Supporting information

New dammarane-type triterpenoid glycosides from Gynostemma burmanicum

Than Thi Kieu My^a, Pham Thanh Ky^a, Pham Thanh Binh^b, Nguyen Phuong Thao^b and Nguyen Tien Dat^c.*

^aDepartment of Pharmacognosy, Hanoi University of Pharmacy, 13-15 Le Thanh Tong, Hanoi, Vietnam; ^bInstitute of Marine Biochemistry, Vietnam Academy of Science and Technology (VAST), 18-Hoang Quoc Viet, Caugiay, Hanoi, Vietnam; ^cCenter for Research and Technology Transfer, VAST, 18-Hoang Quoc Viet, Cau Giay, Hanoi, Vietnam

Corresponding author: Nguyen Tien Dat: ngtiend@imbc.vast.vn

ABSTRACT

The chemical composition of *Gynostemma burmanicum* King ex Chakrav. was investigated for the first time in this study. Nine dammarane glycosides (1–9) were isolated from the EtOH extract of the aerial parts of *G. burmanicum*. Their structures were elucidated by 1D and 2D NMR spectroscopic interpretation as well as by chemical studies. The new compounds were 3β ,20*S*-dihydroxydammar-24-ene-3-O- β -D-glucopyranosyl-20-O-[β -D-xylopyranosyl-(1 \rightarrow 6)- β -D-glucopyranoside] (1), 3β ,12 β ,20*S*-trihydroxydammar-24-ene-3-O- β -D-xylopyranosyl-20-O-[β -D-xylopyranosyl-(1 \rightarrow 6)- β -D-glucopyranoside] (2), and 12-oxo- 3β ,20*S*-dihydroxydammar-24-ene-3-O-[β -D-glucopyranosyl-(1 \rightarrow 6)- β -D-glucopyranosyl-(

KEYWORDS: Gynostemma burmanicum, Cucurbitaceae, dammarane saponin

Cotents

Figure S1. ¹ H NMR spectrum (CD ₃ OD, 500 MHz) of 1
Figure S2. ¹³ C NMR spectrum (CD ₃ OD, 125 MHz) of 1
Figure S3. COSY spectrum (CD ₃ OD, 500 MHz) of 1
Figure S4. HMQC spectrum (CD ₃ OD, 500 MHz) of 1
Figure S5. HMBC spectrum (CD ₃ OD, 500 MHz) of 1
Figure S6. NOESY spectrum (CD ₃ OD, 500 MHz) of 1
Figure S7. ¹ H NMR spectrum (CD ₃ OD, 500 MHz) of 2
Figure S8. ¹³ C NMR spectrum (CD ₃ OD, 125 MHz) of 1
Figure S9. COSY spectrum (CD ₃ OD, 500 MHz) of 2
Figure S10. HMQC spectrum (CD ₃ OD, 500 MHz) of 2
Figure S11. HMBC spectrum (CD ₃ OD, 500 MHz) of 2
Figure S12. NOESY spectrum (CD ₃ OD, 500 MHz) of 2
Figure S13. ¹ H NMR spectrum (CD ₃ OD, 500 MHz) of 3
Figure S14. ¹³ C NMR spectrum (CD ₃ OD, 125 MHz) of 3
Figure S15. COSY spectrum (CD ₃ OD, 500 MHz) of 3
Figure S16. HMQC spectrum (CD ₃ OD, 500 MHz) of 3
Figure S17. HMBC spectrum (CD ₃ OD, 500 MHz) of 3
Figure S18. Key HMBC (\rightarrow) and COSY $()$ of 1-3 and NOESY $(<-)$
correlations of 2

