Peptidic supramolecular assemblies enhance chemotherapy for colorectal cancer
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Colorectal cancer (CRC) has emerged as a frequently diagnosed disease. Advanced CRCs are highly heterogenous and often evade treatments, building drug resistance and reoccurring.

Fusobacterium nucleatum (F. nucleatum) is among the most prevalent bacterial species in colorectal cancer tissues. The increase of F. nucleatum not only promotes and establishes CRC, but also induces metastasis and drug resistance. Targeting such cancer-associated microorganisms might provide a new method to abolish drug resistance and ameliorate poor chemotherapy outcomes.

In a study published in Science Advances, Gao Yuan's group from the National Center for Nanoscience and Technology (NCNST) of the Chinese Academy of Sciences (CAS) provided an antibiotic-free strategy to inhibit specific bacterial strains for enhanced CRC treatment by designing a nitroreductase (NTR)-instructed supramolecular self-assembly system that can inhibit F. nucleatum and sensitize CRC for chemotherapy.

The researchers synthesized an atypical NTR-responsive motif by introducing additional hydrophilic carboxyl group to the nitrobenzene moiety. The conjugation of the NTR-responsive motif to a suitable peptide sequence yielded an assembly precursor. The precursors underwent NTR catalyzed transformation and formed ordered nanofibers both in vitro and in living systems.

These nanofibers rather than the individual molecules exhibited antibacterial activity against F. nucleatum. The local introduction of NTR-instructed assemblies significantly alleviated bacteria-induced oxaliplatin (OXA) resistance effect and eventually efficiently inhibited tumor growth.

Then, the researchers confirmed that the inhibition of F. nucleatum contributed to the enhanced chemotherapy outcomes. They declared that supramolecular self-assembly could serve as an abiotic strategy to interrupt bacteria-mammalian cell interactions and enhance CRC therapy.

"Such a strategy should be applicable to improve other cancer therapies if the tumors bear the risk of microbial infections," said Gao. Taking E. coli associated drug-resistant pancreatic cancer as an example, in such a model, E. coli will metabolize the anticancer drug Gemcitabine into its inactive form, resulting in the failure of chemotherapy against pancreatic cancer.

More information: Jiali Chen et al, Nitroreductase-instructed supramolecular assemblies for microbiome regulation to enhance colorectal cancer treatments, Science Advances (2022). DOI: 10.1126/sciadv.add2789