

Supporting Information

Carbon dots up-regulate heme oxygenase-1 expression towards the acute lung injury therapy

Bo Wang^{a,§}, Peipei Liu^{a,§}, Hui Huang^{a,*}, Xiting Wang^a, Mengling Zhang^a, Jian Huang^b, Fang Lu^{c,*}, Jian Chen^{a,d*}, Yang Liu^a and Zhenhui Kang^{a,c,*}

^a Institute of Functional Nano & Soft Materials (FUNSOM), Jiangsu Key Laboratory for Carbon-Based Functional Materials & Devices, Soochow University, 199 Ren'ai Road, Suzhou, 215123, Jiangsu, China.

^b School of Biology & Basic Medical Science, Soochow University, 199 Ren'ai Road, Suzhou, 215123, Jiangsu, China.

^c Macao Institute of Materials Science and Engineering, Macau University of Science and Technology, Taipa 999078, Macau SAR, China.

^d Chinese Institute for Brain Research, Research Unit of Medical Neurobiology, Chinese Academy of Medical Sciences, Beijing, 102206, China.

^e School of Life Sciences, Beijing University of Chinese Medicine, Beijing 100029, China.

§ These authors contributed equally to this work.

*E-mail: hhuang0618@suda.edu.cn; lufang@bucm.edu.cn; chenjian@cibr.ac.cn;

zhkang@suda.edu.cn

1. Experiments section

Synthesis of CDs

50 g L-ascorbic acid powder was dissolved in 1 L ultrapure water and then the two electrodes (graphite rod, 99.99%, Alfa Aesar Co. Ltd.) were applied with 0.1 A direct current power supply. After 3 weeks, the solution was filtered with slow-speed quantitative filter paper and then the product was dialyzed in a 500 Da dialysis bag against deionized water to remove excess L-ascorbic.

2. Additional figures

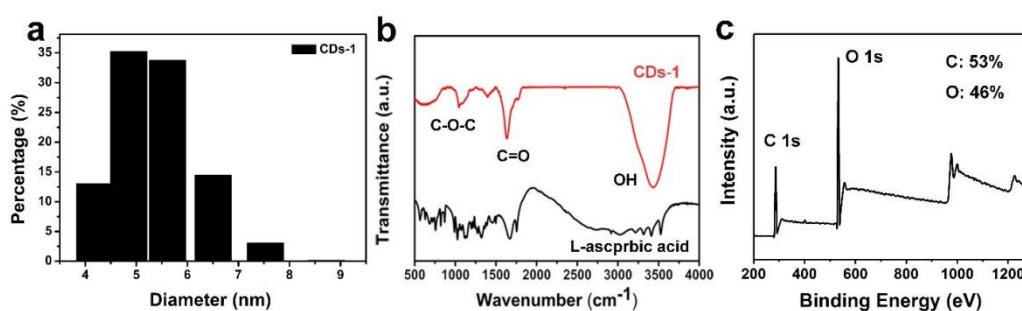


Figure S1. (a) Diameter distributions of CDs-1. (b) FTIR spectra of CDs-1 and L-ascorbic acid (red and black traces, respectively). (c) XPS full scan spectrum of CDs-1.

