

Unsaturation (U.S.): 0.0 - 15.0

Observed m/z Int% Err[ppm / mmu] U.S. Composition
1 168.0658 79.68 -1.6 / -0.3 4.5 C8 H10 N O3
2 +14.3 / +2.4 0.0 C5 H12 O6

Figure S1. CIMS and HRCIMS spectra of compound 6 in CDCl₃.

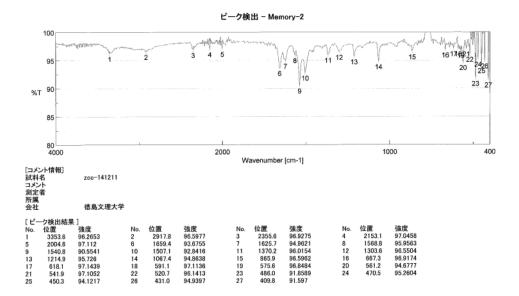


Figure S2. IR spectra of compound 6 in CDCl₃.

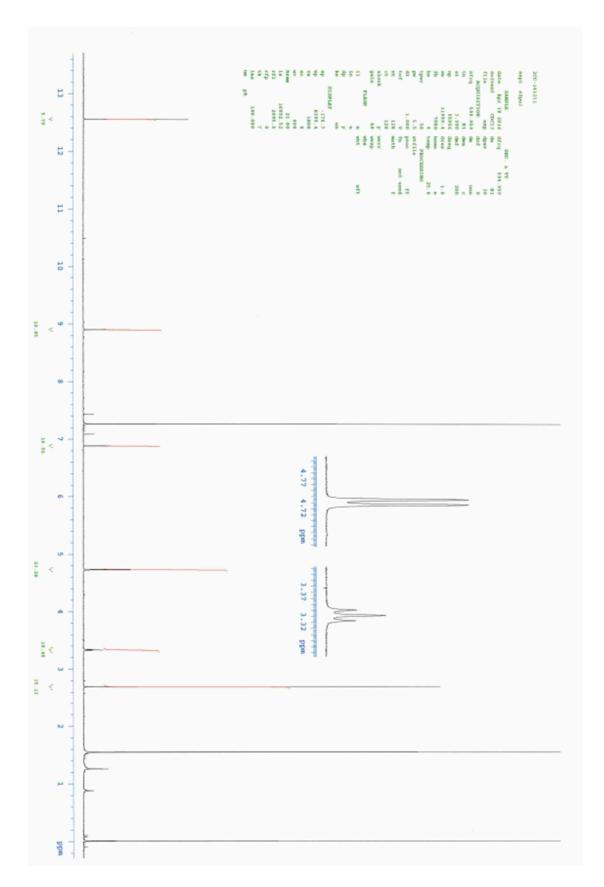


Figure S3. ¹H spectra of compound 6 in CDCl₃.

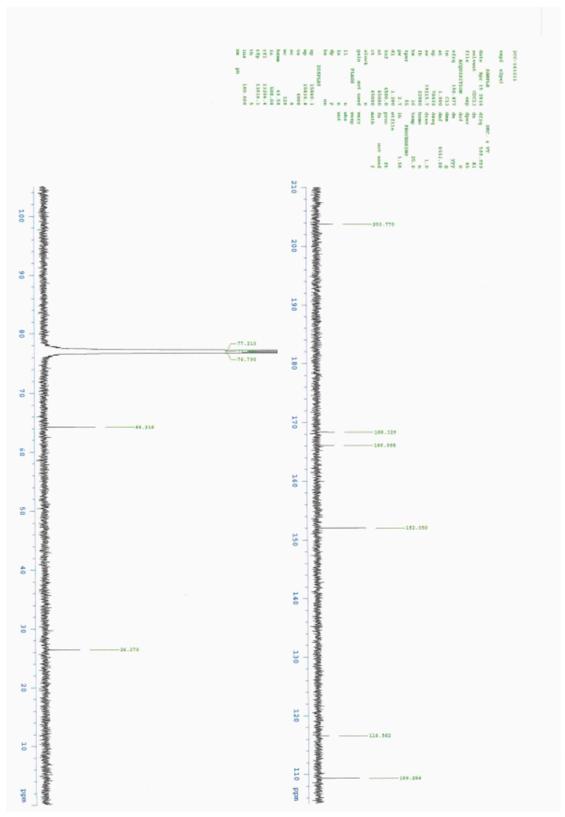


Figure S4. ¹³C spectra of compound 6 in CDCl₃.

