Supporting Information

Antidepressant effects of Sulforaphane (SFN) and its derivatives SLL-III-9 and SLL-III-120 and their potential underlying mechanisms based on the microbiotagut-brain axis

Lili Chen,^{abc} Huihui Cao,^a Xin Zhang,^a Xintong Du,^a Yang Guan,^a Mei Li,^a Alan K Chang,^d Xianran He,^e Xiaolong Li ^f and Xiuli Bi*ab

Affiliations

^aCollege of Life Science, Liaoning University, Shenyang, 110036, China

bShenyang Key Laboratory of Chronic Disease Occurrence and Nutrition Intervention,
College of Life Sciences, Liaoning University, Shenyang, 110036, China.

^cCollege of Mathematics and Statistics, Liaoning University, Shenyang, 110036, China ^dCollege of Life and Environmental Sciences, Wenzhou University, Wenzhou 325035, China

^eInstitute for Interdisciplinary Research, Jianghan University, Wuhan Economic and Technological Development Zone, Wuhan 430056, China

^fShenzhen Fushan Biological Technology Co., Ltd, Kexing Science Park A1 1005, Nanshan Zone, Shenzhen 518057, China

*Corresponding Author

Xiuli Bi, Ph.D

Professor

College of Life Science, Liaoning University, Shenyang

66 Chongshan Road, Shenyang, 110036, P. R. China.

Tel: +86-24-62202232

E-mail address: xiulibi@lnu.edu.cn; xiulibi@gmail.com

Figure S1. ¹H NMR of SLL-III-9.

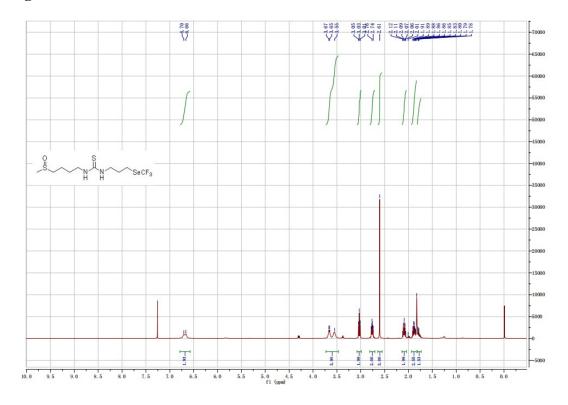


Figure S2. 1 H NMR of SLL-III-120.

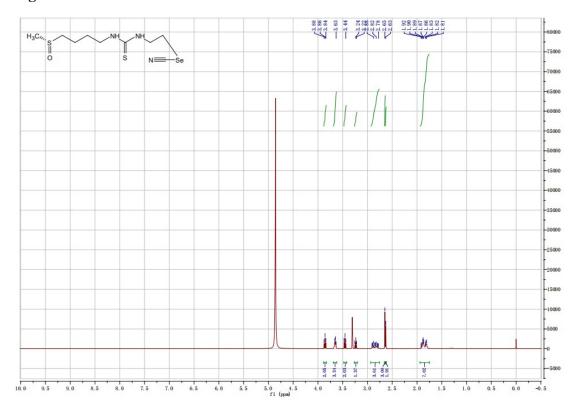


Figure S3. ¹³C NMR of SLL-III-9.

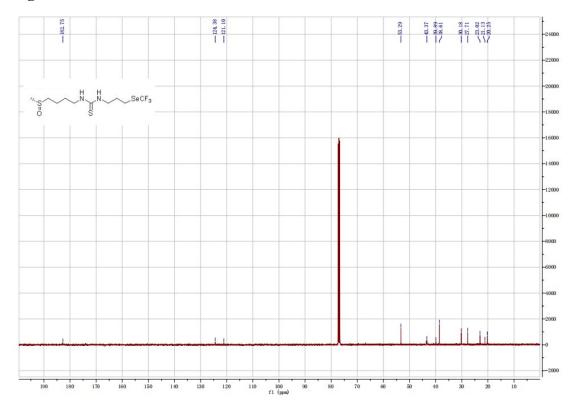


Figure S4. ¹³C NMR of SLL-III-120.

