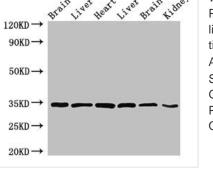


🕜 Tel: +1-301-363-4651 🛛 🗵 Email: cusabio@cusabio.com 📀 Website: www.cusabio.com 🍙

## **MBP** Antibody

Product Code	CSB-PA013551LA01HU
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	P02686
Immunogen	Recombinant Human Myelin basic protein (1-197AA)
Raised In	Rabbit
Species Reactivity	Human, Mouse, Rat
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:100-1:500, IF:1:50-1:500
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Purification Method	>95%, Protein G purified
Isotype	lgG
Clonality	Polyclonal
Alias	Myelin basic protein (MBP) (Myelin A1 protein) (Myelin membrane encephalitogenic protein), MBP
Immunogen Species	Homo sapiens (Human)
Research Area	Neuroscience
Target Names	MBP
Image	$120\text{KD} \rightarrow \frac{\text{Bre^{3T}}_{1}\text{1}^{\text{se^{T}}}\text{Re^{3T}}_{1}\text{1}^{\text{se^{T}}}\text{Re^{3T}}_{1}\text{1}^{\text{Re^{T}}}\text{Re^{3T}}_{1}\text{1}^{\text{Re^{T}}}}{\text{Ne^{T}}} Western Blot$ Positive WB detected in: Rat brain tissue, Rat liver tissue, Mouse heart tissue, Mouse liver

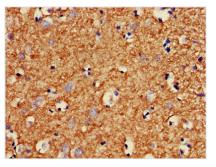


tissue, Mouse brain tissue, Mouse kidney tissue All lanes: MBP antibody at 4µg/ml Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 34, 22, 21, 19, 18 kDa Observed band size: 34 kDa

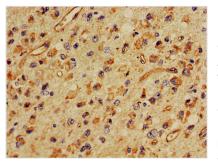
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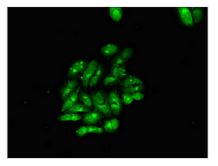
🕜 Tel: +1-301-363-4651 🛛 🖂 Email: cusabio@cusabio.com 🛛 🥭 Website: www.cusabio.com 🌘



IHC image of CSB-PA013551LA01HU diluted at 1:400 and staining in paraffin-embedded human brain tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



IHC image of CSB-PA013551LA01HU diluted at 1:400 and staining in paraffin-embedded human glioma performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.



Immunofluorescent analysis of Hela cells using CSB-PA013551LA01HU at dilution of 1:100 and Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L)