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Plasma-catalytic direct splitting of CO₂ to CO over oxygen-deficient Mo-doped CeO₂ at ambient conditions was achieved. Developing new catalytic technologies involving plasma opens the door to unlocking the potential of using captured CO₂ to make valuable chemicals under mild conditions.

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Plasma-enhanced direct conversion of CO₂ to CO over oxygen-deficient Mo-doped CeO₂⁺

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Plasma CO₂ splitting to CO over oxygen-deficient Mo-doped CeO₂ under mild conditions was investigated for the first time, showing ~20 times higher CO₂ conversion compared to pure CeO₂, which can be attributed to the increased oxygen vacancies (V_O) and the formation of Ce³⁺-V_O-Mo on the catalyst surface. Importantly, V_O sites showed excellent catalytic stability.

Converting CO₂ to value-added fuels and chemicals has been considered as a promising route in CO₂ utilization. Significant efforts have been devoted to the chemical transformation of CO₂, including thermal catalysis,¹ photocatalysis,^{2,3} electrocatalysis,^{4,5} and plasma catalysis.^{6,7} Direct splitting of CO₂ to CO without using any reductant is attractive for CO₂ conversion, as CO is an important chemical feedstock for the synthesis of a range of chemicals and fuels. However, this reaction has to overcome a strong thermodynamic barrier (CO₂ \rightarrow CO + 1/2O₂, ΔH_{298K} = 280 kJ mol⁻¹ = 2.9 eV molecule⁻¹) to break the C=O bond, since CO₂ is very stable. Ultrahigh temperatures (2000 K) are often required to activate CO₂ (Fig. S1, ESI⁺).

In recent years, using non-thermal plasmas (NTPs) for the activation of inert molecules with strong chemical bonds (*e.g.*, CO_2 , CH_4 and N_2) under mild conditions has attracted significant interest, including CO_2 reforming with CH_4 to oxygenates, ^{8,9} CO_2 hydrogenation to methanol,¹⁰ and ammonia synthesis.¹¹ NTP is rich in energetic electrons with a typical energy of 1–10 eV, which is sufficient to activate reactants into chemically reactive species, enabling thermodynamically unfavorable reactions (*e.g.* CO_2 splitting) to proceed at ambient conditions.

Up until now, studies on plasma CO_2 splitting to CO mainly focused on the optimization of operating parameters, ^{12–16} and

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only a few catalysts (*e.g.*, Ni/SiO₂, NiO/TiO₂ and Rh/TiO₂) have been evaluated (Table S1, ESI†).^{17–20} Recently, surface oxygen vacancies (V_O) have been suggested to be the active sites in plasma-catalytic CO₂ splitting to CO. Mei *et al.* reported that higher CO₂ conversion was achieved when coupling plasma with BaTiO₃, which can be explained by the formation of more V_O sites on the surfaces of BaTiO₃ compared with TiO₂.²⁰ Chen *et al.* found the V_O on Ni/TiO₂ contributed to the enhanced CO₂ dissociation.¹⁸ In fact, the coupling of catalysts and plasma is a promising strategy to improve CO₂ conversion and energy efficiency. However, the knowledge in selection of appropriate catalysts for highly efficient CO₂ splitting to CO using NTP was still very limited until now.

Herein, CO₂ splitting to CO over M-doped CeO₂ catalysts (M = Fe, Co, Ni, Cu, Cr, V, Mn or Mo) has been carried out in a dielectric barrier discharge (DBD) reactor (Fig. S2–S5, ESI†). CeO₂ not only serves as a support to anchor and disperse the metal oxide particles but also generates V_O active sites through the interaction with metal oxides. Significant differences were observed among the M-doped CeO₂ catalysts in terms of CO₂ conversion, and the Mo-doped CeO₂ exhibited the best activity in CO₂ conversion (Fig. S3, ESI†). Therefore, Mo-doped CeO₂ has been selected for further studies.

Comprehensive catalyst characterization was carried out to understand the physicochemical properties of Mo-doped CeO₂. As shown in Fig. 1, the X-ray diffraction (XRD) pattern of CeO₂ exhibits characteristic peaks of a cubic fluorite phase (JCPDS, 34-0394). For Mo-doped CeO₂, small peaks of α -MoO₃ (JCPDS, 05-0508) and Mo₄O₁₁ (JCPDS, 05-0337) are observed, revealing the coexistence of Mo(vi) and Mo(v) species. Notably, the characteristic peaks of CeO₂ downshift compared to pure CeO₂, which suggests that Mo doping leads to the expansion of the CeO₂ unit cell. Usually, inserting Mo ions into CeO₂ induces shrinkage of the CeO₂ unit cell since the radius of Mo ions is much smaller than that of Ce ions, resulting in upshifting of CeO₂ peaks, rather than downshifting. Thus, Mo ions do not insert into the CeO₂ unit cell, and there might be other reasons responsible for this downshift. X-ray photoelectron spectroscopy

