Boosting cancer therapy efficiency via photoinduced radical production†

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Current cancer therapy has been restricted by the hypoxic microenvironment of tumors, especially for strongly oxygen-dependent photodynamic therapy. To defeat tumor hypoxia, an oxygen-irrelevant radical nanogenerator, PI/FBC, is developed by the co-assembly of iodized polymer PI and NIR photosensitizer FBC, and further evaluated as a remote controllable free radical generation platform for enhancing antitumor efficiency. The PI/FBC radical nanogenerator can be excited by NIR light to produce ROS through transfer of energy to oxygen and induce the formation of toxic iodine radicals via electron transfer to PI. Notably, unlike conventional tumor treatments, such a radical nanogenerator is controllable and insensitive to oxygen concentration. Moreover, benefiting from the strong NIR emission of FBC, the distribution of the PI/FBC radical nanogenerator can be monitored without incorporating other imaging agents. This PI/FBC radical nanogenerator treatment will no doubt broaden the family of antitumor strategies by using non-oxygen radicals, which is significant for reference in the development of promising anticancer agents.

Introduction

Hypoxia is a typical microenvironment feature in most solid tumors, resulting from the imbalance between oxygen supply and demand.1–2 A hypoxic microenvironment makes cancer cells resistant to cancer treatments, especially for oxygen-consuming therapies, for instance, photodynamic therapy.3–6 To enhance the therapeutic efficiency, great efforts have been dedicated to overcoming the bottleneck arising from hypoxia,7 such as increasing the oxygen concentration,8–10 reducing the oxygen consumption,11,12 or combining multiple therapeutic methods.13–20 Nano-delivery systems provide a promising potency in overcoming hypoxia of tumors, but face challenges such as poor repeatability and potentially adverse toxicity.21,22 Recently, hypoxia-tolerant photosensitizers have provided a powerful tool for hypoxic tumor treatment.23–26 These photosensitizers could produce toxic reactive oxygen species (ROS) such as superoxide (O2−), hydroxyl (·OH) and hydroperoxyl (HO2) through a photo-induced electron transfer mechanism. However, the produced toxic species are related to oxygen, indicating that the photodynamic therapeutic efficiency of these photosensitizers still relies on oxygen concentration although they require much lower oxygen concentration than traditional photosensitizers.27 To defeat tumor hypoxia, it would be promising to develop an entirely oxygen-independent platform with remote controllable ability.

Oxygen-irrelevant free radicals are also highly reactive species, which have been extensively employed to synthesize compounds and polymers.28–31 Benefiting from their efficient generation even under a hypoxic environment, radical nanogenerators based on free radical initiators have been developed to overcome tumor hypoxia for amplifying antitumor efficiency.32–36 Aside from these radical nanogenerators which focus mainly on thermolabile radical generators, production of free radicals via a photo-redox reaction may be another promising approach for hypoxic tumor treatment. Photo-induced radical reactions with excellent controllability have been an appealing option for catalytic organic synthesis and controllable polymerization.37–40 For cancer therapy, Bu and his co-workers developed a ·Cl nano-generator by decorating a SiO2-coated upconversion nanoparticle (UCNP) with AgCl/AgCl heterodots outside, in which ·Cl generation can be triggered by the short wavelength emission of the UCNP upon NIR illumination.41 In consideration of the energy conversion efficiency, it is a great advantage for efficient oxygen-irrelevant radical formation directly via a NIR photosensitizer upon illumination.

