



SUPPLEMENTARY MATERIAL

***In vitro* anticancer potential of *Jasione montana* and its main components against human amelanotic melanoma cells**

Aleksandra Maria Juszcak, Robert Czarnomysy, Jakub Władysław Strawa, Marijana Zovko-Končić, Krzysztof Bielawski and Michał Tomczyk

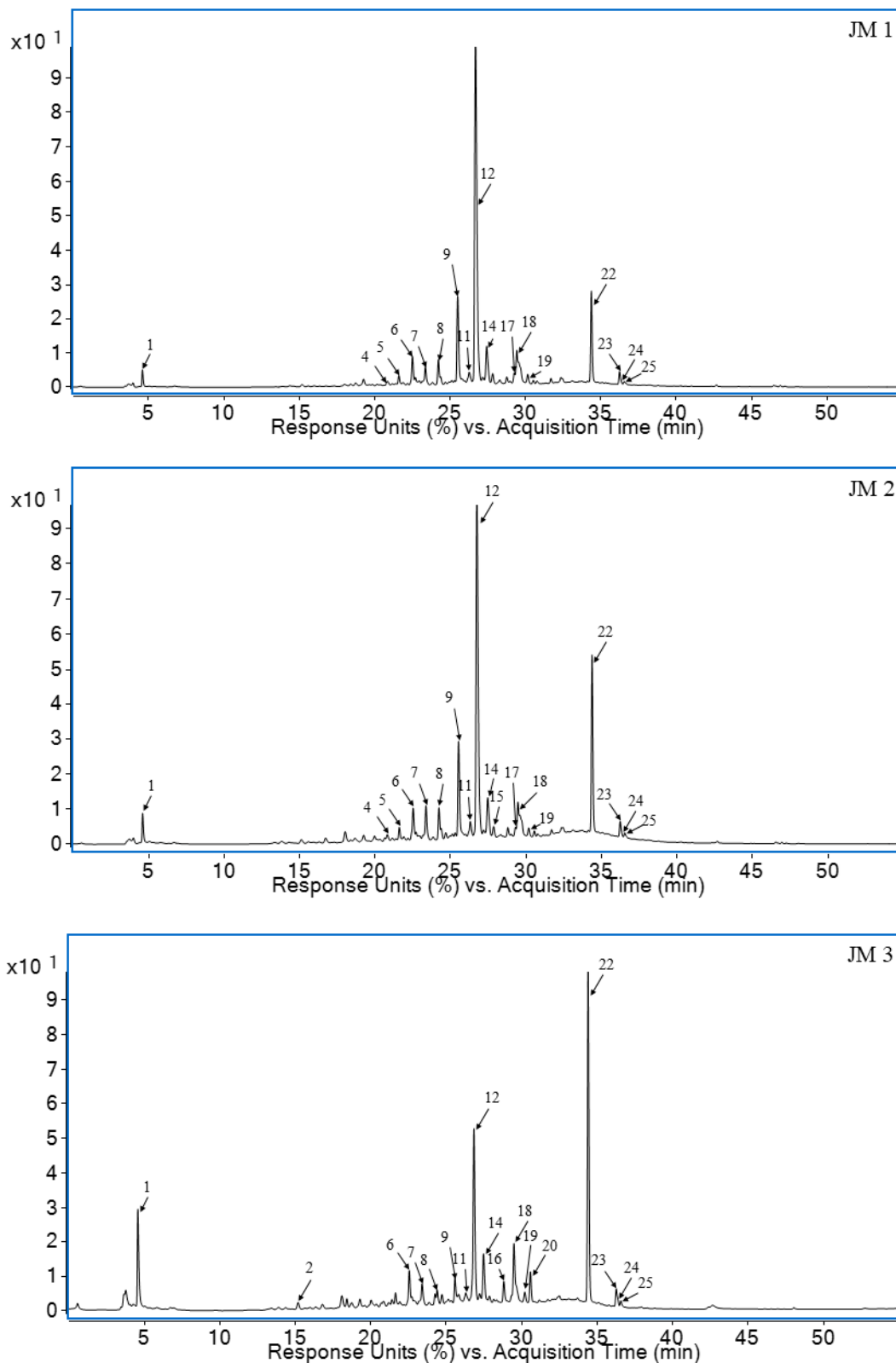


Figure S1. The qualitative assessment of *J. montana* extracts (JM1–JM3).
UV-VIS chromatogram ($\lambda=280$ nm) obtained by LC-PDA–MS.

