

## Product datasheet for **RC215223L4V**

### FECH (NM\_001012515) Human Tagged ORF Clone Lentiviral Particle

#### Product data:

|                           |  |
|---------------------------|--|
| Product Type:             | Lentiviral Particles   |
| Product Name:             | FECH (NM_001012515) Human Tagged ORF Clone Lentiviral Particle   |
| Symbol:                   | FECH   |
| Synonyms:                 | EPP; EPP1; FCE   |
| Mammalian Cell Selection: | Puromycin  |
| Vector:                   | pLenti-C-mGFP-P2A-Puro (PS100093)  |
| Tag:                      | mGFP   |
| ACCN:                     | NM_001012515   |
| ORF Size:                 | 1287 bp  |
| ORF Nucleotide Sequence:  | The ORF insert of this clone is exactly the same as(RC215223).   |
| 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. <a href="#">More info</a> |
| 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:                   | <a href="#">NM_001012515.1</a>   |
| RefSeq Size:              | 3835 bp  |
| RefSeq ORF:               | 1290 bp  |
| Locus ID:                 | 2235   |
| UniProt ID:               | <a href="#">P22830</a>   |
| Cytogenetics:             | 18q21.31   |
| Protein Families:         | Druggable Genome   |
| Protein Pathways:         | Metabolic pathways, Porphyrin and chlorophyll metabolism   |



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**MW:** 48.63 kDa

**Gene Summary:** The protein encoded by this gene is localized to the mitochondrion, where it catalyzes the insertion of the ferrous form of iron into protoporphyrin IX in the heme synthesis pathway. Mutations in this gene are associated with erythropoietic protoporphyria. Two transcript variants encoding different isoforms have been found for this gene. A pseudogene of this gene is found on chromosome 3.[provided by RefSeq, May 2010]