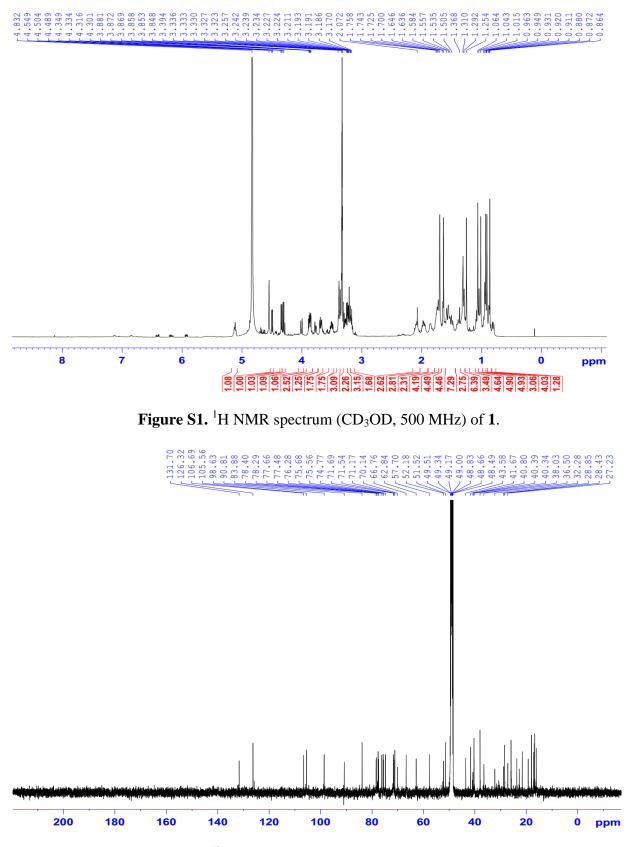


Figure S2. ¹³C NMR spectrum (CD₃OD, 125 MHz) of 1.

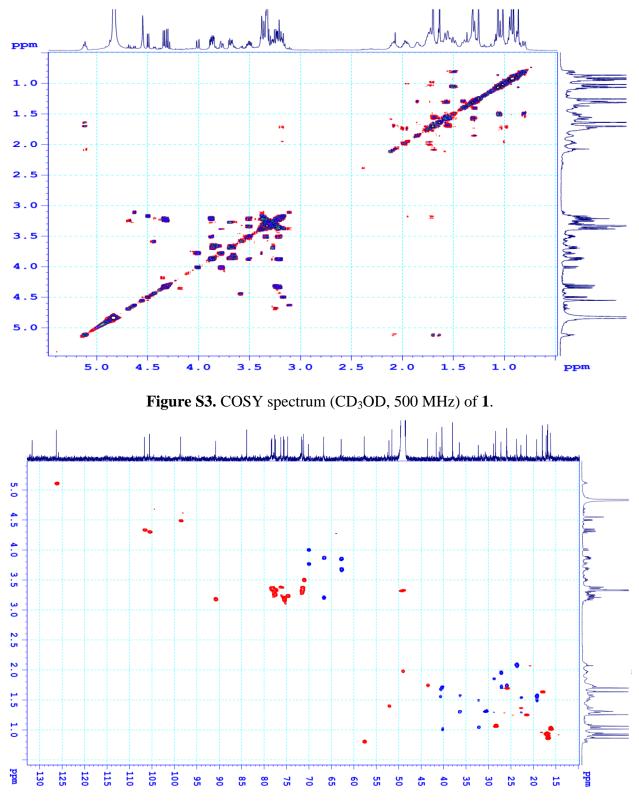


Figure S4. HMQC spectrum (CD₃OD, 500 MHz) of 1.

