

Product datasheet for **SC122501**

MITF (BC012503) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	MITF (BC012503) Human Untagged Clone
Tag:	Tag Free
Symbol:	MITF
Synonyms:	bHLHe32; CMM8; MI; WS2; WS2A
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene sequence for BC012503 edited
 CTTTGCCAGTCCATCTTCAAATTGGAATTATAGAAAGTAGAGGGAGGGATAGTCTACCGT
 CTCTCACTGGATTGGTGCCACCTAAAACATTGTTATGCTGGAAATGCTAGAATATAATCA
 CTATCAGGTGCAGACCCACCTCGAAAACCCACCAAGTACCACATACAGCAAGCCCAACG
 GCAGCAGGTAAAGCAGTACCTTTCTACCACTTTAGCAAATAAACATGCCAACCAAGTCTT
 GAGCTTGCCATGTCCAAACCAGCCTGGCGATCATGTCATGCCACCGGTGCCGGGGAGCAG
 CGCACCCAACAGCCCATGGCTATGCTTACGCTTAACTCCAAGTGTAAAAAGAGTTTAT
 GAAGCAGTGAGAAATGCAGAGAGAGGAGAAGGGGAGGTGGAAAAGGAAAAGCAAAAATAGA
 AGAGGTGTGGGACATGCTGTTTAGAAGTCCGCTTGTGTGAATGTCTGGAATATTATTT
 TTATTTCTCCCTGAGTTGGGGGAAGAAAGAATGGAATATGCATGGATGGATTTGAATCAT
 ATAGCACATGAGACTTTAACGGAAACGCAAAGGTTTAATTGCTGGATACATTCTGTTTCA
 TAATAAAATTGCCACTGCCCGTTAAATCTGCTTTGGTGAAGGCTGGATTGGAAACAAGAC
 TCAAACCTACCTCAAGCTAATTGGTGCATCAAAATTTGCAGCATACAAATACCTGAGAGC
 TGTGATTTAATGCTCATTATTTCCAAATTATGAGATGATGAGCTTCATCTCAATGGGATT
 TACCGTACTATGGACTATGAAGTGTATGCAAAATTCGGAGGCAACTTTTCTAGAGTTGG
 ATTTGATTTAATTTCTAGAGGGACTAAAATCTTTGCCCTATGCCCAAACCAACTGCTTT
 ATTTTCTCTACCCAAATTTGTCATCTAGCAAGATGATTTGACACAAGTCTTCTTTCAT
 TATTTTCATTTTTGGTCAGATCCACTTTGTTTGAAGCTTAGTTCATCTTGTGCTGTG
 CCATCAGCTTTGTGTGAACAGGTCATTAATAAAGTCATTTGCAAATCCAAAAA
 AA



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5' Read Nucleotide Sequence:	>OriGene 5' read for BC012503 unedited GGGCGGTTCAAATTTGTATACGACTCACTATGGGCGGCCGCGATTCCCGGATATCGTCCG ACCCACGCGTCCGCTTTGCCAGTCCATCTTCAAATTGGAATTATAGAAAGTAGAGGGAGG GATAGTCTACCGTCTCTCACTGGATTGGTGCCACCTAAAACATTGTTATGCTGGAATGC TAGAATAATACTACTATCAGGTGCAGACCCACCTCGAAAACCCCAAGTACCACATAC AGCAAGCCCAACGGCAGCAGGTAAGCAGTACCTTTCTACCACTTAGCAAATAAACATG CCAACCAAGTCCAGCTTGCCATGTCCAAACAGCCTGGCGATCATGTCATGCCACCGG TGCCGGGAGCAGCGCACCCAACAGCCCATGGCTATGCTTACGCTTAACTCCAACCTGTG AAAAAGAGTTTATGAAGCAGTGAGAATGCAGAGAGAGGAGAAGGGGAGGTGAAAAAGGAA AAGCAAAAATAGAAGAGGTGTGGGACATGCTGTTTAGAAGTTCGCTTGTGTGAATGTC TGAATATTATTTTTATTCTCCCTGAGTTGGGGGAAGAAAGAATGGAATATGCATGGAT GGATTTGAATCATATAGCACATGAGACTTTAACGGAAACGCAAAGGTTAATTGCTGGAT ACATTCTGTTTCATAATAAAATTGCCACTGCCCGTTAAATCTGCTTTGGTGAAGGCTGGA TTGGAAACAAGACTCANACTACCTTCAAGCTAATTGGTGCATCAAATTTGCAGCATACA AATACCTGAGAGCTGTGATTTAATGCTCATTATTTCCAATTATGAGATGATGAGCTTCAT CTCAATGGGATTTACCGTACTATGGGACTATGAAGTGTTTATGCCAATTCGGAGGCAACT TTNCTAGATTGGNA
Restriction Sites:	Please inquire
ACCN:	BC012503
Insert Size:	1082 bp
OTI Disclaimer:	Our molecular clone sequence data has been matched to the reference identifier above as a point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative RNA splicing form or single nucleotide polymorphism (SNP).
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:	BC012503.1
RefSeq Size:	1082 bp
Locus ID:	4286
Cytogenetics:	3p13
Protein Families:	Druggable Genome, Transcription Factors
Protein Pathways:	Melanogenesis, Melanoma, Pathways in cancer

Gene Summary:

The protein encoded by this gene is a transcription factor that contains both basic helix-loop-helix and leucine zipper structural features. The encoded protein regulates melanocyte development and is responsible for pigment cell-specific transcription of the melanogenesis enzyme genes. Heterozygous mutations in the this gene cause auditory-pigmentary syndromes, such as Waardenburg syndrome type 2 and Tietz syndrome. [provided by RefSeq, Aug 2017]