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(XPS) was employed to analyze surface properties of Mo-doped CeO₂. The deconvoluted Ce 3d XPS spectra are presented in Fig. 2(a). The peaks labeled as v, v'', v''' and u, u'', u''' are assigned to $3d_{5/2}$ and $3d_{3/2}$ electrons of Ce^{4+} , respectively, while the peaks of v', u' and $v_o\!,\,u_o$ correspond to $3d_{5/2}$ and $3d_{3/2}$ electrons of Ce³⁺, respectively.²¹ Clearly, Ce³⁺ exists in CeO₂ and Mo-doped CeO₂, suggesting the formation of V_O in both samples. More importantly, the proportion of Ce³⁺ in Mo-doped CeO_2 is 40.2%, higher than that in CeO_2 (30.0%) (Fig. S6 and S7, ESI[†]). This finding suggests that Mo doping induces partial transformation of Ce4+ to Ce3+ and creates more Vo on Modoped CeO₂. As reconfirmed by the O 1s XPS spectra in Fig. 2(b) and Fig. S6 (ESI^{\dagger}), a higher surface V_O (30.4%) is achieved in the Mo-doped CeO_2 in comparison to pure CeO_2 (21.0%). Furthermore, the transformation of Ce⁴⁺ to Ce³⁺ leads to the expansion of the CeO₂ unit cell since the ion radius of Ce³⁺ (1.23 Å) is higher than that of Ce^{4+} (0.97 Å),²² which explains the reason for the downshifting of CeO₂ peaks in Fig. 1. More interestingly, the



Fig. 2 XPS and Raman spectra of CeO₂ and as-prepared Mo-doped CeO₂ (a) Ce 3d, (b) O 1s, (c) Mo 3d and (d) Raman spectra (V_O , O_C and O_L represent oxygen vacancy, chemisorbed oxygen species and oxygen lattice, respectively).

binding energies of Ce 3d shift significantly towards higher values after Mo doping, revealing that the electron density of the surface CeO₂ species is lower in Mo-doped CeO₂ compared with pure CeO₂, which might be induced by the electron transfer from Ce to Mo, due to the higher electronegativity of Mo. These results suggest different properties of V_O sites in the form of $Ce^{3+}-V_O$ in CeO_2 and $Ce^{3+}-V_O$ -Mo in Mo-doped CeO_2 , as well as the strong interaction between Mo and CeO₂, which agrees with the results of H_2 -temperature programmed reduction (H_2 -TPR), scanning electron microscopy (SEM) and transmission electron microscopy (TEM) (Fig. S8-S11, ESI⁺). Fig. 2(c) shows the deconvoluted Mo 3d spectra, in which the Mo-doped CeO₂ sample exhibits typical doublet peaks of Mo⁶⁺ with an energy gap of *ca.* 3.1 eV, indicating the formation of MoO_3 in Mo-doped CeO₂.²³ The two smaller peaks observed, however, are identified to be $3d_{3/2}$ and $3d_{5/2}$ electrons of Mo(v), demonstrating the formation of the non-stoichiometric MoO_{3-x} ,²³ which is consistent with the Mo_4O_{11} species confirmed by the XRD analysis (Fig. 1). Fig. 2(d) shows the Raman spectra of CeO₂ and Mo-doped CeO₂. For pure CeO_2 , the intense band at 465 cm⁻¹ is well-indexed to the typical F2g modes of a cubic CeO_2 fluorite structure, and the weak bands at 262, 597 and 1171 cm⁻¹ are assigned to V_o, reconfirming the XPS results in Fig. 2(a and b).²³ For Mo-doped CeO₂, the emerging Raman bands at 673, 824 and 997 cm^{-1} are assigned to MoO₃ crystallites.²⁴ However, the band at 955 cm⁻¹ is associated with Mo suboxides $(MoO_{3-x})^{24,25}$ These results indicate that the valence state of Mo in suboxides is Mo⁵⁺. Moreover, the presence of Mo⁵⁺ and Ce³⁺ indicates that more V_O sites are created through the strong interaction between Mo and CeO₂, as well as the calcining atmosphere with deficient oxygen and rich energetic Ar species, which agrees with the results reported by Chen et al.18

Fig. 3 shows the effect of different operating conditions on CO_2 conversion. Clearly, no reaction occurred without plasma (catalyst only, 400 °C). The conversion of CO_2 was 3.8% in the plasma reaction without a catalyst (plasma only). In the plasma reaction coupled with pure CeO_2 , the CO_2 conversion dropped to 1.2%, which suggests that pure CeO_2 is unfavorable for CO_2 splitting to CO despite CeO_2 being O-deficient. Using Mo-doped



Fig. 3 Effect of reaction temperature and additive gas on CO₂ conversion (CO₂ flow rate 40 ml min⁻¹, SEI 20 kJ L⁻¹, molar ratio 4:1 and 30 °C for CO₂/Ar and CO₂/N₂; catalyst only 400 °C).

