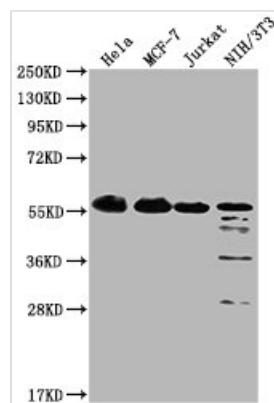




# PKM Monoclonal Antibody

<b>Product Code</b>	CSB-MA018072A0m
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P14618
<b>Immunogen</b>	Recombinant Human Pyruvate kinase PKM protein (2-531AA)
<b>Raised In</b>	Mouse
<b>Species Reactivity</b>	Human, Rat, Mouse, Rabbit
<b>Tested Applications</b>	ELISA, WB, IHC, IF, FC, IP; Recommended dilution: WB: 1:4000-1:256000, IHC: 1:200-1:500, IF: 1:150-1:300, FC: 1:50-1:200, IP: 1µl-4µl
<b>Relevance</b>	Glycolytic enzyme that catalyzes the transfer of a phosphoryl group from phosphoenolpyruvate (PEP) to ADP, generating ATP. Stimulates POU5F1-mediated transcriptional activation. Plays a general role in caspase independent cell death of tumor cells. The ratio between the highly active tetrameric form and nearly inactive dimeric form determines whether glucose carbons are channeled to biosynthetic processes or used for glycolytic ATP production. The transition between the 2 forms contributes to the control of glycolysis and is important for tumor cell proliferation and survival.
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, PH 7.4
<b>Purification Method</b>	>95%, Protein G purified
<b>Isotype</b>	IgG1
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Monoclonal Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Gene Names</b>	PKM
<b>PubMed ID</b>	6C3C7

## Image



### Western Blot

Positive WB detected in: HeLa whole cell lysate, MCF-7 whole cell lysate, Jurkat whole cell lysate, NIH/3T3 whole cell lysate

All lanes: PKM antibody at 1:4000

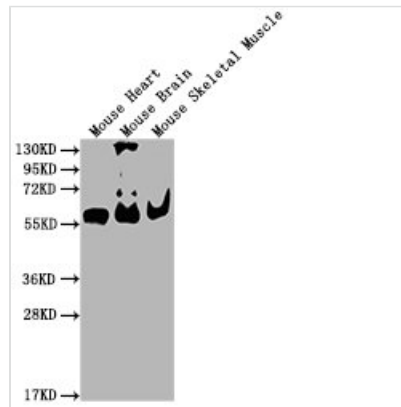
Secondary

Goat polyclonal to Mouse IgG at 1/10000 dilution

Predicted band size: 58 kDa

Observed band size: 58 KDa

Exposure time: 1min


**Western Blot**

Positive WB detected in: Mouse Heart tissue, Mouse Brain tissue, Mouse Skeletal Muscle tissue

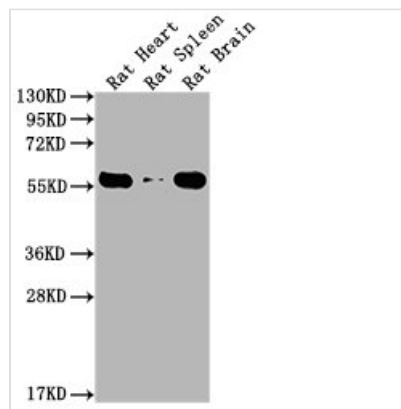
All lanes: PKM antibody at 1:4000

Secondary

Goat polyclonal to Mouse IgG at 1/10000 dilution

Predicted band size: 58 kDa

Observed band size: 58 kDa


**Western Blot**

Positive WB detected in: Rat Heart tissue, Rat Spleen tissue, Rat Brain tissue

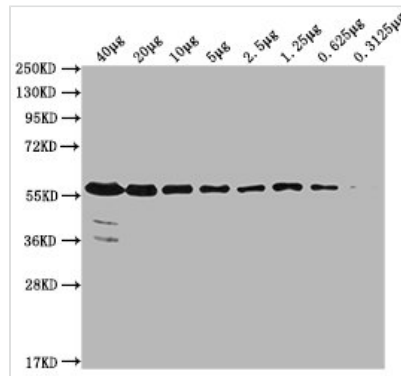
All lanes: PKM antibody at 1:4000

Secondary

Goat polyclonal to Mouse IgG at 1/10000 dilution

Predicted band size: 55-60 kDa

Observed band size: 55-60 kDa


**Western Blot**

Positive WB detected in: MCF-7 whole cell lysate at 40µg, 20µg, 10µg, 5µg, 2.5µg, 1.25µg, 0.625µg, 0.3125µg