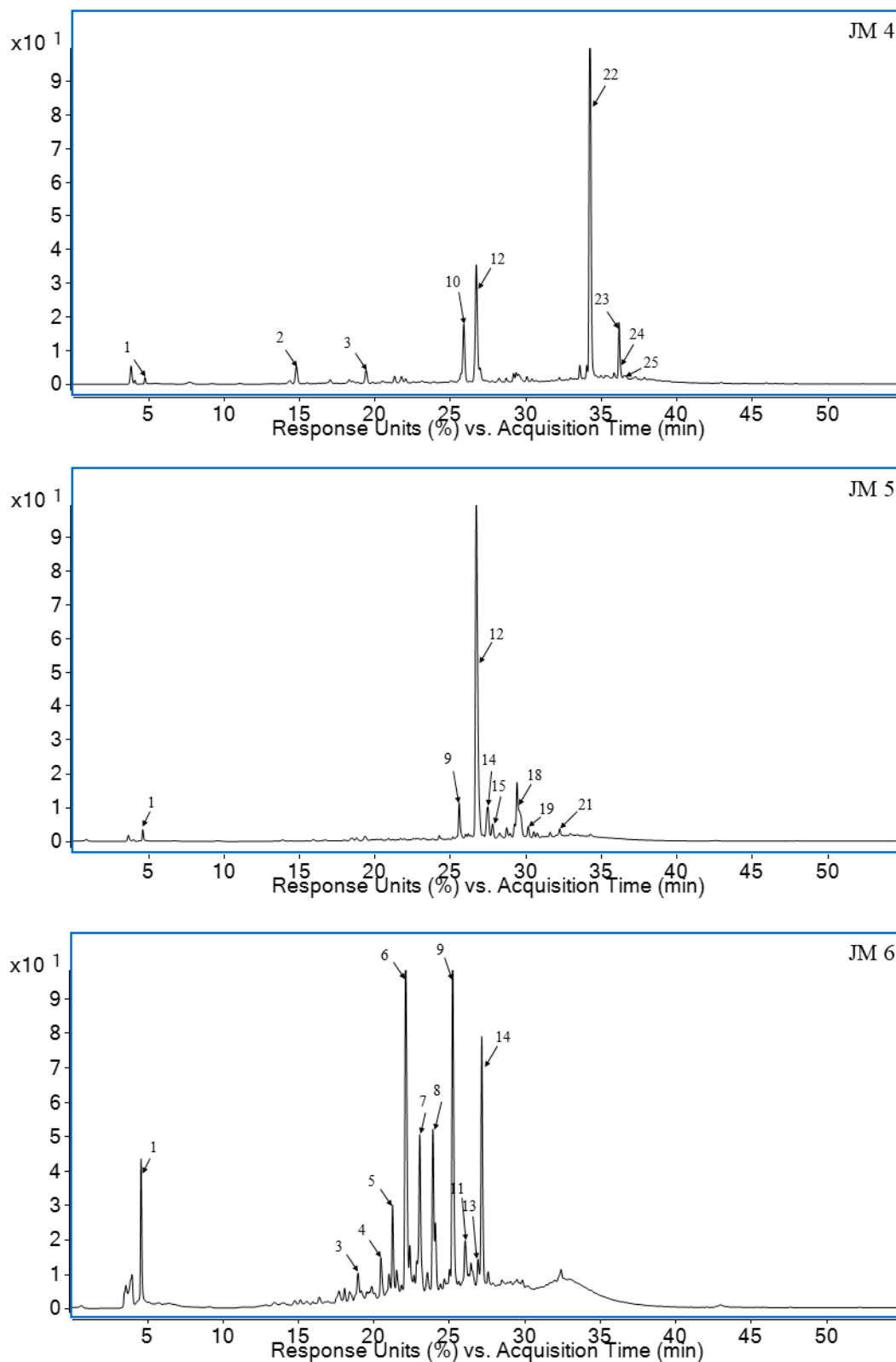


Figure S2. The qualitative assessment of *J. montana* fractions (JM4–JM6).
UV-VIS chromatogram ($\lambda=280$ nm) obtained by LC-PDA–MS.

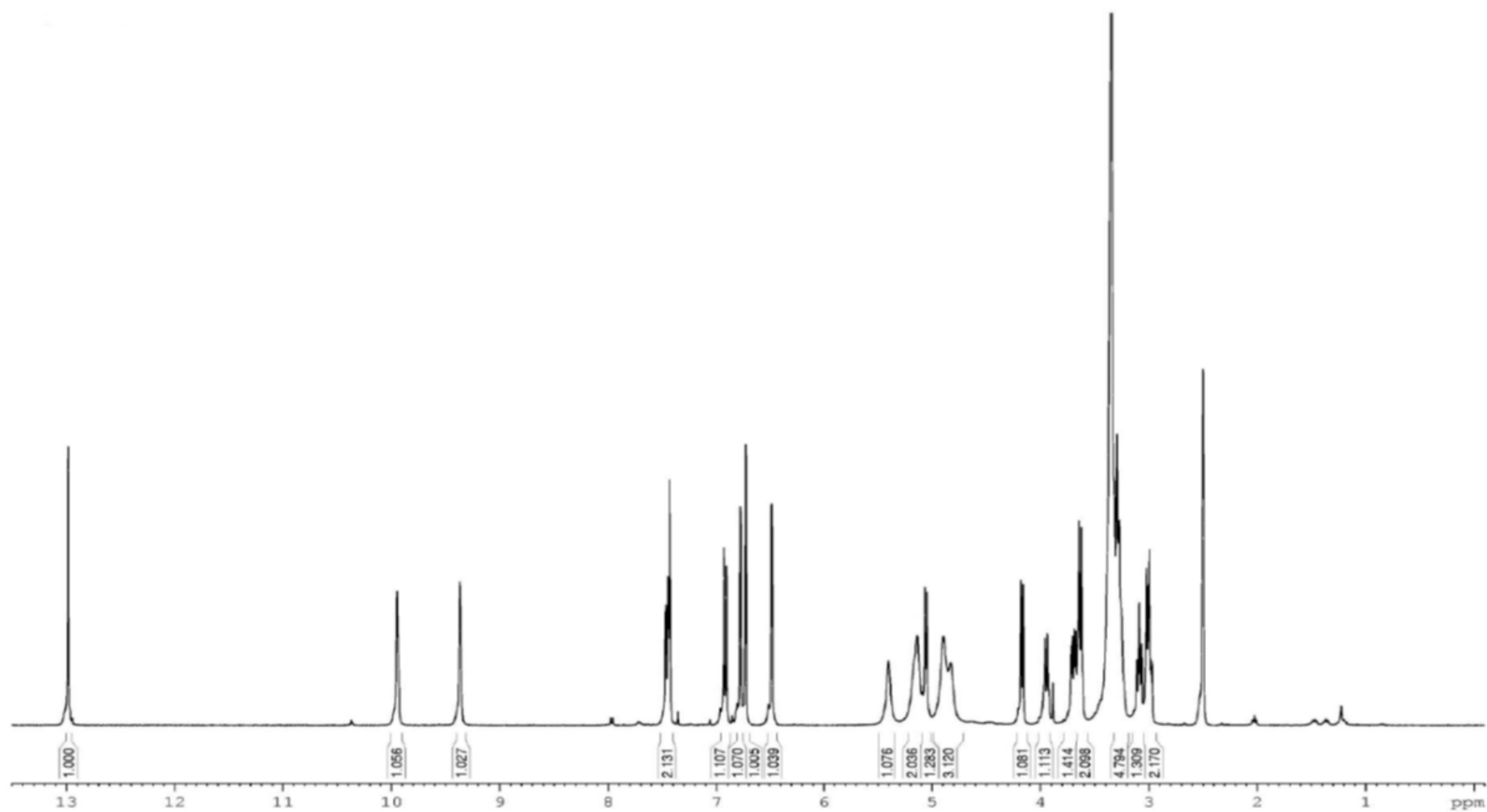


Figure S3. ^1H NMR spectrum (400.15 MHz, $\text{DMSO}-d_6$) of compound 9.

