Microwave-Assisted Synthesis of Mono- and Disubstituted 4-Hydroxyacetophenone Derivatives via Mannich Reaction: synthesis, XRD and HS-analysis

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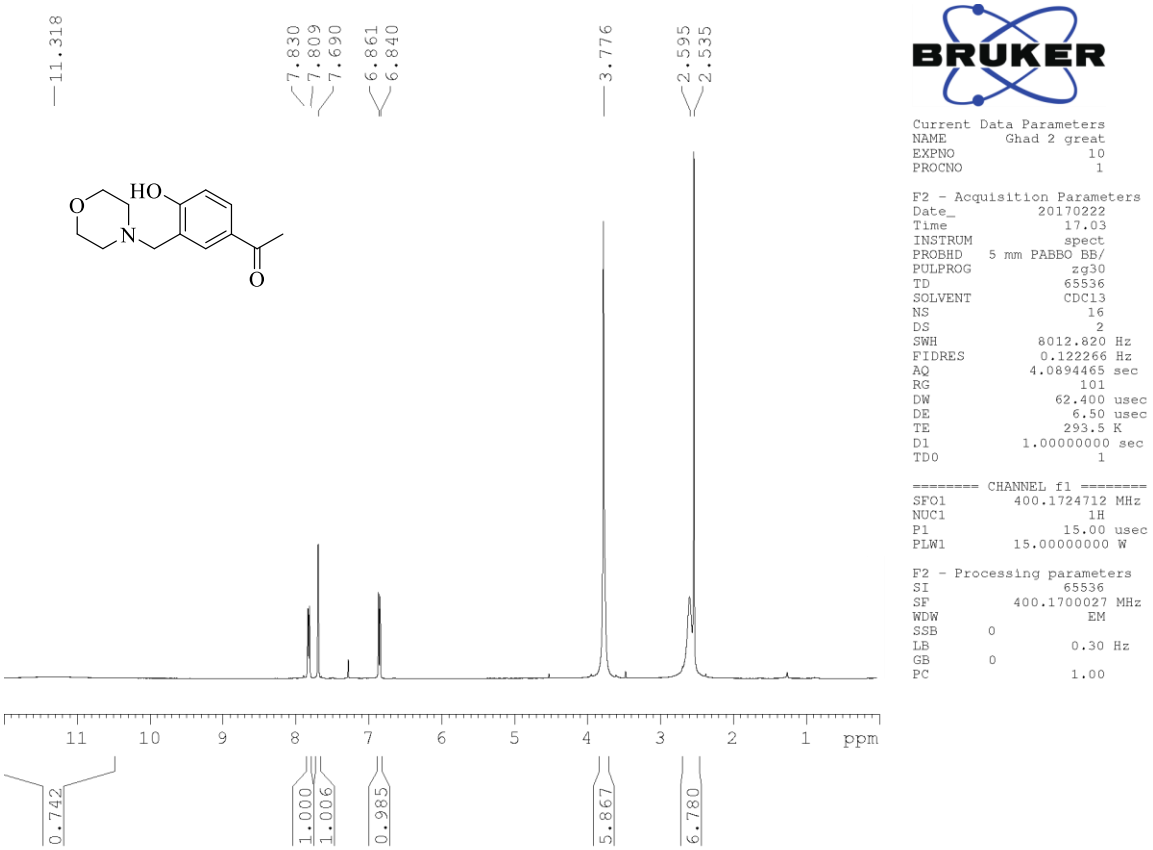


Figure S1. ^{1}HNMR spectrum of compound 2a in $CDCl_3$.

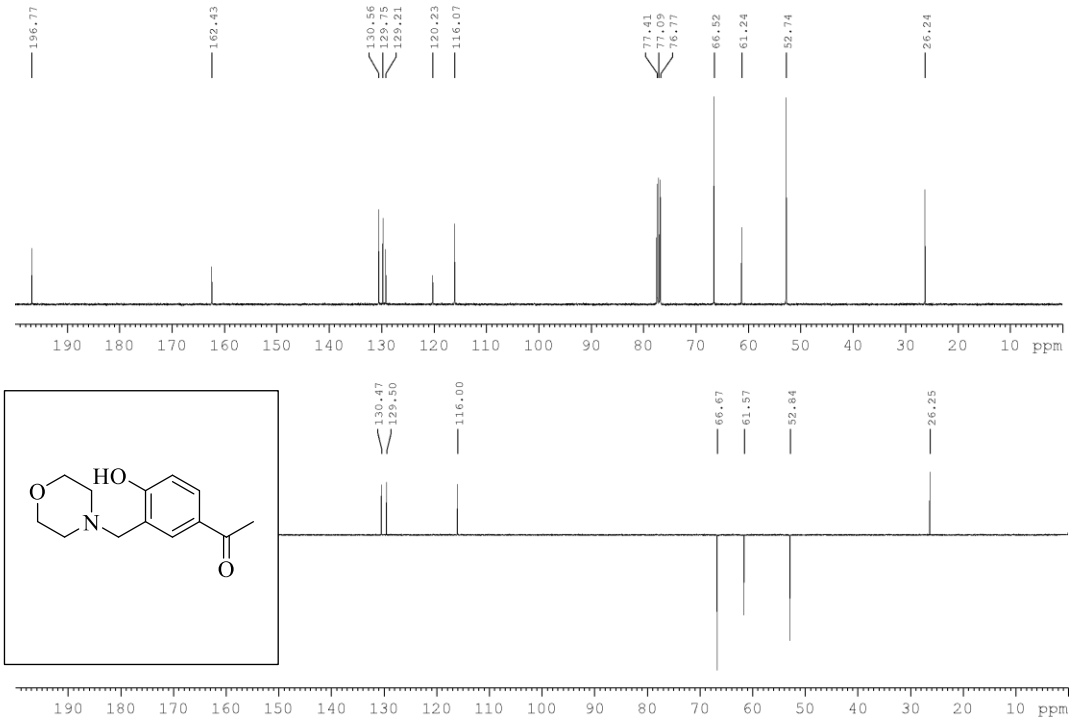


Figure S2. ^{13}C NMR &Dept 135 spectrum of compound 2a in $CDCl_3$.