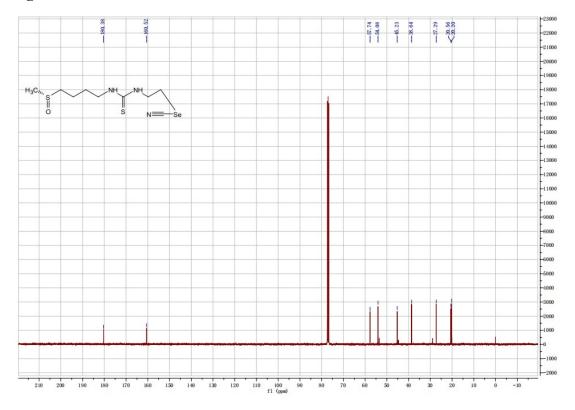


Figure S5. ¹⁹F NMR of SLL-III-9.

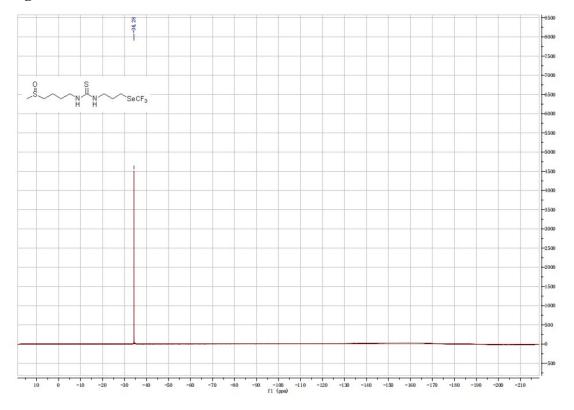


Figure S6. MS of SLL-III-120.

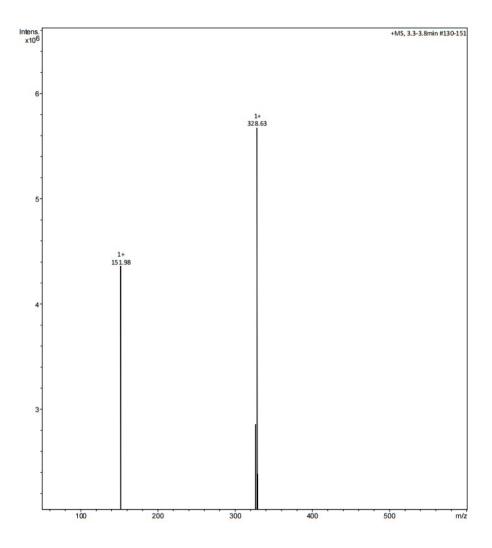


Figure S7. A mouse model of depression was validated by behavioral experiments.

After mice were raised in isolation and given stress stimuli for 13 weeks, depression-related indicators (A) sucrose preference test, (B) tail suspension test, and (C) forced swimming test were performed to detect the successful establishment of the depressed mouse model. **P<0.01 compared to control group.

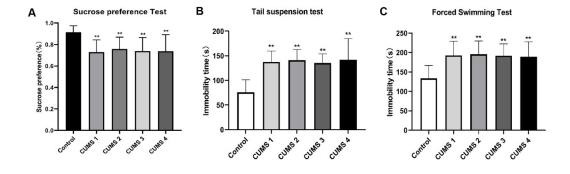


Figure S8. The effect of different concentration of SFN, SLL-III-9, and SLL-III-120 in LPS-activated BV2 cells. After BV2 cells were treated with 1 μ g/mL LPS for 24 h to establish neuroinflammatory cell models, different concentrations of SFN, SLL-III-9, or SLL-III-120 were added to the culture medium for 24 h. The culture medium and cells were collected, and Griess assay, MTT assay and real-time PCR was performed to analysis (A) the production of nitrate, (B) cell viability and (C) iNOS mRNA expression. *P<0.05, **P<0.01, ***P<0.001 compared to the LPS group and **P<0.05, ***P<0.01, ***P<0.001 compared to control group.