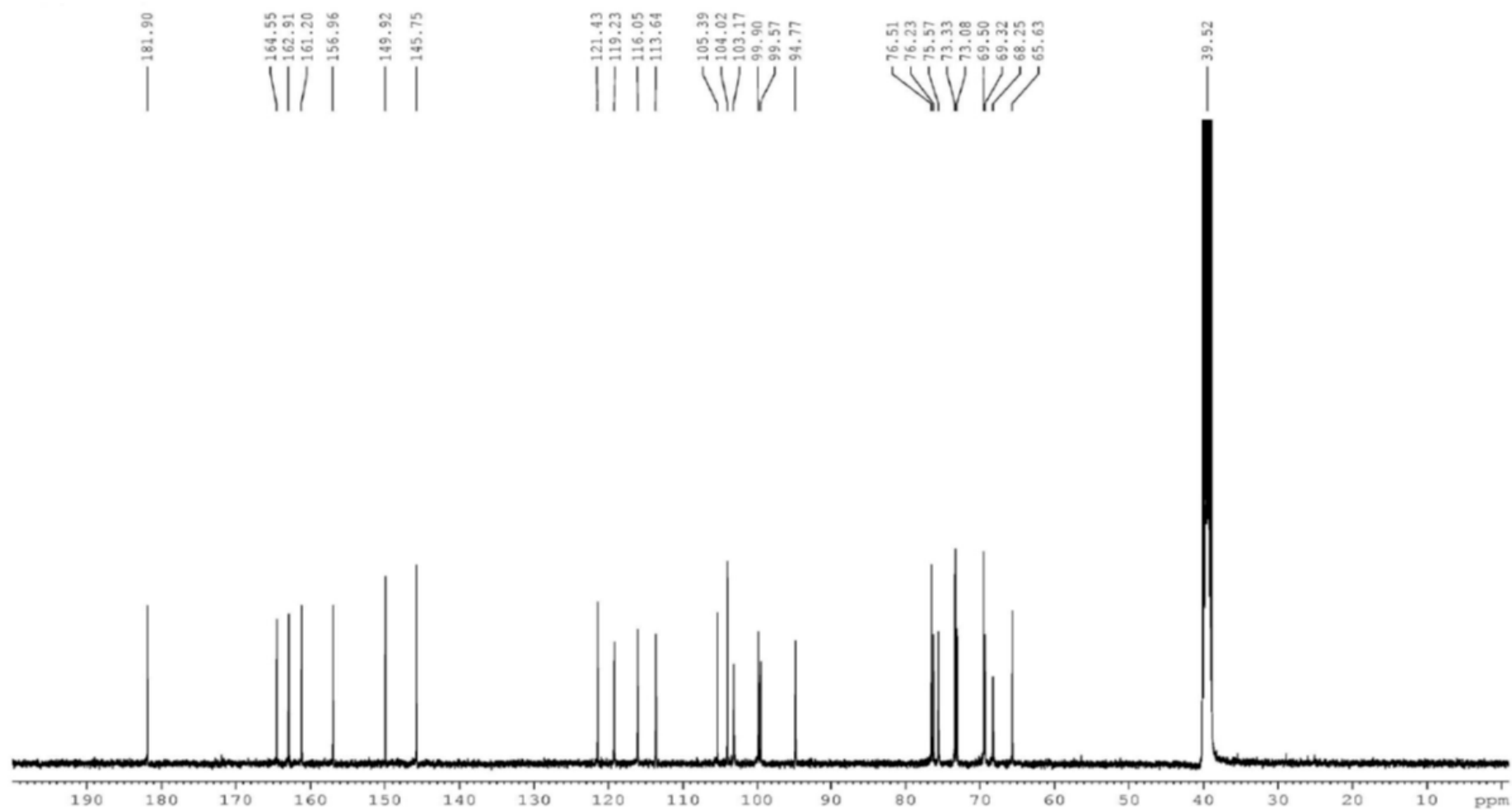


Figure S4. ^{13}C NMR spectrum (400.15 MHz, $\text{DMSO}-d_6$) of compound **9**.

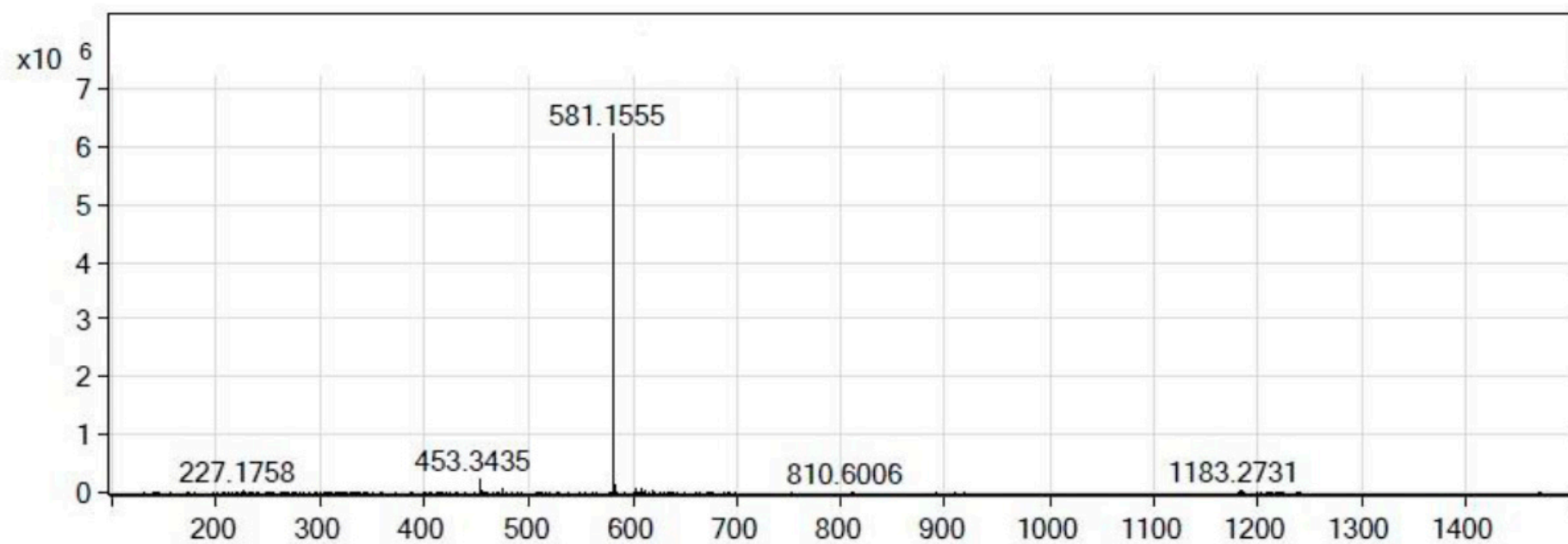


Figure S5. The MS spectrum of compound **9** in positive ion mode.

