

DNAxPAb

Hard-to-Find
Antibody

SHMT1 DNAxPab

Catalog # H00006470-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human SHMT1 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MTMPVNGAHKDADLWSSHDKMLAQPLKDSDEVYNIKKESNRQRVGLELIASENFASRAVLEAL GSCLNNKYSEGYPGQRYGGTEFIDELETLCQKRALQAYKLDPQCWGVNVQPYSGSPANFAVYT ALVEPHGRIMGLDLPDGGHLTHGFMTDKKKISATSIFFESMPYKVNPD TGYN YDQLEENARLFHP KLIAGTSCYSRNLEYARLRKIADENGAYLMADMAHISGLVAAGVVPSPFEHCHVTTTTHKTLRGC RAGMIFYRKGVKSVDPKTGKEILYNLESLINSAVFPGLQGGPHNHAIGVAVALKQAMTLEFKVYQ HQVVANCRALSEALTELGYKMTGGSDNHLILVDLRSGTDGGRAEKVLEACSIACNKNTCPGDR SALRPSGLRLGTPALTSRGLLEKDFQKVAHFHHRGIELTLQIQSDTGVRATLKEFKERLAGDKYQAA VQALREEVESFASLFPLPLPDF
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

Gene Info — SHMT1

Entrez GeneID [6470](#)

GeneBank Accession# [NM_004169.3](#)

Protein Accession# [NP_004160.3](#)

Gene Name SHMT1

Gene Alias CSHMT, MGC15229, MGC24556, SHMT

Gene Description serine hydroxymethyltransferase 1 (soluble)

Omim ID [182144](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes the cellular form of serine hydroxymethyltransferase, a pyridoxal phosphate-containing enzyme that catalyzes the reversible conversion of serine and tetrahydrofolate to glycine and 5,10-methylene tetrahydrofolate. This reaction provides one carbon units for synthesis of methionine, thymidylate, and purines in the cytoplasm. This gene is located within the Smith-Magenis syndrome region on chromosome 17. Alternative splicing of this gene results in 2 transcript variants encoding 2 different isoforms. Additional transcript variants have been described, but their biological validity has not been determined. [provided by RefSeq]

Other Designations 14 kDa protein|cytoplasmic serine hydroxymethyltransferase

Pathway

- [Cyanoamino acid metabolism](#)
- [Glycine](#)
- [Metabolic pathways](#)
- [Methane metabolism](#)
- [One carbon pool by folate](#)

Disease

- [Adenocarcinoma](#)
- [Adenoma](#)
- [Aortic Aneurysm](#)
- [Arthritis](#)
- [Atherosclerosis](#)
- [Autistic Disorder](#)
- [Bone Neoplasms](#)
- [Brain Ischemia](#)
- [Breast cancer](#)
- [Breast Neoplasms](#)
- [Carcinoma](#)
- [Cardiovascular Diseases](#)
- [Cleft Lip](#)
- [Cleft Palate](#)
- [Colorectal Neoplasms](#)
- [Disease Models](#)
- [Ductus Arteriosus](#)
- [Epilepsy](#)
- [Esophageal Neoplasms](#)
- [Folic Acid Deficiency](#)
- [Gastrointestinal Diseases](#)
- [Genetic Predisposition to Disease](#)
- [Head and Neck Neoplasms](#)
- [Heart Defects](#)

- [Hematologic Diseases](#)
- [Hyperhomocysteinemia](#)
- [Hypoplastic Left Heart Syndrome](#)
- [Infant](#)
- [Kidney Diseases](#)
- [Leukemia](#)
- [Lung Neoplasms](#)
- [Lymphoma](#)
- [Mental Disorders](#)
- [Microsatellite Instability](#)
- [Migraine Disorders](#)
- [Migraine with Aura](#)
- [Neoplasms](#)
- [Neural Tube Defects](#)
- [Osteosarcoma](#)
- [Ovarian cancer](#)
- [Ovarian Neoplasms](#)
- [Pregnancy Complications](#)
- [Premature Birth](#)
- [Prostate cancer](#)
- [Prostatic Neoplasms](#)
- [Schizophrenia](#)
- [Spinal Dysraphism](#)
- [Startle Reaction](#)
- [Stomach Neoplasms](#)

- [Stroke](#)
- [Urinary Bladder Neoplasms](#)
- [Vitamin B 12 Deficiency](#)
- [Werner syndrome](#)