

TDG mouse monoclonal antibody (hybridoma)

Catalog # H00006996-M

Size Up to 5 Clones

Specification

Product Description	Mouse monoclonal antibody raised against a full-length recombinant TDG.
Immunogen	TDG (AAH37557.1, 1 a.a. ~ 410 a.a) full-length recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.
Sequence	MEAENAGSYSLQQAQAFYTFPFQQLMAEAPNMAVVNEQQMPVEVPAPAPAQEPVQEAPKGRKRKPRTEPKQPVEPKPVESKKSGKSAKSKEKQEKITDTFKVKRKVDRFNGVSEAELLTKTLPDILTFNLDVIIGINPGLMAAYKGHHYPGPGNHFWKCLFMSGLSEVQLNHMDHTLPGKYGIGFTNMVERTTPGSKDLSSKEFREGGRILVQKLQKYQPRIAVFNGKCIYEIFSKEVFGVKVKNLEFGLQPHKIPDTETLCYGMPSSSARCAQFPRAQDKVHYYIKLDLRDQLKGIERNMDVQEVQYTFDLQLAQEDAKKMAVKEEKYDPGYEAAAYGGAYGENPCSSSEPCGFSSNGLIESVELRGESAFSGIPNGQWMTQSFTDQIPSFNSHCGTQEQEEESHA
Host	Mouse
Reactivity	Human
Quality Control Testing	Antibody reactivity and specificity confirmed by ELISA and Western Blot.
Deliverables	Up to 5 positive hybridoma clones will be delivered to customer in the cryotube format.
Note	Customer should check the viability of the hybridomas within one month from the date of receipt. Fee -for-service of long term hybridoma storage can be performed upon customer's request.

Applications

- Western Blot (Transfected lysate)

[Protocol Download](#)

- Western Blot (Recombinant protein)

[Protocol Download](#)

- ELISA

Gene Info — TDG

Entrez GeneID [6996](#)

GeneBank Accession# [BC037557.1](#)

Protein Accession# [AAH37557.1](#)

Gene Name TDG

Gene Alias -

Gene Description thymine-DNA glycosylase

Omim ID [601423](#)

Gene Ontology [Hyperlink](#)

Gene Summary

The protein encoded by this gene belongs to the TDG/mug DNA glycosylase family. Thymine-DNA glycosylase (TDG) removes thymine moieties from G/T mismatches by hydrolyzing the carbon-nitrogen bond between the sugar-phosphate backbone of DNA and the mispaired thymine. With lower activity, this enzyme also removes thymine from C/T and T/T mispairings. TDG can also remove uracil and 5-bromouracil from mispairings with guanine. This enzyme plays a central role in cellular defense against genetic mutation caused by the spontaneous deamination of 5-methylcytosine and cytosine. This gene may have a pseudogene in the p arm of chromosome 12. [provided by RefSeq]

Other Designations G/T mismatch-specific thymine DNA glycosylase

Pathway

- [Base excision repair](#)

Disease

- [Breast cancer](#)
- [Carcinoma](#)
- [DNA Damage](#)
- [Genetic Predisposition to Disease](#)
- [Graft vs Host Disease](#)

- [Kidney Failure](#)
- [Lung Neoplasms](#)
- [Meningeal Neoplasms](#)
- [Meningioma](#)
- [Multiple Sclerosis](#)
- [Tobacco Use Disorder](#)
- [Urinary Bladder Neoplasms](#)
- [Werner syndrome](#)