

Product datasheet for **RG219519**

ATRX (NM_138270) Human Tagged ORF Clone

Product data:

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|---------------------------|---|
| Product Type: | Expression Plasmids |
| Product Name: | ATRX (NM_138270) Human Tagged ORF Clone |
| Tag: | TurboGFP |
| Symbol: | ATRX |
| Synonyms: | JMS; MRX52; RAD54; RAD54L; XH2; XNP; ZNF-HX |
| Mammalian Cell Selection: | Neomycin |
| Vector: | pCMV6-AC-GFP (PS100010) |
| E. coli Selection: | Ampicillin (100 ug/mL) |
| ORF Nucleotide Sequence: | >RG219519 representing NM_138270 Red=Cloning site Blue=ORF Green=Tags(s) |

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
GCC**CGATCGCC**

ATGACCGCTGAGCCCATGAGTGAAAGCAAGTTGAATACATTGGTGCAGAAGCTTCATGACTTCCTTGAC
ACTCATCAGAAGAATCTGAAGAAACAAGTTCTCCTCCAGACTTGCAATGAATCAAAACACAGATAAAAT
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TGGTATGCAGCCACCACCATACAGCGTGACCCACCCCAATGAGAAGCAAAAATCCAGGACCTTCCCAA
GGGAAATCAATG

ACGCGTACGCGGCCGCTCGAG - GFP Tag - GTTTAA

Protein Sequence: >RG219519 representing NM_138270
 Red=Cloning site Green=Tags(s)

MTAEPMSSEKLNLT VQKLHDFLAHSSESEETSSPPRLAMNQNTDKISGSGSNSDMMENSKEEGTSSSEK
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 NHFKD SIYRHPSLQVLICKNCFKYMSDDISRDSGMDQCRWCAEGGNLICDFCHNAFCCKKILRNL
 GRKELSTIMDENNQWYCYICHPEPLLDLVTACNSVFENLEQLLQQNKKKIKVDSEKSNKVEHTSRFSPK
 KTSNCSNGEKKLDDSCSGSVTYSYSALIVPKEMIKKAKKLIETTANMNSYVFKLQATDNSEISSATK
 LRQLKAFKSVLADIKKAHLALEEDLNSEFRAMDAVNKEKNTKEHKVIDAKFETKARKGEKPCALEKKDIS
 KSEAKLSRKQVDSEHMQNVPTEEQRTNKSTGGEHKKSDRKEEPQYEPANTSEDLDMDIVSVPSSVPEDI
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 ETNPVTSNSDEECNETVKEKQKLSVPVRKKDKRNSSDSAIDNPKNPKLPKSKQSETVDQNSDSEMLAIL
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 VTENLVLSSHTGFCQSSGDEALSKSVPVTVD DDDDDNDPENRIAKKMLLEEIKANLSSDEGDSSDPEE
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 VKTKKSPGSGCILAHCMLGKTLQVVSFLHTVLLCDKLDLDFSTALVVCPLNTALNWMNEFEKWQEGKDD
 EKLEVSELATVKRPQERSYMLQRWQEDGGVMIIGYEMYRNLAQGRNVKSRKLEIFNKALVDPGPDFVVC
 DEGHILKNEASAVSKAMNSIRRRRIILTGTPLNQLIEYHCMVNF IKENLLGSIKEFRNRFINPIQNGQ
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 WNPSYDIQSI FRVYRFGQTKPVYVYRFLAQGTMEDKIYDRQVTKQSLSFRVVDQQQVERHFTMNELTELY
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 EAEKGLTMRFNIPGTNLPPVFSNSQTPYIPFNLGALSAMSNQLEDLINQGREKVV EATNSVTAVRIQ
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 NQQQQQMTYQQATLGHLMMPKPPNLIMNPSNYQQIDMRGMYQPVAGGMQPPPLQRAPPPMRSKNPGPSQ
 GKSM

TRTRPLE - GFP Tag - V

Restriction Sites: Sgfl-MluI

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| OTI Disclaimer: | The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. More info |
| OTI Annotation: | This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene. |
| Components: | The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water). |
| Reconstitution Method: | <ol style="list-style-type: none">1. Centrifuge at 5,000xg for 5min.2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.3. Close the tube and incubate for 10 minutes at room temperature.4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C. |
| RefSeq: | NM_138270.2 |
| RefSeq Size: | 10216 bp |
| RefSeq ORF: | 7365 bp |
| Locus ID: | 546 |
| UniProt ID: | P46100 |
| Cytogenetics: | Xq21.1 |
| Domains: | SNF2_N, DEAD, helicase_C |
| Protein Families: | Druggable Genome, Transcription Factors |
| Gene Summary: | The protein encoded by this gene contains an ATPase/helicase domain, and thus it belongs to the SWI/SNF family of chromatin remodeling proteins. This protein is found to undergo cell cycle-dependent phosphorylation, which regulates its nuclear matrix and chromatin association, and suggests its involvement in the gene regulation at interphase and chromosomal segregation in mitosis. Mutations in this gene are associated with X-linked syndromes exhibiting cognitive disabilities as well as alpha-thalassemia (ATRX) syndrome. These mutations have been shown to cause diverse changes in the pattern of DNA methylation, which may provide a link between chromatin remodeling, DNA methylation, and gene expression in developmental processes. Multiple alternatively spliced transcript variants encoding distinct isoforms have been reported. [provided by RefSeq, Jul 2017] |