

OriGene Technologies, Inc.

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Product datasheet for TP701127

FOXL2 Mutant (C134W) Human Recombinant Protein

Product data:

| Product Type: | Mutant Proteins |
|--|--|
| Description: | Purified mutant recombinant protein of Human forkhead box L2 (FOXL2), mutation atIC134WI |
| Species: | Human |
| Expression Host: | HEK293T |
| Expression cDNA Clone or AA Sequence: | A DNA sequence from TrueORF clone, RC209127, encoding the full-length of FOXL2(C134W) |
| Tag: | Myc-DDK |
| Predicted MW: | 38.6 kDa |
| Concentration: | >0.05 µg/µL as determined by microplate Bradford method |
| Purity: | > 80% as determined by SDS-PAGE and Coomassie blue staining |
| Buffer: | 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol |
| Note: | For culture applications, please filter before use. Note that you may experience some loss of protein during the filtration process. |
| Storage: | Store at -80°C after receiving vials. |
| Stability: | Stable for at least 12 months from receipt of products under proper storage and handling conditions. Avoid repeated freeze-thaw cycles. |
| RefSeq: | <u>NP 075555.1</u> |
| Locus ID: | 668 |
| RefSeq Size: | 2744 |
| Cytogenetics: | 3q22.3 |
| RefSeq ORF: | 1128 |
| Synonyms: | BPES; BPES1; PFRK; PINTO; POF3 |
| Summary: | This gene encodes a forkhead transcription factor. The protein contains a fork-head DNA- binding domain and may play a role in ovarian development and function. Expansion of a polyalanine repeat region and other mutations in this gene are a cause of blepharophimosis syndrome and premature ovarian failure 3. [provided by RefSeq, Jul 2016] |



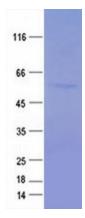
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GRIGENE FOXL2 Mutant (C134W) Human Recombinant Protein – TP701127

Protein Families:

Druggable Genome, Transcription Factors

Product images:



Purified recombinant protein FOXL2 (C134W) was analyzed by SDS-PAGE gel and Coomossie Blue Staining.

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