SZABO SCANDIC

Part of Europa Biosite

Produktinformation

- Forschungsprodukte & Biochemikalien
- Zellkultur & Verbrauchsmaterial
- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!

Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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SPG7 FISH Probe

Catalog # : FA0378 **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)

Applications

Fluorescent In Situ Hybridization (Cell)

Protocol Download

Gene Information

Entrez GeneID:	6687
Gene Name:	SPG7
Gene Alias:	CAR,CMAR,FLJ37308,MGC126331,MGC126332,PGN,SPG5C
Gene Description:	spastic paraplegia 7 (pure and complicated autosomal recessive)
Omim ID:	602783, 607259
Gene Ontology:	Hyperlink
Gene Summary:	This gene encodes a nuclear-encoded mitochondrial metalloprotease protein that is a member of the AAA (ATPases associated with a variety of cellular activities) protein family. Members of this protein family share an ATPase domain and have roles in diverse cellular processes including membrane trafficking, intracellular motility, organelle biogenesis, protein folding, and proteolysis. Two transcript variants encoding distinct isoforms have been identified for this gene. Mutations associated with this gene cause autosomal recessive spastic paraplegia 7. [provided by RefSeq
Other Designations:	cell adhesion regulator,cell matrix adhesion regulator,paraplegin, isoform 1,spastic paraplegia 7

Application Image

Fluorescent In Situ Hybridization (Cell)

Related Disease

Disease Progression Genetic Predisposition to Disease Kidney Failure, Chronic
Motor Neuron Disease Multiple Sclerosis Paraparesis, Spastic
Spastic Paraplegia, Hereditary