'Drug factory' implants eliminate ovarian, colorectal cancer in mice
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Rice University bioengineer Omid Veiseh with a vial of bead-like implants his lab invented to serve as anti-cancer drug factories. Veiseh and colleagues showed the implants could eradicate ovarian and colorectal cancer in mice in as little as six days. Credit: "Jeff Fitlow/Rice University"

Rice University bioengineers have shown they can eradicate advanced-stage ovarian and colorectal cancer in mice in as little as six days with a treatment that could be ready for human clinical trials later this year.

The researchers used implantable "drug factories" the size of a pinhead to deliver continuous, high doses of interleukin-2, a natural compound that activates white blood cells to fight cancer. The drug-producing beads can be implanted with minimally invasive surgery. Each contains cells engineered to produce interleukin-2 that are encased in a protective shell.

The treatment and animal test results are described online today in a Science Advances study co-authored by Omid Veiseh, Amanda Nash and colleagues from Rice, the University of Texas MD Anderson Cancer Center, the University of Virginia and others.

Veiseh, an assistant professor of bioengineering whose lab produced the treatment, said human clinical trials could begin as soon as this fall because one of his team's key design criteria was helping cancer patients as quickly as possible. The team chose only components that had previously proven safe for use in humans, and it has demonstrated the safety of the new treatment in multiple tests.

"We just administer once, but the drug factories keep making the dose every day, where it's needed until the cancer is eliminated," Veiseh said. "Once we determined the correct dose—how many factories we needed—we were able to eradicate tumors in 100% of animals with ovarian cancer and in seven of eight animals with colorectal cancer."

Rice University bioengineering graduate student Amanda Nash is creating cell-encapsulation technologies to treat cancer. Nash and colleagues created the technology with an eye toward rapid approval for human clinical trials, using only components that have previously proven safe for use in humans. Credit: Jeff Fitlow/Rice University
In the newly published study, researchers placed drug-producing beads beside tumors and within the peritoneum, a sac-like lining that supports intestines, ovaries and other abdominal organs. Placement within this cavity concentrated interleukin-2 within tumors and limited exposure elsewhere.

"A major challenge in the field of immunotherapy is to increase tumor inflammation and anti-tumor immunity while avoiding systemic side effects of cytokines and other pro-inflammatory drugs," said study co-author Dr. Amir Jazaeri, professor of gynecologic oncology and reproductive medicine at MD Anderson. "In this study, we demonstrated that the 'drug factories' allow regulatable local administration of interleukin-2 and eradication of tumor in several mouse models, which is very exciting. This provides a strong rationale for clinical testing."

Interleukin-2 is a cytokine, a protein the immune system uses to recognize and fight disease. It is an FDA-approved cancer treatment, but Nash, a graduate student in Veiseh's group and the study's lead author, said the drug factories provoke a stronger immune response than existing interleukin-2 treatment regimens because the beads deliver higher concentrations of the protein directly to tumors.

"If you gave the same concentration of the protein through an IV pump, it would be extremely toxic," Nash said. "With the drug factories, the concentration we see elsewhere in the body, away from the tumor site, is actually lower than what patients have to tolerate with IV treatments. The high concentration is only at the tumor site."

Nash said the same general approach used in the study could be applied to treat cancers of the pancreas, liver, lungs and other organs. The drug factories could be placed next to tumors and within the linings that surround those organs and most others, she said. And if a different cytokine is needed to target a specific form of cancer, the beads can be loaded with engineered cells that make that immunotherapeutic compound.

The bead's outer shell shields its cytokine-producing cells from immune attacks. The shells are made of materials the immune system recognizes as foreign objects but not as immediate threats, and Veiseh's lab leveraged that in its design.

"We found foreign body reactions safely and robustly turned off the flow of cytokine from the capsules within 30 days," he said. "We also showed we could safely administer a second course of treatment should it become necessary in the clinic."

Avenge Bio, a Massachusetts-based startup co-founded by Veiseh, has licensed the cytokine-factory technology from Rice. Additional co-authors
include Maria Jarvis, Samira Aghlara-Fotovat, Sudip Mukherjee, Andrea Hernandez, Andrew Hecht, Yufei Cui, Shirin Nouraein, Jared Lee, David Zhang and Oleg Igoshin of Rice; Peter Rios, Sofia Ghani, Ira Joshi and Douglas Isa of CellTrans Inc.; Chunyu Xu and Weiyi Peng of the University of Houston; Rahul Sheth of MD Anderson; and José Oberholzer of both CellTrans Inc. and the University of Virginia.


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