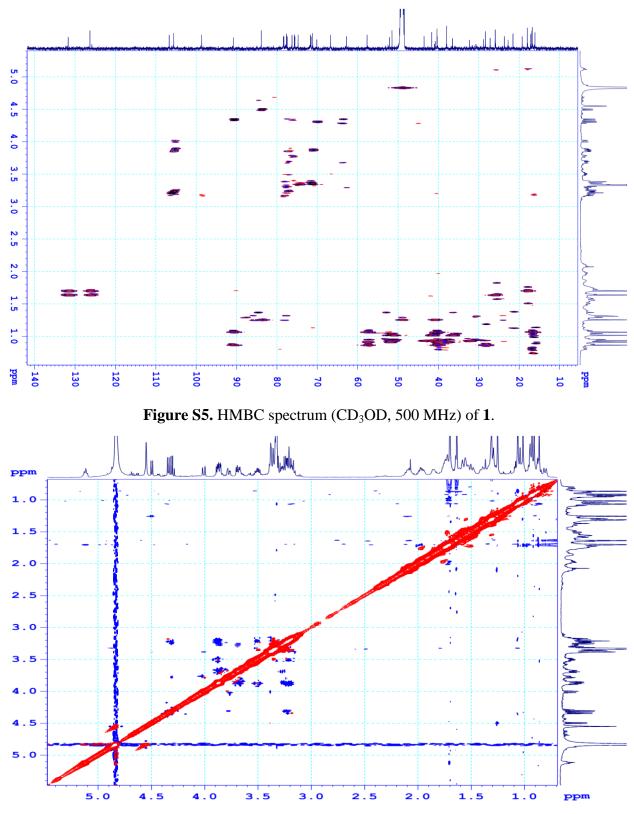


Figure S6. NOESY spectrum (CD₃OD, 500 MHz) of 1.

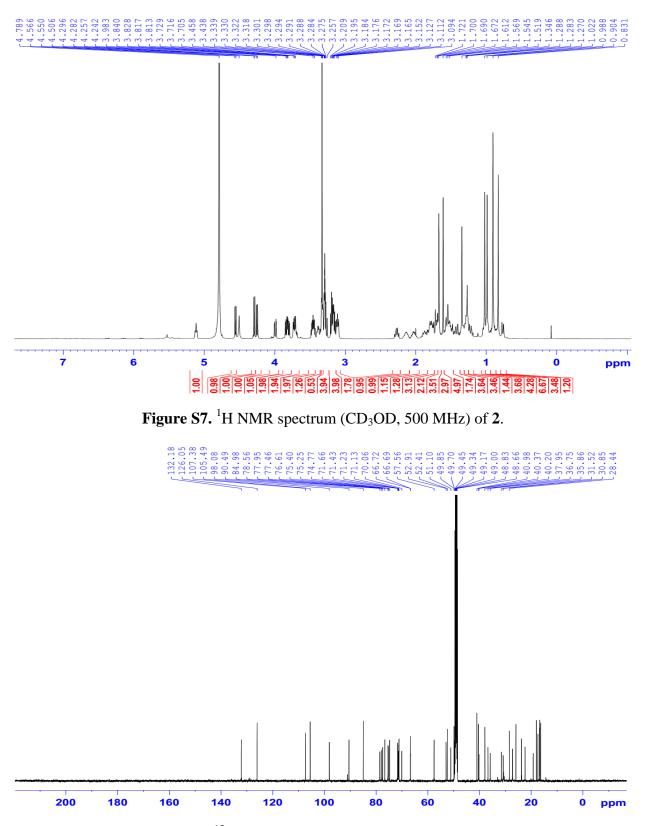
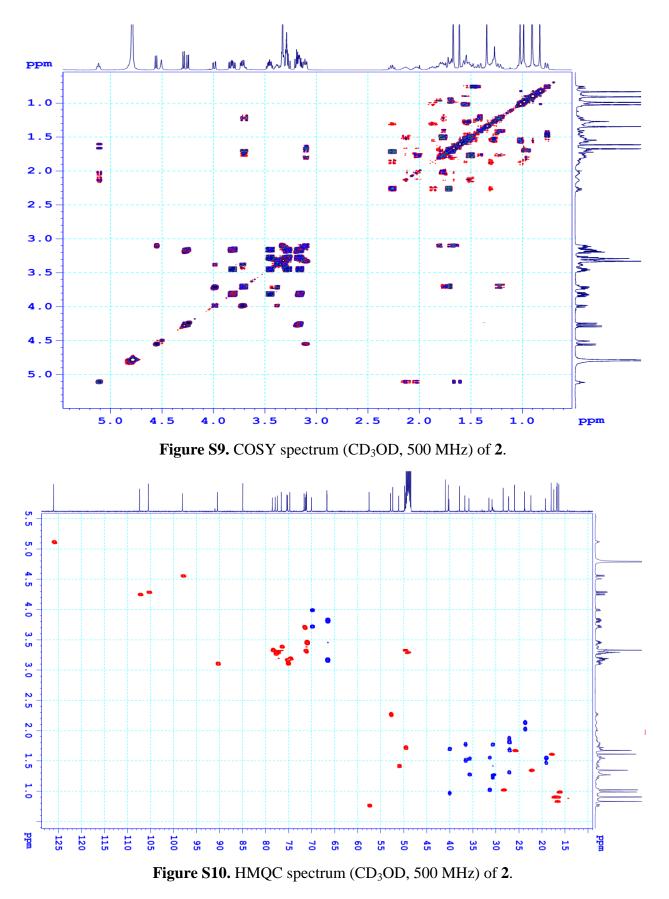


Figure S8. ¹³C NMR spectrum (CD₃OD, 125 MHz) of 1.



