e at -20C	CYP11A1 (D8F4F) Rabbit mAb		Cell Signaling		
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## For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: WB, IP, IF-IC	Reactivity: H M R	Sensitivity: Endogenous	<b>MW (kDa):</b> 50	Source/Isotype: Rabbit IgG	UniProt ID: #P05108	Entrez-Gene Id: 1583		
Product Usage	Aŗ	plication		Dilution				
Information	W	Western Blotting			1:1000			
	Im	munoprecipitation			1:50	0		
	Im	Immunofluorescence (Immunocytochemistry)			1:400 - 1:800			
Storage	Sup 0.0	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. <i>Do not aliquot the antibody.</i>						
Specificity / Sensitivity		CYP11A1 (D8F4F) Rabbit mAb recognizes endogenous levels of total CYP11A1 protein.						
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human CYP11A1 protein.						
Background		In steroidogenic tissues, such as the adrenal cortex, testis, ovary, and placenta, all steroids are synthesized from the common precursor cholesterol. Two families of steroidogenic enzymes, cytochrome P450 hydroxylase enzymes (CYP) and hydroxylaseroid dehydrogenases (HSD), catalyze the production of most steroidogenic <i>CYP</i> gene family (1). The cytochrome P450scc (cholesterol side-chain cleavage enzyme) encoded by <i>CYP11A1</i> catalyzes the first and rate-limiting step in steroidogenesis, conversion of cholesterol into pregnenolone (2).						
		adrenal cortex, testis, and ovary, CYP11A1 expression is regulated by the cAMP-PKA pathway (6), and the transcription factor SF1/NR5A1 has been shown to play a central role in mediating the cAMP signal on the <i>CYP11A1</i> promoter within steroidogeneic cells of the adrenal cortex and gonads (7). Defects in CYP11A1 are the cause of adrenal insufficiency congenital with 46, XY sex reversal (AICSR), which is a rare disorder that can present as acute adrenal insufficiency in infancy or childhood (8,9).						
Background Refere	ences 1. N 2. F 3. H 4. H 5. H 6. H 7. V 8. T 9. K	lelson, D.R. et al. (19 Richards, J.S. et al. (19 Ianukoglu, I. and Jef Ianukoglu, I. et al. (1 Ianukoglu, I. et al. (1 Ianukoglu, I. et al. (1 Iu, M.C. et al. (1991) Vatanabe, N. et al. (1 Gajima, T. et al. (2001 Catsumata, N. et al. (	993) DNA Cell Bi 1987) Recent Pro icoate, C.R. (198 981) J Biol Chen 981) J Biol Chen 981) J Biol Chen 0 Biochem J 274 1994) Eur J Biocl 1994) Eur J Biocl 1 J Clin Endocrin 2002) J Clin End	ol 12, 1-51. bg Horm Res 43, 231-76 0) J Biol Chem 255, 305 n 256, 4329-35. n 256, 4321-8. ( Pt 3), 813-7. nem 222, 825-34. ol Metab 86, 3820-5. iocrinol Metab 87, 3808-	5. 7-61. 13.			
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).						
Western Blot Buffer	r IMP 0.1%	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		WB: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry)						
Cross-Reactivity Ke	ey H:h X:X GP:	H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected						

Trademarks and Patents CYP11A1 (D8F4F) Rabbit mAb (#14217) Datasheet Without Images Cell Signaling Technology

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