

## Product datasheet for RC215751L4V

## OriGene Technologies, Inc.

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## EDA (NM\_001005612) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

Product Name: EDA (NM 001005612) Human Tagged ORF Clone Lentiviral Particle

Symbol: EDA

Synonyms: ECTD1; ED1-A1; ED1-A2; EDA-A1; EDA-A2; EDA1; EDA2; HED; HED1; ODT1; STHAGX1;

TNLG7C; XHED; XLHED

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-mGFP-P2A-Puro (PS100093)

Tag: mGFP

**ACCN:** NM\_001005612

ORF Size: 1158 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(RC215751).

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

**RefSeq:** <u>NM 001005612.2</u>

RefSeq Size:5281 bpRefSeq ORF:1161 bpLocus ID:1896

UniProt ID: Q92838

Cytogenetics: Xq13.1

**Protein Families:** Druggable Genome, Secreted Protein, Transmembrane





## EDA (NM\_001005612) Human Tagged ORF Clone Lentiviral Particle - RC215751L4V

**Protein Pathways:** Cytokine-cytokine receptor interaction

MW: 40.8 kDa

**Gene Summary:** The protein encoded by this gene is a type II membrane protein that can be cleaved by furin

to produce a secreted form. The encoded protein, which belongs to the tumor necrosis factor

family, acts as a homotrimer and may be involved in cell-cell signaling during the

development of ectodermal organs. Defects in this gene are a cause of ectodermal dysplasia,

anhidrotic, which is also known as X-linked hypohidrotic ectodermal dysplasia. Several transcript variants encoding many different isoforms have been found for this gene.

[provided by RefSeq, Jul 2008]