Herein, we develop a biocompatible free radical nanogenerator (PI/FBC) by the co-assembly of iodized block copolymer POEGMA-b-PIEMA (PI) and tetrafluorophenyl bacteriochlorin (FBC) photosensitizer with strong NIR absorbance for hypoxia-irrelevant phototherapy against tumor hypoxia. As shown in Scheme 1, FBC as a NIR photosensitizer can not only produce ROS under NIR illumination, but also induce the production of highly cytotoxic iodine free radicals, endowing the PI/FBC...
nanogenerator with superior remote controllability. Moreover, iodide as the radical initiator was installed on the polymer chain by reversible addition–fragmentation chain transfer (RAFT) polymerization, avoiding unpredictable side effects suffered from the leakage of initiators. Besides, POEGMA as the hydrophilic block can form a hydrophilic shell on the surface of the PI/FBC nanogenerator, effectively improving the biocompatibility of the PI/FBC radical nanogenerator and extending the circulation time. More importantly, the strong NIR absorption of FBC is beneficial to realize deep penetration against skin tissue without severe side effects. Thus, the PI/FBC radical nanogenerator may lead to the new trend of free radical-induced tumor therapy, providing a reference for the development of non-oxygen free radicals for future hypoxia antitumor therapy.

Results and discussion

Synthesis of PI/FBC nanoparticles

NIR absorption photosensitizer FBC was firstly synthesized according to our previous work (Fig. S1, ESI†). Then, iodine-containing monomer 2-iodoethyl methacrylate was prepared via an esterification reaction between methacyryloxy chloride and 2-iodoethanol (Fig. S2, ESI†). Before POEGMA-b-PIEMA was synthesized through RAFT polymerization, a POEGMA macroRAFT agent was prepared by the homopolymerization of OEGMA according to the route as illustrated in Scheme S1 (ESI†). The successful synthesis of POEGMA and POEGMA-b-PIEMA was confirmed by 1H NMR spectroscopy (Fig. S3 and S4, ESI†) and gel permeation chromatography (GPC, Fig. S5, ESI†), respectively. Afterwards, PI/FBC nanoparticles were fabricated by the co-assembly of POEGMA-b-PIEMA and FBC. The transmission electron microscopy (TEM) image shown in Fig. 1a reveals the well-defined spherical morphology of PI/FBC nanoparticles. The hydrodynamic size distribution of PI/FBC nanoparticles was determined by dynamic light scattering (DLS, Fig. 1b). Compared with PI nanoparticles (directly formed by the self-assembly of POEGMA-b-PIEMA block polymers), the size of the PI/FBC nanoparticles did not increase but decreased slightly after FBC was incorporated, which may have resulted from the more concentrated hydrophobic moieties of FBC. The zeta potential value of PI/FBC nanoparticles was below $-10$ mV in comparison to that of PI nanoparticles, which not only suggested the successful encapsulation of FBC, but also showed good colloidal stability as displayed in Fig. 1c. The photophysical properties of PI/FBC nanoparticles were evaluated by utilizing UV-vis and fluorescence spectrophotometers (Fig. 1d). The UV-vis spectra of PI/FBC nanoparticles indicated that FBC was successfully encapsulated in the nanoparticles, in agreement with the DLS and zeta potential results. Besides, owing to the efficient encapsulation of FBC, the strong NIR emission of FBC can also be observed in the fluorescence spectrum of PI/FBC nanoparticles, endowing PI/FBC nanoparticles with excellent imaging performance. Additionally, iodine-free PC/FBC nanoparticles as the control were prepared by encapsulating FBC with a block copolymer, poly(ethylene glycol)-block-poly(lactic acid) (PEG-b-PLA), and the detailed characterization is shown in Fig. S6–S8 (ESI†).

