

AS038

Leader in Biomolecular Solutions for Life Science



HRP-conjugated Donkey anti-Rabbit IgG (H+L)

Catalog No.: AS038 **12 Publications**

Basic Information

Observed MW

42kDa

Calculated MW

Category

Secondary Antibody

Applications

WB,IHC-P

Cross-Reactivity

Conjugate

HRP

Background

Secondary antibodies are affinity-purified antibodies which will work with target-specific primary antibody in the detection, sorting or purification of its specified target. Secondary antibodies offer increased versatility enabling users to use many detection systems (e.g. HRP, AP, fluorescence). They can also provide greater sensitivity through signal amplification as multiple secondary antibodies. Most commonly, secondary antibodies are generated by immunizing the host animal (different from host species of primary antibody) with a pooled population of normal immunoglobulins from the host species of primary antibody and can be further purified and modified (i.e. antibody fragmentation, label conjugation, etc.) to ensure well-characterized specificity to corresponding normal immunoglobulins.

Recommended Dilutions

WB 1:5000 - 1:100000

IHC-P 1:50 - 1:200

Immunogen Information

Gene ID Swiss Prot

Immunogen

Rabbit IgG

Synonyms

Contact

 www.abclonal.com

Product Information

Source

Donkey

Isotype

Horseradish peroxidase
conjugated IgG

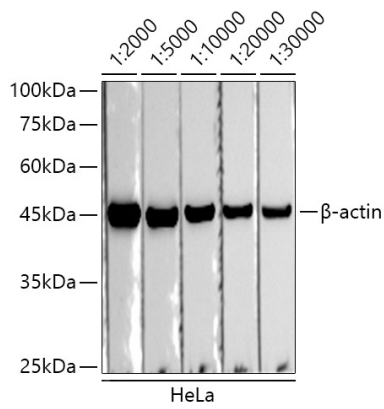
Purification

Affinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles.
Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.

Validation Data



Western blot analysis of lysates from HeLa cells, using β -actin antibody as the primary antibody.

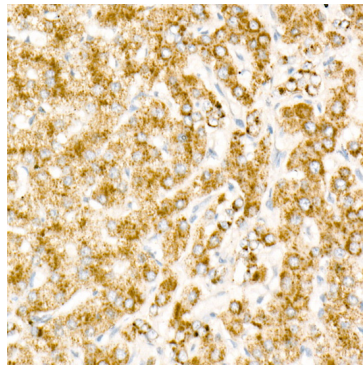
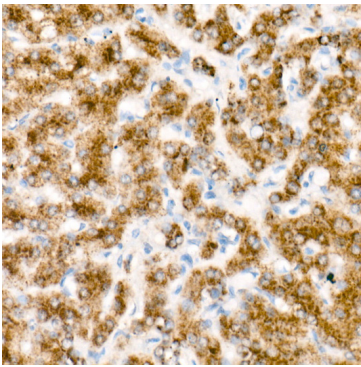
Secondary antibody: HRP Donkey Anti-Rabbit IgG (H+L) antibody (AS038) at 1:2000-1:30000 dilution

Lysates/proteins: 25 μ g per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 30s.



Immunohistochemistry analysis of paraffin-embedded Human liver (primary antibody is acox1) using HRP Donkey Anti-Rabbit IgG (H+L) (AS038) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.

Immunohistochemistry analysis of paraffin-embedded Human liver (the primary antibody is acox1, And the control secondary antibody is stained) using HRP Donkey Anti-Rabbit IgG (H+L) (AS038) at dilution of 1:100 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.