Catalytic wet peroxide degradation of acrylonitrile wastewater by ordered mesoporous Ag/CeO$_2$: synthesis, performance and kinetics

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Ordered mesoporous Ag/CeO$_2$ catalysts have been successfully synthesized by a microwave assisted soft template method. The morphology, structure and chemical composition of the catalyst were characterized by XRD, N$_2$ adsorption–desorption, SEM, EDS, TEM and XPS. The study of catalytic performance and reaction kinetics of organic matter degradation in acrylonitrile wastewater were elucidated by temporal evolution of intermediates and final products detected by GC/MS analysis along with a continuous flow experiment study. The results show that the Ag/CeO$_2$ has an ordered mesoporous structure, the specific surface area is 91.4–118.2 m$^2$ g$^{-1}$ and the average pore size is 12.63–16.86 nm. 0.4-Ag/CeO$_2$ showed the best catalytic performance, the COD removal rate reached 94.6%, which was 30% higher than that of CeO$_2$. The degradation is in accordance with the second-order reaction kinetics of the Arrhenius empirical model and Langmuir–Hinshelwood kinetic model. However the latter fits better, and the linear correlation coefficient $R^2$ is more than 0.98, which describes the adsorption catalytic mechanism of Ag/CeO$_2$. According to the analysis by GC/MS, the organic compounds in acrylonitrile wastewater oxidized into intermediate compounds and other small compounds, then are further oxidized into carbon dioxide and water. The catalytic activity of Ag/CeO$_2$ was the result of the combination of Lewis acid–base position of CeO$_2$ and redox cycle of Ce$^{3+}$/Ce$^{4+}$.

**Introduction**

Acrylonitrile is an important chemical raw material, and its production wastewater is toxic and harmful due to high organic compound concentrations such as those of acrylonitrile, acetonitrile and N-heterocyclic compounds.$^1,2$ At present, the treatment methods of acrylonitrile wastewater include biological methods, distillation methods, incineration methods and catalytic oxidation methods, etc.$^3,5$. The effluent COD after biological treatment is very high due to the low biodegradability of the –CN group. The distillation method has a poor effect on the separation of components with similar boiling points. The cost of incineration is high and the equipment is corroded seriously. The catalytic oxidation method has low catalytic efficiency and serious dissolution of metal ions. Catalytic wet peroxide (CWPO) which uses H$_2$O$_2$ as an oxidant, can effectively degrade organic matter at ambient temperature.$^6$ The design and development of catalysts become the key to improve the removal of pollutants.

Due to the unique valence electron configuration, CeO$_2$ is reduced from Ce$^{4+}$ to Ce$^{3+}$, resulting in the formation of oxygen vacancies on the surface of CeO$_2$, which promote the number of active centers. The ability of store and release oxygen improves the catalytic activity of CeO$_2$.$^7,8$ The redox cycle between Ce$^{3+}$ and Ce$^{4+}$ is similar to that of Fe$^{2+}$/Fe$^{3+}$ in Fenton catalytic oxidation.$^{9,10}$ In addition to redox, CeO$_2$ also has unique acid–base properties. CeO$_2$ is a metal oxide with Lewis acid and base sites, which is a typical acid–base catalyst.$^{11}$ In the past two decades, many nanocomposites containing CeO$_2$ have been prepared, it can be divided into the following categories, metal/CeO$_2$, metal oxide/CeO$_2$, CeO$_2$/support, such as Co/CeO$_2$, Ni/CeO$_2$, Au/Cu/CoO,$^{16}$ MnO$_2$/CeO$_2$, CuO/CoO,$^{18}$ CeO$_2$/SiO$_2$, CeO$_2$/g-C$_3$N$_4$, CeO$_2$/Al$_2$O$_3$, Ni–La$_2$O$_3$–CeO$_2$/SBA-15,$^{21}$ Ni–CeO$_2$/graphene,$^{22}$ and so on. Metal or metal oxide supported on CeO$_2$ carrier by impregnation method, or doped in the lattice of CeO$_2$ in the form of metal cation to form metal–Ce solid solution, which can improve the catalytic performance of the catalyst.$^{24–26}$ Compared with platinum, palladium, gold and other precious metals, silver is stable and relatively cheaper, which makes silver more practical in industrial applications.$^{25,27}$ In particular, silver nanoparticles have been proved to be promising catalysts, and many excellent catalysts based on silver nanoparticles have been reported, such as Ag/TiO$_2$, Ag/SBA-15,$^{28,29}$ Ag/Al$_2$O$_3$, Ag/g-C$_3$N$_4$ (ref. 34 and 35) and so on. Also some scholars have successfully synthesized Ag/CeO$_2$ catalysts.$^{36–38}$ Most of them were prepared by hydrothermal or coprecipitation methods, and the Ag/CeO$_2$ prepared was used for catalytic
degradation of \( p \)-nitrophenol,\textsuperscript{36} naphthalene,\textsuperscript{39} formaldehyde,\textsuperscript{40} propene,\textsuperscript{41} etc. Therefore, it is necessary to further study the application potential of Ag/CeO\(_2\) in catalytic degradation of organic compounds.\textsuperscript{42}

