



# DHHC-15 Monoclonal Antibody

<b>Catalog No</b>	YP-mAb-01651
<b>Isotype</b>	IgG
<b>Reactivity</b>	Human;Rat;Mouse;
<b>Applications</b>	WB
<b>Gene Name</b>	ZDHHC15
<b>Protein Name</b>	Palmitoyltransferase ZDHHC15
<b>Immunogen</b>	The antiserum was produced against synthesized peptide derived from human ZDHHC15. AA range:288-337
<b>Specificity</b>	DHHC-15 Monoclonal Antibody detects endogenous levels of DHHC-15 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>	ZDHHC15; Palmitoyltransferase ZDHHC15; Zinc finger DHHC domain-containing protein 15; DHHC-15
<b>Observed Band</b>	38kD
<b>Cell Pathway</b>	Golgi apparatus membrane ; Multi-pass membrane protein . Cell junction, synapse, postsynaptic density .
<b>Tissue Specificity</b>	Expressed in placenta, liver, lung, kidney, heart and brain.
<b>Function</b>	catalytic activity:Palmitoyl-CoA + protein-cysteine = S-palmitoyl protein + CoA.;disease:Defects in ZDHHC15 are the cause of mental retardation X-linked type 91 (MRX91) [MIM:300577]. Mental retardation is characterized by significantly sub-average general intellectual functioning associated with impairments in adaptative behavior and manifested during the developmental period. Non-syndromic mental retardation patients do not manifest other clinical signs.;domain:The DHHC domain is required for palmitoyltransferase activity.;function:Palmitoyltransferase specific for GAP43 and DLG4/PSD95.;PTM:Autopalmitoylated.;similarity:Belongs to the DHHC palmitoyltransferase family.;similarity:Contains 1 DHHC-type zinc finger.;tissue specificity:Expressed in placenta, liver, lung, kidney, heart and brain.;
<b>Background</b>	The protein encoded by this gene belongs to the DHHC palmitoyltransferase family. Mutations in this gene are associated with mental retardatio X-linked type



91 (MRX91). Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2009],

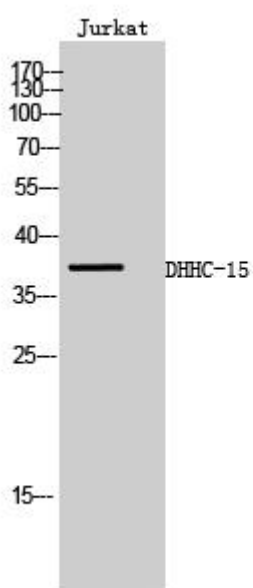
**matters needing attention**

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 DHHC-15 Monoclonal Antibody