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Product datasheet for RC211985L3V

Plasma Kallikrein 1B (KLKB1) (NM_000892) Human Tagged ORF Clone Lentiviral Particle

Product data:

| Product Type: | Lentiviral Particles |
|------------------------------|---|
| Product Name: | Plasma Kallikrein 1B (KLKB1) (NM_000892) Human Tagged ORF Clone Lentiviral Particle |
| Symbol: | Plasma Kallikrein 1B |
| Synonyms: | KLK3; PKK; PKKD; PPK |
| Mammalian Cell Selection: | Puromycin |
| Vector: | pLenti-C-Myc-DDK-P2A-Puro (PS100092) |
| Tag: | Myc-DDK |
| ACCN: | NM_000892 |
| ORF Size: | 1914 bp |
| ORF Nucleotide Sequence: | The ORF insert of this clone is exactly the same as(RC211985). |
| OTI Disclaimer: | The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. <u>More info</u> |
| OTI Annotation: | This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene. |
| RefSeq: | <u>NM 000892.2</u> |
| RefSeq Size: | 2245 bp |
| RefSeq ORF: | 1917 bp |
| Locus ID: | 3818 |
| UniProt ID: | <u>P03952</u> |
| Cytogenetics: | 4q35.2 |
| Domains: | APPLE, Tryp_SPc, PAN |
| Protein Families: | Druggable Genome, Protease |



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| | Plasma Kallikrein 1B (KLKB1) (NM_000892) Human Tagged ORF Clone Lentiviral Particle – RC211985L3V |
|----------------|--|
| Protein Pathwa | s: Complement and coagulation cascades |
| MW: | 71.37 kDa |
| Gene Summary | This gene encodes a glycoprotein that participates in the surface-dependent activation of blood coagulation, fibrinolysis, kinin generation and inflammation. The encoded preproprotein present in plasma as a non-covalent complex with high molecular weight kininogen undergoes proteolytic processing mediated by activated coagulation factor XII to generate a disulfide-linked, heterodimeric serine protease comprised of heavy and light chains. Certain mutations in this gene cause prekallikrein deficiency. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jan 2016] |

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