In this study, we report a soft template method assisted by microwave for preparing of ordered mesoporous Ag/CeO\(_2\) catalyst by a combination of the block copolymer and the redox reaction between Ag\(^+\) and Ce\(^3+\). And its catalytic performance in CWPO system, reaction kinetics and degradation mechanism of organic compounds in acrylonitrile wastewater were systematically studied.

**Materials and methods**

**Materials**

All reagents used were of analytical grade without further purification as follows: triblock copolymer F127 (EO\(_{106}\)PO\(_{70}\)EO\(_{106}\), \( M_\text{av} = 12 \, 600 \), Sigma-Aldrich); cerium nitrate (Ce(NO\(_3\))\(_3\)-6H\(_2\)O, Aladdin), hydrogen peroxide solution (H\(_2\)O\(_2\), 30 wt\%, Sinopharm Chemical Reagent Co., Ltd), silver nitrate (AgNO\(_3\), Sinopharm Chemical Reagent Co., Ltd), sodium hydroxide (NaOH, Sinopharm Chemical Reagent Co., Ltd). Test wastewater sample was taken from a acrylonitrile chemical plant in Fushun, the initial COD value was 1500 mg L\(^{-1}\). The concentration of organic matter in wastewater was expressed by COD index.

**Catalysts preparation**

5 mmol Ce (NO\(_3\))\(_3\)-6H\(_2\)O and 2 mmol AgNO\(_3\) were added into a mixture of 20 mL ethanol and 2 g F127, the pH value was adjusted to 10 by NaOH solution, the mixture was stirred for 60 min at room temperature, then the mixture was reacted in a microwave reactor at 100 °C for 60 min, the gel was dried in oven at 120 °C for 10 hours, then calcined in muffle furnace at 550 °C for 5 h and finally mesoporous Ag/CeO\(_2\) was obtained. During different AgNO\(_3\) dosage, the molar ratio of Ag to Ce is 0.4:1, 0.8:1 and 1.0:1, respectively, which recorded as 0.4-Ag/CeO\(_2\), 0.8-Ag/CeO\(_2\), 1.0-Ag/CeO\(_2\).

In the same way, 5 mmol Ce (NO\(_3\))\(_3\)-6H\(_2\)O was added into the mixed solution of ethanol and F127, the mesoporous CeO\(_2\) was obtained.

**Characterization**

The D8 advance X-ray diffractometer of Brooke spectroscopic instrument company, Germany, using a radiation source Cu (K\(_\alpha\) = 1.54178 nm, 40 kV and 15 mA) with a scanning rate of 10 (°) min\(^{-1}\), continuous scanning mode, wide-angle scanning range is 5–90° and small-angle scanning range is 0.5–8.0°. Autosorb-iQ2-MP automatic physical static analyzer of Cantor instrument company, the liquid nitrogen temperature is 77 K. SU8010 field emission scanning electron microscope of Hitachi company, Japan, with accelerating voltage of 15 kV and working distance of WD = 4 mm. JEM-2100F high resolution transmission electron microscope of Japan Electronics Co., Ltd., accelerating voltage 200 kV. ESCALAB 250Xi photoelectron spectrometer of Thermo Scientific company, USA, uses monochromatic Al target as X-ray source. Agilent 7890a/5975c GC/MS instrument of Agilent Technology Co., Ltd. of the United States, the carrier gas is high-purity nitrogen, HP-5MS column.