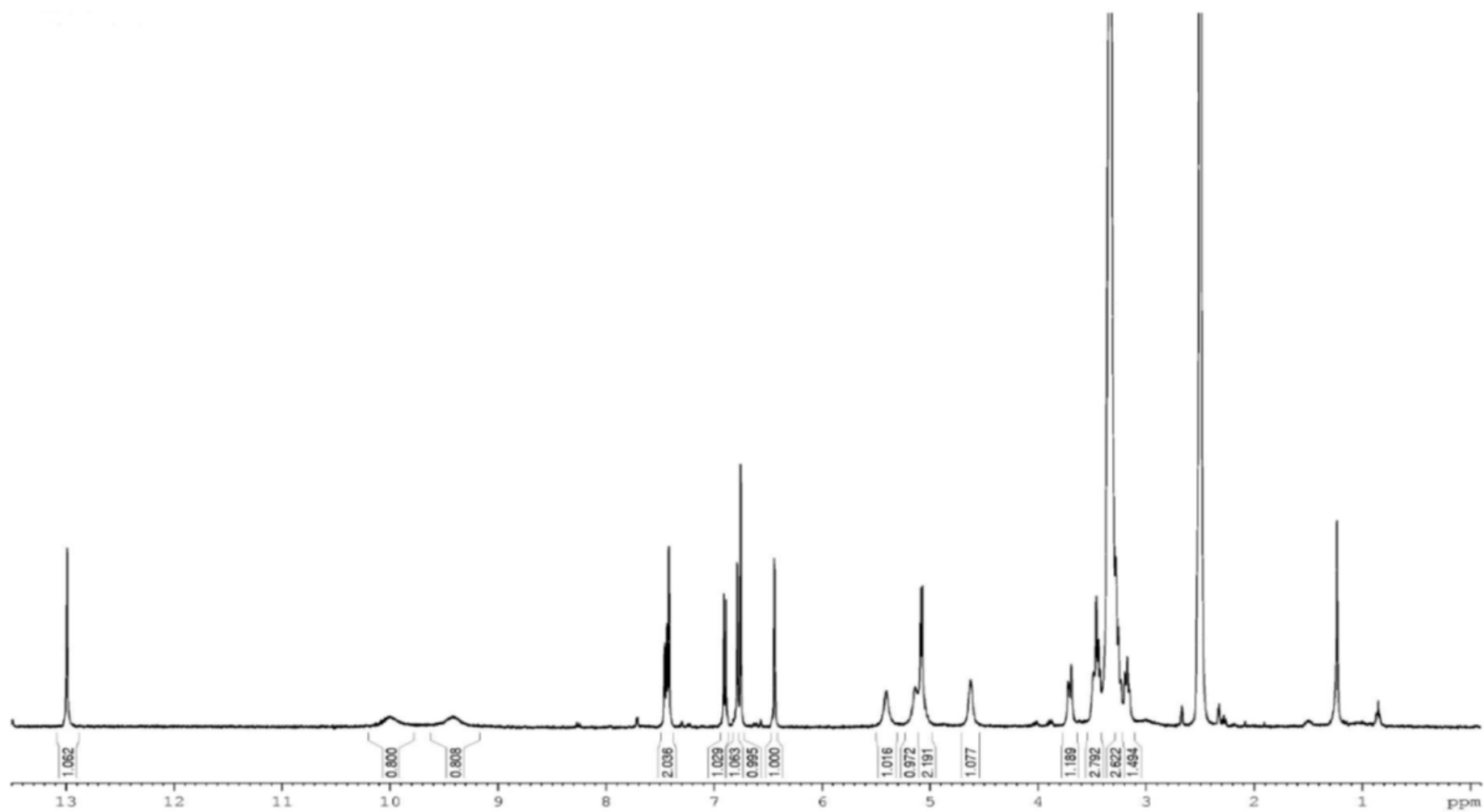


Figure S6. ^1H NMR spectrum (400.15 MHz, $\text{DMSO}-d_6$) of compound 12.

