abcam

Product datasheet

Human AICDA peptide ab13718

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Description

Product name Human AICDA peptide

Purity > 90 % HPLC.

Animal free No

Nature Synthetic

Species Human

Specifications

Our Abpromise guarantee covers the use of ab13718 in the following tested applications.

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

Applications

Blocking - Blocking peptide for Anti-AlCDA antibody - ChIP Grade (ab5197)

Form

Liquid

Additional notes

- 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.

Preparation and Storage

Stability and Storage

Shipped at $4\,^{\circ}\text{C}$. Upon delivery aliquot and store at -20 $^{\circ}\text{C}$ or -80 $^{\circ}\text{C}$. Avoid repeated freeze / thaw

cycles.

Information available upon request.

General Info

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Function RNA-editing deaminase involved in somatic hypermutation, gene conversion, and class-switch

recombination. Required for several crucial steps of B-cell terminal differentiation necessary for

efficient antibody responses.

Tissue specificity Strongly expressed in lymph nodes and tonsils.

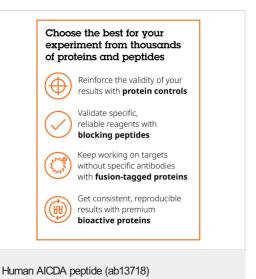
Involvement in diseaseDefects in AICDA are the cause of hyper-lgM immunodeficiency syndrome type 2 (HIGM2)

[MIM:605258]; also known as hyper-lgM syndrome 2. HIGM2 is an autosomal recessive disorder characterized by normal or elevated serum lgM levels with absence of lgG, lgA, and lgE, resulting in a profound susceptibility to bacterial infections. HIGM2 causes the absence of lg class switch recombination (CSR), the lack of lg somatic hypermutations, and lymph node hyperplasia caused

by the presence of giant germinal centers.

Sequence similarities Belongs to the cytidine and deoxycytidylate deaminase family.

Images



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