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DATA SHEET

Rabbit Anti-Histone H3 K27M Mutant/H3K27M [RM192]: RM0106

Intended Use: For Research Use Only

Description: Histone H3 is one of the five main histone proteins involved in the structure of chromatin in eukaryotic cells. Featuring a main globular domain and a long N-terminal tail, H3 is involved with the structure of the nucleosomes of the 'beads on a string' structure. The N-terminal tail of histone H3 protrudes from the globular nucleosome core and can undergo several different types of epigenetic modifications that influence cellular processes. These modifications include the covalent attachment of methyl or acetyl groups to lysine and arginine amino acids and the phosphorylation of serine or threonine. Histone variant H3 is typically enriched in active chromatin.

Specifications:

Clone:	RM192
Source:	Rabbit
Isotype:	IgG
Reactivity:	All
Immunogen:	A peptide corresponding to Histone H3 K27M mutant
Localization:	Nucleus
Formulation:	Protein A affinity Antibody in PBS pH7.4, containing BSA and ≤ 0.09% sodium azide (NaN3)
Storage:	Store at 2°- 8°C
Applications:	IHC, ELISA, ICC/IF, WB

Package:

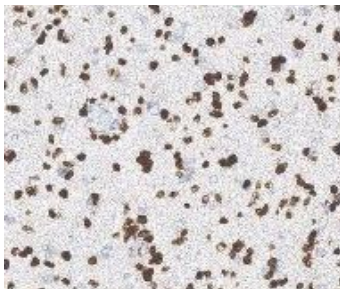
Description	Catalog No.	Size
Histone H3 K27M Mutant/H3K27M Concentrated	RM0106	1 ml

IHC Procedure*:

Brain with Histone H3 K27M mutant, 293T cells transfected with a DNA construct encoding Histone H3 K27M mutant

Concentrated Dilution:	100-1000
Pretreatment:	Tris EDTA pH9.0, 15 minutes using Pressure Cooker, or 30-60 minutes using water bath at 95°-99°C
Incubation Time and Temp:	30-60 minutes @ RT
Detection:	Refer to the detection system manual

* Result should be confirmed by an established diagnostic procedure.



FFPE human brain tumor stained with anti-Histone H3 K27M using DAB

References:

1. BRAF Fusion Analysis in Pilocytic Astrocytomas: KIAA1549-BRAF 15-9 Fusions Are More Frequent in the Midline Than Within the Cerebellum. Faulkner, et al. Journal of neuropathology and experimental neurology 74: 867-72, 2015.
2. Specific detection of methionine 27 mutation in histone 3 variants (H3K27M) in fixed tissue from high-grade astrocytomas. Bechet, D; et al. Acta neuropathologica 128:733-41, 2014.
3. A sensitive and specific histopathologic prognostic marker for H3F3A K27M mutant pediatric glioblastomas. Venneti, S; et al. Acta neuropathologica 128: 743-53, 2014.
4. Inhibition of PRC2 activity by a gain-of-function H3 mutation found in pediatric glioblastoma. Lewis, Peter W, et al. Science, 340: 857-61, 2013.

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