ARTICLE IN PRESS

Chemical Engineering Journal xxx (xxxx) xxxx



Contents lists available at ScienceDirect

Chemical Engineering Journal



journal homepage: www.elsevier.com/locate/cej

Excellent performance of porous carbon from urea-assisted hydrochar of orange peel for toluene and iodine adsorption

Kangxin Xiao^a, Huan Liu^{a,b,*}, Yang Li^a, Guangyan Yang^a, Yijie Wang^a, Hong Yao^a

^a State Key Laboratory of Coal Combustion, School of Energy and Power Engineering, Huazhong University of Science and Technology, Wuhan 430074, China ^b Department of New Energy Science and Engineering, School of Energy and Power Engineering, Huazhong University of Science and Technology, Wuhan 430074, China

HIGHLIGHTS

G R A P H I C A L A B S T R A C T

- Highly porous carbon was obtained by activation of urea-assisted hydrochar.
 Highly porous carbon possessed ultra-
- high surface area up to $3053 \text{ m}^2/\text{g}$.
- N-containing groups involving activation of hydrochar facilitated the porosity.
- Good liner relation of between variations of surface area and N content was found.
- Highly porous carbon showed excellent capacities for toluene and iodine adsorption.

ARTICLE INFO

Keywords: Orange peel waste Urea-assisted hydrochar Porous carbon Nitrogen-containing group Toluene Iodine



ABSTRACT

In order to promote the physicochemical properties and the pollutants adsorption capacities of porous carbon, a novel urea-assisted hydrothermal method was developed to produce N-doped hydrochar as an ideal precursor from orange peel waste. Different from the modification technologies for nitrogen doping, the highly porous carbon prepared by KOH activation of the urea-assisted hydrochar contained extremely low N content. But such sample showed ultra-high specific surface area up to $3053 \text{ m}^2/\text{g}$, with 63% and 39% higher than the porous carbons through conventional N-free method and hydrochar-urea mixing activation method respectively. In-depth study revealed that during 600-800 °C activation procedure, the N-containing groups including pyridinic-N, pyridonic-N and graphitic-N in the urea-assisted hydrochar reacted sufficiently with K₂CO₃, an intermediate of KOH activation, and generated NH₃, creating richer porous structures in carbon. Particularly, there was a good linear relationship between the relative increment of specific surface area and the relative reduction of N content in porous carbon when the activation temperature increasing. This connection can be served as a reference to increase the porosity of porous carbon through the activation of N-doped carbonaceous precursor. As for pollutants adsorption, the highly porous carbon from urea-assisted hydrochar possessed excellent toluene adsorption capacity up to 724 mg/g and iodine adsorption capacity up to 2252 mg/g.

1. Introduction

Porous carbon materials have shown great potentials for pollution control in air and water environment, due to their large surface area, structure tenability and excellent thermal stability [1–3]. Generally, the high cost of porous carbon is one of the major challenges for its commercial applications [4]. Therefore, organic solid waste with rich carbon and low inorganic contents (e.g. fruit peel and coconut shell)

E-mail address: huanliu@hust.edu.cn (H. Liu).

https://doi.org/10.1016/j.cej.2019.122997

Received 29 August 2019; Received in revised form 26 September 2019; Accepted 27 September 2019 1385-8947/ @ 2019 Elsevier B.V. All rights reserved.

^{*} Corresponding author at: State Key Laboratory of Coal Combustion, School of Energy and Power Engineering, Huazhong University of Science and Technology, Wuhan 430074, China.