



Product Information Sheet

Polyclonal Anti- serine/threonine kinase 11, STK11/LKB1

Catalogue No. PA1355 Immunogen

A synthetic peptide corresponding to a sequence at the C-terminal of

human STK11 (421-433 aa), different from the mouse sequence by

one amino acid.

Ig type rabbit IgG

Purity

Size 100µg/vial Immunogen affinity purified.

Specificity Application

No cross reactivity with other

Lot No. 01310120255124

proteins.

Human,rat,mouse

Recommended application

Western blot

Immunohistochemistry(P)

	Concen-	Tested	Concluded	Antigen
	tration	Species	Species	Retrieval
WB	1ug/ml	Hu, Rat	Ms	-
IHC-P	1ug/ml	Hu, Rat, Ms	-	By Heat
IHC-F	-	-	-	-
ICC	-	1	-	-

Other applications have not been tested.

Optimal dilutions should be determined by end user.

Contents

Each vial contains 5mg BSA, 0.9mg NaCl, 0.2mg Na_2HPO_4 , 0.05mg Thimerosal, 0.05mg NaN_3 .

Reconstitution

0.2ml of distilled water will yield a concentration of 500µg/ml.

To reorder contact us at:

Antagene, Inc. Storage

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Toll Free: 1(866)964-2589 email: Info@antageneinc.com

At -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for longer time.

BACKGROUND

Serine/threonine kinase 11 or LKB1 is a protein kinase which in humans is encoded by the *STK11* gene. The *STK11/LKB1* gene, which encodes a member of the serine/threonine kinase, regulates cell polarity and functions as a tumour suppressor Smith et al. (1999) found that the mouse Lkb1 gene encodes a protein showing strong sequence similarity to human LKB1. Karuman et al. (2001) demonstrated that LKB1 physically associates with p53 (191170) and regulates specific p53-dependent apoptosis pathways. Jenne et al. (1998) determined that the STK11 gene extends over 23 kb of genomic DNA and is composed of 9 exons, which are transcribed in telomere-to-centromere direction. Smith et al. (1999) found that the mouse Lkb1 gene consists of 10 exons covering approximately 15 kb.

REFERENCE

1.Jenne DE, Reimann H, Nezu J, Friedel W, Loff S, Jeschke R, Müller O, Back W, Zimmer M (January 1998). "Peutz-Jeghers syndrome is caused by mutations in a novel serine threonine kinase".