

Resource Summary Report

Generated by [RRID](#) on Aug 2, 2026

CD79B (D7V2F) Rabbit mAb

RRID:AB_2800254

Type: Antibody

Proper Citation

Cell Signaling Technology Cat# 96024, RRID:AB_2800254

Antibody Information

URL: http://antibodyregistry.org/AB_2800254

Proper Citation: Cell Signaling Technology Cat# 96024, RRID:AB_2800254

Target Antigen: Ig-beta

Host Organism: rabbit

Clonality: monoclonal

Comments: Applications: W, IP, IHC-Bond, IHC-P

Antibody Name: CD79B (D7V2F) Rabbit mAb

Description: This monoclonal targets Ig-beta

Target Organism: h, m

Clone ID: Clone D7V2F

Antibody ID: AB_2800254

Vendor: Cell Signaling Technology

Catalog Number: 96024

Record Creation Time: 20250820T010420+0000

Record Last Update: 20250820T095716+0000

Ratings and Alerts

No rating or validation information has been found for CD79B (D7V2F) Rabbit mAb.

No alerts have been found for CD79B (D7V2F) Rabbit mAb.

Data and Source Information

Source: [Antibody Registry](#)

Usage and Citation Metrics

We found 7 mentions in open access literature.

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

Geng L, et al. (2025) Systematic profiling reveals betaine as an exercise mimetic for geroprotection. *Cell*.

Stahl D, et al. (2025) CSF1R+ myeloid-monocytic cells drive CAR-T cell resistance in aggressive B cell lymphoma. *Cancer cell*, 43(8), 1476.

Lei J, et al. (2025) Senescence-resistant human mesenchymal progenitor cells counter aging in primates. *Cell*, 188(18), 5039.

Choi J, et al. (2024) Molecular targets of glucocorticoids that elucidate their therapeutic efficacy in aggressive lymphomas. *Cancer cell*, 42(5), 833.

Ma S, et al. (2024) Spatial transcriptomic landscape unveils immunoglobulin-associated senescence as a hallmark of aging. *Cell*, 187(24), 7025.

Scheich S, et al. (2023) Targeting N-linked Glycosylation for the Therapy of Aggressive Lymphomas. *Cancer discovery*, 13(8), 1862.

Sevdali E, et al. (2022) BAFFR activates PI3K/AKT signaling in human naive but not in switched memory B cells through direct interactions with B cell antigen receptors. *Cell reports*, 39(13), 111019.