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# Promotion of SO<sub>2</sub> resistance of Ce–La/TiO<sub>2</sub> denitrification catalysts by V doping

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Conventional cerium-based denitrification catalysts show good catalytic activity at moderate and high temperatures, but their denitrification performance may be decreased due to poisoning by SO<sub>2</sub> in the flue gas. In this paper, V was introduced into Ce–La/TiO<sub>2</sub> catalysts by a ball-milling method, and the effects of the V content on catalyst denitrification performance and SO<sub>2</sub> resistance were investigated. Fourier-transform diffuse reflectance *in situ* infrared spectroscopy was used to examine the denitrification mechanism and evaluate the catalysts for surface acidity, redox characteristics, and SO<sub>2</sub> adsorption. After introducing V, Brønsted acids played the dominant role in the catalytic reaction by increasing the number of acidic sites on the catalyst surface, adsorbing NH<sub>3</sub> to participate in the reaction, and improving the sulfur resistance by inhibiting SO<sub>2</sub> poisoning. The Ce<sup>3+</sup> and O ratio on the catalyst surface were also enhanced by V doping, which reduced interactions between SO<sub>2</sub> and the primary metal oxide active ingredients. The modified catalyst inhibited the formation of sulfate species on the catalyst surface and prevented the generation of additional nitrate species on the surface, which protected the main active sites. After V doping, the NH<sub>3</sub>–SCR reaction on the catalyst surface followed the Langmuir–Hinshelwood mechanism.

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## 1 Introduction

NO<sub>x</sub> is a major atmospheric pollutant whose emissions cause acid rain, photochemical smog, and ozone layer depletion.<sup>1</sup> Selective catalytic reduction (SCR) has a wide range of denitrification efficiencies, does not produce secondary contamination due to NH<sub>3</sub> escape, and is widely used in various plants and municipal waste facilities.<sup>2</sup> Cerium-based catalysts are one of the commonly used catalysts in SCR, which is due to the fact that Ce atoms contain a special arrangement of electrons outside the nucleus and have excellent oxidizing properties and oxygen storage capacity. CeO<sub>2</sub> can be converted from Ce<sup>4+</sup> to Ce<sup>3+</sup> when cerium ions generate oxygen vacancies,<sup>3</sup> giving the catalyst excellent redox properties and thermal stability. Cerium promotes the conversion of nitrate species and ammonia species, which facilitates the conversion of NH<sub>3</sub> and NO to reactive intermediates on the catalyst surface, thus promoting the NH<sub>3</sub>–SCR reaction.

Usually, the flue gas in coal-fired boilers contains SO<sub>2</sub>, which inevitably poisons the catalyst and reduces its denitrification performance. The reasons for SO<sub>2</sub> poisoning can be classified

into three categories: (1) the competitive adsorption of SO<sub>2</sub> with reactants inhibits the formation of reactive intermediates; (2) the reaction of SO<sub>2</sub> with NH<sub>3</sub> to generate (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> and NH<sub>4</sub>HSO<sub>4</sub>, which covers active sites on the catalyst surface and reduces the specific surface area of the catalyst; (3) metal active sites are sulfated by SO<sub>2</sub> to create metal sulfates, which results in the loss of the active components and results in irreversible deactivation. For Ce-based catalysts, SO<sub>2</sub> can be adsorbed and oxidized to SO<sub>3</sub> by CeO<sub>2</sub> to generate Ce(SO<sub>4</sub>)<sub>2</sub> or Ce<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub> or react with NH<sub>3</sub> to form stable (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> or NH<sub>4</sub>HSO<sub>4</sub>, the NO conversion of CeO<sub>2</sub>/TiO<sub>2</sub> catalysts was close to 100% above 300 °C and remained stable for 12 h. However, the value decreased significantly after 12 h, which was below 40% after 48 h.<sup>4</sup> For Mn-based catalysts, the NO conversion of fresh samples of MnO<sub>2</sub>-based catalysts was 100% at 130–150 °C. After SO<sub>2</sub> poisoning, the NO conversion decreased sharply from 100% to 53% at 130 °C and did not increase when the reaction temperature was increased to 150 °C.<sup>5</sup> Ma *et al.*<sup>6</sup> prepared Cu<sub>0.02</sub>Fe<sub>0.2</sub>W<sub>0.02</sub>TiO<sub>x</sub> by doping W to Cu<sub>0.02</sub>Fe<sub>0.2</sub>TiO<sub>x</sub> catalysts, the NO conversion of which maintained at 100% after passing water and sulfur at 240 °C for 10 h. Jiang *et al.*<sup>7</sup> prepared Mn–Fe(0.4)–Ce(0.4)/ACN catalysts, which showed a NO<sub>x</sub> conversions higher than 95% at 100–250 °C, and remained around 91% at 175 °C for 32 h under 100 ppm SO<sub>2</sub> and 10 vol% H<sub>2</sub>O.

Early studies on SO<sub>2</sub> poisoning catalysts mostly concentrated on vanadium-based catalysts. In the temperature range of 180 to 400 °C, Li *et al.*'s<sup>8</sup> investigation of the distribution of SO<sub>2</sub>

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oxidation products during the  $\text{NH}_3$ -SCR process over  $\text{V}_2\text{O}_5/\text{TiO}_2$  catalysts discovered that the amount of gaseous  $\text{SO}_3$  products increased with temperature in this range. The presence of  $\text{O}_2$  in flue gas promoted the oxidation of  $\text{SO}_2$  and the formation of gaseous  $\text{SO}_3$  on the  $\text{V}_2\text{O}_5/\text{TiO}_2$  catalysts. On the catalyst surface, the major pathway for  $\text{SO}_2$  oxidation comprises oxidizing  $\text{SO}_2$  to generate adsorbed  $\text{SO}_3$ , which either desorbs to form gaseous  $\text{SO}_3$  or interacts with  $\text{NH}_3$  to form sulfate. Sulfite is created when a tiny amount of  $\text{SO}_2$  interacts with  $\text{NH}_3$ ; this product was then quickly oxidized to  $(\text{NH}_4)_2\text{SO}_4$ . At 180 °C, the amount of deposition products peaked, while below 230 °C, more  $\text{NH}_4^+$  emerged. Additives that also strengthened the water and sulfur resistance of the rare-earth-based catalysts helped them conduct denitrification even better. Many researchers have prepared binary-loaded metal oxide catalysts using transition metals and doping with rare earth elements such as  $\text{CoO}$ ,<sup>9</sup>  $\text{Cr}_2\text{O}_3$ ,<sup>10</sup> and  $\text{Sm}_2\text{O}_3$ .<sup>11</sup> This method boosted the denitrification activity of the original Ce-based catalysts as well as their resistance to water and sulfur.

In this paper, we prepared a Ce-La-V/TiO<sub>2</sub> catalyst *via* a previously reported ball milling method.<sup>12</sup> The effect of the V introduction on the denitrification performance of the catalyst as well as on the water and sulfur resistance was tested. It was found that the NO conversion still maintained 85% after 52 h of water and sulfur feeding. After stopping the water and sulfur,

1.4 and 2.3). In addition, the catalyst was poisoned by passing 300 ppm  $\text{SO}_2$  and 5%  $\text{H}_2\text{O}$  on top of a mixture of 500 ppm NO, 500 ppm  $\text{NH}_3$ , 5%  $\text{H}_2\text{O}$  and  $\text{N}_2$ , which was allowed to stabilize and then reacted continuously for 4 h to obtain a poisoned sample, which was labeled as 10Ce-2La-*x*V/Ti-S.

## 2.2 Catalysts activity test

First, the prepared catalysts were ball-milled and attached to porous cordierite cylinders (commercial grade,  $\phi$  20 × *L* 50). After drying in an oven, the catalyst-loaded cordierite cylinders were placed in a quartz tube furnace to test the catalytic activity of the catalyst. The catalyst was loaded with a weight of 1 g. The gas mixture consisted of 500 ppm NO, 500 ppm  $\text{NH}_3$ , 300 ppm  $\text{SO}_2$ , 5%  $\text{H}_2\text{O}$ , and 3%  $\text{O}_2$ , with  $\text{N}_2$  as the equilibrium gas. The total gas flow rate was 2500 mL min<sup>−1</sup> and the corresponding GHSV was 150 000 mL g<sup>−1</sup> h<sup>−1</sup>. The flow of all the gases was controlled by a mass flowmeter. NO and NO<sub>2</sub> concentrations at the outlet were monitored in real time using an ECOM flue gas analyzer (Germany). NO conversion, N<sub>2</sub> selectivity and SO<sub>2</sub> conversion were calculated from eqn (1)–(3):

$$\text{NO conversion (\%)} = \frac{[\text{NO}]_{\text{in}} - [\text{NO}]_{\text{out}}}{[\text{NO}]_{\text{in}}} \times 100\% \quad (1)$$

$$\text{N}_2 \text{ selectivity (\%)} = \frac{[\text{NO}]_{\text{in}} + [\text{NH}_3]_{\text{in}} - [\text{NO}_2]_{\text{out}} - 2[\text{N}_2\text{O}]_{\text{out}} - [\text{NO}]_{\text{out}} - [\text{NH}_3]_{\text{out}}}{[\text{NO}]_{\text{in}} + [\text{NH}_3]_{\text{in}} - [\text{NO}]_{\text{out}} - [\text{NH}_3]_{\text{out}}} \times 100\% \quad (2)$$

the NO conversion recovered to 99%. In addition, the mechanism of the resistance to  $\text{SO}_2$  toxicity at high temperatures was elucidated using various physicochemical characterization techniques and reaction kinetics analysis.

