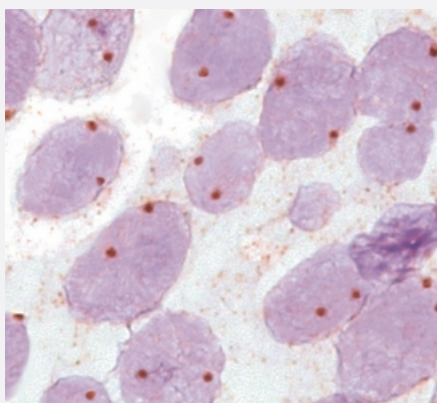


# MYCN CISH Probe

Catalog # CG0022      Size 400 uL

## Applications



### Chromogenic *In Situ* Hybridization (FFPE Tissue)

Normal nuclei each with two MYCN signals

## Specification

<b>Product Description</b>	MYCN CISH Probe is designed for the qualitative detection human MYCN gene in formalin-fixed, paraffin-embedded tissue or cells by chromogenic <i>in situ</i> hybridization (CISH).
<b>Reactivity</b>	Human
<b>Recommend Usage</b>	The product is ready-to-use. No reconstitution, mixing, or dilution is required. Bring probe to room temperature (18-25°C) and mix briefly before use.
<b>Supplied Product</b>	<p>Reagent Provided:</p> <p>Digoxigenin-labeled oligonucleotides targeting sequences mapping in 2p24.3* (chr2:15,846,046-16,213,717) harboring the MYCN gene.</p> <p>2. Formamide based hybridization buffer.</p> <p>*According to Human Genome Assembly GRCh37/hg19</p>
<b>Probe Position</b>	
<b>Regulatory Status</b>	For research use only (RUO)
<b>Storage Instruction</b>	Store at 2-8°C in an upright position. Return to storage conditions immediately after use.

## Note

The probe is intended to be used in combination with the CISH Implementation Kit 2 (Catalog #: [KA5366](#)), which provides necessary reagents for specimen pretreatment and post-hybridization processing.

Hybridization signals of digoxigenin-labeled polynucleotides appear as brown- to dark brown colored distinct dots (MYCN gene).

Normal situation: In interphases of normal cells or cells without an amplification involving the MYCN gene locus, two distinct dot-shaped brown signals per nucleus appear.

Aberrant situation: In a cell with amplification of the MYCN gene locus or aneuploidy of chromosome 2, multiple copies of the brown signal or brown signal clusters will be observed.

## Interpretation of Result

## Applications

- Chromogenic *In Situ* Hybridization (FFPE Tissue)

Normal nuclei each with two MYCN signals

## Gene Info — MYCN

Entrez GeneID	<a href="#">4613</a>
Gene Name	MYCN
Gene Alias	MODED, N-myc, NMYC, ODED, bHLHe37
Gene Description	v-myc myelocytomatosis viral related oncogene, neuroblastoma derived (avian)
Omim ID	<a href="#">164280</a> <a href="#">164840</a> <a href="#">602585</a>
Gene Ontology	<a href="#">Hyperlink</a>
Gene Summary	This gene is a member of the MYC family and encodes a protein with a basic helix-loop-helix (bHLH) domain. This protein is located in the nucleus and must dimerize with another bHLH protein in order to bind DNA. Amplification of this gene is associated with a variety of tumors, most notably neuroblastomas. [provided by RefSeq]
Other Designations	N-myc proto-oncogene protein neuroblastoma MYC oncogene neuroblastoma-derived v-myc avian myelocytomatosis viral related oncogene oncogene NMYC pp65/67 v-myc avian myelocytomatosis viral related oncogene, neuroblastoma derived v-myc myelocytomatosis viral

## Disease

- [Kidney Neoplasms](#)

- [Wilms Tumor](#)