

## Product datasheet for **AP14206PU-N**

### **BBS4 (Center) Rabbit Polyclonal Antibody**

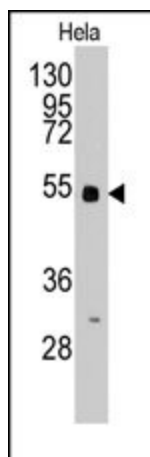
#### **Product data:**

<b>Product Type:</b>	Primary Antibodies
<b>Applications:</b>	WB
<b>Recommended Dilution:</b>	ELISA: 1/1,000. Western blotting: 1/50 - 1/100.
<b>Reactivity:</b>	Human
<b>Host:</b>	Rabbit
<b>Isotype:</b>	Ig
<b>Clonality:</b>	Polyclonal
<b>Immunogen:</b>	This antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide selected from the Center region of human BBS4.
<b>Specificity:</b>	This antibody reacts to BBS4.
<b>Formulation:</b>	PBS with 0.09% (W/V) sodium azide State: Purified State: Liquid purified Ig
<b>Concentration:</b>	lot specific
<b>Purification:</b>	Prepared by Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS
<b>Conjugation:</b>	Unconjugated
<b>Storage:</b>	Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing.
<b>Stability:</b>	Shelf life: one year from despatch.
<b>Gene Name:</b>	Bardet-Biedl syndrome 4
<b>Database Link:</b>	<a href="#">Entrez Gene 585 Human Q96RK4</a>
<b>Background:</b>	BBS4 contains tetratricopeptide repeats (TPR), similar to O-linked N-acetylglucosamine transferase. Mutations in the gene encoding this protein have been observed in patients with Bardet-Biedl syndrome type 4. BBS4 may play a role in pigmentary retinopathy, obesity, polydactyly, renal malformation and mental retardation.
<b>Synonyms:</b>	Bardet-Biedl syndrome 4 protein



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## Product images:



Western blot analysis of anti-BBS4 Pab in HeLa cell line lysates (35ug/lane). BBS4 (arrow) was detected using the purified Pab (1:60 dilution).