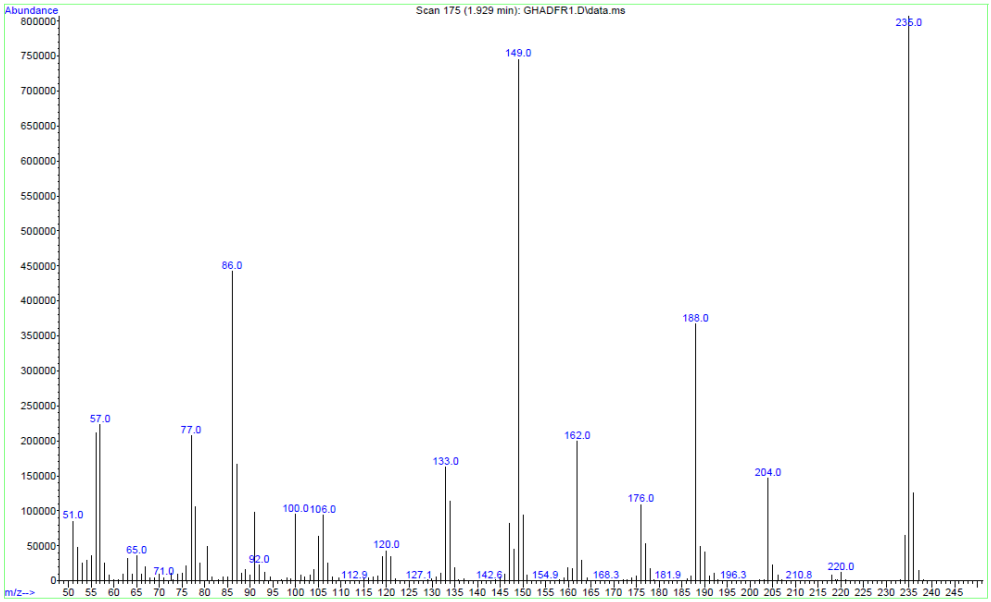


Figure S3. MS spectrum of compound 2a

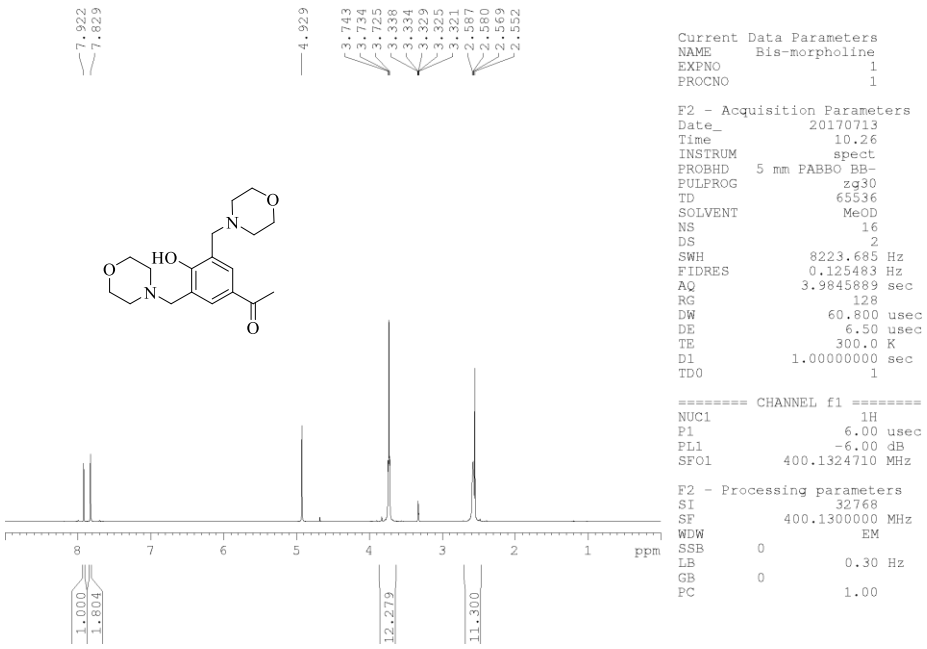


Figure S4. [1]HNMR spectrum of compound 2b in Methanol-d_4

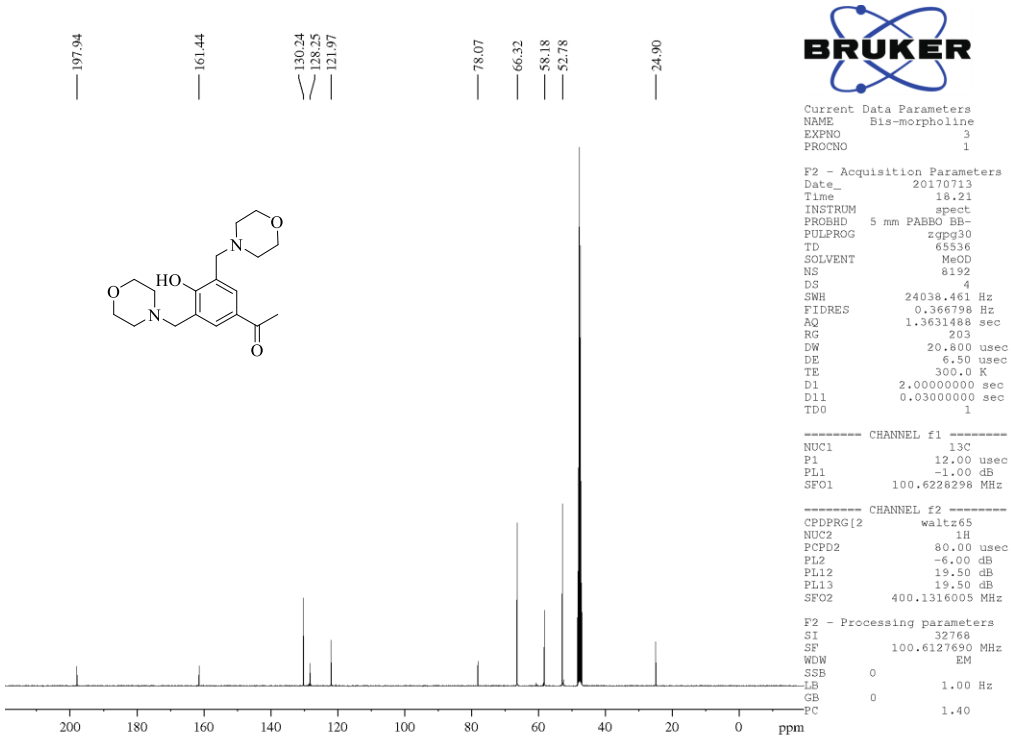


Figure S5. [13]C NMR spectrum of compound 2b in Methanol-d_4