Reactive species production of PI/FBC nanoparticles

PI/FBC nanoparticles were expected to produce both reactive oxygen species and iodine free radicals upon NIR illumination.
And then we determined the extracellular reactive species (RS) production of PI/FBC nanoparticles before in vitro antitumor performance. DPBF as a total ROS probe was employed to measure the ROS production of PI/FBC nanoparticles under NIR irradiation. As shown in Fig. 2b, S9 and S10 (ESI†), PI/FBC and PC/FBC nanoparticles can both cause the absorbance decrease of DPBF with the irradiation time extension, indicating that a significant amount of ROS was generated by the nanoparticles. It is noteworthy that the absorbance decrease of DPBF of PI/FBC nanoparticles was modest compared to that of PC/FBC nanoparticles, which may be induced by the iodine free radical production of PI/FBC nanoparticles. The iodine free radical production competes with the ROS generation to consume excited photosensitizers, resulting in the decrease of ROS formation. To verify the hypothesis, 2,2’-amino-di(2-ethyl-benzothiazoline sulphonic acid-6) ammonium salt (ABTS) as a radical probe was adopted to determine the production of free radicals. We can observe that the UV-vis spectrum of ABTS of PC/FBC nanoparticles changed slightly after NIR illumination (Fig. 2c); however, a significant increase was found in that of PI/FBC nanoparticles (Fig. 2d). To further confirm the production of free radicals, we also used a radical trapping agent, TEMPO, to capture the produced iodine free radical under NIR illumination (Fig. S11, ESI†). The peak at 2.8 ppm increased with irradiation time extension, suggesting the successful formation of iodine radicals.

**Intracellular uptake and photoactivity**

Encouraged by the exciting extracellular consequence, in vitro antitumor performance was evaluated on 4T1 cells. A confocal laser scanning microscope (CLSM) was employed to determine the cellular internalization and cytotoxic species production of nanoparticles. As shown in Fig. 3a, bright red fluorescence can be observed around the cell nucleus (blue fluorescence), indicating that nanoparticles can efficiently enter cancer cells. Subsequently, 2,7-dichlorofluorescein diacetate (DCFH-DA), a fluorescent probe, was used to determine the production of oxidizing toxic species as shown in Fig. 3b. Compared to PI and PC/FBC nanoparticles, strong green fluorescence of DCF generated via oxidation of reactive species to DCFH-DA was
detected after treatment with NIR irradiation in the presence of PI/FBC nanoparticles, indicating that the DCF formation resulted from the toxic species produced by PI/FBC nanoparticles after NIR irradiation. There was almost no green fluorescence in PI nanoparticle treated cells with NIR illumination, which may reveal that PI nanoparticles are biocompatible.

MTT assay was employed to further evaluate the antitumor efficiency under normal and hypoxic conditions (FBC concentration: 0–8 µg mL⁻¹) (Fig. 3c–f). From Fig. 3c and e, it can be observed that there is no significant cytotoxicity against cells for PI, PC/FBC and PI/FBC nanoparticles, indicating that all the nanoparticles have good biocompatibility. It is noteworthy that, with NIR irradiation (750 nm, 50 mW cm⁻², 10 min), PI/FBC nanoparticles could efficiently induce cell death in a concentration-dependent manner, even in a hypoxic environment (Fig. 3d and f). However, the PC/FBC and PI nanoparticle treated cells did not end their days obviously. These results may be associated with the modest or disabled production of toxic species of PC/FBC or PI nanoparticles, respectively, which is in line with the foregoing consequence of intracellular DCF fluorescence detected by CLSM. In addition, we also determined the cytotoxicity of PC/FBC nanoparticles at a higher FBC concentration (0–32 µg mL⁻¹) under an elevated laser power density (100 mW cm⁻², 15 min) (Fig. S12, ESI†). The results revealed that the cell survival decreased with the increase of PC/FBC concentration, which further confirmed that the high cell survival of PC/FBC in the previous experiment resulted from the limited ROS generation.

**In vivo imaging and biodistribution, photodynamic therapy and bio-safety**

Based on the remarkable in vitro photo-induced toxicity against cancer cells, an in vivo performance study was carried out on nude mice bearing tumors. 4T1 tumor-bearing mice were obtained by subcutaneously injecting 200 µL of 5 × 10⁶ tumor cells. For in vivo distribution monitoring via fluorescence imaging, PI/FBC nanoparticles were injected into mice via the tail vein after the xenograft tumor volume reached 100 mm³. The fluorescence imaging was monitored in real time via detecting the emission at 750 nm after FBC was excited with 510 nm light. As shown in Fig. 4a and S13 (ESI†), the fluorescence signal at the tumor gradually increased with time extension, suggesting that PI/FBC could be efficiently enriched in tumors by the enhanced permeability and retention (EPR) effect of tumors. The biodistribution results also distinctly demonstrated that PI/FBC nanoparticles were primarily accumulated in the tumor site 24 h post-injection.