**Catalytic wet peroxidation experiments**

1) **Catalytic performance test.** A certain amount of prepared catalyst and H\(_2\)O\(_2\) (30 wt\%) were added into 50 mL of acrylonitrile wastewater sample which COD was diluted to 500 mg L\(^{-1}\). After stirring for 60 min, put the solution into the polytetrafluoroethylene reaction tank, react at a certain temperature for a period of time. After centrifugation, the COD concentration of degraded acrylonitrile wastewater was determined by rapid digestion spectrophotometry.

2) **Reaction kinetics test.** A customized catalyst evaluation system was used as the reaction device. 1 g 0.4-Ag/CeO\(_2\) was added into the reactor to form a fixed bed reactor. At the same time, the dosage of H\(_2\)O\(_2\) and acrylonitrile wastewater was 10–30 mL min\(^{-1}\) controlled by pump and flowmeter. Adjust the reaction temperature from 80 °C to 120 °C, the wastewater flowed out of the reactor and was determined by rapid digestion spectrophotometry and GC/MS.

**Results and discussion**

X-ray diffraction analysis

The phase and crystal structure of the catalyst was investigated by the X-ray diffraction (XRD). The typical XRD patterns of pure CeO\(_2\) and Ag/CeO\(_2\) catalysts is shown in Fig. 1a, the peaks recorded at angles (2\(\theta\)) of 28.6°, 33.2°, 47.6°, 56.6°, 59.4°, 69.8°,
76.9° and 79.3° are corresponding to plane (111), (200), (220) and (311) of CeO$_2$ with face-centered cubic (FCC) structure. While 38.2°, 44.1°, 64.4° and 77.4° are corresponding to plane (111), (200), (220) and (311) of Ag structure. All the peaks in the pattern are indexed for face-centered cubic structure, and the lattice constant calculated from XRD data is closely agreement with Joint Committee of Powder Diffraction Data (JCPDS) (card no: 34-0394 and 65-2871). 43,44 The small angle diffraction pattern of the samples are shown in Fig. 1b. The sharp peak near 0.5° indicates that the sample has good crystallinity and ordered mesoporous structure. The CeO$_2$ meso-structure avoids a severe breakdown throughout the Ag nanoparticles supporting step.

N$_2$ adsorption–desorption analysis

The N$_2$ adsorption–desorption isotherms of the prepared samples is shown in Fig. 2. According to the IUPAC classification, 45 the isotherms belong to IV isotherms, which are typical cage structure characteristics, indicating that the loaded samples have mesoporous structure. There is an obvious H2 type hysteresis in the range of $P/P_0$ between 0.6 and 0.9, which indicating that Ag was successfully loaded on CeO$_2$ without changing the original mesoporous structure of CeO$_2$. The pore size is calculated by BJH model, specific surface area and pore volume are calculated by BET method. The structural parameters of the samples are listed in Table 1. The noticeable change of the porous structure of such Ag/CeO$_2$ samples as compared to CeO$_2$ may be due to the Ag$^{+}$–Ce$^{3+}$ redox reaction during the microwave assisted hydrothermal reaction. 46 The comparison of the porous structure of these catalysts allows the conclusion that Ag is predominantly localized in the pores. With the increase of Ag doping, Ag is accumulated and blocked in the pores, which leading to a considerable decrease of both $S_{BET}$ and $V_p$ values. The specific surface area of 0.4-Ag/CeO$_2$ is larger than that of 0.8-Ag/CeO$_2$. Moreover, low Ag loading can reduce the cost of catalyst. In general, Ag/CeO$_2$ synthesized by microwave-assisted synthesis has large specific surface area and relatively large pore size.

Microstructural analysis

The morphology and element mapping analysis of 0.4-Ag/CeO$_2$ by SEM and EDS are shown in Fig. 3. The Fig. 3a reveals that the mesoporous Ag/CeO$_2$ consists of spheres with diameter of 10–20 nm. The Ag nanoparticles are encapsulated by CeO$_2$ mesostructure and randomly distributed throughout the entire mesoporous CeO$_2$ framework. Fig. 3b–d shows that Ce, O and Ag are uniformly distributed in the samples.

