

Precision design of nanomedicines to target tumor microenvironment and overcome low chemosensitivity for pancreatic cancer treatment



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报告摘要:

In spite of the promising therapeutic potential exhibited by the numerous antitumor nanomaterials, it remains a major challenge for particulate-based therapeutics to effectively transport into solid tumors, especially in stromal enriched tumors. The recent progress on understanding the critical roles and the underlying mechanisms of tumor microenvironment on tumor rapid growth and metastasis has made targeting the tumor microenvironment becomes a feasible strategy to improve the effectiveness of nanomaterials-based cancer diagnosis and therapy. Expression of stromal fibroblasts-specific enzymes and pathological changes of tumor vasculature are the most remarkable hallmarks of tumor microenvironment in almost all types of tumors and can be considered as good specific triggers or targets for design of broad-spectrum and local-environment responsive functional nanomaterial-based platform. Here we report novel biomimetic nanostructures based on oligopeptide selfassemblies that could quickly response and regulate the key components of tumor microenvironment. With such a strategy, the optimal formulations were developed, showing the high activation efficacy and antitumor efficacy. The tailor-made self-assembled biomolecule nanomaterials have the potential to be used in early and late stages of solid tumors, especially for stromal enriched solid tumors, which is expected to be of crucial importance for clinical tumor therapeutics. Although additional research is urgent needed to develop robust methods for targeting and regulating nanomaterials to tumor sites and the supporting environment, the applications of tumor microenvironment-based nanotechnology for safer and more effective antitumor nanomedicine have so far been proven to be successful and will eventually revolutionize the current landscape of cancer therapy.

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