

APC/C:Cdh1-mediated degradation of Skp2

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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

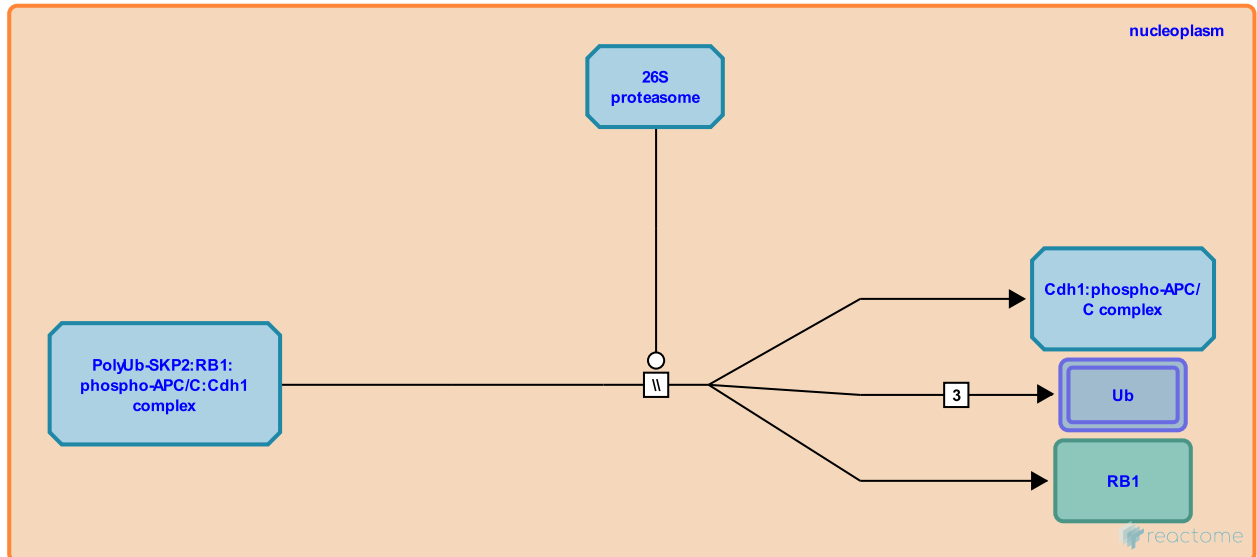
This document contains 1 reaction ([see Table of Contents](#))

APC/C:Cdh1-mediated degradation of Skp2 [↗](#)

Stable identifier: R-HSA-188191

Type: omitted

Compartments: nucleoplasm



SKP2 is degraded by the anaphase promoting complex/Cyclosome and its activator FZR1 (Cdh1) [APC/C(Cdh1)] (Bashir et al, 2004; Wei et al, 2004). The tight regulation of APC/C(Cdh1) activity ensures the timely elimination SKP2 and, thus, plays a critical role in controlling the M/G1 transition (mitotic exit). APC/C:Cdh1-mediated degradation of SKP2 depends on RB1, as RB1 recruits SKP2 to the APC/C:Cdh1 complex, by simultaneously interacting with SKP2 and FZR1. RB1 does not undergo APC/C:Cdh1-mediated ubiquitination (Binne et al. 2007).

Literature references

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Editions

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