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## Role of major and brain-specific *Sgce* isoforms in the pathogenesis of myoclonus-dystonia syndrome

Jianfeng Xiao<sup>a</sup>, Satya R. Vemula<sup>a</sup>, Yi Xue<sup>a</sup>, Mohammad M. Khan<sup>a</sup>, Francesca A. Carlisle<sup>d</sup>, Adrian J. Waite<sup>d</sup>, Derek J. Blake<sup>d</sup>, Ioannis Dragatsis<sup>c</sup>, Yu Zhao<sup>a</sup>, and Mark S. LeDoux<sup>a,b,\*</sup>

<sup>a</sup>Department of Neurology, University of Tennessee Health Science Center, Memphis, TN, 38163, USA

<sup>b</sup>Department of Anatomy and Neurobiology, University of Tennessee Health Science Center, Memphis, TN, 38163, USA

<sup>c</sup>Department of Physiology, University of Tennessee Health Science Center, Memphis, TN, 38163, USA

<sup>d</sup>MRC Centre for Neuropsychiatric Genetics and Genomics, Cardiff University School of Medicine, Cathays, Cardiff, CF24 4HQ, Great Britain

### Abstract

Loss-of-function mutations in *SGCE*, which encodes  $\epsilon$ -sarcoglycan ( $\epsilon$ -SG), cause myoclonus--dystonia syndrome (OMIM159900, DYT11). A “major”  $\epsilon$ -SG protein derived from CCDS5637.1 (NM\_003919.2) and a “brain-specific” protein, that includes sequence derived from alternative exon 11b (CCDS47642.1, NM\_001099400.1), are reportedly localized in post- and pre-synaptic membrane fractions, respectively. Moreover, deficiency of the “brain-specific” isoform and other isoforms derived from exon 11b may be central to the pathogenesis of DYT11. However, no animal model supports this hypothesis. Gene-trapped ES cells (CMHD-GT\_148G1-3, intron 9 of NM\_011360) were used to generate a novel *Sgce* mouse model (C57BL/6J background) with markedly reduced expression of isoforms derived from exons 3' to exon 9 of NM\_011360. Among those brain regions analyzed in adult (2 month-old) wild-type (WT) mice, cerebellum showed the highest relative expression of isoforms incorporating exon 11b. Homozygotes (*Sgce*<sup>Gt(148G1)Cmhd/Gt(148G1)Cmhd</sup> or *Sgce*<sup>Gt/Gt</sup>) and paternal heterozygotes (*Sgce*<sup>m+/pGt</sup>, m-maternal, p-paternal) showed 60 to 70% reductions in expression of total *Sgce*. Although expression of the major (NM\_011360) and brain-specific (NM\_001130189) isoforms was markedly reduced, expression of short isoforms was preserved and relatively small amounts of chimeric  $\epsilon$ -SG/ $\beta$ -galactosidase fusion protein was produced by the *Sgce* gene-trap locus. Immunoaffinity purification followed by mass spectrometry assessments of *Sgce*<sup>m+/pGt</sup> mouse brain using pan- or brain-specific  $\epsilon$ -SG antibodies revealed significant reductions of  $\epsilon$ -SG and

\*Corresponding author: University of Tennessee Health Science Center, Department of Neurology, 855 Monroe Avenue, Suite 415 Link Building, Memphis, TN, 38163, USA, Tel.: 901-448-1662, mledoux@uthsc.edu.

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other interacting sarcoglycans. Genome-wide gene-expression data using RNA derived from adult *Sgce*<sup>m+/pGt</sup> mouse cerebellum showed that the top up-regulated genes were involved in cell cycle, cellular development, cell death and survival, while the top down-regulated genes were associated with protein synthesis, cellular development, and cell death and survival. In comparison to WT littermates, *Sgce*<sup>m+/pGt</sup> mice exhibited “tiptoe” gait and stimulus-induced appendicular posturing between Postnatal Days 14 to 16. Abnormalities noted in older *Sgce*<sup>m+/pGt</sup> mice included reduced body weight, altered gait dynamics, and reduced open-field activity. Overt spontaneous or stimulus-sensitive myoclonus was not apparent on the C57BL/6J background or mixed C57BL/6J-BALB/c and C57BL/6J-129S2 backgrounds. Our data confirm that mouse *Sgce* is a maternally imprinted gene and suggests that short *Sgce* isoforms may compensate, in part, for deficiency of major and brain-specific *Sgce* isoforms.

## Keywords

*Sgce*; Dystonia; Myoclonus; Sarcoglycans; Gene Trap

## 1. Introduction

Myoclonus-dystonia syndrome is a clinically and genetically heterogeneous disorder characterized by myoclonic jerks affecting mainly proximal muscles of the upper extremities, neck and trunk. Mutations (missense, nonsense, splicing, insertions, and deletions) in *SGCE* represent a major cause of the myoclonus-dystonia syndrome (MDS, DYT11, OMIM-159900)(Zimprich et al., 2001, Aichhorn et al., 2006) (Asmus et al., 2005) and are believed to cause disease via loss-of-function mechanisms (Kinugawa et al., 2009, Xiao et al., 2013). The onset of the disorder is usually in the first or second decade of life. Females tend to have earlier onset and greater probability of leg involvement than males (Raymond et al., 2008). *SGCE* is a maternally imprinted gene and MDS is inherited in an autosomal dominant fashion with reduced penetrance (Zimprich et al., 2001). *SGCE* encodes the epsilon member of the sarcoglycan (SG) family ( $\epsilon$ -SG), single pass transmembrane proteins that are components of the dystrophin-glycoprotein complex (DGC). In cardiac and skeletal muscle, the DGC connects the actin cytoskeleton to the extracellular matrix.

In humans and mice, *SGCE/Sgce* harbors several alternatively spliced exons (Fig. 1) (Yokoi et al., 2005, Nishiyama et al., 2004, Ritz et al., 2011). In humans, a “brain-specific” isoform that incorporates alternative exon 11b (CCDS47642.1, NM\_001099400.1, NP\_001123661.1) shows high expression in Purkinje cells and neurons of the cerebellar dentate nucleus with significantly lower levels in the globus pallidus, striatum and substantia nigra (Ritz et al., 2011). In mice, the “major”  $\epsilon$ -SG protein derived from NM\_011360 (NP\_035490.3) and the “brain-specific” isoform are reportedly localized in post- and pre-synaptic membrane fractions, respectively (Nishiyama et al., 2004). Mice may also express another “brain-specific” isoform encoded by an elongated exon 11b (exon 11c) (Yokoi et al., 2005). Proteins derived from transcripts containing either exon 11b or 11c encode proteins with C-terminal PDZ-binding motifs (Yokoi et al., 2005). Therefore, it has been suggested that the brain-specific isoforms may be central to the pathogenesis of MDS. However,

although 11b was not screened in several published cohorts, no disease-associated mutations in terminal exons of *SGCE* (exons 11, 11b and 12) have been reported in patients with MDS (Table 1). Furthermore, the capacity of  $\epsilon$ -SG to tolerate deleterious variants encoded by 11b is difficult to estimate given that 11b was not captured in many large whole-exome sequencing projects (Table S1). On the other hand, there are relatively few reported sequence variants in the terminal exons of *SGCE* (Table S1) which encode the C-terminus PDZ-binding cytoplasmic domain of  $\epsilon$ -SG.

Several mouse models have been used to explore the biology of  $\epsilon$ -SG and pathobiology of MDS. Overexpression of wild-type (WT)  $\epsilon$ -SG in mice resulted in substitution of  $\epsilon$ -SG for  $\alpha$ -SG in muscle DGC without overt behavioral or morphological consequences (Imamura et al., 2005). Paternal heterozygous null mice (*Sgce*<sup>m+/p-</sup>) were reported to exhibit myoclonus, increased slips on a beam-walking test, hyperactivity, and higher levels of striatal dopamine (DA) and dopamine metabolites 3,4-dihydroxyphenylacetic acid (DOPAC) and homovanillic acid (HVA) (Yokoi et al., 2006). In contrast, mice with conditional knock-out (KO) of *Sgce* in Purkinje cells showed subtle motor learning deficits but did not exhibit myoclonus or robust motor abnormalities (Yokoi et al., 2012a). To gain additional insights into  $\epsilon$ -SG function and the pathogenesis of MDS due to mutations in *SGCE*, we developed and characterized a novel *Sgce* gene-trap mouse model that disrupts expression of brain isoforms derived from exon 11b and other exons 3' to exon 9.

## 2. Materials and methods

### 2.1. Developmental expression of *Sgce*

All mouse experiments were performed in accordance with the National Institutes of Health's Guidelines for the Care and Use of Laboratory Animals and approved by our Institutional Animal Care and Use Committee. Expression patterns of *Sgce* transcripts were established in WT mice with relative quantitative reverse transcriptase-PCR (QRT-PCR) using tissues from cerebral cortex, cerebellum, hippocampus, ventral midbrain, striatum, thalamus, spinal cord, and liver. We examined 6 mice (3 male, 3 female) at 9 developmental time points (E15, P1, P7, P14, P21, P30, 2 mo, 6 mo, and 1 yr). Dissection of all brain regions which were readily identifiable in older mice was not possible at E15 and P1. SYBR Green-based QRT-PCR was performed with Ambion's RETROscript® Reverse Transcription Kit and a LightCycler® 480 System (Roche, Indianapolis, IN, USA). One primer pair (Table S2, 85F and 85R) was designed to examine expression of all isoforms (Fig. 1). Expression of other isoforms was established by placing primers in exons 9 and 10 (98F and 98R) or exons 9 and 11b (E9F and E11bR) with  $\beta$ -actin as the endogenous control (Fig. 1 and Table S2). Data was normalized to E15 cerebral cortex. Detailed methods are provided in a previous publication from our laboratory (Xiao et al., 2012).

