PAPER

Check for updates

Electrocatalytic carboxylation of halogenated compounds with mesoporous silver electrode materials†

Si-Li Shan, Cheng-Jie Jiang, Yu-Ting Liu, Jing-Jie Zhang, Huan Wang* and Jia-Xing Lu [ID] *

Mesoporous silver materials are used as electrocatalysts for halogenated compounds. The mesoporous silver materials have uniform mesoporous size (8 nm), large specific surface area (12 m^2 g^{-1}), high pore volume (0.07 cm^3 g^{-1}), and a good 3D network structure of the metallic silver skeleton. The results show that the prepared materials exhibit high performance in electrocatalytic carboxylation of halogenated compounds to acid (78%).

The main factor of global warming and the greenhouse effect is the increase of concentration of carbon dioxide (CO_2).[1,2] There are two ways to solve the problem of CO_2 concentration. The first method is to establish a low-carbon economic development model. The second method is to convert CO_2 into useful organic chemicals.[3,4] As human production and life are inseparable from the use of fossil fuels, the problem of CO_2 emissions will still exist. Therefore, today's society urgently pays attention to how to effectively transform and utilize CO_2; in addition, the conversion of CO_2 into productive products has attracted widespread attention.[5,6] CO_2 conversion is a challenge because of its high chemical and thermodynamic stability. Through electrochemical means, CO_2 can be activated at normal temperature and pressure.[7] Therefore, from the perspective of environmental protection and technical means, electro-carboxylation is a simple and easy method for CO_2 immobilization. Electrocatalytic carboxylation can be applied to a variety of substrates, such as olefins,[8] alkynes,[9] alcohols,[10] aldehydes,[11] ketones,[12] epoxides,[13] imines[14] and organic halides.[15] Halogenated compounds are toxic, carcinogenic and difficult to biodegrade. Electrocarboxylation of halogenated compounds not only immobilizes CO_2 and reduces environmental pollution, but also converts them into useful compounds. More importantly, several of them are fine chemicals with industrial use, and some of them can produce anti-influenza drugs, such as benzoic acid, sodium benzoate, benzyl benzoate *etc*.[16]

Silver shows remarkable electrocatalytic activity when it is used for electrocarboxylation of organic halides.[17] In addition, silver nanoparticles have been used in electrocatalytic reactions.

In the field of electrocatalysis using silver nanoparticles, researchers have made great efforts. Firstly, a silver nanoparticle-modified electrode was successfully prepared and applied to electrocatalytic reduction of benzyl chloride.[18] At the same time, through the electrochemical detection method, the catalytic activity of the nano silver modified electrode was compared with the silver electrode, and the catalytic activity of the modified electrode after repeated use was discussed. Simonet *et al.* electrodeposited silver nanoparticles on glassy carbon electrode, and explored a variety of organic halogenated compounds to study the universality of nano silver-modified electrodes in electrocatalytic reduction of organic halides.[19]

On the other hand, numerous research efforts have been reported to control the morphology of metal nanoporous materials to obtain optimized activity in electrocatalysis. Mesoporous silver material has the advantages of large specific surface area, large pore volume, regular mesoporous structure, good chemical inertia, good ductility, good thermal stability and good conductivity, so it has been widely concerned. These properties are beneficial to heterogeneous catalytic reactions and other applications, such as adsorption/separation, fuel cells and biosensors.[20] However, mesoporous silver materials are mainly used to enhance the reproducibility and sensitivity of Raman scattering signals,[21] and are rarely used in electrocarboxylation. How to improve the yield of electrocarboxylation by using mesoporous silver electrode and expand the application range of mesoporous silver materials is worthy of our efforts.

In this study, we used hard template method to synthesize mesoporous silver for electrocatalytic carboxylation of halogenated compounds. Different mesoporous silver materials are obtained by adjusting the amount of silver nitrate solution.