## 2 Experimental

### 2.1 Catalysts preparation

According to the chemical formula of  $\text{Ce}_{10}\text{La}_2\text{V}_x\text{Ti}_{88-x}$  (where *x* is the mass ratio of  $\text{V}_2\text{O}_5$  per 100 parts of catalyst), cerium carbonate, lanthanum carbonate, ammonium metavanadate and titanium dioxide were added and put in the nylon ball milling jar, appropriate amount of deionized water and zirconia balls were added. The volume of the ball milling jar was 530 mL, the inner diameter was 94.5 mm, the height was 93.5 mm, the weight of zirconia balls was 95 g, and the volume of water was 100 mL. Then the slurry was ball milled in QM-Z/B planetary ball mill, the rotational speed and time were set (usually 500 rpm, 1 h), and the direction of rotation was changed once per 20 min, separation of the slurry from the zirconia balls at the end of the process, and the slurry was dried in an oven at 105 °C for 12 h. After drying, the slurry was calcined at 500 °C for 4 h to obtain the desired catalysts, and the corresponding catalysts were notated as 10Ce-2La-*x*V/Ti (*x* = 0.1, 0.3, 0.6, 0.9,

$$\text{SO}_2 \text{ conversion (\%)} = \frac{[\text{SO}_2]_{\text{in}} - [\text{SO}_2]_{\text{out}}}{[\text{SO}_2]_{\text{in}}} \times 100\% \quad (3)$$

where the subscript 'in' and 'out' represent the inlet and outlet concentrations of each gas at steady state, respectively.

In order to visualize the differences in the catalytic activity of the samples more intuitively, the tests were conducted at a higher GHSV (150 000 mL g<sup>−1</sup> h<sup>−1</sup>), keeping the NO conversion rate lower than 30%, and the surface activation energies of the different catalysts were calculated. Assuming that the  $\text{NH}_3$ -SCR catalytic reaction is not controlled by diffusion, the SCR reaction rate normalized by the specific surface area of the catalyst can be calculated according to eqn (4)<sup>13,14</sup>

$$\text{Rate (mol m}^{-2} \text{ s}^{-1}) = \frac{X_{\text{NO}} \times Q \times C_f}{V_m \times W \times S_{\text{BET}}} \quad (4)$$

where  $X_{\text{NO}}$  is the NO conversion (%) at different temperatures, *Q* is the volumetric flow (2500 mL min<sup>−1</sup>),  $V_m$  is the molar volume of gas under standard conditions (22.4 L mol<sup>−1</sup>),  $C_f$  is the fed NO concentration (500 ppm), *W* is the weight of the catalyst (g), and  $S_{\text{BET}}$  is the specific surface area of the catalyst (m<sup>2</sup> g<sup>−1</sup>).

Turn over frequency (TOF) can be calculated to compare the catalytic rate of different catalysts. Keeping the NO conversion below 30% (to ensure that all catalytically active sites are



operational), the TOF value of NO on the active center Ce is calculated from eqn (5):<sup>15,16</sup>

$$\text{TOF} = \frac{P_V/RT\alpha}{m_{\text{cat}}\beta_{\text{Ce}}/M_{\text{Ce}}} \quad (5)$$

where  $P$  is the standard atmospheric pressure ( $1.01 \times 10^5$  Pa),  $v$  is the flow of NO,  $R$  is the molar gas constant ( $8.314 \text{ J mol}^{-1} \text{ K}^{-1}$ ),  $T$  is the test temperature (K),  $\alpha$  is the NO conversion (%),  $m_{\text{cat}}$  is the weight of the catalyst (g),  $\beta_{\text{Ce}}$  is the loading ratio of Ce (%) calculated using XPS data, and  $M_{\text{Ce}}$  is the molecular weight of cerium ( $140.1 \text{ g mol}^{-1}$ ).

### 2.3 Characterization of catalysts

The physical composition of the samples was examined using a Thermo ARL SCINTAG X'TRA X-ray diffraction analyzer. Cu K- $\alpha$  rays (wavelength 0.154056 nm) were used as its X-ray source, with a tube voltage of 40 kV and a tube current of 40 mA. Sample surface area, pore volume and pore size were measured by  $\text{N}_2$  adsorption and desorption using a Biaoode SSA-7300 automatic pore size and surface area analyzer. X-ray photoelectron spectroscopy (XPS) was carried out using a Thermo Scientific K-Alpha electron spectrometer equipped with an Al K $\alpha$  ( $h\nu = 1486.6 \text{ eV}$ ) radiation at an operating voltage of 12 kV and a reference pressure of  $3 \times 10^{-7}$  mbar. Temperature programmed desorption/reduction (TPD/TPR) experiments were carried out on a chemisorption apparatus (Micromeritics, AutoChem II 2920). TG and DSC quantities were measured using a simultaneous thermal analyzer (PerkinElmer STA-8000).  $\text{N}_2$  was used as the carrier gas, and the samples were heated up to  $100^\circ\text{C}$  at a heating rate of  $10^\circ\text{C min}^{-1}$  to pretreat moisture and other impurities on their surface, and then heated up to  $800^\circ\text{C}$  at a heating rate of  $10^\circ\text{C min}^{-1}$ . The *in situ* IR tests were performed on an *in situ* FTIR spectrometer (Tensor 27). The tests were consisted by four parts:  $\text{NH}_3$  adsorption,  $\text{NO} + \text{O}_2$  adsorption,  $\text{NO} + \text{O}_2$  reaction with pre-adsorbed  $\text{NH}_3$  and  $\text{NH}_3$  reaction with pre-adsorbed  $\text{NO} + \text{O}_2$ .

## 3 Results and discussion

### 3.1 Characterization of catalysts physical properties

**3.1.1 XRD.** The physical phases present in the catalysts were analyzed by X-ray diffraction. As shown in Fig. 1, all samples contained peaks near  $25.2^\circ$ ,  $37.7^\circ$ ,  $48.0^\circ$ ,  $53.9^\circ$ , and  $54.9^\circ$ , which belonged to anatase  $\text{TiO}_2$  (JCPDS no. 21-1272). In addition, weak diffraction peaks of  $\text{CeO}_2$  (JCPDS no. 43-1002) were detected near  $28.6^\circ$ ,  $33.0^\circ$ , which indicated that some  $\text{CeO}_2$  was present on the catalyst surface.  $\text{La}_2\text{O}_3$  and  $\text{V}_2\text{O}_5$  diffraction peaks were not observed for all samples, indicating that they are poorly crystallized and highly dispersed on the surface of the catalysts, which allows the active components to interact more fully and thus enhance the catalytic activity.<sup>17</sup> The absence of neither crystallographic changes nor the appearance of new diffraction peaks in the poisoned samples compared with the fresh samples may be due to the low content of sulfate species generated on the catalyst surface or that they were amorphous.

Fig. 2 shows the  $\text{N}_2$  adsorption-desorption isotherms and BJH pore size distribution. As shown in Fig. 2(a), all samples

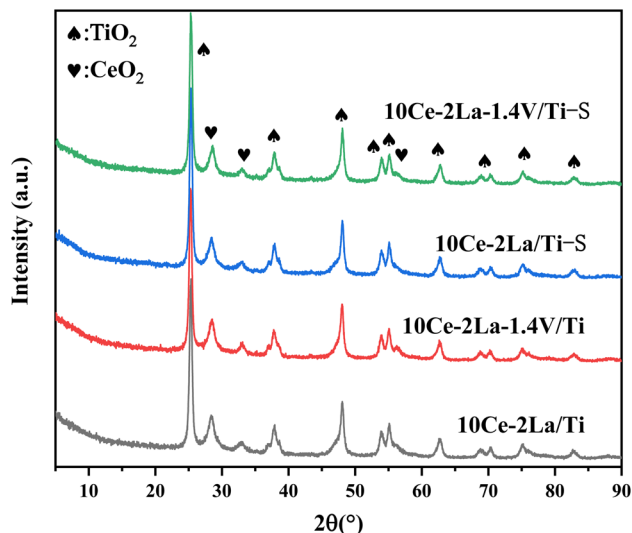


Fig. 1 XRD spectra of 10Ce-2La- $x$ V/Ti series catalysts before and after poisoning.

