

Product datasheet for SC311756

LHX4 (AK096250) Human Untagged Clone

Product data:

Product Type: Expression Plasmids Product Name: LHX4 (AK096250) Human Untagged Clone Tag: Tag Free Symbol: LHX4 Vector: pCMV6 series >NCBI ORF sequence for AK096250, the custom clone sequence may differ by one or more **Fully Sequenced ORF:** nucleotides **Restriction Sites:** Please inquire ACCN: AK096250 **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. The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube **Components:** 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:** AK096250.1 **RefSeq Size:** 2655 bp **RefSeq ORF:** 2655 bp



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	LHX4 (AK096250) Human Untagged Clone – SC311756
Locus ID:	89884
Cytogenetics:	1q25.2
Protein Families:	Druggable Genome, Transcription Factors
Gene Summary:	This gene encodes a member of a large protein family which contains the LIM domain, a unique cysteine-rich zinc-binding domain. The encoded protein is a transcription factor involved in the control of differentiation and development of the pituitary gland. Mutations in this gene cause combined pituitary hormone deficiency 4. [provided by RefSeq, Dec 2010]

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