

Product datasheet for SC312058

OriGene Technologies, Inc.

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NBEAL2 (AK074036) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: NBEAL2 (AK074036) Human Untagged Clone

Tag: Tag Free Symbol: NBEAL2

Synonyms: BDPLT4; GPS
Vector: pCMV6 series

Fully Sequenced ORF: >NCBI ORF sequence for AK074036, the custom clone sequence may differ by one or more

nucleotides

Restriction Sites: Please inquire

ACCN: AK074036

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).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

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: 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: <u>AK074036.1</u>, <u>BAB84862.1</u>

RefSeq Size: 4422 bp





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 RefSeq ORF:
 4422 bp

 Locus ID:
 23218

 Cytogenetics:
 3p21.31

Gene Summary: The protein encoded by this gene contains a beige and Chediak-Higashi (BEACH) domain and

multiple WD40 domains, and may play a role in megakaryocyte alpha-granule biogenesis. Mutations in this gene are a cause of gray platelet syndrome. [provided by RefSeq, Dec 2011]