All lanes: PKM antibody at 1:4000

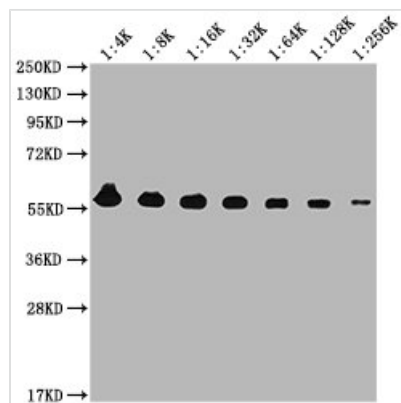
Secondary

Goat polyclonal to Mouse IgG at 1/10000 dilution

Predicted band size: 58 kDa

Observed band size: 58 kDa

Exposure time: 5min


**Western Blot**

Positive WB detected in: MCF-7 whole cell lysate

All lanes: PKM antibody at 1:4000, 1:8000, 1:16000, 1:32000, 1:64000, 1:128000, 1:256000

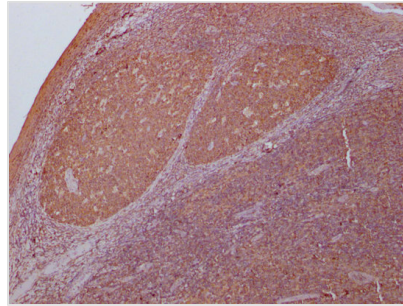
Secondary

Goat polyclonal to Mouse IgG at 1/10000 dilution

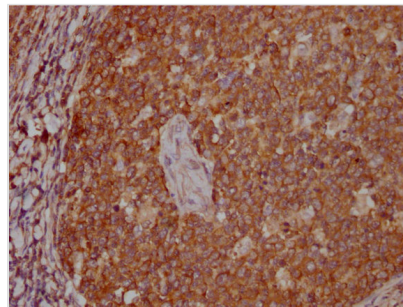
Predicted band size: 58 kDa

Observed band size: 58 kDa

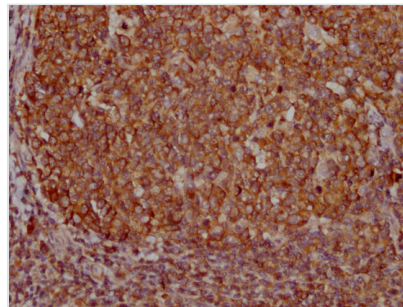
Exposure time: 5min



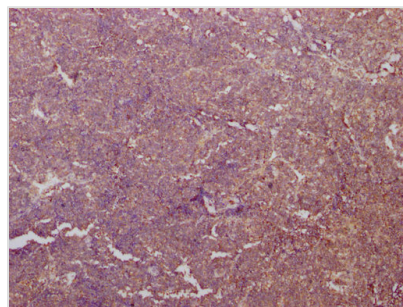
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human tonsil tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. 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.



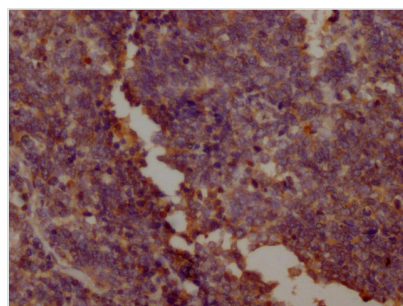
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human tonsil tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. 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-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human tonsil tissue performed on a Leica Bond™ 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.



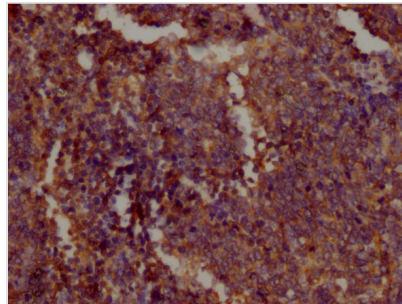
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human lung cancer tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min 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.



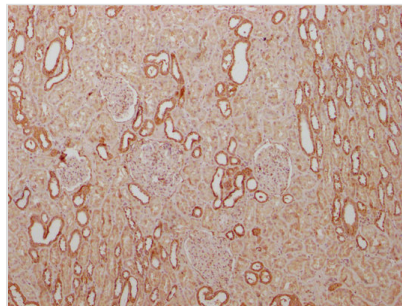
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human lung cancer tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a



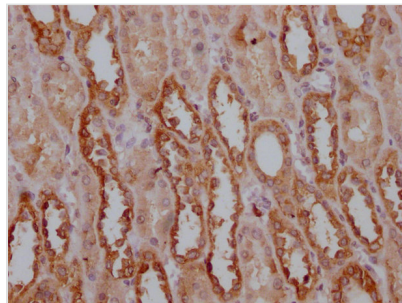
biotinylated secondary antibody and visualized using an HRP conjugated SP system.



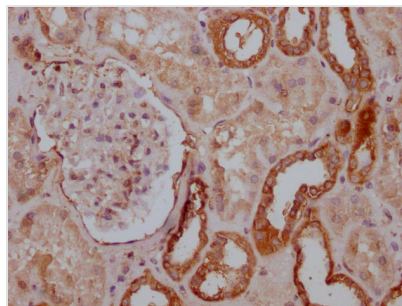
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human lung cancer tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min 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.



