

Product datasheet for TA380680S

RAF1 Rabbit Polyclonal Antibody

Product data:

OriGene Technologies, Inc.

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Product Type:	Primary Antibodies
Applications:	IP, WB
Recommended Dilution:	WB,1:500 - 1:2000 IP,1:50 - 1:100
Reactivity:	Human
Modifications:	Phospho S259
Host:	Rabbit
lsotype:	IgG
Clonality:	Polyclonal
Immunogen:	A synthetic phosphorylated peptide around S259 of human Phospho-Raf1-S259 (NP_002871.1).
Formulation:	Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.
Concentration:	lot specific
Purification:	Affinity purification
Conjugation:	Unconjugated
Storage:	Store at -20°C. Avoid freeze / thaw cycles.
Stability:	Shelf life: one year from despatch.
Predicted Protein Size:	73kDa/75kDa
Gene Name:	Raf-1 proto-oncogene, serine/threonine kinase
Database Link:	<u>Entrez Gene 5894 Human</u> <u>P04049</u>



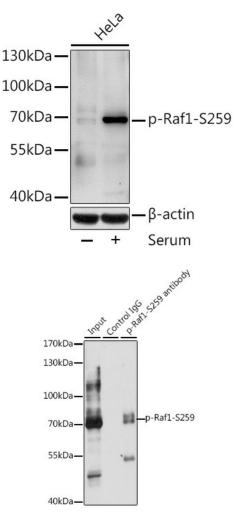
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GRIGENE RAF1 Rabbit Polyclonal Antibody – TA380680S

Background:This gene is the cellular homolog of viral raf gene (v-raf). The encoded protein is a MAP kinase
kinase kinase (MAP3K), which functions downstream of the Ras family of membrane
associated GTPases to which it binds directly. Once activated, the cellular RAF1 protein can
phosphorylate to activate the dual specificity protein kinases MEK1 and MEK2, which in turn
phosphorylate to activate the serine/threonine specific protein kinases, ERK1 and ERK2.
Activated ERKs are pleiotropic effectors of cell physiology and play an important role in the
control of gene expression involved in the cell division cycle, apoptosis, cell differentiation
and cell migration. Mutations in this gene are associated with Noonan syndrome 5 and
LEOPARD syndrome 2.

Synonyms: C-RAF; CRAF; NS5; RAF; Raf-1

Product images:



Western blot analysis of extracts of HeLa cells, using Phospho-Raf1-S259 antibody ([TA380680]) at 1:1000 dilution. HeLa cells were treated by 10% FBS after serum-starvation overnight. |Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) at 1:10000 dilution. |Lysates/proteins: 25ug per lane. |Blocking buffer: 3% BSA.

Immunoprecipitation analysis of 200ug extracts of HeLa cells, using 3 ug Phospho-Raf1-S259 pAb ([TA380680]). Western blot was performed from the immunoprecipitate using Phospho-Raf1-S259 pAb ([TA380680]) at a dilition of 1:1000. HeLa cells were treated by 10% FBS at 37°C for 30 minutes after serum-starvation overnight.

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