

B6.129S6-Gt(ROSA)26Sor^{tm1.1}(R26-60-DR5-TA-Cerulean)Mgn/Vu

This is a retinoic acid responsive CFP reporter allele. In this allele, the Rosa26 promoter was modified using RMCE, replacing DNA sequences from -60 to +81 with a multimerized retinoic acid response element (DR5) fused to a TATA box.

Keywords: [Rosa26^{R26-60-DR5-TA-Cerulean}](#) [Mgn](#) [Rosa26](#) [RMCE](#) [Cerulean](#)

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Mouse Information

Common Name	Rosa26 ^{R26-60-DR5-TA-Cerulean}
Research Applications	<i>Not provided</i>
MMRRC ID	<i>Not provided</i>
Jackson Laboratories Stock No	<i>Not provided</i>
VCMR ID	KV
Additional Strain Information	Through homologous recombination in ES cells, a 5.165 kb region of the Rosa26 gene containing exon 1 was replaced by a floxed tk-neo cassette, a puromycin-delta-TK fusion gene driven by the mouse phosphoglycerol kinase promoter (puroR-delta-TK) and a neomycin resistant gene driven by the bacterial EM7 promoter (EM7neo) flanked by minimal (34 bp) tandemly oriented lox71 and lox2272 sites (Cre-recombinase recognition sequences).

Genetic Alteration

Mutation #1: RMCE Targeted Mutagenesis	
Type of Allele	Gene Replacement
Targeted Gene	Name: gene trap ROSA 26, Philippe Soriano Symbol: Gt(ROSA)26Sor NCBI: 14910
Allele	Name: targeted mutation 1.1 Symbol: Gt(ROSA)26Sor ^{tm1.1} (R26-60-DR5-Cerulean)Mgn
Description of Targeting Vector	Through homologous recombination in ES cells, a 5.165 kb region of the Rosa26 gene containing exon 1 was replaced by a floxed tk-neo cassette, a puromycin-delta-TK fusion gene driven by the mouse phosphoglycerol kinase promoter (puroR-delta-TK) and a neomycin resistant gene driven by the bacterial EM7 promoter (EM7neo) flanked by minimal (34 bp) tandemly oriented lox71 and lox2272 sites (Cre-recombinase recognition sequences).

Vector Genbank File	pRosa26.LCA.gb
Allele Map	<i>Not Provided</i>
PCR Genotyping Protocol	R2660_and_228_genotyping_protocol.docx
Type of Allele	Gene Replacement
Exchanged Cassette Gene Name	(CFP)
Exchanged Cassette Allele Name	Rosa26{tm1.1(R26-DR5-TA-Cerulean)}
Description of Exchange Vector	In the R26-60-DR5-TA-Cerulean exchange vector a native Rosa26 gene sequence from -60 to +81 is replaced by a retinoic acid response element (DR5) fused to a TATA-Cerulean(CFP) fluorescent reporter. The vector also contains RMCE-enabling Lox66/Lox2272 sites and Pgk-hygromycin resistance cassette flanked by tandem FRT sites, enabling excision of the cassette after germline transmission.
Exchanged Cassette Genbank File	R26-60-DR5-TA-Cerulean.gb
PCR Genotyping Protocol	<i>Not provided</i>
Citations	<p>Publication</p> <p>Partial promoter substitutions generating transcriptional sentinels of diverse signaling pathways in embryonic stem cells and mice. (2012) <i>Dis Model Mech</i> 5: 956-66 (Added 11/6/2013) PMID: 22888097</p>

Background Strain Information

Strain Type	Congenic Strain
Chimera/Founder Genetic Background	129S6/SvEvTac
Current Genetic Background	C57BL/6J
Number of Generations Backcrossed	2

Strain Description

Germline 129S6 chimeras were backcrossed to C57BL/6J for two generations.


96.97% C57BL/6J at cryopreservation.

Cryopreserved in 2010.

Attachments

 [r2660dr5tacerh_pcr_protocol.doc](#) - Added on July 27, 2010 at 1:55 PM by Jill Lindner

PCR protocol for genotyping mice

 [R26-60-DR5-TA-Cerulean.png](#) - Added on July 19, 2010 at 10:17 AM by Mark Magnuson

Rosa26-60-DR5-TA-Cerulean gene targeting and RMCE strategy.