showed typical H3 hysteresis loops at higher relative pressures in the latter half ( $P/P_0 = 0.7-1.0$ ), which is usually associated with capillary coalescence in the catalyst.<sup>18</sup> The curves were a type IV isotherm according to the IUPAC classification, indicating that the catalysts had mesoporous structures. From Fig. 2(b), it is clear that all samples showed a wide range of pore sizes, further confirming that the catalysts are mesoporous and are made up of a variety of irregularly shaped particles stacked together. In addition, the specific surface area, total pore volume, and average pore size obtained from  $\text{N}_2$  adsorption-desorption isotherms were calculated (Table 1). These values were slightly reduced after adding V because it favored sintering. After poisoning, the catalyst's surface area, average pore volume, and total pore volume were drastically decreased, demonstrating that sulfate species had coated the catalyst's surface.<sup>19</sup>

**3.1.2 TEM.** Fig. 3 shows the TEM images, HR-TEM images and elemental distributions of Ce, La, Ti and V for 10Ce-2La/Ti and 10Ce-2La-1.4V/Ti. From Fig. 3(a) and (c), it can be seen that the catalysts are porous structures formed by the stacking of nanoparticles with uneven sizes, which can exhibit good catalytic activity. In the HR-TEM images in Fig. 3(b) and (d), the crystal spacing of 0.35 nm corresponds to the (101) facets of anatase  $\text{TiO}_2$ , 0.19 nm corresponds to the (220) crystalline facets of  $\text{CeO}_2$ , and 0.17 nm can be indexed to the (202) crystalline facets of  $\text{TiO}_2$ . Lattice fringes of lanthanum oxide and vanadium oxides were not found in any of the catalysts, confirming that they existed in an amorphous structure. It can also be observed that the elemental distribution of Fig. 3(e-h) and (i-k) shows that the elements Ce, La, Ti, and V are all uniformly distributed in the catalyst.<sup>20</sup>

### 3.2 Catalyst surface acidity

$\text{NH}_3$ -TPD was used to investigate the acidic properties of the catalyst surface (Fig. 4). While the peaks of medium and strong



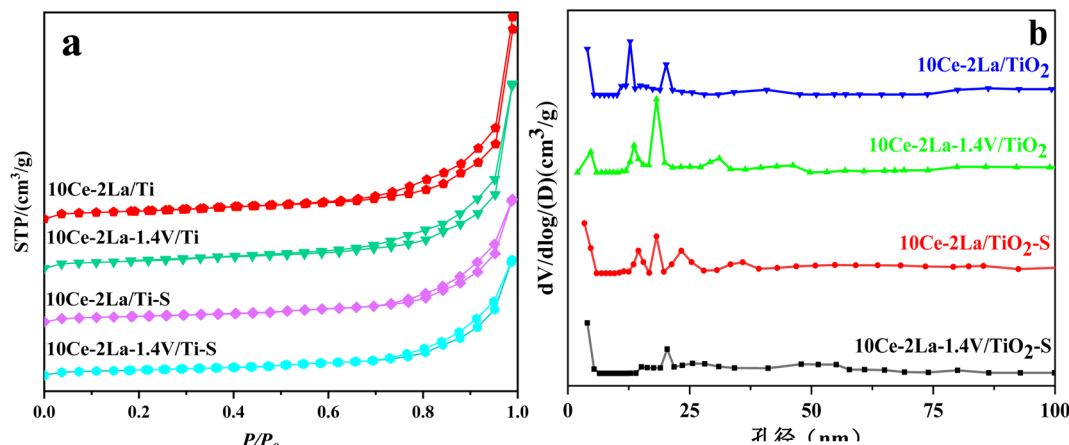


Fig. 2 (a) N<sub>2</sub> adsorption-desorption isotherms and (b) BJH pore size distributions of 10Ce-2La-xV/Ti series catalysts before and after poisoning.

Table 1 Structural information of 10Ce-2La-xV/Ti series catalysts

Samples	Surface area (m <sup>2</sup> g <sup>-1</sup> )	Pore volume (cm <sup>3</sup> g <sup>-1</sup> )	Diameter of hole (nm)
10Ce-2La/Ti	74.13	0.062	21.96
10Ce-2La-1.4V/Ti	71.81	0.056	16.55
10Ce-2La/Ti-S	57.51	0.037	15.73
10Ce-2La-1.4V/Ti-S	54.30	0.035	15.65

acidic sites in the temperature range of 200–500 °C were associated with the desorption of NH<sub>4</sub><sup>+</sup> from Brønsted acidic sites, the peaks of weak acidic sites below 200 °C corresponded to physisorbed NH<sub>3</sub> on the catalyst. The peaks of strong acidic sites >500 °C were attributed to the desorption of liganded NH<sub>3</sub> on Lewis acidic sites.<sup>21</sup> By comparison, we found that the weak acidic site peaks did not change much after the introduction of V. However, the peak intensities and peak areas of the strong acidic sites in the catalysts increased significantly and the

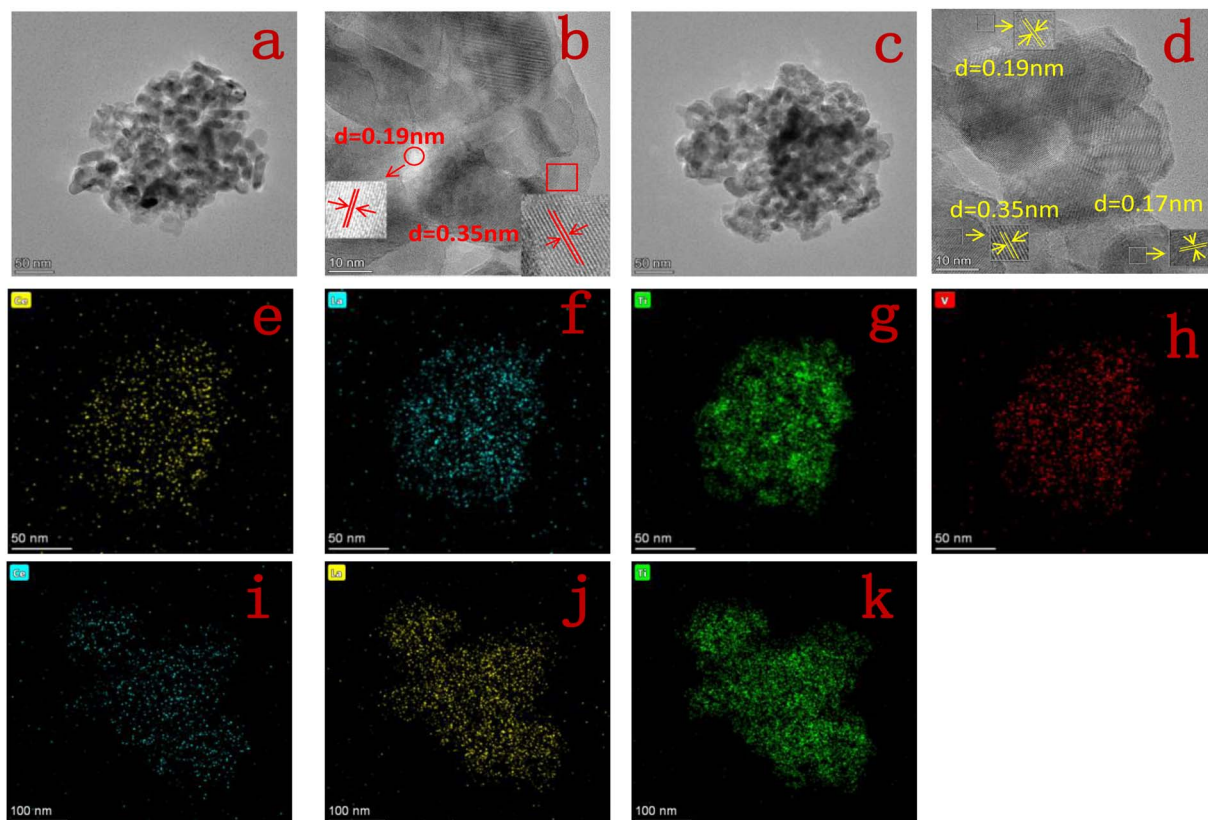


Fig. 3 (a and b) TEM and HR-TEM image of 10Ce-2La/Ti; (c and d) TEM and HR-TEM image of 10Ce-2La-1.4V/Ti; (e-h) elemental distribution of Ce, La, Ti, V of 10Ce-2La-1.4V/Ti; (i-k) elemental distribution of Ce, La, and Ti of 10Ce-2La/Ti.



