SONY

Product Data Sheet

PerCP/Cy5.5 anti-H2A.X-Phosphorylated (Ser139)

Catalog # / Size: 3667065 / 25 tests
3667070 / 100 tests

Clone: 2F3

Isotype: Mouse IgG1, κ

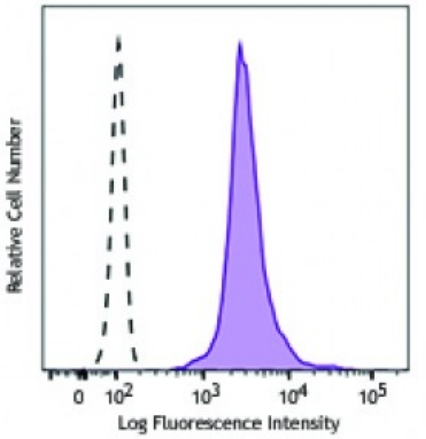
Immunogen: Modified peptide

Reactivity: Human,Mouse

Preparation: The antibody was purified by affinity chromatography and conjugated with PerCP/Cy5.5 under optimal conditions. The solution is free of unconjugated PerCP/Cy5.5 and unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).

Concentration: Lot-specific



Nocodozole treated HeLa cells (24 hours) were fixed and permeabilized with cold 70% ethanol, then intracellularly stained with anti-H2A.X Phospho (Ser139) (clone 2F3) PerCP/Cy5.5 (filled histogram) or mouse IgG1, κ PerCp/Cy5.5 isotype control (open

Applications:

Applications: Flow Cytometry

Recommended Usage: Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

* PerCP/Cy5.5 has a maximum absorption of 482 nm and a maximum emission of 690 nm.

Application Notes: **Additional reported applications (for the relevant formats of this clone) include:** immunohistochemistry on paraffin embedded sections2, immunofluorescence microscopy[3-9], Western blotting [10-12], and flow cytometry[1,13]. Clone 2F3 cross-reacts with mouse4.

Intracellular staining protocol for Anti-H2A.X-Phosphorylated (Ser139) Antibody for Flow Cytometry

1. Prepare 70% absolute ethanol. Chill solution by storing at -20°C.
2. Prepare cells of interest.
3. Wash 1X: resuspend with PBS, then pellet cells by centrifugation (250Xg, 5min)
4. Discard the supernatant and vortex to loosen cell pellet.
5. Add pre-cooled 70% ethanol drop by drop, while vortexing.
6. Incubate at -20°C for 60 minutes.
7. Wash 3X with BioLegend Cell Staining Buffer and resuspend the cells at 0.5-1 X 10^7 cells/ml in the cell staining buffer.
8. Perform immunofluorescent staining for flow cytometry.

Application References:
1. Jha JC, *et al.* 2013. *J. Virol.* 87:5255. (FC) PubMed
2. Akbay A, *et al.* 2008. *Am J Pathol.* 173:536. (IHC) PubMed

3. Mochizuki K, *et al.*2008.*J cell Sci.*121:2148. (IF) PubMed
4. Xiao R, *et al.* 2007. *Mol Cell Biol.*27:5393. (IF) PubMed
5. Rossi DJ, *et al.* 2007. *Nature*. 447:725. (IF) PubMed
6. Loidl J, *et al.* 2009. *Mol Cell Biol.* 20:2048. (IF) PubMed
7. Beels L, *et al.* 2009. *Circulation.* 120:1903. (IF) PubMed
8. Suzuki K, *et al.* 2010. *Nucleic Acids Res*. 38:e129. (IF) PubMed
9. Lukaszewicz A. 2010. *Chromasoma* Apr 27. [Epub ahead of print] (IF) PubMed
10. Yamada C, *et al.* 2010 *J. Biol. Chem.* 285:16693. (WB) PubMed
11. Bu Y, *et al.* 2010, *Biochem Biophys Res Commun.* 397:157. (WB) PubMed
12. Massignan T, *et al.* 2010. *J. Biol Chem.* 285:7752. (WB) PubMed
13. Banath JP, *et al.* 2010. *BMC Cancer* 10:4 (FC)
14. Zhang M., *et al.* 2011. *Cancer Res.* 23:7155. PubMed
15. Kuefner MA, *et al.* 2012. *Radiology* 264:59. PubMed
16. Yoshihara Y, *et al.* 2012. *Biochem Biophys Res Cmmun.* 421:57. PubMed
17. Titus S, *et al.* 2013. *Sci Transl Med.* 13:21. PubMed
18. Crown KN, *et al.* 2013. *G3.* 6:1927. PubMed
19. Schenkwein D, *et al.* 2013. *Nucleic Acids Res.* 41:e61. PubMed
20. Zhadanova NS, *et al.* 2014. *Mol Cell Biol.* 34:2786. PubMed
21. Horrell SA, *et al.* 2014. *Eukaryot Cell.* 13:1300. PubMed
22. Maya-Mendoza A, *et al.* 2015. *Mol Oncol.* 9:601. PubMed

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Description: H2A.X is a 14 kD basal histone and a member of the H2 histone family. This nuclear protein is synthesized in the G1 and S phase of the cell cycle and is known to be important for DNA repair and maintaining genomic stability and for recombination between immunoglobulin switch regions. H2A.X becomes phosphorylated on serine 139 after double-stranded DNA breaks. Phosphorylated H2A.X promotes DNA repair and maintains genomic stability. The 2F3 monoclonal antibody reacts with phosphorylated human H2A.X (Ser139) and has been shown to be useful for Western blotting, immunofluorescence and flow cytometry.

Antigen References:

1. Mannironi C, *et al.*1989. *Nucleic Acids Res.* 17:9113.
2. Celeste A, *et al.* 2002. *Science* 296:922.
3. Bassing CH, *et al.* 2002. *Proc. Natl. Acad. Sci. USA* 99:8173.
4. Reina-San-Mar