

New monoclonal antibody developed that can target proteins inside cancer cells

March 13 2013

Researchers have discovered a unique monoclonal antibody that can effectively reach inside a cancer cell, a key goal for these important anticancer agents, since most proteins that cause cancer or are associated with cancer are buried inside cancer cells. Scientists from Memorial Sloan-Kettering Cancer Center and Eureka Therapeutics have collaborated to create the new human monoclonal antibody, which targets a protein associated with many types of cancer and is of great interest to cancer researchers.

Unlike other human therapeutic [monoclonal antibodies](#), which can target only proteins that remain on the outside of [cancer cells](#), the new monoclonal antibody, called ESK1, targets a protein that resides on the inside of the cell.

ESK1 is directed at a protein called WT1, which is overexpressed in a range of leukemias and other cancers including myeloma and breast, ovarian, and [colorectal cancers](#). WT1 is a high priority target for [cancer drugs](#) because it is an oncogenic protein, meaning that it supports the formation of [cancer](#). In addition, it is found in few healthy cells, so there are less likely to be side effects from drugs that target it.

"This is a new approach for attacking WT1, an important cancer target, with an antibody therapy. This is something that was previously not possible," said David A. Scheinberg, MD, PhD, Chair of the Sloan-Kettering Institute's [Molecular Pharmacology](#) and Chemistry Program and an inventor of the antibody. "There has not been a way to make

small [molecule drugs](#) that can inhibit WT1 function. Our research shows that you can use a monoclonal antibody to recognize a cancer-associated protein inside a cell, and it will destroy the cell."

The first studies of the antibody are showing promise in preclinical research as a treatment for leukemia as reported March 13, 2013, in *Science Translational Medicine*.

"ESK1 represents a paradigm change for the field of [human monoclonal antibody](#) therapeutics," said Cheng Liu, PhD, President and Chief Executive Officer of Eureka Therapeutics. "This research suggests that human [antibody therapy](#) is no longer limited to targeting proteins present outside cancer cells, but can now target proteins within the cancer cell itself."

ESK1 was engineered to mimic the functions of a T cell receptor, a key component of the immune system. T cells have a receptor system that is designed to recognize proteins that are inside the cell. As proteins inside the cell get broken down as part of regular cellular processes, molecules known as HLA molecules carry fragments of those proteins—known as peptides—to the surface. When T cells recognize certain peptides as abnormal, the T cell kills the diseased cell.

In the current study, the investigators showed that ESK1 alone was able to recognize WT1 peptides and kill cancer cells in the test tube and also in mouse models for two different types of human leukemia. "We were surprised that the antibody worked so well on its own," said Dr. Scheinberg, senior author of the paper. "We had originally expected that we might need to use the antibody as a carrier to deliver a drug or a radioactive therapy to kill the cancer cells, but this was not necessary."

Additional studies must be done in the laboratory before ESK1 is ready to be tested in patients. But the monoclonal antibody was engineered to

be fully human, which should speed the time it takes to move the drug into the clinic. Researchers expect that the first clinical trials, for leukemia, could begin in about a year.

The antibody was developed under a collaborative effort between Memorial Sloan-Kettering and Eureka, which have jointly filed for patent protection.

Provided by Memorial Sloan-Kettering Cancer Center

Citation: New monoclonal antibody developed that can target proteins inside cancer cells (2013, March 13) retrieved 19 September 2024 from <https://medicalxpress.com/news/2013-03-monoclonal-antibody-proteins-cancer-cells.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.