### 2.2. Generation of *Sgce* gene-trap mice

One line of ES cells on a 129X1/SvJ  $\times$  129S1/Sv background and generated using the Gep-SD5 vector (clone number CW509161 [CMHD-GT\_148G1-3], Fig. 1) was obtained from The Center for Modeling Human Disease (CMHD, Ontario, Canada; [www.cmhd.ca/genetrap/vectors.html](http://www.cmhd.ca/genetrap/vectors.html)). The Gep-SD5 vector does not contain an internal ribosome entry

sequence. Due to the rules of nonsense-mediated decay, the selection transcript is only produced if the insertion site is close to the polyA near the 3' end of the gene. The polyA sequence of the reporter gene (*lacZ*) is deleted which results in an unstable trapped transcript leading to hypomorphic mutants. DNA was extracted from cultured ES cells and used to confirm correct targeting of *Sgce*. Gene-trapped ES cells were injected into the blastocoel cavity of E3.5 C57BL/6 embryos using standard procedures. Male chimeras were bred to C57BL/6J female mice to establish coat color and germline transmission. The insertion site of the gene trap was identified with long-range PCR and confirmed with Sanger sequencing. We established that the terminal exons (3' to exon 9) of *Sgce* were disrupted in homozygotes (*Sgce*<sup>Gt(148G1)Cmhd/Gt(148G1)Cmhd</sup> or *Sgce*<sup>Gt/Gt</sup>) and paternal heterozygotes (*Sgce*<sup>m+/pGt</sup>, m-maternal, p-paternal). Male *Sgce*<sup>m+/pGt</sup> mice were crossed to C57BL/6J female mice for 10 generations. Three primers (Table S2) were used for PCR-based genotyping with one primer located within the gene trap vector (*Sgce\_KO\_V2L*) and the two other primers flanking the gene-trap insertion within intron 9 of *Sgce* (NM\_011360). The Mouse Direct PCR kit from Biotool (Houston, TX, USA) was used for genotyping with the following cycling conditions: 95°C for 5 min; 35 cycles at 95°C for 10 s, 60°C for 30 s, and 72°C for 30 s; and final at 72°C for 7 min. The WT allele yields a 552 bp amplicon whereas the gene-trap mutant allele generates a 393 bp amplicon. *Sgce*<sup>m+/pGt</sup> mice on the C57BL/6J background were crossed to BALB/c and 129S2 mice for 5 or more generations to determine if the occurrence of myoclonus depended on genetic background.

### 2.3. *Sgce* expression in gene-trap mice

Relative levels of mouse *Sgce* mRNA were determined in the cerebral cortex and cerebellum of 1-month-old WT, *Sgce*<sup>m+/pGt</sup>, *Sgce*<sup>mGt/p+</sup>, and *Sgce*<sup>mGt/pGt</sup> mice (n = 3/genotype) using 3 primer pairs targeting different regions of *Sgce* (Fig. 1, Table S2). Sanger sequencing with two reverse primers in *lacZ* and one forward primer in exon 9 (*Sgce\_E9F*) was used to establish in frame splicing of exon 9 to the Gep-SD5 vector sequence, and expression of *Sgce/lacZ* fusion transcript(s) was quantified with a forward primer in exon 9 (*Sgce\_E9F*) and reverse primer in *lacZ* ( $\beta$ -Gal\_R2, Table S2, Fig. 1). The predicted molecular weight of a chimeric  $\epsilon$ -SG/ $\beta$ -gal fusion protein derived from the major  $\epsilon$ -SG isoform and the Gep-SD5  $\beta$ -galactosidase was calculated at ~159 kDa using two independent algorithms (Stothard, 2000, Bjellqvist et al., 1993). Of note, the chimeric protein harbors 25 amino acids between the end of  $\epsilon$ -SG and the first methionine of  $\beta$ -galactosidase (Fig. S1). Mouse  $\beta$ -actin was used as the endogenous control. To ascertain spatial expression patterns, relative levels of mouse *Sgce* mRNA were determined in 6 brain regions (cerebral cortex, cerebellum, hippocampus, striatum, thalamus, and midbrain), cervical spinal cord, and liver harvested from 8 adult mice (3-month old, 4 males and 4 females) of WT, *Sgce*<sup>m+/pGt</sup>, and *Sgce*<sup>mGt/pGt</sup> genotypes. SYBR Green-based QRT-PCR was performed using 3 primer pairs targeting different regions of *Sgce* (Fig. 1, Table S2). Mouse  $\beta$ -actin was used as the endogenous control. *Sgce*<sup>mGt/p+</sup> mice were not included in region of interest analyses since they showed no significant expression differences from WT mice in cerebral cortex and cerebellum (Table S3).

## 2.4. Immunoaffinity purification (IAP) and mass spectrometry analyses

Antibodies used for IAP and Western blotting included pan-e-SG antibody esg3788, pan-e-SG antibody esg3790, and brain-specific antibody esg2-1358 (Waite et al., 2016). The esg2-1358 antibody was generated against a short peptide (NH<sub>2</sub>-C-QRFEVNGIPEERKLTEAMSL-COOH) derived exclusively from exon 11b (Waite et al., 2016). The pan-e-SG antibodies esg3788 and esg3790 were raised in rabbits immunized with a thioredoxin fusion protein containing the entire C-terminal intracellular domain of mouse e-SG (NP\_035490.3) that was used to generate previous e-SG antibodies (Esapa et al., 2007). The anti- $\beta$ -dystroglycan monoclonal antibody MANDAG2, developed by G.E. Morris, was obtained through the Developmental Studies Hybridoma Bank. Whole brain tissues from C57BL/6J WT and *Sgce*<sup>m+/pGt</sup> (all 3-month-old male) were snap frozen in liquid nitrogen and homogenized in digitonin (Merck Chemicals, Darmstadt, Germany) lysis buffer [1% digitonin (w/v), 150mM NaCl, 50mM Tris pH 8.0, 1mM EGTA, 1mM sodium orthovanadate and cOmplete™ Protease Inhibitor Cocktail (Roche, West Sussex, UK)], clarified and pre-cleared on protein A-agarose beads as previously described (Esapa et al., 2007). Lysates were then incubated with antibody-conjugated protein A-agarose beads overnight at 4°C. Proteins were eluted from the beads via incubation in 2° non-reducing Laemmli buffer at 60°C for 30 minutes followed by boiling at 95°C for 5 minutes. DTT was added to the elution at a final concentration of 50 mM, and the elution was boiled at 95°C for 5 min. An aliquot of each IAP elution was analyzed via Western blot for enrichment of e-SG and  $\beta$ -dystroglycan. The remaining eluate was resolved on a 4–12% gradient NuPAGE Novex Bis-Tris gel (Invitrogen, Loughborough, UK) for 10 min, and fixed in 50% (v/v) methanol with 4% (v/v) acetic acid for 15 min. The gel was stained using colloidal Coomassie InstantBlue protein stain (Expedeon, Swavesey, UK) for 1 hr, then destained in ultrapure water before each elution was excised in its entirety as two gel plugs. Gel plugs were sent for in-gel tryptic digest followed by Velos Orbitrap mass spectrometry at the Advanced Mass Spectrometry Facility (University of Birmingham-England). Data analysis was carried out using Proteome Discoverer (Thermo Scientific, Loughborough, UK) with the SEQUEST search algorithm.

## 2.5. X-gal staining

X-gal staining was performed with a LacZ Detection Kit for Tissues (InvivoGen San Diego, CA) following the manufacturer's protocol. Briefly, adult (3-month old) *Sgce*<sup>m+/pGt</sup> mice were transcardially perfused with normal saline followed by 4% paraformaldehyde in 0.1M phosphate buffered saline (PBS) with 2 mM MgCl<sub>2</sub> (pH 7.4). Brains were then placed into a cryoprotection solution (30% sucrose in 0.1M phosphate buffer [PB] with 2 mM MgCl<sub>2</sub>) overnight. Coronal and sagittal cryosections (40  $\mu$ m) were acquired and rinsed in PBS (2mM MgCl<sub>2</sub>, pH 7.4). Floating sections were then incubated with the staining solution (6 mM potassium ferricyanide, 6 mM potassium ferrocyanide, 2 mM MgCl<sub>2</sub>, 0.02% Igepal, 0.01% sodium deoxycholate, and 1 mg/ml X-gal solution in PBS) for 48 hrs at 37°C. Sections then rinsed with PBS, mounted, dehydrated and coverslipped.

## 2.6. Biogenic monoamines

Striatum was acutely harvested from 3 male and 3 female WT and *Sgce*<sup>m+/pGt</sup> mice at 3-months of age. All subsequent processing was completed by the Vanderbilt University Neurochemistry Core. Tissue samples were homogenized with a dismembrator in a solution (pH 3.8) containing 0.1M trichloroacetic acid (TCA), 0.01 M sodium acetate, 0.0001 M EDTA, 5 ng/ml isoproterenol (as internal standard) and 10.5 % methanol. Ten microliters of homogenates were collected for assay of protein concentrations. Samples were then spun in a microcentrifuge at 10,000 g for 20 min and the supernatant was removed for monoamine analysis. Biogenic amines were determined by high-pressure liquid chromatography utilizing an Antec Decade II (oxidation: 0.65) electrochemical detector operated at 33°C. Supernatants (20 µl) were injected using a Water 2707 autosampler onto a Phenomenex Kintex (2.6 u, 100 A) C18 HPLC column (100 × 4.60 mm). Biogenic amines were eluted with a mobile phase consisting of 89.5% 0.1M TCA, 0.01 M sodium acetate, 0.0001 M EDTA and 10.5 % methanol (pH 3.8). Solvents were delivered at 0.6 ml/min using a Waters 515 HPLC pump. Using this HPLC solvent the following biogenic amines were eluted in the following order: noradrenaline, 3, 4-dihydroxyphenylacetic acid (DOPAC); dopamine (DA); 5-hydroxyindoleacetic acid (5-HIAA); homovanillic acid (HVA); 5-hydroxytryptophan (5-HT); and 3-methoxytyramine (3-MT). HPLC control and data acquisition were managed by Empower software.

Protein concentrations were determined with a BCA Protein Assay Kit (ThermoFisher Scientific, Waltham, MA, USA). Ten microliter volumes of tissue homogenate were distributed onto a 96-well plate and 200 µl of mixed BCA reagent (25 ml of Protein Reagent A is mixed with 500 µl of Protein Reagent B) was added. The plate was then incubated for 2 hrs at room temperature for color development. A BSA standard curve was simultaneously run. Absorbance was measured with a POLARstar Omega plate reader (BMG LABTECH, Cary, NC, USA).

