

## **Product datasheet for SC328418**

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## SLC29A3 (NM\_001174098) Human Untagged Clone

**Product data:** 

**Product Type:** Expression Plasmids

**Product Name:** SLC29A3 (NM\_001174098) Human Untagged Clone

Tag: Tag Free Symbol: SLC29A3

Synonyms: ENT3; HCLAP; HJCD; PHID

Mammalian Cell

Selection:

None

Vector: pCMV6-XL5

E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM\_001174098, the custom clone sequence may differ by one or

more nucleotides

**Restriction Sites:** Please inquire ACCN: NM 001174098

**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).



## SLC29A3 (NM\_001174098) Human Untagged Clone - SC328418

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>NM 001174098.1</u>, <u>NP 001167569.1</u>

RefSeq Size: 2283 bp
RefSeq ORF: 777 bp
Locus ID: 55315
UniProt ID: Q9BZD2
Cytogenetics: 10q22.1

**Protein Families:** Transmembrane

**Gene Summary:** This gene encodes a nucleoside transporter. The encoded protein plays a role in cellular

uptake of nucleosides, nucleobases, and their related analogs. Mutations in this gene have been associated with H syndrome, which is characterized by cutaneous hyperpigmentation and hypertrichosis, hepatosplenomegaly, heart anomalies, and hypogonadism. A related disorder, PHID (pigmented hypertrichosis with insulin-dependent diabetes mellitus), has also been associated with mutations at this locus. Alternatively spliced transcript variants have

been described.[provided by RefSeq, Mar 2010]

Transcript Variant: This variant (2) uses an alternate splice site in the 3' coding region, which results in a frameshift, compared to variant 1. It encodes isoform b, which is shorter than

isoform a.