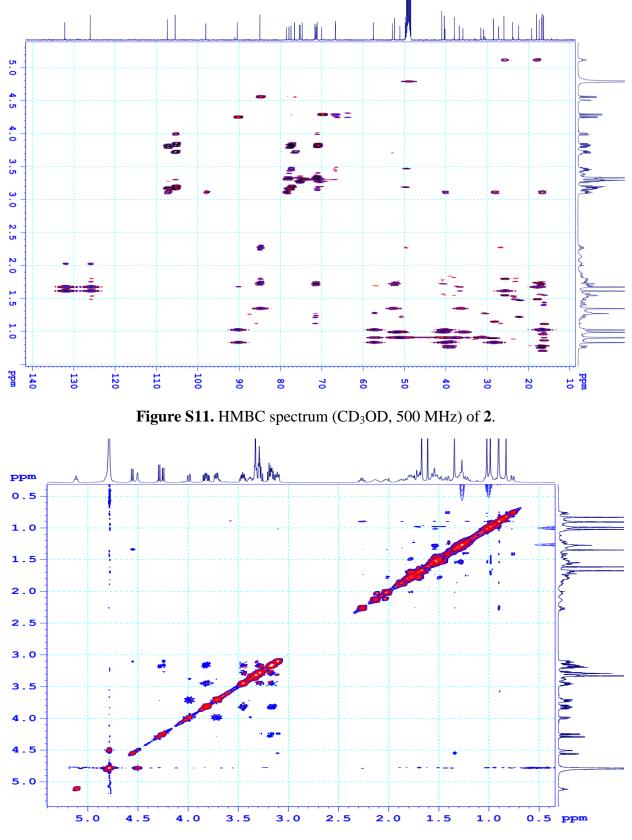


Figure S12. NOESY spectrum (CD₃OD, 500 MHz) of 2.

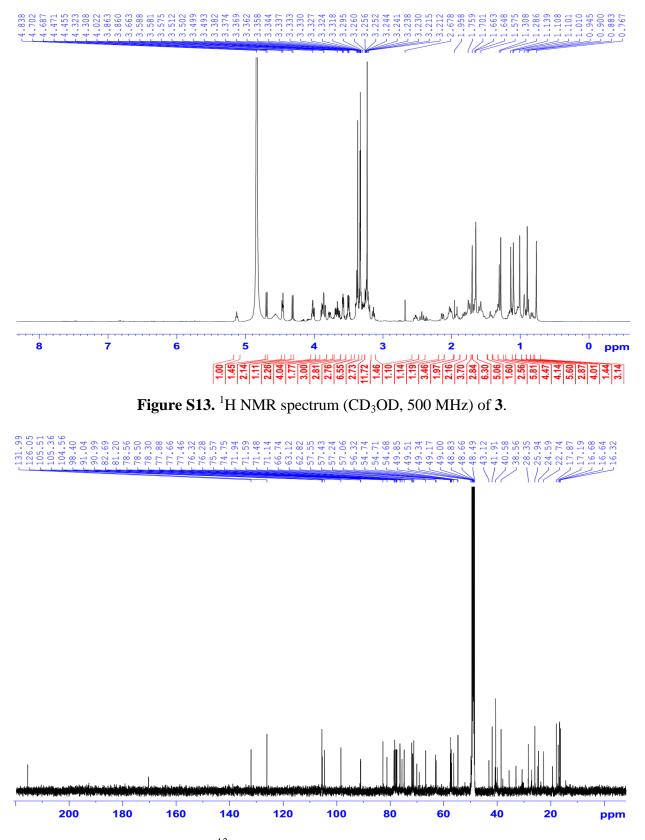


Figure S14. ¹³C NMR spectrum (CD₃OD, 125 MHz) of 3.

