

Recombinant Human FTO protein ab109039

2 References

Description	
Product name	Recombinant Human FTO protein
Purity	> 90 % SDS-PAGE. ab109039 is 0.2µm filtered.
Endotoxin level	< 1.000 Eu/µg
Expression system	Escherichia coli
Accession	<u>Q9C0B1</u>
Protein length	Full length protein
Animal free	No
Nature	Recombinant
Species	Human
Sequence	KRTPAEEREREAKKLRLLLEELEDTWLPYLTPKDDEFYQQ WQLKYPKLIL REASSVSEELHKEVQEAFLLHKGCLFRDLVRIQGDLL TPVSRILIGN PGCTYKYLNTRLFTVPWPVKGSNIKHTAEIAACETFLKL NDYLQIETI QALEELAAKEKANEDAVPLCMSADFPRVGMGSSYNGQD EVDIKSRAAYNV TLLNFMDPQKMPYLKEEYPFGMGKMAVSWHHDENLVDR SAVAVYSYSCG PEEESEDDSHLEGRDPDIWHVGFKISWDIETPGLAIPHQ GDCYFMLDDL NATHQHCVLAGSQPRFSSTHRVAECSTGTLDYILQRCQLA LQNVCDVDN DDVSLKSFEPAVLKQGEEIHNEVEFEWLRQFWFQGNRYR KCTDWWCQPM QLEALWKKMEGVTVNAVLHEVKREGLPVEQRNEILTALAS LTARQNLRRE WHARCQSRIARTLPADQKPECRPYWEKDDASMPPLPFDL TDIVSELRGQLL EAKP
Predicted molecular weight	65 kDa including tags
Amino acids	2 to 505
Tags	His tag N-Terminus

Specifications

Our **Abpromise guarantee** covers the use of **ab109039** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
pH: 8.20
Constituents: 0.869% Tris HCl, 0.87% Sodium chloride

General Info

Function Dioxygenase that repairs alkylated DNA and RNA by oxidative demethylation. Has highest activity towards single-stranded RNA containing 3-methyluracil, followed by single-stranded DNA containing 3-methylthymine. Has low demethylase activity towards single-stranded DNA containing 1-methyladenine or 3-methylcytosine. Has no activity towards 1-methylguanine. Has no detectable activity towards double-stranded DNA. Requires molecular oxygen, alpha-ketoglutarate and iron. Contributes to the regulation of the global metabolic rate, energy expenditure and energy homeostasis. Contributes to the regulation of body size and body fat accumulation.

Tissue specificity Ubiquitously expressed, with relatively high expression in adrenal glands and brain; especially in hypothalamus and pituitary.

Involvement in disease Defects in FTO are the cause of growth retardation developmental delay coarse facies and early death (GRDDCFED) [MIM:612938]. The disease consists of a severe children multiple congenital anomaly syndrome with death by the age of 3 years. All affected individuals had postnatal growth retardation, microcephaly, severe psychomotor delay, functional brain deficits, and characteristic facial dysmorphism. In some patients, structural brain malformations, cardiac defects, genital anomalies, and cleft palate were also observed.

Sequence similarities Belongs to the fto family.

Domain The 3D-structure of the Fe2OG dioxygenase domain is similar to that of the Fe2OG dioxygenase domain found in the bacterial DNA repair dioxygenase alkB and its mammalian orthologs, but sequence similarity is very low. As a consequence, the domain is not detected by protein signature databases.

Cellular localization Nucleus.

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