

MLL5 Antibody (N-term) Blocking Peptide
Synthetic peptide
Catalog # BP14173a**Specification**

MLL5 Antibody (N-term) Blocking Peptide - Product InformationPrimary Accession [Q8IZD2](#)**MLL5 Antibody (N-term) Blocking Peptide - Additional Information****Gene ID** 55904**Other Names**

Histone-lysine N-methyltransferase 2E, Lysine N-methyltransferase 2E, Myeloid/lymphoid or mixed-lineage leukemia protein 5, KMT2E, MLL5

Format

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

Precautions

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

MLL5 Antibody (N-term) Blocking Peptide - Protein Information**Name** KMT2E**Synonyms** MLL5**Function**

Associates with chromatin regions downstream of transcriptional start sites of active genes and thus regulates gene transcription (PubMed: [23629655](http://www.uniprot.org/citations/23629655), PubMed: [24130829](http://www.uniprot.org/citations/24130829), PubMed: [23798402](http://www.uniprot.org/citations/23798402)). Chromatin interaction is mediated via the binding to tri-methylated histone H3 at 'Lys-4' (H3K4me3) (PubMed: [24130829](http://www.uniprot.org/citations/24130829), PubMed: [23798402](http://www.uniprot.org/citations/23798402)). Key regulator of hematopoiesis involved in terminal myeloid differentiation and in the regulation of hematopoietic stem cell (HSCs) self-renewal by a mechanism that involves DNA methylation (By similarity). Also acts as an important cell cycle regulator, participating in cell cycle regulatory network machinery at multiple cell cycle stages including G1/S transition, S phase progression and mitotic entry (PubMed: [14718661](http://www.uniprot.org/citations/14718661), PubMed: [18573682](http://www.uniprot.org/citations/18573682), PubMed: [19264965](http://www.uniprot.org/citations/19264965)).

target="_blank">19264965, PubMed:23629655). Recruited to E2F1 responsive promoters by HCFC1 where it stimulates tri-methylation of histone H3 at 'Lys-4' and transcriptional activation and thereby facilitates G1 to S phase transition (PubMed:23629655). During myoblast differentiation, required to suppress inappropriate expression of S-phase-promoting genes and maintain expression of determination genes in quiescent cells (By similarity).

Cellular Location

Chromosome. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome. Nucleus speckle. Note=Absent from the nucleolus (PubMed:14718661). Localizes to chromosome during interphase and to centrosomes during mitosis (PubMed:23798402). Dissociation from mitotic chromosome is likely due to histone H3 phosphorylation on 'Thr-3' and 'Thr-6' (PubMed:23798402). [Isoform NKp44L]: Cytoplasm. Cell membrane; Peripheral membrane protein

Tissue Location

Widely expressed in both adult and fetal tissues (PubMed:12101424, PubMed:23958951). Highest levels of expression observed in fetal thymus and kidney and in adult hematopoietic tissues, jejunum and cerebellum (PubMed:12101424, PubMed:23958951). Isoform NKp44L: Not detected on circulating cells from healthy individuals, but is expressed on a large panel of tumor and transformed cells (PubMed:23958951).

MLL5 Antibody (N-term) Blocking Peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

MLL5 Antibody (N-term) Blocking Peptide - Images

MLL5 Antibody (N-term) Blocking Peptide - Background

This gene is a member of the myeloid/lymphoid or mixed-lineage leukemia (MLL) family and encodes a protein with an N-terminal PHD zinc finger and a central SET domain. Overexpression of the protein inhibits cell cycle progression. Alternative transcriptional splice variants have been characterized. [provided by RefSeq].

MLL5 Antibody (N-term) Blocking Peptide - References

Liu, J., et al. J. Biol. Chem. 285(27):20904-20914(2010) Fujiki, R., et al. Nature 459(7245):455-459(2009) Cheng, F., et al. Int. J. Biochem. Cell Biol. 40(11):2472-2481(2008) Sun, X.J., et al. PLoS ONE 3 (1), E1499 (2008) :Olsen, J.V., et al. Cell 127(3):635-648(2006)