

## OriGene Technologies, Inc.

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## Product datasheet for TL310390V

## Phospholipase A2 IIA (PLA2G2A) Human shRNA Lentiviral Particle (Locus ID 5320)

## **Product data:**

| Product Type: | shRNA Lentiviral Particles  |
|---------------|---|
| Product Name: | Phospholipase A2 IIA (PLA2G2A) Human shRNA Lentiviral Particle (Locus ID 5320)  |
| Locus ID:     | 5320  |
| Synonyms:     | MOM1; PLA2; PLA2B; PLA2L; PLA2S; PLAS1; sPLA2   |
| Vector:       | pGFP-C-shLenti (TR30023)  |
| Format:       | Lentiviral particles  |
| Components:   | PLA2G2A - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1<br>scramble control), 0.5 ml each, >10^7 TU/ml.   |
| RefSeq:       | <u>NM 000300, NM 001161727, NM 001161728, NM 001161729, NM 000300.1, NM 000300.2, NM 000300.3, NM 001161727.1, NM 001161728.1, NM 001161729.1, BC005919, BM708748, NM 000300.4, NM 001161727.2, NM 001161728.2</u>  |
| UniProt ID:   | <u>P14555</u>   |
|               |   |
| Summary:      | The protein encoded by this gene is a member of the phospholipase A2 family (PLA2). PLA2s constitute a diverse family of enzymes with respect to sequence, function, localization, and divalent cation requirements. This gene product belongs to group II, which contains secreted form of PLA2, an extracellular enzyme that has a low molecular mass and requires calcium ions for catalysis. It catalyzes the hydrolysis of the sn-2 fatty acid acyl ester bond of phosphoglycerides, releasing free fatty acids and lysophospholipids, and thought to participate in the regulation of the phospholipid metabolism in biomembranes. Several alternatively spliced transcript variants with different 5' UTRs have been found for this gene. [provided by RefSeq, Sep 2009] |



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|                            | Phospholipase A2 IIA (PLA2G2A) Human shRNA Lentiviral Particle (Locus ID 5320) – TL310390V   |
|----------------------------|--|
| Performance<br>Guaranteed: | OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples. |
|                            | For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with   |

(90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

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