The TEM characterization results of 0.4-Ag/CeO$_2$ is shown in Fig. 4. Fig. 4a clearly reveals that the prepared Ag/CeO$_2$ is composed of nanospheres with 10–20 nm, and the dark parts are Ag particles. Many different lattice fringes can be seen from Fig. 4b, the lattice spacing of CeO$_2$ and Ag can be identified. The fringes with $d = 0.31$ nm matches the CeO$_2$(111) facet, and the fringes with $d = 0.24$ nm matches the Ag(111) facet. In the channel of mesoporous CeO$_2$, Ag$^{+}$ precursors are reduced to silver atoms, which nucleate and grow into silver nanostructures. Ag nanoparticles are encapsulated by CeO$_2$ mesoporous structure and distributed in the whole mesoporous CeO$_2$ framework randomly. The mesoscopic structure of CeO$_2$ avoids the destruction of the framework during the doping process of Ag nanoparticles.

XPS spectra analysis

The XPS analysis results are shown in Fig. 5. As shown in Fig. 5a, Ce, O and Ag are contained in the 0.4-Ag/CeO$_2$, which is consistent with the scanning results of EDS. From Fig. 5b, the two characteristic peaks of Ag 3d$^{3/2}$ and Ag 3d$^{5/2}$ are symmetrically distributed at the binding energies of 374.1 eV and 368.3 eV, respectively. 48,49 This is due to the shift of the peak caused by the charge transfer of Ag, which proves that the formation of metallic Ag NPs (Ag$^0$). From Fig. 5c, the characteristic peaks of Ce$^{4+}$ 3d$^{3/2}$ and Ce$^{4+}$ 3d$^{5/2}$ appeared at 916.3 eV and 897.8 eV, and the characteristic peaks of Ce$^{3+}$ 3d$^{3/2}$ and Ce$^{3+}$ 3d$^{5/2}$ appeared at 901.3 eV and 882.6 eV, respectively. 48 Since the oxygen defect structure of cerium compounds is dynamic and may change spontaneously with changes in physical parameters such as oxygen partial pressure, temperature and the presence of other ions, it is reasonable that Ce$^{3+}$ and Ce$^{4+}$ exist in Ag/CeO$_2$ at the same time. 48,49 Fig. 5d shows the O1s spectrum of the sample with a wide spectrum. The characteristic O1s peaks caused by Ce–O bond and H–O bond are located at 529.6 eV and 531.9 eV, respectively. The binding energy of 529.6 eV is lattice oxygen O$^{2-}$ – Ce$^{4+}$ in Ag/CeO$_2$ sample. With the reduction of
CeO$_2$, the binding energy of lattice oxygen moves to higher position. The binding energy of 531.9 eV is caused by the adsorption of H$_2$O. The interaction between Ag and CeO$_2$ is that Ag species contribute valence electrons to CeO$_2$ as electron donors, and CeO$_2$ is reduced by electron acceptor, and oxygen on CeO$_2$ surface will also reverse flow to Ag nanoparticles. Ag doping increases the oxygen defects in CeO$_2$ lattice and increases the concentration of adsorbed oxygen on the surface. The oxygen defect of CeO$_2$ can stabilize the oxidation state of Ag species and improve the redox ability of Ag/CeO$_2$.

Catalytic wet peroxide performance

0.4-Ag/CeO$_2$, 0.8-Ag/CeO$_2$, 1.0-Ag/CeO$_2$ and CeO$_2$ were selected as catalyst to compare their catalytic performance. The dosage of catalyst was 0.25 g, acrylonitrile wastewater was diluted to COD concentration of 500 mg L$^{-1}$, pH value was 6, H$_2$O$_2$ (30 wt%) dosage was 8 mL, reaction temperature was 100 $^\circ$C, reaction time was 20, 40, 60, 80, 100 and 120 min respectively. The COD removal rate is shown in Fig. 6.

