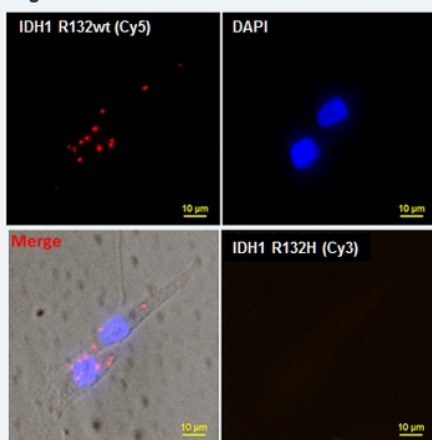


mutaFISH™ IDH1 R132H R132wt IDH2 R172K R172wt RNA Probes

Catalog # FP0012 Size 1 Probe Set

Applications

Fig. 1

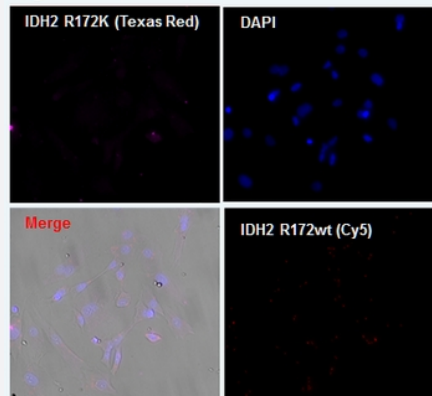


mutation specific, Fluorescence *In Situ* Hybridization (Cells)

Fig.1 mutaFISH™ staining was performed *in situ* in human SH-SY5Y cells. IDH1 R123H point mutation was not detected via orange signal (Cy3), IDH1 R123 wildtype was detected via red signal.

Fig.2 mutaFISH™ staining was performed *in situ* in human SH-SY5Y cells. IDH2 R172K point mutation was detected via purple signal (Texas Red), IDH2 R172 wildtype was detected via red signal.

Fig. 2



Specification

Product Description

mutaFISH™ IDH1 R132H R132wt IDH2 R172K R172wt RNA Probes is designed to detect human IDH1 R132H gene mutation and IDH2 R172K gene mutation on single strand RNA in cells using padlock probe and *in situ* rolling-circle amplification technology.

Reactivity

Human

Supplied Product

Content:

1. RT IDH1 R132 Primer
2. RT IDH2 R172 Primer
3. mutaFISH™ IDH1 R132H RNA Probe
4. mutaFISH™ IDH1 R132wt RNA Probe
5. mutaFISH™ IDH2 R172K RNA Probe
6. mutaFISH™ IDH2 R172wt RNA Probe
7. Detection Probe-Aqua 431
8. Detection Probe-Texas Red X
9. Detection Probe-6-HEX

Technology

[mutaFISH™ \(mutation-specific Fluorescence *In Situ* Hybridization\)](#)

Comparison

[FISH Probes vs mutaFISH™ Probes](#)

Fluorophore

Aqua 431 (Excitation Peak (nm): 431; Emission Peak (nm): 480)
Texas Red X (Excitation Peak (nm): 595; Emission Peak 613)
6-HEX (Excitation Peak (nm): 533; Emission Peak (nm): 559)

Probe Position

Regulatory Status

For research use only (RUO)

Storage Instruction

Store at -20°C.
Aliquot to avoid repeated freezing and thawing.

Note

We recommend mutaFISH™ RNA Accessory Kit (Catalog #: [KA4915](#)) which provides necessary reagents and enzymes for *in situ* reverse transcription, RNA digestion, mutaFISH™ hybridization, ligation and amplification prior to mutaFISH™.

Video

Applications

- mutation specific, Fluorescence *In Situ* Hybridization (Cells)

Fig.1 mutaFISH™ staining was performed *in situ* in human SH-SY5Y cells. IDH1 R123H point mutation was not detected via orange signal (Cy3), IDH1 R123 wildtype was detected via red signal.

