

**FOXO3 Antibody (Center)**  
**Affinity Purified Rabbit Polyclonal Antibody (Pab)**  
**Catalog # AP14166c****Specification**

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**FOXO3 Antibody (Center) - Product Information**

Application	IF, WB, IHC-P,E
Primary Accession	<a href="#">O43524</a>
Other Accession	<a href="#">O9WVH4</a> , <a href="#">NP_963853.1</a> , <a href="#">NP_001446.1</a>
Reactivity	Human
Predicted	Mouse
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	71277
Antigen Region	224-253

**FOXO3 Antibody (Center) - Additional Information****Gene ID** 2309**Other Names**

Forkhead box protein O3, AF6q21 protein, Forkhead in rhabdomyosarcoma-like 1, FOXO3, FKHL1, FOXO3A

**Target/Specificity**

This FOXO3 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 224-253 amino acids from the Central region of human FOXO3.

**Dilution**IF~~1:10~50  
WB~~1:1000  
IHC-P~~1:10~50  
E~~Use at an assay dependent concentration.**Format**

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

**Storage**

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions**

FOXO3 Antibody (Center) is for research use only and not for use in diagnostic or therapeutic procedures.

**FOXO3 Antibody (Center) - Protein Information**

**Name** FOXO3 ([HGNC:3821](#))

**Function** Transcriptional activator that recognizes and binds to the DNA sequence 5'-[AG]TAAA[TC]A-3' and regulates different processes, such as apoptosis and autophagy (PubMed:[10102273](#), PubMed:[16751106](#), PubMed:[21329882](#), PubMed:[30513302](#)). Acts as a positive regulator of autophagy in skeletal muscle: in starved cells, enters the nucleus following dephosphorylation and binds the promoters of autophagy genes, such as GABARAP1L, MAP1LC3B and ATG12, thereby activating their expression, resulting in proteolysis of skeletal muscle proteins (By similarity). Triggers apoptosis in the absence of survival factors, including neuronal cell death upon oxidative stress (PubMed:[10102273](#), PubMed:[16751106](#)). Participates in post-transcriptional regulation of MYC: following phosphorylation by MAPKAPK5, promotes induction of miR-34b and miR-34c expression, 2 post-transcriptional regulators of MYC that bind to the 3'UTR of MYC transcript and prevent its translation (PubMed:[21329882](#)). In response to metabolic stress, translocates into the mitochondria where it promotes mtDNA transcription (PubMed:[23283301](#)). In response to metabolic stress, translocates into the mitochondria where it promotes mtDNA transcription. Also acts as a key regulator of chondrogenic commitment of skeletal progenitor cells in response to lipid availability: when lipids levels are low, translocates to the nucleus and promotes expression of SOX9, which induces chondrogenic commitment and suppresses fatty acid oxidation (By similarity). Also acts as a key regulator of regulatory T-cells (Treg) differentiation by activating expression of FOXP3 (PubMed:[30513302](#)).

**Cellular Location**

Cytoplasm, cytosol. Nucleus Mitochondrion matrix. Mitochondrion outer membrane; Peripheral membrane protein; Cytoplasmic side. Note=Retention in the cytoplasm contributes to its inactivation (PubMed:[10102273](#), PubMed:[15084260](#), PubMed:[16751106](#)). Translocates to the nucleus upon oxidative stress and in the absence of survival factors (PubMed:[10102273](#), PubMed:[16751106](#)) Translocates from the cytosol to the nucleus following dephosphorylation in response to autophagy-inducing stimuli (By similarity). Translocates in a AMPK-dependent manner into the mitochondrion in response to metabolic stress (PubMed:[23283301](#), PubMed:[29445193](#)). Serum deprivation increases localization to the nucleus, leading to activate expression of SOX9 and subsequent chondrogenesis (By similarity). {ECO:0000250|UniProtKB:Q9WVH4, ECO:0000269|PubMed:[10102273](#), ECO:0000269|PubMed:[15084260](#), ECO:0000269|PubMed:[16751106](#), ECO:0000269|PubMed:[23283301](#), ECO:0000269|PubMed:[29445193](#)}

