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Supplementary Material

Microwave-assisted synthesis of ruthenium(II) complexes with TMSA (trimethylsilylacetylene) as inhibitors against the migration of breast cancer cells

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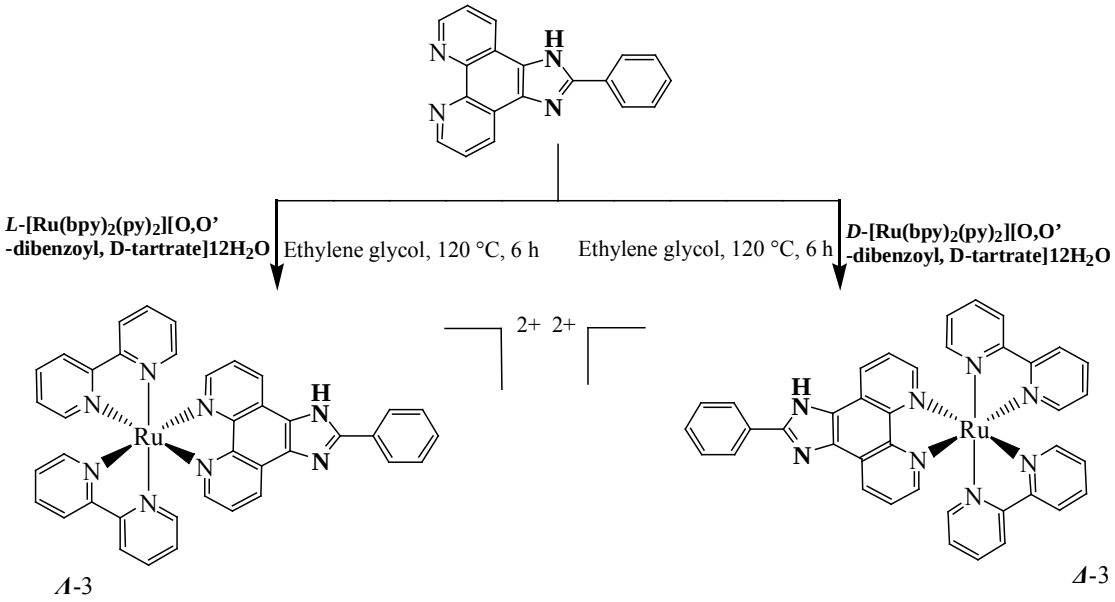


Scheme S1 The synthesis route of chiral ruthenium(II) complexes ***Λ***-3 and ***Δ***-3.

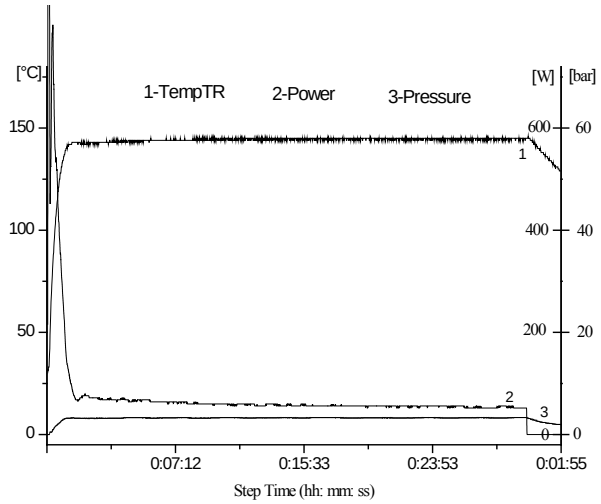


Figure S1 Reaction profile of L0627 in acetonitrile at 140 °C for 30 min. Temperature (1), Power (2), and Pressure (3).

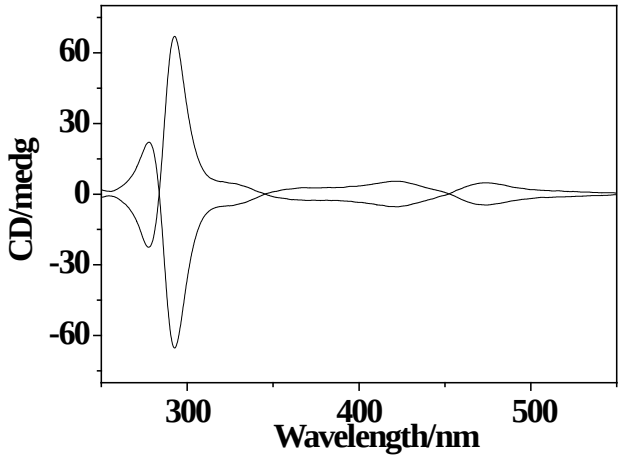


Figure S2 The CD spectra of ruthenium(II) complexes ***Λ***-1 and ***Δ***-1.

