

PtpA binds ATP6V1H

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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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Reactome database release: 88

This document contains 1 reaction ([see Table of Contents](#))

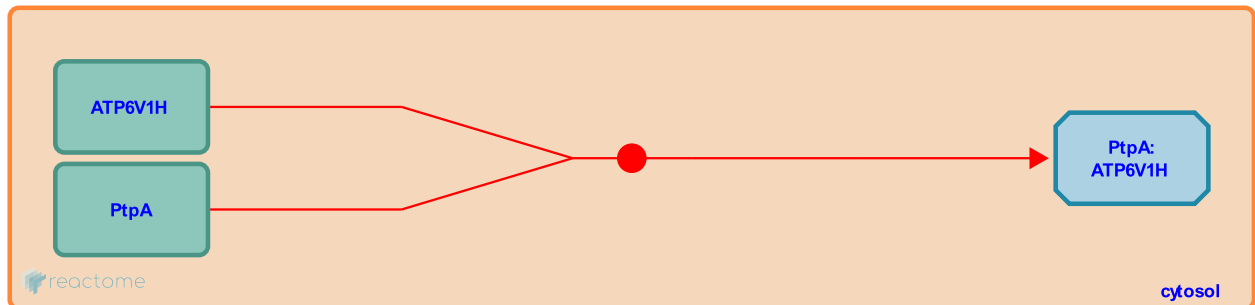
PtpA binds ATP6V1H [↗](#)

Stable identifier: R-HSA-9636397

Type: binding

Compartments: cytosol

Diseases: tuberculosis



The H subunit of human cytosolic ATPase (ATP6V1H) binds to the Mtb probable low molecular weight protein-tyrosine phosphatase (ptpA), directly impairing ATPase trafficking to the phagosome (Wong et al. 2011).

Literature references

Hmama, Z., Sun, J., Av-Gay, Y., Bach, H., Wong, D. (2011). Mycobacterium tuberculosis protein tyrosine phosphatase (PtpA) excludes host vacuolar-H⁺-ATPase to inhibit phagosome acidification. *Proc. Natl. Acad. Sci. U.S.A.*, 108, 19371-6. [↗](#)

Editions

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