Fig.2 mutaFISH™ staining was performed *in situ* in human SH-SY5Y cells. IDH2 R172K point mutation was detected via purple signal (Texas Red), IDH2 R172 wildtype was detected via red signal.

Gene Info — IDH1

Entrez GeneID [3417](#)

Gene Name IDH1

Gene Alias IDCDC, IDH, IDP, IDPC, PICD

Gene Description isocitrate dehydrogenase 1 (NADP+), soluble

Omim ID [147700](#)

Gene Ontology [Hyperlink](#)

Gene Summary

Isocitrate dehydrogenases catalyze the oxidative decarboxylation of isocitrate to 2-oxoglutarate. These enzymes belong to two distinct subclasses, one of which utilizes NAD(+) as the electron acceptor and the other NADP(+). Five isocitrate dehydrogenases have been reported: three NAD(+)-dependent isocitrate dehydrogenases, which localize to the mitochondrial matrix, and two NADP(+)-dependent isocitrate dehydrogenases, one of which is mitochondrial and the other predominantly cytosolic. Each NADP(+)-dependent isozyme is a homodimer. The protein encoded by this gene is the NADP(+)-dependent isocitrate dehydrogenase found in the cytoplasm and peroxisomes. It contains the PTS-1 peroxisomal targeting signal sequence. The presence of this enzyme in peroxisomes suggests roles in the regeneration of NADPH for intraperoxisomal reductions, such as the conversion of 2, 4-dienoyl-CoAs to 3-enoyl-CoAs, as well as in peroxisomal reactions that consume 2-oxoglutarate, namely the alpha-hydroxylation of phytanic acid. The cytoplasmic enzyme serves a significant role in cytoplasmic NADPH production. [provided by RefSeq]

Other Designations NADP+-specific ICDH|NADP-dependent isocitrate dehydrogenase, cytosolic|NADP-dependent isocitrate dehydrogenase, peroxisomal|oxalosuccinate decarboxylase

Pathway

- [Biosynthesis of alkaloids derived from histidine and purine](#)
- [Biosynthesis of alkaloids derived from ornithine](#)
- [Biosynthesis of alkaloids derived from shikimate pathway](#)
- [Biosynthesis of alkaloids derived from terpenoid and polyketide](#)

- [Biosynthesis of phenylpropanoids](#)
- [Biosynthesis of plant hormones](#)
- [Biosynthesis of terpenoids and steroids](#)
- [Citrate cycle \(TCA cycle\)](#)
- [Glutathione metabolism](#)
- [Metabolic pathways](#)
- [Reductive carboxylate cycle \(CO₂ fixation\)](#)

Disease

- [Adenoma](#)
- [Astrocytoma](#)
- [Blast Crisis](#)
- [Brain Neoplasms](#)
- [Carcinoma](#)
- [Chronic Disease](#)
- [Cleft Lip](#)
- [Cleft Palate](#)
- [Disease Progression](#)
- [Disease Susceptibility](#)
- [Genetic Predisposition to Disease](#)
- [Glioblastoma](#)
- [Glioma](#)
- [Hematologic Diseases](#)
- [HIV Infections](#)
- [Leukemia](#)
- [Lung Neoplasms](#)

- [Lymphoma](#)
- [Melanoma](#)
- [Monosomy](#)
- [Myelodysplastic Syndromes](#)
- [Myeloproliferative Disorders](#)
- [Neoplasm Metastasis](#)
- [Nervous System Neoplasms](#)
- [Neuroectodermal Tumors](#)
- [Oligodendroglioma](#)
- [Pancreatic cancer](#)
- [Pancreatic Neoplasms](#)
- [Polycythemia Vera](#)
- [Primary Myelofibrosis](#)
- [Recurrence](#)
- [Skin Neoplasms](#)
- [Supratentorial Neoplasms](#)
- [Thrombocythemia](#)
- [Thyroid Neoplasms](#)