

Retroviruses: Structure, Function, and Impact on Human Health

Sophia White*

Department of Immunology and Infectious Diseases, University of Harvard, United States

DESCRIPTION

Retroviruses are a unique class of viruses that utilize an enzyme called reverse transcriptase to convert their RNA genome into DNA, which then integrates into the host's genome. This characteristic allows retroviruses to establish persistent infections, making them particularly challenging to eradicate. Among the most well-known retroviruses is the Human Immunodeficiency Virus (HIV), which causes Acquired Immunodeficiency Syndrome (AIDS). However, retroviruses encompass a broader category of viruses, many of which have significant implications for both human and animal health. This article provides a comprehensive examination of retroviruses, their structure, replication mechanisms, classification, impact on health, and ongoing research efforts. Retroviruses belong to the family Retroviridae, which includes several genera that infect a wide range of hosts, from humans to animals. Unlike most viruses that replicate through direct RNA or DNA synthesis, retroviruses undergo reverse transcription, a unique process that allows them to integrate into the host genome permanently. Single-stranded RNA genome unlike double-stranded DNA viruses, retroviruses carry their genetic material as RNA. Reverse transcriptase enzyme converts RNA into complementary DNA (cDNA). Integration into host DNA the viral DNA integrates into the host genome using an enzyme called integrase. Persistent infection once integrated, the viral genome can remain dormant or be activated for replication. Retroviruses share a similar structural composition, which consists of envelope a lipid bilayer derived from the host cell membrane, containing viral glycoproteins essential for binding and entry into host cells. Capsid A protective protein shell that encloses the viral RNA and enzymes. RNA Genome consists of two identical copies of single-stranded RNA. Enzymes reverse Transcriptase converts viral RNA into DNA. Integrase facilitates the integration of viral DNA into the host genome. Protease helps process viral proteins necessary for new virus formation. Retroviruses follow a unique replication cycle that

involves six key stages attachment and Entry the virus binds to specific receptors on the host cell membrane, facilitating entry via fusion or endocytosis. Reverse Transcription once inside the host cell, reverse transcriptase converts the viral RNA into DNA. Integration the newly formed viral DNA is transported into the nucleus and integrated into the host genome by integrase. Transcription and Translation the host cell's machinery transcribes and translates the viral genome, producing viral proteins. Assembly newly synthesized viral components assemble into immature virions. Maturation and Release protease processes the viral proteins, leading to the formation of mature, infectious virions that bud off from the host cell. Retroviruses are classified into two subfamilies lentivirus (e.g., HIV-1, HIV-2): Causes slow, progressive infections. Deltaretrovirus (e.g., Human T-cell leukemia virus, HTLV) associated with leukemia and neurological diseases. Gammaretrovirus (e.g., Murine leukemia virus) mainly found in rodents but has implications for cancer research. Betaretrovirus (e.g., Jaagsiekte sheep retrovirus) primarily infects animals. Alpharetrovirus (e.g., Rous sarcoma virus) involved in oncogenesis (cancer formation). Typically cause asymptomatic infections in primates and are not associated with disease in humans. Retroviruses have significant implications for human health, particularly in the context of infectious diseases and cancer. HIV is the most well-known retrovirus and the causative agent of AIDS. HIV targets CD4+ T cells, essential components of the immune system, progressively weakening immune defenses. If untreated, HIV infection leads to AIDS, a condition characterized by severe immune suppression and vulnerability to opportunistic infections.

ACKNOWLEDGEMENT

None.

CONFLICT OF INTEREST

The author declares there is no conflict of interest.

Received:	02-December-2024	Manuscript No:	IPJHRV-25-22484
Editor assigned:	04-December-2024	PreQC No:	IPJHRV-25-22484 (PQ)
Reviewed:	18-December-2024	QC No:	IPJHRV-25-22484
Revised:	23-December-2024	Manuscript No:	IPJHRV-25-22484 (R)
Published:	30-December-2024	DOI:	10.21767/21767-9676.10.4.33

Corresponding author Sophia White, Department of Immunology and Infectious Diseases, University of Harvard, United States, E-mail: white44@gmail.com

Citation White S (2024) Retroviruses: Structure, Function, and Impact on Human Health. J HIV Retrovirus. 10:33.

Copyright © 2024 White S. 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.