It can be seen from Fig. 6 that CeO$_2$ based catalysts exhibit good catalytic performance, among which 0.4-Ag/CeO$_2$ has the best performance. The COD removal rate reaches 94.6%, which is about 30% higher than that of CeO$_2$ alone. With the increase of Ag doping, the COD removal rate decreases. This is because too much Ag will lead to the reduction of Ag/CeO$_2$ particle size and the formation of aggregates. The aggregates are heavy and dense, which leads to the reduction of active sites on catalyst surface. This is consistent with the N$_2$ adsorption–desorption characterization results.
The circulation of Ce$^{3+}$/Ce$^{4+}$ in CeO$_2$/H$_2$O$_2$ system promotes H$_2$O$_2$ to decomposition of 'OH and O$_2^{-}$ active species to degrade organic compounds.$^{25,33}$ The doping of noble metal Ag makes Ag ions enter the lattice of CeO$_2$ and replace the original cerium atom, resulting in new oxygen vacancies or local lattice distortion.$^{34}$ These effects will cause the transformation of crystal form, which is consistent with the results of XPS analysis. The oxygen vacancy and Ce$^{4+}$ reduction center can be used as the active sites of the reaction, and the increase of oxygen vacancy improves the catalytic activity of the catalyst.

Catalyst reusability study

It is very necessary to evaluate the activity and stability of the catalyst for practical application. The reusability of the catalyst was studied under optimized operating conditions (i.e., dosage of 0.4-Ag/CeO$_2$ was 0.25 g, pH = 6, H$_2$O$_2$ (30 wt%) dosage was 8 mL, temperature was 100 °C, reaction time 120 min and acrylonitrile wastewater was diluted to COD concentration of 500 mg L$^{-1}$). The results are shown in Fig. 7. In the first cycle, maximum removal of COD was reached 94.3%. Herein, after each cycle used catalyst was separated by centrifugation and then washed with deionized water subsequently, catalyst was dried at 120 °C for 6 h, and then the catalyst was calcined in muffle furnace for 2 h in order to remove the adsorbed refractory organic matter. The obtained catalyst was further reused in the next experiment. So as compared with 1st cycle removal efficiency of COD dropped gradually, i.e., 91.5%, 87.2%, 84.8%, 82.3% and 80.1% in 2nd to 6th cycle, respectively. Then, the removal rate dropped further, only 53.6% for the tenth time. Kumar A’s team reported perovskite-like catalyst La$_{0.5}$Ce$_{0.5}$FeO$_3$ synthesized by sol-gel method and peroxide acrylonitrile from aqueous solution showed the acrylonitrile removal efficiency gradually dropped from 90.11% to 69.92% during four cycles.$^{35}$ The drop of COD removal efficiency in each cycle could be attributed to various reasons: such as leaching of active metal ions (Ag or Ce) from the catalyst surface leads to suppress the active sites,$^{36}$ carbon deposition on the catalyst surface that causes inhibited 'OH radicals production$^{46}$ and the collapse of mesoporous structure reduces the active sites.$^{24}$

Reaction kinetics

Due to the complex composition of acrylonitrile wastewater and a variety of intermediate products during the degradation
process, it is impossible to analyze the degradation kinetics of each organic matter separately. Therefore, the degradation kinetics of organic matter was studied by the monitoring the change of COD concentration of acrylonitrile wastewater. The Arrhenius empirical model and Langmuir–Hinshelwood model were used to analyze the degradation process of organic compounds.

**Arrhenius empirical model**

According to the relevant literature, Arrhenius empirical model was adopt, which was assumed that the degradation process of acrylonitrile wastewater by 0.4-Ag/CeO$_2$ conforms to the second-order kinetic equation, then the reaction rate equation is as follow (1):

$$\frac{d[C]}{dt} = A \exp \left( \frac{E_a}{RT} \right) [C]^a [O]^b$$  (1)

where: $A$ is the pre-exponential factor; [C] is the concentration of COD, mg L$^{-1}$; $E_a$ is the activation energy of the reaction, kJ mol$^{-1}$; $R$ is the gas constant; [O] is the concentration of oxidant, mol L$^{-1}$; $t$ is the reaction time, s; $a$ and $b$ are the reaction order.

The oxidation process is only controlled by the concentration of reactants and reaction temperature. It is assumed that the oxidant H$_2$O$_2$ is excessive and can be approximately regarded as a constant. Then eqn (1) can be simplified as follow (2):

$$\frac{d[COD]}{dt} = k[COD]^2$$  (2)

For a continuous reactor, the two sides of the equation can be integrated between $t_1 = 0$, $t_2 = t$, then eqn (3) obtained.

$$\frac{1}{[COD]} - \frac{1}{[COD]_0} = kt$$  (3)

where: [COD] is the COD concentration of solution after reaction $t$ min, mg L$^{-1}$; [COD]$_0$ is the initial concentration of organic matter, mg L$^{-1}$.

