

Resource Summary Report

Generated by [NIF](#) on Sep 15, 2026

Ki67 Antibody (SP6)

RRID:AB_2314700

Type: Antibody

Proper Citation

Thermo Fisher Scientific Cat# MA1-90584, RRID:AB_2314700

Antibody Information

URL: http://antibodyregistry.org/AB_2314700

Proper Citation: Thermo Fisher Scientific Cat# MA1-90584, RRID:AB_2314700

Target Antigen: Ki67

Host Organism: rabbit

Clonality: monoclonal

Comments: Discontinued; Applications: Immunohistochemistry (Paraffin) (1:100), Western Blot (1:200); Reactive Species: Human, Rat

Antibody Name: Ki67 Antibody (SP6)

Description: This monoclonal targets Ki67

Target Organism: human

Clone ID: SP6

Defining Citation: [PMID:22315167](#)

Antibody ID: AB_2314700

Vendor: Thermo Fisher Scientific

Catalog Number: MA1-90584

Record Creation Time: 20231110T042044+0000

Record Last Update: 20260829T234243+0000

Ratings and Alerts

No rating or validation information has been found for Ki67 Antibody (SP6).

Warning: Discontinued at Thermo Fisher Scientific
Discontinued; Applications: Immunohistochemistry (Paraffin) (1:100), Western Blot (1:200); Reactive Species: Human, Rat

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 [NIF](#) .

Lee R, et al. (2022) Synthetic Essentiality of Tryptophan 2,3-Dioxygenase 2 in APC-Mutated Colorectal Cancer. *Cancer discovery*, 12(7), 1702.

Duan X, et al. (2021) An airway organoid-based screen identifies a role for the HIF1 α -glycolysis axis in SARS-CoV-2 infection. *Cell reports*, 37(6), 109920.

Xu R, et al. (2019) OLIG2 Drives Abnormal Neurodevelopmental Phenotypes in Human iPSC-Based Organoid and Chimeric Mouse Models of Down Syndrome. *Cell stem cell*, 24(6), 908.

Momcilovic M, et al. (2018) The GSK3 Signaling Axis Regulates Adaptive Glutamine Metabolism in Lung Squamous Cell Carcinoma. *Cancer cell*, 33(5), 905.

Dubail J, et al. (2016) Impaired ADAMTS9 secretion: A potential mechanism for eye defects in Peters Plus Syndrome. *Scientific reports*, 6, 33974.