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## Datasheet for ABIN185018 anti-ECT2 antibody (N-Term)

3 Images



### Overview

Quantity:	100 µg
Target:	ECT2
Binding Specificity:	N-Term
Reactivity:	Human
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This ECT2 antibody is un-conjugated
Application:	ELISA, Flow Cytometry (FACS), Immunofluorescence (IF)

## Product Details

Purpose:	ECT2
Immunogen:	Peptide with sequence AENSVLTSTTGRT-C, from the N Terminus of the protein sequence according to NP_060568.3.
Sequence:	AENSVLTSTT GRT
Isotype:	lgG
Specificity:	This antibody is expected to recognise NP_001245244.1 (iso a), NP_060568.3 (iso b), NP_001336023.1 (iso c), NP_001336026.1 (iso d), NP_001336030.1 (iso e), NP_001336031.1 (iso f), NP_001336032.1 (iso g)
Cross-Reactivity:	Human
Purification:	Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity

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## Product Details

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Grade:

#### Verified

## Target Details

Target:	ECT2
Alternative Name:	ECT2 (ECT2 Products)
Background:	ECT2, epithelial cell transforming sequence 2 oncogene, FLJ10461, epithelial cell transforming sequence 2 oncogene protein
Gene ID:	1894
NCBI Accession:	NP_060568
Pathways:	Neurotrophin Signaling Pathway, Cell-Cell Junction Organization

## Application Details

Application Notes:	Peptide ELISA: antibody detection limit dilution 1:128000.
Comment:	Immunofluorescence: Strong expression of the protein seen in the nuclei and cytoplasm of
	A431 and U2OS cells. Recommended concentration: 10µg/ml.
	Flow Cytometry: Flow cytometric analysis of A431 cells. Recommended conc
Restrictions:	For Research Use only

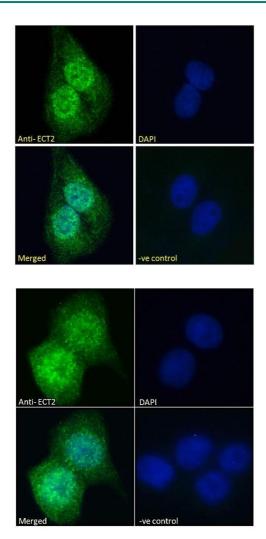
## Handling

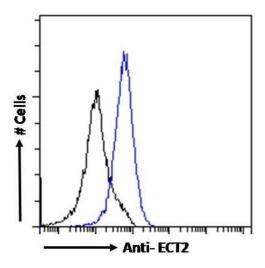
Format:	Liquid
Concentration:	0.5 mg/mL
Buffer:	Supplied at 0.5 mg/mL in Tris saline, 0.02 % sodium azide, pH 7.3 with 0.5 % bovine serum albumin.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Handling Advice:	Minimize freezing and thawing.
Storage:	-20 °C

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Aliquot and store at -20°C, with minimal freeze/thawing. A working aliquot may be refrigerated at 4°C for a few weeks and still remain viable.

#### Images





#### Immunofluorescence

**Image 1.** (ABIN185018) Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15 % Triton. Primary incubation 1hr (10  $\mu$ g/mL) followed by Alexa Fluor 488 secondary antibody (2  $\mu$ g/mL), showing strong nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10  $\mu$ g/mL) followed by Alexa Fluor 488 secondary antibody (2  $\mu$  g/mL).

#### Immunofluorescence

**Image 2.** (ABIN185018) Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15 % Triton. Primary incubation 1hr (10  $\mu$ g/mL) followed by Alexa Fluor 488 secondary antibody (2  $\mu$ g/mL), showing strong nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10  $\mu$ g/mL) followed by Alexa Fluor 488 secondary antibody (2  $\mu$  g/mL).

#### **Flow Cytometry**

**Image 3.** (ABIN185018) Flow cytometric analysis of paraformaldehyde fixed A431 cells (blue line), permeabilized with 0.5 % Triton. Primary incubation 1hr ( $10 \mu g/mL$ ) followed by Alexa Fluor 488 secondary antibody ( $1 \mu g/mL$ ). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.

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