

Limited elongation of the HIV-1 transcript

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Introduction

Reactome is open-source, open access, manually curated and peer-reviewed pathway database. Pathway annotations are authored by expert biologists, in collaboration with Reactome editorial staff and cross-referenced to many bioinformatics databases. A system of evidence tracking ensures that all assertions are backed up by the primary literature. Reactome is used by clinicians, geneticists, genomics researchers, and molecular biologists to interpret the results of high-throughput experimental studies, by bioinformaticians seeking to develop novel algorithms for mining knowledge from genomic studies, and by systems biologists building predictive models of normal and disease variant pathways.

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Literature references

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- Fabregat, A., Jupe, S., Matthews, L., Sidiropoulos, K., Gillespie, M., Garapati, P. et al. (2018). The Reactome Pathway Knowledgebase. *Nucleic Acids Res, 46*, D649-D655.
- Fabregat, A., Korninger, F., Viteri, G., Sidiropoulos, K., Marin-Garcia, P., Ping, P. et al. (2018). Reactome graph database: Efficient access to complex pathway data. *PLoS computational biology, 14*, e1005968.

Reactome database release: 88

This document contains 1 reaction (see Table of Contents)

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Limited elongation of the HIV-1 transcript **₹**

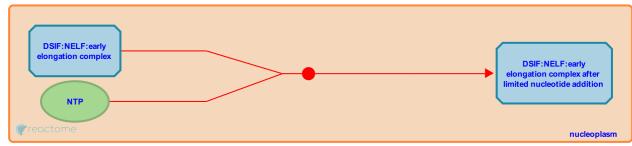
Stable identifier: R-HSA-167087

Type: binding

Compartments: nucleoplasm

Diseases: Human immunodeficiency virus infectious disease

Inferred from: Addition of nucleotides leads to transcript elongation (Homo sapiens)



In the absence of Tat, transcriptional elongation beyond position +59 does not occur (Kao et al., 1987).

Literature references

Peterlin, BM., Luciw, PA., Kao, SY., Calman, AF. (1987). Anti-termination of transcription within the long terminal repeat of HIV-1 by tat gene product. *Nature*, 330, 489-93.

Editions

2005-07-27	Authored	Matthews, L., Rice, AP.
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