

Thymidylate Synthase (D26G11) Rabbit



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IP	<b>Reactivity:</b> H M R Hm Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 30	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P04818	Entrez-Gene Id: 7298
Product Usage Information		<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:100	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Thymidylate Synthase (D26G11) Rabbit mAb detects endogenous levels of total Thymidylate Synthase protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to central residues of human Thymidylate Synthase protein.				
Background		The methylation of deoxyuridine monophosphate (dUMP) to deoxythymidine monophosphate (dTMP) is an essential step in the formation of thymine nucleotides (1,2, reviewed in 3). This process is catalyzed by thymidylate synthase (TS or TYMS), a homodimer composed of two 30 kDa subunits. TS is an intracellular enzyme that provides the sole <i>de novo</i> source of thymidylate, making it a required enzyme in DNA biosynthesis with activity highest in proliferating cells (1). Being the exclusive source of dTMP, investigators have concluded that TS is also an important target for anticancer agents such as 5-fluorouracil (5-FU) (1-5). 5-FU acts as a TS inhibitor and is active against solid tumors such as colon, breast, head, and neck. Research studies have demonstrated that patients with metastases expressing lower levels of TS have a higher response rate to treatment with 5-FU than patients with tumors that have increased levels of TS (5). Researchers continue to investigate TS expression in different types of cancers (6-10).				
Background References		1. Johnston, P.G. et al. (1991) <i>Cancer Res</i> 51, 6668-76. 2. Aschele, C. et al. (2002) <i>Ann Oncol</i> 13, 1882-92. 3. Jackman, A.L. and Calvert, A.H. (1995) <i>Ann Oncol</i> 6, 871-81. 4. Van Triest, B. et al. (2000) <i>J Histochem Cytochem</i> 48, 755-60. 5. Johnston, P.G. et al. (1994) <i>J Clin Oncol</i> 12, 2640-7. 6. Kwon, H.C. et al. (2007) <i>Ann Oncol</i> 18, 504-9. 7. Allegra, C.J. et al. (2002) <i>J Clin Oncol</i> 20, 1735-43. 8. Allegra, C.J. et al. (2003) <i>J Clin Oncol</i> 21, 241-50. 9. Tsourouflis, G. et al. (2008) <i>Dig Dis Sci</i> 53, 1289-96. 10. Kim, S.H. et al. (2009) <i>Am J Clin Oncol</i> 32, 38-43.				
Species React	ivity	Species reactivity is de	etermined by testir	g in at least one approve	ed application (e.g.,	western blot).
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**Western Blot Buffer** 

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** 

W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key

H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey

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