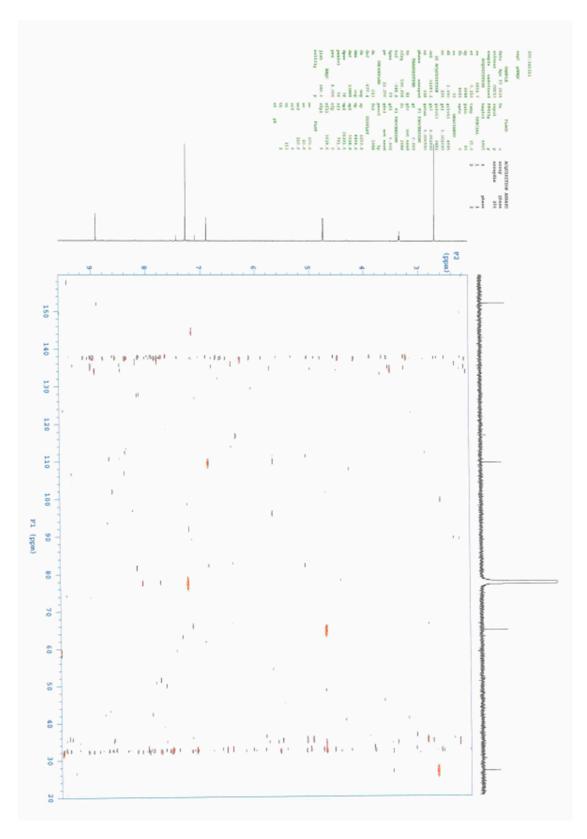


Figure S5. HSQC spectra of compound 6 in CDCl₃.

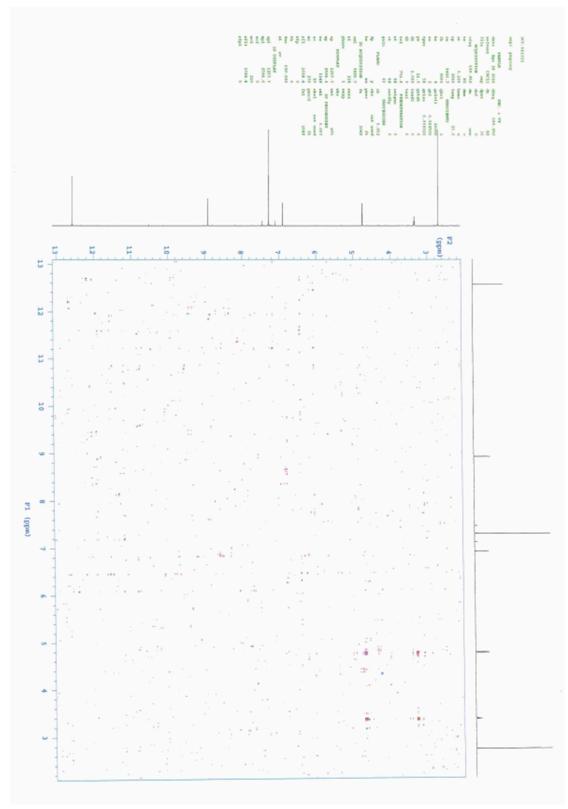


Figure S6. COSY spectra of compound 6 in CDCl₃.

