Resource Summary Report

Generated by FDI Lab - SciCrunch.org on May 24, 2025

BRM (D9E8B) XP® Rabbit mAb

RRID:AB_2797783
Type: Antibody

Proper Citation

(Cell Signaling Technology Cat# 11966, RRID:AB_2797783)

Antibody Information

URL: http://antibodyregistry.org/AB_2797783

Proper Citation: (Cell Signaling Technology Cat# 11966, RRID:AB_2797783)

Target Antigen: SMARCA2

Host Organism: rabbit

Clonality: monoclonal

Comments: Applications: W, IP, IHC-P, IF-IC, ChIP

Antibody Name: BRM (D9E8B) XP® Rabbit mAb

Description: This monoclonal targets SMARCA2

Target Organism: h, mk

Clone ID: Clone D9E8B

Antibody ID: AB_2797783

Vendor: Cell Signaling Technology

Catalog Number: 11966

Record Creation Time: 20241017T001355+0000

Record Last Update: 20241017T015312+0000

Ratings and Alerts

No rating or validation information has been found for BRM (D9E8B) XP® Rabbit mAb.

No alerts have been found for BRM (D9E8B) XP® Rabbit mAb.

Data and Source Information

Source: Antibody Registry

Usage and Citation Metrics

We found 8 mentions in open access literature.

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

Sasaki M, et al. (2025) Efficacy of CBP/p300 Dual Inhibitors against Derepression of KREMEN2 in cBAF-Deficient Cancers. Cancer research communications, 5(1), 24.

Ng J, et al. (2024) Molecular and Pathologic Characterization of YAP1-Expressing Small Cell Lung Cancer Cell Lines Leads to Reclassification as SMARCA4-Deficient Malignancies. Clinical cancer research : an official journal of the American Association for Cancer Research, OF1.

Duplaquet L, et al. (2024) Mammalian SWI/SNF complex activity regulates POU2F3 and constitutes a targetable dependency in small cell lung cancer. Cancer cell, 42(8), 1352.

Bhat KP, et al. (2024) CRISPR activation screens identify the SWI/SNF ATPases as suppressors of ferroptosis. Cell reports, 43(6), 114345.

Bolomsky A, et al. (2024) IRF4 requires ARID1A to establish plasma cell identity in multiple myeloma. Cancer cell, 42(7), 1185.

Liu W, et al. (2023) RNF138 inhibits late inflammatory gene transcription through degradation of SMARCC1 of the SWI/SNF complex. Cell reports, 42(2), 112097.

Kofink C, et al. (2022) A selective and orally bioavailable VHL-recruiting PROTAC achieves SMARCA2 degradation in vivo. Nature communications, 13(1), 5969.

Lazar JE, et al. (2020) Global Regulatory DNA Potentiation by SMARCA4 Propagates to Selective Gene Expression Programs via Domain-Level Remodeling. Cell reports, 31(8), 107676.