

Supporting Information

ROS-mediated multi-targeting anticancer mechanisms of copper

(II) 2-hydroxy-1-naphthaldehyde complexes

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X-ray crystallography C1 and C2

Single crystals of C1 and C2 were selected and analyzed by Burker APEX-II CCD diffractometer. The crystals were kept at 296.15K during data collection. The structures were solved by using Olex2. The crystals data of two complexes are listed in tableS1. The selected bond lengths (Å) and bond angles (°) of crystal structure are listed in Table S1, S2.

Table S1. Selected bond length (Å) in complexes C1 and C2.

Identification Cod	C1 Bond Length (Å)	C2 Bond Length (Å)
Cu1-O1	1.895	1.917
Cu1-N1	1.946	2.023
Cu1-N2	2.009	2.023
Cu1-N3	----	1.927
Cu1-Br1	----	2.8458
Cu1-Cl1	2.236	----

Table S2. Selected bond angles (°) in complexes C1 and C2.

C1		C2	
Identification Code	Bond Angles (°)	Identification Code	Bond Angles (°)
O-Cu-Cl1	89.93	O-Cu-Br	108.14
O-Cu-N2	173.69	N2-Cu-Br	95.61
O-Cu-N1	91.76	N2-Cu-O	87.53
N2-Cu-Cl1	95.82	N3-Cu-Br	88.34
N1-Cu-Cl1	176.72	N3-Cu-O	91.45
N2-Cu-N1	82.63	N3-Cu-N2	176.04
		N1-Cu-Br	93.22
		N1-Cu-O	157.68
		N1-Cu-N3	82.56

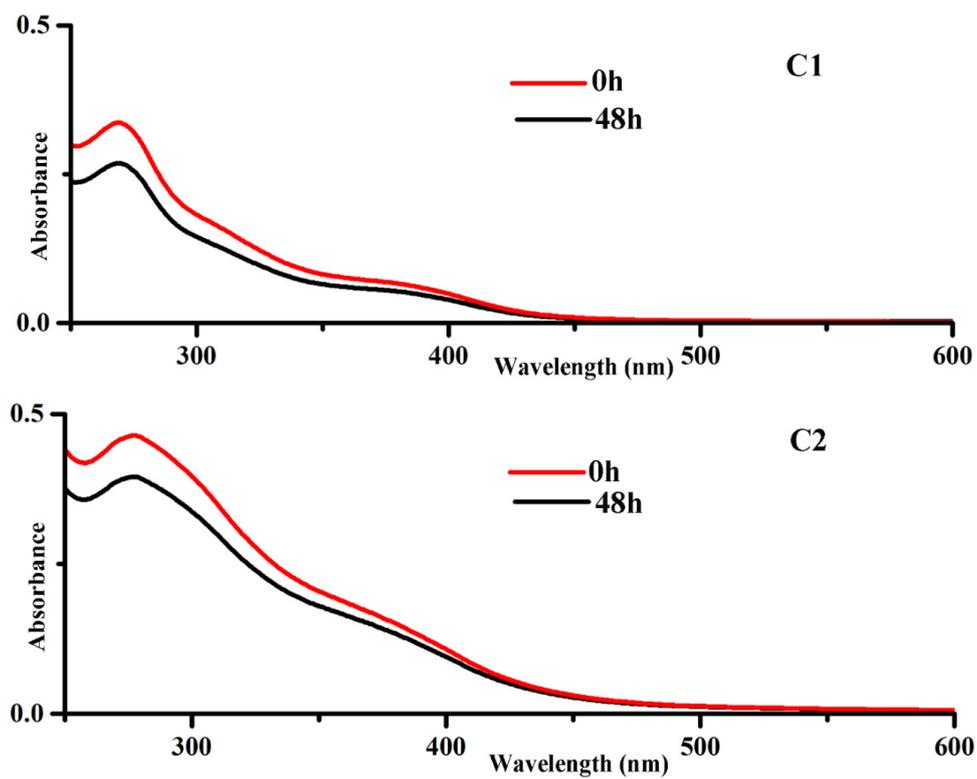


Figure S1. UV-Vis spectra of the complexes' stability (C1, C2).

