

Product Data Sheet

Purified anti-Histone H2A

Catalog # / Size: 619401 / 50 µl (5 Western blots)

619402 / 200 µl (20 Western blots)

Clone: Poly6194 Isotype: Rabbit IgG

Immunogen: Peptide mixture containing amino acid resides 1-5 and 81-96 of histone H2A

Reactivity: Human, others not tested

Preparation: The antibody was purified by antigen-affinity chromatography.

Formulation: This antibody is provided in phosphate-buffered solution, pH 7.2, containing

0.09% sodium azide and 50% glycerol.

Storage: Upon receipt, store frozen at -20° C.



Applications: WB - Quality tested

Recommended Usage: Each lot of this antibody is quality control tested by Western blotting. Western

blotting, suggested working dilution(s): Use 10 µl per 5 ml antibody dilution buffer for each mini-gel. It is recommended that the reagent be titrated for

optimal performance for each application.

Description: Histone H2A is a 14 kD nuclear protein that is a component of an octamer

containing pairs of each of four core histones (H2A, H2B, H3, H4). The core histones create nucleosome structure of chromosomal fiber in eukaryotes

and are dynamic in gene regulation. The histones exhibit cell

cycle-dependent transcriptional regulation. Histone H2A can be modified by phosphorylation and acetylation; phosphorylation of H2A correlates with mitotic chromosome condensation. Histone H2A interacts with Histones H3B, H3, and H4 as well as HIV-1 Tat peptides. The Poly6194 antibody has been

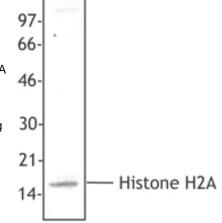
shown to be useful for the Western blotting of human histone 2A.

Antigen References: 1. McGhee JD, et al. 1980. Ann. Rev. Biochem. 49:1115.

3. Benedetti A, et al. 2002. Biochem. J. 367:505.

2. Nakamura TM, et al. 2004. Mol. Cell. Biol. 24:6215.

Related Products: Product Clone Application Purified anti-H2A.X-Phosphorylated (Ser139) 2F3 Purified anti-Histone H3 Poly6019 WB, IF Purified anti-Histone H4 Poly6020 Purified anti-H2A.X Poly6133 WB, IF HRP Donkey anti-rabbit IgG (minimal x-reactivity) Polv4064



Jurkat cell extract was resolved by electrophoresis, transferred to nitrocellulose, and probed with rabbit anti-Histone H2A polyclonal antibody. Proteins were visualized using a donkey anti-rabbit secondary conjugated to HRP and a chémiluminescence detection system.

IF, IHC, WB, ICFC

ELIŚA, IHC, WB