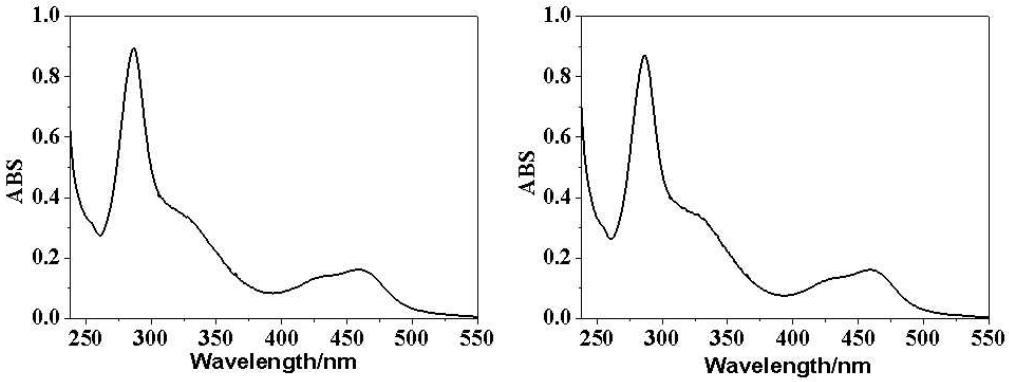
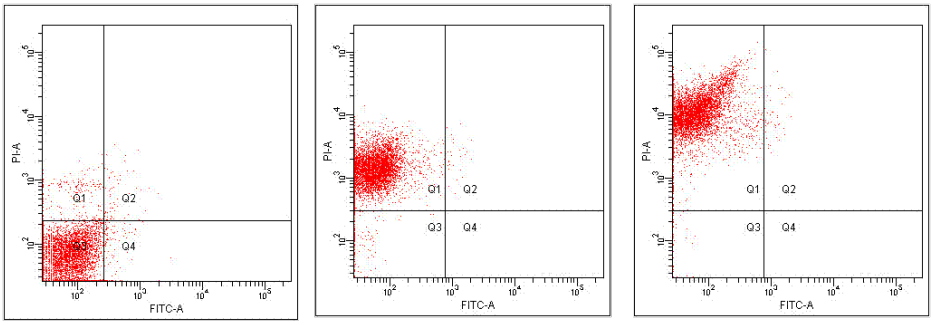


Figure S3 The electronic spectra of ruthenium(II) complexes ***Λ***-1 and ***Δ***-1.

Table S1 The inhibitory activity (IC_{50}/μM) of chiral ruthenium(II) complexes and cisplatin against selected cell lines.

Comp.	Inhibitory Activity /IC_{50}			
	MDA-MB-231	**EC-1**	**HepG2**	**HaCat**
***Λ*-1**	32.1±1.2	88.0±2.6	135.7±1.0	120.7±1.0
***Δ*-1**	36.9±1.6	113.4±1.0	113.9±0.8	171.9±0.5
***Λ*-3**	93.1±0.9	578.2±0.4	132.1±0.5	197.2±2.3
***Δ*-3**	276.9±0.9	593.7±1.0	145.7±0.5	213.9±1.8
Cisplatin	36.1±0.8	/	7.23±2.4	7.48±1.3



Control ***Λ*-1** ***Δ*-1**

Figure S4. FCM analysis of the apoptosis of MDA-MB-231 cells treated with complexes ***Λ***-1 and ***Δ***-1 after 24 h. The effect of both complexes (20 μM) on MDA-MB-231 cells apoptosis was determined by flow cytometry. Untreated (control) cells treated for 24 h were harvested, fixed, stained with Annexin V/PI, and assessed for cell apoptosis distribution by flow cytometric analysis.