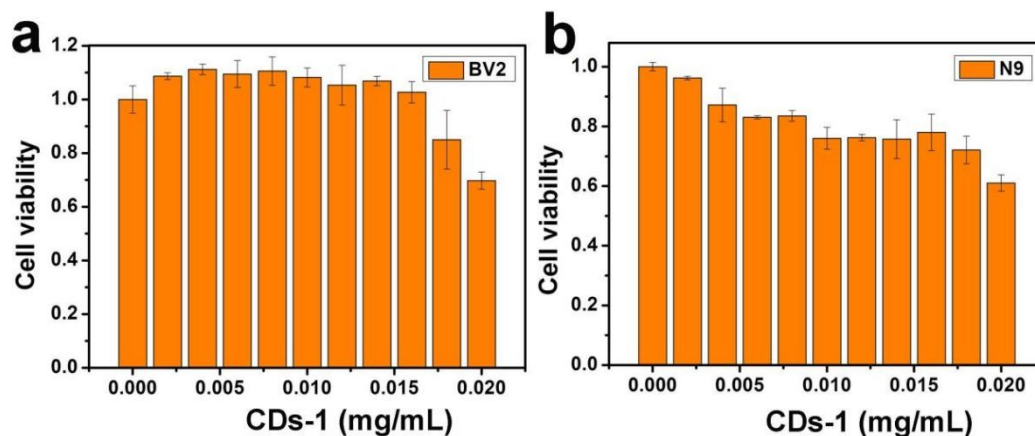


Figure S2. Viability of BV2 (a) and N9 (b) cells after 24 h treatment with different concentrations of CDs-1 as calculated from CCK-8 assay.

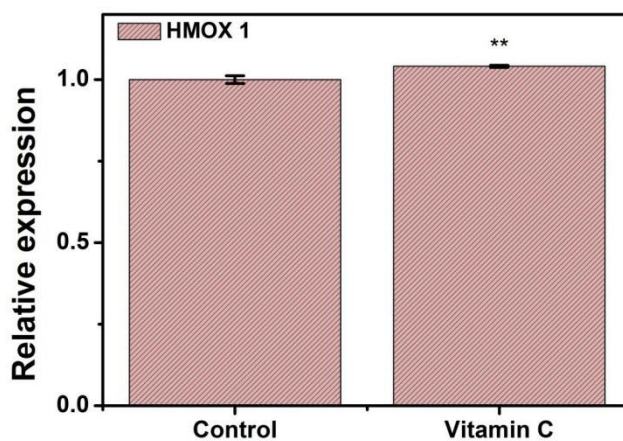


Figure S3. Effect of L-ascorbic acid (VC) on HMOX 1 gene expression in RAW264.7 cells. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$, The mean and standard deviation are represented by column and error bar.

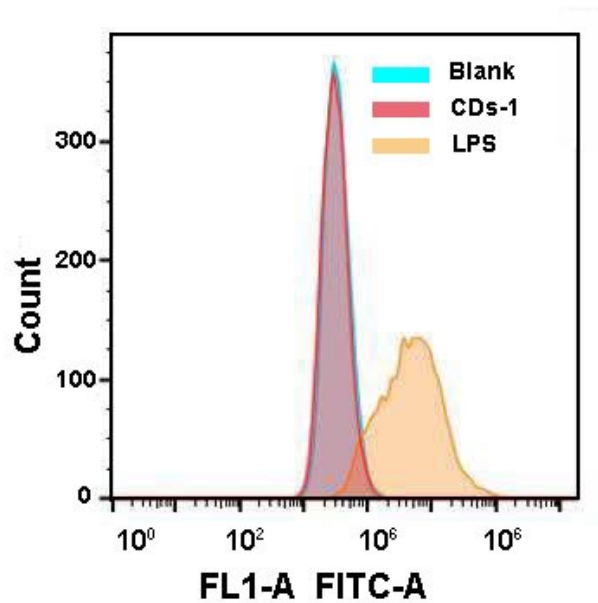


Figure S4. The reactive oxygen species (ROS) level of RAW264.7 cells treated with CDs-1 and LPS respectively was tested by flow cytometry. Untreated RAW264.7 was set as blank.

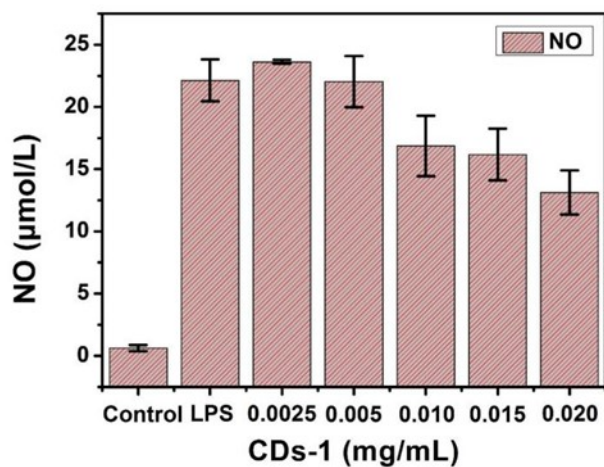


Figure S5. Level of NO in RAW264.7 treated with different concentrations of CDs-1.

