

Product datasheet for RC224278L4V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Fibrinogen alpha chain (FGA) (NM_000508) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Fibrinogen alpha chain (FGA) (NM 000508) Human Tagged ORF Clone Lentiviral Particle

Symbol: FGA
Synonyms: Fib2

Mammalian Cell Puromycin

Selection:

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

ACCN: NM_000508

ORF Size: 2598 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC224278).

Sequence:

Domains:

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. More info

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 000508.3</u>

 RefSeq Size:
 3655 bp

 RefSeq ORF:
 2601 bp

 Locus ID:
 2243

 UniProt ID:
 P02671

 Cytogenetics:
 4q31.3

Protein Families: Druggable Genome, Secreted Protein

FBG





Fibrinogen alpha chain (FGA) (NM_000508) Human Tagged ORF Clone Lentiviral Particle – RC224278L4V

Protein Pathways: Complement and coagulation cascades

MW: 94.97 kDa

Gene Summary: This gene encodes the alpha subunit of the coagulation factor fibrinogen, which is a

component of the blood clot. Following vascular injury, the encoded preproprotein is proteolytically processed by thrombin during the conversion of fibrinogen to fibrin.

Mutations in this gene lead to several disorders, including dysfibrinogenemia,

hypofibrinogenemia, afibrinogenemia and renal amyloidosis. Alternative splicing results in multiple transcript variants, at least one of which encodes an isoform that undergoes

proteolytic processing. [provided by RefSeq, Jan 2016]