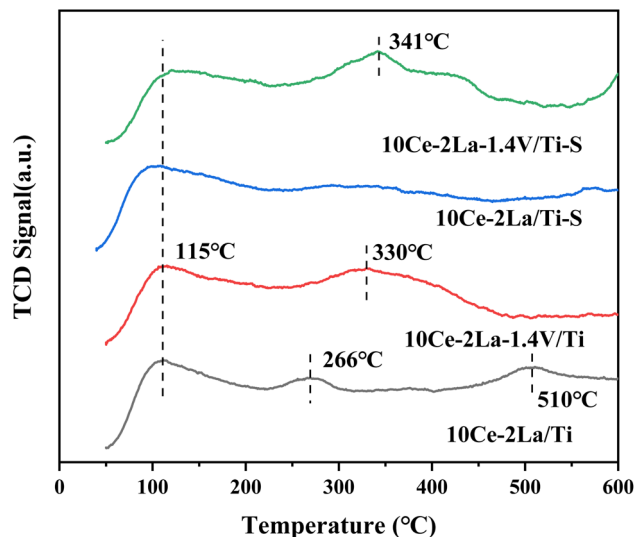


Fig. 4  $\text{NH}_3$ -TPD spectra of 10Ce-2La- $x$ V/Ti series catalysts before and after poisoning.

detachment peaks moved toward higher temperatures, which may be due to the interaction of the different active components, which in turn led to an increase in the length of the metal-oxygen coordination bonds and the surface acidity.<sup>22</sup> Meanwhile the decrease in the peak area of the strong acidic sites indicates that the medium-strong and weak acidic sites play a greater role in the catalytic activity of the catalyst during the catalytic reaction. In addition, it was observed that the peak intensity of the weak acidic sites of the 10Ce-2La/Ti catalyst became smaller after poisoning, which may be due to the reaction of  $\text{SO}_2$  on the surface of the catalyst to generate sulfate to reduce the number of acidic sites, which led to the weakening of the catalyst's ability to adsorb  $\text{NH}_3$ . In contrast, the peak position of catalyst 10Ce-2La-1.4V/Ti-S not only shifted toward high temperature, but also the peak intensity and peak area of the medium-strong acidic sites increased instead, indicating that the sulfation of the catalyst increased the  $\text{NH}_3$  uptake by the medium, which is one of the reasons that the catalyst exhibits good water and sulfur.<sup>23</sup>

### 3.3 Catalyst redox properties

XPS was used to investigate the chemical environment of the catalyst surface elements. The XPS spectra of catalyst Ce 3d are shown in Fig. 5(a), in which ten peaks were fitted to the spectra of all the samples. These were identified with "U" and "V" for the  $3d_{5/2}$  and  $3d_{3/2}$  spin-orbit components of Ce, respectively, which were located at 879 eV ( $V_0$ ), 885 eV ( $V_2$ ), 900 eV ( $U_0$ ), and 905 eV ( $U_2$ ). These belong to the  $3d^{10}4f^1$  initial electronic state of  $\text{Ce}^{3+}$ , and the remaining peaks were related to the  $3d^{10}4f^0$  state of  $\text{Ce}^{4+}$ . The existence of the double peaks of  $V_0/U_0$  and  $V_2/U_2$  suggests the presence of partially-reduced oxygen vacancies in the catalyst.<sup>24</sup> The fitted integrated peak area ratio of  $\text{Ce}^{3+}$  and total Ce was used to determine the relative content of  $\text{Ce}^{3+}$  on the catalyst surface, as shown in Table 2 by the formula  $\text{Ce}^{3+} (\%) = (S_{V_0} + S_{V_2} + S_{U_0} + S_{U_2}) / (S_U + S_V) \times 100\%$ .<sup>25</sup> After

introducing V, Ce and La concentrations on the catalyst surface considerably rose, proving that V interacted with the active ingredients and had an impact on how many atoms were present there. The higher proportion of  $\text{Ce}^{3+}$ , was attributed to an increase in the Ce concentration and also to the movement of free carriers on the catalyst surface towards the  $\text{CeO}_2$  surface, which had a higher binding energy (eV).<sup>26</sup> A large fraction of  $\text{Ce}^{3+}$  on the catalyst surface enhanced the formation of oxygen vacancies, unsaturated chemical bonds, and charge imbalance, all of which boost the NO oxidation reaction.<sup>27,28</sup> Fig. 5(b) depicts the catalyst's La 3d XPS spectra, in which La  $3d_{5/2}$  and La  $3d_{3/2}$  peaks emerged in all spectra at 834 eV and 37 eV, as well as at 852 eV and 855 eV, respectively.<sup>29</sup>

Fig. 5(c) shows the O 1s XPS spectrum of the catalyst, in which peaks belonging to lattice oxygen and surface unsaturated oxygen (labeled  $O_\beta$  and  $O_\alpha$ ), can be observed at positions 528.7–529 eV and 530.1–530.7 eV. Because surface unsaturated oxygen  $O_\alpha$  has a higher migration rate than lattice oxygen  $O_\beta$ ,  $O_\alpha$  increased the number of surface oxygen vacancies, which were more favorable to the SCR reaction.<sup>30</sup> The relative atomic percentages of these oxygen species on the catalyst surface were estimated from the areas of the fitted peaks, as listed in Table 2. The  $O_\alpha / (O_\alpha + O_\beta)$  ratio on the catalyst surface significantly increased after adding V, which is consistent with the Ce 3d XPS analysis and is one of the reasons for the good activity of the catalyst. Fig. 5(d) showed the V 2p XPS spectra before and after catalyst poisoning, in which the  $V^{4+}$  peak was located at 515.2 eV, and the  $V^{5+}$  peaks were located at 516.4 eV and 523.1 eV. This indicates that V on the catalyst surface mainly existed as  $V^{5+}$ .<sup>31</sup> The ratio of  $V^{4+} / (V^{4+} + V^{5+})$  on the catalyst surface increased significantly after poisoning.  $\text{Ce}^{3+}$  also increased significantly, but the concentrations of Ce and V, the main active components on the surface of the catalyst, decreased. This was because  $\text{SO}_2$  reacted with the active components to produce sulfate in the presence of  $\text{O}_2$ .<sup>7</sup> Similarly, the significant increase in the  $O_\alpha / (O_\alpha + O_\beta)$  ratio on the surface after catalyst poisoning was because the higher  $\text{Ce}^{3+}$  ratio promoted the creation of more surface oxygen vacancies. This was because oxygen from sulfate species was formed on the sample surface after sulfation, as well as chemisorbed oxygen.<sup>32</sup> Fig. 5(e) shows the S 2p XPS spectra of the catalyst before and after poisoning, in which peaks belonging to the spin-orbit components of  $S^{6+} 2p_{3/2}$  and  $S^{6+} 2p_{1/2}$  can be observed at 167.7–168 eV and 168.8–169 eV after fitting. This implies that S in +6 valence state was mostly present on the catalyst surface as  $\text{SO}_4^{2-}$ .<sup>33,34</sup> After poisoning, the XPS peaks of all elements on the catalyst changed to higher binding energies, showing that the electron cloud density around each element was lowered to varying degrees. All were involved in the sulfation reaction. Peaks of Ce, La, O, Ti, V, S and C were observed in the full spectrum of the catalyst in Fig. 5(f).

As shown in Fig. 6, the redox properties of catalysts before and after 10Ce-2La/Ti and 10Ce-2La-1.4V/Ti poisoning were tested using  $\text{H}_2$ -TPR. Measurements were performed using CuO as the standard, and the hydrogen consumption for the reduction peaks between 400 °C and 700 °C was quantitatively calculated (Table 3). 10Ce-2La/Ti displayed a large  $\text{H}_2$  reduction



