

Resource Summary Report

Generated by [RRID](#) on Aug 7, 2026

Phospho-p53 (Ser392) Antibody

RRID:AB_331462

Type: Antibody

Proper Citation

Cell Signaling Technology Cat# 9281, RRID:AB_331462

Antibody Information

URL: http://antibodyregistry.org/AB_331462

Proper Citation: Cell Signaling Technology Cat# 9281, RRID:AB_331462

Target Antigen: Phospho-p53 (Ser392)

Host Organism: rabbit

Clonality: polyclonal

Comments: Applications: W. Consolidation on 10/2018: AB_10079167, AB_10828088, AB_331462.

Antibody Name: Phospho-p53 (Ser392) Antibody

Description: This polyclonal targets Phospho-p53 (Ser392)

Target Organism: h, m, mouse, human

Antibody ID: AB_331462

Vendor: Cell Signaling Technology

Catalog Number: 9281

Record Creation Time: 20250820T040337+0000

Record Last Update: 20250820T175551+0000

Ratings and Alerts

No rating or validation information has been found for Phospho-p53 (Ser392) Antibody.

No alerts have been found for Phospho-p53 (Ser392) Antibody.

Data and Source Information

Source: [Antibody Registry](#)

Usage and Citation Metrics

We found 6 mentions in open access literature.

Listed below are recent publications. The full list is available at [RRID](#) .

Yan X, et al. (2026) Activation of T Cell-Intrinsic p53 by Acetylation Elicits Antitumor Immunity to Boost Cancer Immunotherapy. Cancer discovery, 16(1), 155.

Xiong S, et al. (2025) Dependence on Mdm2 for Mdm4 inhibition of p53 activity. Cancer letters, 621, 217622.

Li H, et al. (2023) Transgelin Promotes Glioblastoma Stem Cell Hypoxic Responses and Maintenance Through p53 Acetylation. Advanced science (Weinheim, Baden-Wurttemberg, Germany), e2305620.

Darr J, et al. (2020) iTAG-RNA Isolates Cell-Specific Transcriptional Responses to Environmental Stimuli and Identifies an RNA-Based Endocrine Axis. Cell reports, 30(9), 3183.

Gong L, et al. (2019) p53 Protects Cells from Death at the Heatstroke Threshold Temperature. Cell reports, 29(11), 3693.

Brady OA, et al. (2018) The transcription factors TFE3 and TFEB amplify p53 dependent transcriptional programs in response to DNA damage. eLife, 7.