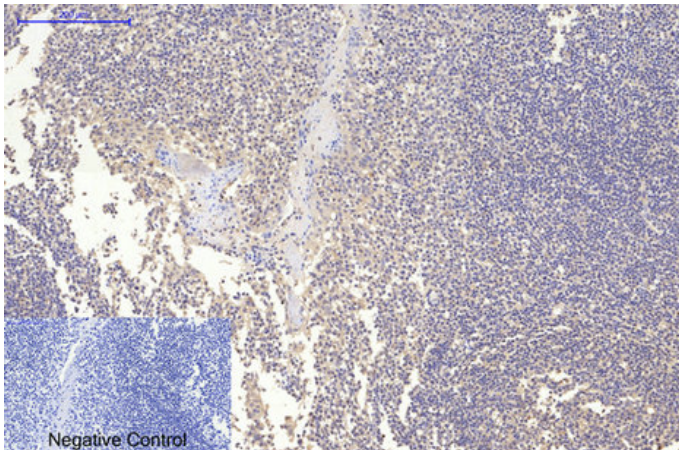




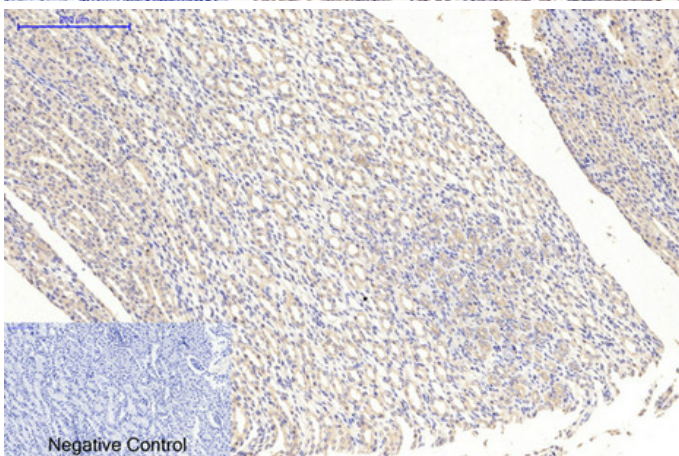
## MAP2 Monoclonal Antibody(7D4)

Catalog_no :	<u>AM3067</u>
Applications :	<u>IHC-P,IF</u>
Reactivity :	<u>Human,Mouse,Rat</u>
Category :	<u>抗原抗体</u>
Size :	<u>100µg/50µg</u>
Gene_name :	<u>MAP2</u>
Protein_name :	<u>Microtubule-associated protein 2</u>
Humangene_id	<u><a href="#">4133</a></u>
:	<u></u>
Humanswissprot	<u><a href="#">P11137</a></u>
_no :	<u></u>
Mouseswissprot	<u><a href="#">P20357</a></u>
_no :	<u></u>
Ratswissprot_no	<u><a href="#">P15146</a></u>
:	<u></u>
Immunogen :	<u>Synthetic Peptide of MAP2</u>
Specificity :	<u>The antibody detects endogenous MAP2 proteins.</u>
Formulation :	<u>PBS, pH 7.4, containing 0.02% sodium azide as Preservative and 50% Glycerol.</u>
Source :	<u>Mouse</u>
Dilution :	<u>IHC 1:200 IF 1:50-200</u>
Purification :	<u>The antibody was affinity-purified from mouse ascites by affinity-chromatography using specific immunogen.</u>
Storage_stability	<u>-20°C/1 year</u>
:	<u></u>
Msds :	<u>MSDS_Antibody.pdf</u>
Other_name :	<u>MAP2; Microtubule-associated protein 2; MAP-2</u>

