

Ornithine Aminotransferase Human

Description: Ornithine Aminotransferase Human Recombinant produced in E.Coli is a single, non-glycosylated polypeptide chain containing 408 amino acids (33-439 a.a.) and having a molecular weight of 45.2kDa. The Ornithine Aminotransferase is purified by proprietary chromatographic techniques.

Catalog #: ENPS-479

For research use only.

Synonyms: DKFZp781A11155, HOGA, OATASE, Ornithine aminotransferase mitochondrial, Ornithine--oxo-acid aminotransferase, OAT, OKT, GACR.

Source: Escherichia Coli.

Physical Appearance: Sterile filtered colorless solution.

Amino Acid Sequence: MTVQGPPTSD DIFEREYKYG AHNYHPLPVA LERGKGIYLW
DVEGRKYFDF LSSYSANVQG HCHPKIVNAL KSQVDKLTLT SRAFYNNVLG EYEEYITKLF
NYHKVLPMT GVEAGETACK LARKWGYTVK GIQKYKAKIVAAGNFWGRT LSAISSSTDP
TSYDGFQPFM PGFDIIPYND LPALERALQD PNVAAFMVEP IQGEAGVVVP DPGYLMGVRE
LCTRHQVLFI ADEI

Purity: Greater than 90.0% as determined by SDS-PAGE.

Formulation:

The Ornithine Aminotransferase protein solution contains 20mM Tris, pH-8, and 10% glycerol.

Stability:

Store at 4°C if entire vial will be used within 2-4 weeks. Store, frozen at -20°C for longer periods of time. For long term storage it is recommended to add a carrier protein (0.1% HSA or BSA). Avoid multiple freeze-thaw cycles.

Usage:

NeoBiolab's products are furnished for LABORATORY RESEARCH USE ONLY. The product may not be used as drugs, agricultural or pesticidal products, food additives or household chemicals.

Introduction:

Ornithine Aminotransferase is a mitochondrial enzyme which is an important factor that converts arginine and ornithine into the major excitatory and inhibitory neurotransmitters glutamate and GABA. Ornithine Aminotransferase mutations result in a deficiency that cause the autosomal recessive eye disease Gyrate Atrophy.

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