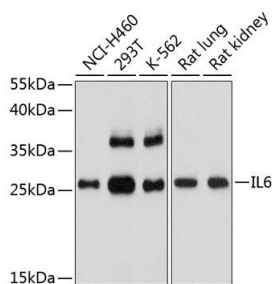


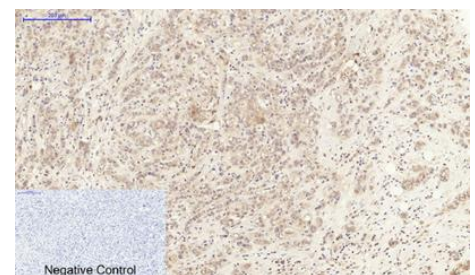
<b>Cat. No:</b>	MAB-94635
<b>Conjugate:</b>	Unconjugated
<b>Size:</b>	100 ug
<b>Clone:</b>	I621
<b>Concentration:</b>	1mg/ml
<b>Host:</b>	Rabbit
<b>Isotype:</b>	IgG
<b>Immunogen:</b>	The antiserum was produced against synthesized peptide derived from the Internal region of human IL6. AA range:131-180
<b>Reactivity:</b>	Hu,Ms, Rt  Western Blot: 1/500 - 1/2000.
<b>Applications:</b>	Immunohistochemistry (paraffin-embedded tissues): 1:100-300 Immunofluorescence: 1:100-300 ELISA: 1/20000
<b>Molecular Weight:</b>	23kDa
<b>Purification:</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Synonyms:</b>	IL6; IFNB2; Interleukin-6; IL-6; B-cell stimulatory factor 2; BSF-2; CTL differentiation factor; CDF; Hybridoma growth factor; Interferon beta-2; IFN-beta-2
<b>Background:</b>	This gene encodes a cytokine that functions in inflammation and the maturation of B cells. In addition, the encoded protein has been shown to be an endogenous pyrogen capable of inducing fever in people with autoimmune diseases or infections. The protein is primarily produced at sites of acute and chronic inflammation, where it is secreted into the serum and induces a transcriptional inflammatory response through interleukin 6 receptor, alpha. The functioning of this gene is implicated in a wide variety of inflammation-associated disease states, including susceptibility to diabetes mellitus and systemic juvenile rheumatoid arthritis. Alternative splicing results in multiple transcript variants.
<b>Form:</b>	Liquid
<b>Buffer:</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Storage:</b>	Store at -20°C. Avoid freeze / thaw cycles.



Western blot analysis of extracts of various cell lines, using IL6 antibody at 1:1000



Western Blot analysis of K562 cells using IL-6 Antibody. Antibody was diluted at 1:1000.

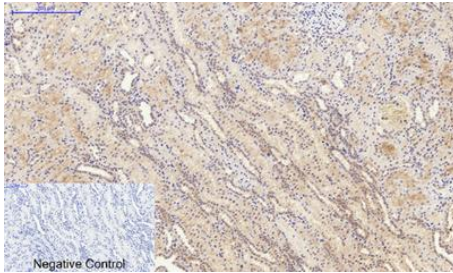


Immunohistochemical analysis of paraffin-embedded Human-breast-cancer tissue. 1,IL-6 Antibody was diluted at

dilution.  
Secondary antibody: HRP Goat Anti-  
Rabbit IgG  
(H+L) at 1:10000 dilution.  
Lysates/proteins: 25ug per lane.  
Blocking buffer: 3% nonfat dry milk in  
TBST.  
Detection: ECL West Pico Plus.  
Exposure time: 90s.

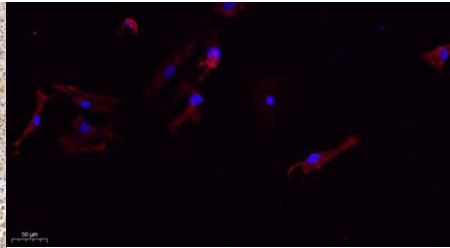
Secondary antibody was diluted at  
1:20000

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 Human-kidney  
tissue.

1,IL-6 Antibody 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(



Immunofluorescence analysis of A549.  
1,primary Antibody(red) was diluted at  
1:200(4°C overnight).  
2, Goat Anti Rabbit IgG (H&L) - Alexa  
Fluor 594 Secondary  
antibody was diluted at 1:1000(room  
temperature, 50min).  
3, Picture B: DAPI(blue) 10min.