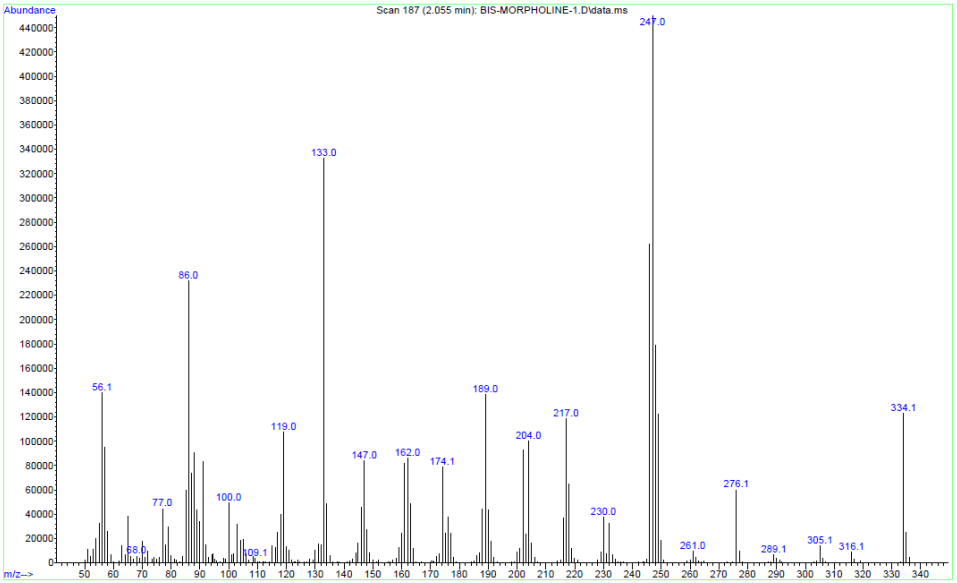


Figure S6. MS spectrum of compound 2b

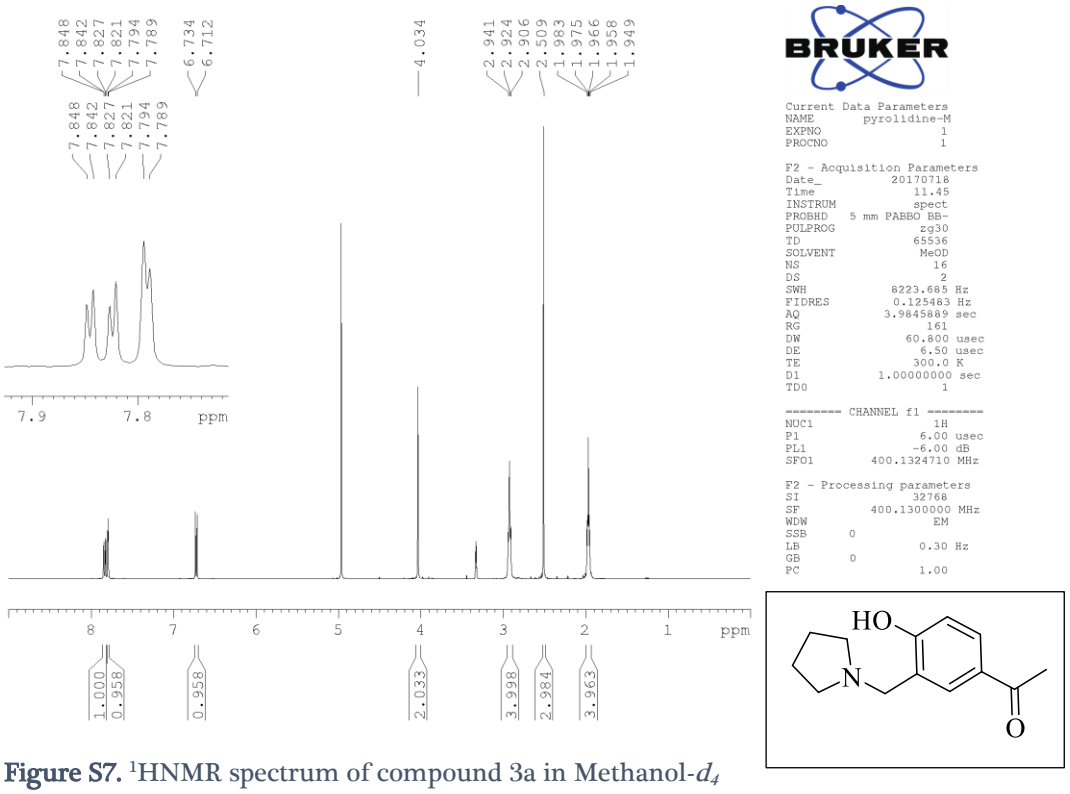


Figure S7. [1]HNMR spectrum of compound 3a in Methanol-d_4

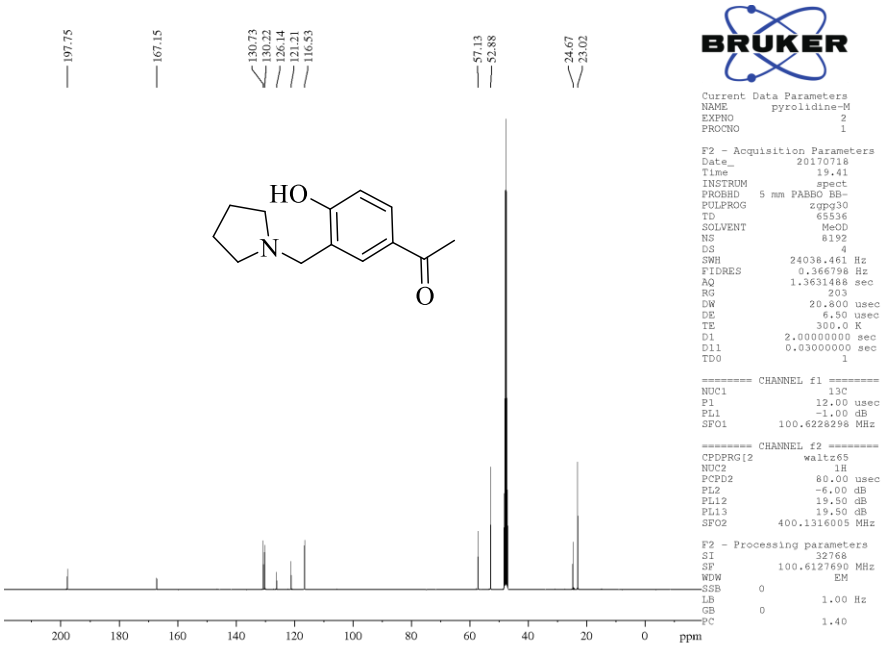


