.

Although the surface area of α -nFe₂O₃ is the smallest, it still shows the best catalytic performance, so we infer that the difference in their catalytic activity in AP decomposition is mainly attributed to the difference in the charge distribution of the nuclei of different iron isotopes. The s-orbital is spherically symmetric when the nuclear charge is regarded as a point charge, but it's not a point charge under the real condition. The difference in the number of neutrons in isotopes leads to the difference in the nuclear forms of different atoms of the same element, and the distribution of the nuclear charge also varies.²⁵ The distribution difference of nuclear charge with the change of nuclear form makes the nuclear field an equipotential surface consistent with the distribution of nuclear charge. Then for the s-orbital electrons, for example, instead of having a spherical symmetry distribution, they have a probability distribution along the equipotential surface of the nuclear field. For the p- and d-orbitals, there exist the same distortion to a certain degree for the same isotopes, but different for different isotopes. When separated isotopes are utilized as materials, the distortion of the same isotope is completely coincident, resulting in a significant cluster effect.²⁶ Based on the theories of isotope effect and the data obtained in this work, we infer that the natural α -Fe₂O₃ is weakened by the overlapping of distorted orbitals, while the separated isotopic α -Fe₂O₃ shows a remarkable cluster effect because of the overlapping of distorted orbitals, which would reflect in the band gap energy, that is, the band gap energy of oxides of natural elements is larger than that of oxides of separated isotopes as the characterization results. The band gap energies of different separated isotopes are apparently distinct due to the different structures of orbital distortion. In this work, band gap energy is a clue to detect the catalytic difference between isotopic catalysts. We also expect to

find out other parameters or methods to more directly as well as precisely reflect the difference.

All the experimental results indicate that the difference of the energy state of isotopomers when utilizing as catalysts is big enough to be taken into consideration at the chemical level. The law of it is not clear at present but the discovery of the catalytic isotope effect in this system is meaningful. Further studies on the difference of catalytic activity between separated isotopic and natural abundance oxides would not only explore the influence of nuclear charge distribution on the molecular states of extra nuclear electrons but also be possible to obtain a new kind of catalyst, namely isotopic catalyst.

Author contributions

Songtao Xiao, the secondary corresponding author (Songtao Xiao and Guoan Ye contributed equally to this work), has made substantial contributions to the design of the work and been visualizing during the whole work. He also has drafted the work. Yubing Xue has also drafted the work, made contributions to the acquisition and interpretation of data, and substantively revised the manuscript. Jing Zhao contributed the project administration. Xiechun Liu contributed the formal analysis of data. Haifeng Cong contributed the data curation and acquisition. Tian Lan has been providing the necessary resources. Dashu Pan contributed the analysis of the data. Lingyu Wang contributed resources and data curation. Xiaojuan Liu contributed the review and editing work. Guoan Ye, the secondary corresponding author (Songtao Xiao and Guoan Ye contributed equally to this work), contributed the funding acquisition and methodology. Yinggen Ouyang, the first corresponding author, has made contributions to the funding acquisition, conceptualization, and supervision.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 F. Soddy, *J. Chem. Soc. Trans.*, 1911, **99**, 72–83.
- 2 P. Krumbiegel, *Isotopes Environ. Health Stud.*, 2011, **47**, 1–17.
- 3 A. L. Buchachenko, *J. Phys. Chem. B*, 2013, **117**, 2231–2238.
- 4 W. W. Cleland, *Arch. Biochem. Biophys.*, 2005, **433**, 2–12.
- 5 C. Gunnemann, D. W. Bahnemann and P. K. J. Robertson, *ACS Omega*, 2021, **6**, 11113–11121.
- 6 K. Fujiki, Y. Kanayama, S. Yano, *et al.*, *Chem. Sci.*, 2019, **10**, 1936–1944.
- 7 P. E. Yankwich and M. Calvin, *J. Chem. Phys.*, 1949, **17**, 109–110.



8 J. Bigeleisen and M. Wolfsberg, *J. Chem. Phys.*, 1955, **23**, 1535–1539.

9 J. F. Marlier, *Accounts Chem. Res.*, 2001, **34**, 283–290.

10 U. Weisflog, P. Krumbiegel and H. Hübner, *Isot. Environ. Health Stud.*, 1970, **6**, 285–287.

11 H. Gang, L. Xinggui, L. Jianhua, *et al.*, *Rare Met. Mater. Eng.*, 2011, **40**, 2010–2013.

12 E. M. Simmons and J. F. Hartwig, *Angew. Chem., Int. Ed.*, 2012, **51**, 3066–3072.

13 Y. Zhou and J. Wang, *Sci. China Chem.*, 2016, **46**, 573–578.

14 S. Chaturvedi and P. N. Dave, *J. Saudi Chem. Soc.*, 2013, **17**, 135–149.

15 B. Shu-hong, C. Si-yu and F. Lu, *Chin. J. Energ. Mater.*, 2021, **29**, 460–470.

16 H. Li, Y. Chen, X. Zhao and S. Deng, *Chin. J. Inorg. Chem.*, 2018, **34**, 1670–1676.

17 F. Wang, X. F. Qin, Y. F. Meng, *et al.*, *Mater. Sci. Semicond. Process.*, 2013, **16**, 802–806.

18 X. Li, S. Li, H. Hu, T. Sun and S. Yang, *Catal. Lett.*, 2022, **152**, 3479–3488.

19 D. T. M. Phan, T. Häger and W. Hofmeister, *J. Raman Spectrosc.*, 2017, **48**, 453–457.

20 T. Yamashita and P. Hayes, *Appl. Surf. Sci.*, 2008, **254**, 2441–2449.

21 T. J. Tauc, R. Grigorovici and A. Vancu, *Phys. Status Solidi*, 1966, **15**, 627–637.

22 C. Laberty and A. Navrotksy, *Geochim. Cosmochim. Acta*, 1988, **62**, 2905–2913.

23 V. V. Boldyrev, *Thermochim. Acta*, 2006, **443**, 1–36.

24 K. Gao, G. Li, Y. Luo, *et al.*, *J. Therm. Anal. Calorim.*, 2014, **118**, 43–49.

25 J. Bigeleisen, *J. Am. Chem. Soc.*, 1966, **118**, 3676–3680.

26 S. Zhang, Q. Liu, M. Tang and Y. Liu, *ACS Earth Space Chem.*, 2020, **4**, 420–433.