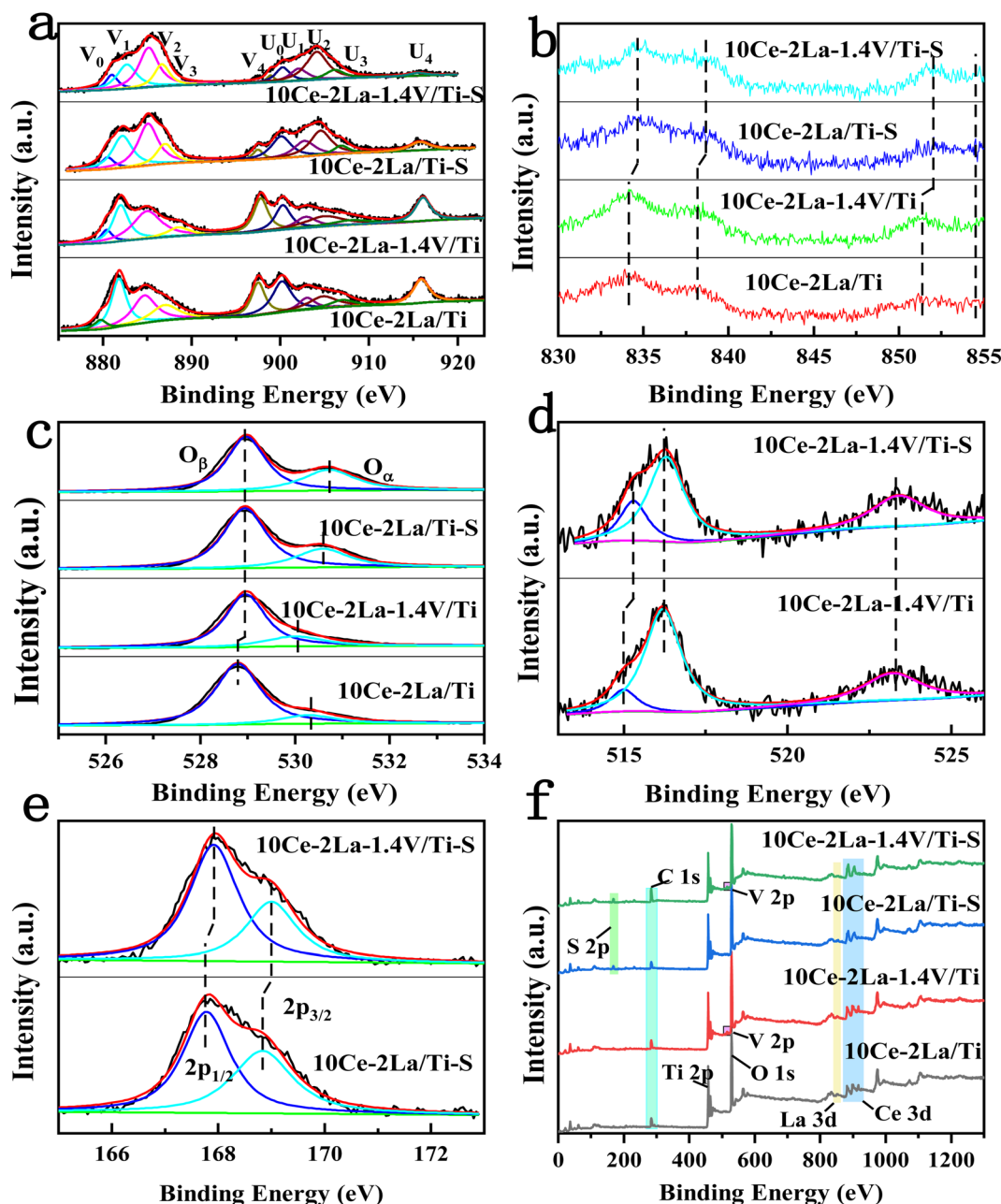


Fig. 5 XPS spectra of 10Ce-2La/Ti and 10Ce-2La-1.4V/Ti catalysts and before and after poisoning. (a) Ce 3d; (b) La 3d; (c) O 1s; (d) V 2p; (e) S 2p; (f) full spectrum.

peak, which was attributed to the reduction of surface-ligated unsaturated  $\text{Ce}^{4+}$ , which played an important role in the oxidation reaction.<sup>35</sup> After introducing V, the catalyst reduction

peaks shifted toward lower temperatures, and the  $\text{H}_2$  consumption increased significantly ( $1.37 \rightarrow 1.51 \text{ mmol g}^{-1}$ ). This indicates strong interactions between V oxides and Ce

Table 2 Surface element concentrations and  $\text{Ce}^{3+}$ ,  $\text{O}_\alpha$ , and  $\text{V}^{4+}$  ratios of catalysts

Samples	Surface atomic concentration (at%)						$\text{Ce}^{3+}/\text{Ce}$	$\text{O}_\alpha/\text{O}$	$\text{V}^{4+}/\text{V}$
	O	Ce	La	V	S	Ti			
10Ce-2La/Ti	71.94	2.74	0.28	—	—	25.04	42.14%	14.34%	—
10Ce-2La-1.4V/Ti	70.58	3.42	0.40	1.21	—	24.92	45.60%	21.40%	9.98%
10Ce-2La/Ti-S	70.53	2.39	0.25	—	3.75	23.08	53.49%	27.51%	—
10Ce-2La-1.4V/Ti-S	69.84	2.92	0.40	0.72	4.63	21.50	57.30%	34.07%	18.83%



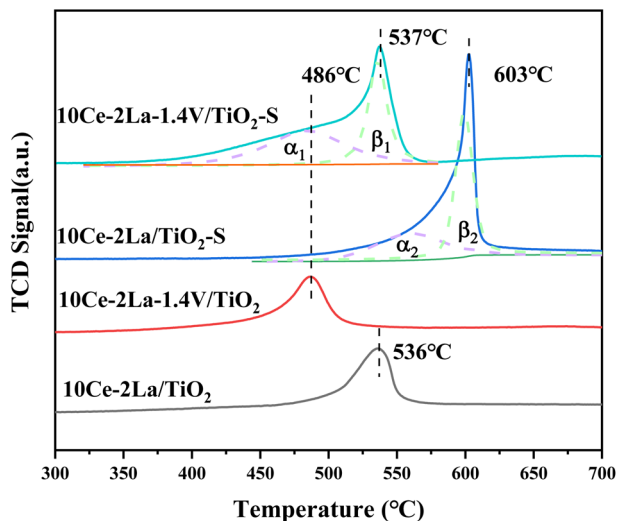


Fig. 6  $\text{H}_2$ -TPR spectra of 10Ce-2La-xV/Ti series catalysts.

Table 3 Catalyst  $\text{H}_2$ -TPR peak positions and corresponding  $\text{H}_2$  consumption

Samples	Peak position (°C)	$\text{H}_2$ consumption ( $\text{mmol g}^{-1}$ )
10Ce-2La/Ti	536	1.37
10Ce-2La-1.4V/Ti	486	1.51
10Ce-2La/Ti-S	603	2.05
10Ce-2La-1.4V/Ti-S	537	2.54

oxides on the catalyst, which enhanced the reducing properties of the catalyst, which promoted catalytic cycling during SCR reaction.<sup>36</sup> A strong peak was observed at 603 °C in the poisoned 10Ce-2La/Ti-S sample, which was shifted to a higher temperature relative to the fresh sample. This shift was primarily due to the reduction of  $\text{S}^{6+}$  to  $\text{S}^{4+}$  in sulfate, along with the reduction of certain metal oxides.<sup>37</sup> To study the degree of metal oxide reduction in the catalyst 10Ce-2La-1.4V/Ti-S, the peaks were divided into  $\alpha$ - and  $\beta$ -peaks by fitting, and the fitted area ratio was used to calculate the amount of hydrogen consumed for each peak. The  $\alpha$  peak was associated with the reduction of metal oxides on the catalyst surface, while the  $\beta$  peak was related to the reduction process of sulfate species.<sup>36</sup> The position of the metal oxide reduction peak in 10Ce-2La-1.4V/Ti-S was essentially unchanged, and the hydrogen consumption of  $\alpha_1$  was larger than that of  $\alpha_2$  ( $1.54 > 0.89 \text{ mmol g}^{-1}$ ). That of  $\beta_1$  was less than that of  $\beta_2$  ( $1.00 < 1.157 \text{ mmol g}^{-1}$ ), indicating that co-doping V with 10Ce-2La/Ti limited the interactions between  $\text{SO}_2$  and active metal oxide.

### 3.4 $\text{SO}_2$ desorption and reaction

$\text{SO}_2$ -TPD was used to investigate the adsorption capacity of the catalyst toward  $\text{SO}_2$ . As shown in Fig. 7(a), the catalyst contained two main desorption peaks in the temperature ranges of 400–460 °C and 600–670 °C. The low-temperature peaks were related to the desorption of  $\text{SO}_2$  from the catalyst surface,<sup>38</sup> while the high-temperature peaks were related to the desorption of  $\text{SO}_2$

