

STAT1,STAT3,STAT6 phosphorylation

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03/05/2024

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

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Reactome database release: 88

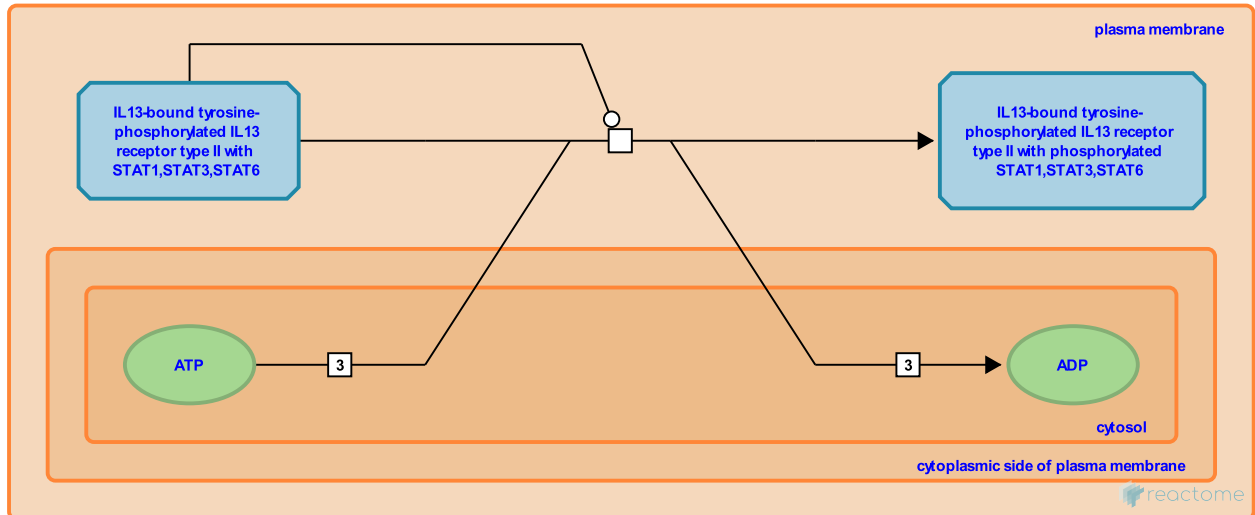
This document contains 1 reaction ([see Table of Contents](#))

STAT1,STAT3,STAT6 phosphorylation ↗

Stable identifier: R-HSA-6788582

Type: transition

Compartments: plasma membrane, cytosol, extracellular region



Once bound to the Interleukin-13 (IL13) type II receptor, Signal transducer and activator of transcription 3 (STAT3) is tyrosine phosphorylated by Janus kinase 2 (JAK2), while STAT1 and STAT6 are phosphorylated by Non-receptor tyrosine kinase 2 (TYK2) (Bhattacharjee et al. 2013).

Literature references

Yakubenko, VP., Mulya, A., Cathcart, MK., Kundu, S., Bhattacharjee, A., Shukla, M. (2013). IL-4 and IL-13 employ discrete signaling pathways for target gene expression in alternatively activated monocytes/macrophages. *Free Radic. Biol. Med.*, 54, 1-16. ↗

Editions

2015-07-01	Authored	Jupe, S.
2016-09-02	Edited	Jupe, S.
2016-09-02	Reviewed	Leibovich, SJ.