

## Bioengineering and the Fight Against Cancer **Yan R Kizaki\***

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### Opinion

Cancer is one of the top causes of death in the United States and throughout the world. According to the National Cancer Institute, experts anticipated that almost 1.7 million people in the United States would be diagnosed with cancer in 2016. Approximately 14 million persons were diagnosed with cancer worldwide in 2012. Cancer is a disease caused by the growth and multiplication of damaged cells in the body. There are many different types of cancer that can occur in different places of the body. The prostate, breasts, lungs, and colon are some of the most prevalent places where cancer can occur.

Radiotherapy, chemotherapy, and surgery are among the most prevalent cancer treatments, according to the National Cancer Institute. A lot of patients will get a mix of all of these treatments. While these treatments are beneficial for many individuals, scientists are always looking for novel ways to treat and cure advanced and aggressive cancers.

Immunotherapy is a relatively new type of treatment that has shown a lot of promise in recent years. Immunotherapy is a method in which cancer is attacked using a patient's own immune system.

Before delving into what immunotherapy is, it's vital to take a closer look at the immune system, the body's first line of defence against disease. The immune system is an umbrella word for a group of cells and other substances that attack and kill invading pathogens, safeguarding a person from illness. T-cells are special cells that help fight illness as part of the immune system. T-cells are divided into two types, those that detect and warn of sickness, and those that target and eradicate the problem.

When cells become infected with a sickness such as a common virus—they use specific proteins known as Major Histocompatibility Complex proteins to signal to the immune system that they are vulnerable. T-cells are frequently good in eliminating sickness, but they are often ineffective at recognising cancer cells. This is due to cancer cells' ability to utilise a variety of tactics to avoid being detected by T-cell receptors. As a result, T-cells frequently pass cancer cells unnoticed, allowing the malignancy to thrive and spread unabated.

The goal of immunotherapy is to find a mechanism for T-cells to recognise cancer cells and then destroy them. Immunotherapy as a cancer-fighting technique has shown tremendous promise, despite the fact that it is still in its infancy. A handful of clinical trials have resulted in patients being successfully treated for cancers that were previously deemed deadly.

Editorial office, British Biomedical Bulletin,  
United Kingdom

\*Corresponding author: Yan R Kizaki

✉ mikakos687@gmail.com

Editorial office, British Biomedical Bulletin,  
United Kingdom.

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There are currently two major techniques for researchers to utilise immunotherapy: medications named checkpoint inhibitors by the BBC, and genetic manipulation of a patient's T-cells in a lab. It's critical to examine both approaches more closely.

Checkpoint inhibitors are a type of medicine that helps to prevent cancer cells from being able to hide from the immune system. As a result, the T-cells are able to begin battling the tumour. Some doctors are now using checkpoint inhibitor medications to treat qualified patients with specific types of cancer after the US Food and Drug Administration gave them the go-ahead. Checkpoint inhibitors are most commonly used to treat malignancies of the lung, bladder, and kidney. They've been shown to be useful in the treatment of certain types of lymphoma.

While this method is still in its infancy and has only been tested in a few clinical trials, the findings have been generally positive. T-cell alteration, occurs when a patient's own T-cells are transformed in a lab to help them fight cancer more successfully. The cells are reinserted into the patient after the changes have been performed, and the cancer is targeted. Due to multiple clinical trials demonstrating its efficacy in successfully treating specific forms of aggressive blood cancer, such as leukaemia, this technique has sparked a lot of interest and enthusiasm among researchers.

T-cell modification treatment was utilised on a small number of terminally ill leukaemia patients who had exhausted all other therapeutic options. "Essentially, what this approach does is it genetically reprograms the T-cell to seek out, recognise, and eliminate the patient's tumour cells."