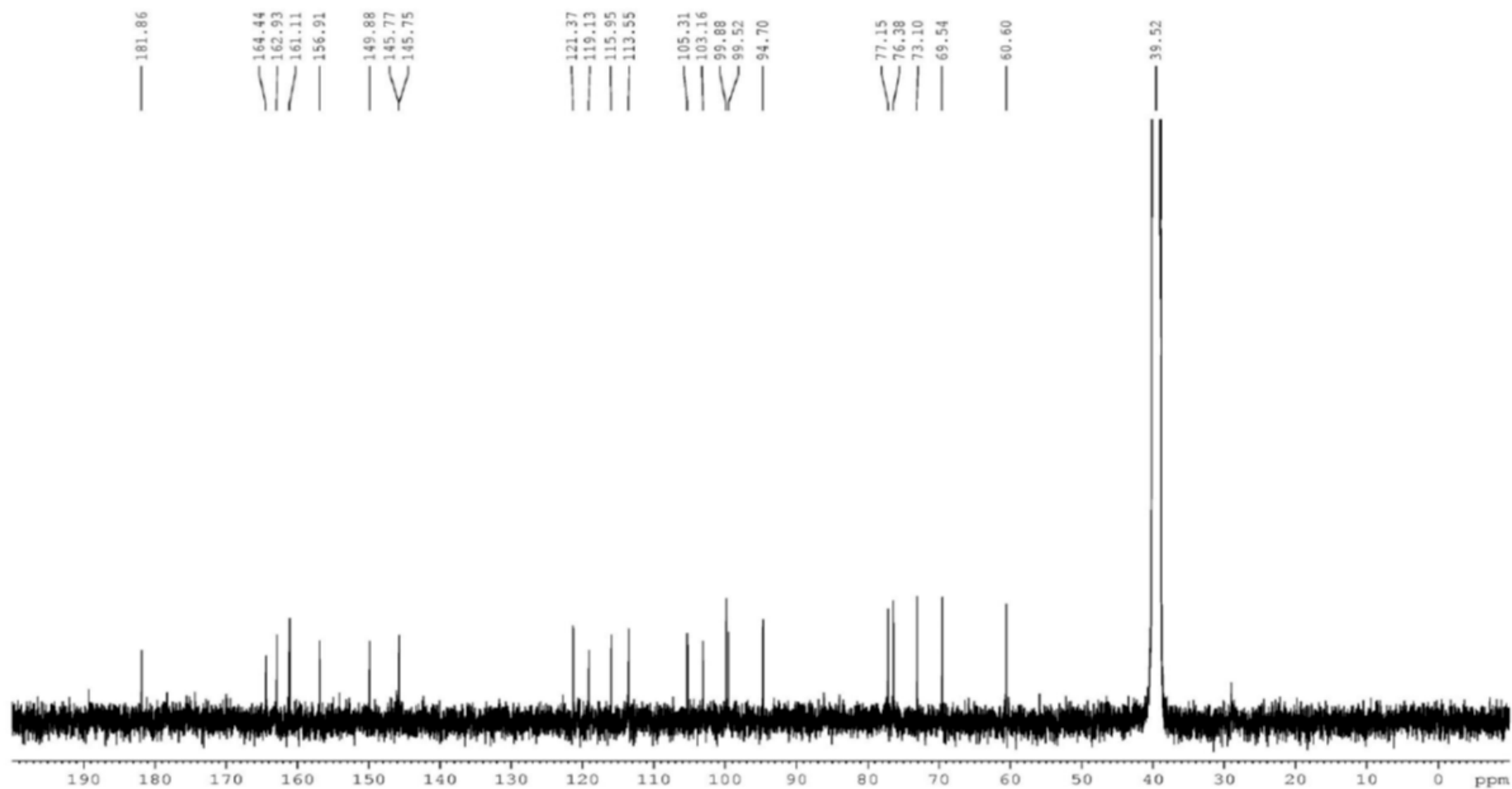


Figure S7. ¹³C NMR spectrum (400.15 MHz, DMSO-*d*₆) of compound 12.

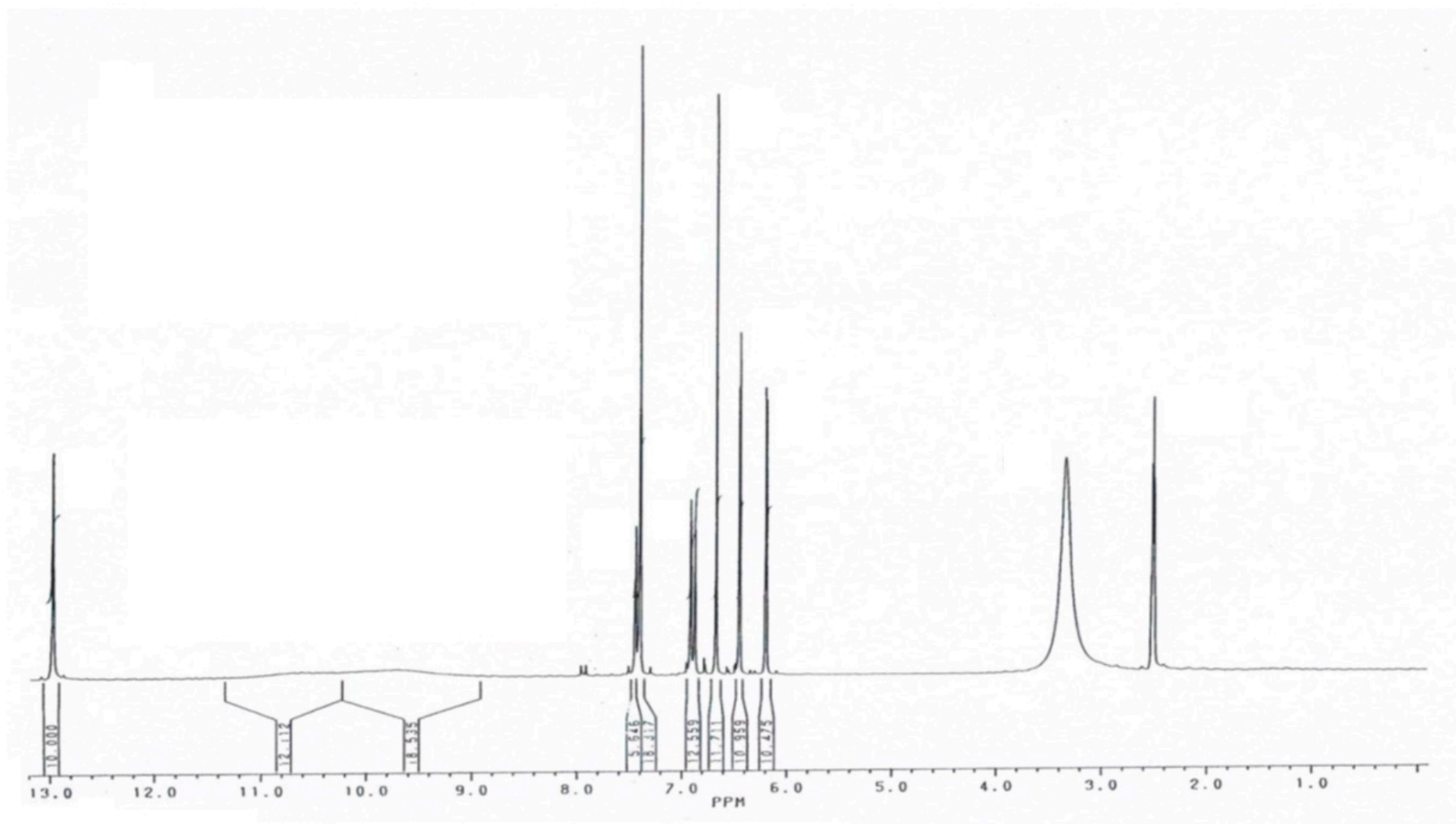


Figure S8. ^1H NMR spectrum (400.3 MHz, $\text{DMSO-}d_6$) of compound 22.