## 2.7. Behavioral assessments

Adult (3-month-old) *Sgce*<sup>m+/pGt</sup> mice and gender-matched WT littermates were subjected to a battery of motor and behavioral examinations including open-field activity, rotarod, vertical rope climbing, raised-beam task, grip strength, gait analysis (DigiGait™), dominance tube, and cross-maze test as described in a recent publication from our laboratory (Xiao et al., 2016). Mice were weighted weekly. Video recordings and righting reflex assays were performed prior to weaning in independent groups of WT and *Sgce*<sup>m+/pGt</sup> mice.

WT and *Sgce* mutant mice were observed for the presence of spontaneous and stimulus-induced (audiogenic [100 dB, 40 ms, 1 Hz]; tail, trunk and limb displacement with a microspatula; snout tap; and tail pinch) abnormal movements (tonus, clonus, tremor, twisting, bobbing, wagging, sustained flexion, and sustained extension; Table S4) on a daily basis from Postnatal Day 1 (P1) through P30 and then 3X weekly through 6 months of age. Although assessments were qualitative, abnormal movements were most apparent between P10 and P20, particularly P12 to P16.

An initial cohort of 10 Postnatal Day 14 (P14) mice (all males, 5 WT and 5 *Sgce*<sup>m+/pGt</sup>) was videotaped for 5 min. Each mouse was placed in a clear rectangular arena (18 cm x 30 cm) and subjected to limb, tail and truncal displacements with a microspatula. The videotapes were scored by 3 raters blinded to genotype using an adaption of a previously published rating scale (Raïke et al., 2012). This ordinal scale was utilized to score all abnormal movements in five body regions (face, neck, trunk, forelimbs, and hindlimbs): 0 - absent abnormal movement, 1- slight and intermittent, 2 - mild and common or moderate and intermittent, 3 - moderate and common, or 4 - marked and prolonged (Table S4). In addition, specific types of involuntary movements (tonus, clonus, tremor, twisting, bobbing, wagging, sustained extension, and sustained flexion) were rated in binary fashion as absent (0) or present (1).

A second cohort of 20 P16 mice (WT [n = 10] and *Sgce*<sup>m+/pGt</sup> [n = 10]), (5 males and 5 females for each genotype) was videotaped for a total of 6 min of open-field behavior acquired in 1-min bins and scored by 3 blinded raters. Mice were placed alone in a clear plastic cage (18 cm × 30 cm) for videotaping and returned to their home cages between video sessions.

## 2.8. Whole-genome gene-expression analysis

Total RNA from mouse cerebellum was isolated from 6 adult (10 month-old) *Sgce*<sup>m+/pGt</sup> mice (3 males and 3 females) and 6 age- and gender-matched WT controls (Ambion™ TRI Reagent®, ThermoFisher Scientific). The quality of total RNA was assessed with a NanoDrop® ND-1000 spectrophotometer (NanoDrop Technologies) and RNA integrity was verified with an Agilent 2100 Bioanalyzer using the Agilent RNA 6000 Nano kit. Whole-genome gene-expression data was generated with the Affymetrix GeneChip® Mouse Gene 2.0 ST Array (Santa Clara, CA, USA). This array was designed using data from RefSeq (release 51), Ensembl (release 65) and lincRNA db and provides comprehensive coverage of over 30,000 mRNA transcripts and 2000 lincRNA transcripts. The 2.0 ST Array was designed with a median of 22 25-mer probes per transcript. Each array includes background antigenic probes, poly-A controls and hybridization controls. Target RNA was first reverse transcribed into cDNA, followed by in-vitro transcription to generate biotin-labelled cRNA for subsequent hybridization. Hybridized target cRNA were stained with streptavidin phycoerythrin and scanned using an Affymetrix GeneArray Scanner.

Data were processed using Affymetrix Expression Console software that incorporates the Robust Multi-array Average (RMA) normalization algorithm. Genes were annotated using Affymetrix MoGene 2.0 ST, V.1, release 35 annotation files from NetAffx™ server of Affymetrix. The RMA-normalized .chp files were summarized further with GeneSpring GX® 13.1.1 (Agilent® Technologies, Santa Clara, CA). Scatter plots were used to assess the reproducibility of gene expression within genotypes. A *t*-test statistic was used in identifying significantly dysregulated genes ( $p < 0.05$ ) between WT and *Sgce*<sup>m+/pGt</sup> mice. Differential gene expression was also analyzed using a False Discovery Rate (FDR) corrected  $p < 0.05$  (Benjamini and Heller, 2008). A heat map created using unsupervised hierarchical clustering with average linkage and Euclidean distance and a volcano plot were used to visualize differential gene expression. Using WebGestalt (WEB-based GENE SeT AnaLysis Toolkit)

and Ingenuity Pathway Analysis (IPA), we investigated the effects of differentially regulated genes on pathways (KEGG, Kyoto Encyclopedia of Genes and Genomes) and molecular/cellular networks (Zhang et al., 2005).

To validate data obtained with the Affymetrix GeneChip<sup>®</sup> Mouse Gene 2.0 ST Array, we selected 6 up-regulated and 6 down-regulated genes moderately expressed in cerebellum, and 3 genes related to the DGC, for QRT-PCR. Using TaqMan<sup>®</sup> probes, QRT-PCR was performed on the Roche LightCycler<sup>®</sup> 480 system with primers designed using the Roche Universal ProbeLibrary Assay Design Center. A total of 16 cerebellar RNA samples from WT and *Sgce*<sup>m+/pGt</sup> mice (8 males and 8 females in each group, including the 6 samples used for whole-genome gene-expression analysis) were employed for validation with  $\beta$ -actin as the endogenous control. Technical triplicates were performed for all samples and median values were utilized for statistical analysis.

## 2.9. Statistics

ANOVA with post-hoc tests was used to determine the effects of genotype and gender on parametric behavioral measures. The Mann-Whitney test was used to determine the effects of genotype within gender for a non-parametric behavioral measure (slips on the raised beam task). Two-tailed *t*-tests were used to establish the effects of genotype on monoamine levels in striatum. Fisher's exact test was used to determine the effects of genotype on the results of dominance tube testing. An alpha ( $\alpha$ ) of 0.05 was chosen for statistical significance.

## 3. Results

### 3.1 Expression of *Sgce* is developmentally regulated

Overall, brain expression of total (Fig. 1C) and brain-specific (Fig. 1E) *Sgce* increased with increasing postnatal age ( $p < 0.0001$ , for both). There was a significant effect of region on expression of all *Sgce* isoforms (primers 85F and 85R), "long" isoforms (primers 98F and 98R), and the brain-specific isoform derived, in part, from exon 11b ( $p < 0.0001$ , for all). The highest expression levels for all isoforms and the brain-specific isoform were detected in 2-month-old mouse brain (Fig. 1). In general, cerebellar levels of all isoforms and the brain-specific isoform were 2 to 3 fold higher than hippocampus and cerebral cortex. In older adult mice (1 yr), expression of all isoforms and the brain-specific isoform were similar in cerebellum and striatum. The brain-specific isoform was not detected in liver and total *Sgce* mRNA was very low in this non-neural tissue.

### 3.2 *Sgce* is maternally imprinted

Male and female heterozygous and homozygous mice were fertile and pups of all genotypes and both genders were born at normal Mendelian ratios. As seen in Table 2, there were significant effects of gender ( $F_{1,59} = 135.2$ ,  $p < 0.0001$ ) and genotype ( $F_{1,59} = 78.1$ ,  $p < 0.0001$ ) on weights in 3-month-old mice (Table 2). Overall, *Sgce*<sup>m+/pGt</sup> mice were 10 to 15% smaller than their WT littermates.



Overall, long isoforms comprised roughly 70% of all isoforms in *Sgce*<sup>+/+</sup> mice. More exclusively, the brain-specific isoform derived from exon 11b only comprised 10% of total *Sgce* (Table 3). Total expression of *Sgce* transcripts was reduced by approximately 60 to 70% in *Sgce*<sup>m+/pGt</sup> and *Sgce*<sup>mGt/pGt</sup> mice and unchanged in *Sgce*<sup>mGt/p+</sup> mice (Tables 3 and S3). Sanger sequencing using a forward primer in exon 9 (*Sgce*\_E9F) and two reverse primers in *lacZ* ( $\beta$ \_Gal\_R1 and  $\beta$ \_Gal\_R2) yielded *Sgce/lacZ* fusion amplicons (716 and 329 bp, respectively). QRT-PCR showed that long isoforms including the brain-specific isoform were not expressed in *Sgce*<sup>m+/pGt</sup> and *Sgce*<sup>mGt/pGt</sup> mice (Table 3). However, expression of short *Sgce* isoforms and *Sgce/lacZ* fusion transcript(s) was preserved. Expression levels of fusion transcript(s) detected with random decamers were about twice the levels obtained with oligo(dT) primers (Table 3 and S3).

### 3.3 *Sgce*<sup>m+/pGt</sup> mice express reduced levels of $\epsilon$ -SG

$\epsilon$ -SG has been previously shown to form part of dystrophin-associated protein complexes in the brain that contain  $\beta$ -,  $\delta$ - and  $\zeta$ -sarcoglycans and  $\beta$ -dystroglycan (Waite et al., 2016). Therefore, we performed IAP and mass spectrometry analyses to investigate whether these complexes were disrupted in the *Sgce*<sup>m+/pGt</sup> mice compared to WT littermates. IAPs were performed with two different antibodies to enrich for total  $\epsilon$ -SG (esg3788) or the brain-specific  $\epsilon$ -SG isoform specifically (esg2-1358). Purification of all  $\epsilon$ -SG isoforms was verified by Western blot analysis using an  $\epsilon$ -SG polyclonal antibody (Figs. 2 and S2). Both the ubiquitous and brain-specific  $\epsilon$ -SG isoforms were enriched in the esg3788 IAP and the brain-specific  $\epsilon$ -SG isoforms in the esg2-1358 IAP from WT brain tissue. However, there was a major reduction in detectable  $\epsilon$ -SG in both IAPs from the *Sgce*<sup>m+/pGt</sup> tissue leaving a residual immunoglobulin-related signal of approximately 50 kDa. Western blot of the IAPs using an anti- $\beta$ -dystroglycan antibody demonstrated co-purification of the DGC component  $\beta$ -dystroglycan in the esg3788 IAP from WT tissue and to a lesser extent in the esg2-1358 IAP. By contrast, there was only trace detection of  $\beta$ -dystroglycan in the esg3788 IAP from *Sgce*<sup>m+/pGt</sup> tissue that was absent in the esg2-1358 IAP. With high exposure, a putative  $\epsilon$ -SG/ $\beta$ -gal band was seen at ~159 kDa in the esg3788 IAP from *Sgce*<sup>m+/pGt</sup> tissue (Fig. S2).