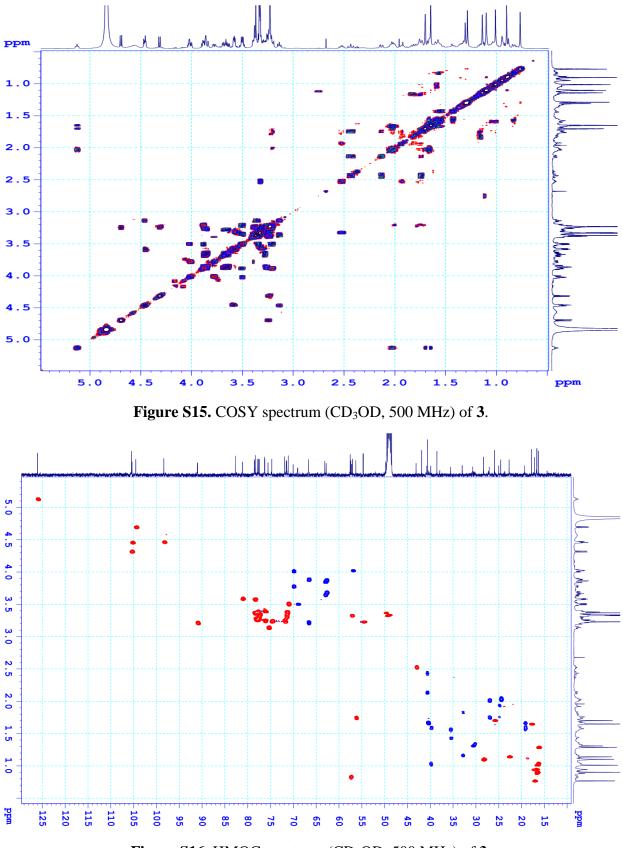


Figure S16. HMQC spectrum (CD₃OD, 500 MHz) of 3.

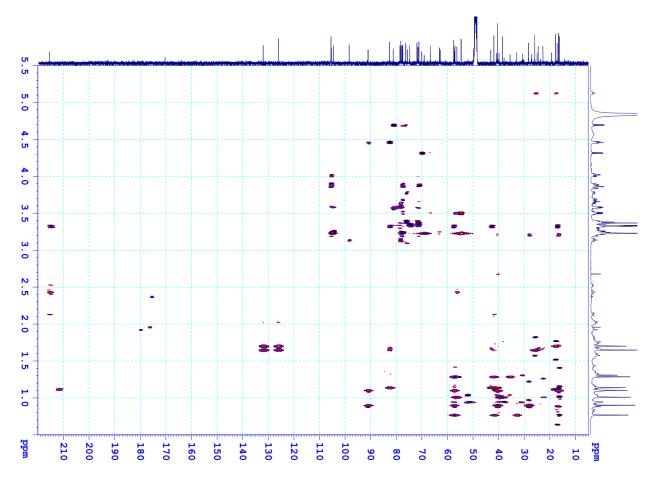
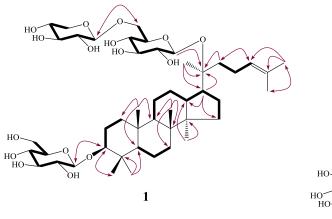
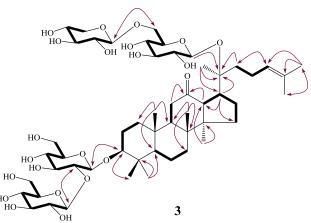


Figure S17. HMBC spectrum (CD₃OD, 500 MHz) of 3.





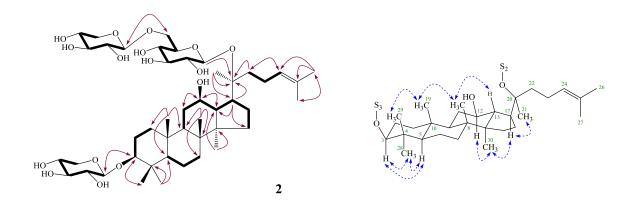


Figure S18. Key HMBC (\rightarrow) and COSY (\longrightarrow) of 1-3 and NOESY (<^{...}>) correlations of 2