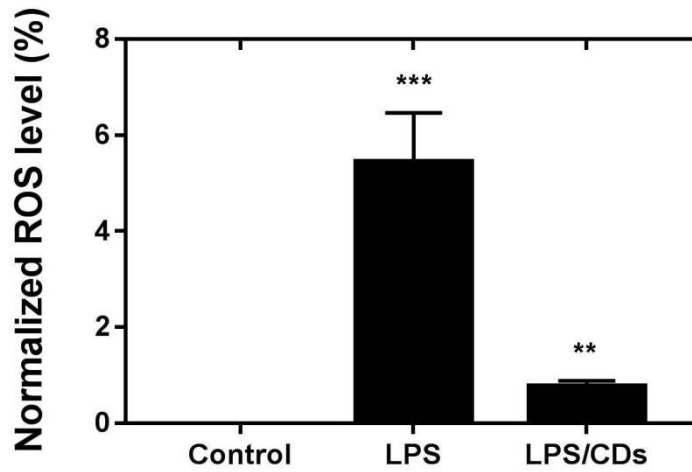


Figure S6. Normalized cellular ROS level in RAW264.7 cells in Control group, LPS group and CDs-1-treated-group respectively. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$, The mean and standard deviation are represented by column and error bar.

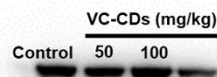


Figure S7. The proteins expression of GAPDH in lung tissue of mice treated with CDS-1 for 7 days, the concentrations of CDs-1 were 0, 50 and 100 mg/kg, respectively.

