

Product datasheet for TP505914

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

Galt (NM 016658) Mouse Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Purified recombinant protein of Mouse galactose-1-phosphate uridyl transferase (Galt), with C-

terminal MYC/DDK tag, expressed in HEK293T cells, 20ug

Species: Mouse

Expression Host: HEK293T

Expression cDNA >MR205914 protein sequence Clone or AA

Red=Cloning site Green=Tags(s)

Sequence:

MAATFRASEHQHIRYNPLQDEWVLVSAHRMKRPWQGQVEPQLLKTVPRHDPLNPLCPGATRANGEVNPHY DGTFLFDNDFPALQPDAPDPGTSDHPLFRAEAARGVCKVMCFHPWSDVTLPLMSVPEIRAVIDAWASVTE ELGAQYPWVQIFENKGAMMGCSNPHPHCQVWASSFLPDIAQREERSQQTYHSQHGKPLLLEYGHQELLRK ERLVLTSEHWIVLVPFWAVWPFQTLLLPRRHVRRLPELNPAERDDLASIMKKLLTKYDNLFETSFPYSMG WHGAPTGLKTGATCDHWQLHAHYYPPLLRSATVRKFMVGYEMLAQAQRDLTPEQAAERLRALPEVHYCLA

QKDKETAAIA

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

C-MYC/DDK Tag:

Predicted MW: 41.2 kDa

Concentration: >0.05 µg/µL as determined by microplate BCA 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

For testing in cell culture applications, please filter before use. Note that you may experience Note:

some loss of protein during the filtration process.

Storage: Store at -80°C after receiving vials.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and handling

conditions. Avoid repeated freeze-thaw cycles.

NP 057867 RefSeq:

Locus ID: 14430

UniProt ID: Q03249, A2AMS3





Galt (NM_016658) Mouse Recombinant Protein - TP505914

RefSeq Size: 2000

Cytogenetics: 4 22.07 cM

RefSeq ORF: 1083

Synonyms: AW553376

Summary: The protein encoded by this gene is the second enzyme in the Leloir pathway, the metabolic

pathway for D-galactose catabolism. It catalyzes the conversion of galactose-1-phosphate and

uridine diphosphate-glucose to glucose-1-phosphate and uridine diphosphate galactose. Deficiency of this enzyme causes the genetic metabolic disorder galactosemia. Mice lacking this protein accumulate high levels of galactose and galactose-1 phosphate but are viable and fertile. This protein is negatively regulated through signaling by the polypeptide hormone prolactin, specifically via the short isoform of the prolactin receptor and the transcription factor Forkhead box O3. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2014]