## **Resource Summary Report**

Generated by FDI Lab - SciCrunch.org on Apr 27, 2024

# Phospho-Estrogen Receptor alpha (Ser118) (16J4) Mouse mAb

RRID:AB\_331289 Type: Antibody

#### **Proper Citation**

(Cell Signaling Technology Cat# 2511, RRID:AB\_331289)

#### **Antibody Information**

**URL:** http://antibodyregistry.org/AB\_331289

Proper Citation: (Cell Signaling Technology Cat# 2511, RRID:AB\_331289)

Target Antigen: Phospho-Estrogen Receptor alpha (Ser118) (16J4) Mouse mAb

**Host Organism:** mouse

Clonality: monoclonal

Comments: Applications: W, IHC-P. Consolidation on 10/2018: AB\_10079292,

AB\_10831843, AB\_331289.

Antibody Name: Phospho-Estrogen Receptor alpha (Ser118) (16J4) Mouse mAb

**Description:** This monoclonal targets Phospho-Estrogen Receptor alpha (Ser118) (16J4)

Mouse mAb

Target Organism: human

Antibody ID: AB\_331289

Vendor: Cell Signaling Technology

Catalog Number: 2511

### Ratings and Alerts

No rating or validation information has been found for Phospho-Estrogen Receptor alpha (Ser118) (16J4) Mouse mAb.

No alerts have been found for Phospho-Estrogen Receptor alpha (Ser118) (16J4) Mouse mAb.

#### Data and Source Information

**Source:** Antibody Registry

#### **Usage and Citation Metrics**

We found 5 mentions in open access literature.

Listed below are recent publications. The full list is available at FDI Lab - SciCrunch.org.

Ng ASN, et al. (2022) AKTIP loss is enriched in ER?-positive breast cancer for tumorigenesis and confers endocrine resistance. Cell reports, 41(11), 111821.

Vydra N, et al. (2021) Heat shock factor 1 (HSF1) cooperates with estrogen receptor ? (ER?) in the regulation of estrogen action in breast cancer cells. eLife, 10.

He YH, et al. (2021) ER? determines the chemo-resistant function of mutant p53 involving the switch between lincRNA-p21 and DDB2 expressions. Molecular therapy. Nucleic acids, 25, 536.

Guan J, et al. (2019) Therapeutic Ligands Antagonize Estrogen Receptor Function by Impairing Its Mobility. Cell, 178(4), 949.

Hinohara K, et al. (2018) KDM5 Histone Demethylase Activity Links Cellular Transcriptomic Heterogeneity to Therapeutic Resistance. Cancer cell, 34(6), 939.