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Diagnostik & molekulare Diagnostik



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See the following pages for more information!



Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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NSD1/CEN5q FISH Probe

Catalog # : FA0658

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 (150 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: [64324](#)

Gene Name: NSD1

Gene Alias: ARA267, DKFZp666C163, FLJ10684, FLJ22263, FLJ44628, KMT3B, SOTOS, STO

Gene Description: nuclear receptor binding SET domain protein 1

Omim ID: [117550](#), [130650](#), [277590](#), [601626](#), [606681](#)

Gene Ontology: [Hyperlink](#)

Gene Summary: This gene encodes a protein containing a SET domain, 2 LXXLL motifs, 3 nuclear translocation signals (NLSs), 4 plant homeodomain (PHD) finger regions, and a proline-rich region. The encoded protein enhances androgen receptor (AR) transactivation, and this enhancement can be increased further in the presence of other androgen receptor associated coregulators. This protein may act as a nucleus-localized, basic transcriptional factor and also as a bifunctional transcriptional regulator. Mutations of this gene have been associated with Sotos syndrome and Weaver syndrome. One version of childhood acute myeloid leukemia is the result of a cryptic translocation with the breakpoints occurring within nuclear receptor-binding Su-var, enhancer

of zeste, and trithorax domain protein 1 on chromosome 5 and nucleoporin, 98-kd on chromosome 11. Two transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq]

Other Designations: androgen receptor-associated coregulator 267

Gene Pathway

[Lysine degradation](#)

Related Disease

[Abnormalities, Multiple Attention Deficit Disorder with Hyperactivity](#) [Autistic Disorder](#)
[Chromosome Breakage](#) [Chromosome Deletion](#) [Craniofacial Abnormalities](#)
[Developmental Disabilities](#) [Endocrine System Diseases](#) [Facies Gigantism](#)
[Growth Disorders](#) [Mental Disorders](#) [Mental Retardation](#) [Motor Skills](#)
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