The relationship between COD removal rate and reaction time is shown in Fig. 8, Which the experimental were taken at 333, 353 and 373 K, pH = 6, H$_2$O$_2$ dosage was 30 mL L$^{-1}$.

According to the test data, plot $T$ with $1/[COD] - 1/[COD]_0$, and the results are shown in Fig. 9. The reaction time was controlled by adjusting the wastewater flow rate, when the flow rate of 10, 15, 20, 25, 30 and 35 mL min$^{-1}$ corresponding to the reaction time of 43, 26, 21, 17, 14 and 12 min respectively.

The relevant parameters after linear fitting at different temperatures are shown in Table 2. It can be seen from Table 2 that the linear correlation coefficient $R^2$ of the reaction kinetic equation at different temperatures is more than 0.92, which indicates that the process of CWPO degradation of organic matters in acrylonitrile wastewater by mesoporous Ag/CeO$_2$ conforms to the second-order reaction kinetics of empirical model. The reaction rate constants increase with the increase of temperature, which is consistent with the change of COD removal rate under the same conditions.

**Langmuir–Hinshelwood model**

The reaction kinetics analysis was also carried out through Langmuir–Hinshelwood mechanism of heterogeneous catalytic oxidation, which assumes that the reaction between organic matter and hydrogen peroxide is initiated on the surface of catalyst.$^3$ Many researchers have performed Langmuir–
Hinshelwood, Freundlich and Temkin isotherms for heterogeneous catalytic oxidation.27-41

According to the Langmuir–Hinshelwood mechanism, the heterogeneous catalytic reaction process can be divided into three steps: first, organics and H\textsubscript{2}O\textsubscript{2} molecules are adsorbed on the surface of the catalyst; second, the reaction between reactants to form products; third, the product desorbed from the catalyst surface.42 Based on the Langmuir–Hinshelwood reaction kinetic model, it is assumed that the organic matter in acrylonitrile wastewater reacted with H\textsubscript{2}O\textsubscript{2} on the surface of the catalyst.

As shown in formula (4), the rate \( R \) of heterogeneous catalytic reaction is directly proportional to the surface fraction covered by organic matter and H\textsubscript{2}O\textsubscript{2} molecule in acrylonitrile wastewater.

\[
R = K_s S_{\text{COD}} S_{\text{H}_2\text{O}_2}
\]  

(4)

The competitive and noncompetitive adsorption rates of organics in acrylonitrile wastewater and H\textsubscript{2}O\textsubscript{2} on the catalyst surface can be expressed as formula (5).

\[
R = \frac{K_{\text{ACN}}[\text{COD}]K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2]}{(1 + K_{\text{ACN}}[\text{COD}] + K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2] + K_a[\text{P}])^2}
\]  

(5)

where: \( K_{\text{ACN}} \), \( K_{\text{H}_2\text{O}_2} \), \( K_a \) are adsorption equilibrium constants of organics, H\textsubscript{2}O\textsubscript{2} and product respectively. [COD] is the equilibrium concentration of organic matter in acrylonitrile wastewater.

In heterogeneous catalytic reactions, adsorption and desorption occur fast at equilibrium. Therefore, the concentration of product formation [P] in formula (5) is ignored, and the above equation is simplified to obtain formula (6):

\[
R = \frac{K_{\text{ACN}}[\text{COD}]K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2]}{(1 + K_{\text{ACN}}[\text{COD}] + K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2])^2}
\]  

(6)

Because of the H\textsubscript{2}O\textsubscript{2} is excessive, it can be considered as a constant. Eqn (6) can be simplified by arranging the constant terms together to obtain eqn (7).

\[
R = \frac{ab[\text{COD}]}{(c + b[\text{COD}])^2}
\]  

(7)

where \( a \), \( b \) and \( c \) are \( K_aK_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2] \), \( K_{\text{ACN}} \) and \( (1 + K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2]) \), respectively.

