

Singapore's NUS Medicine researchers engineer bacteria for targeted chemotherapy delivery

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A novel method for delivering chemotherapy drugs directly to tumour sites by utilising the natural interactions between bacteria and cancer cells.



Traditional chemotherapy often poses significant challenges, including severe side effects, damage to healthy tissue and limited efficacy. Researchers at the Yong Loo Lin School of Medicine, National University of Singapore (NUS Medicine) have developed a revolutionary approach to cancer treatment — one that is a more targeted, effective, and less toxic alternative to traditional chemotherapy. This new approach not only improves treatment effectiveness but also significantly reduces the required drug dosage for cancer treatment.

Led by Associate Professor Matthew Chang, researchers at the NUS Synthetic Biology for Clinical and Technological Innovation (SynCTI) and the Synthetic Biology Translational Programme (Syn Bio TRP) at NUS Medicine identified a new method of drug delivery that offers hope for the development of a new clinical treatment for cancer patients.

Prodrugs are inactive molecules that transform into active drugs within the body, particularly in tumour environments, by leveraging unique tumour conditions, such as low oxygen or high acidity, to activate the drug precisely at the cancer site, minimising damage to healthy tissues. However, current prodrug strategies exhibit limited target specificity and frequently depend on macromolecular carriers, which complicates both drug distribution and clearance.

To overcome these limitations, NUS Medicine researchers developed a prodrug delivery method that utilises a commensal *Lactobacillus* strain that binds specifically to cancer cells via a surface molecule called heparan sulfate. These engineered bacteria carry a prodrug that converts to the chemotherapy drug SN-38 at the tumour site.

In preclinical models of nasopharyngeal cancer, the engineered bacteria localised specifically in the tumour and released the chemotherapy drug directly at the cancer site, reducing tumour growth by 67% and increasing the effectiveness of the chemotherapy drug by 54%.

One of the most promising aspects of this research is the potential broader applications across various types of cancer

therapy, as the *Lactobacillus* strain identified by the researchers binds specifically to cancer cells. Lead Researcher Dr Shen Haosheng, Research Fellow at SynCTI said: "By harnessing the affinity between bacteria and cancer cells, we aim to revolutionise chemotherapy delivery. We are evaluating the binding affinity of several microbial strains to multiple cancer cell lines with the aim of developing a versatile delivery system using microbial strains to target chemotherapy drugs to various mucosal cancers, such as colorectal, bladder, stomach, oral, lung, and nasal cancer."