To further verify the antitumor efficiency of PI/FBC nanoparticles, mice bearing 4T1 tumors were randomly divided into seven groups (control, PI, PC/FBC, PI/FBC, PI + L, PC/FBC + L and PI/FBC + L) and separately administered with the corresponding nanoparticles through the tail vein when the tumor volume reached 100 mm³. Tumors in PI + L, PC/FBC + L and PI/FBC + L groups were irradiated with a 750 nm laser for 15 min at a power density of 200 mW cm⁻². Tumor volumes and body weights were recorded to evaluate the tumor suppression effect and biocompatibility of nanoparticles, respectively (Fig. 4b and c). We found that PC/FBC and PI/FBC nanoparticles could significantly inhibit the tumor growth after illumination with a 750 nm laser. It is noteworthy that PI/FBC exhibited a superior inhibition efficiency than PC/FBC, which could be attributed to the oxygen-irrelevant iodine free radicals generated by PI/FBC nanoparticles under illumination. However, PI nanoparticles did not show any antitumor effect, indicating that the oxygen-irrelevant free radical production was triggered by FBC under 750 nm laser irradiation. Without laser illumination, PI, PC/FBC and PI/FBC nanoparticles all showed no obvious difference compared to the control, revealing that they had a good biocompatibility. No significant weight loss was observed in Fig. 4c, suggesting minimal side effects of the nanoparticles. On 15th day, the mice were sacrificed to harvest tumors and primary organs (heart, liver, spleen, lungs and kidneys) for further evaluation of antitumor efficiency. Isolated tumors were weighed and photographed as depicted in Fig. 4d and e. PI/FBC + L treated tumors had the lightest weight and smallest volume, indicating that PI/FBC nanoparticles could efficiently inhibit the tumor growth after illumination with a 750 nm laser. H&E
staining on tissue sections (Fig. 4f and S14, ESI†) also revealed that PI/FBC nanoparticles are highly biocompatible with major organs and could cause significant cell necrosis after being triggered by 750 nm laser irradiation.

Conclusions

In conclusion, a biocompatible hypoxia-tolerant nano-system, PI/FBC, was fabricated by co-assembling an iodized polymer with a NIR photosensitiser, FBC. PI/FBC nanoparticles could efficiently accumulate in tumor sites through the EPR effect, which can be monitored in real time via NIR fluorescence imaging. Upon being excited with an NIR laser, PI/FBC nanoparticles could not only produce cytotoxic ROS by transferring energy to oxygen, but also cause highly toxic iodine free radical production through an oxygen-irrelevant mechanism. Therefore, PI/FBC nanoparticles could facilitate a significantly enhanced antitumor efficiency by conquering the hypoxic tumor environment in a controllable manner. This PI/FBC radical nanogenerator treatment will no doubt broaden the family of antitumor strategies by using nonoxygen free radicals, which is significant for reference in the development of prospective anticancer agents.

Data availability

The electronic supplementary information include experimental detail, NMR data, spectra data and histological data.

Author contributions

Z. Liu conceived the project and designed the experiments. Z. Liu and M. Wu performed the sample preparation, characterization and biological experiment. Z. Liu and M. Wu wrote the paper and contributed equally. All authors discussed the results and commented on the manuscript.

Ethical statement

All animal studies were conducted on male Balb/c nude mice (four to five weeks) in compliance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals approved by the Animal Care and Use Committee of East China University of Science and Technology.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the National Natural Science Foundation of China (No. 22075079 and 21875063) and the Science and Technology Commission of Shanghai Municipality for the Shanghai International Cooperation Program (19440710600).

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