Inactivation of Cyclin A:Cdk2 complexes 

by p27/p21

Coqueret, O., Inga, A., Matthews, L., Pagano, M., Zaccara, S.
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


Inactivation of Cyclin A:Cdk2 complexes by p27/p21

Stable identifier: R-HSA-187934

Type: transition

Compartments: nucleoplasm

During G1, the activity of cyclin-dependent kinases (CDKs) is controlled by the CDK inhibitors (CKIs) CDKN1A (p21) and CDKN1B (p27), thereby preventing premature entry into S phase (Guardavaccaro and Pagano, 2006).

Literature references


Editions

<table>
<thead>
<tr>
<th>Date</th>
<th>Action</th>
<th>Author</th>
</tr>
</thead>
<tbody>
<tr>
<td>2006-09-19</td>
<td>Authored</td>
<td>Pagano, M.</td>
</tr>
<tr>
<td>2006-09-28</td>
<td>Edited</td>
<td>Matthews, L.</td>
</tr>
<tr>
<td>2006-10-06</td>
<td>Reviewed</td>
<td>Coqueret, O.</td>
</tr>
<tr>
<td>2016-02-04</td>
<td>Reviewed</td>
<td>Inga, A., Zaccara, S.</td>
</tr>
</tbody>
</table>