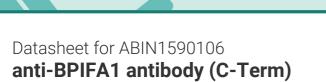
# antibodies -online.com







Image



### Overview

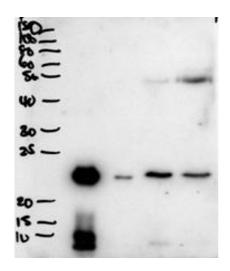
Quantity:	100 μg
Target:	BPIFA1
Binding Specificity:	C-Term
Reactivity:	Human
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This BPIFA1 antibody is un-conjugated
Application:	Western Blotting (WB), ELISA

## **Product Details**

Purpose:	BPIFA1 / PLUNC
Sequence:	NEVLRGLDIT LVHD
Isotype:	IgG
Specificity:	Reported variants represent identical protein: NP_057667.1, NP_001230122.1, NP_570913.1
Cross-Reactivity:	Human
Purification:	Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.
Grade:	Verified

# **Target Details**

Target:	BPIFA1
Alternative Name:	BPIFA1 (BPIFA1 Products)
Background:	BPIFA1, BPI fold containing family A, member 1, LPLUNC3, LUNX, NASG, PLUNC, SPLUNC1,
	SPURT, bA49G10.5, OTTHUMP0000030625, OTTHUMP00000030626,
	OTTHUMP00000030627, ligand-binding protein RYA3, lung-specific protein X, nasopharyngea
	carcinoma-related prote
Gene ID:	51297
NCBI Accession:	NP_057667
Application Details	
Application Notes:	Western Blot: Approx 26 kDa band observed in secretions of Human primary airway cells in
	culture-and in Human Bronchoalveolar Lavage fluid (calculated MW of 26.7 kDa according to
	NP_057667.1). Recommended concentration: 1-3 µg/mL.
	Peptide ELISA: antibody detection limit dilution 1:64000.
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	0.5 mg/mL
Buffer:	Supplied at 0.5 mg/mL in Tris saline, 0.02 % sodium azide, pH 7.3 with 0.5 % bovine serum
	albumin.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which
	should be handled by trained staff only.
Handling Advice:	Minimize freezing and thawing.
Storage:	-20 °C
Storage Comment:	Aliquot and store at -20°C, with minimal freeze/thawing. A working aliquot may be refrigerated
	at 4°C for a few weeks and still remain viable.



## **Western Blotting**

**Image 1.** ABIN1590106 (2 $\mu$ g/ml) staining of secretions from Human primary airway cells in culture (lanes 1 and 2), and in Human Bronchoalveolar Lavage fluid (lanes 3 and 4) . Data obtained from Dr. C Bingle, AURM, University of Sheffield, UK. Primary incubation was 1 hour. Detected by chemiluminescence.