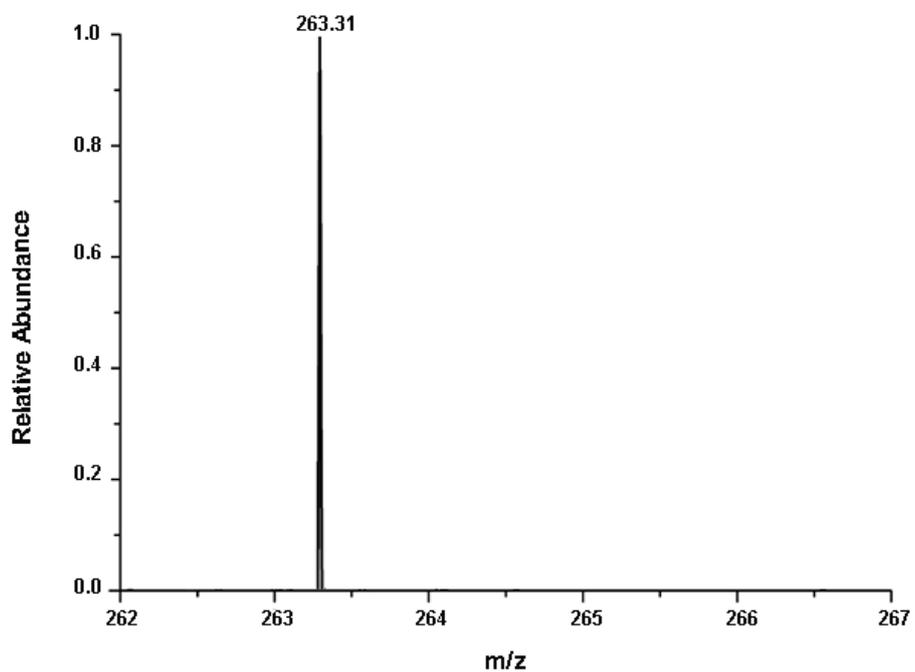


Figure S2. ESI-mass spectrum of the Ligand showing an intense signal at m/z: 263.31 for $C_{17}H_{14}N_2O, [H]^+$.

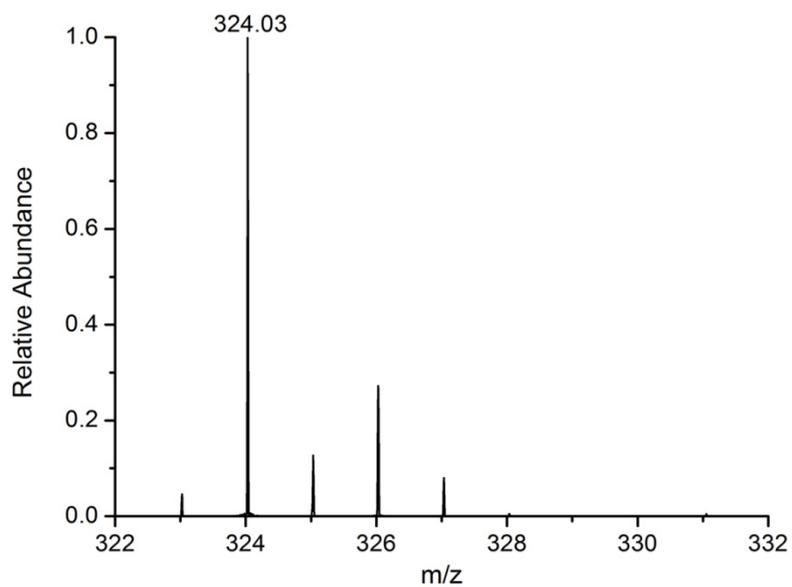


Figure S3. ESI-mass spectrum of C1 showing an intense signal at m/z : 324.03 for $C_{17}H_{13}ClCuN_2O [M - Cl]^+$.