The experimental results show that mesoporous silver with different mesoporous pore sizes has different catalytic

Shanghai Key Laboratory of Green Chemistry and Chemical Processes, School of Chemistry and Molecular Engineering, East China Normal University, Shanghai, 200062, China. E-mail: hwang@chem.ecnu.edu.cn; jxlu@chem.ecnu.edu.cn

Table 1 Structure parameters for mesoporous Ag

Entry	Material	BET surface area[a] ($m^2\ g^{-1}$)	Total pore volume[b] ($cm^3\ g^{-1}$)	Average pore size[c] (nm)
1	mesoAg-1	4	0.04	79.5
2	mesoAg-2	3	0.04	68.9
3	mesoAg-3	12	0.07	8.0
4	mesoAg-4	12	0.06	7.1
5	mesoAg-5	9	0.07	145.1

[a] BET surface areas calculated from the N_2 adsorption branches in the range of relative pressure $(p/p_0) = 0.05$–0.20. [b] Total pore volumes measured at $p/p_0 = 0.99$. [c] Pore diameters obtained from N_2 adsorption branches by BJH method.
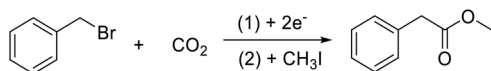
Scheme 1 Electrocarboxylation of benzyl bromide under CO_2 atmosphere.

materials, and the specific surface area and pore volume of mesoAg-3 are the largest. With the decrease of the mass of silver nitrate solution added in the preparation process, the specific surface area of the material first increases and then decreases. The surface of the material is an important place for the reaction, the mesoporous structure has a large specific surface area and pore volume, which can provide a large space for the reaction to be fully carried out, so the catalytic efficiency may be increased. Therefore, it can be inferred that mesoAg-3 has the highest catalytic efficiency.

The electrocarboxylation of halogenated compounds is to break the carbon halide bond in the halide by electrochemical method, so as to remove the halogen atom, and then react with CO_2 to form corresponding carboxylic acid compounds. Now we have been able to carboxylate a variety of organic halides, such as halogenated benzyl, by electrocatalytic method. In order to explore the effect of materials on electrocatalytic carboxylation

of halogenated compounds, benzyl bromide was used as a model compound (Scheme 1).

The yields of electrocarboxylation of benzyl bromide using different mesoporous silver are shown in Table 2. The yield of methyl phenylacetate was 17% on a flat silver electrode (Table 2, entry 1). 78% and 69% yields can be obtained on mesoAg-3 and mesoAg-4, respectively, which is 4.6–4.1 times higher than that on flat silver electrode (Table 2, entries 4 and 5). Using mesoAg-1, mesoAg-2 and mesoAg-5, the yields were 25%, 33% and 34%, respectively (Table 2, entries 2, 3 and 6). The above data show that electrocatalytic properties of the five mesoporous silver materials are better than that of the flat silver, and the yields on mesoAg-3 and mesoAg-4 are higher than those on mesoAg-1, mesoAg-2 and mesoAg-5. MesoAg-3 is the best among the five mesoporous silvers. It is known that the specific surface area and pore volume of mesoAg-3 and mesoAg-4 materials are larger than those of other three materials. These two materials can provide larger space for the full reaction in the catalytic process, so their catalytic efficiency is higher than those of other three materials.

For comparison, the other two silver materials, silver nanoparticles and foam silver, were tested (Table 2, entries 7–8). The yields on the two materials were 36% and 30%, respectively. From Table S2,† the specific surface area of silver nanoparticles

Table 2 The yield of methyl phenylacetate at different mesoporous silver materials[a]

Entry	Substrate	Cathode	Q ($F\ mol^{-1}$)	Yield[b] (%)
1	Benzyl bromide	Silver plate	2	17
2	Benzyl bromide	mesoAg-1	2	25
3	Benzyl bromide	mesoAg-2	2	33
4	Benzyl bromide	mesoAg-3	2	78
5	Benzyl bromide	mesoAg-4	2	69
6	Benzyl bromide	mesoAg-5	2	34
7	Benzyl bromide	Silver nanoparticles	2	36
8	Benzyl bromide	Foam silver	2	30
9	Benzyl bromide	mesoAg-3	1.5	44
10	Benzyl bromide	mesoAg-3	1.75	59
11	Benzyl bromide	mesoAg-3	2.25	78
12	Benzyl bromide	mesoAg-3	2.5	76
13	2-Phenylbromomethyl benzene	mesoAg-3	2	25
14	3-Phenylbenzyl bromide	mesoAg-3	2	36
15	4-Bromomethylbiphenyl	mesoAg-3	2	46

