## **BBS4 Rabbit Polyclonal Antibody**



## **CAB3759**

**Product Information** 

Size:

20uL, 50uL, 100uL, 200uL

**Observed MW:** 

53kDa

Calculated MW:

38kDa/58kDa/59kDa

**Applications:** 

WB IF

Reactivity:

Human

Q96RK4

**Antibody Information** 

**Recommended dilutions:** WB 1:500 - 1:2000 IF 1:50 -

1:200

Source:

Rabbit

Isotype:

**Purification:** Affinity purification

IgG

**Protein Background** 

This gene is a member of the Bardet-Biedl syndrome (BBS) gene family. Bardet-Biedl syndrome is an autosomal recessive disorder characterized by severe pigmentary retinopathy, obesity, polydactyly, renal malformation and mental retardation. The proteins encoded by BBS gene family members are structurally diverse. The similar phenotypes exhibited by mutations in BBS gene family members are likely due to the protein's shared roles in cilia formation and function. Many BBS proteins localize to the basal bodies, ciliary axonemes, and pericentriolar regions of cells. BBS proteins may also be involved in intracellular trafficking via microtubule-related transport. The protein encoded by this gene has sequence similarity to O-linked Nacetylglucosamine (O-GlcNAc) transferases in plants and archaebacteria and in human forms a multi-protein 'BBSome' complex with seven other BBS proteins. Alternate splicing results in multiple transcript variants.

Immunogen information

Gene ID:

585

Uniprot

Synonyms:

BBS4

Immunogen:

Recombinant fusion protein containing a sequence corresponding

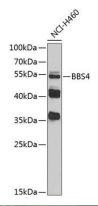
to amino acids 350-519 of human BBS4 (NP\_149017.2).

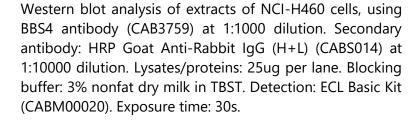
Storage:

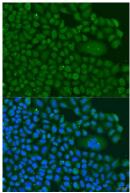
Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.02%

sodium azide, 50% glycerol, pH7.3.

## **Product Images**







Immunofluorescence analysis of U2OS cells using BBS4 antibody (CAB3759) at dilution of 1:100. Blue: DAPI for nuclear staining.