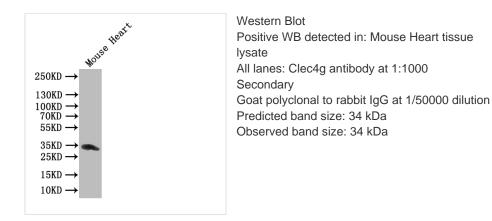


🕜 Tel: +1-301-363-4651 🛛 🗵 Email: cusabio@cusabio.com 📀 Website: www.cusabio.com 🍙

Clec4g Antibody

Product Code	CSB-PA803912ZA01MO
Storage	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
Uniprot No.	Q8BNX1
Immunogen	Recombinant Mus musculus Clec4g protein (52-294aa)
Raised In	Rabbit
Species Reactivity	Mus musculus
Tested Applications	ELISA, WB, IHC, IF; Recommended dilution: WB:1:500-1:5000, IHC:1:50-1:200, IF:1:30-1:100
Form	Liquid
Conjugate	Non-conjugated
Storage Buffer	Preservative: 0.03% Proclin 300 Constituents: 50% Glycerol, 0.01M PBS, pH 7.4
Purification Method	Antigen affinity purification
Isotype	lgG
Clonality	Polyclonal
Product Type	Polyclonal Antibody
Immunogen Species	Mus musculus (Mouse)
Target Names	Clec4g
_	

Image

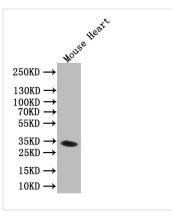


1

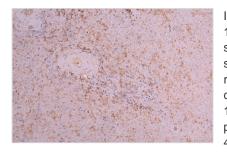
CUSABIO TECHNOLOGY LLC



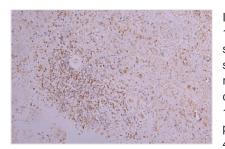
🕜 Tel: +1-301-363-4651 🛛 🖂 Email: cusabio@cusabio.com 🤅 Website: www.cusabio.com 🌘



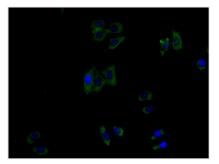
Western Blot Positive WB detected in: Mouse Heart tissue lysate All lanes: Clec4g antibody at 1:1000 Secondary Goat polyclonal to rabbit IgG at 1/50000 dilution Predicted band size: 34 kDa Observed band size: 34 kDa



IHC image of CSB-PA803912ZA01MO diluted at 1:66 and staining in paraffin-embedded human spleen 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 Goat anti-rabbit polymer IgG labeled by HRP and visualized using 0.05% DAB.



IHC image of CSB-PA803912ZA01MO diluted at 1:66 and staining in paraffin-embedded human spleen 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 Goat anti-rabbit polymer IgG labeled by HRP and visualized using 0.05% DAB.



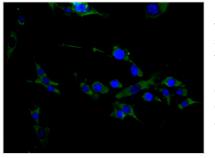
Immunofluorescence staining of Hela cell with CSB-PA803912ZA01MO at 1:30, counterstained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).

2

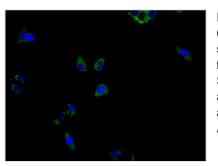


CUSABIO TECHNOLOGY LLC

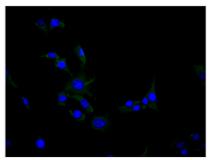
🕜 Tel: +1-301-363-4651 🛛 🖂 Email: cusabio@cusabio.com 🛛 🥭 Website: www.cusabio.com 🌘



Immunofluorescence staining of NIH/3T3 cell with CSB-PA803912ZA01MO at 1:30, counterstained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Immunofluorescence staining of Hela cell with CSB-PA803912ZA01MO at 1:30, counterstained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Immunofluorescence staining of NIH/3T3 cell with CSB-PA803912ZA01MO at 1:30, counterstained with DAPI. The cells were fixed in 4% formaldehyde and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).