The remaining immunoaffinity-purified material from all samples was separated by polyacrylamide gel electrophoresis and stained with colloidal Coomassie blue. Each sample was excised as two gel bands and analysed via mass spectrometry to identify the protein constituents. Multiple high confidence  $\epsilon$ -,  $\beta$ -,  $\delta$ - and  $\zeta$ -sarcoglycan peptides were detected in the esg3788 and esg2-1358 IAPs from WT tissue indicating purification of the entire brain sarcoglycan complex. Only a subset of these peptides was detected in the esg3788 IAP from the *Sgce*<sup>m+/pGt</sup> mice (Table 4). Additionally, the spectral counts for each sarcoglycan were reduced in *Sgce*<sup>m+/pGt</sup> mice versus WT littermates, reflecting a lower abundance of the sarcoglycans in the esg3788 IAP from *Sgce*<sup>m+/pGt</sup> mice (Table S5). However, there was a complete absence of detectable sarcoglycan peptides in the esg2-1358 IAP from *Sgce*<sup>m+/pGt</sup> tissue indicating a loss of brain-specific  $\epsilon$ -SG isoform containing protein complexes.

Both the Western blot and mass spectrometry data from the IAPs indicate a significant reduction in  $\epsilon$ -sarcoglycan protein isoforms in *Sgce*<sup>m+/pGt</sup> mice compared with WT littermates with a complete loss of detectable  $\epsilon$ -SG isoform 2. However, detection of  $\beta$ -

dystroglycan by Western blot and  $\epsilon$ -,  $\beta$ -,  $\delta$ - and  $\zeta$ -sarcoglycans using mass spectrometry in the esg3788 IAP indicate the existence of residual  $\epsilon$ -SG containing sarcoglycan complexes and dystrophin-associated protein complexes in the *Sgce<sup>m+/pGt</sup>* mice.

### 3.4 Sgce is widely expressed in the central nervous system (CNS)

In *Sgce<sup>m+/pGt</sup>* mouse brain, X-gal staining was seen throughout the CNS (Fig. 3). In cerebellum, staining was prominent in the molecular layer with weaker expression in the granule cell layer and cerebellar nuclei (Fig. 3B). In the basal ganglia, staining in the caudate-putamen and lateral globus pallidus was more intense than that seen in the subthalamic nucleus and substantia nigra (Fig. 3C). In cerebral cortex, staining was most prominent within Layer V but also stood out in Layers II and III (Figs. 3A & D). In hippocampus, modest staining intensity was seen in the pyramidal cell layer and dentate gyrus (Fig. 3G).

### 3.5 Sgce<sup>m+/pGt</sup> mice have normal monoamine levels in the striatum

There were no significant effects of gender or genotype on striatal monoamine levels (Table S6). However, noradrenaline levels trended higher in *Sgce<sup>m+/pGt</sup>* mice.

### 3.6. Abnormal gait and appendicular posturing are seen in preweanling Sgce<sup>m+/pGt</sup> mice

No evidence of myoclonus or limb dystonia was noted while *Sgce<sup>m+/pGt</sup>* mice on the C57BL/6J background or mixed C57BL/6JxBABL/c and C57BL/6Jx129S2 backgrounds were routinely observed in their home cages or during open field behavior from the early postnatal period through 1.5 yrs of age. Moreover, myoclonus was not observed in response to audiogenic or tactile stimuli. However, abnormal hindlimb posturing in response to perturbations and “tiptoe” walking were significantly more common in preweanling *Sgce<sup>m+/pGt</sup>* mice (Videos S1 & S2) in comparison with WT littermates. Male and female *Sgce<sup>m+/pGt</sup>* mice had significantly higher scores than WT littermates (Table 5) on the rating scale for abnormal movements (Table S4). With the use of perturbations, the identification of abnormal motor phenotypes showed a low false positive rate. False positive and negative rates were higher with ratings of unperturbed open-field behavior (Table 5).

### 3.7. Sgce<sup>m+/pGt</sup> mice show evidence of mild abnormalities on measures of gait, open-field activity and behavior

In the preweanling period, *Sgce<sup>m+/pGt</sup>* mice and WT littermates showed no righting time differences (Fig. S3). In adult mice, there were significant effects of gender ( $F_{1,60} = 84.1$ ,  $p < 0.0001$ ) and genotype ( $F_{1,60} = 91.0$ ,  $p < 0.0001$ ) on grip strength (Table 2). Male WT mice exhibited greater grip strength than *Sgce<sup>m+/pGt</sup>* littermates (mean difference = 55 g). The effect of genotype was similar in female mice (mean difference = 54 g). However, after normalized by weight, the effect of genotype on grip strength was no longer apparent ( $p = 0.56$ ). There were no effects of genotype on rope climbing or tube dominance. There was a notable effect of genotype ( $F_{1,60} = 15.2$ ,  $p = 0.0003$ ) but no effect of gender or the genotype-gender interaction on cross-maze scores. Male and female *Sgce<sup>m+/pGt</sup>* mice had significantly lower scores than their gender-matched WT littermates on the cross-maze test ( $p < 0.05$ , for both).

*Sgce<sup>m+/pGt</sup>* mice were less active than their WT littermates on multiple parameters open-field activity (Table 2). There were significant genotype\*gender interactions with female mice showing larger effects of genotype on open-field assays of distance traveled, jump count, and ambulatory count ( $p = 0.0001$ ,  $0.0036$ , and  $0.014$ , respectively). There were large effects of genotype on stereotypic counts ( $F_{1,60} = 32.65$ ,  $p < 0.0001$ ) and ambulatory episodes ( $F_{1,60} = 10.89$ ,  $p = 0.0016$ ) but no significant effects of the genotype\*gender interaction.

DigiGait™ analyses showed minor effects of genotype (Table 2) on several gait parameters. Male *Sgce<sup>m+/pGt</sup>* mice had shorter forepaw ( $p = 0.004$ ) and hindpaw ( $p < 0.0001$ ) stride lengths and higher forepaw and hindpaw stride frequencies ( $p < 0.0001$ , for both) than WT gender-matched littermates. These differences did not reach statistical significance in female mice. Forepaw areas were smaller in *Sgce<sup>m+/pGt</sup>* mice than WT littermates ( $F_{1,60} = 17.1$ ,  $p = 0.0001$ ). Overall, hindpaw areas were also smaller in *Sgce<sup>m+/pGt</sup>* mice than WT littermates ( $F_{1,60} = 30.6$ ,  $p < 0.0001$ ) but there was a genotype\*gender interaction ( $F_{1,60} = 18.8$ ,  $p < 0.0001$ ) with male mice showing a more marked effect of the mutant allele. There was a small overall effect of genotype on forepaw stance width ( $F_{1,60} = 4.1$ ,  $p = 0.047$ ). Hindpaw stance width was narrower in *Sgce<sup>m+/pGt</sup>* mice than WT littermates ( $F_{1,60} = 23.33$ ,  $p < 0.0001$ ) although the difference between *Sgce<sup>m+/pGt</sup>* mice and WT females did not reach statistical significance ( $p = 0.060$ ).

On the rotarod, both *Sgce<sup>m+/pGt</sup>* mice and WT littermates showed increased latencies to fall from Day 1 through Day 5 but there was no effect of genotype (Fig. 4A). On the raised-beam task, there were no significant effects of gender or genotype on beam traversal times (Figs. 4B & C). On the 9-mm round beam, male and female *Sgce<sup>m+/pGt</sup>* mice had more slips than gender-matched WT littermates ( $p = 0.016$  and  $p = 0.22$ , respectively). Male and female *Sgce<sup>m+/pGt</sup>* mice also had more slips on the 9-mm square beam than gender-matched WT littermates ( $p = 0.0003$  and  $p = 0.035$ , respectively).

### 3.8. Whole-genome gene expression abnormalities distinguish *Sgce<sup>m+/pGt</sup>* mice from WT littermates

The high linear correlations among scatter plots in male and female WT and *Sgce<sup>m+/pGt</sup>* mice support the reproducibility of our whole-genome gene expression data (Figs. S4 & S5). Differential gene expression was visualized with a heat map (Fig. S6) and volcano plot (Fig. S7). The symmetric volcano plot also attests to the quality of our gene expression analyses. With FDR = 0.05 and fold change (FC) = 2.0, we ended up with only two genes (*Sgce* and *1700016P03Rik*) that were significantly dysregulated (Table 6, Fig. S7). Without correction for familywise error, we found 6 and 5 genes that were down- or up-regulated by 1.5X, respectively (Table 6). At a 1.25X FC, we found 38 and 31 genes that were significantly up- or down-regulated, respectively. QRT-PCR corroborated the direction and relative magnitude of differential expression for 11/12 genes that were assayed (Table 7).

Genes showing the largest up-regulation (Table 6) included the immediate-early gene *Fos* and three members of the nuclear receptor subfamily 4 (*Nr4a3*, *Nr4a2* and *Nr4a1*). Other up-regulated genes encode cell-cycle (*Cdkn1a*) or complement (*Cd55*) proteins or small

nucleolar RNAs (*Snord35a*, *Snora35*, and *Snord14e*). The top down-regulated genes (Table 6) included those encoding an E3-ubiquitin ligase (*Mid1*) and islet cell autoantigen 1 (*Ica1*).