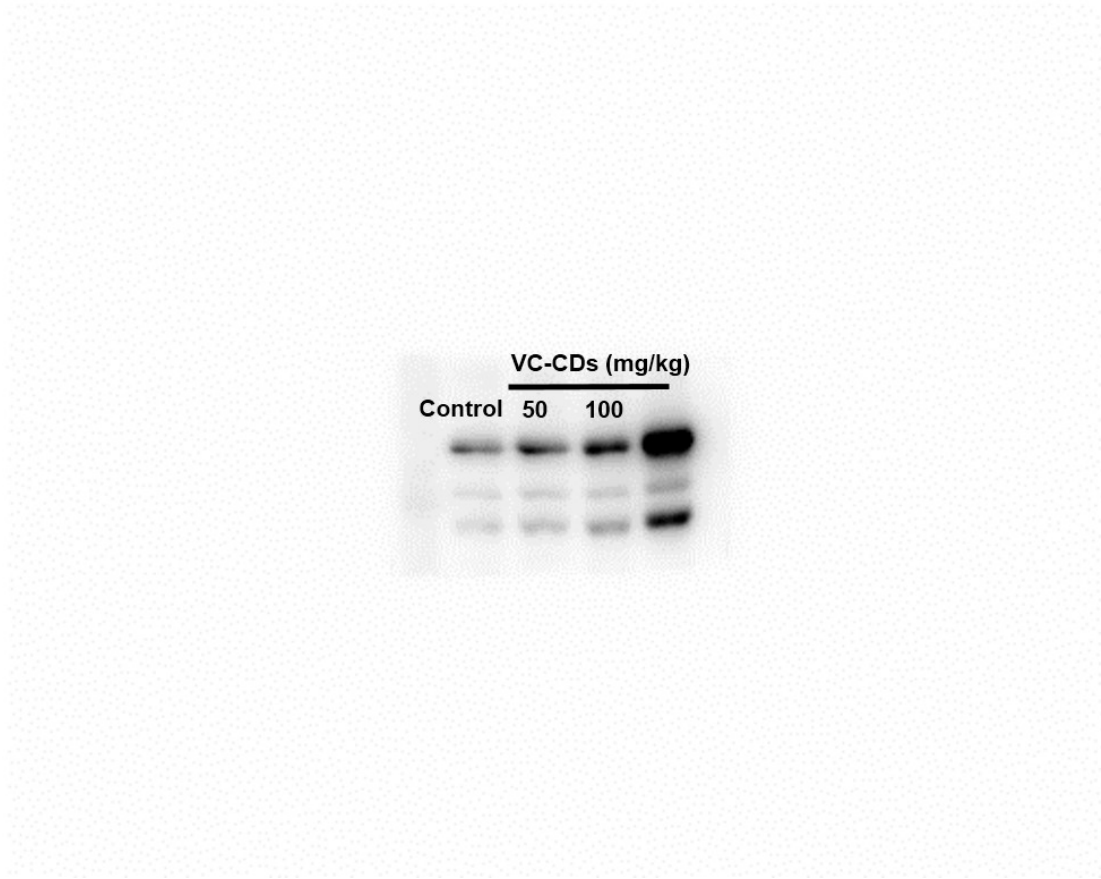


Figure S8. The proteins expression of HMOX 1 in lung tissue of mice treated with CDS-1 for 7 days, the concentrations of CDs-1 were 0, 50 and 100 mg/kg, respectively.

