

# **Product datasheet for LY300508**

#### OriGene Technologies, Inc.

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## Phospholipase C beta 3 (PLCB3) Human Knockdown Lysate

#### **Product data:**

**Product Type:** Knockdown Lysates

**Description:** WB-validated PLCB3 Knockdown HT-1080 Cell Lysate

Species: Human
Tag: Tag Free

**Synonyms:** PLCB3; Phospholipase C Beta 3; 1-Phosphatidylinositol 4,5-Bisphosphate Phosphodiesterase

Beta-3; Phospholipase C, Beta 3 (Phosphatidylinositol-Specific); Phosphoinositide

Phospholipase C-Beta-3; EC 3.1.4.11; Phospholipase C-Beta-3; PLC Beta 3; PLC-Beta-3; SMDCD

**Predicted MW:** 139 kDa

**Components:** 1 vial of 100 ug WT HT-1080 cell lysate

1 vial of 100 ug PLCB3 KD HT-1080 cell lysate

Storage: Store at -20 °C for two years.

**Concentration:** Lot-specific

**Buffer:** IntactProtein Cell-Tissue Lysis buffer

**Locus ID:** 5331

UniProt ID: Q01970

**Protein Families:** Druggable Genome

**Protein Pathways:** Alzheimer's disease, Calcium signaling pathway, Chemokine signaling pathway, Gap junction,

GnRH signaling pathway, Huntington's disease, Inositol phosphate metabolism, Long-term

depression, Long-term potentiation, Melanogenesis, Metabolic pathways,

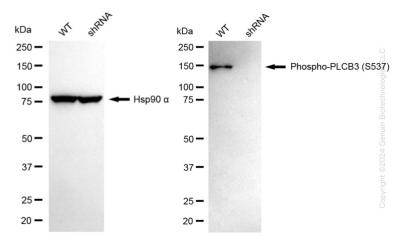
Phosphatidylinositol signaling system, Vascular smooth muscle contraction, Wnt signaling

pathway

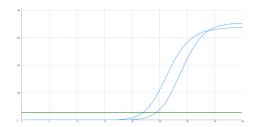




# **Product images:**



Western blotting analysis. PLCB3 protein expression in wild-type (WT) and shRNA knockdown (KD) HT-1080 cells was detected using Western blotting. Hsp90  $\alpha$  served as a loading control. The blots were incubated with primary antibodies against PLCB3 and Hsp90  $\alpha$ , respectively, followed by incubating with HRP-conjugated goat anti-rabbit secondary antibody. Images were developed using FeQ^ $\rm IM$  ECL Substrate Kit.



Genotype	Ct Value
Wild-Type	21.81
Knock-Down	24.25
$\Delta Ct (Ct_{KD}-Ct_{WT})$	2.15
% mRNA Reduction	<b>4</b> 77%

RT-qPCR analysis. HT-1080 cells were infected with PLCB3-specific shRNA lentiviral particles, total RNA was extracted from wild-type and knockdown cells, RT-qPCR was performed using gene-specific primers. ΔCt (CtKD-CtWT) was used to calculate mRNA reduction (%) between wild-type and knockdown cells using the following formula: (1-1/2ΔCt) x 100%.