abcam

Product datasheet

Human LOXL1 peptide ab97928

1 图像

描述

产**品名称** 人LOXL1多肽

纯**度** > 70 % HPLC.

70 - 90% by HPLC

Accession Q08397

无动物成分 No

性质 Synthetic

种属Human预测分子量63 kDa

技术指标

Our Abpromise guarantee covers the use of ab97928 in the following tested applications.

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

应**用** Blocking

形式 Lyophilized

补充说明 - First try to dissolve a small amount of peptide in either water or buffer. The more charged

residues on a peptide, the more soluble it is in aqueous solutions.

- If the peptide doesn't dissolve try an organic solvent e.g. DMSO, then dilute using water or

buffer.

- Consider that any solvent used must be compatible with your assay. If a peptide does not

dissolve and you need to recover it, lyophilise to remove the solvent.

- Gentle warming and sonication can effectively aid peptide solubilisation. If the solution is

cloudy or has gelled the peptide may be in suspension rather than solubilised.

- Peptides containing cysteine are easily oxidised, so should be prepared in solution just prior

to use.

制备和贮存

稳定性和存储 Shipped at 4°C. Store at -20°C.

Information available upon request.

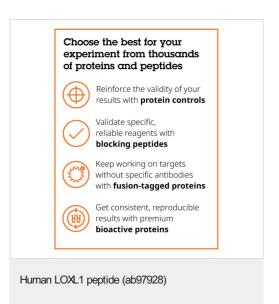
相关性

LOXL1 is a member of the lysyl oxidase gene family. The prototypic member of the family is essential to the biogenesis of connective tissue, encoding an extracellular copper-dependent amine oxidase that catalyses the first step in the formation of crosslinks in collagens and elastin. A highly conserved amino acid sequence at the C-terminus end appears to be sufficient for amine oxidase activity, suggesting that each family member may retain this function. The N-terminus is poorly conserved and may impart additional roles in developmental regulation, senescence, tumor suppression, cell growth control, and chemotaxis to each member of the family. LOXL1 is active on elastin and collagen substrates. Genetic variations in LOXL1 are associated with risk of developing exfoliation syndrome (XFS) [MIM:177650]; also called exfoliation glaucoma (XFG). Exfoliation syndrome (XFS) is characterized by accumulation of abnormal microfibrillar deposits that line the aqueous bathed surfaces of the anterior segment of the eye. The prevalence of XFS increases with age, and a number of studies have pointed to a geographical clustering of XFS, although this condition is found worldwide; reported prevalence rates average about 10 to 20% of the general population over age 60.

细胞定位

Secreted, extracellular space.

图片



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