A large collection of genes directly or indirectly associated with the dystrophin-glycoprotein complex were specifically examined for evidence of differential expression between WT and *Sgce*<sup>m+/pGt</sup> cerebellum (Table S7). Based on BioGPS (biogps.org) and the Allen Brain Atlas ([www.brain-map.org](http://www.brain-map.org)), the majority of these genes show significant expression in cerebellum. The largest, albeit modest, FCs were seen for *Stx1A* (syntaxin 1A) and *Cav3* (caveolin). There was no significant up-regulation of other sarcoglycans ( $\alpha$ ,  $\beta$ ,  $\gamma$ ,  $\delta$ , or  $\zeta$ ).

There were few enriched KEGG pathways for up-regulated genes and none for down-regulated genes. The largest collection of significantly up-regulated genes was associated with systemic lupus erythematosus and the ribosome (Table S8). IPA analysis indicated that a relatively large percentage of up- or down-regulated genes are involved in cell cycle, cellular development, and cell death and survival (Table S9). Regulation of eIF4 and p70S6K signaling and mTOR signaling were the top canonical pathway for up-regulated genes. Significant upstream regulators of dysregulated genes are shown in Table S10 and include *Ihnr1*, *Crem*, and *Creb1* for up-regulated genes, and *Tnfrsf4* and *Nfe2l1* for down-regulated genes. Interactome analysis identified NF- $\kappa$ B and TP53 as the major hubs for up- and down-regulated genes, respectively (Fig. 5). Akt and NFE2L2 were secondary hubs for up- and down-regulated genes, respectively.

## 4. Discussion

### 4.1 Splice variants of SGCE and Sgce

Human *SGCE*, located on the reverse strand of Chr. 7q21.3 (CRCh38/hg38), has 4 Consensus CDS isoforms (Fig. 1) and 4 RefSeq isoforms. Mouse *Sgce*, located on the reverse strand of Chr. 6qA1 (CRCm38/mm10) has 4 Consensus CDS isoforms and 5 RefSeq isoforms. The major isoform (NM\_011360) harbors 11 exons with exon 8, but without exons 1a, 11, and 11b; while the brain-specific isoform (NM\_001130189) also harbors 11 exons with exon 11b, but without exons 1a and 8. The gene-trap insertion site within intron 9 of NM\_011360 is predicted to disrupt all isoforms with the exception of AK132210 (9 exons and without exons 1a, 10, 11, and 12) and ENSMUST00000139029. Total *Sgce* expression was reduced by 60 to 70% in our mouse model which exhibited mild motor and cognitive abnormalities along with stimulus-induced appendicular posturing. Presumably, residual expression of *Sgce* was enough to prevent the appearance of overt myoclonus. Amino acids 1 to 317 of mouse and human  $\epsilon$ -SG are located in the extracellular space with only 23% of the full-length protein located in the cytoplasmic space (amino acids 339 – 437). The few *SGCE* mutations causally-associated with DYT11 and localized to the cytoplasmic domain of  $\epsilon$ -SG are frameshifts and likely associated with nonsense-mediated decay (Valente et al., 2005, Asmus et al., 2005). However, it is possible that the failure to include all alternative exons, including exon 11b, in screening efforts has contributed to the relatively low reported frequency of *SGCE* mutations in subjects with MDS (Ritz et al., 2009, Peall et al., 2014, Valente et al., 2005). Moreover, our data suggest that the long isoforms of *Sgce* are required for entirely normal motor and cognitive functioning, at least in mice.

## 4.2 Long isoforms of *Sgce* are widely expressed in the CNS and enriched in cerebellum

Compatible with data reported by BioGPS (BioGPS.org) and Allen Brain Atlas ([www.brain-map.org](http://www.brain-map.org)), we showed that *Sgce* is widely expressed in mouse brain from the early postnatal period through adulthood and modestly enriched in cerebellum in comparison with other sensorimotor structures such as striatum, thalamus and spinal cord. The expression patterns of *Sgce* identified with X-gal staining in our mouse model are similar to those reported in rats with *in situ* hybridization (Xiao and LeDoux, 2003). For the data described herein,  $\beta$ -gal staining was derived from a chimeric e-SG/ $\beta$ -gal fusion protein and appears to represent expression of the major isoform and/or other protein isoforms that are derived from exon 10 and exons 3' to exon 10. The  $\beta$ -gal staining does not incorporate expression of the shortest *Sgce* isoforms. IAP-mass spectrometry and QRT-PCR showed that the major and brain-specific isoforms of *Sgce* were eliminated in *Sgce*<sup>m+/pGt</sup> mice. However, short isoforms terminating 5' to intron 9 and small amounts of a chimeric e-SG/ $\beta$ -gal fusion protein were preserved and may have compensated, at least in part, for absence of the major and brain-specific isoforms.

## 4.3 *Sgce*<sup>m+/pGt</sup> mice exhibit mild behavioral and motor abnormalities

As reported previously in humans and *Sgce* null mice, our *Sgce* mutant mice generated with gene-trap technology showed clear-cut maternal imprinting such that *Sgce* expression from both *Sgce*<sup>mGt/pGt</sup> and *Sgce*<sup>m+/pGt</sup> mice was absent (Yokoi et al., 2005). Male and female *Sgce*<sup>m+/pGt</sup> mice were smaller than WT littermates and showed several abnormalities on behavioral and motor testing. Male and female *Sgce*<sup>m+/pGt</sup> mice showed reduced scores on the cross-maze test. Based on the results of rope climbing, rotarod and open-field activity, it is unlikely that the significantly lower scores on the cross-maze test were due to motor deficits. Instead, the cross-maze scores are a likely manifestation of increased anxiety in *Sgce*<sup>m+/pGt</sup> mice although contributions from cognitive deficits cannot be excluded (Salimov et al., 1995, Salimov et al., 1996). Anxiety is a well-known psychiatric co-morbidity in human myoclonus-syndrome due to mutations in *SGCE* (Peall et al., 2013, Peall et al., 2016). Accordingly, it is possible that deleterious sequence variants localized to terminal exons of *SGCE* could be risk factors for human anxiety even if not associated with dystonia and/or myoclonus.

Dystonia or dystonic-like posturing was not described in previously published murine models of DYT11 (Yokoi et al., 2005, Yokoi et al., 2006, Yokoi et al., 2012a, Yokoi et al., 2012b). Stimulus-induced flexion posturing was seen in our preweanling *Sgce*<sup>m+/pGt</sup> mice mainly between P14 and P16, a key period for development of cerebellar Purkinje cells and cerebellar cortex (Kano and Hashimoto, 2012). In theory, Purkinje cells and/or local area networks could be unstable until additional maturation has been completed. In this regard, stimulus-induced dystonia, seen in humans with paroxysmal kinesigenic dyskinesias due to mutation in *PRRT2*, often becomes manifest during childhood (Erro et al., 2014), and transient dystonia of infancy usually manifests between 5 and 10 months of age and then gradually disappears after 3 months to 5 years (Bonnet et al., 2010). Given that dystonia remains an inexact clinical diagnosis, both in mice and men, we have restricted our description to flexion posturing. We could not illicit abnormal posturing in postweanling mice or extensor posturing in preweanling mice. *Sgce*<sup>m+/pGt</sup> mice also exhibited a “tiptoe”

gait with raised caudal trunk and narrowed hindlimb stance widths. The narrowed hindlimb stance widths persisted into adulthood (Table 2). Tiptoe walking is not specific to dystonia or *Sgce*<sup>m+/pGt</sup> mice and has also been described in Yoshimura mice, a model of chronic cervical myelopathy (Wang et al., 2014). However, early gait abnormalities may be common in humans with *SGCE* mutations (Asmus et al., 2005). In one study, for example, all *SGCE* mutations carriers had a gait disorder, unsteadiness or frequent falls before 18 months of age (Asmus et al., 2005).

Our gene-trap *Sgce* mouse model showed some similarities and a few important differences from previously reported *Sgce* null and conditional KO mice, as well as *Tor1a-Sgce* double null mice. All *Sgce* models have shown impaired performance on the raised-beam task with increased numbers of slips (Yokoi et al., 2006, Yokoi et al., 2012a, Yokoi et al., 2012b, Yokoi et al., 2010). *Sgce* null mice had increased vertical open-field motor activity, anxiety-like behavior, myoclonus and increased striatal dopamine (Yokoi et al., 2006). In comparison, our gene-trap model also showed evidence of anxiety but exhibited reduced vertical open-field motor activity, had no evidence of myoclonus and striatal dopamine levels were normal. In addition, slips on the raised-beam task are not specific to *Sgce* models and have been reported in *Tor1a*, *Thap1*, and *Ciz1* models of dystonia (Yokoi et al., 2015, Ruiz et al., 2015, Xiao et al., 2016, Zhao et al., 2008). Increased slips on the raised beam task have been reported in heterozygous *Tor1a* null mice and hMT1 transgenic mice that overexpress mutant human torsinA (Zhao et al., 2008, Dang et al., 2005, Song et al., 2014).

#### 4.4 Gene expression abnormalities in *Sgce*<sup>m+/pGt</sup> mice

IAP-mass spectrometry showed that brain-specific e-SG interacts with  $\beta$ -SG,  $\delta$ -SG, and  $\zeta$ -SG. Gene-expression data did not show evidence for their compensatory up-regulation in *Sgce*<sup>m+/pGt</sup> mice. Furthermore, there was no apparent up-regulation for other components of the DGC at the transcript level. Given the modest phenotypes in our *Sgce*<sup>m+/pGt</sup> mice, it seems that short isoforms of e-SG were able to partially compensate for loss of the major and brain-specific isoforms and other long isoforms.

Based on IPA analysis, it appears that up- and/or down-regulation of genes in various cell cycle, cellular development, and cell death and survival pathways served to compensate for e-SG deficiency in our mouse model. However, we cannot exclude the possibility that some of these changes in gene expression were not due to novel effects of mutant e-SG/ $\beta$ -gal fusion protein(s) that resulted from insertion of the Gep-SD5 gene-trap vector into intron 9 of *Sgce*. Although a single-pass transmembrane cell-surface protein seemingly involved in synaptic transmission, our gene expression data suggests that e-SG shows functional overlap with other dystonia proteins such as TAF1, CIZ1, torsinA, THAP1, and G $\alpha$ (olf) which play direct or indirect roles in cell cycle control, DNA repair and neurodevelopment (LeDoux et al., 2013, Xiao et al., 2016).