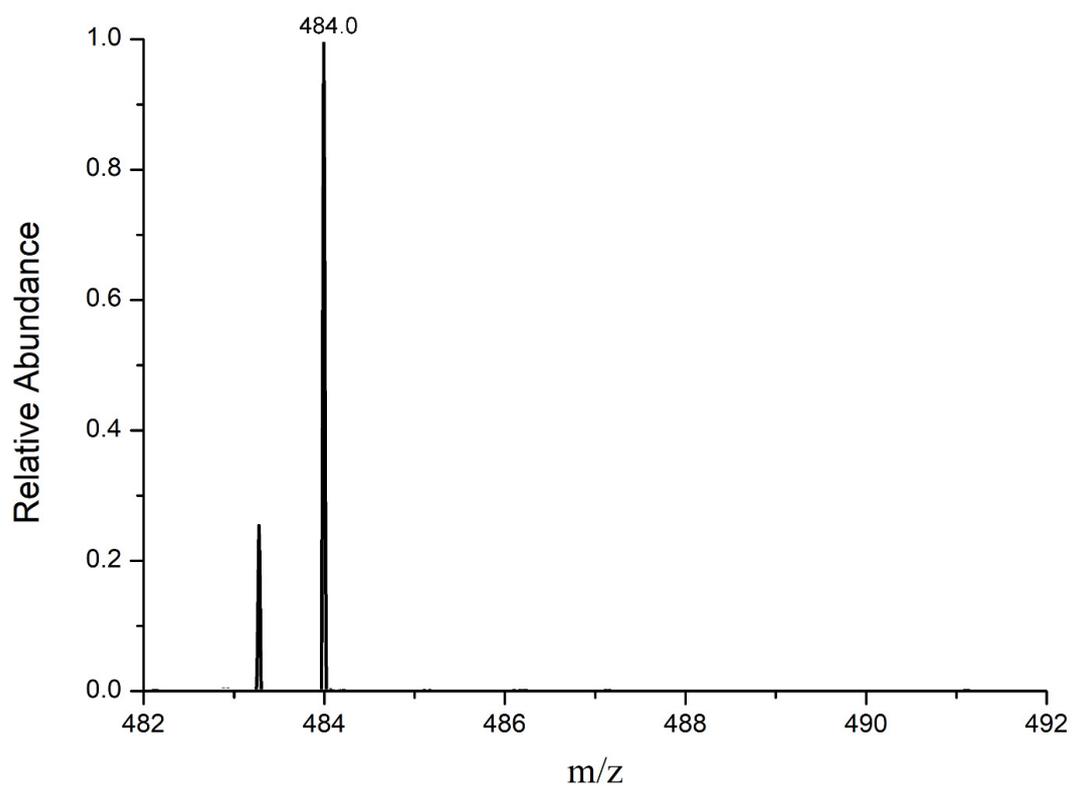


Figure S4. ESI-mass spectrum of the C2 showing an intense signal at m/z : 484.0 for $C_{22}H_{18}BrCuN_3O[M+H]^+$.

