

Release of PLCG from FCGR

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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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- Fabregat, A., Jupe, S., Matthews, L., Sidiropoulos, K., Gillespie, M., Garapati, P. et al. (2018). The Reactome Pathway Knowledgebase. *Nucleic Acids Res*, 46, D649-D655. [↗](#)
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Reactome database release: 88

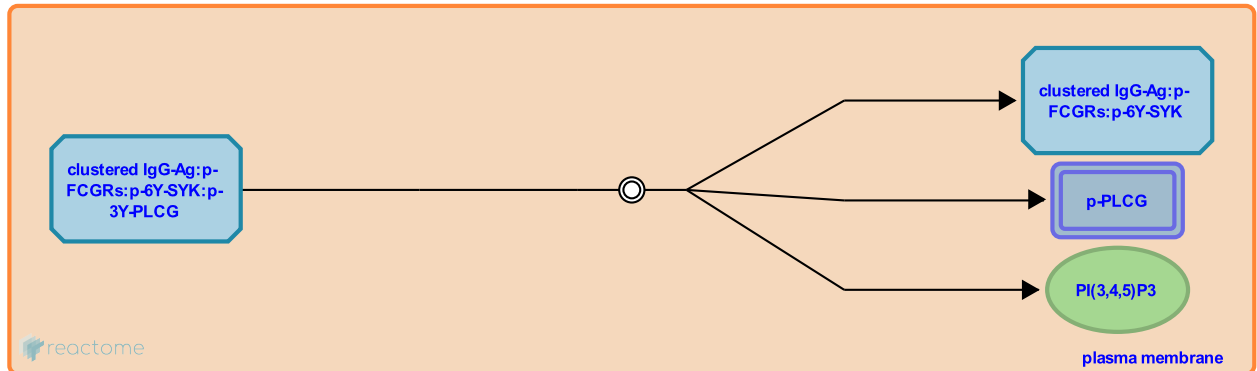
This document contains 1 reaction ([see Table of Contents](#))

Release of PLCG from FCGR [↗](#)

Stable identifier: R-HSA-2029272

Type: dissociation

Compartments: plasma membrane, cytosol, extracellular region



Activated PLCG translocates to the plasma membrane and interacts with the inositol ring of the membrane bound phosphatidylinositol 4,5-bisphosphate (PIP₂) with its PH domain. The active enzyme promotes intracellular signaling by catalysing the hydrolysis of PIP₂ to generate the second messengers IP₃ and DAG.

Literature references

Ji, Q., Carpenter, G. (1999). Phospholipase C-gamma as a signal-transducing element. *Exp Cell Res*, 253, 15-24. [↗](#)

Editions

2012-01-04	Authored, Edited	Garapati, P V.
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