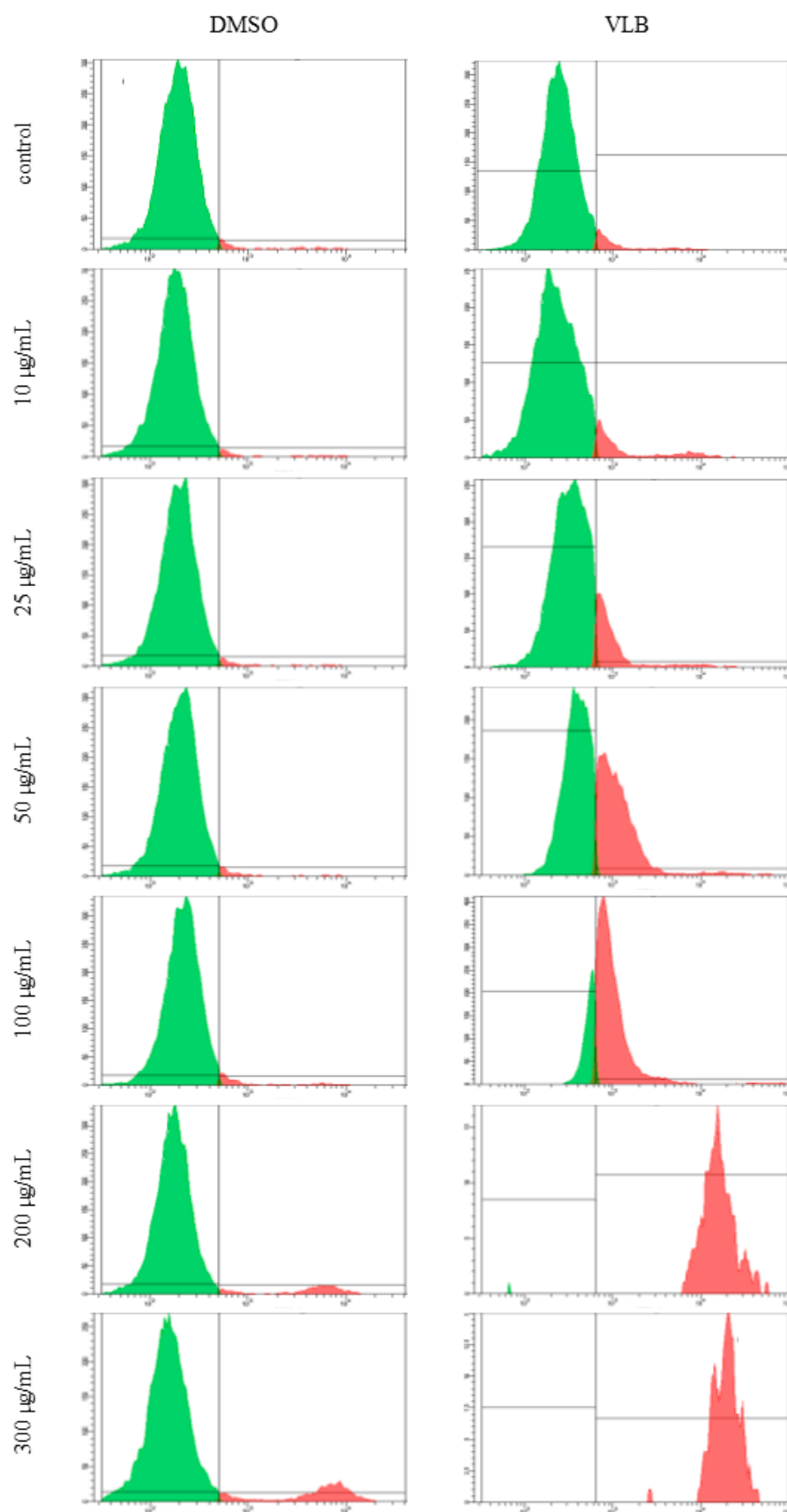


Figure S9. Flow cytometric analysis of cytotoxicity of C32 melanoma cells after 24 h of incubation with DMSO and VLB (10, 25, 50, 100, 200, and 300 µg/mL) comparable with untreated control by the Fixable Viability Stain assay.

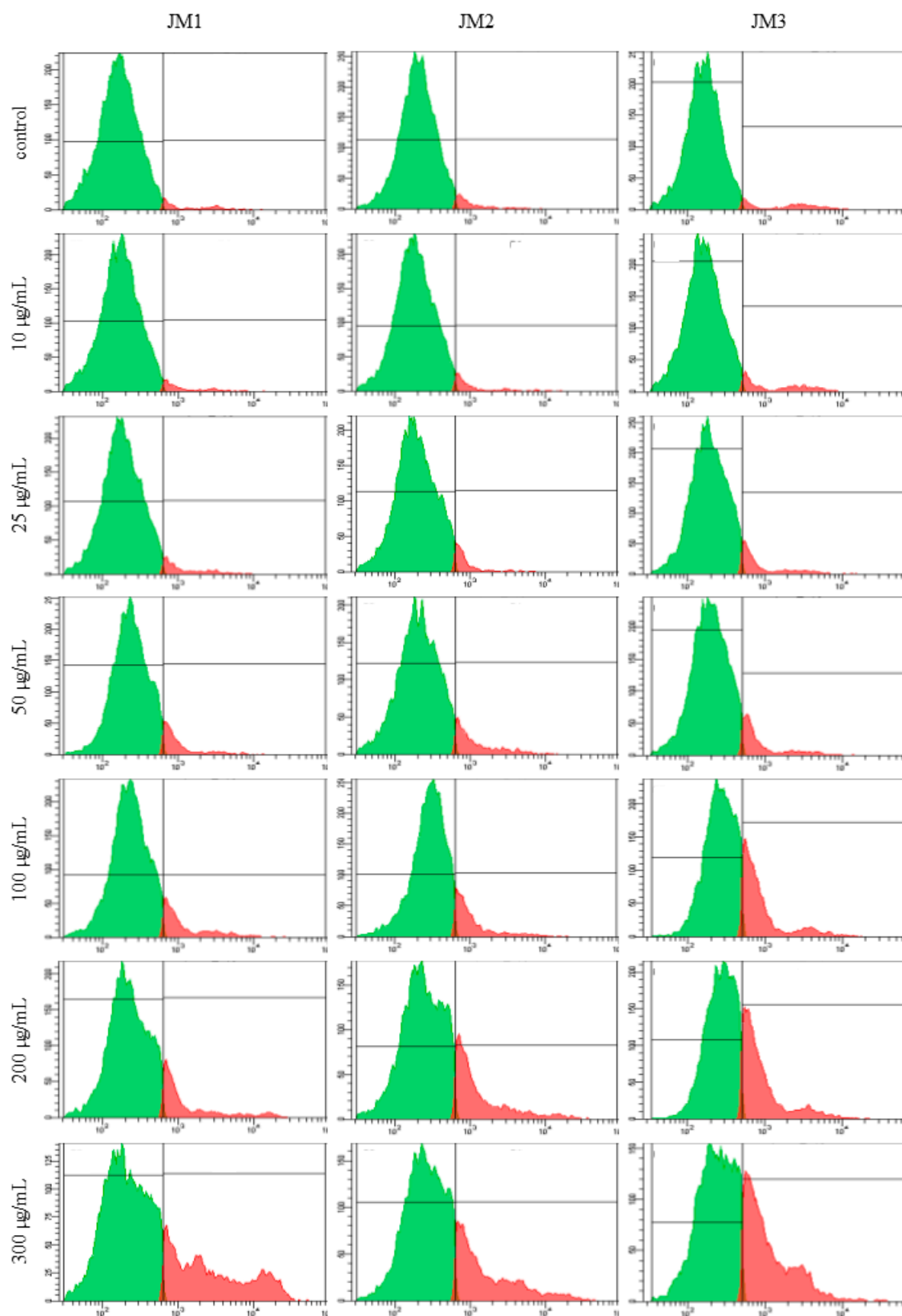


Figure S10. Flow cytometric analysis of cytotoxicity of C32 melanoma cells after 24 h of incubation with **JM1–JM3** (10, 25, 50, 100, 200, and 300 µg/mL) comparable with untreated control by the Fixable Viability Stain assay.

