# ANTIBODIES ONLINE

# Datasheet for ABIN967410 anti-ABL1 antibody

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## Overview

Quantity:	0.1 mg	
Target:	ABL1	
Reactivity:	Human, Mouse	
Host:	Mouse	
Clonality:	Monoclonal	
Conjugate:	This ABL1 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunofluorescence (IF), Immunoprecipitation (IP)	

# Product Details

Brand:	BD Pharmingen™	
Immunogen:	Recombinant Mouse Abl Gag Fusion Protein	
Clone:	8E9	
lsotype:	lgG1	
Cross-Reactivity:	Mouse (Murine)	
Characteristics:	<ol> <li>Since applications vary, each investigator should titrate the reagent to obtain optimal results.</li> <li>Please refer to us for technical protocols.</li> <li>Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.</li> </ol>	
Purification:	The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.	

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Target Details	
Target:	ABL1
Alternative Name:	Abl (ABL1 Products)
Background:	The proto-oncogene c-abl was first isolated from the mouse genome as a gene with similarity
	to the v-abl oncogene of Abelson murine leukemia virus. The c-abl gene encodes a protein
	tyrosine kinase that is localized in the cytoplasm and nucleus. The c-abl protein shares several
	common features with other cytoplasmic tyrosine kinases, including the src-homology domains
	2 (SH2) and 3 (SH3). The SH2 domain is believed to bind specifically to tyrosine residues of
	other proteins. The function of the SH3 domain is still unclear. Unique to the c-abl tyrosine
	kinase is a large C-terminal segment which seems to be essential for its biological function,
	since mice homozygous for a C-terminal deletion of c-abl have multiple defects at birth. The C-
	terminal fragment of c-abl contains a DNA-binding domain, and the DNA-binding affinity of this
	domain seems to be regulated by phosphorylation of critical serine/threonine residues. The c-
	abl proto-oncogene can be activated in a variety of ways. For example, in Philadelphia
	chromosome (Ph1)-positive leukemias the c-abl proto-oncogene on chromosome 9 becomes
	fused to the bcr gene on chromosome 22, and bcr-abl fusion proteins are produced. Ph1-
	positive cells express either the a-typical 210 kDa bcr-abl fusion protein or a smaller 185 kDa
	bcr-abl fusion protein. The bcr sequences activate the c-abl tyrosine kinase by deregulating its
	expression, and actin filament-binding function associated with c-abl is also activated.
	Expression of bcr-abl fusion proteins in vitro leads to transformation of pre-B lymphoid cells
	supporting their role as an oncogene. The phosphorylated form of c-abl is observed at ${\sim}145$
	kDa on SDS/PAGE. The 8E9 clone has been reported to react with an epitope in the tyrosine
	kinase domain of murine abl proteins [Wang et al.]. This antibody is routinely tested by western
	blot analysis.
Molecular Weight:	145 kDa

Pathways:Apoptosis, Regulation of Muscle Cell Differentiation, Platelet-derived growth Factor ReceptorSignaling, Lipid Metabolism

# **Application Details**

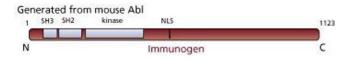
Comment:	Related Products: ABIN967389
Restrictions:	For Research Use only
Handling	
Format:	Liquid

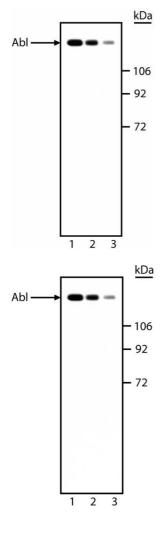
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Concentration:	0.5 mg/mL
Buffer:	Aqueous buffered solution containing $\leq 0.09$ % sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C
Storage Comment:	Store undiluted at 4° C.
Publications	
Product cited in:	Guo, Lian, Xian, Lee, Deisseroth, Stass, Champlin, Talpaz, Wang, Arlinghaus: "BCR-ABL protein
	expression in peripheral blood cells of chronic myelogenous leukemia patients undergoing
	therapy." in: <b>Blood</b> , Vol. 83, Issue 12, pp. 3629-37, (1994) (PubMed).
	Guo, Hirsch-Ginsberg, Xian, Stass, Champlin, Giralt, McCredie, Campbell, Arlinghaus: "Acute
	lymphoid leukemia molecular phenotype in a patient with benign-phase chronic myelogenous
	leukemia." in: Hematologic pathology, Vol. 7, Issue 2, pp. 91-106, (1993) (PubMed).
	Guo, Wang, Arlinghaus: "Detection of BCR-ABL proteins in blood cells of benign phase chronic
	myelogenous leukemia patients." in: <b>Cancer research</b> , Vol. 51, Issue 11, pp. 3048-51, (1991) (
	PubMed).
	Wang: "Negative regulation of c-abl tyrosine kinase by its variable N-terminal amino acids." in:
	Oncogene research, Vol. 3, Issue 3, pp. 293-8, (1989) (PubMed).
	Kipreos, Lee, Wang: "Isolation of temperature-sensitive tyrosine kinase mutants of v-abl
	oncogene by screening with antibodies for phosphotyrosine." in: Proceedings of the National
	Academy of Sciences of the United States of America, Vol. 84, Issue 5, pp. 1345-9, (1987) (
	PubMed).

#### Image 1.





#### Western Blotting

**Image 2.** Western blot analysis of Abl. Lysate from A-431 human epidermal carcinoma cells was probed with anti-Abl (clone 8E9, ABIN967410) and titrated between 1 myg/ml and 0.04 myg/ml (lanes 1-3). Abl is identified at ~145 kDa.

### Western Blotting

Image 3.

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