

#99891 Store at -20°C

Homologous Recombination (HR) DNA Repair Antibody Sampler Kit

1 Kit (9 x 20 microliters)



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Product Includes	Product #	Quantity	Mol. Wt	Isotype/Source
ATM (D2E2) Rabbit mAb	2873	20 µl	350 kDa	Rabbit IgG
Phospho-ATM (Ser1981) (D25E5) Rabbit mAb	13050	20 µl	350 kDa	Rabbit IgG
Rad51 (D4B10) Rabbit mAb	8875	20 µl	37 kDa	Rabbit IgG
BRCA1 (A8X9F) Rabbit mAb	14823	20 µl	220 kDa	Rabbit IgG
BRCA2 (D9S6V) Rabbit mAb	10741	20 µl	380 kDa	Rabbit IgG
Rad54 (D4W3Z) Rabbit mAb	15016	20 µl	84 kDa	Rabbit IgG
p95/NBS1 (D6J5I) Rabbit mAb	14956	20 µl	95 kDa	Rabbit IgG
Phospho-p95/NBS1 (Ser343) Antibody	3001	20 µl	95 kDa	Rabbit
CtIP (D76F7) Rabbit mAb	9201	20 µl	110 kDa	Rabbit IgG
Anti-rabbit IgG, HRP-linked Antibody	7074	100 µl		Goat

Please visit cellsignal.com for individual component applications, species cross-reactivity, dilutions, protocols, and additional product information.

Description

The Homologous Recombination (HR) DNA Repair Antibody Sampler Kit provides an economical means of detecting proteins involved in HR DNA repair. The kit includes enough antibodies to perform two western blot experiments with each primary antibody.

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. *Do not aliquot the antibody.*

Background

DNA double-strand breaks (DSBs) are potentially hazardous lesions that can be induced by ionizing radiation (IR), radiomimetic chemicals, or DNA replication inhibitors. Cells recognize and repair DSBs via two distinct but partly overlapping signaling pathways, non-homologous end joining (NHEJ) and homologous recombination (HR). DSBs that arise during S or G2 phase are repaired via HR, using the replicated sister chromatid as a repair template (1). Activation of ATM by autophosphorylation on Ser1981 occurs in response to exposed DNA DSBs. ATM regulates various responses to DNA damage, including regulation of HR factors (2). Rad51 recombinase polymerizes and forms a filament along single-stranded DNA, mediating HR with the help of auxiliary proteins, including Rad54 and BRCA2 (3). BRCA2 has been shown to be required for localization of Rad51 to sites of DSBs, and cells lacking BRCA1 and BRCA2 cannot repair DSBs through HR (4). NBS1 is critical for HR, and requires CDK-dependent association with CtIP and subsequent phosphorylation by ATM at Ser278 and Ser343 (5-6).

Background References

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3. Sung, P. et al. (2003) *J Biol Chem* 278, 42729-32.
4. Tutt, A. and Ashworth, A. (2002) *Trends Mol Med* 8, 571-6.
5. Wang, H. et al. (2013) *PLoS Genet* 9, e1003277.
6. Wen, J. et al. (2013) *Oncogene* 32, 4448-56.

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