Supporting Information

Specific Oxide Nanoclusters Enhance Intracellular ROS for Cancer Targeted Therapy

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Figure S1. MTT assay. U87 cells were incubated 24h with Ir ions at different concentrations while the Fe ions concentration is constant 0.1mM/L.



Figure S2. *In situ* biosynthesized NCs in U87 cells. A) TEM image of IrO₂ NCs, B) TEM image of iron oxide NCs.



Figure S3. UV-vis absorption. UV-vis absorption spectrum of the *in situ* biosynthesized IrO₂ and iron oxide NCs

Table S1. Sequence of primers used in this study.

Gene	Primer	Sequence
Bcl-2	F	GTGACTTCCGATCAGGAAGG
	R	CTTCCAGACATTCGGAGACC
Вах	F	AGTAACATGGAGCTGCAGAGG
	R	ATGGTTCTGATCAGTTCCGG
p53	F	TGTCATGGCGACTGTCCAGC
	R	GCTCGACGCTAGGATCTGAC
Cytochrome c	F	GAGCGGGGAGTGTTCGTTGT
	R	GTCTGCCCCTTTCTTCCTTCT
Casp-3	F	CATGGAAGCGAATCAATGGACT
	R	CTGTACCAGACCGAGATGTCA
Casp-9	F	ACTTTCCCAGGTTTTGTTTCCT
	R	GAAATTAAAGCAACCAGGA
GAPDH	F	CCCACTAACATCAAATGGGG
	R	CCTTCCACAATGCAAAGTT



Figure S4. **Effect of biosynthesized IrO₂ and iron oxide NCs on the expression of apoptotic genes**. Graphs with qRT-PCR analysis data for Bcl2, Bax, Caspase-3, Caspase-9 and Cytochrome c. Expression levels were normalized by GAPDH



Figure S5. Biosynthesized IrO₂ and iron oxide NCs can affect cell migration ability of U87 and HepG2 cells. A) Biosynthesized NCs inhibit migration in U87 and HepG2 cells analyzed by Transwell assay. B-C) Quantification of migrated cells.



Figure S6. Effect of as-biosynthesized NCs on cell migration. A) U87 and HepG2 cells after treatment in vitro wound healing assay. Photographs were taken at hour 0, and 48 h, respectively, after the wound was made. B-C) The wound healing assay was expressed as relative wound width.



Figure S7. H&E Staining. The tumor tissue stained by hematoxylin and eosin.



Figure S8. H&E Staining. The vital organs stained by hematoxylin and eosin.