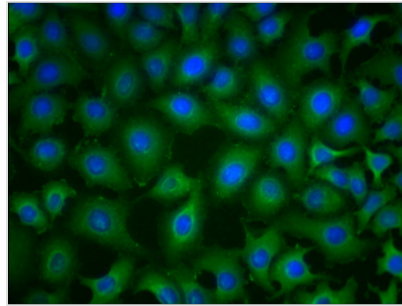
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human kidney tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. 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.



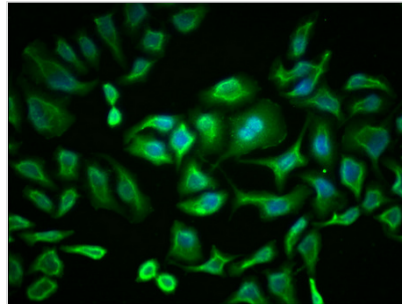
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human kidney tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. 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.



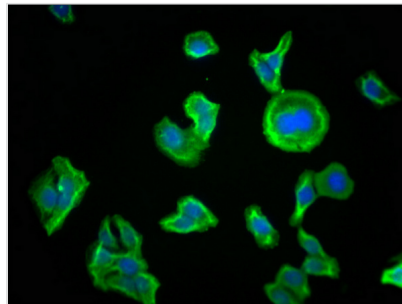
IHC image of CSB-MA018072A0m diluted at 1:400 and staining in paraffin-embedded human kidney tissue performed on a Leica Bond™ system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. 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.



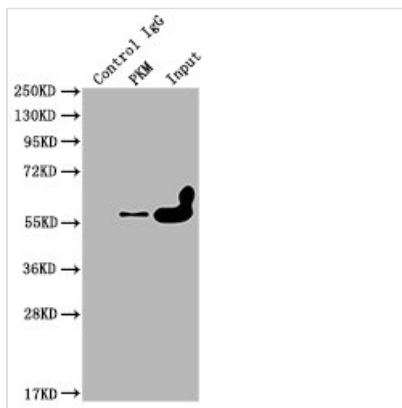
Immunofluorescence staining of A549 cells with CSB-MA018072A0m at 1:230, counter-stained with DAPI. The cells were blocked in 10% normal Goat Serum and then incubated with the primary antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



Immunofluorescence staining of HeLa cells with CSB-MA018072A0m at 1:230, counter-stained with DAPI. The cells were blocked in 10% normal Goat Serum and then incubated with the primary antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



Immunofluorescence staining of HepG2 cells with CSB-MA018072A0m at 1:230, counter-stained with DAPI. The cells were blocked in 10% normal Goat Serum and then incubated with the primary antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



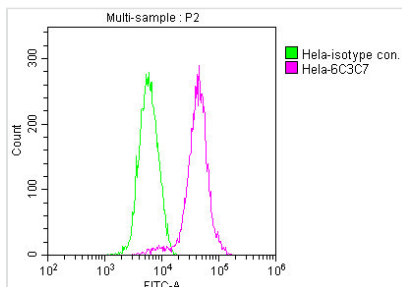
Immunoprecipitating PKM in HeLa whole cell lysate

Lane 1: Mouse control IgG instead of CSB-MA018072A0m in HeLa whole cell lysate.

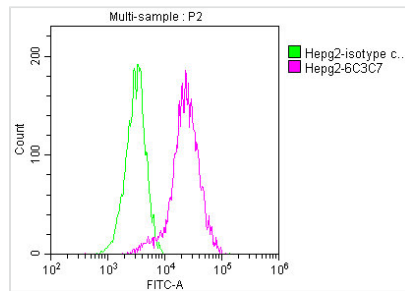
Lane 2: CSB-MA018072A0m (1?) + HeLa whole cell lysate (500?g)

Lane 3: HeLa whole cell lysate (10?g)

For western blotting, the blot was detected with CSB-MA018072A0m at 1:2000, and a HRP-conjugated Protein G antibody was used as the secondary antibody at 1:2000



Overlay histogram showing HeLa cells stained with CSB-MA018072A0m (red line) at 1:100. The cells were incubated in 1x PBS /10% normal goat serum to block non-specific protein-protein interactions followed by primary antibody for 1 h at 4°C. The secondary antibody used was FITC goat anti-mouse IgG(H+L) at 1/200 dilution for 1 h at 4°C. Isotype control antibody (green line) was used under the same conditions. Acquisition of >10,000 events was performed.



Overlay histogram showing HepG2 cells stained with CSB-MA018072A0m (red line) at 1:100. The cells were incubated in 1x PBS /10% normal goat serum to block non-specific protein-protein interactions followed by primary antibody for 1 h at 4°C. The secondary antibody used was FITC goat anti-mouse IgG(H+L) at 1/200 dilution for 1 h at 4°C. Isotype control antibody (green line) was used under the same conditions. Acquisition of >10,000 events was performed.