

H3f3b Antibody (C-Term)
Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP21780b**Specification**

H3f3b Antibody (C-Term) - Product Information

Application	WB,E
Primary Accession	P84244
Reactivity	Mouse
Host	Rabbit
Clonality	polyclonal
Isotype	Rabbit IgG
Calculated MW	15328

H3f3b Antibody (C-Term) - Additional Information**Gene ID** 15078;15081**Other Names**

Histone H33, H3f3a, H33a

Target/Specificity

This H3f3b antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 103-136 amino acids from mouse H3f3b.

Dilution

WB~~1:2000

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

H3f3b Antibody (C-Term) is for research use only and not for use in diagnostic or therapeutic procedures.

H3f3b Antibody (C-Term) - Protein Information**Name** H3-3a {ECO:0000250|UniProtKB:P84243}

Function Variant histone H3 which replaces conventional H3 in a wide range of nucleosomes in active genes. Constitutes the predominant form of histone H3 in non-dividing cells and is incorporated into chromatin independently of DNA synthesis. Deposited at sites of nucleosomal displacement throughout transcribed genes, suggesting that it represents an epigenetic imprint of

transcriptionally active chromatin. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling.

Cellular Location

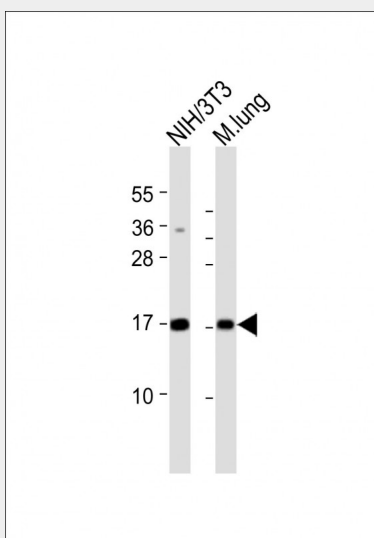
Nucleus. Chromosome.

H3f3b Antibody (C-Term) - Protocols

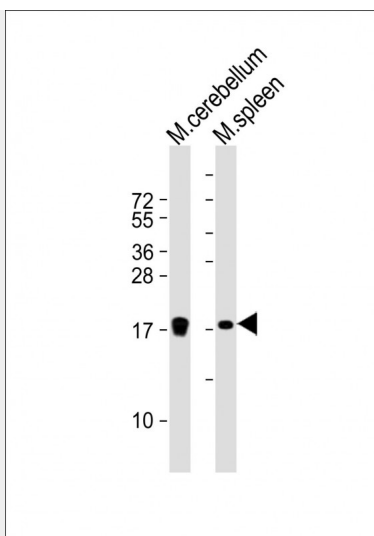
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

H3f3b Antibody (C-Term) - Images



All lanes : Anti-H3f3b Antibody (C-Term) at 1:8000 dilution Lane 1: NIH/3T3 whole cell lysate Lane 2: mouse lung lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 15 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



All lanes : Anti-H3f3b Antibody (C-Term) at 1:2000 dilution Lane 1: mouse cerebellum lysate Lane 2: mouse spleen lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 15 kDa Blocking/Dilution buffer: 5% NFDM/TBST.

H3f3b Antibody (C-Term) - Background

Variant histone H3 which replaces conventional H3 in a wide range of nucleosomes in active genes. Constitutes the predominant form of histone H3 in non-dividing cells and is incorporated into chromatin independently of DNA synthesis. Deposited at sites of nucleosomal displacement throughout transcribed genes, suggesting that it represents an epigenetic imprint of transcriptionally active chromatin. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling.

H3f3b Antibody (C-Term) - References

Hraba-Renevey S.,et al.Nucleic Acids Res. 17:2449-2461(1989).
Bramlage B.,et al.Differentiation 62:13-20(1997).
Lopez-Alanon D.M.,et al.DNA Cell Biol. 16:639-644(1997).
Carninci P.,et al.Science 309:1559-1563(2005).
Mancini P.,et al.J. Mol. Evol. 59:458-463(2004).