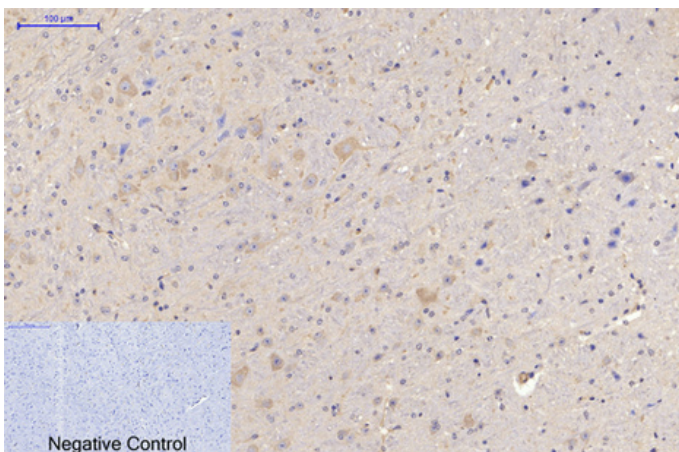
## Product Images



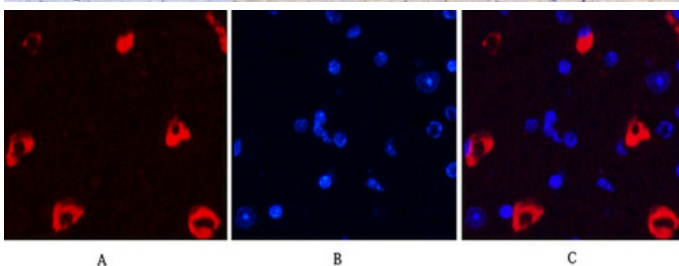
Immunohistochemical analysis of paraffin-embedded Human-Tonsil tissue. 1,MAP2 Monoclonal Antibody(7D4) was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.



Immunohistochemical analysis of paraffin-embedded Rat-kidney tissue. 1,MAP2 Monoclonal Antibody(7D4) was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.



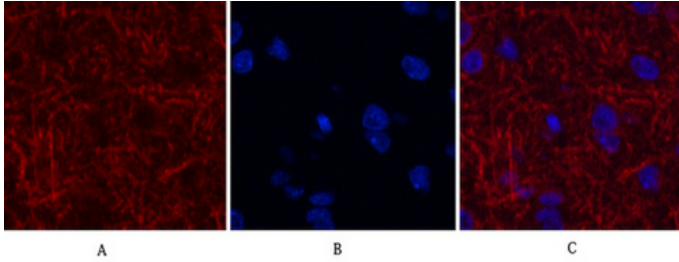
Immunohistochemical analysis of paraffin-embedded Mouse-brain tissue. 1,MAP2 Monoclonal Antibody(7D4) was diluted at 1:200(4°C,overnight). 2, Sodium citrate pH 6.0 was used for antibody retrieval(>98°C,20min). 3,Secondary antibody was diluted at 1:200(room temperature, 30min). Negative control was used by secondary antibody only.



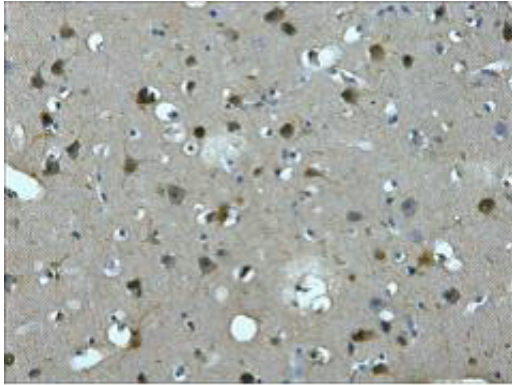
Immunofluorescence analysis of Mouse-brain tissue. 1,MAP2 Monoclonal Antibody(7D4)(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labeled Secondary antibody was diluted at 1:300(room temperature, 50min).3, Picture B: DAPI(blue) 10min. Picture A:Target. Picture B: DAPI. Picture C: merge of A+B

Immunofluorescence analysis of Rat-brain tissue. 1,MAP2 Monoclonal Antibody(7D4)(red) was diluted at 1:200(4°C,overnight). 2, Cy3 labeled Secondary antibody

