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Supporting Information

Inhibition of proliferation and migration of tumor cells through lipoic

acid-modified oligoethylenimine-mediated p53 gene delivery

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Figure S1. Gel retardation assay of OEI1800 and LA-OEI with plasmid pEGFP-N3 at different mass ratios.



Figure S2. The transfection of HeLa cells with the plasmid pEGFP-N3 using OEI as the carrier at mass ratios of 7.5, 10.0, 15.0, 20.0, 25.0, 30.0, 40.0 and 50.0, respectively (a-h). The scale bar is 200 μm.



Figure S3. *In vitro* transfection efficiency analysis of OEI1800 (A) and LA-OEI (B) at different mass ratios, using the transfection of pGL-3 plasmid as a model. The data were presented as mean value \pm SD of three experiments.



Figure S4. TEM images of OEI/pEGFP-N3 (a) and LA-OEI/pEGFP-N3 (b) nanocomplexes at a mass ratio of 4.0. The scale bar is 500 nm.



Figure S5. Cell viability analysis of the carriers LA-OEI and OEI1800 at different concentrations through MTT method. The data were presented as mean value \pm SD of three experiments (ns, not significant; *p<0.05; **p<0.01).



Figure S6. Induction of anti-proliferative effect by the carriers-mediated p53 transfection in PC-3 cells through MTT method. The data were presented as mean value \pm SD of three experiments (ns, not significant; *p<0.05; **p<0.01).



Figure S7. Live/Dead staining of HeLa cells after p53 transfection mediated by different carriers, in which living and dead cells were stained to green and red, respectively. The scale bar is $100 \mu m$.



Figure S8. Inhibition of colony formation after p53 transfection mediated by different carriers (A) and the quantitative analysis (B): (a) control; (b) OEI1800; (c) LA-OEI; (d) OEI/p53; and (e) LA-OEI/p53. The scale bar is 200 μm.



Figure S9. The activity analysis of caspase-3, -8 and -9 after p53 delivery mediated by different carriers.



Figure S10. Mitochondrial membrane potential analysis of HeLa cells after p53 transfection mediated by different carriers, using JC-1 probe. The scale bar is $100 \mu m$.



Figure S11. Wound healing assay for the anti-migration effect induced by the carriersmediated p53 transfection. The scale bar is 200 μ m, and data were presented as mean value \pm SD of three experiments.

Table S1. Hydrodynamic diameter and zeta potential of LA-OEI/pEGFP-N3 and OEI/pEGFP-N3 nanocomplexes at different mass ratios. Data were presented as mean value \pm SD of three experiments.

Entry	Mass ratio	Hydrodynamic diameter	Polydispersity	Zeta potential
		(nm)	index	(mV)
LA-OEI/pEGFP-N3	1.0	274.3 ± 4.6	0.301	$+12.3 \pm 1.0$
	2.0	221.8 ± 4.3	0.290	$+17.9 \pm 1.2$
	3.0	155.6 ± 1.0	0.244	$+26.8\pm0.5$
	4.0	105.5 ± 0.9	0.152	$+30.5\pm0.3$
	5.0	91.8 ± 2.6	0.144	$+33.7\pm0.7$
OEI/pEGFP-N3	1.0	389.5 ± 8.7	0.294	+6.1 ± 0.2
	2.0	332. 2 ± 5.6	0.266	$+6.5 \pm 0.2$
	3.0	299.2 ± 1.8	0.272	$+18.3\pm0.3$
	4.0	186.8 ± 0.4	0.195	$+25.5\pm0.5$
	5.0	129.6 ± 0.7	0.155	$+28.0\pm0.5$