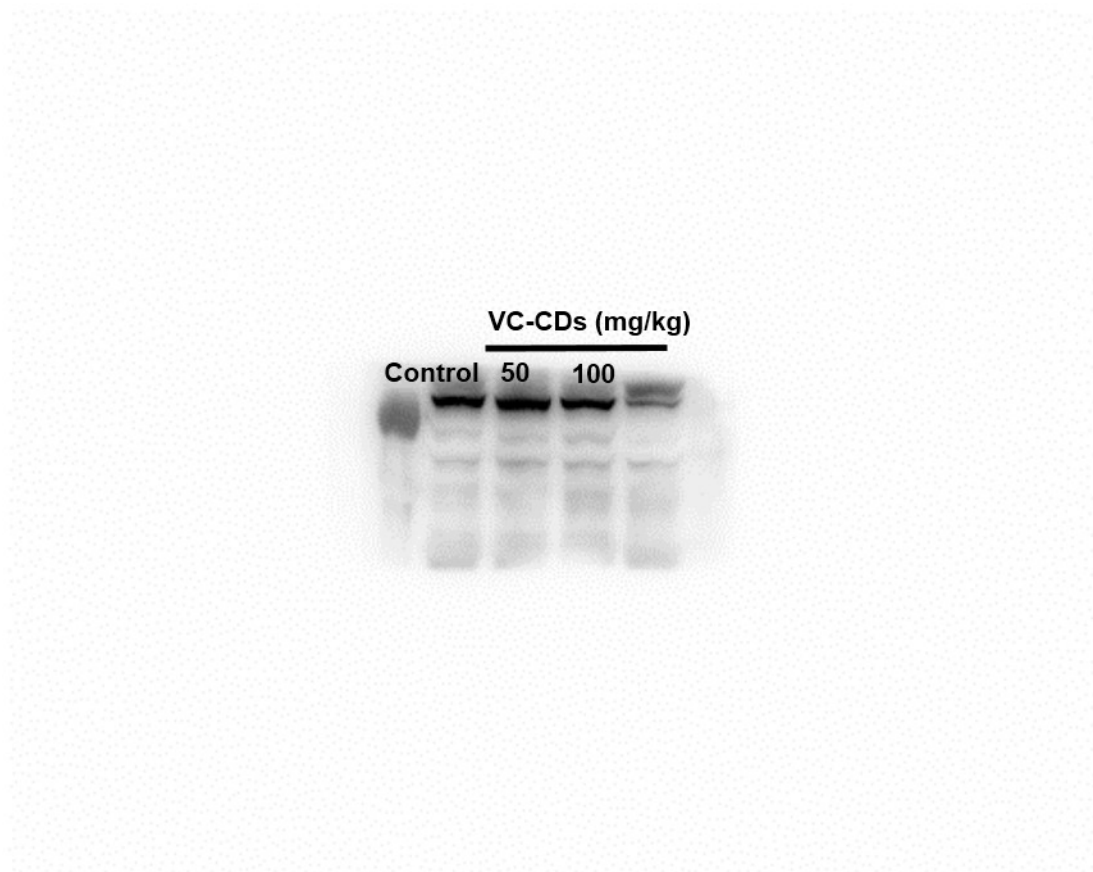


Figure S9. The proteins expression of NRF 2 in lung tissue of mice treated with CDS-1 for 7 days, the concentrations of CDs-1 were 0, 50 and 100 mg/kg, respectively.

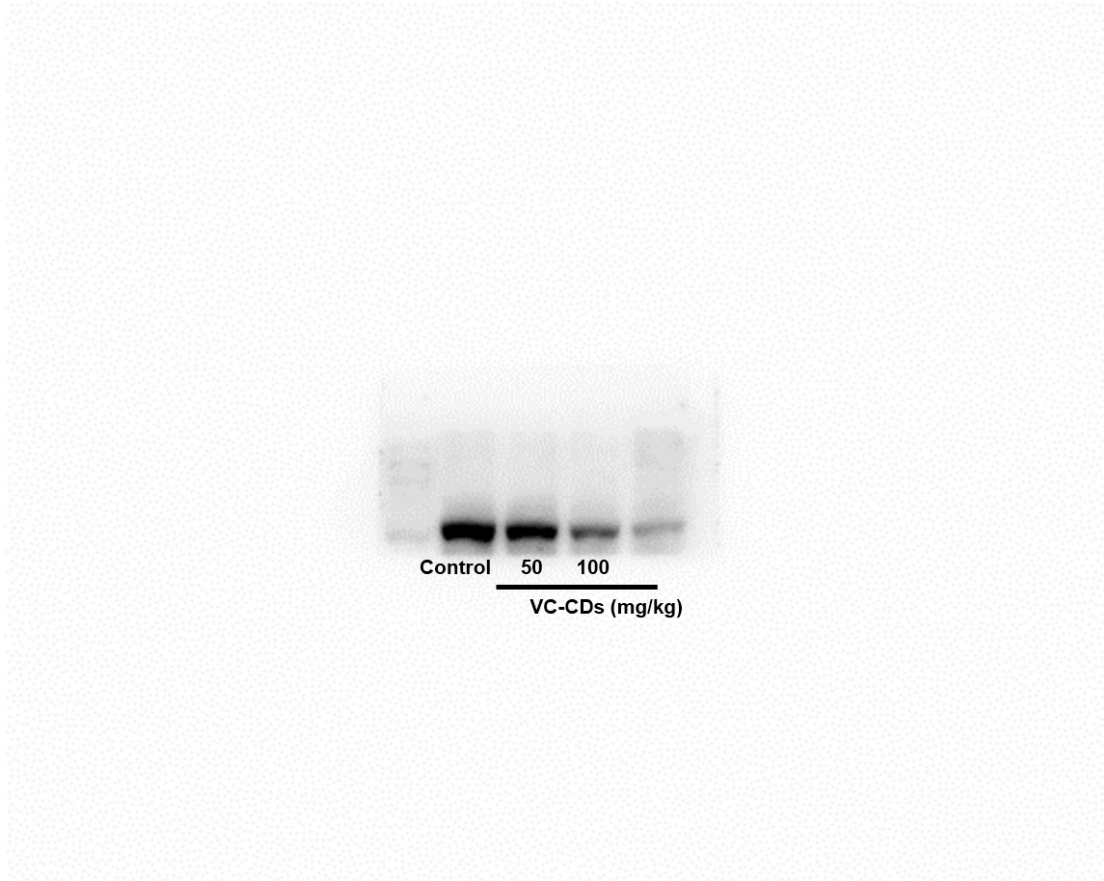


Figure S10. The proteins expression of BACH 1 in lung tissue of mice treated with CDS-1 for 7 days, the concentrations of CDs-1 were 0, 50 and 100 mg/kg, respectively.