

SHC1 bound to integrin alphaIIb beta3 is phosphorylated somehow

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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

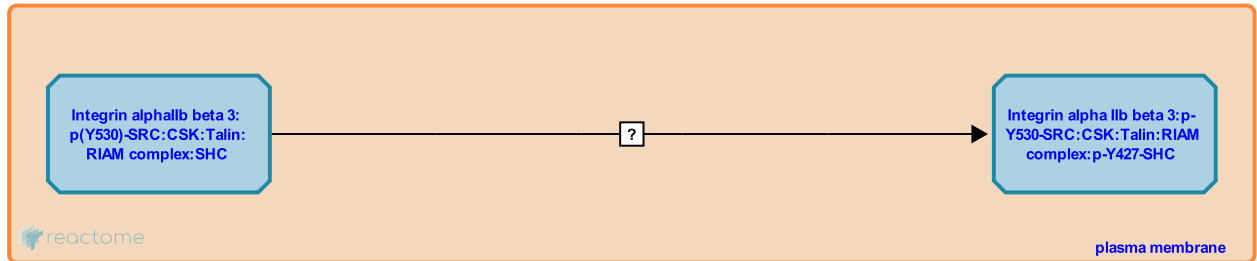
This document contains 1 reaction ([see Table of Contents](#))

SHC1 bound to integrin alphaIIb beta3 is phosphorylated somehow ↗

Stable identifier: R-HSA-443905

Type: uncertain

Compartments: plasma membrane



In a mechanism that is presumed to be analogous to signaling of SHC downstream of the insulin and TrkA receptors, SHC becomes phosphorylated and dissociates from the integrin alphaIIb beta3 complex.

Literature references

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Phillips, DR., Cowan, KJ., Law, DA. (2000). Identification of shc as the primary protein binding to the tyrosine-phosphorylated beta 3 subunit of alpha IIbeta 3 during outside-in integrin platelet signaling. *J Biol Chem*, 275, 36423-9. ↗

Editions

2009-09-04	Authored	Akkerman, JW.
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