8

CeO₂ instead of CeO₂, however, results in a significant increase of CO₂ conversion by a factor of 9 at 400 °C. In addition, the Mo-doped CeO₂ showed stable CO₂ conversion for at least 10 h (Fig. S12, ESI[†]). Interestingly, the reaction performance can be further improved by using a lower reaction temperature (30 °C) and an additive gas (Ar or N₂). This promotional effect was more pronounced when adding N2. The highest CO2 conversion of 23.2% and energy efficiency of 14.3% were achieved in the plasma splitting of CO2 with N2 addition over Mo-doped CeO2 at 30 °C (Table S1, ESI[†]), while the corresponding formation rate of CO and O_2 was 24.9 mmol h⁻¹ and 12.4 mmol h⁻¹, respectively (Table S2, ESI⁺). The optimal energy efficiency achieved in this study is comparable to that reported in previous works (Table S1, ESI[†]).

Regarding surface reactions, the improved activity over Modoped CeO₂, on one hand, is mainly attributed to the increased formation of V_{Ω} (Fig. 2 and Fig. S6, ESI^{\dagger}), since V_{Ω} serve as adsorption centers for CO₂ dissociative adsorption,^{17,26} *i.e.*, $CO_2 + V_O \rightarrow O_L/O_C + CO$. On the other hand, the promoted performance originates from the different properties of Vo in the forms of $Ce^{3+}-V_{O}$ and $Ce^{3+}-V_{O}$ -Mo due to strong interaction between Mo and CeO₂ and the higher electronegativity of Mo compared with Ce (Fig. 1, 2 and Fig. S8, ESI⁺), which leads to a stronger binding strength of CO₂ with the V_O of Ce³⁺–V_O–Mo. As discussed above, V_O is the active site for CO₂ activation; thus, the stability of V_{Ω} greatly influences the subsequent catalytic cycle. It is well recognized that oxygen can be produced in plasma CO₂ splitting, and the produced O species could poison the catalyst through filling V_O sites to form stable lattice oxygen species, resulting in termination of the catalytic cycle.

Therefore, the spent Mo-doped CeO₂ catalysts were further characterized by XPS, Raman, XRD and H2-TPR (Fig. 4). Interestingly, compared with the fresh catalyst, the spent Mo-doped CeO_2 catalysts show an increased amount of Ce^{3+} and MoO_{3-x} , which can be confirmed by the higher intensities of Ce³⁺ peaks

(b) (1) fresh

CeO,

Ce (2) using at 400 °C a-MoO (3) using at 30 °C with N (a.u.) Counts (s) MoO Intensity (3) (1) fresh (2) (2)(2) using at 400 °C (1) (1) (3) using at 30 °C with N. 910 900 890 880 870 1200 1000 800 600 400 920 Binding energy (eV) Raman shift (cm⁻¹ (C) CeO, (d) using at 30 °C with N₂ α-MoO. consumption (a.u.) Intensity (a.u.) ↓ Mo.O. fresh using at 30 ŕ frest

Ce3d

Fig. 4 (a) XPS spectra, (b) Raman spectra, (c) XRD patterns, (d) H₂-TPR profiles of Mo-doped CeO₂ before and after reaction.

300 400 500 600

Reduction temperature (°C)

700 800 and the Raman band at ~950 cm⁻¹ in Fig. 4(a and b), respectively. The enhancement effect is more pronounced for the catalyst used at 400 °C. Correspondingly, more Vo sites were created in the high-temperature reaction (Table S3, ESI⁺). In addition, the color of the catalyst changed from gray/blue (MoO_{3-x}) to yellow (MoO_{3-y}) with x > y after the reaction at 30 °C, while no visible changes were observed on the phase structure of Mo-doped CeO₂ before and after the reaction (30 $^{\circ}$ C), as shown in Fig. 4(c). These results indicate that some O atoms from CO_2 splitting are adsorbed onto the catalyst, as seen by the slightly increased H_2 consumption in Fig. 4(d). Even so, the V_O concentration of Mo-doped CeO₂ used at 30 °C remained at a similar level with that of the fresh sample (Table S3, ESI[†]). These findings suggest that Vo-rich Modoped CeO₂ is stable after the plasma reaction, and hightemperature reactions facilitate the formation and recovery of V_o sites, resulting from accelerating recombinative desorption of adsorbed O atoms.²⁷

