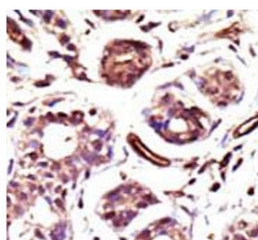


Ubiquitin-protein ligase E3A Antibody / UBE3A (F54766)

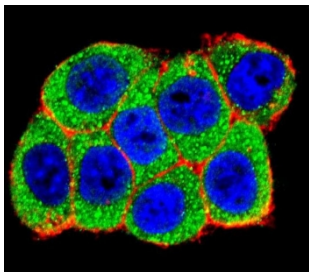
Catalog No.	Formulation	Size
F54766-0.4ML	In 1X PBS, pH 7.4, with 0.09% sodium azide	0.4 ml
F54766-0.08ML	In 1X PBS, pH 7.4, with 0.09% sodium azide	0.08 ml

[Bulk quote request](#)

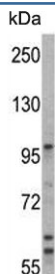
Availability	1-3 business days
Species Reactivity	Human
Format	Purified
Host	Rabbit
Clonality	Polyclonal (rabbit origin)
Isotype	Rabbit Ig
Purity	Purified
UniProt	Q05086
Localization	Cytoplasmic, nuclear
Applications	Flow Cytometry : 1:10-1:50 (1x10 ⁶ cells) Immunofluorescence : 1:10-1:50 Immunohistochemistry (FFPE) : 1:10-1:50 Western Blot : 1:500-1:1000
Limitations	This Ubiquitin-protein ligase E3A antibody is available for research use only.



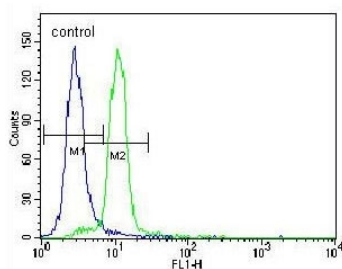
IHC testing of FFPE human breast cancer tissue with Ubiquitin-protein ligase E3A antibody. HIER: steam section in pH6 citrate buffer for 20 min and allow to cool prior to staining.



Immunofluorescent staining of human HeLa cells with Ubiquitin-protein ligase E3A antibody (green), DAPI nuclear stain (blue) and anti-Actin (red).



Western blot testing of human T-47D cell lysate with Ubiquitin-protein ligase E3A antibody. Predicted molecular weight ~100 kDa.



Flow cytometry testing of human HeLa cells with Ubiquitin-protein ligase E3A antibody; Blue=isotype control, Green= Ubiquitin-protein ligase E3A antibody.

Description

UBE3A interacts with the E6 protein of the cancer-associated human papillomavirus types 16 and 18. The E6/E6-AP complex binds to and targets the p53 tumor-suppressor protein for ubiquitin-mediated proteolysis. It is an E3 ubiquitin-protein ligase which accepts ubiquitin from an E2 ubiquitin-conjugating enzyme in the form of a thioester and then directly transfers the ubiquitin to targeted substrates. It can target itself for ubiquitination in vitro and efficiently promotes its own degradation in vivo. It appears that only unmodified E6-AP molecules can bind efficiently to p53 in the presence of the HPV E6 oncoprotein. UBE3A binds UBQLN1 and UBQLN2. Defects in UBE3A are a cause of Angelman syndrome (AS) [MIM:105830]; also known as 'happy puppet syndrome'. AS is characterized by features of severe motor and intellectual retardation, microcephaly, ataxia, frequent jerky limb movements and flapping of the arms and hands, hypotonia, hyperactivity, hypopigmentation, seizures, absence of speech, frequent smiling and episodes of paroxysmal laughter, and an unusual facies characterized by macrostomia, a large mandible and open-mouthed expression, a great propensity for protruding the tongue ('tongue thrusting'), and an occipital groove. UBE3A contains 1 HECT-type E3 ubiquitin-protein ligase domain.

Application Notes

The stated application concentrations are suggested starting points. Titration of the Ubiquitin-protein ligase E3A antibody may be required due to differences in protocols and secondary/substrate sensitivity.

Immunogen

A portion of amino acids 836-865 from the human protein was used as the immunogen for the Ubiquitin-protein ligase E3A antibody.

Storage

Aliquot the Ubiquitin-protein ligase E3A antibody and store frozen at -20°C or colder. Avoid repeated freeze-thaw cycles.