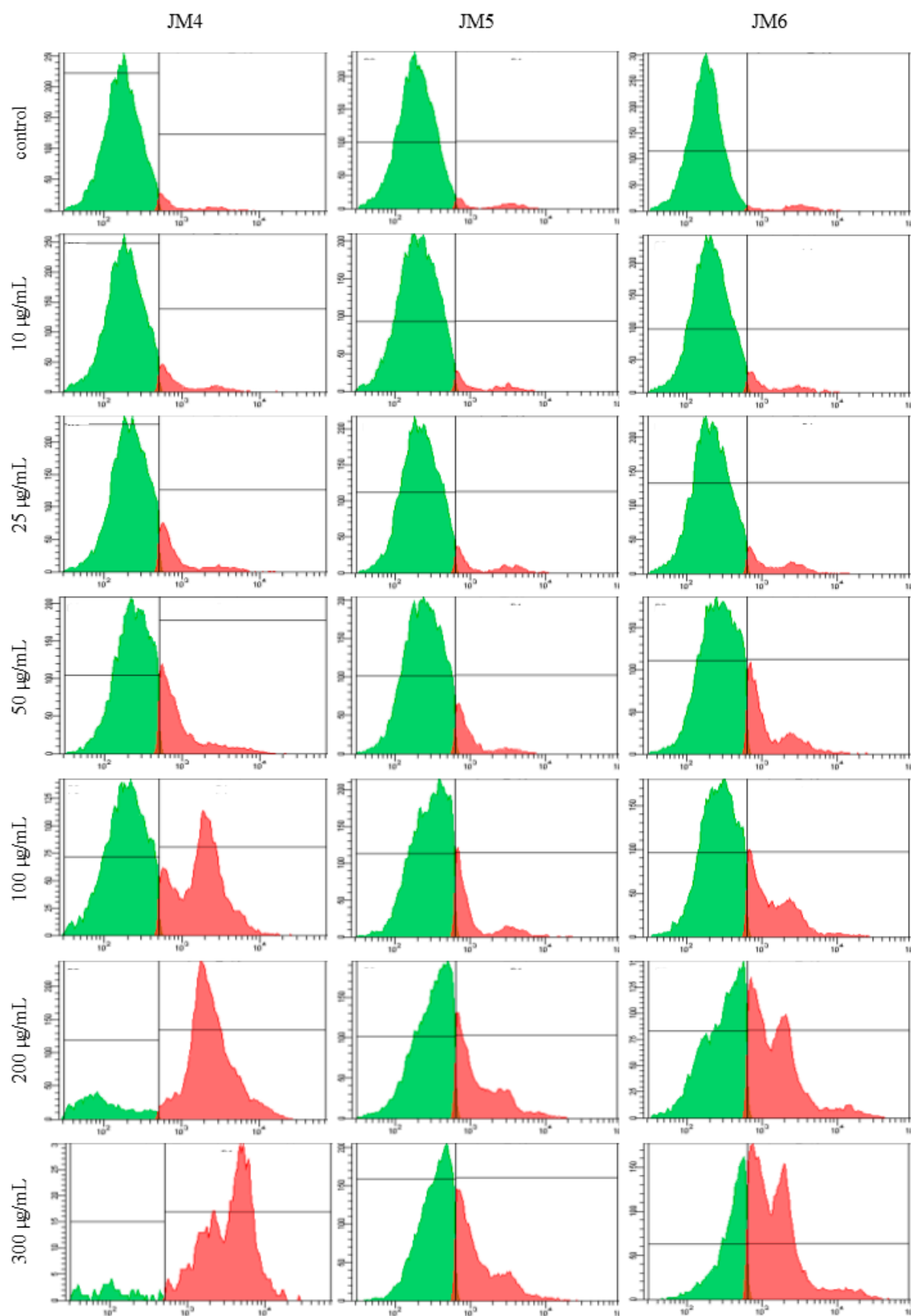


Figure S11. Flow cytometric analysis of cytotoxicity of C32 melanoma cells after 24 h of incubation with **JM4–JM6** (10, 25, 50, 100, 200, and 300 µg/mL) comparable with untreated control by the Fixable Viability Stain assay.