Several relatively distinct groups of genes showed up-regulation (*Nr4a3*, *Fos*, *Nr4a2*, *Nr4a1*, *Cdkn1a*, *Cd55*, *Snord35a*, *Snora35* and *Snord14e*) or down-regulation (*Mid1* and *Icaf1*). Intermittent subclinical involuntary movements and/or anxiety could be responsible for differential up-regulation of the immediate-early genes *Fos*, *Nr4a3*, *Nr4a2*, and *Nr4a1* in *Sgce*<sup>m+/pGt</sup> mice (Huguet et al., 2016). CDKN1A is an essential cell-cycle protein that

interacts with the dystonia-associated protein CIZ1 (LeDoux et al., 2013, Xiao et al., 2012) and CD55 regulates the complement system, an important player in neuronal differentiation (Stephan et al., 2012). *Snord35a*, *Snora35* and *Snord14e* encode small nucleolar RNAs, a class of RNA molecules that contribute to modifications of rRNA and tRNAs such as methylation and pseudouridylation. MID1 functions as a ubiquitin ligase and regulator of mTOR signaling and plays an important role in neurodevelopment and neurodegeneration. Lack of MID1 causes abnormal development of the cerebellum (Lancioni et al., 2010).

Top up-regulated canonical pathways (eIF4, p7056K, mTOR) and IPA nodes (NF- $\kappa$ B, Akt) are related to intracellular pathways regulating the cell cycle. The Akt/mTOR component of this intracellular network has been shown to regulate neurite outgrowth in cerebellar granule cells (Okada et al., 2011) and the morphology of Purkinje cells (Thomanetz et al., 2013). The major hubs for down-regulated genes, TP53 and NFE2L2, can also be readily linked to cell cycle control and neurodevelopment. In cerebellum, elevation of NFE2L2 (alias NRF2) levels is beneficial in a mouse model of Alexander disease caused by autosomal mutation in GFAP (LaPash Daniels et al., 2012). TP53 regulates CDKN1A at the G1/S cell-cycle checkpoint. In aggregate, gene expression studies suggest that compensatory up- or down-regulation of genes related to neurodevelopment and neuronal morphology may compensate for loss of major and brain-specific e-SG isoforms.

## 5. Conclusions

*Sgce<sup>m+/pGt</sup>* mice show germline absence of major and brain-specific isoforms harboring exons 3' to intron 9. *Sgce* was maternally-imprinted in our mouse model. *Sgce<sup>m+/pGt</sup>* mice exhibited tiptoe walking and occasional stimulus-induced appendicular flexion posturing in the preweanling period from P10 to P20. Adult *Sgce<sup>m+/pGt</sup>* mice showed increased slips on a raised-beam task, anxiety-like behavioral abnormalities, and narrowed hindlimb stance widths but no myoclonus or differences in striatal monoamines in comparison with WT littermates. Overall, adult *Sgce<sup>m+/pGt</sup>* mice were less active than WT littermates. There was no evidence for up-regulation of other sarcoglycans in *Sgce<sup>m+/pGt</sup>* mice. Genes differentially expressed between *Sgce<sup>m+/pGt</sup>* mice and WT littermates were involved in cell cycle, cellular development, and cell death and survival. Our gene-trap mice provide novel insights into the biology and pathobiology of e-SG.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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### Highlights

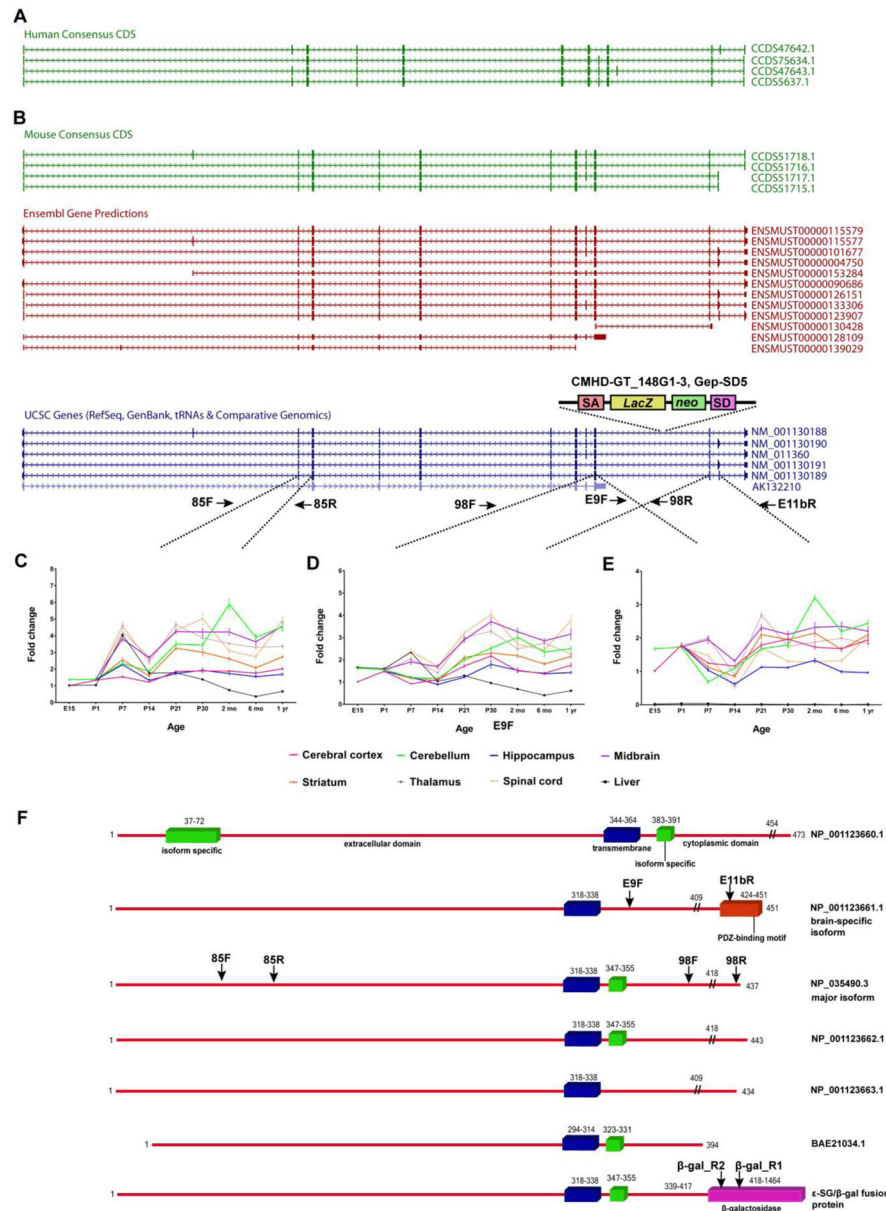
We generated a gene-trap mouse model with reduced expression of e-sarcoglycan.

*Sgce* mutant mice exhibit a tiptoe gait and appendicular posturing.

*Sgce* mutant mice have altered gait dynamics and reduced open-field activity.

In the cerebellum, dysregulated genes were involved in cell cycle and development.

Short *Sgce* isoforms compensate for deficiency of major and brain-specific isoforms.



**Fig. 1.** (A) Human *SGCE* Consensus CDS isoforms. (B) Mouse *Sgce* isoforms derived from Consensus CDS, Ensembl gene predictions and UCSC genes showing the location of the CMHD-GT\_14G1-3 Gep-SD5 gene trap. Three distinct primer pairs (C, 85F/85R; D, 98F/98R; and E, E9F/E11bR) were designed to examine *Sgce* expression in eight different tissues across nine developmental time points. (F) Protein structures of the major (NP\_035490.3), longest (NP\_001123660.1), brain-specific (NP\_001123661), and shortest (BAE21034.1) mouse isoforms, and predicted chimeric  $\epsilon$ -SG/ $\beta$ -gal fusion protein derived from the major isoform. Blue bars, transmembrane domain. Green bars, isoform specific domains. Red bar, protein sequence derived from exon 11b that includes a PDZ-binding motif. Pink bar,  $\beta$ -galactosidase, predicted sites of  $\epsilon$ -SG/ $\beta$ -gal protein fusion. Also shown

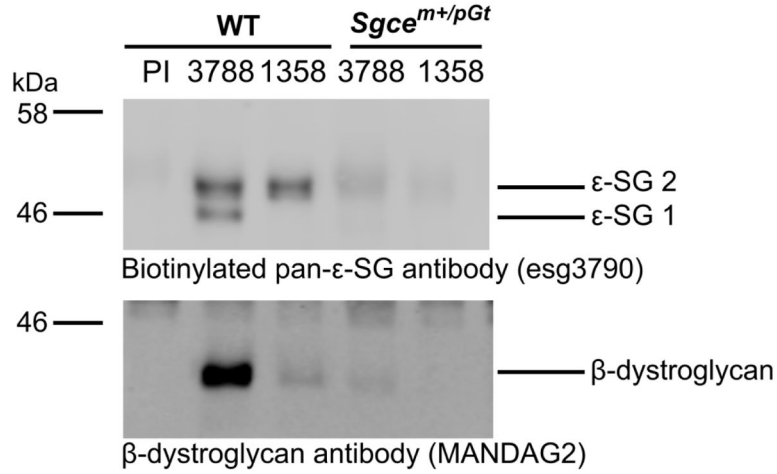
are the approximate locations of primers that would amplify the corresponding regions of *Sgce* and hybrid transcripts. SA, splice acceptor. *lacZ*, gene encoding  $\beta$ -galactosidase. *neo*, neomycin resistance gene. SD, splice donor.

