abcam

Product datasheet

Anti-ErbB3 / HER3 antibody [SGP1] ab20162

3 References

Overview

Product name Anti-ErbB3 / HER3 antibody [SGP1]

Description Mouse monoclonal [SGP1] to ErbB3 / HER3

Host species Mouse

Tested applications

Suitable for: IP, Flow Cyt

Species reactivity

Reacts with: Human

Predicted to work with: Mouse, Rat, Chimpanzee

Immunogen Recombinant full length protein corresponding to Human ErbB3/ HER3. 'c-erb B3 protein from

transfected human kidney fibroblasts purified by wheatgerm lectin affinity chromatography'

Epitope Extracellular domain.

General notesThe Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Myeloma

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -20°C or -80°C. Avoid freeze /

thaw cycle.

Storage buffer Preservative: 0.02% Sodium azide

NS0

Constituent: 99.98% PBS

Purity Protein A/G purified

Clonality Monoclonal

Clone number SGP1

Isotype IgG1

Light chain type kappa

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Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab20162 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IP		Use at an assay dependent concentration.
Flow Cyt		Use at an assay dependent concentration. <u>ab170190</u> - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.

Target

Function Binds and is activated by neuregulins and NTAK.

Tissue specificity Epithelial tissues and brain.

Involvement in diseaseDefects in ERBB3 are the cause of lethal congenital contracture syndrome type 2 (LCCS2)

[MIM:607598]; also called Israeli Bedouin multiple contracture syndrome type A. LCCS2 is an autosomal recessive neurogenic form of a neonatally lethal arthrogryposis that is associated with atrophy of the anterior horn of the spinal cord. The LCCS2 syndrome is characterized by multiple joint contractures, anterior horn atrophy in the spinal cord, and a unique feature of a markedly distended urinary bladder. The phenotype suggests a spinal cord neuropathic etiology.

Sequence similaritiesBelongs to the protein kinase superfamily. Tyr protein kinase family. EGF receptor subfamily.

Contains 1 protein kinase domain.

Developmental stageOverexpressed in a subset of human mammary tumors.

Domain The cytoplasmic part of the receptor may interact with the SH2 or SH3 domains of many signal-

transducing proteins.

Post-translational Ligand-binding increases phosphorylation on tyrosine residues and promotes its association with

modifications the p85 subunit of phosphatidylinositol 3-kinase.

Cellular localization Secreted and Cell membrane.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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