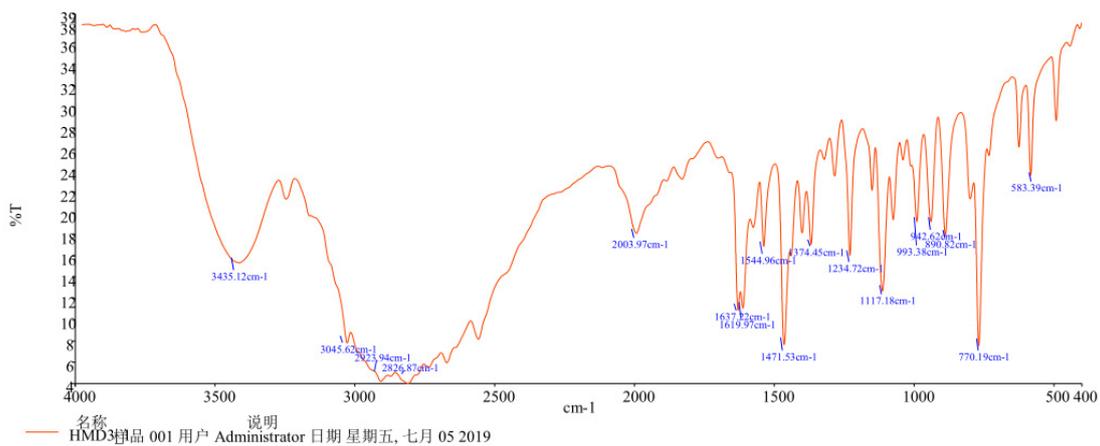


Figure S5. IR spectrum of the Ligand.

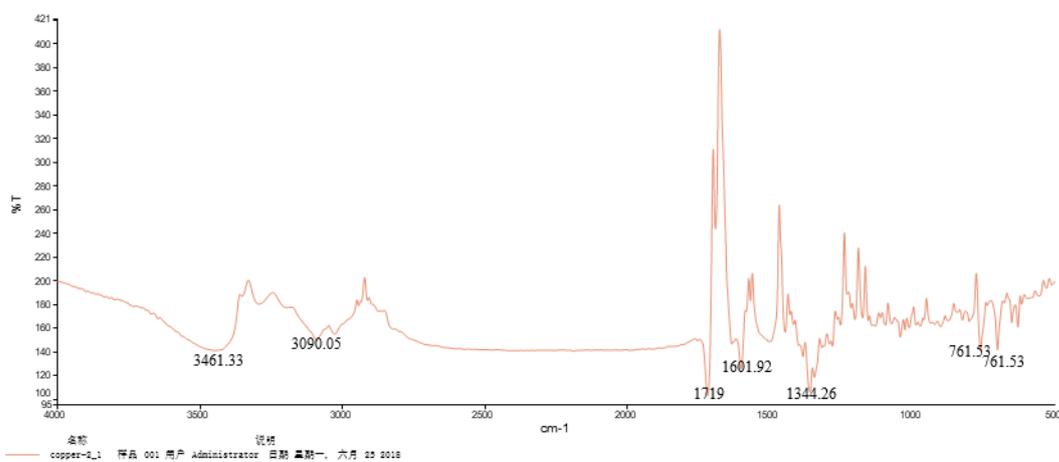


Figure S6. IR spectrum of Cu(II) complex (C1).

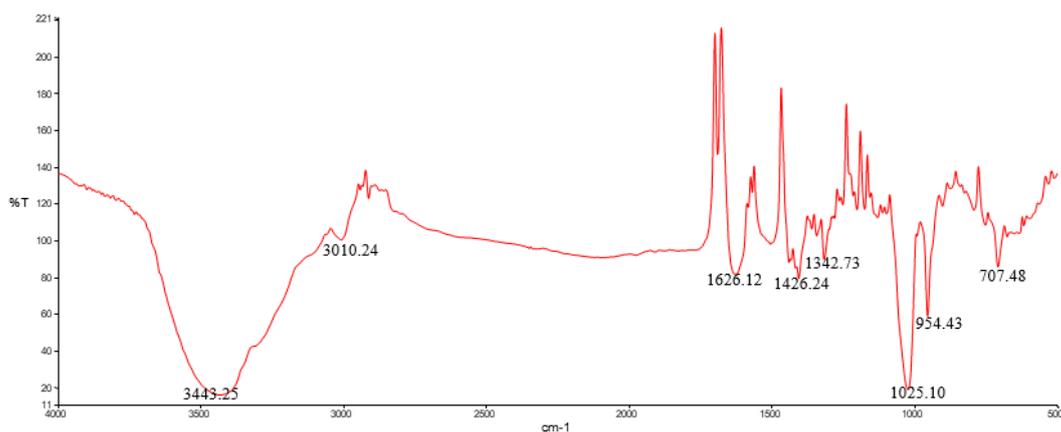


Figure S7. IR spectrum of Cu(II) complex (C2).