Figure S8. ^{13}C NMR spectrum of compound 3a in Methanol-d_4

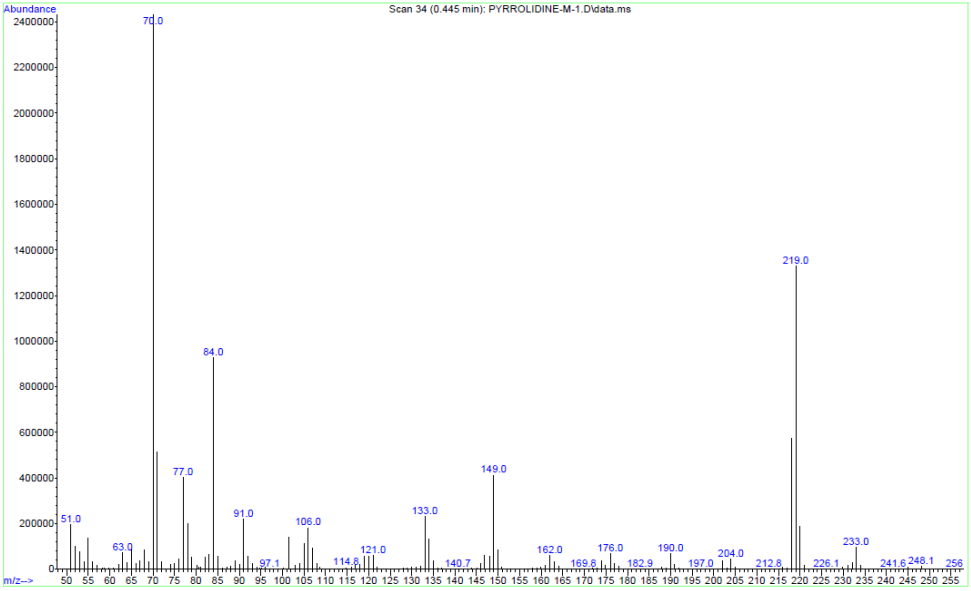


Figure S9. MS spectrum of compound 3a

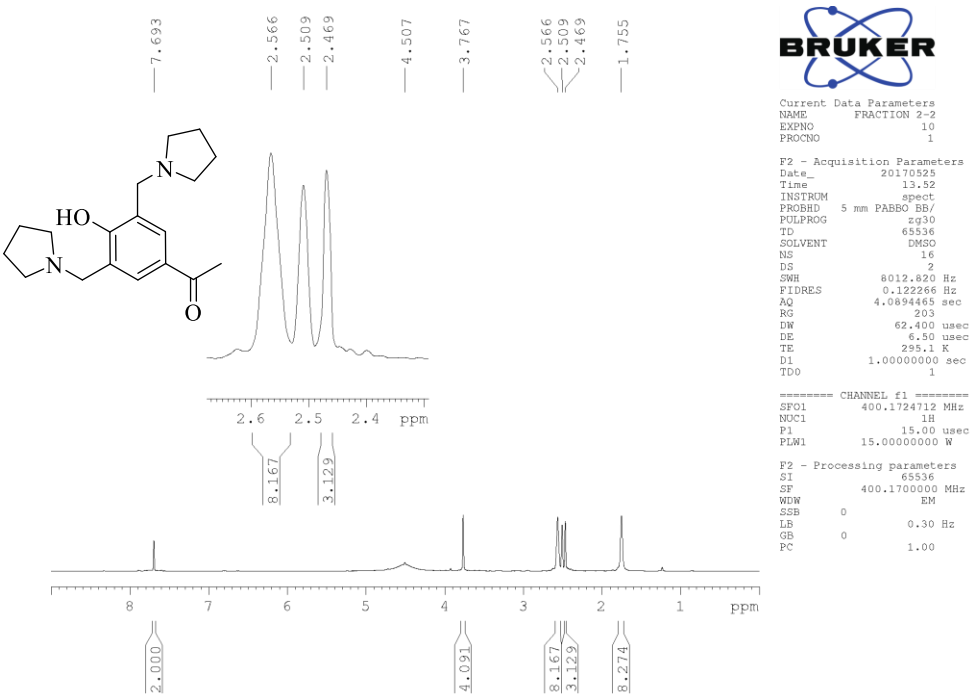


Figure S10. [1]HNMR spectrum of compound 3b in DMSO-d_6

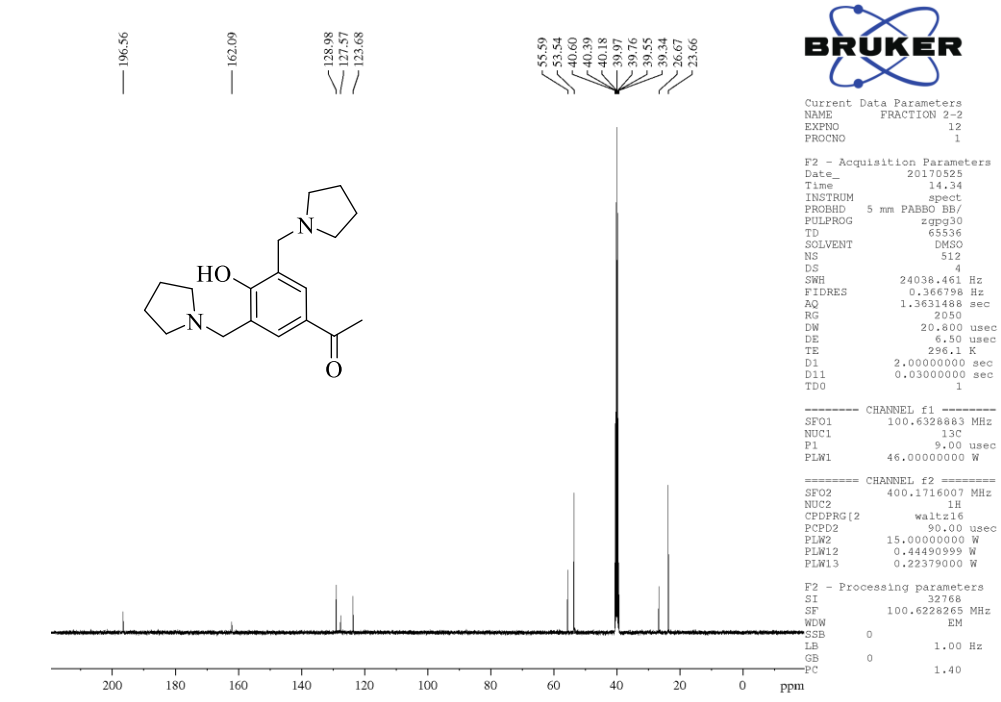