In addition to surface reactions, gas-phase reactions also play a crucial role in the plasma-catalytic process. In a pure CO_2 DBD, CO₂ splitting to CO mainly proceeds through the electron impact dissociation of CO_2 ($CO_2 + e \rightarrow CO + O + e$), which can be confirmed by plasma chemical kinetic modeling,²⁸ as well as the formation of CO bands and O atomic lines detected by optical emission spectra of CO₂ DBD (Fig. 5(a)). As shown in Fig. 5(a), the presence of strong N₂ ($C^3\Pi_u \rightarrow B^3\Pi_g, B^3\Pi_g \rightarrow A^3\Sigma_u^+$) molecular bands and Ar atomic lines suggests the formation of excited nitrogen species (N_2^*) and metastable Ar species (Ar*).^{29,30} These species create an additional reaction route for CO_2 dissociation (N₂* (or Ar*) + CO₂ \rightarrow CO + O + N₂ (or Ar)), supported by the increased intensity of O atomic lines and CO bands when adding N2 or Ar, which contributes to the enhanced CO_2 conversion. Furthermore, 150 ppm NO_x was detected by Fourier transform infrared (FTIR) in the case of N₂ addition (Fig. S13, ESI^{\dagger}), revealing that N₂ can be regarded as an alternative scavenger of O species.12 The elimination of partial O species can effectively limit the reverse reaction, *i.e.*, O + CO + $M \rightarrow CO_2 + M$, and thus enhance the CO_2 conversion. This could explain why adding N_2 has a stronger promotion on the CO_2 conversion compared with Ar.

Interaction between reactive species in the gas phase and catalyst is also crucial in plasma-catalytic reactions.³¹ Compared to plasma only, packing Mo-doped CeO₂ into the



Fig. 5 (a) Emission spectra of CO₂ plasma, (b) possible pathways of plasma-catalytic CO₂ splitting over V_O-rich catalyst.

(a)

30

2Theta (deg.)

20

discharge zone decreased the current (Fig. S14, ESI⁺), lowering the contribution of the gas-phase reactions. In this case, the CO₂ conversion, however, was still improved, which might be attributed to the interaction between active species and Vo-rich Mo-doped CeO₂, accelerating V_O recovery. Therefore, a possible reaction mechanism is proposed in Fig. 5(b). Firstly, CO₂ is activated to species of CO_2^+ , $CO_{2(v)}$, CO and O radicals as shown in Fig. 5(a) (step 1). Then, the energetic CO_2 -species are adsorbed on the Vo sites to decrease their internal energy (step 2). After that, the V_{O} sites have the potential to grasp the O atom of the adsorbed CO_2 molecule, which weakens the C=O bond, producing adsorbed CO and O (step 3). Subsequently, the adsorbed CO desorbs as the final CO product. While the adsorbed O mainly desorbs from Vo sites in the form of O2 through reacting with the active O radicals produced in the gas phase ($O_g + O_{ad} - V_O \rightarrow O_{2,g} + V_O$), *i.e.*, Eley-Rideal (E-R) mechanism. Meanwhile, Vo sites recover completing the catalytic cycle (step 4). Using isotope trace analysis, we demonstrated the desorption of Nad through an E-R reaction in plasma-catalytic NH₃ decomposition.³¹ Therefore, the desorption of O_{ad} through E-R reaction is also expected. Note that, too strong of a CO_2-V_0 bond makes it easy to split the CO₂ molecule, but the corresponding desorption of adsorbed O is difficult. By contrast, too weak of a CO_2 -V_O bond means it could be hard to split CO_2 , although the adsorbed O can easily desorb from the catalyst surface. Therefore, a catalyst with a proper binding strength between V_O sites and CO_2 benefits the conversion of CO_2 and favors the catalytic cycle.

In conclusion, plasma-catalytic CO2 splitting over M-doped CeO_2 catalysts (M = Fe, Co, Ni, Cu, Cr, V, Mn or Mo) has been investigated. Mo-doped CeO₂ exhibited the best activity; this is attributed to the increased oxygen vacancies created by strong interaction between Mo and CeO₂, as well as the calcining atmosphere being oxygen-deficient and rich in Ar metastable species. Furthermore, oxygen vacancies were stable during the reaction, which is ascribed to the interaction between active O produced in the gas phase and the adsorbed O on the oxygen vacancy site, resulting in desorbing as O2 molecules and recovering oxygen vacancy sites. These findings suggest that introducing proper doping on CeO₂ offers a potential route to tune properties of oxygen vacancy in CeO₂. Additionally, adding N₂ and Ar into the plasma process enhanced the CO₂ conversion, especially when adding N₂. This promotional effect is mainly attributed to the new reaction routes induced by the presence of metastable species. We found that N2 can be used as an O scavenger to forward the chemical equilibrium and inhibit the reverse reaction to form CO₂.

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Conflicts of interest

There are no conflicts to declare.

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