[a] General conditions: c(substrate) = 0.1 mol L^{-1}, DMF = 15 mL, c(TEABr) = 0.1 mol L^{-1}, j = 2 mA cm^{-2}, T = 25 °C, P_{CO_2} = 1 atm, Mg as anode, electric charge = 2.0 F mol^{-1}. [b] GC yield.

and foam silver is 1.94 and 0.04 m^2 g^{-1}, respectively. As described above, the yield on the mesoAg-3 material is 78% and the specific surface area of mesoAg-3 material is 12 m^2 g^{-1}. The specific surface area of mesoAg-3 is the largest, and the specific surface area of foam silver is the smallest. Correspondingly, the catalytic efficiency of mesoAg-3 is the highest, and the catalytic efficiency of foamed silver is the lowest. This shows that a material with a small specific surface area cannot provide enough space for the reaction and has low catalytic efficiency.

Then the influence of electric charge amount was studied (Table 2, entries 4, 9–12). When the charge was in the range of 1.5–2 F mol^{-1}, the yield of methyl phenylacetate was increased with the increase in charge. When the charge exceeded 2 F mol^{-1}, the yield of methyl phenylacetate remained 78%. Moreover, the electrocarboxylation yields of other aromatic halides, 2-phenylbromomethyl benzene, 3-phenylbenzyl bromide and 4-bromomethylbiphenyl, were 25%, 36% and 46%, respectively (Table 2, entries 13–15). This indicates that mesoAg-3 electrode can effectively electrocatalytic carboxylation of aromatic halides.

After the reaction, the composite electrode was cleaned with solvent and used again in the next step under the same conditions. As shown in Fig. 4, even after 7 runs, the activity of the mesoporous silver materials does not decrease significantly, and the yield of methyl phenylacetate can be maintained at about 70%. In order to further study the stability of mesoAg-3 electrode, it was characterized by SEM after 7 times of use. As shown in Fig. 5, some silver particles in the used mesoAg-3 material are slightly larger than those before electrolysis. This may be due to the instability of the silver outside the template, it will agglomerate in the electrolysis process. The N$_2$ adsorption desorption isotherms and the corresponding BJH pore size distribution curves of mesoAg-3 electrode materials were shown in Fig. S4.† According to the N$_2$ adsorption desorption isotherms, there is obvious IV adsorption isotherm and H1 hysteresis loop in the specific pressure range of 0.45–0.75 before and after reuse, indicating that the material still has mesoporous structure after repeated use for 7 times. According to the BJH diagram, the peak appears at 8 nm before and after reuse, which indicates that the material is still a typical mesoporous material after 7 times of use. It can be seen from the data in Table S3† that the specific surface area of the material after reuse decreases. Due to the slight agglomeration, the specific
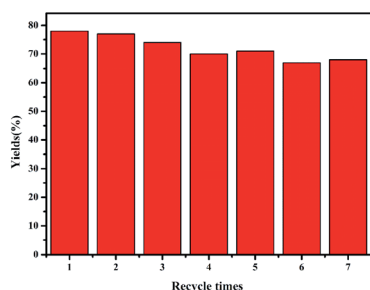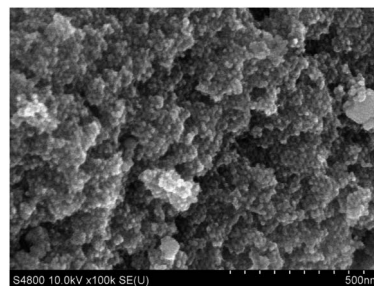

Fig. 5 SEM characterization of mesoporous silver cathode after bulk electrolysis.

surface area of the material after reuse decreases, resulting in a slight decrease in the electrolytic yield.