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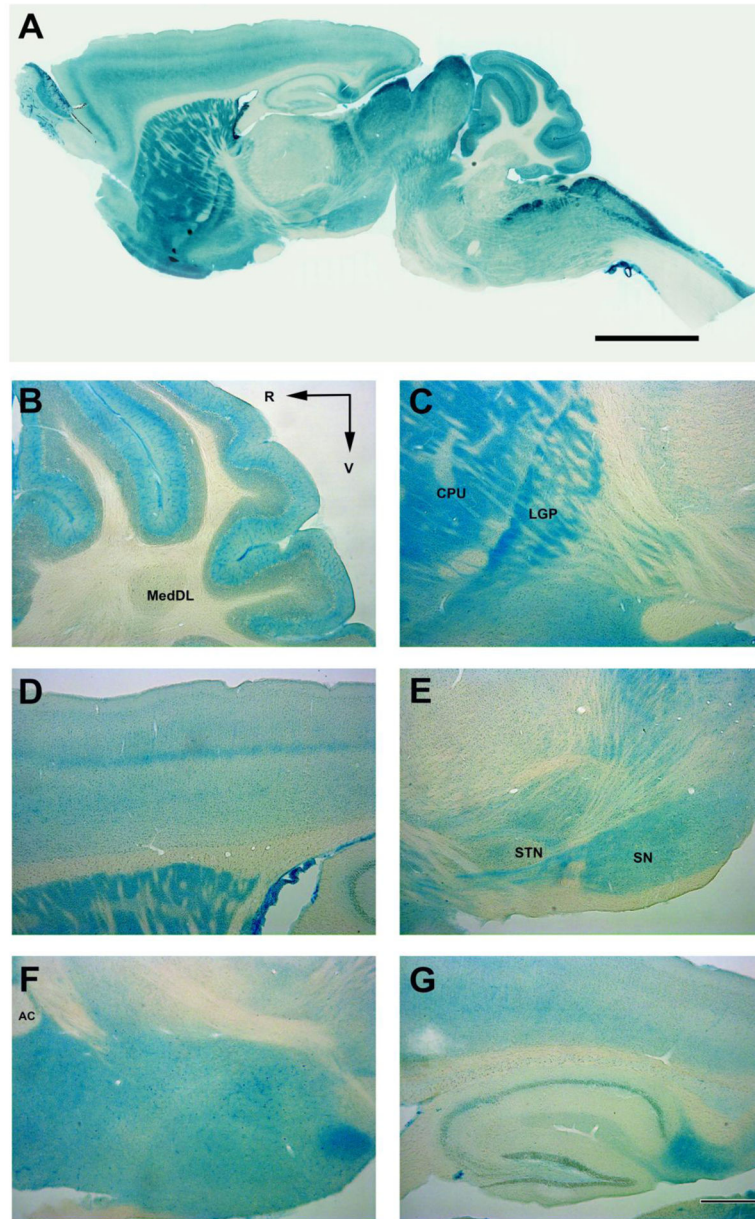
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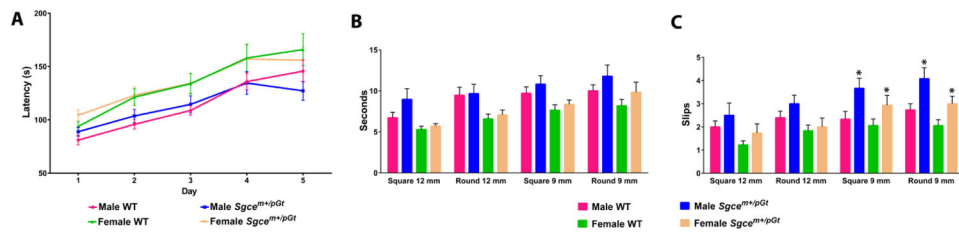
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**Fig. 2.** Western blots of immunoaffinity purifications (IAPs) of ε-SG-containing complexes from WT C57BL/6J and *Sgce*<sup>m+/pGt</sup> mice. The left three lanes contain proteins isolated from WT brain tissue whereas the right two lanes contain protein from *Sgce*<sup>m+/pGt</sup> brain tissue. Lanes are labelled with the antibody used to isolate proteins: PI, preimmune IgG; 3788, esg3788 anti-ε-SG; 1358, esg2-1358 anti-ε-SG brain-specific isoform. The biotinylated anti-ε-SG antibody detects both the major isoform (lower band, ε-SG 1) and brain-specific isoform (upper bands, ε-SG 2) of ε-SG in IAPs using the esg3788 anti-ε-SG antibody, and only the brain-specific isoform in IAPs using the esg2-1358 anti-ε-SG brain-specific isoform antibody. Western blot using the anti-β-dystroglycan antibody MANDAG2 showing co-purifications of β-dystroglycan in the esg3788 and esg-1358 IAPs from WT tissue. β-dystroglycan is severely reduced in the esg3788 IAP from *Sgce*<sup>m+/pGt</sup> brain tissue and absent in the esg-1358 IAP.

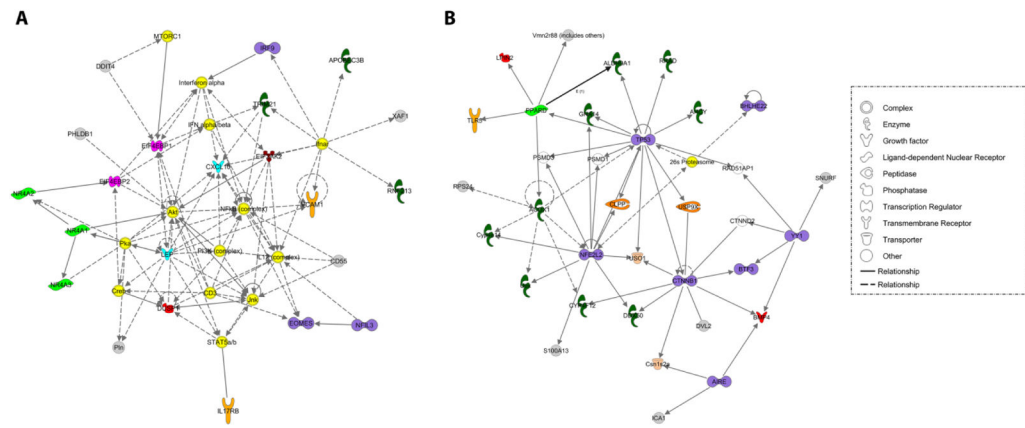


**Fig. 3.** X-gal expression in *Sgce*<sup>m+/pGt</sup> mice. A parasagittal section (A) of *Sgce*<sup>m+/pGt</sup> mouse brain, showing intense reaction in the gray matter of cerebellar cortex, striatum and cerebral cortex. (B) Cerebellum. (C) Striatum. (D) Cerebral cortex. (E) Ventral midbrain. (F) Hypothalamus. (G) Hippocampus. R, rostral; V, ventral; AC, anterior commissure; MedDL, medial cerebellar nucleus, dorsolateral protuberance; CPU, caudate putamen; LGP, lateral globus pallidus; STN, subthalamic nucleus; SN, substantia nigra. Scale bar = 400  $\mu$ m.

**Fig. 4.**

Performance of *Sgce*<sup>m+/pGt</sup> mice (n = 15 males and 15 females) and WT littermates (n = 18 males and 16 females) on the rotarod (A) and raised beam tasks (B and C). Beam traversal times (B) and slips (C) were recorded for 4 different beams. \*significant effect of genotype within gender ( $p < 0.05$ )





**Fig. 5.** Ingenuity interactome analysis of up-regulated (A) and down-regulated (B) genes in *Sgce*<sup>m+/pGt</sup> mouse cerebellum.

**Table 1***SGCE* variant screening in myoclonus-dystonia syndrome

<b>Study</b>	<b>Subjects screened (positive)</b>	<b>Exon 11b screened</b>	<b>Mutation locations</b>
Han et al (2003)	7 families (3)	No	Exons 3 and 7
Schüle et als (2004)	10 families (2)	No	Exons 2 and 7
Valente EM et al. (2005)	58 index patients (6)	No	Exons 3, 4, 5 and 9
Tezenas du Montcel et al. (2006)	76 index patients (16)	Yes	Exons 2, 3, 4, 6, 7, and 9
Gerrits et al. (2006)	31 index patients (7)	No	Exons 2, 3, 5, 6, 7
Nardocci N et al. (2008)	11 families (9)	No	Exons 3, 4, 6, and 7
Ritz K et al. (2009)	86 index patients (13)	No	Exons 2, 3, 5, 6, and 7
Asmus F et al. (2009)	23 index patients (7)	Yes	Exons 3, 6, 7, and 10
Carecchio M et al. (2013)	46 new index patients (8)	No	Exons 3, 4, 6, and 9
Peall KJ et al. (2014)	89 index patients (19)	Yes	Exons 3, 5, 6, 7 and 9

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**Table 2**

Effects of genotype and gender on weight and behavioral measures in 3-month-old mice. Ambulatory count, the total number of X + Y photo beam breaks while in ambulatory movement status. Stereotypic count, any partial-body movements that occur within the ambulatory box such as grooming, head-weaving or scratching. Vertical count, number of periods of continuous Z photo beam breaks. Jump count, the number of times that the mouse leaves the photo beam array for a period of time. Ambulatory episodes, the number of times the mouse has started moving after the resting delay has expired. Values are means  $\pm$  SEM except for dominance tube.

