



# HCCS Monoclonal Antibody

<b>Catalog No</b>	YP-mAb-02651
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Mouse;Monkey
<b>Applications</b>	WB
<b>Gene Name</b>	HCCS
<b>Protein Name</b>	Cytochrome c-type heme lyase
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from human Cytochrome c-type Heme Lyase. AA range:81-130
<b>Specificity</b>	HCCS Monoclonal Antibody detects endogenous levels of HCCS protein.
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source</b>	Monoclonal, Mouse,IgG
<b>Purification</b>	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Dilution</b>	WB 1:500-1:2000
<b>Concentration</b>	1 mg/ml
<b>Purity</b>	≥90%
<b>Storage Stability</b>	-20°C/1 year
<b>Synonyms</b>	HCCS; CCHL; Cytochrome c-type heme lyase; CCHL; Holocytochrome c-type synthase
<b>Observed Band</b>	31kD
<b>Cell Pathway</b>	Mitochondrion inner membrane . Membrane ; Lipid-anchor .
<b>Tissue Specificity</b>	Brain,Liver,Ovary,
<b>Function</b>	catalytic activity:Holocytochrome c = apocytochrome c + heme.,disease:Defects in HCCS are a cause of microphthalmia syndromic type 7 (MCOPS7) [MIM:309801]; also known as microphthalmia with linear skin defects (MLS) or MIDAS syndrome. Microphthalmia is a clinically heterogeneous disorder of eye formation, ranging from small size of a single eye TO complete bilateral absence of ocular tissues (anophthalmia). In many cases, microphthalmia/anophthalmia occurs in association with syndromes that include non-ocular abnormalities. MCOPS7 is a disorder characterized by unilateral or bilateral microphthalmia, linear skin defects in affected females, and in utero lethality for males. Skin defects are limited to the face and neck, consisting of areas of aplastic skin that heal with age to form hyperpigmented areas. Additional features in female patients include agenesis of the corpus callosum, scle



## Background

holocytochrome c synthase(HCCS) Homo sapiens The protein encoded by this gene is an enzyme that covalently links a heme group to the apoprotein of cytochrome c. Defects in this gene are a cause of microphthalmia syndromic type 7 (MCOPS7). Three transcript variants encoding the same protein have been found for this gene. [provided by RefSeq, Jan 2010],

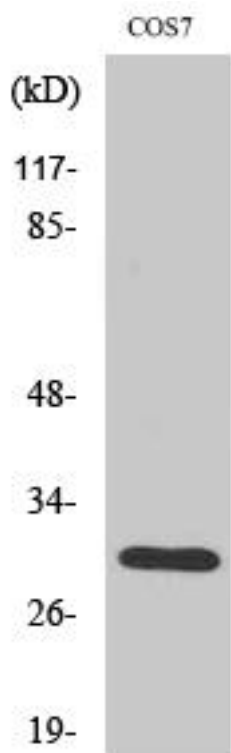
## matters needing attention

Avoid repeated freezing and thawing!

## Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

## Products Images



Western Blot analysis of various cells using HCCS Monoclonal Antibody