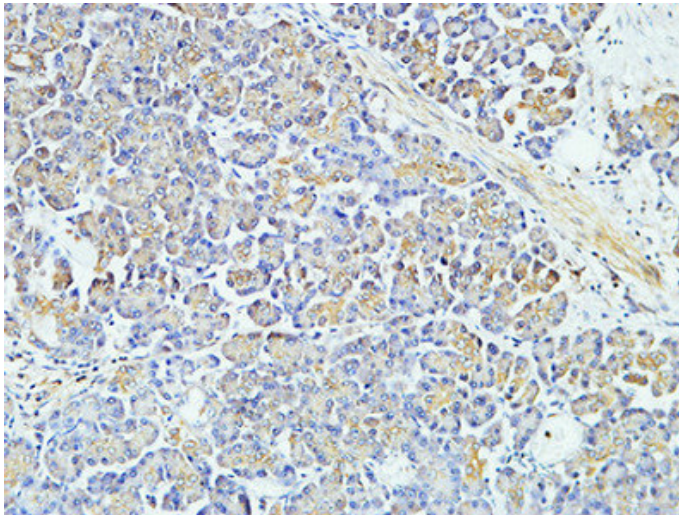




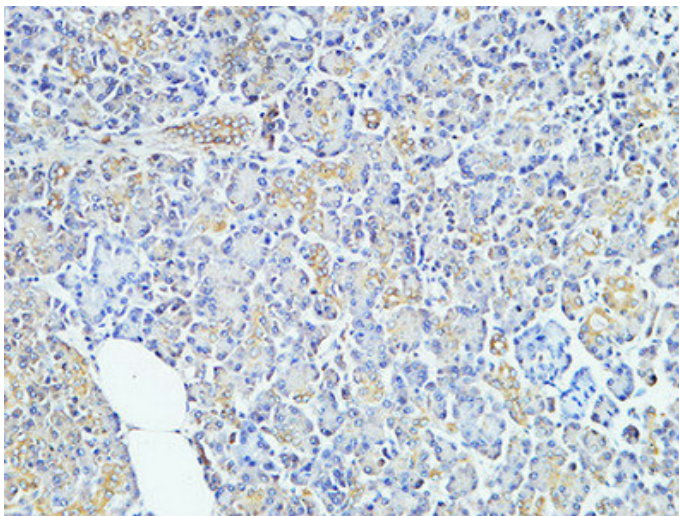
was diluted at 1:300(room temperature, 50min).3,  
Picture B: DAPI(blue) 10min. Picture A:Target. Picture B:  
DAPI. Picture C: merge of A+B



IHC staining of Human brain tissue paraffin-embedded,  
diluted at 1:200.

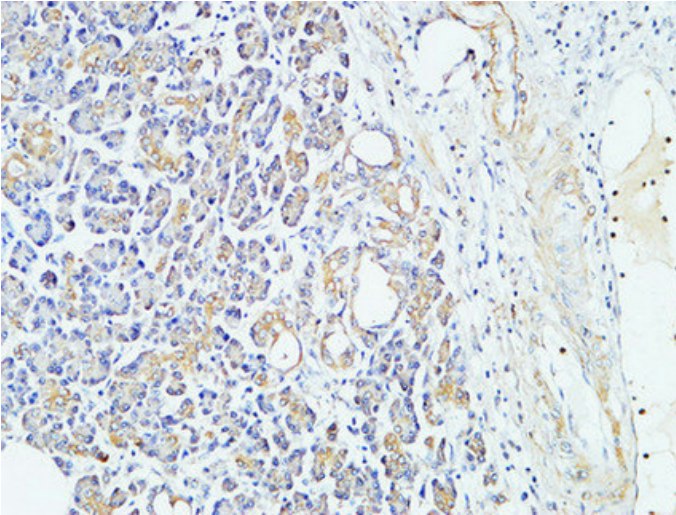


Immunohistochemical analysis of paraffin-embedded  
Human pancreas. 1, Antibody was diluted at  
1:400(4°,overnight). 2, High-pressure and temperature  
EDTA, pH8.0 was used for antigen retrieval. 3,Secondary  
antibody was diluted at 1:200(room temperature,  
30min).

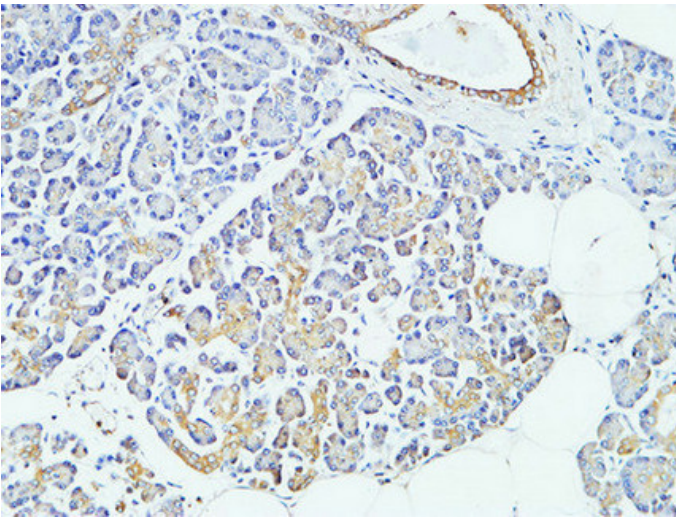


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Immunohistochemical analysis of paraffin-embedded Human pancreas. 1, Antibody was diluted at 1:400(4°, overnight). 2, High-pressure and temperature EDTA, pH8.0 was used for antigen retrieval. 3, Secondary antibody was diluted at 1:200(room temperature, 30min).