# [One Step] C1QBP Antibody Kit



Catalog No.: RK05628

## **Basic Information**

#### **Observed MW**

#### **Calculated MW**

## Category

Primary antibody

# **Background**

## Component

Catalog No.	<b>Product Name</b>	Applications	Cross-Reactivity
AS014	HRP-conjugated Goat anti-Rabbit IgG (H+L)	ELISA,WB,IHC-P,DB	
A1883	GC1q R/C1QBP Rabbit	ELISA,WB,IF/ICC	Human, Mouse, Rat

# **Recommended Dilutions**

**AS014 ELISA** 1:5000 - 1:10000

**A1883 WB** 1:500 - 1:2000

For more information please visit www.abclonal.com

## **Product Information**

Source Isotype

**Purification** 

Storage

## **Contact**

<b>a</b>	400-999-6126
×	cn.market@abclonal.com.cn
$\overline{f e}$	www.abclonal.com.cn

# **Immunogen Information**

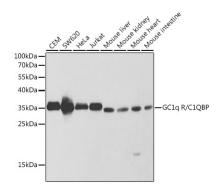
Gene ID Swiss Prot

## **Immunogen**

Rabbit IgG

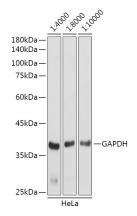
Recombinant fusion protein containing a sequence corresponding to amino acids 74-282 of human GC1q R/GC1q R/C1QBP (NP\_001203.1).

## **Synonyms**



Western blot analysis of various lysates using GC1q R/C1QBP Rabbit pAb (A1883) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins:  $25\mu g$  per lane.

Blocking buffer: 3% nonfat dry milk in TBST.



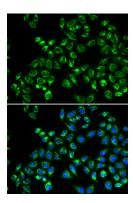
Western blot analysis of lysates from HeLa cells, using GAPDH (AC001) antibody as the primary antibody at dilution of 1:80000.

Secondary antibody: using HRP Goat Anti-Rabbit lgG (H+L) antibody (AS014) at 1:4000-1:10000 dilution. Lysates/proteins: 25 $\mu$ g per lane.

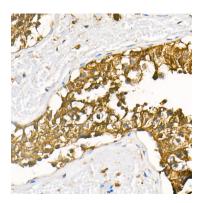
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 3s.



Immunofluorescence analysis of HeLa cells using GC1q R/C1QBP Rabbit pAb (A1883). Secondary antibody: Cy3 Goat Anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.



Immunohistochemistry analysis of HRP Goat Anti-Rabbit IgG (H+L) in paraffinembedded human testis tissue using HRP Goat Anti-Rabbit IgG (H+L) (AS014) at a dilution of 1:100 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.