



# COTL1 rabbit pAb

Cat No.:ES17297

For research use only

## Overview

<b>Product Name</b>	COTL1 rabbit pAb
<b>Host species</b>	Rabbit
<b>Applications</b>	WB
<b>Species Cross-Reactivity</b>	Human; Mouse;Rat
<b>Recommended dilutions</b>	WB 1: 500-2000
<b>Immunogen</b>	Synthesized peptide derived from human COTL1 AA range: 29-79
<b>Specificity</b>	This antibody detects endogenous levels of COTL1 at Human/Mouse/Rat
<b>Formulation</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Storage</b>	Store at -20°C. Avoid repeated freeze-thaw cycles.
<b>Protein Name</b>	COTL1
<b>Gene Name</b>	COTL1 CLP
<b>Cellular localization</b>	Cytoplasm . Cytoplasm, cytoskeleton . Nucleus .
<b>Purification</b>	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
<b>Clonality</b>	Polyclonal
<b>Concentration</b>	1 mg/ml
<b>Observed band</b>	
<b>Human Gene ID</b>	23406
<b>Human Swiss-Prot Number</b>	Q14019
<b>Alternative Names</b>	
<b>Background</b>	This gene encodes one of the numerous actin-binding proteins which regulate the actin cytoskeleton. This protein binds F-actin, and also interacts with 5-lipoxygenase, which is the first committed enzyme in leukotriene biosynthesis. Although this gene has been reported to map to chromosome 17 in the Smith-Magenis syndrome region, the best alignments for this gene are to chromosome 16. The Smith-Magenis syndrome

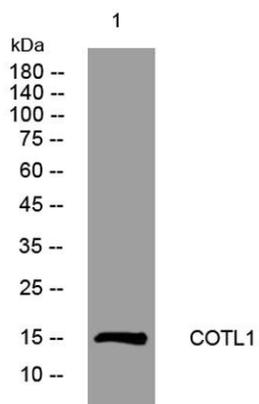




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region is the site of two related pseudogenes.  
[provided by RefSeq, Jul 2008],

Western blot analysis of lysates from HCT116 cells,  
primary antibody was diluted at 1:1000, 4° over night



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