

## **Product datasheet for SC334214**

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## IGF2 (NM\_001291862) Human Untagged Clone

## **Product data:**

**Product Type:** Expression Plasmids

**Product Name:** IGF2 (NM\_001291862) Human Untagged Clone

Tag: Tag Free Symbol: IGF2

Synonyms: C11orf43; GRDF; IGF-II; PP9974; SRS3

**Vector:** pCMV6 series

Fully Sequenced ORF: >NCBI ORF sequence for NM\_001291862, the custom clone sequence may differ by one or

more nucleotides

AAGACCCCGCCCACGGGGGCGCCCCCCCAGAGATGGCCAGCAATCGGAAGTGA

**Restriction Sites:** Sgfl-Mlul

**ACCN:** NM 001291862

**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:** 

- 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 001291862.2, NP 001278791.1

RefSeq Size: 4710 bp RefSeq ORF: 543 bp Locus ID: 3481 **UniProt ID:** P01344 Cytogenetics: 11p15.5

**Protein Families:** Druggable Genome, ES Cell Differentiation/IPS, Secreted Protein

This gene encodes a member of the insulin family of polypeptide growth factors, which are **Gene Summary:** 

> involved in development and growth. It is an imprinted gene, expressed only from the paternal allele, and epigenetic changes at this locus are associated with Wilms tumour, Beckwith-Wiedemann syndrome, rhabdomyosarcoma, and Silver-Russell syndrome. A readthrough INS-IGF2 gene exists, whose 5' region overlaps the INS gene and the 3' region overlaps this gene. Alternatively spliced transcript variants encoding different isoforms have

been found for this gene. [provided by RefSeq, Oct 2010]

Transcript Variant: This variant (5) differs in the 5' UTR exon, compared to variant 1. Variants 1, 2, 4 and 5 encode the same isoform (1). Sequence Note: This RefSeg 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.