

Product datasheet for **TL320625**

ARK5 (NUAK1) Human shRNA Plasmid Kit (Locus ID 9891)

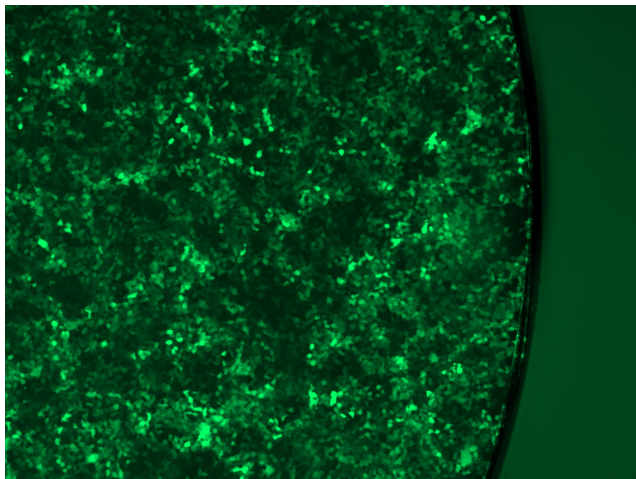
Product data:

Product Type:	shRNA Plasmids
Product Name:	ARK5 (NUAK1) Human shRNA Plasmid Kit (Locus ID 9891)
Locus ID:	9891
Synonyms:	ARK5
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	NUAK1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 9891). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	NM_014840 , NM_014840.1 , NM_014840.2 , BC152462 , BC160165
UniProt ID:	O60285

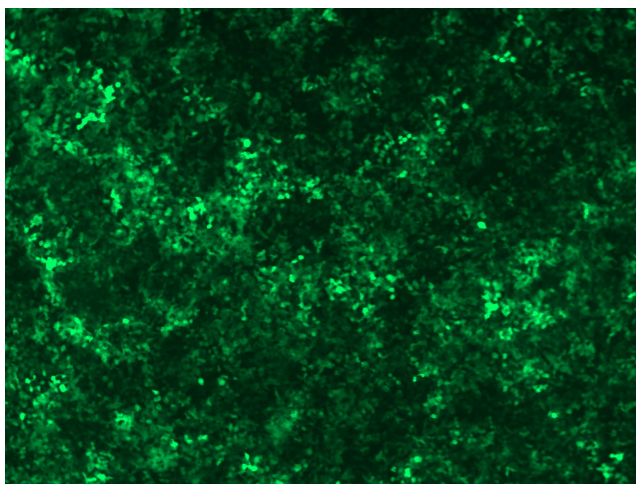


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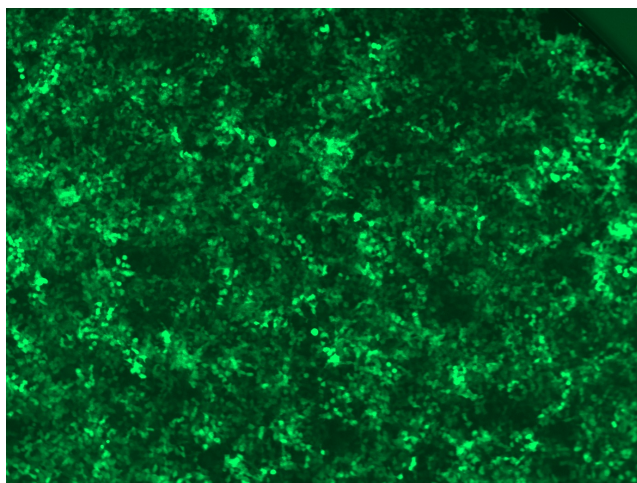
Summary:	<p>Serine/threonine-protein kinase involved in various processes such as cell adhesion, regulation of cell ploidy and senescence, cell proliferation and tumor progression. Phosphorylates ATM, CASP6, LATS1, PPP1R12A and p53/TP53. Acts as a regulator of cellular senescence and cellular ploidy by mediating phosphorylation of 'Ser-464' of LATS1, thereby controlling its stability. Controls cell adhesion by regulating activity of the myosin protein phosphatase 1 (PP1) complex. Acts by mediating phosphorylation of PPP1R12A subunit of myosin PP1: phosphorylated PPP1R12A then interacts with 14-3-3, leading to reduced dephosphorylation of myosin MLC2 by myosin PP1. May be involved in DNA damage response: phosphorylates p53/TP53 at 'Ser-15' and 'Ser-392' and is recruited to the CDKN1A/WAF1 promoter to participate to transcription activation by p53/TP53. May also act as a tumor malignancy-associated factor by promoting tumor invasion and metastasis under regulation and phosphorylation by AKT1. Suppresses Fas-induced apoptosis by mediating phosphorylation of CASP6, thereby suppressing the activation of the caspase and the subsequent cleavage of CFLAR. Regulates UV radiation-induced DNA damage response mediated by CDKN1A. In association with STK11, phosphorylates CDKN1A in response to UV radiation and contributes to its degradation which is necessary for optimal DNA repair (PubMed:25329316).[UniProtKB/Swiss-Prot Function]</p>
shRNA Design:	<p>These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.</p>
Performance Guaranteed:	<p>OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.</p> <p>For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).</p>

Product images:

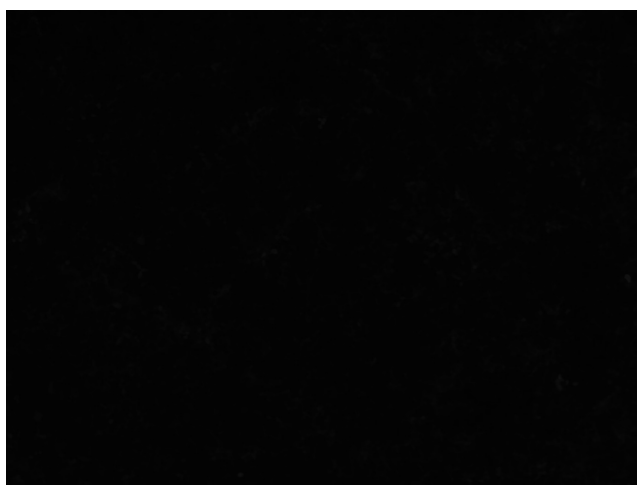
GFP signal was observed under microscope at 48 hours after transduction of TL320625A virus into HEK293 cells. TL320625A virus was prepared using lenti-shRNA TL320625A and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of TL320625B virus into HEK293 cells. TL320625B virus was prepared using lenti-shRNA TL320625B and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL320625C] virus into HEK293 cells. [TL320625C] virus was prepared using lenti-shRNA [TL320625C] and [TR30037] packaging kit.



GFP signal was observed under microscope at 48 hours after transduction of [TL320625D] virus into HEK293 cells. [TL320625D] virus was prepared using lenti-shRNA [TL320625D] and [TR30037] packaging kit.