

Product datasheet for SC201861

BLM (NM_000057) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: BLM (NM_000057) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: BLM

Synonyms: BS; MGRISCE1; RECQ2; RECQL2; RECQL3

ACCN: NM_000057

Insert Size: 917 bp

Insert Sequence: >SC201861 3'UTR clone of NM_000057

The sequence shown below is from the reference sequence of NM_000057. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

ATAAAAATATGATAAAACCA

CAACCTGCCATCACGAGATTTCGATTCCACCGCCGC

Restriction Sites: Sgfl-Rsrll

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).



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BLM (NM_000057) Human 3' UTR Clone - SC201861

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.

RefSeq: <u>NM 000057.4</u>

Summary: The Bloom syndrome is an autosomal recessive disorder characterized by growth deficiency,

microcephaly and immunodeficiency among others. It is caused by homozygous or compound heterozygous mutation in the gene encoding DNA helicase RecQ protein on chromosome 15q26. This Bloom-associated helicase unwinds a variety of DNA substrates including Holliday junction, and is involved in several pathways contributing to the

maintenance of genome stability. Identification of pathogenic Bloom variants is required for

heterozygote testing in at-risk families. [provided by RefSeq, May 2020]

Locus ID: 641 MW: 34.5