

# Product datasheet for SR311396

## GJC2 Human siRNA Oligo Duplex (Locus ID 57165)

### **Product data:**

### OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product Type:siRNA Oligo DuplexesPurity:HPLC purifiedQuality Control:Tested by ESI-MSSequences:Available with shipmentStability:One year from date of shipment when stored at -20°C.# of transfections:Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final conc. 10 nM).Note:Single siRNA duplex (10nmol) can be ordered.RefSeq:MM 202435UniProt ID:O35442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) -3 unique 27mer siRNA duplexes -2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex -2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer -2 mlSummary:Shis gene encodes a gap junction protein, and pi junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in pripheral myelination in humans. Defects in this gene are the cause of autosomal recessive pelizeus-Merzbacher-like disease-1. [provided by RefSeq, Jul 2008]	i loudet dutu.	
Quality Control:Tested by ESI-MSSequences:Available with shipmentStability:One year from date of shipment when stored at -20°C.# of transfections:Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final conc. 10 nM).Note:Single siRNA duplex (10nmol) can be ordered.RefSeq:NM 020435UniProt ID:Q5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Product Type:	siRNA Oligo Duplexes
Sequences:Available with shipmentStability:One year from date of shipment when stored at -20°C.# of transfections:Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final conc. 10 nM).Note:Single siRNA duplex (10nmol) can be ordered.RefSeq:NM 020435UniProt ID:Q5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Purity:	HPLC purified
Stability:One year from date of shipment when stored at -20°C.# of transfections:Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final conc. 10 nM).Note:Single siRNA duplex (10nmol) can be ordered.RefSeq:NM 020435UniProt ID:Q5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Quality Control:	Tested by ESI-MS
# of transfections:Approximately 330 transfections/2nmol in 24-well plate under optimized conditions (final conc. 10 nM).Note:Single siRNA duplex (10nmol) can be ordered.RefSeq:NM 020435UniProt ID:Q5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Sequences:	Available with shipment
conc. 10 nM).Note:Single siRNA duplex (10nmol) can be ordered.RefSeq:NM 020435UniProt ID:Q5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Stability:	One year from date of shipment when stored at -20°C.
RefSeq:NM 020435UniProt ID:Q5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	# of transfections:	
UniProt ID:O5T442Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Note:	Single siRNA duplex (10nmol) can be ordered.
Synonyms:CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	RefSeq:	<u>NM 020435</u>
Components:GJC2 (Human) - 3 unique 27mer siRNA duplexes - 2 nmol each (Locus ID 57165) Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	UniProt ID:	<u>Q5T442</u>
Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol Included - SR30005, RNAse free siRNA Duplex Resuspension Buffer - 2 mlSummary:This gene encodes a gap junction protein. Gap junction proteins are members of a large family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Synonyms:	CX46.6; Cx47; GJA12; HLD2; LMPH1C; LMPHM3; PMLDAR; SPG44
family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive	Components:	Included - SR30004, Trilencer-27 Universal Scrambled Negative Control siRNA Duplex - 2 nmol
	Summary:	family of homologous connexins and comprise 4 transmembrane, 2 extracellular, and 3 cytoplasmic domains. This gene plays a key role in central myelination and is involved in peripheral myelination in humans. Defects in this gene are the cause of autosomal recessive



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2021 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

# GJC2 Human siRNA Oligo Duplex (Locus ID 57165) - SR311396Performance<br/>Guaranteed:OriGene guarantees that at least two of the three Dicer-Substrate duplexes in the kit will<br/>provide at least 70% or more knockdown of the target mRNA when used at 10 nM<br/>concentration by quantitative RT-PCR when the TYE-563 fluorescent transfection control<br/>duplex (cat# SR30002) indicates that >90% of the cells have been transfected and the HPRT<br/>positive control (cat# SR30003) provides 90% knockdown efficiency.For non-conforming siRNA, requests for replacement product must be made within ninety<br/>(90) days from the date of delivery of the siRNA kit. To arrange for a free replacement with<br/>newly designed duplexes, please contact Technical Services at techsupport@origene.com.<br/>Please provide your data indicating the transfection efficiency and measurement of gene<br/>expression knockdown compared to the scrambled siRNA control (quantitative RT-PCR data<br/>required).

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2021 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US