Figure S11. [13]C NMR spectrum of compound 3b in DMSO-d_6

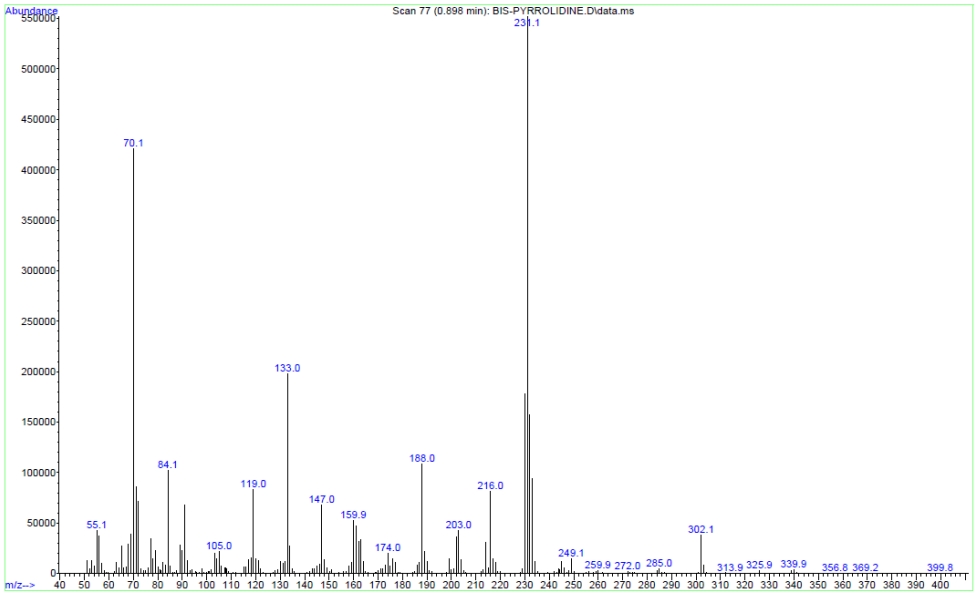


Figure S12. MS spectrum of compound 3b

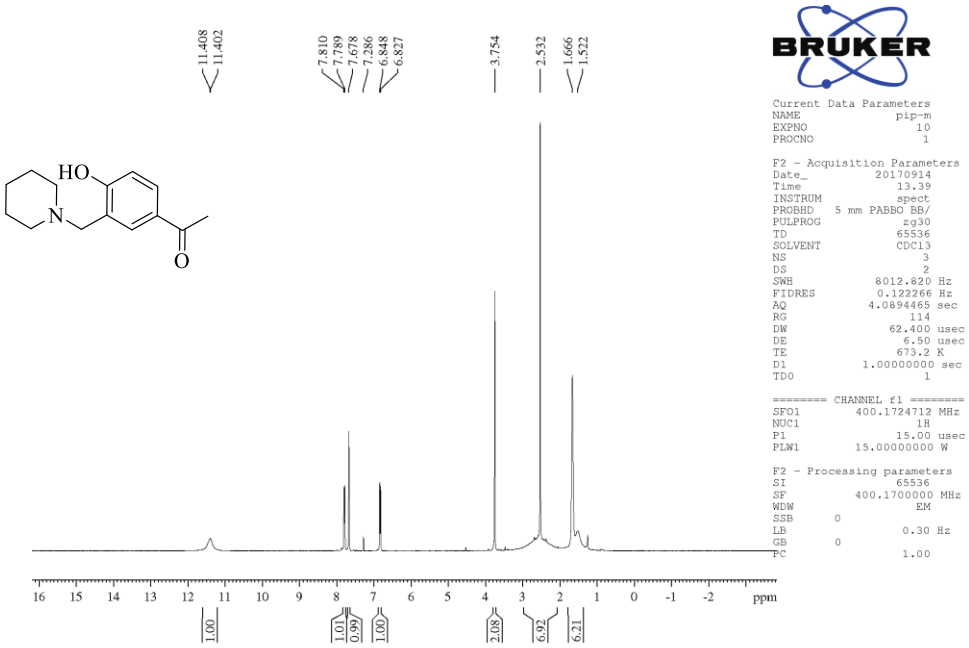


Figure S13. ^{1}HNMR spectrum compound 4a in $CDCl_3$.

