

# The role of t cells in adaptive immunity: Guardians of immune memory.

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## Introduction

The human immune system is a remarkable and intricate defense network that stands as the body's sentinel against pathogens and diseases. At the heart of this formidable system are the T cells, essential soldiers in the armory of adaptive immunity [1]. These cells are responsible for orchestrating and fine-tuning the immune response, effectively serving as the guardians of immune memory. In this article, we delve into the pivotal role of T cells in adaptive immunity, exploring their various subsets, functions, and the profound impact they have on our health and well-being [2].

## The two faces of adaptive immunity

The immune system is broadly categorized into two branches: innate and adaptive immunity. While innate immunity provides a rapid, nonspecific first-line defense, adaptive immunity is a more sophisticated and specific response that develops over time. At the heart of adaptive immunity are the T cells, lymphocytes that orchestrate a highly tailored and targeted response against invading pathogens [3].

T cells are a diverse group of immune cells with a vast repertoire of receptors that can recognize specific antigens. This diversity allows T cells to respond to a wide range of pathogens, from viruses and bacteria to cancer cells. T cells are typically classified into two main groups: CD4<sup>+</sup> T cells and CD8<sup>+</sup> T cells. CD4<sup>+</sup> T cells, also known as helper T cells, play a central role in orchestrating the immune response. They assist in activating other immune cells, including B cells, which produce antibodies, and cytotoxic CD8<sup>+</sup> T cells [4].

CD8<sup>+</sup> T cells, on the other hand, are cytotoxic T cells responsible for directly attacking infected cells. They recognize and destroy cells that have been compromised by intracellular pathogens, such as viruses [5].

One of the remarkable functions of T cells is their ability to recognize antigens presented to them by antigen-presenting cells (APCs), such as dendritic cells. APCs process and display fragments of pathogen-derived antigens on their cell surface, allowing T cells to recognize these antigens and initiate an immune response [6].

## Immune memory and T cell persistence

T cells are crucial for the development of immunological memory. When T cells encounter a pathogen, they can differentiate into memory T cells that "remember" the pathogen. This memory allows for a faster and more robust

immune response upon reencountering the same pathogen, a principle upon which vaccination is based [7].

While T cells are essential for effective immunity, their dysregulation can lead to autoimmune diseases. In autoimmune conditions, T cells mistakenly target and attack the body's own tissues. Understanding T cell dysregulation is critical for developing therapies to mitigate autoimmune diseases [8].

## The future of T cell immunotherapy

T cells are also at the forefront of cutting-edge therapies, such as CAR-T cell therapy. In this approach, T cells are genetically engineered to target cancer cells, revolutionizing the treatment of some forms of cancer. T cells stand as the vanguards of adaptive immunity, orchestrating a highly specific and tailored response to pathogens while laying the foundation for immunological memory [9]. Their diverse roles and functions make them indispensable in the body's defense against diseases, and they continue to be a subject of intensive research with the potential to transform immunotherapies, vaccination strategies, and our understanding of immunity [10].

## References

1. Moderbacher CR, Ramirez SI, Dan JM, et al. Antigen-specific adaptive immunity to SARS-CoV-2 in acute COVID-19 and associations with age and disease severity. *Cell*. 2020;183(4):996-1012.
2. Williams A, Flavell RA, Eisenbarth SC. The role of NOD-like receptors in shaping adaptive immunity. *Curr Opin Immunol*. 2010;22(1):34-40.
3. Veigl SJ. Adaptive immunity or evolutionary adaptation? Transgenerational immune systems at the crossroads. *Biology & Philosophy*. 2022;37(5):41.
4. Ardura MI, Banchereau R, Mejias A, et al. Enhanced monocyte response and decreased central memory T cells in children with invasive *Staphylococcus aureus* infections. *PLoS one*. 2009;4(5):e5446.
5. Grifoni A, Weiskopf D, Ramirez SI, et al. Targets of T cell responses to SARS-CoV-2 coronavirus in humans with COVID-19 disease and unexposed individuals. *Cell*. 2020;181(7):1489-501.
6. Zuccotti GV, Vigano A, Borelli M, et al. Modulation of innate and adaptive immunity by lactoferrin in human immunodeficiency virus (HIV)-infected, antiretroviral

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- therapy-naïve children. *Int J Antimicrob Agents*. 2007;29(3):353-5.
7. Nish SA, Schenten D, Wunderlich FT, et al. T cell-intrinsic role of IL-6 signaling in primary and memory responses. *elife*. 2014;3:e01949.
  8. Luo L, Li X, Zhang J, et al. Enhanced immune memory through a constant photothermal-metabolism regulation for cancer prevention and treatment. *Biomaterials*. 2021;270:120678.
  9. Illingworth J, Butler NS, Roetynck S, et al. Chronic exposure to *Plasmodium falciparum* is associated with phenotypic evidence of B and T cell exhaustion. *J Immunol*. 2013;190(3):1038-47.
  10. Soares AP, Kwong Chung CK, et al. Longitudinal changes in CD4<sup>+</sup> T-cell memory responses induced by BCG vaccination of newborns. *J Infect Dis*. 2013;207(7):1084-94.