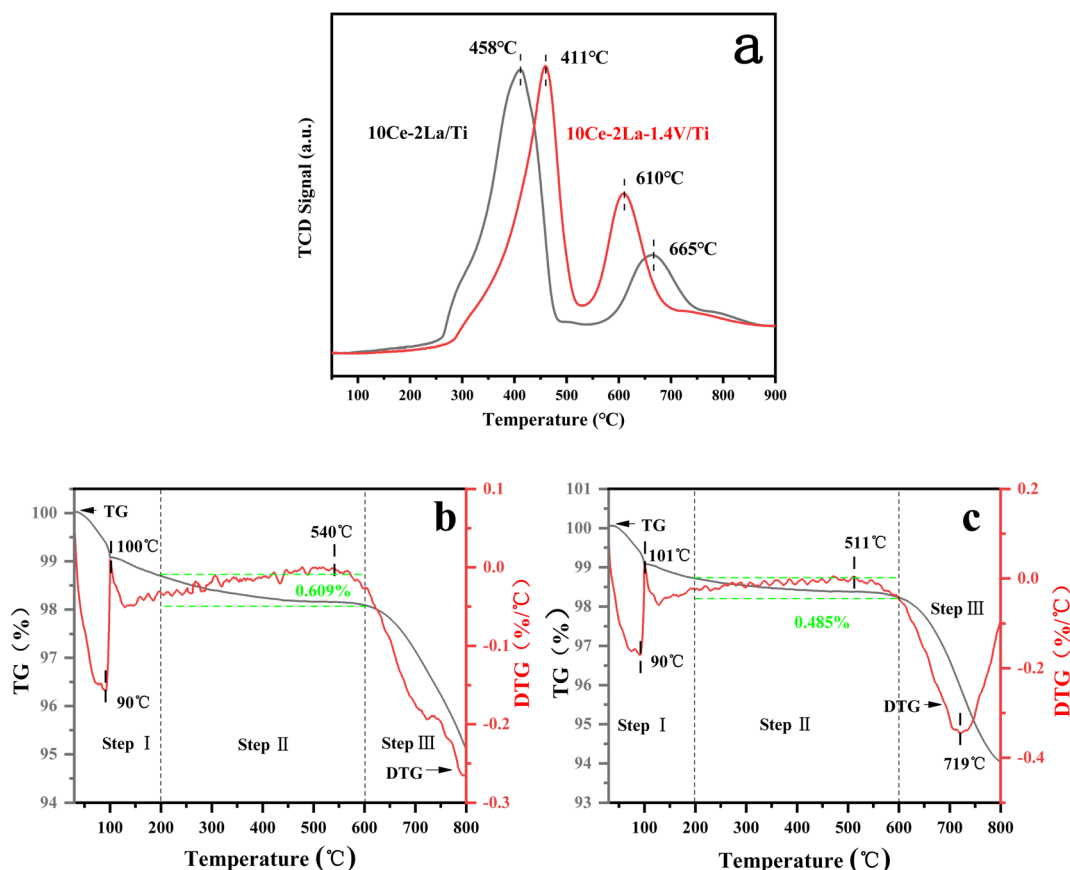


Fig. 7  $\text{SO}_2$ -TPD curves of fresh catalysts and TG/DTG curves of poisoned catalysts. (a)  $\text{SO}_2$ -TPD curves; (b) 10Ce-2La/Ti; (c) 10Ce-2La-1.4V/Ti.



generated by the decomposition of sulfate species. The peak of 10Ce-2La/Ti at 665 °C represented the decomposition of cerium sulfate.<sup>39</sup> In comparison, the desorption peak of 10Ce-2La/Ti at low temperature has a larger peak area, implying that more SO<sub>2</sub> molecules are adsorbed on the surface than 10Ce-2La-1.4V/Ti, the addition of V reduced the amount of SO<sub>2</sub> molecules adsorbed on the surface of the catalyst in order to alleviate competition between SO<sub>2</sub> and NO adsorption, and to prevent the occupation of more active sites. The addition of V enhanced the strength of the catalyst's high-temperature peak, indicating that SO<sub>2</sub> molecules were adsorbed onto V<sub>2</sub>O<sub>5</sub> species, shielding the catalyst's major active sites.<sup>40</sup>

To study the substances generated on the surface of the samples during the catalytic reaction, the poisoned samples were analyzed using thermogravimetric analysis, and the results are shown in Fig. 7(b) and (c). The two samples' weight reduction procedure was separated into three steps: step I (40–200 °C) was mostly ascribed to the desorption of water from the samples; step II (200–600 °C) was attributed to the breakdown of ammonium sulfate and ammonia bisulfate on the catalyst surface; and step III (600–800 °C) to the decomposition of

sulfates in the samples.<sup>36,41</sup> 10Ce-2La-1.4V/Ti-S showed an absorption peak related to the decomposition of sulfate (511 °C) and an exothermic peak related to the phase transition of the composite metal oxide (719 °C) above 500 °C. The 10Ce-2La/Ti-S catalyst had a higher decomposition temperature (540 °C), which was attributed to the low decomposition temperature of VOSO<sub>4</sub> generated by SO<sub>2</sub> with V. The 10Ce-2La/1.4V/Ti-S catalyst had the highest decomposition temperature (540 °C). In addition, the weight loss of ammonium sulfate or ammonia bisulfate on the surface of the 10Ce-2La-1.4V/Ti-S catalyst was larger (0.609%) than that of 10Ce-2La/Ti-S (0.485%). It is shown that the introduction of V protects the main active component of the catalyst from reacting with SO<sub>2</sub> and inhibits the formation of sulfate species on the catalyst surface, thus enhancing the water and sulfur resistance of the catalyst.

### 3.5 *In situ* infrared spectroscopy of catalysts

Fig. 8(a) and (b) show the *in situ* DRIFTS spectra of catalysts 10Ce-2La/Ti and 10Ce-2La-1.4V/Ti after adsorbing NH<sub>3</sub> (500 ppm) for 30 min. The peaks near 954 (960) cm<sup>-1</sup> for both samples were related to weakly-adsorbed gas-phase NH<sub>3</sub>.<sup>42,43</sup>

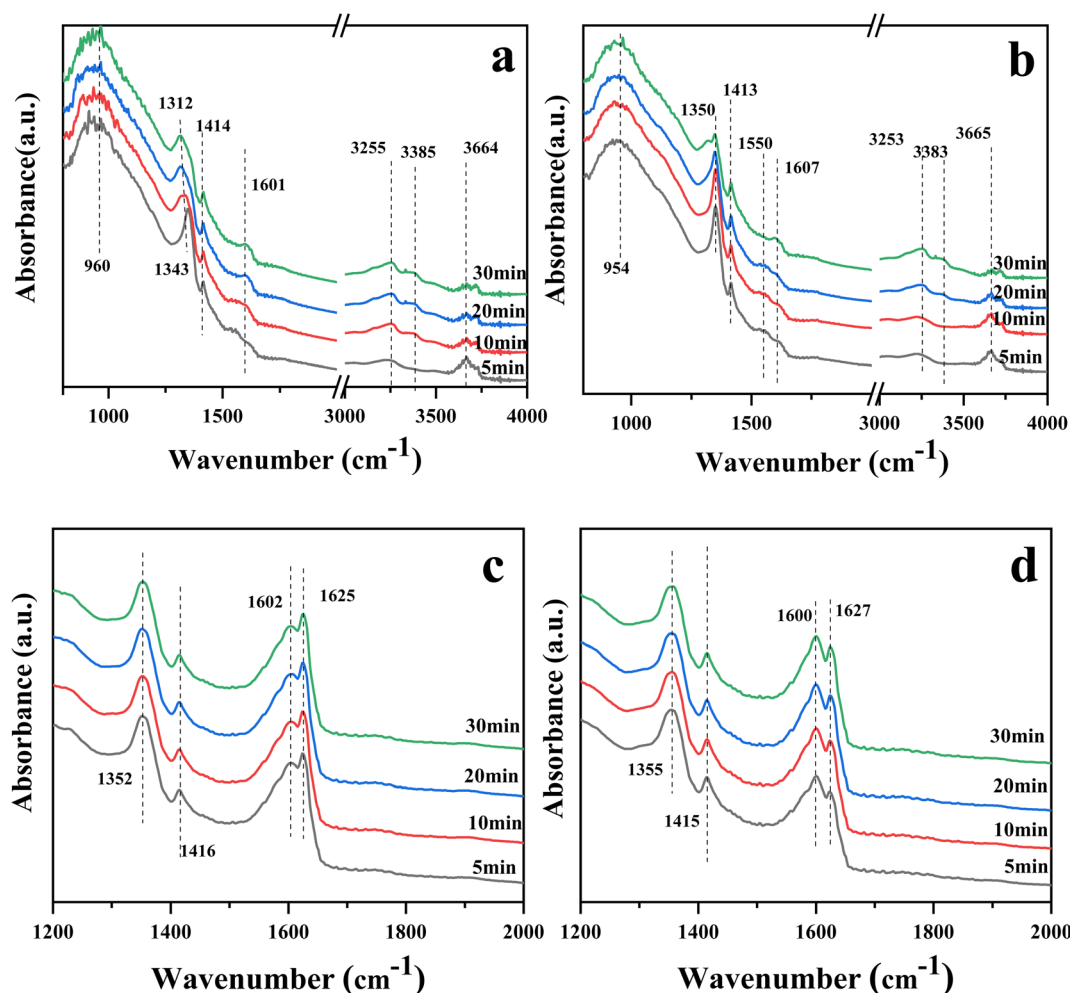


Fig. 8 *In situ* DRIFTS spectra of NH<sub>3</sub> or NO + O<sub>2</sub> adsorption on catalysts. (a) NH<sub>3</sub> adsorption on 10Ce-2La/Ti, (b) NH<sub>3</sub> adsorption on 10Ce-2La-1.4V/Ti, (c) NO + O<sub>2</sub> adsorption on 10Ce-2La/Ti, and (d) NO + O<sub>2</sub> adsorption on 10Ce-2La-1.4V/Ti.





Absorption peaks near 1312–1343 (1350)  $\text{cm}^{-1}$  and 1414 (1413)  $\text{cm}^{-1}$  belonged to  $\text{NH}_4^+$  at Brønsted acid sites.<sup>44,45</sup> The asymmetric bending vibration of  $\text{NH}_3$  coupled to Lewis acid sites was responsible for the lower absorption peak at 1601 (1607)  $\text{cm}^{-1}$ .<sup>44</sup> The peak at 3253–3385  $\text{cm}^{-1}$  was attributed to the stretching vibration of  $\text{NH}_3$  at the Lewis acid sites.<sup>46</sup> 10Ce–2La–1.4V/Ti shows a weak adsorption band near 1550  $\text{cm}^{-1}$ , which is due to the oxidation of  $\text{NH}_3$  on the catalyst surface by surface-active oxygen to generate an important intermediate product,  $-\text{NH}_2$ , in the catalytic reaction  $-\text{NH}_2$  facilitates the conversion of NO, and promotes the smooth progress of the SCR reaction.<sup>47</sup> In comparison, the intensity of the peaks related to the Brønsted acidic sites increased after the introduction of V, and  $\text{NH}_3$  was more easily activated. This indicates that Brønsted acids dominated the reaction, and the addition of V increased the proportion of Brønsted acids.

