

# SHOX(Texas Red)/CENXp(FITC) FISH Probe

Catalog # FA0615      Size 200 uL

## Specification

<b>Product Description</b>	Made to order FISH probes for identification of gene amplification using Fluorescent In Situ Hybridization Technique. ( <a href="#">Technology</a> ).
<b>Origin</b>	Human
<b>Source</b>	Genomic DNA
<b>Reactivity</b>	Human
<b>Notice</b>	We <b>strongly recommend</b> the customer to use FFPE FISH PreTreatment Kit 1 (Catalog #: <a href="#">KA2375</a> or <a href="#">KA2691</a> ) for the pretreatment of Formalin-Fixed Paraffin-Embedded (FFPE) tissue sections.
<b>Regulation Status</b>	For research use only (RUO)
<b>Supplied Product</b>	DAPI Counterstain (1500 ng/mL ) 250 uL
<b>Storage Instruction</b>	Store at 4°C in the dark.

## Applications

- Fluorescent In Situ Hybridization (Cell)

[Protocol Download](#)

## Gene Info — SHOX

<b>Entrez GeneID</b>	<a href="#">6473</a>
<b>Gene Name</b>	SHOX
<b>Gene Alias</b>	GCFX, PHOG, SHOXY, SS
<b>Gene Description</b>	short stature homeobox

**Omim ID** [127300 249700 312865 400020 604271](#)

**Gene Ontology** [Hyperlink](#)

**Gene Summary** This gene belongs to the paired homeobox family and is located in the pseudoautosomal region 1 (PAR1) of X and Y chromosomes. Defects in this gene are associated with idiopathic growth retardation and in the short stature phenotype of Turner syndrome patients. This gene is highly conserved across species from mammals to fish to flies. Alternatively spliced transcript variants encoding different isoforms have been noted for this gene. [provided by RefSeq]

**Other Designations** OTTHUMP00000022821|OTTHUMP00000022822|growth control factor, X-linked|pseudoautosomal homeobox-containing osteogenic

## Disease

- [Growth Disorders](#)