## 5648

**Limited Uses** 

force or effect.

## CA9 (D10C10) Rabbit mAb



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

<b>Applications:</b> W, IHC-P	Reactivity:	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 35-58	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q16790	Entrez-Gene Id: 768
Product Usage Information		<b>Application</b> Western Blotting Immunohistochemistry (Paraffin)			<b>Dilution</b> 1:1000 1:50 - 1:200	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
		For a carrier free (BSA and azide free) version of this product see product #83980.				
Specificity/Sensitivity		CA9 (D10C10) Rabbit mAb recognizes endogenous level of total CA9 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Asp361 of human CA9 protein.				
Background		Carbonic anhydrases (CA) are a family of ancient zinc metalloenzymes found in almost all living organisms. All CA can be divided into 3 distinct classes ( $\alpha$ , $\beta$ , and $\gamma$ ) that evolved independently and have no significant homology in sequence and overall folding. All functional CA catalyze the reversible hydration of $CO_2$ into $HCO_3$ and $H^+$ and contain a zinc atom in the active sites essential for catalysis. There are many isoforms of CA in mammals and they all belong to the $\alpha$ class (1,2). CA9 is a member of alpha class, a plasma membrane protein with the catalytic domain in the extracellular space. Its expression is restricted to very few normal tissues (mainly the gastrointestinal tract) (2). CA9 expression is strongly induced by hypoxia and down-regulated by the wildtype von Hippel–Lindau (VHL) tumor suppressor protein. CA9 expression is increased in many types of tumor, especially in solid hypoxic tumors with a poor responsiveness to the conventional radio-and/or chemotherapies; CA9 is considered as a tumor hypoxia marker and a promising target for cancer therapeutic intervention (3-5).				
Background References		<ol> <li>Smith, K.S. et al. (1999) Proc Natl Acad Sci USA 96, 15184-9.</li> <li>Tripp, B.C. et al. (2001) J Biol Chem 276, 48615-8.</li> <li>Potter, C.P. and Harris, A.L. (2003) Br J Cancer 89, 2-7.</li> <li>Winum, J.Y. et al. (2009) Anticancer Agents Med Chem 9, 693-702.</li> <li>De Simone, G. and Supuran, C.T. (2010) Biochim Biophys Acta 1804, 404-9.</li> </ol>				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
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 IHC-P: Immunohistochemistry (Paraffin)				
Cross-Reactivity Key		H: Human				
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