The DRIFTS spectra of the catalysts 10Ce–2La/Ti and 10Ce–2La–1.4V/Ti after 30 minutes of NO (500 ppm) +  $\text{O}_2$  adsorption are shown in Fig. 8(c) and (d). Nitrate bridged ions and free nitrate ions, respectively, were responsible for the bands at 1355 (1352)  $\text{cm}^{-1}$  and 1600 (1602)  $\text{cm}^{-1}$ .<sup>48</sup> The absorption peak

belonging to the nitrite species  $\text{trans-N}_2\text{O}_2^{2-}$  was located at 1415  $\text{cm}^{-1}$ . Weakly adsorbed gas-phase  $\text{NO}_2$  was thought to be active in the band at 1625–1627  $\text{cm}^{-1}$  because it may react quickly with gaseous  $\text{NH}_3$ .<sup>45</sup>

Fig. 9(a) and (b) show the DRIFTS profiles of the catalysts 10Ce–2La/Ti and 10Ce–2La–1.4V/Ti after 30 min of pre-adsorption of  $\text{NH}_3$  (500 ppm) followed by NO (500 ppm) +  $\text{O}_2$  (3%) adsorption for 30 min. In the  $\text{NH}_3$  adsorption curve, the peaks near 1317 and 1415 (1417)  $\text{cm}^{-1}$  belonged to  $\text{NH}_4^+$  on the Brønsted acid sites, and the intensity of the peak at 1415 (1417)  $\text{cm}^{-1}$  did not change greatly with the passage of NO +  $\text{O}_2$ . This was attributed to the coincidence of the peak at 1600 (1604)  $\text{cm}^{-1}$  for the  $\text{NH}_3$  and Lewis acid sites with the stronger peaks of rapidly-generated nitrate at 1604 and 1625  $\text{cm}^{-1}$  after flowing NO +  $\text{O}_2$ . This indicates that pre-adsorbed  $\text{NH}_3$  did not directly react with gas-phase NO and that the  $\text{NH}_3$ -SCR did not involve gas-phase NO. With the passage of NO +  $\text{O}_2$ , the intensity of the 1415 (1417)  $\text{cm}^{-1}$  peak did not change significantly, because, at this time, the 1600 (1604)  $\text{cm}^{-1}$  peaks of  $\text{NH}_3$  and Lewis acid sites overlapped with those of the stronger nitrate peaks at 1604 and 1625  $\text{cm}^{-1}$ . This suggests that pre-adsorbed

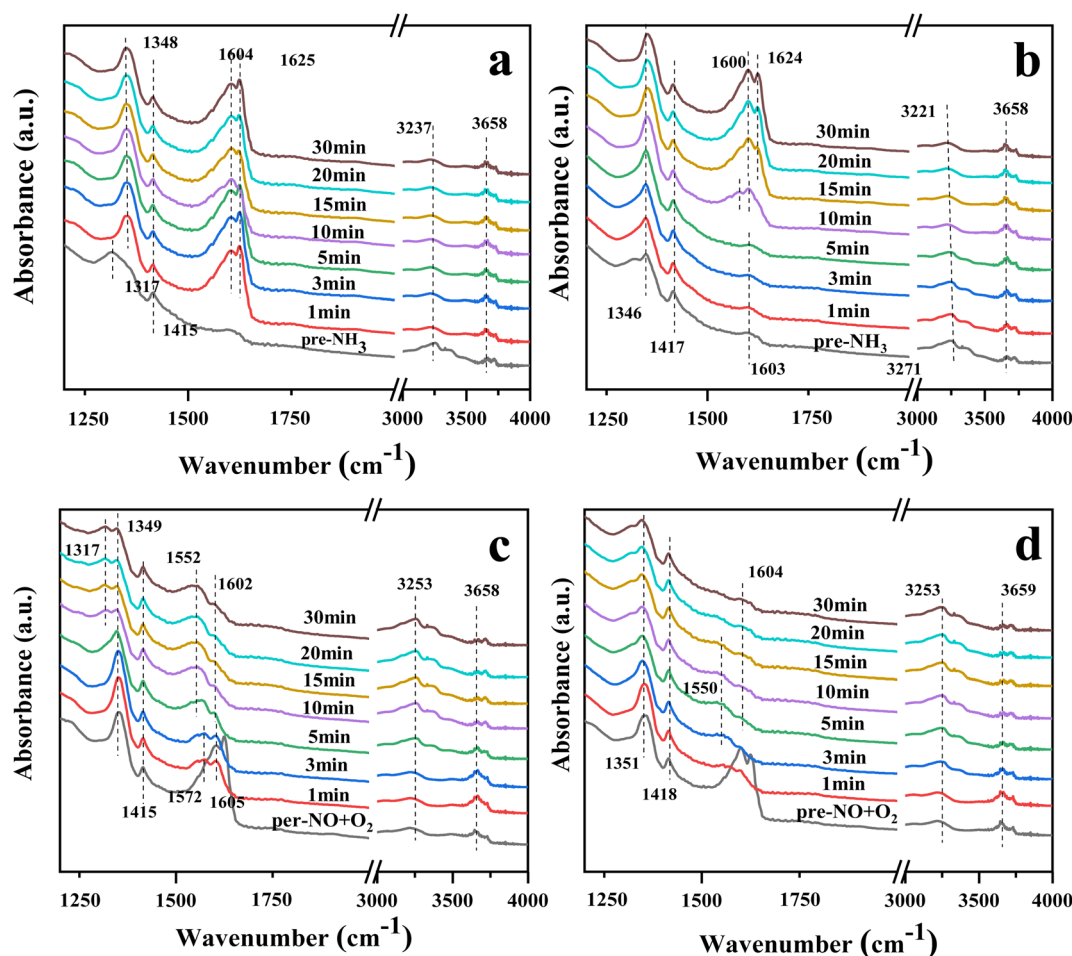


Fig. 9 *In situ* DRIFTS spectra of  $\text{NH}_3(\text{NO} + \text{O}_2)$  reaction with pre-adsorbed  $\text{NO} + \text{O}_2(\text{NH}_3)$  on the catalyst. (a) Reaction of  $\text{NO} + \text{O}_2$  on 10Ce–2La/Ti with pre-adsorbed  $\text{NH}_3$ ; (b) reaction of  $\text{NO} + \text{O}_2$  on 10Ce–2La–1.4V/Ti with pre-adsorbed  $\text{NH}_3$ ; (c) reaction of  $\text{NH}_3$  on 10Ce–2La/Ti with pre-adsorbed  $\text{NO} + \text{O}_2$ ; (d) reaction of  $\text{NH}_3$  on 10Ce–2La–1.4V/Ti with pre-adsorbed  $\text{NO} + \text{O}_2$  reaction.



