Electronic Supplementary Material (ESI) for Biomaterials Science. This journal is © The Royal Society of Chemistry 2022

Figures S1-S9 in supporting information

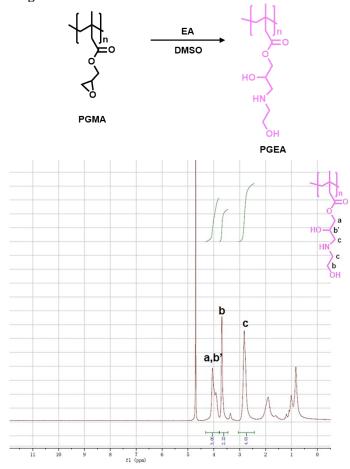


Fig. S1. Synthetic route and typical <sup>1</sup>H NMR (400 MHz) spectra in D<sub>2</sub>O of PGEA polycation.

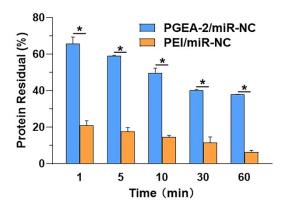


Fig. S2. Protein absorption assay of PGEA-2/miR-NC and PEI/miR-NC complexes treated with excess BSA. (\*p < 0.05)

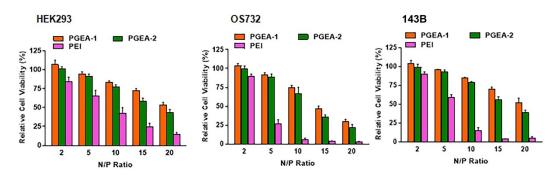
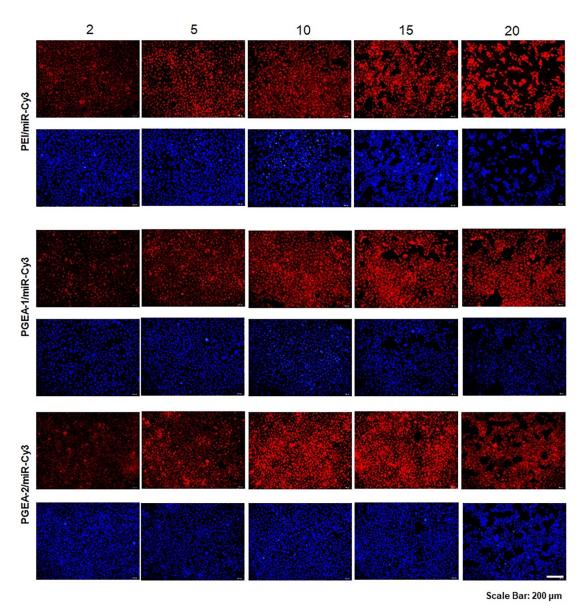
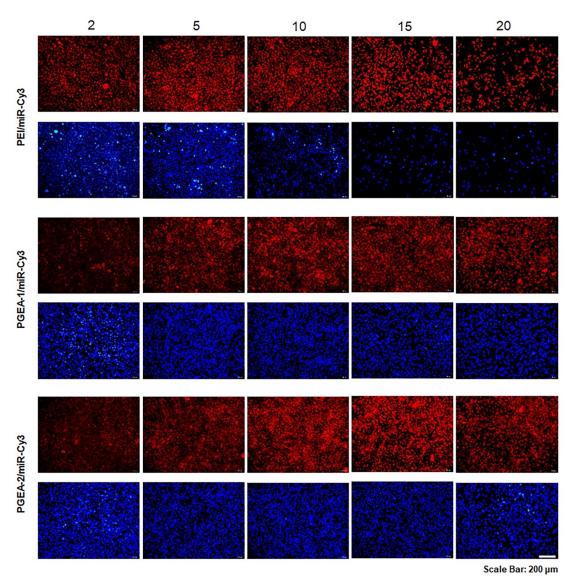


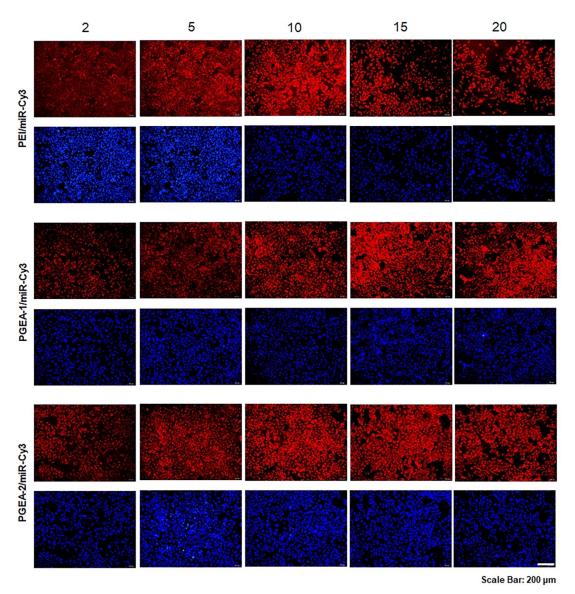
Fig. S3. Cytotoxicities in various cell lines after 24 h incubation with different vectors.



**Fig. S4**. Transfection efficiencies mediated by different vectors in HEK293 cells (Red: miR-Cy3; Blue: nucleus labeled with DAPI).



**Fig. S5**. Transfection efficiencies mediated by different vectors in OS732 cells (Red: miR-Cy3; Blue: nucleus labeled with DAPI).



**Fig. S6**. Transfection efficiencies mediated by different vectors in 143B cells (Red: miR-Cy3; Blue: nucleus labeled with DAPI).

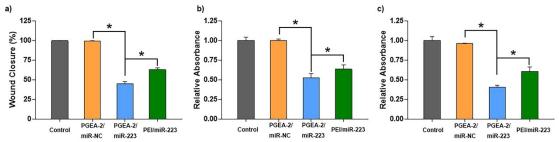


Fig. S7. Statistical analysis of a) wound healing, b) invasion and c) clonogenic assays in different treatment groups. (\*p < 0.05)

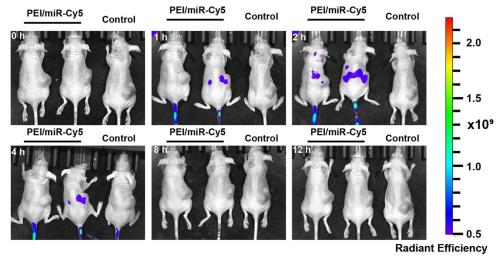
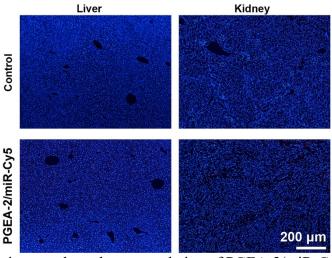


Fig. S8. The accumulation of PEI mediated miRNA delivery in mice with osteosarcoma.



**Fig. S9**. Representative images show the accumulation of PGEA-2/miR-Cy5 system in mice liver and kidney after 12 h (Blue: necleus stained with DAPI; Red: miR-Cy5).