

Disintegration of matrix layer

Aiken, C., Bukrinsky, M., Gopinathrao, G., Iordanskiy, S.

European Bioinformatics Institute, New York University Langone Medical Center, Ontario Institute for Cancer Research, Oregon Health and Science University.

The contents of this document may be freely copied and distributed in any media, provided the authors, plus the institutions, are credited, as stated under the terms of <u>Creative Commons Attribution 4.0 International (CC BY 4.0)</u> <u>License</u>. For more information see our <u>license</u>.

14/05/2024

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.

The development of Reactome is supported by grants from the US National Institutes of Health (P41 HG003751), University of Toronto (CFREF Medicine by Design), European Union (EU STRP, EMI-CD), and the European Molecular Biology Laboratory (EBI Industry program).

Literature references

- Fabregat, A., Sidiropoulos, K., Viteri, G., Forner, O., Marin-Garcia, P., Arnau, V. et al. (2017). Reactome pathway analysis: a high-performance in-memory approach. *BMC bioinformatics, 18,* 142. 7
- Sidiropoulos, K., Viteri, G., Sevilla, C., Jupe, S., Webber, M., Orlic-Milacic, M. et al. (2017). Reactome enhanced pathway visualization. *Bioinformatics*, 33, 3461-3467. A
- 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. *オ*

This document contains 1 reaction (see Table of Contents)

Disintegration of matrix layer ↗

Stable identifier: R-HSA-173642

Type: transition

Compartments: cytosol

Diseases: Human immunodeficiency virus infectious disease



After fusion of the viral membrane with the target cell membrane, the viral core, which is surrounded by a layer of Matrix (p17) proteins, is exposed to the cytoplasm. Disintegration of the Matrix layer allows for the conical-shaped viral core to be fully released, and allow for viral capsid dissociation and eventually reverse transcription. Dissociation of the Matrix layer is not well characterized, but is believed to occur due to disruption of protein-protein interactions as a result of the conditions of the cytoplasm (including pH), which differ from that of the internal viral structure.

Literature references

- Orenstein, J., Zhang, H., Pomerantz, RJ., Dornadula, G. (2000). Morphologic changes in human immunodeficiency virus type 1 virions secondary to intravirion reverse transcription: evidence indicating that reverse transcription may not take place within the intact viral core. *J Hum Virol*, *3*, 165-72.
- Fassati, A., Goff, SP. (2001). Characterization of intracellular reverse transcription complexes of human immunodeficiency virus type 1. *J Virol, 75*, 3626-35. *¬*

Editions

2006-02-17	Edited	Gopinathrao, G.
2006-04-03	Authored	Iordanskiy, S., Bukrinsky, M.
2006-10-31	Reviewed	Aiken, C.