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Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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EGR2 FISH Probe

Catalog # : FA0253

規格 : [200 uL]

List All

Specification

Product Description:	Made to order FISH probes for identification of gene amplification using Fluorescent In Situ Hybridization Technique. (Technology)
Supplied Product:	DAPI Counterstain (1500 ng/mL) 250 uL
Storage Instruction:	Store at 4°C in the dark.
Origin:	Human
Source:	Genomic DNA
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)

Application Image

Fluorescent In Situ Hybridization (Cell)

Applications

Fluorescent In Situ Hybridization (Cell)

 [Protocol Download](#)

Gene Information

Entrez GeneID:	1959
Gene Name:	EGR2
Gene Alias:	AT591,CMT1D,CMT4E,DKFZp686J1957,FLJ14547,KROX20
Gene Description:	early growth response 2 (Krox-20 homolog, Drosophila)
Omim ID:	129010 , 145900 , 605253 , 607678
Gene Ontology:	Hyperlink
Gene Summary:	The protein encoded by this gene is a transcription factor with three tandem C2H2-type zinc fingers. Defects in this gene are associated with Charcot-Marie-Tooth disease type 1D (CMT1D), Charcot-Marie-Tooth disease type 4E (CMT4E), and with Dejerine-Sottas syndrome (DSS). Multiple transcript variants encoding two different isoforms have been found for this gene. [provided by RefSeq]
Other Designations:	KROX-20, Drosophila, homolog (early growth response-2),Krox-20 homolog,OTTHUMP00000062133,early growth response 2 protein

Related Disease

[Alzheimer Disease](#) [Alzheimer disease](#) [Cardiovascular Diseases](#)
[Charcot-Marie-Tooth Disease](#) [Deafness](#) [Deafness](#) [Diabetes Complications](#)
[Genetic Predisposition to Disease](#)
[Hereditary Central Nervous System Demyelinating Diseases](#)
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