DHFR Antibody	T C	Cell Signaling TECHNOLOGY®	
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#	3 Trask Lane Danvers Mas	sachusetts 01923 USA	
For Research Use Only. Not for Use in Diagnostic Procedures.			

Applications: W	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 22	Source/Isotype: Rabbit	UniProt ID: #P00374	Entrez-Gene Id 1719	
Product Usage Information		Application Western Blotting			Dilution 1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. <i>Do not aliquot the antibody.</i>					
Specificity/Sensit	ivity	DHFR Antibody recognizes endogenous levels of total DHFR protein. This antibody does not cross-read with DHFRL1 protein.					
Source / Purificat	ion	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human DHFR protein. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		Dihydrofolate reductase (DHFR) catalyzes tetrahydrofolate regeneration through the reduction of dihydrofolate using NADPH as a cofactor (1). As a key enzyme in folate metabolism, DHFR is ubiquitously expressed in the mitochondria, and is essential for the synthesis of purines, pyrimidines, and some amino acids (2). DHFR is capable of translational autoregulation by binding within the coding region of its own mRNA sequence to repress cellular DHFR protein levels (3). Mutations in the DHFR gene are known to cause inborn errors of folate metabolism resulting in megaloblastic anemia, pancytopenia, and severe cerebral folate deficiency (4). Because tetrahydrofolate is essential for DNA synthesis, cell growth, and proliferation, DHFR is a target of chemotherapeutic agents (e.g., methotrexate and pemetrexed) used in the treatment of many cancer types (5). Increased expression of DHFR has also been identified as a potential mechanism for tumor resistance to methotrexate, and therefore has been utilized as a clinical biomarker to predict patient responsiveness to folate antagonists (6,7).					
Background Refe	rences	1. Davies, J.F. et al. (1990) <i>Biochemistry</i> 29, 9467-79. 2. Schnell, J.R. et al. (2004) <i>Annu Rev Biophys Biomol Struct</i> 33, 119-40. 3. Ercikan-Abali, E.A. et al. (1997) <i>Biochemistry</i> 36, 12317-22. 4. Banka, S. et al. (2011) <i>Am J Hum Genet</i> 88, 216-25. 5. Vander Heiden, M.G. and DeBerardinis, R.J. (2017) <i>Cell</i> 168, 657-69. 6. Nakano, M. et al. (2017) <i>J Biol Chem</i> 292, 4873-84. 7. Organista-Nava, J. et al. (2018) <i>Oncol Lett</i> 15, 8405-11.					
Species Reactivit	y	Species reactivity is d	letermined by testin	g in at least one approve	ed application (e.g.,	western blot).	
Western Blot Buf	fer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.					
Applications Key		W: Western Blotting					
Cross-Reactivity I	۲ey	H: Human M: Mouse R: Rat Mk: Monkey					
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