Supporting information materials of

Synergetic anticancer activity of gold porphyrin appended to phenyl tin malonate organometallic complexes

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Figure S1. Overlay of the IR spectra of compounds **1** and **SnPh**₂ recorded in solid state by ATR-IR spectroscopy. The asterisks indicate the carboxylate antisymmetric (v_{as}) and symmetric (v_s) and Sn-O stretching bands.



Figure S2. Overlay of the IR spectra of compounds 1 and Sn_2Ph_6 recorded in solid state by ATR-IR spectroscopy. The asterisks indicate the O-C=O antisymmetric (v_{as}) and symmetric (v_s) and Sn-O stretching bands.



Figure S3. Overlay of the IR spectra of compounds SnPh₂ and AuP-SnPh₂ recorded in solid state by ATR-IR spectroscopy. The asterisks indicate the carboxylate antisymmetric (v_{as}) and symmetric (v_s) and Sn-O stretching bands.



Figure S4. Overlay of the IR spectra of compounds Sn_2Ph_6 and $AuP-Sn_2Ph_6$ recorded in solid state by ATR-IR spectroscopy. The asterisks indicate the carboxylate antisymmetric (v_{as}) and symmetric (v_s) and Sn-O stretching bands.



Figure S5. Overlay of the UV-Vis. absorption spectra of compounds Sn₂Ph₆ and Sn₂Ph₆ recorded in dichloromethane.



Figure S6. ¹H NMR spectrum of SnPh₂ recorded in DMSO-d₆.



Figure S7. ¹H NMR spectrum of Sn₂Ph₆ recorded in DMSO-d₆.



Figure S8. ¹H NMR spectrum of AuP-SnPh₂ recorded in DMSO-d₆.



Figure S9. ¹H NMR spectrum of AuP-Sn₂Ph₆ recorded in DMSO-d₆.



Figure S10. ES-MS spectrum of SnPh₂.



Figure S11. Zoom on the experimental (up) and calculated (down) isotopic fragment M+Na⁺ of ES-MS spectrum of **SnPh**₂



Figure S12. ES-MS spectrum of AuP-SnPh₂.



Figure S13. Zoom on the experimental (up) and calculated (down) isotopic fragment M⁺ of ES-MS spectrum of **AuP-SnPh**₂.



Figure S14. ES-MS spectrum of AuP-Sn₂Ph₆.



Figure S15. Zoom on the experimental (up) and calculated (down) isotopic fragment M⁺ of ES-MS spectrum of **AuP-Sn₂Ph**₆