For the non competitive adsorption process of organics and H\textsubscript{2}O\textsubscript{2}, the expression of reaction rate can be written as formula (8).

\[
R = \frac{K_{\text{ACN}}[\text{COD}]K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2]}{(1 + K_{\text{ACN}}[\text{COD}])(1 + K_{\text{H}_2\text{O}_2}[\text{H}_2\text{O}_2])}
\]  

(8)

For a constant concentration of H\textsubscript{2}O\textsubscript{2}, expression (8) can be written as a linear Langmuir isotherm as formula (9).

\[
R(Q_e) = \frac{ab[\text{COD}]}{(c + b[\text{COD}])^2}
\]  

(9)

where: \( Q_e \) is the equilibrium amount of adsorbate in the adsorbent (mg g\textsuperscript{-1}). Eqn (9) is simplified to linear equation to form formula (10).

\[
\frac{[\text{COD}]}{Q_e} = \frac{[\text{COD}]}{a} + \frac{1}{ab}
\]  

(10)

The Langmuir equilibrium constant \( \text{K}_{\text{ACN}} \) is derived from the curves of \([\text{COD}]Q_e\) and \([\text{COD}]\) at 333 K, 353 K and 373 K, as shown in Fig. 10. The isotherm constants and \( R^2 \) values are listed in Table 2. It can be seen from Table 2 that the linear correlation coefficient \( R^2 \) is above 0.98. By comparing the linear correlation coefficient \( R^2 \) between Langmuir–Hinshelwood model and Arrhenius model, it is concluded that Langmuir–Hinshelwood model is more consistent with the CWPO reaction process. In the Ag/CeO\textsubscript{2}/H\textsubscript{2}O\textsubscript{2} system, Ce\textsuperscript{3+} reacts with H\textsubscript{2}O\textsubscript{2} to produce \cdot\text{OH} and peroxide species, which degrade the organic compounds adsorbed on the surface of CeO\textsubscript{2}.

**Degradation pathway and mechanism**

Under the premise of catalyst adsorption equilibrium, the water samples with reaction time of 0 (raw water), 17, 21, 28 and

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**Table 2  Fitting parameter of dynamic equation**

| Model                        | Temperature/K | \( R^2 \) | Linear equation               |
|-----------------------------|---------------|------------|-------------------------------|
| Arrhenius empirical model   | 373           | 0.9326     | \( y = 7.221 \times 10^{-5}x - 2.530 \times 10^{-4} \) |
|                             | 353           | 0.9279     | \( y = 7.053 \times 10^{-5}x - 2.127 \times 10^{-4} \) |
|                             | 333           | 0.938      | \( y = 6.745 \times 10^{-5}x - 1.832 \times 10^{-4} \) |
| Langmuir–Hinshelwood model  | 373           | 0.9859     | \( y = 0.2367x - 138.96 \) |
|                             | 353           | 0.9861     | \( y = 0.2525x - 155.49 \) |
|                             | 333           | 0.9921     | \( y = 0.2719x - 176.57 \) |

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**Fig. 10  Fitting results of organics adsorption in Langmuir–Hinshelwood model.**
43 min were taken for GC/MS determination respectively. The cluster analysis of the determination results is shown in Table 3.

According to the GC/MS analysis results, 40 kinds of organic matter were detected, the main organic compounds in acrylonitrile wastewater are nitriles, such as 3-cyanopyridine, cis-1,2-dicyanitrile cyclobutane and succinonitrile; phenolic compounds, such as 4-methylphenol, 4-methoxyphenol, 4-(1,1,3,3-tetramethylbutyl)phenol, etc. Nitriles and phenols accounted for about 80% of the organic matter content in the raw wastewater. According to the changes of the number and content of various kinds of organic matter in Table 3, the degradation trend diagram of organic matter is shown in Fig. 11 and transformation paths of various organic matter is shown in Fig. 12.

