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Product datasheet for SC206230

Nephronophthisis (NPHP1) (NM_207181) Human 3' UTR Clone

Product data:

Product Type:	3' UTR Clones
Product Name:	Nephronophthisis (NPHP1) (NM_207181) Human 3' UTR Clone
Symbol:	Nephronophthisis
Synonyms:	JBTS4; NPH1; SLSN1
Mammalian Cell Selection:	Neomycin
Vector:	pMirTarget (PS100062)
ACCN:	NM_207181
Insert Size:	473 bp
Insert Sequence:	>SC206230 3'UTR clone of NM_207181 The sequence shown below is from the reference sequence of NM_207181. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC TTGGGTGAAATGAGAAAGAATGCAGTGTGACAGTGGCAGCCTCTAGCCCTCAGCTTCCCACGGAATCAG ATGGATCCTCCACGATTACGTGAATAAAATGATGGAACCAAAAATCACTGTCACTTTACAACTTAGGTT TTACTCTTTTCTTTCTACAGACCATATTTTTAAAGAAATGTTTATACAATAATTTAAATATTTTTTAAA ACCATAAAATAAATTTTTATAAGGAATACTGTTATATCTAAATTTAAACAGTATTTATTTTTTCAAAAA CAGCTACTTAAGTTAATGGTATAGATTTCTATAAAAGCAAGATTTTGTCAAAAACTAAATTTATGATTA TTCAAGAAAGTGAAAAAAACAACCTACAGAATGGGAAAACATATTTGCAAATCATCTAACTGATAAAGG TCTAGTATCCAAAATATTTAAATTTATGAGTGTTAATAAAATTTATCTTGTTCAATGAA ACGCGTAAGCGGCCGCGGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG
Restriction Sites:	SgfI-MluI
OTI Disclaimer:	Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences , e.g., single nucleotide polymorphisms (SNPs).
Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.

View online »

RefSeq: NM_207181.4

Summary: This gene encodes a protein with src homology domain 3 (SH3) patterns. This protein interacts with Crk-associated substrate, and it appears to function in the control of cell division, as well as in cell-cell and cell-matrix adhesion signaling, likely as part of a multifunctional complex localized in actin- and microtubule-based structures. Mutations in this gene cause familial juvenile nephronophthisis type 1, a kidney disorder involving both tubules and glomeruli. Defects in this gene are also associated with Senior-Loken syndrome type 1, also referred to as juvenile nephronophthisis with Leber amaurosis, which is characterized by kidney and eye disease, and with Joubert syndrome type 4, which is characterized by cerebellar ataxia, oculomotor apraxia, psychomotor delay and neonatal breathing abnormalities, sometimes including retinal dystrophy and renal disease. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

Locus ID: 4867

MW: 18.8