

Resource Summary Report

Generated by [dkNET](#) on Sep 3, 2026

Phospho-eNOS (Ser1177) (C9C3) Rabbit mAb

RRID:AB_823493

Type: Antibody

Proper Citation

Cell Signaling Technology Cat# 9570, RRID:AB_823493

Antibody Information

URL: http://antibodyregistry.org/AB_823493

Proper Citation: Cell Signaling Technology Cat# 9570, RRID:AB_823493

Target Antigen: Phospho-eNOS (Ser1177)

Host Organism: rabbit

Clonality: monoclonal

Comments: Applications: W, IP, E-P. Consolidation: AB_2298588.

Antibody Name: Phospho-eNOS (Ser1177) (C9C3) Rabbit mAb

Description: This monoclonal targets Phospho-eNOS (Ser1177)

Target Organism: Human, Bovine, Pig

Clone ID: C9C3

Antibody ID: AB_823493

Vendor: Cell Signaling Technology

Catalog Number: 9570

Alternative Catalog Numbers: 9570S, 9570L

Record Creation Time: 20231110T043212+0000

Record Last Update: 20260829T011423+0000

Ratings and Alerts

No rating or validation information has been found for Phospho-eNOS (Ser1177) (C9C3) Rabbit mAb.

No alerts have been found for Phospho-eNOS (Ser1177) (C9C3) Rabbit 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 [dkNET](#) .

Efentakis P, et al. (2024) Implications and hidden toxicity of cardiometabolic syndrome and early-stage heart failure in carfilzomib-induced cardiotoxicity. British journal of pharmacology, 181(16), 2964.

Hu Z, et al. (2024) ROUX-en-Y gastric bypass surgery improves metabolic syndrome-related erectile dysfunction in mice via the IRS-1/PI3K/AKT/eNOS pathway. Sexual medicine, 12(2), qfae029.

Zhang Q, et al. (2020) Chloroquine differentially modulates coronary vasodilation in control and diabetic mice. British journal of pharmacology, 177(2), 314.

Mori Y, et al. (2018) Glucose-Dependent Insulinotropic Polypeptide Suppresses Peripheral Arterial Remodeling in Male Mice. Endocrinology, 159(7), 2717.

Passaro D, et al. (2017) Increased Vascular Permeability in the Bone Marrow Microenvironment Contributes to Disease Progression and Drug Response in Acute Myeloid Leukemia. Cancer cell, 32(3), 324.