The War on Cancer: Targeting the Root of Cancer Cells

Heng Chernov*

Department of Oncology, Wayne State University School of Medicine, MI 48201, USA

Introduction

One of the key factors that makes cancer so challenging is its inherent heterogeneity. Each cancer type, and even individual tumors, exhibit a unique set of genetic mutations, cellular behaviours, and responses to treatments. This complexity is partly responsible for the difficulty in identifying universally effective therapies. However, modern research has shifted focus to understanding cancer at a deeper, molecular level. Rather than only targeting the tumor itself, scientists are beginning to probe the roots of cancer its very origins. By investigating the genetic and environmental factors that contribute to the initiation and development of cancer, we can begin to develop targeted therapies that attack cancer cells at their most fundamental level, potentially providing more effective and less toxic treatment options.

Description

Cancer arises from a series of genetic mutations that disrupt the normal functioning of cells. These mutations often affect key genes that regulate the cell cycle, DNA repair mechanisms, and cell death pathways. Mutations in oncogenes, which drive cell proliferation, and tumor suppressor genes, which inhibit cell growth, can lead to uncontrolled cell division. Over time, these rogue cells accumulate additional mutations that enable them to evade immune detection, resist treatment, and metastasize to other parts of the body. This process of tumorigenesis is driven by a combination of genetic predisposition and environmental factors, such as exposure to carcinogens, lifestyle choices, and infections [1,2].

The traditional approach to cancer treatment has been to target the tumor itself, either through surgery, radiation, or chemotherapy. These methods aim to destroy the cancer cells directly but often fail to eliminate the root causes of the disease. Surgery can remove visible tumors, but it cannot address the microscopic cancer cells that may have already spread to other parts of the body. Radiation and chemotherapy, while effective at shrinking tumors, also harm healthy cells, leading to side effects that can be debilitating and life-threatening. Moreover, cancer cells often develop resistance to these treatments, rendering them less effective over time. As we continue the fight against cancer, it is important to remember that cancer is not a single disease but a collection of related diseases. There are more than 100 different types of cancer, each with its own set of challenges and treatment considerations [3].

In recent years, there has been a growing emphasis on targeted therapies, which aim to pinpoint specific molecular targets within cancer cells. These therapies can block the signalling pathways that cancer cells rely on for survival and growth. For example, tyrosine kinase inhibitors, which target enzymes involved in cancer cell signalling, have shown promise in treating certain types of cancer, such as chronic myelogenous leukemia and non-small cell lung cancer. Another class of targeted therapies involves monoclonal antibodies, which are designed to bind to specific proteins on the surface of cancer cells, either marking them for destruction by the immune system or blocking their

*Address for Correspondence: Heng Chernov, Department of Oncology, Wayne State University School of Medicine, MI 48201, USA; E-mail: hengchernov@gmail. com

Copyright: © 2025 Chernov H. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Received: 01 January, 2025, Manuscript No. jmhmp-25-162185; Editor Assigned: 04 January, 2025, PreQC No. P-162185; Reviewed: 15 January, 2025, QC No. Q-162185; Revised: 21 January, 2025, Manuscript No. R-162185; Published: 28 January, 2025, DOI: 10.37421/2684-494X.2025.10.268

ability to grow and divide. Vaccines, such as the Human Papillomavirus (HPV) vaccine, which protects against certain types of cervical and other cancers, offer hope for reducing cancer risk in the future.

However, targeting cancer cells at the molecular level is not without its challenges. One of the biggest hurdles is the fact that cancer cells are often genetically unstable, meaning that they accumulate mutations at a rapid rate. This genetic instability can create a moving target for treatment, as the molecular makeup of a tumor can change over time. Moreover, some cancer cells may exist in a dormant state, evading treatment while lying hidden within the body. These dormant cells are often responsible for cancer recurrence after the initial treatment has appeared successful. To truly eradicate cancer, it may be necessary to target these dormant cells and prevent their reawakening. Public health campaigns aimed at reducing smoking, promoting physical activity, and encouraging regular screenings for cancers such as breast, cervical, and colorectal cancer have proven effective in reducing cancer incidence [4].

One promising approach in cancer therapy involves reprogramming the tumor microenvironment to make it less conducive to cancer growth. This can involve inhibiting the formation of new blood vessels, a process known as angiogenesis, or reactivating the immune system to recognize and attack cancer cells. Immunotherapy, which uses the body's immune system to target cancer cells, has been one of the most exciting advancements in cancer treatment in recent years. Checkpoint inhibitors, which block the signals that prevent immune cells from attacking cancer, have shown remarkable success in cancers such as melanoma, lung cancer, and bladder cancer. Despite the advances in molecular therapies, prevention remains a critical aspect of the war on cancer. Many cancers are caused by environmental factors, such as tobacco use, exposure to carcinogens, and poor dietary habits [5].

Another innovative approach is the use of gene therapy to directly modify cancer cells or the surrounding tissues. By introducing new genetic material into cancer cells, scientists can potentially correct the mutations that drive tumorigenesis or reprogram the cells to behave in a more normal, controlled manner. Gene editing tools, such as CRISPR-Cas9, have opened up new possibilities for precisely modifying the genetic code of cancer cells. While still in its early stages, gene therapy holds immense promise for treating cancers that have not responded to traditional therapies.

Conclusion

The war on cancer is far from over, but the progress made in recent years offers hope for a future where cancer is no longer a death sentence. By targeting the root causes of cancer, whether through genetic, molecular, or environmental interventions, we are taking significant steps toward understanding and defeating this complex and elusive disease. While the road ahead may be long, the advancements in cancer research and treatment provide a clear path forward, one that promises a future where cancer is no longer the devastating force it once was. This diversity makes it difficult to find a one-size-fits-all solution, but it also underscores the importance of personalized medicine. By tailoring treatments to the individual patient's genetic makeup and the specific characteristics of their tumor, doctors can improve the likelihood of success and minimize side effects.

Acknowledgement

None.

Conflict of Interest

None.

References

- 1. Heng, Henry HQ. "Cancer genome sequencing: The challenges ahead." *Bioessays* 29 (2007): 783-794.
- Heng, Henry HQ. "The genome-centric concept: Resynthesis of evolutionary theory." *Bioessays* 31 (2009): 512-525.

- McClellan, Jon and Mary-Claire King. "Genetic heterogeneity in human disease." Cell 141 (2010): 210-217.
- Iourov, Ivan Y., Svetlana G. Vorsanova, Yuri B. Yurov and Sergei I. Kutsev. "Ontogenetic and pathogenetic views on somatic chromosomal mosaicism." *Genes* 10 (2019): 379.
- 5. Nurse, Paul. "Biology must generate ideas as well as data." *Nature* 597 (2021): 305.

How to cite this article: Chernov, Heng. "The War on Cancer: Targeting the Root of Cancer Cells." *J Mol Hist Med Phys* 10 (2025): 268.