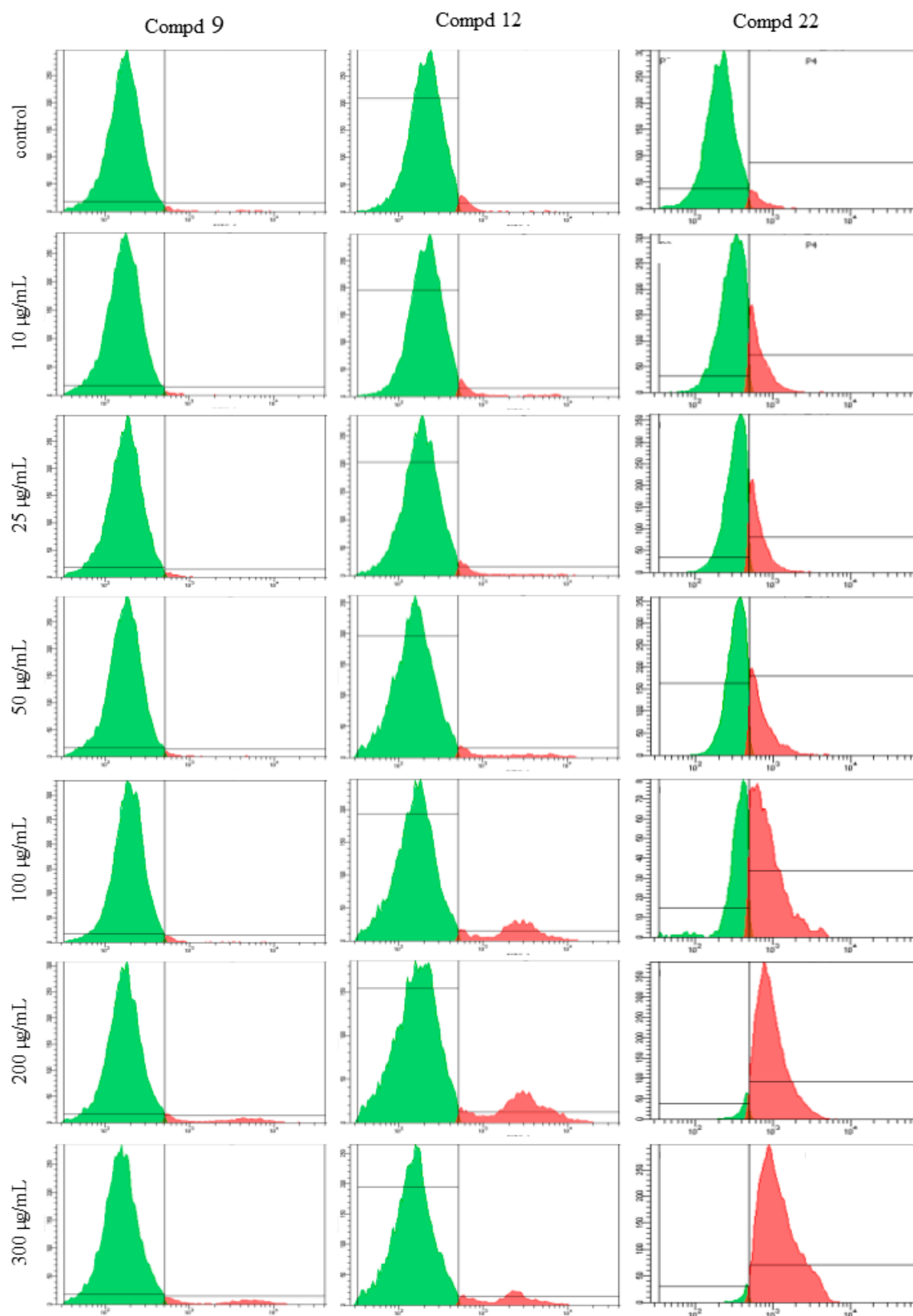


Figure S12. Flow cytometric analysis of cytotoxicity of C32 melanoma cells after 24 h of incubation with compound **9**, **12**, and **22** (10, 25, 50, 100, 200, and 300 µg/mL) comparable with untreated control by the Fixable Viability Stain assay.

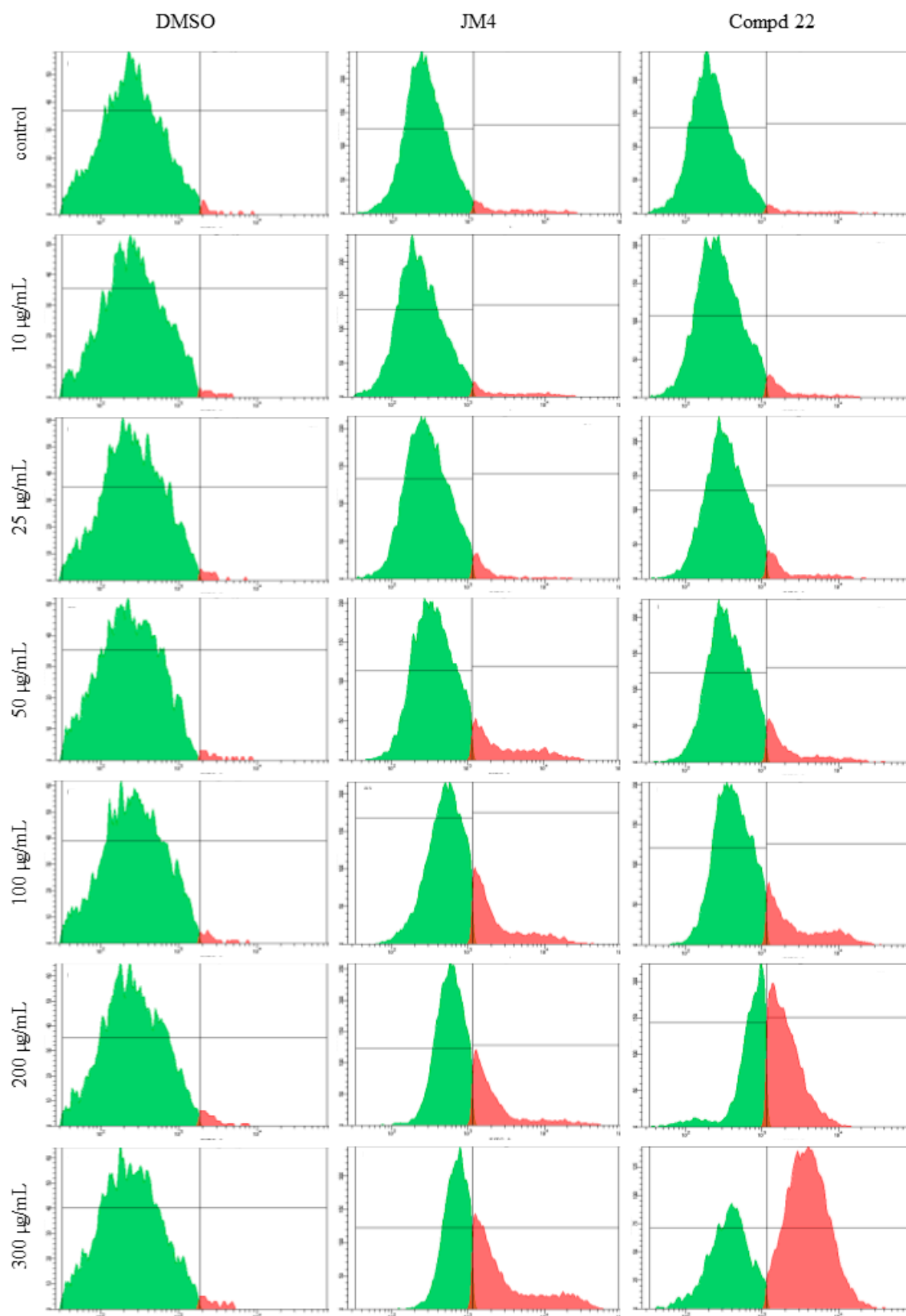


Figure S13. Flow cytometric analysis of cytotoxicity of normal human fibroblasts cells after 24 h of incubation with DMSO, **JM4**, and compound **22** (10, 25, 50, 100, 200, and 300 µg/mL) comparable with untreated control by the Fixable Viability Stain assay.