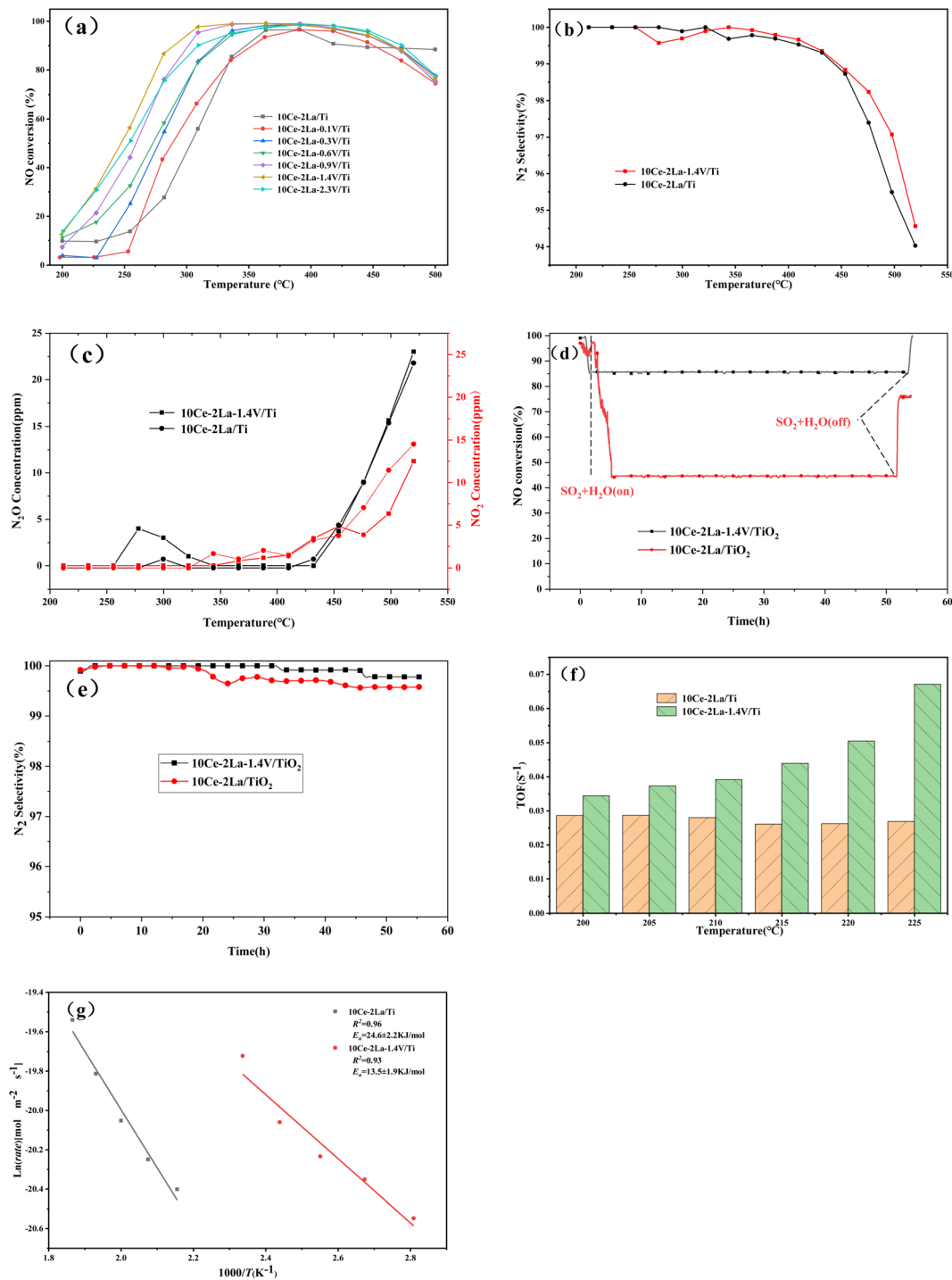


Fig. 10 Catalytic performance versus temperature curves of 10Ce-2La-xV/Ti series catalysts. (a) NO removal efficiency versus temperature transformation curves at different weight ratios V; (b) N<sub>2</sub> selectivity at different temperatures; (c) effect of temperature on the concentration of NO<sub>2</sub> and N<sub>2</sub>O production; (d) stability test of the catalysts for water and sulfur resistance; (e) N<sub>2</sub> selectivity at different time; (f) TOF values (200–225 °C). (g) Arrhenius plots.

NH<sub>3</sub> did not react directly with gas-phase NO and that the NH<sub>3</sub>-SCR reaction did not proceed *via* the Eley-Rideal mechanism.

Fig. 9(c) and (d) show the DRIFTS spectra of the catalysts 10Ce-2La/Ti and 10Ce-2La-1.4V/Ti pre-adsorbed with NO (500 ppm) + O<sub>2</sub> (3%) for 30 min and then allowed to adsorb NH<sub>3</sub> (500 ppm) for 30 min. Prior to the passage of NH<sub>3</sub>, peaks attributed

to nitrates were observed at 1349 (1351) cm<sup>-1</sup>, 1624 (1625) cm<sup>-1</sup>, 1415 (1418) cm<sup>-1</sup>, and 1600 (1604) cm<sup>-1</sup>, similar to those in Fig. 8(a) and (b).<sup>49</sup> The nitrate peaks at 1602 and 1625 cm<sup>-1</sup> disappeared rapidly upon the passage of NH<sub>3</sub>, it indicates that adsorbed NO can react with NH<sub>3</sub> in the adsorbed state, so the reaction mechanism is Langmuir-Hinshelwood mechanism. In

an L-H reaction mechanism, the reaction orders of both NO and  $\text{NH}_3$  are zero.<sup>50</sup>

### 3.6 Catalyst denitrification performance testing

Fig. 10(a) shows the NO conversion *versus* temperature curves of the catalysts after adding different ratios of V. The overall activity of the catalyst was improved after adding V, and the starting temperatures were all lower. As the weight percent of V increases, the catalyst denitrification efficiency improved and then became worse, and the temperature window gradually became wider. The optimal catalytic efficiency was obtained when the weight ratio of  $\text{V}_2\text{O}_5$  was 1.4, and the denitrification efficiency remained above 80% in the temperature range of 270–490 °C. The NO conversion reached a maximum of 99% at 330 °C, and the catalytic activity was greatly improved compared with the original rare-earth-based catalyst. In addition to catalytic activity,  $\text{N}_2$  selectivity is another important index for evaluating a catalyst's performance. As the temperature of the reaction increased,  $\text{N}_2\text{O}$  and  $\text{NO}_2$  were produced, causing the  $\text{N}_2$  selectivity to slowly decrease, as shown in Fig. 10(b) and (c). In the reaction temperature interval, the  $\text{N}_2$  selectivity and  $\text{N}_2\text{O}$  concentration of catalyst was slightly improved by the addition of V. The lower  $\text{N}_2\text{O}$  generation resulted in better overall  $\text{N}_2$  selectivity than the original catalyst.

The stability of the catalyst was tested by passing 300 ppm  $\text{SO}_2$  under constant test conditions and the results were shown in Fig. 10(d). After 2.5 h of the water and sulfur feeding, the NO conversion of 10Ce–2La/Ti decreased from the initial 97% to 44%, which maintained almost unchanged for 52 h. If the feeding of water and sulfur was cut off, the NO conversion recovered to 78%. Compared with 10Ce–2La/TiO<sub>2</sub>, the V-doped catalyst had a higher NO conversion in the presence of water and sulfur, which remained above 85% and recovered to 99% after shutting down water and sulfur. Fig. 10(e) shows the  $\text{N}_2$  selectivity comparison in the water and sulfur resistance stabilization experiments. Both the two catalysts showed a high  $\text{N}_2$  selectivity over than 99% and the  $\text{N}_2$  selectivity was not influenced by water and sulfur. It can also seen that the  $\text{N}_2$  selectivity was slightly improved by the addition of V. The turnover frequency (TOF) value calculated from eqn (5) is shown in Fig. 10(f). The 10Ce–2La–1.4V/Ti catalyst displayed better TOF values at each temperature, demonstrating that single Ce atoms on the catalytic surface were the primary active sites and displayed the best intrinsic activity under identical circumstances. Additionally, the reaction rate of NO on each square meter of a catalyst within the investigated temperature range was calculated according to eqn (4). It was plotted as  $\ln(\text{rate})$  against  $1000/T$ , and the Arrhenius activation energy of the reaction was determined by fitting. As shown in Fig. 10(g), the surface activation energy of 10Ce–2La/Ti was 24.6 kJ mol<sup>−1</sup>, which was higher than that of 10Ce–2La–1.4V/Ti (13.5 kJ mol<sup>−1</sup>). The lower activation energy suggests an improved denitrification efficiency.

## 4. Conclusion

V-doped 10Ce–2La–xV/Ti catalysts were prepared by a ball milling method, and it was found that the rare earth Ce exerted

a synergistic effect with V. The composite carrier-type oxides exhibited a good activity window and denitrification performance, as well as excellent water and sulfur resistance when the weight ratio of  $\text{V}_2\text{O}_5$  was 1.4. The prepared catalysts showed mesoporous structures, and the ball milling method enhanced the interactions between different active components, resulting in good dispersion of the active components. V doping improved the surface acidity of the catalysts, and Brønsted acids on the surface of the catalysts played the dominant role. The greater number of acid sites adsorbed more  $\text{NH}_3$  that participated in the reaction and also prevented  $\text{SO}_2$  poisoning. The  $\text{Ce}^{3+}$  and O ratios on the catalyst surface were also enhanced by the addition of V, which was significant for the SCR- $\text{NH}_3$  reaction. V limited the interactions between  $\text{SO}_2$  and the main active component (metal oxides), reduced the amount of  $\text{SO}_2$  adsorbed on the catalyst surface, and inhibited the formation of sulfate species on the catalyst surface. These all improved the sulfur resistance of the catalyst. The  $\text{NH}_3$ -SCR on the catalyst surface proceeded *via* the Langmuir–Hinshelwood mechanism, with Brønsted acid sites playing the dominant role. The better denitrification activity and sulfur resistance of 10Ce–2La–xV/Ti catalysts make them promising for various applications.

## Author contributions

Yang Liu: writing – original draft, data curation, formal analysis, investigation, methodology. Na Wang: conceptualization, data curation, formal analysis, funding acquisition, investigation, methodology, project administration, supervision, validation, writing – review & editing. Huidong Xie: investigation, methodology, project administration, supervision. Yepeng Sun: data curation, formal analysis. Liang Zhang: investigation, methodology. Kaiyue Yang: investigation, methodology. Chang Yang: data curation, formal analysis. Chengmin Ge: conceptualization, funding acquisition, supervision, validation.

## Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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