Conclusions

Electrocarboxylation of benzyl bromide can be achieved by using mesoporous silver for the first time under galvanostatic conditions. Through a series of experiments, it is found that the mesoporous silver prepared by adding appropriate amount of AgNO$_3$ solution has the most regular structure and the highest electrocatalytic performance. When the amount of AgNO$_3$ solution is too much or too little, the mesoporous structure of the prepared materials is stacked, and the electrocatalytic performance is lower than that of the former. The electrocatalytic effect of mesoporous silver was studied under different electrolytic charge and reactants. After 7 runs, the activity of the mesoporous silver materials does not decrease significantly, and the yield of methyl phenylacetate can be maintained at about 70% which indicates that mesoAg-3 electrode has the advantages of good stability.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Notes and references

1 C. M. Sánchez-Sánchez, V. Montiel, D. A. Tryk, *et al.*, *Pure Appl. Chem.*, 2001, **73**, 1917–1927.
2 M. Aresta and A. Dibenedetto, *Catal. Today*, 2004, **98**, 455–462.
3 T. Sakakura, J. C. Choi and H. Yasuda, *Chem. Rev.*, 2007, **107**, 2365–2387.
4 M. Mikkelsen, M. Jørgensen and F. C. Krebs, *Energy Environ. Sci.*, 2010, **3**, 43–81.
5 C. T. Dinh, T. Burdyny, M. G. Kibria, *et al.*, *Science*, 2018, **360**, 783–787.

Fig. 4 Reuse of mesoporous silver cathode for electrocarboxylation of bromobenzyl.

6 S. Gao, Y. Lin, X. Jiao, Y. Sun, Q. Luo, W. Zhang, D. Li, J. Yang and Y. Xie, *Nature*, 2016, **529**, 68–71.

7 B. L. Chen, H. W. Zhu, Y. Xiao, Q. L. Sun, H. Wang and J. X. Lu, *Electrochem. Commun.*, 2014, **42**, 55–59.

8 S. N. Steinmann, C. Michel, R. Schwiedernoch, M. Wu and P. Sautet, *J. Catal.*, 2016, **343**, 240–247.

9 H. Wang, G. Zhang, Y. Liu, Y. Luo and J. Lu, *Electrochem. Commun.*, 2007, **9**, 2235–2239.

10 G. Q. Yuan, H. F. Jiang and C. Lin, *Tetrahedron*, 2008, **64**, 5866–5872.

11 M. Feroci, M. A. Casadei, M. Orsini, L. Palombi and A. Inesi, *J. Org. Chem.*, 2003, **68**, 1548–1551.

12 A. P. Doherty, *Electrochim. Acta*, 2002, **47**, 2963–2967.

13 A. Gennaro, A. A. Isse, J. M. Saveant, M. G. Severin and E. Vianello, *J. Am. Chem. Soc.*, 1996, **118**, 7190–7196.

14 K. H. M. Cardozo, R. Vessecchi, G. Ricardo, *et al.*, *J. Brazil. Chem. Soc.*, 2009, **20**, 1625–1631.

15 U. Hess and R. Thiele, *J. Prakt. Chem.*, 1982, **324**, 385–399.

16 P. P. Luo, Y. T. Zhang, *et al.*, *Catalysts*, 2017, **7**, 274.

17 A. A. Isse, L. Falciola, P. R. Mussini and A. Gennaro, *Chem. Commun.*, 2006, **3**, 344–346.

18 A. A. Isse, S. Gottardello, C. Maccato and A. Gennaro, *Electrochem. Commun.*, 2006, **8**, 1707–1712.

19 J. Simonet, *J. Appl. Electrochem.*, 2009, **39**, 1625–1632.

20 D. J. Guo and Y. Ding, *Electroanalysis*, 2012, **24**, 2035–2043.

21 C. F. Tian, J. Li and C. S. Ma, *Nanoscale*, 2015, **7**, 12318–12324.

22 J. K. Shon, S. S. Kong, J. M. Kim, C. H. Ko, *et al.*, *Chem. Commun.*, 2009, **45**, 650–652.