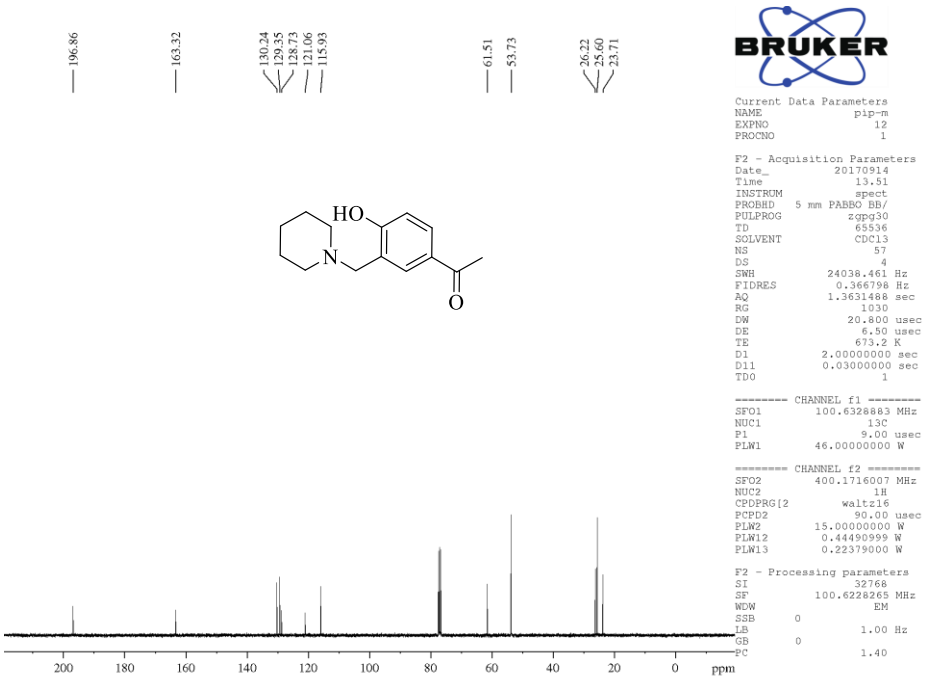


Figure S14. ^{13}C NMR spectrum compound 4a in $CDCl_3$.

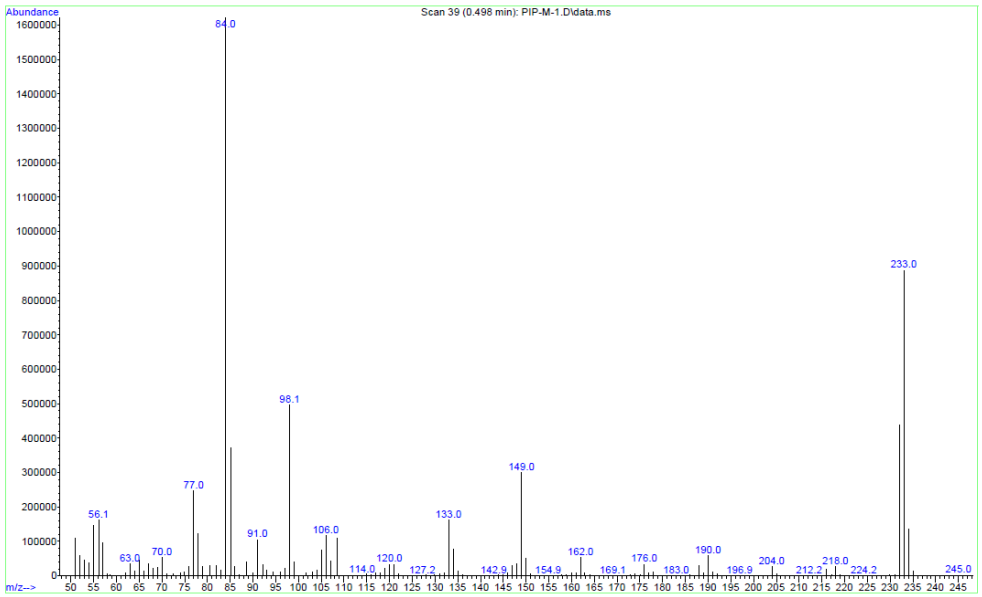
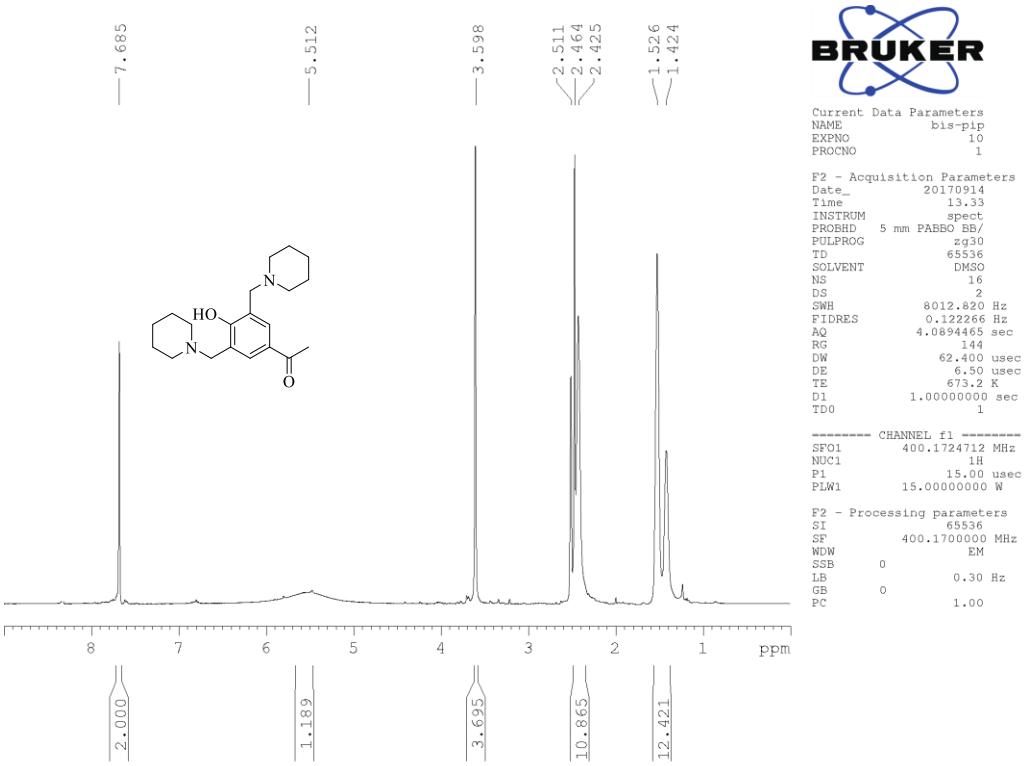


