

MLXPL rabbit pAb

Catalog_no :	<u>AT6797</u>
Applications :	<u>WB</u>
Reactivity :	<u>Human, Mouse,Rat</u>
Category :	<u>抗原抗体</u>
Size :	<u>100µg/50µg/20µg</u>
Gene_name :	<u>MLXIPL BHLHD14 MIO WBSCR14</u>
Protein_name :	<u>MLXPL</u>
Humangene_id	<u>51085</u>
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Humanswissprot	<u>Q9NP71</u>
_no :	
Mousegene_id :	<u>58805</u>
Mouseswissprot	<u>Q99MZ3</u>
_no :	
Ratgene_id :	<u>171078</u>
Ratswissprot_no	<u>Q8VIP2</u>
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Immunogen :	<u>Synthesized peptide derived from human MLXPL</u>
Specificity :	<u>This antibody detects endogenous levels of MLXPL at Human/Mouse/Rat</u>
Formulation :	<u>Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.</u>
Source :	<u>Rabbit</u>
Dilution :	<u>WB 1 : 500-2000</u>
Purification :	<u>The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.</u>
Concentration :	<u>1 mg/ml</u>
Storage_stability	<u>-20°C/1 year</u>
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Background :	<u>This gene encodes a basic helix-loop-helix leucine zipper transcription factor of the Myc/Max/Mad superfamily. This protein forms a heterodimeric complex and binds and activates, in a glucose-dependent manner, carbohydrate response element (ChoRE)</u>



motifs in the promoters of triglyceride synthesis genes. The gene is deleted in Williams-Beuren syndrome, a multisystem developmental disorder caused by the deletion of contiguous genes at chromosome 7q11.23. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Dec 2015],
