Electronic Supplementary Information

Synthesis, characterisation, and *in vitro* anticancer activity of catalytically active indole-based half-sandwich complexes.

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Figure S1: ¹H NMR stability studies

Figures S2 – S9: ¹H and ¹³C NMR spectra of complexes **1 – 4**

Figures S10 – S13: High-resolution ESI mass spectrum of complexes 1 – 4

Figure S14: IC₅₀ graphs for the ind-py ligand against PNT2, A2780, and A2780cisR.



Figure 1. ¹H NMR (400 MHz) stability studies of complex **1** (A), **2** (B), **3** (C), **4** (D), and Ind-Py (E) in MeOD/D₂O (1:1 v/v, 1.1 mM, 298 K) and complexes **1** (E), **2** (F), **3** (G) and **4** (H) in *d*₆-DMSO (1.1 mM, 298 K). The low signal-to-noise ratio for (B) is due to the poor solubility of complex **2** in MeOD/D₂O.



Figure S3. $^{\rm 13}{\rm C}$ NMR spectrum of complex ${\bf 1}$



Figure S5. ¹³C NMR spectrum of complex **2**



Figure S7. ¹³C NMR spectrum of complex **3**



Figure S9. ¹³C NMR spectrum of complex **4**



Figure S10. High-resolution ESI mass spectrum of complex 1



Figure S11. High-resolution ESI mass spectrum of complex 2



Figure S12. High-resolution ESI mass spectrum of complex 3



Figure S13. High-resolution ESI mass spectrum of complex 4



Figure S14. IC $_{50}$ graphs for the ind-py ligand against PNT2, A2780, and A2780cisR.