Figure S15. MS spectrum of compound 4a

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DS                          2
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FIDRES               0.122266 Hz
AQ                 4.0894465 sec
RG                        144
DW                   62.400 usec
DE                    6.50 usec
TE                    673.2 K
D1                1.00000000 sec
TD0                         1

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NUC1                       1H
P1                      15.00 usec
PLW1            15.00000000 W

F2 - Processing parameters
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Figure S16. [1]HNMR spectrum of compound 4b in DMSO-d_6

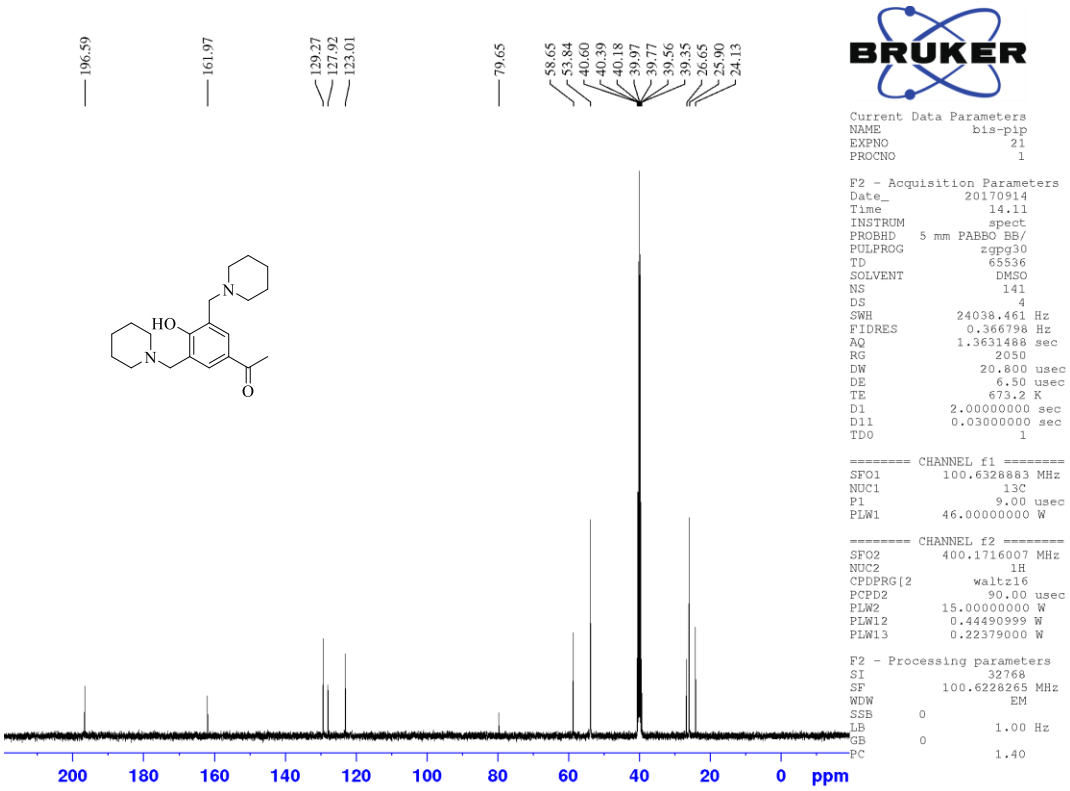

Figure S17. ^{13}C NMR spectrum of compound 4b in DMSO-d_6.

