## ARC (D7Q3G) Rabbit mAb (PE Conjugate)



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For Research Use Only. Not for Use in Diagnostic Procedures.

Source/Isotype: Applications: Reactivity: Sensitivity: **UniProt ID:** Entrez-Gene Id: FC-FP Н Endogenous Rabbit IgG #O60936 8996 **Product Usage** Application Dilution Information 1:50

Flow Cytometry (Fixed/Permeabilized)

Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the **Storage** antibody. Protect from light. Do not freeze.

Specificity / Sensitivity ARC (D7Q3G) Rabbit mAb (PE Conjugate) recognizes endogenous levels of total ARC protein.

Source / Purification Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro125 of human ARC protein, specific to a region encoded by isoform 2 of the

NOL3 gene.

This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct **Product Description** 

flow cytometric analysis in human cells. This antibody is expected to exhibit the same species cross-

reactivity as the unconjugated ARC (D7Q3G) Rabbit mAb #38916.

**Background** 

Apoptosis repressor with caspase recruitment domain (ARC), also independently identified as muscleenriched cytoplasmic protein (MYP), is a CARD domain protein that regulates apoptosis (1). The ARC protein CARD domain is highly homologous to those in other cell death regulators, including caspase-2, caspase-9, RAIDD, and Apaf-1 (2). The NOL3 gene encodes both the cytoplasmic ARC protein and a 30 kDa nucleolar protein (Nop30) that is involved in RNA splicing. ARC is encoded from isoform 2 of NOL3, while isoform 1 produced by alternative splicing encodes Nop30. Both ARC and Nop30 proteins share common amino-terminal sequences (3). Research studies show that ARC can bind to caspase-8 and caspase-2 and inhibit apoptosis through extrinsic pathways that involve the receptor proteins Fas, TNFR1, and DR3 (1). Additional research indicates that the ARC anti-apoptotic mechanism may include both intrinsic (mitochondrial) and extrinsic (death receptor) pathways (4). In addition to binding caspases, ARC can disrupt the interaction with the death domains of Fas and FADD, which inhibits death-inducing signaling complex (DISC) assembly. The CARD domain of ARC can inhibit intrinsic apoptosis through binding to the pro-apoptotic Bax protein (5). Phosphorylation of ARC at Thr149 by CK2 is required for targeting of ARC to the mitochondria (6). ARC is able to suppress necroptosis, a programmed pathway of necrosis triggered by blocking the recruitment of RIP1 to TNFR1 (7). Expression of ARC protein is predominantly seen in terminally differentiated cells under normal conditions and is markedly induced in a variety of cancers including pancreatic, colorectal, breast, lung, glioblastoma, liver, kidney, melanoma, and acute myeloid leukemia (1, 8-12).

## **Background References**

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- 3. Stoss, O. et al. (1999) J Biol Chem 274, 10951-62.
- 4. Nam, Y.J. et al. (2004) Mol Cell 15, 901-12.
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- 8. Mercier, I. et al. (2008) Cell Cycle 7, 1640-7.
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## **Species Reactivity**

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

1/1/24, 2:36 PM ARC (D7Q3G) Rabbit mAb (PE Conjugate) (#89210) Datasheet Without Images Cell Signaling Technology

**Applications Key** 

FC-FP: Flow Cytometry (Fixed/Permeabilized)

**Cross-Reactivity Key** 

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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