Supplementary data

Chemical synthesis, inhibitory activity and molecular mechanism of 1deoxynojirimycin–chrysin as a potent α-glucosidase inhibitor

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Fig. S1. ¹H-NMR spectrum (400 MHz) of compound 1 in DMSO-*d*₆.



Fig. S2. ¹³C-NMR spectrum (101 MHz) of compound 1 in DMSO- d_6 .



Fig. S3. ¹H-NMR spectrum (400 MHz) of compound 2 in DMSO-*d*₆.



Fig. S4. ¹³C-NMR spectrum (101 MHz) of compound 2 in DMSO-*d*₆.



Fig. S5. ¹H-NMR spectrum (600 MHz) of compound **3** in CDCl₃.



Fig. S6. ¹³C-NMR spectrum (150 MHz) of compound 3 in CDCl₃.



Fig. S8. ¹³C-NMR spectrum (101 MHz) of compound 4 in DMSO-*d*₆.



Fig. S9. HRMS spectrum of compound 4.



Fig. S10. HPLC characterization of compound 4.



Fig. S11. ¹H-NMR spectrum (600 MHz) of compound 5 in DMSO-*d*₆.



Fig. S12. ¹³C-NMR spectrum (150 MHz) of compound 5 in DMSO-*d*₆.



Fig. S13. HRMS spectrum of compound 5.



Fig. S14. HPLC characterization of compound 5.



Fig. S11. ¹H-NMR spectrum (600 MHz) of compound 6 in DMSO-*d*₆.



Fig. S12. ¹³C-NMR spectrum (150 MHz) of compound 6 in DMSO-*d*₆.



Fig. S17. HRMS spectrum of compound 6.



Fig. S18. HPLC characterization of compound 6.