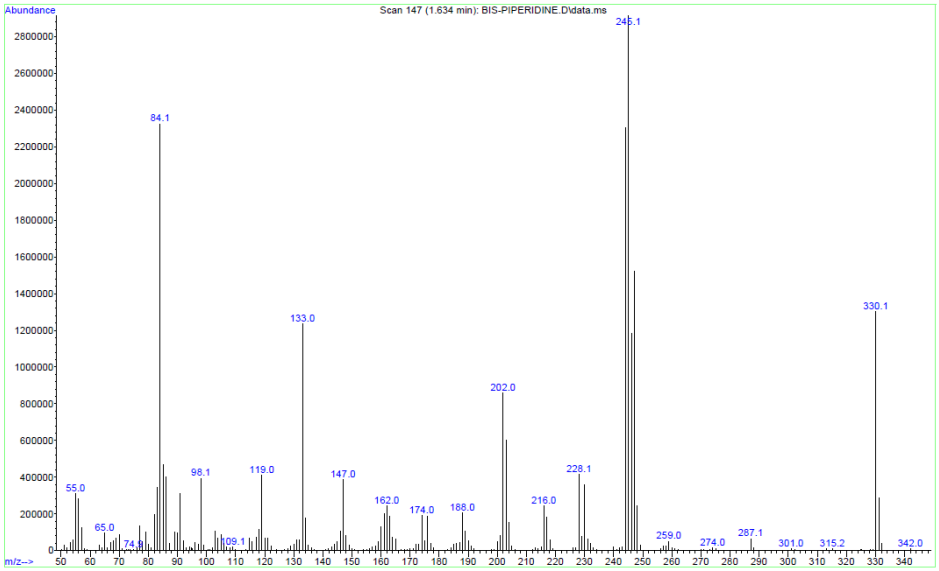

Figure S18. MS spectrum of compound 4b

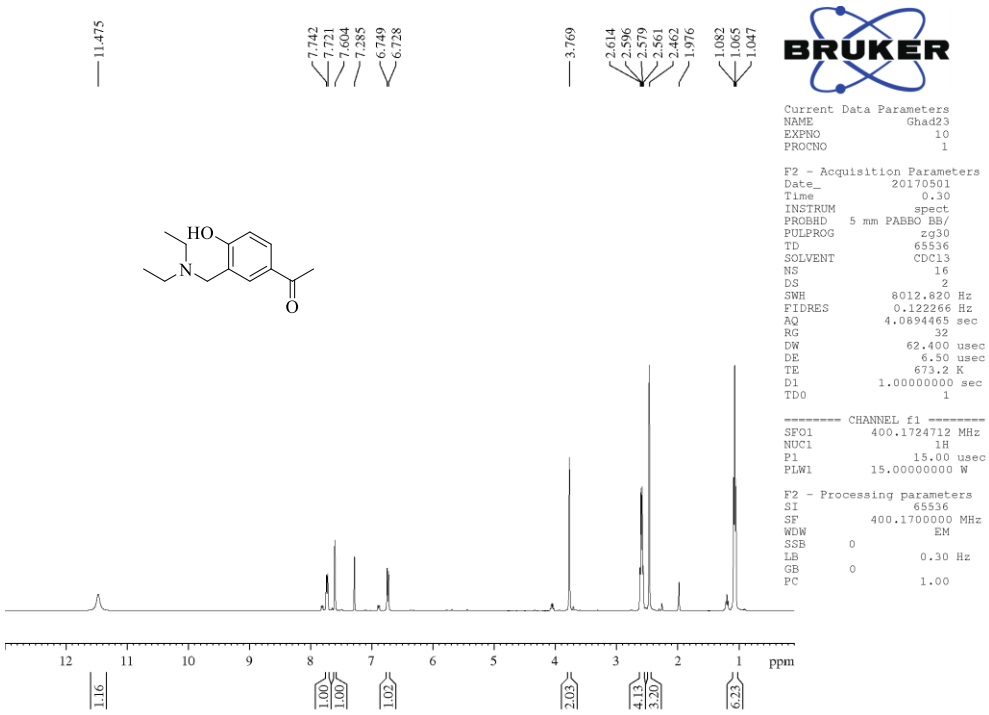

Figure S19. ^{1}HNMR spectrum of compound 5a in $CDCl_3$.

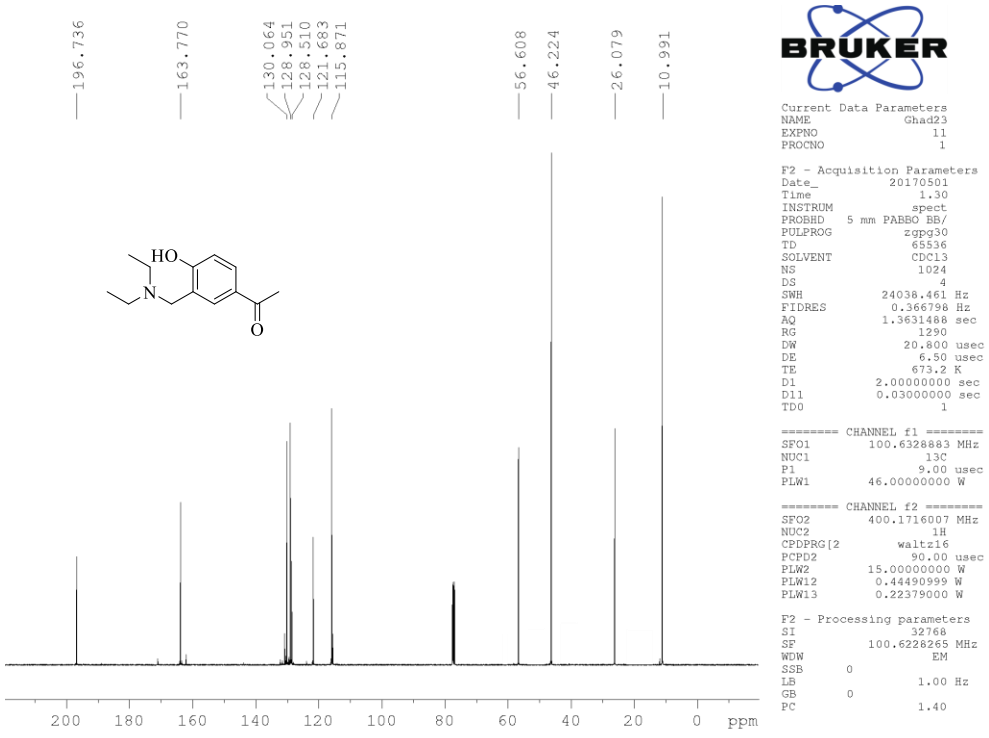

Figure S 20. ^{13}C NMR spectrum of compound 5a in $CDCl_3$.

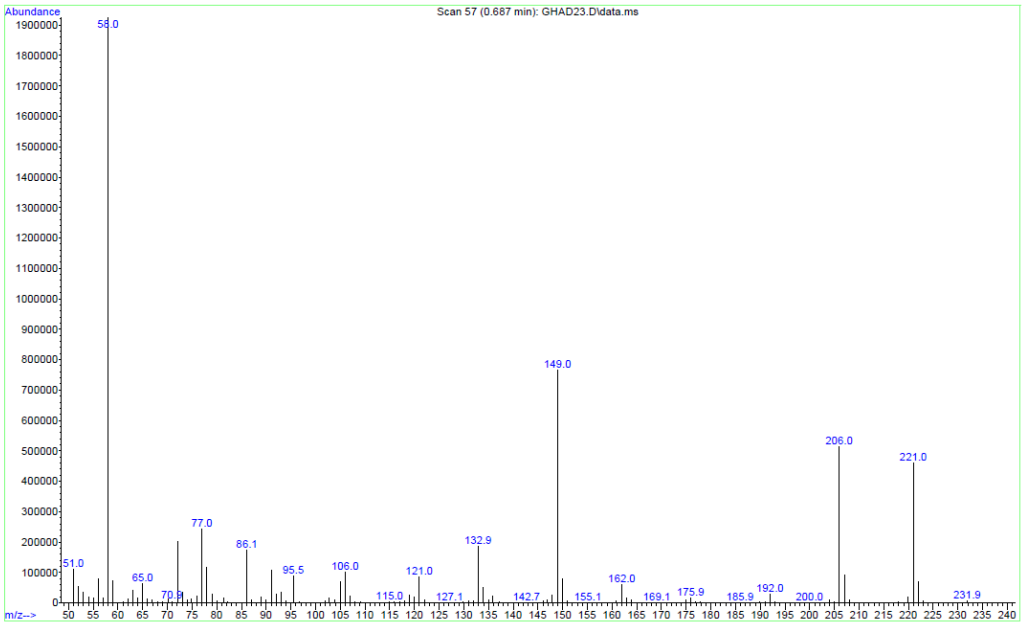

Figure S 21. MS spectrum of compound 5a