According to Fig. 11, the content of nitriles and phenolic compounds decreased significantly, the degradation rate of nitrile compounds is 86.9%, and the degradation rate of phenolic compounds is 96.2%. The results show that Ag/CeO₂ catalyst can effectively degrade nitriles and phenols in CWPO system. The degradation trends of amino compounds, alkanes, organic acids and esters are generally opposite to that of nitriles and phenols. It is possible that phenol may be converted to alkanes and organic esters, while nitrile may be converted to amino compounds during CWPO degradation. Combined with the data in Table 3, it can be seen that after 43 minutes of reaction, the content of alkanes gradually increased to 39 times of raw water, and the content of organic acids and esters reached 49 times of raw water. The content of N-containing heterocycles increased sharply at 28 min, while the content of nitriles is very low at this time, indicating that N-containing heterocycles are transformed from nitriles. The low content of alcohols, ketones and ethers may be due to the fact that they are converted from phenols and then oxidized to acids by OH. A small amount of sulfur compounds in raw water were oxidized to thioether. It can be inferred that, under the catalysis of Ag/CeO₂, the organic matter undergoes a chemical reaction in the CWPO system. These organic compounds oxidized into intermediate compounds and other small compounds, then these intermediate compounds get further oxidized into carbon.

### Table 3  Cluster analysis degraded organics

| Reaction time/min | Organics | Nitriles | Phenols | N-containing heterocycles | Amino compounds | Alcohol ketone ethers | Alkanes | Organic acids and esters | Sulfur compounds |
|-------------------|----------|----------|---------|---------------------------|----------------|----------------------|---------|--------------------------|----------------|
| 0                 | Number   | 13       | 3       | 6                         | 4              | 2                    | 5       | 4                        | 3              |
|                   | Content% | 47.57    | 33.01   | 4.51                      | 5.04           | 1.33                 | 0.9     | 0.57                     | 7.07           |
| 17                | Number   | 12       | 2       | 3                         | 2              | 4                    | 10      | 1                        | 0              |
|                   | Content% | 41.52    | 40.83   | 4.97                      | 0.84           | 4.71                 | 6.9     | 0.24                     | 0              |
| 21                | Number   | 7        | 4       | 2                         | 3              | 1                    | 6       | 3                        | 0              |
|                   | Content% | 23.76    | 15.80   | 4.48                      | 10.64          | 8.37                 | 26.96   | 9.98                     | 0              |
| 28                | Number   | 9        | 1       | 8                         | 3              | 0                    | 8       | 3                        | 0              |
|                   | Content% | 22.29    | 1.51    | 28.01                     | 13.19          | 0                    | 27.01   | 7.97                     | 0              |
| 43                | Number   | 4        | 2       | 4                         | 4              | 2                    | 6       | 6                        | 0              |
|                   | Content% | 6.23     | 1.90    | 7.51                      | 19.91          | 1.12                 | 35.26   | 28.07                    | 0              |

![Fig. 11](image1.png)  
Trend diagram of organic degradation.

![Fig. 12](image2.png)  
Diagram of organic matter organic transformation path.
dioxide and water, which is consistent with the conclusion of Kumar A’s study.55

The Ag/CeO2 exhibits strong catalytic activity, which is due to the activation of electrophilic and nucleophilic by Lewis acid center (Ce4+) and acid–base pair (Ce–O), respectively. However, 3-cyanopyridine and 4-cyanopyridine did not transform during the degradation and transformation of nitriles, indicating that Ag/CeO2 has no activity against these two compounds. The above analysis shows that Ag/CeO2 catalyst not only promotes H2O2 to produce more ‘OH, but also promotes the chemical conversion between organic compounds in acrylonitrile wastewater. The catalytic activity of Ag/CeO2 is the result of weak Lewis acid, strong basicity and unique redox properties of CeO2.

Conclusions

In summary, the ordered mesoporous Ag/CeO2 with specific surface area of 91.4–118.2 m² g⁻¹ and pore diameter of 12.63–16.86 nm was successfully prepared by microwave-assisted soft template method. The CWPO degradation test was carried out with 0.4-Ag/CeO2 as the catalyst, and the COD removal rate reached 94.6%. Ag doping improved the catalytic activity of CeO2 and achieved good stability. The degradation process of organic matter with Ag/CeO2 as catalyst conformed to the Langmuir model, which describes the adsorption catalytic mechanism of Ag/CeO2 in detail. The catalytic activity of Ag/CeO2 is the result of the interaction of weak Lewis acid, strong alkalinity and unique redox properties of CeO2, which provides theoretical basis for the application of CeO2 based catalyst in CWPO system of other complex organic industrial wastewater.

Conflicts of interest

The authors declare that they have no conflict of interest.

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