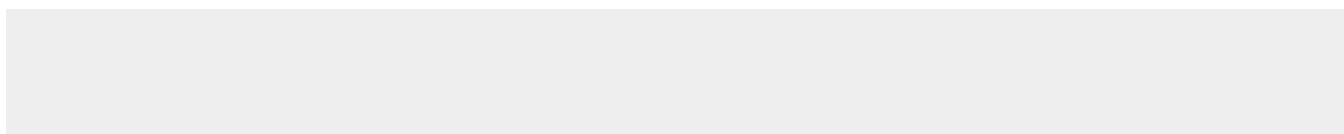
**Tissue Location**

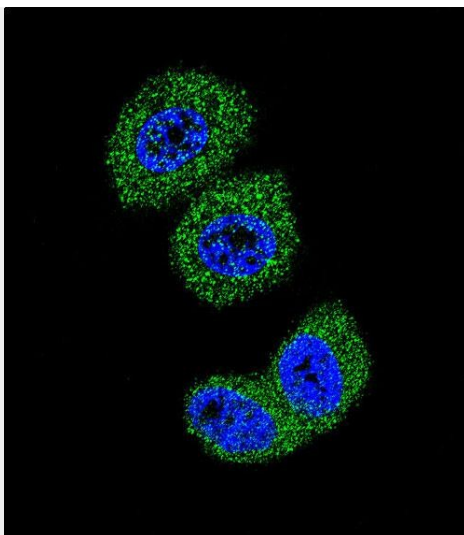
Ubiquitous..

**FOXO3 Antibody (Center) - Protocols**

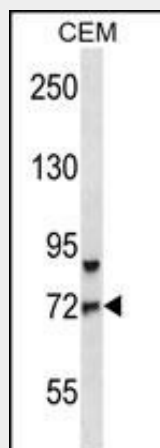
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

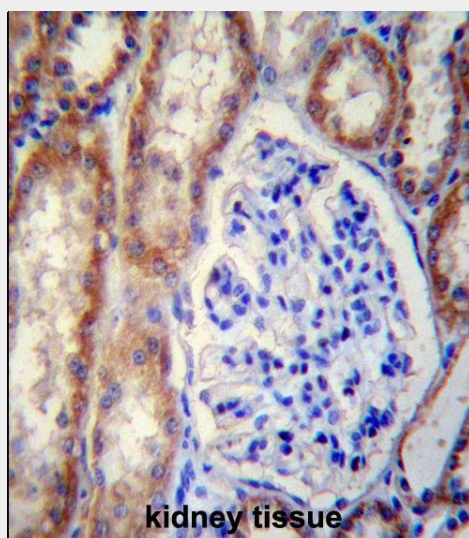
**FOXO3 Antibody (Center) - Images**



Confocal immunofluorescent analysis of FOXO3 Antibody (Center)(Cat#AP14166c) with MCF-7 cell followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green).DAPI was used to stain the cell nuclear (blue).



FOXO3 Antibody (Center) (Cat. #AP14166c) western blot analysis in CEM cell line lysates (35ug/lane).This demonstrates the FOXO3 antibody detected the FOXO3 protein (arrow).



FOXO3 Antibody (Center) (Cat. #AP14166c)immunohistochemistry analysis in formalin fixed and

paraffin embedded human kidney tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of ANKS1B FOXO3 Antibody (Center) for immunohistochemistry. Clinical relevance has not been evaluated.

#### **FOXO3 Antibody (Center) - Background**

This gene belongs to the forkhead family of transcription factors which are characterized by a distinct forkhead domain. This gene likely functions as a trigger for apoptosis through expression of genes necessary for cell death. Translocation of this gene with the MLL gene is associated with secondary acute leukemia. Alternatively spliced transcript variants encoding the same protein have been observed.

#### **FOXO3 Antibody (Center) - References**

Zhuo de, X., et al. J. Biol. Chem. 285(41):31491-31501(2010)  
Tudzarova, S., et al. EMBO J. 29(19):3381-3394(2010)  
Shimada, M., et al. Hum. Genet. 128(4):433-441(2010)  
Mikse, O.R., et al. Cancer Res. 70(15):6205-6215(2010)  
Chen, J., et al. PLoS ONE 5 (8), E12293 (2010) :