

Product datasheet for **SC330728**

ASH2L (NM_001261832) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: ASH2L (NM_001261832) Human Untagged Clone
Tag: Tag Free
Symbol: ASH2L
Synonyms: ASH2; ASH2L1; ASH2L2; Bre2
Vector: pCMV6-Entry (PS100001)
Fully Sequenced ORF: >SC330728 representing NM_001261832.
Blue=Insert sequence Red=Cloning site Green=Tag(s)

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ATGGATACTCAGGCGGGCTCCGTGGATGAAGAGAATGGCCGACAGTTGGGTGAGGTAGAGCTGCAATGT
GGGATTTGTACAAAATGGTTCACGGCTGACACATTTGGCATAGATACCTCATCTGTCTACCTTTTCATG
ACCAACTACAGTTTTTCATTGCAACGTCTGCCATCACAGTGGGAATACCTATTTCTCCGGAAGCAAGCA
AACTTGAAGGAAATGTGCCTTAGTGCTTTGGCCAACCTGACATGGCAGTCCCGAACACAGGATGAACAT
CCGAAGACAATGTTCTCCAAGATAAAGGATATTATACCATTTATTGATAAATACTGGGAGTGCATGACA
ACCAGACAGAGACCTGGGAAAATGACTTGGCCAAATAACATTGTTAAAACAATGAGTAAAGAAAGAGAT
GTATTTCTGGTAAAGGAACACCCAGATCCAGGCAGTAAAGATCCAGAAGAAGATTACCCCAAATTTGGA
CTTTTGGATCAGGACCTTAGTAACATTGGTCTGCTTATGACAACCAAAAACAGAGCAGTGTGTGTCT
ACTAGTGGGAATTTAAATGGGGGAATTGCAGCAGGAAGCAGCGGAAAAGGACGAGGAGCCAAGCGCAAA
CAGCAGGATGGAGGGACCACAGGGACCACCAAGAAGGCCCGGAGTGACCCCTTTGTTTTCTGCTCAGCGC
CTTCCCTCATGGCTACCCATTGGAACACCCGTTTAAACAAGATGGCTATCGGTATATTCTAGCTGAG
CCTGATCCGCACGCCCTGACCCCGAGAAGCTGGAACCTGACTGCTGGGCAGGAAAACCTATTCTGGA
GACCTCTACAGAGCCTGCTTGTATGAACGGTTTTGTTAGCCCTACATGATCGAGCTCCCCAGTTAAAG
ATCTCAGATGACCGGCTGACTGTGGTTGGAGAGAAGGGCTACTCTATGGTGAGGGCCTCATGGAGTA
CGGAAAGGTGCCTGGTATTTTAAAATCACTGTGGATGAGATGCCACCAGATACCGCTGCCAGACTGGGT
TGGTCCCAGCCCCTAGGAAACCTTCAAGCTCCTTTAGGTTATGATAAATTTAGCTATTCTTGGCGGAGC
AAAAGGGAACCAAGTCCACCAGTCCATTGGCAAACACTACTCTTCTGGCTATGGACAGGGAGACGTC
CTGGGATTTTATATTAATCTTCTGAAGACACAGAGACAGCCAAGTATTGCCAGACACATACAAAGAT
AAGGCTTTGATAAAATCAAGAGTTATTTGTATTTTGGAGAAAAGACTTTGTGGATAAAGCAGAGAAG
AGCCTGAAGCAGACTCCCATAGTGAGGTTTCCATTAACCTTTGGACCATGCTTCAAGTATCTCCGAAAG
GATCTCACTTACCGCCCTATGAGTGACATGGGCTGGGGCCCGTGGTAGAGCACACCCTGGCTGACGTC
TTGTATCAGTGGAGACAGAAGTGGATGGGAGGCGCAGTCCCCATGGGAACCTGA
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Restriction Sites: SgfI-MluI
ACCN: NM_001261832
Insert Size: 1506 bp



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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:	NM_001261832.1
RefSeq Size:	2956 bp
RefSeq ORF:	1506 bp
Locus ID:	9070
UniProt ID:	Q9UBL3
Cytogenetics:	8p11.23
Protein Families:	Druggable Genome, Transcription Factors
MW:	56.5 kDa
Gene Summary:	<p>Component of the Set1/Ash2 histone methyltransferase (HMT) complex, a complex that specifically methylates 'Lys-4' of histone H3, but not if the neighboring 'Lys-9' residue is already methylated. As part of the MLL1/MLL complex it is involved in methylation and dimethylation at 'Lys-4' of histone H3. May function as a transcriptional regulator. May play a role in hematopoiesis.[UniProtKB/Swiss-Prot Function]</p> <p>Transcript Variant: This variant (3) differs in the 5' UTR, initiates translation at a downstream, in-frame start codon and lacks an exon in the 3' coding region, but maintains the reading frame, compared to variant 1. The encoded isoform (c) is shorter than isoform a. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.</p>