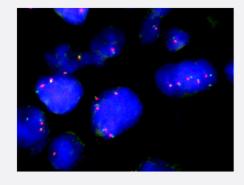


UBE3A/CEN15q FISH Probe

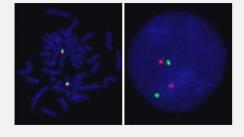
Catalog # FG0025 Size 200 uL, 100 uL

Applications



Fluorescent *In Situ* Hybridization (Formalin/PFA-fixed paraffin-embedded sections)

Human breast cancer (FFPE) stained with UBE3A/CEN15q FISH Probe. Human breast cancer showed no UBE3A gene amplification



Hybridization position of the probes on the chromosome.

Hybridization position of the probes on the chromosome.

Specification

Product Description

Labeled FISH probes for identification of gene amplification using Fluorescent In Situ Hybridization T echnique. (<u>Technology</u>).



Product Information

Probe 1	Name: UBE3A Size: Approximately 370kb Fluorophore: Texas Red Location: 15q11.2
Probe 2	Name: CEN15q Size: Approximately 680kb Fluorophore: FITC Location: 15q11.2
Probe Gap	The gap between two probes is approximately 5,700 kb.
Origin	Human
Source	Genomic DNA
Reactivity	Human
Form	Liquid
Notice	We strongly recommend the customer to use FFPE FISH PreTreatment Kit 1 (Catalog #: KA2375 or KA2691) for the pretreatment of Formalin-Fixed Paraffin-Embedded (FFPE) tissue sections.
Regulation Status	For research use only (RUO)
Quality Control Testing	Representative images of normal human cell (lymphocyte) stain with the dual color FISH probe. The I eft image is chromosomes at metaphase, and the right image is an interphase nucleus.
Supplied Product	DAPI Counterstain (1500 ng/mL) 125 uL for each 100 uL FISH Probe
Storage Instruction	Store at 4°C in the dark.
Note	Hybridization position of the probes on the chromosome. Hybridization position of the probes on the chromosome.

Applications

• Fluorescent In Situ Hybridization (Cell)

Protocol Download

• Fluorescent *In Situ* Hybridization (Formalin/PFA-fixed paraffin-embedded sections)

Human breast cancer (FFPE) stained with UBE3A/CEN15q FISH Probe. Human breast cancer showed no UBE3A gene amplification

Protocol Download



Gene Info — UBE3A	
Entrez GenelD	7337
Gene Name	UBE3A
Gene Alias	ANCR, AS, E6-AP, EPVE6AP, FLJ26981, HPVE6A
Gene Description	ubiquitin protein ligase E3A
Omim ID	<u>105830</u> <u>601623</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes an E3 ubiquitin-protein ligase, part of the ubiquitin protein degradation system . This imprinted gene is maternally expressed in brain and biallelically expressed in other tissues. Maternally inherited deletion of this gene causes Angelman Syndrome, characterized by severe m otor and intellectual retardation, ataxia, hypotonia, epilepsy, absence of speech, and characteristic facies. The protein also interacts with the E6 protein of human papillomavirus types 16 and 18, resulting in ubiquitination and proteolysis of tumor protein p53. Alternative splicing of this gene results in three transcript variants encoding three isoforms with different N-termini. Additional transcript variants have been described, but their full length nature has not been determined. [provided by RefSeq
Other Designations	CTCL tumor antigen se37-2 human papilloma virus E6-associated protein oncogenic protein-ass ociated protein E6-AP

Pathway

• <u>Ubiquitin mediated proteolysis</u>

Disease

- Attention Deficit Disorder with Hyperactivity
- Autistic Disorder
- Genetic Predisposition to Disease
- Mental Disorders
- NARP
- Tuberculosis