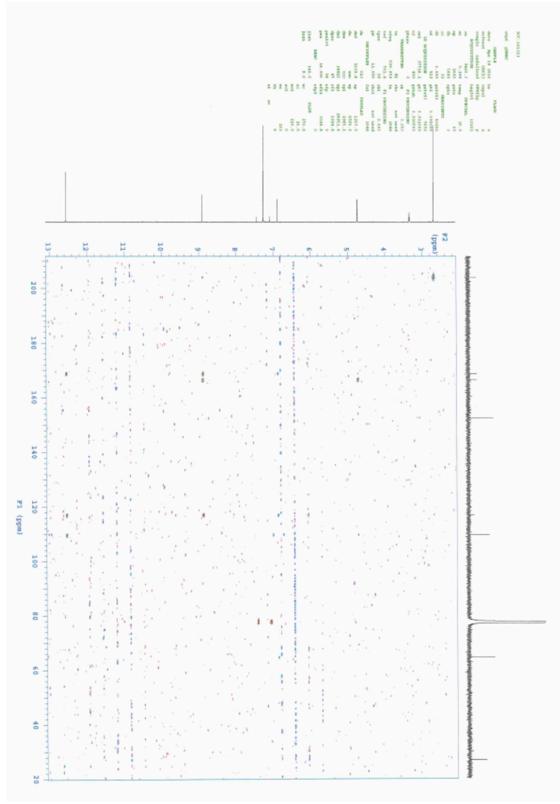


Figure S7. HMBC spectra of compound 6 in CDCl₃.

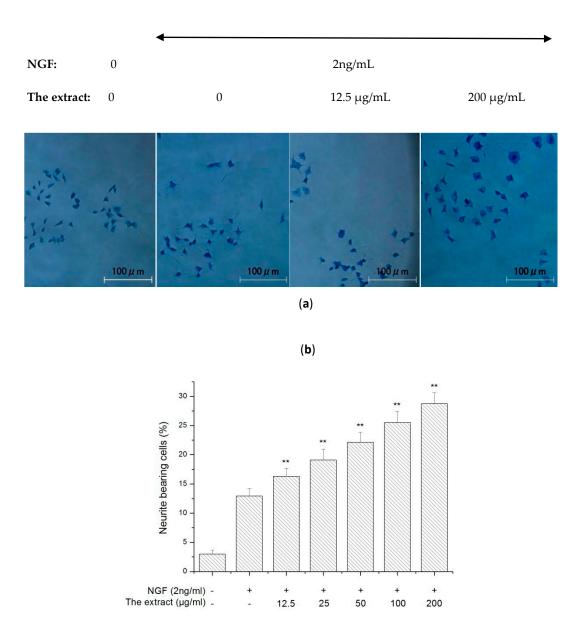


Figure S8. Effect of MeOH extract from the mycelium of *H. erinaceus* on promoting neurite outgrowth in NGF-induced PC12 cells. PC12 cells were seeded in collagen-coated 24-well plates in normal serum medium for 24 h, then changed to low serum medium (2% HS and 1% FBS) and exposure to vehicle (0.1% DMSO) as a negative control, NGF (2 ng/mL) as a positive control, or NGF (2 ng/mL) + the extract (12.5–200 μ g/mL) for another 96 h. Scale bar: 100 μ M. (a) Cell morphology was observed and photographed as described in the analysis of neurite outgrowth. (b) Neurite-bearing cells were measured as described in thre analysis of neurite outgrowth. Data are expressed as the mean \pm SD from three independent experiments. **p < 0.01 represents significant differences compared with NGF-treated PC12 cells (positive control) (ANOVA followed by Dunnett's test).

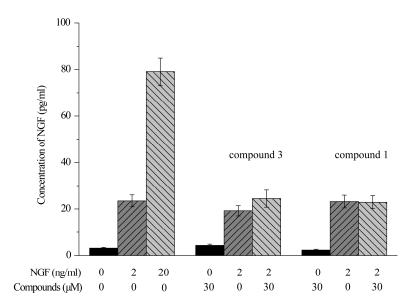


Figure S9. Effects of compounds 1 and 3 on the stimulation of NGF secretion in PC12 cells. PC12 cells were incubated with 30 μ M compounds with or without NGF (2 ng/mL) for 96 h and then the NGF level in the conditioned medium was measured by NGF beta rat ELISA Kit (Thermo Fisher Scientific Inc.: Waltham, MS, USA) following the manufacturer's instructions. Recombinant rat beta-NGF was used as standards.