Supporting Information

Nuclease activity and protein-binding properties of a novel tetranuclear thiosemicarbazide Pt(II) complex

Jia Shao, Wei-Guo Bao, He Tian, Bing Li, Xiao-Fei Zhao, Xin Qiao, Jing-Yuan Xu*

Tianjin Key Laboratory on Technologies Enabling Development of Clinical Therapeutics and Diagnostics (Theranostics), School of Pharmacy, Tianjin Medical University, Tianjin 300070, P. R. China.

* E-mail: xujingyuan@tmu.edu.cn

Figure captions:

Figure S1 IR spectrum of HAm4M.

Figure S2 IR spectrum of complex 1.

- Figure S3 (a) ESI-MS spectrum of [Pt₄(Am4M)₄] (1) showing the parent ion peak in MeOH.
 The peak corresponds to M+H⁺ species. (b) ESI-MS spectrum of [Pt₄(Am4M)₄] (1) showing an enlarged view.
- **Figure S4** Packing diagram of **1** viewed along the crystallographic [101] showing (a) weak coordinated bonds and (b) extensive π - π stackings.
- Figure S5 Absorption spectral traces of 1 in the buffer solution at five different time points (0 min, 30 min, 60 min, 90min and 3h). There was no significant spectral change during the spectral measurements.
- Figure S6 Gel electrophoresis diagrams showing the cleavage of SC pBR322 DNA (40 ng) incubated with 1 in a buffer containing 10% DMF at 37 °C for 2 h. Lane 1: DNA control; Lanes 2–8: DNA + 1 (1, 2, 3, 4, 5, 6 and 7 μM, respectively.)
- Figure S7 Gel electrophoresis diagrams showing the cleavage of SC pUC19 DNA (100 ng) incubated with 1 at 37 °C for 1 h. D₂O was used for dilution of 1 to a volume of 20 μL for lane 3 and 5. Lane 1: DNA control; Lane 2: DNA + 1 (4 μM in H₂O); Lane 3: DNA + 1 (4 μM in D₂O); Lane 4: DNA + 1 (5 μM in H₂O); Lane 5: DNA + 1 (5 μM in D₂O).

Figure S8 Fluorescence spectra of BSA $[c(BSA) = 3 \times 10^{-5} \text{ M}]$ in the various concentrations of **1** $[c(\text{complex 1}) = 0-5.96 \times 10^{-6} \text{ M}]$ at 288 K (a) and 308 K (b), respectively.

Table S1 The cleavage of SC pBR322 DNA (40 ng) by $[Cu({}^{t}BuPhimp)(Cl)]$ and 1 (μ M) at 37

°C for 2 h. (- negitive, + positive)



Figure S1 IR spectrum of HAm4M.



Figure S2 IR spectrum of complex 1.







(b)

Figure S3 (a) ESI-MS spectrum of $[Pt_4(Am4M)_4]$ (1) showing the parent ion peak in MeOH. The peak corresponds to M+H⁺ species; (b) ESI-MS spectrum of $[Pt_4(Am4M)_4]$ (1) showing an enlarged view.



(a)



Figure S4 Packing diagram of **1** viewed along the crystallographic [101] showing (a) weak coordinated bonds and (b) extensive π - π stackings.



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Figure S7 Gel electrophoresis diagrams showing the cleavage of SC pUC19 DNA (100 ng) incubated with **1** at 37 °C for 1 h. D₂O was used for dilution of **1** to a volume of 20 μ L for lane 3 and 5. Lane 1: DNA control; Lane 2: DNA + **1** (4 μ M in H₂O); Lane 3: DNA + **1** (4 μ M in D₂O); Lane 4: DNA + **1** (5 μ M in H₂O); Lane 5: DNA + **1** (5 μ M in D₂O).



(a)



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Table S1 The cleavage of SC pBR322 DNA (40 ng) by [Cu(^t BuPhimp)(Cl)] and 1 (μ M) at 37 ^c	'C for
2 h. (- negitive, + positive)		

	[Cu(^t BuPhimp)(Cl)]	SC disappear	1	SC disappear
	(μM)		(µM)	
1	0	0	1	-
2	0	0	3	+
3	5	-	5	+
4	10	-	7	+
5	20	-		
6	50	+		
7	100	+		