	Male		Female	
	<i>Sgce</i> <sup>+/+</sup> (n=18)	<i>Sgce</i> <sup>m+/pGt</sup> (n=15)	<i>Sgce</i> <sup>+/+</sup> (n=16)	<i>Sgce</i> <sup>m+/pGt</sup> (n=15)
Weight (g)	31.2 $\pm$ 0.7	26.3 $\pm$ 0.5 *	24.9 $\pm$ 0.4	20.2 $\pm$ 0.4 *
Grip strength (g)	329.2 $\pm$ 6.9	274.3 $\pm$ 5.8 *	276.4 $\pm$ 4.5	222.1 $\pm$ 4.8 *
Grip strength/weight	10.6 $\pm$ 0.3	10.4 $\pm$ 0.2	11.2 $\pm$ 0.2	11.0 $\pm$ 0.3
Dominance tube	53.3%	46.7%	49.0%	47.1%
Cross maze score (%)	33.5 $\pm$ 2.3	25.4 $\pm$ 2.0 *	33.1 $\pm$ 2.5	23.5 $\pm$ 2.3 *
Rope climbing (s)	5.4 $\pm$ 0.5	4.5 $\pm$ 0.4	4.1 $\pm$ 0.6	4.3 $\pm$ 0.4
Rope climbing/weight	0.17 $\pm$ 0.01	0.17 $\pm$ 0.01	0.16 $\pm$ 0.02	0.21 $\pm$ 0.02
Open field activity				
<i>Distance traveled (cm)</i>	1858.4 $\pm$ 52.0	1594.7 $\pm$ 30.4 *	2455.7 $\pm$ 53.2	1798.7 $\pm$ 43.3 *
<i>Ambulatory count</i>	897.4 $\pm$ 32.7	782.1 $\pm$ 18.5 *	1168.4 $\pm$ 31.4	904.2 $\pm$ 31.3 *
<i>Stereotypic Count</i>	2596.8 $\pm$ 102.3	2245.1 $\pm$ 42.1 *	2607.0 $\pm$ 49.5	2183.5 $\pm$ 56.8 *
<i>Vertical count</i>	189.9 $\pm$ 11.2	187.5 $\pm$ 15.0	135.3 $\pm$ 10.7	101.4 $\pm$ 10.2 *
<i>Jump count</i>	35.1 $\pm$ 2.9	38.3 $\pm$ 2.7	45.9 $\pm$ 4.1	29.5 $\pm$ 2.4 *
<i>Average velocity (cm/s)</i>	36.2 $\pm$ 1.5	34.4 $\pm$ 1.6	43.5 $\pm$ 1.9	45.2 $\pm$ 2.1
<i>Ambulatory episodes</i>	85.4 $\pm$ 6.0	75.9 $\pm$ 6.4	112.9 $\pm$ 6.2	85.2 $\pm$ 3.7 *
DigiGait™				
<i>Propel (s) Forelimb</i>	0.127 $\pm$ 0.003	0.116 $\pm$ 0.003 *	0.116 $\pm$ 0.004	0.110 $\pm$ 0.004
<i>Propel (s) Hindlimb</i>	0.195 $\pm$ 0.003	0.168 $\pm$ 0.002 *	0.175 $\pm$ 0.004	0.165 $\pm$ 0.002 *
<i>Stride length (cm) Forelimb</i>	7.06 $\pm$ 0.13	6.38 $\pm$ 0.10 *	6.56 $\pm$ 0.15	6.44 $\pm$ 0.11
<i>Stride length (cm) Hindlimb</i>	7.22 $\pm$ 0.09	6.45 $\pm$ 0.08 *	6.75 $\pm$ 0.10	6.57 $\pm$ 0.06
<i>Stride Frequency (steps/s) Forelimb</i>	2.87 $\pm$ 0.04	3.29 $\pm$ 0.03 *	2.92 $\pm$ 0.06	3.04 $\pm$ 0.05
<i>Stride Frequency (steps/s) Hindlimb</i>	2.81 $\pm$ 0.04	3.09 $\pm$ 0.04 *	2.86 $\pm$ 0.05	3.00 $\pm$ 0.02 *
<i>Stance width (cm) Forelimb</i>	1.58 $\pm$ 0.02	1.49 $\pm$ 0.03 *	1.44 $\pm$ 0.03	1.40 $\pm$ 0.03
<i>Stance width (cm) Hindlimb</i>	2.60 $\pm$ 0.05	2.35 $\pm$ 0.04 *	2.30 $\pm$ 0.02	2.21 $\pm$ 0.02 *
<i>Step angle (deg) Forelimb</i>	64.87 $\pm$ 1.83	65.81 $\pm$ 1.74	67.79 $\pm$ 1.49	65.63 $\pm$ 1.66
<i>Step angle (deg) Hindlimb</i>	54.77 $\pm$ 1.62	55.93 $\pm$ 0.99	61.04 $\pm$ 1.32	59.54 $\pm$ 1.33
<i>Paw Area (cm<sup>2</sup>) Forelimb</i>	0.23 $\pm$ 0.01	0.20 $\pm$ 0.01 *	0.22 $\pm$ 0.01	0.19 $\pm$ 0.01 *

	Male		Female	
	<i>Sgce</i> <sup>+/+</sup> (n=18)	<i>Sgce</i> <sup>m+/pGt</sup> (n=15)	<i>Sgce</i> <sup>+/+</sup> (n=16)	<i>Sgce</i> <sup>m+/pGt</sup> (n=15)
<i>Paw Area (cm<sup>2</sup>) Hindlimb</i>	0.46 ± 0.01	0.38 ± 0.01 *	0.41 ± 0.01	0.40 ± 0.01

\*  $p < 0.05$ , for effect of genotype within gender.

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**Table 3**

*Sgce* expression in 3-month-old gene-trap mice using relative quantitative reverse transcriptase PCR (QRT-PCR). Primers 1 (85F/85R), all isoforms. Primers 2 (98F/98R), long isoforms. Primers 3 (E9F/E1 IbR), brain-specific isoform. Primers 4 (E9F/β-gal\_R2), *Sgce/lacZ* fusion transcript. Values are referenced to wild-type (*Sgce*<sup>+/+</sup>) cerebral cortex and presented as means ± standard error of the mean (SEM) (n = 8 mice/genotype).

Tissue	<i>Sgce</i> <sup>+/+</sup>				<i>Sgce</i> <sup>mi/lpCr</sup>				<i>Sgce</i> <sup>mi/lpCr</sup>			
	Primers 1	Primers 2	Primers 3	Primers 4	Primers 1	Primers 2	Primers 3	Primers 4	Primers 1	Primers 2	Primers 3	Primers 4
Cerebral cortex	1.00 ± 0.02	0.70 ± 0.05	0.09 ± 0.01	0.00 ± 0.00	0.33 ± 0.04	0.00 ± 0.00	0.00 ± 0.00	0.15 ± 0.04	0.32 ± 0.03	0.00 ± 0.00	0.00 ± 0.00	0.18 ± 0.03
Cerebellum	2.70 ± 0.09	1.38 ± 0.08	0.17 ± 0.03	0.00 ± 0.00	1.09 ± 0.08	0.00 ± 0.00	0.00 ± 0.00	0.35 ± 0.03	0.99 ± 0.08	0.00 ± 0.00	0.00 ± 0.00	0.53 ± 0.04
Hippocampus	0.92 ± 0.02	0.62 ± 0.05	0.07 ± 0.01	0.00 ± 0.00	0.27 ± 0.04	0.00 ± 0.00	0.00 ± 0.00	0.15 ± 0.03	0.24 ± 0.02	0.00 ± 0.00	0.00 ± 0.00	0.14 ± 0.03
Midbrain	1.77 ± 0.05	1.27 ± 0.06	0.12 ± 0.01	0.00 ± 0.00	0.80 ± 0.09	0.00 ± 0.00	0.00 ± 0.00	0.38 ± 0.05	0.71 ± 0.06	0.00 ± 0.00	0.00 ± 0.00	0.30 ± 0.04
Striatum	1.27 ± 0.02	0.86 ± 0.05	0.11 ± 0.02	0.00 ± 0.00	0.46 ± 0.04	0.00 ± 0.00	0.00 ± 0.00	0.24 ± 0.03	0.39 ± 0.03	0.00 ± 0.00	0.00 ± 0.00	0.21 ± 0.03
Thalamus	1.47 ± 0.04	1.05 ± 0.04	0.10 ± 0.01	0.00 ± 0.00	0.60 ± 0.07	0.00 ± 0.00	0.00 ± 0.00	0.29 ± 0.03	0.57 ± 0.04	0.00 ± 0.00	0.00 ± 0.00	0.27 ± 0.04
Spinal cord	1.38 ± 0.02	1.11 ± 0.07	0.07 ± 0.00	0.00 ± 0.00	0.59 ± 0.05	0.00 ± 0.00	0.00 ± 0.00	0.25 ± 0.03	0.53 ± 0.03	0.00 ± 0.00	0.00 ± 0.00	0.23 ± 0.03
Liver	0.39 ± 0.02	0.28 ± 0.03	0.00 ± 0.00	0.00 ± 0.00	0.01 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.01 ± 0.00	0.01 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.01 ± 0.00

**Table 4**

Unique sarcoglycan peptides identified in wild-type and *Sgce*<sup>m+/pGt</sup> whole brain e-SG immunoaffinity purifications (IAPs). For each IAP (wild-type versus *Sgce*<sup>m+/pGt</sup> tissue, esg3788 antibody versus esg2-1358 antibody), all unique tryptic sarcoglycan peptides detected via mass spectrometry are included in the table. The e-SG2 brain-specific peptides detected in wild-type tissue are in bold. Abbreviations: IAP, immunoaffinity purification; esg3788, anti-e-SG antibody 3788; esg2-1358, anti-e-SG2 antibody 1358.

Protein	esg3788 IAP		esg2-1358 IAP	
	Wild-type	<i>Sgce</i> <sup>m+/pGt</sup>	Wild-type	<i>Sgce</i> <sup>m+/pGt</sup>
<b>e-SG</b>	TPYSDGVLYGSPTAENVGKPTIIEITAYNRR			
	QVSTYQEVVR	QVSTYQEVVR	QVSTYQEVVR	
	EVENPQNQLR		EVENPQNQLR	None
	<b>FEVNGIPEER</b>			
	<b>KLTEAmSL</b>			
<b>β-SG</b>	RNENLVITGNNQPIVFQQTGTTK	RNENLVITGNNQPIVFQQTGTTK	RNENLVITGNNQPIVFQQTGTTK	
	LPSSSSGDQSGSGDWVR		LPSSSSGDQSGSGDWVR	
	THNILFSTDYETHEFHLPSTGK			
	TSITSDIGmQFFDPR	TSITSDIGmQFFDPR	TSITSDIGmQFFDPR	None
	LcMcADGTLFK		LcMcADGTLFK	
	GNEGVFIMGK		GNEGVFIMGK	
<b>δ-SG</b>	GVEINAEAGNMEAIcR		GVEINAEAGNMEAIcR	
	LEGDSEFLQPLYAK	LEGDSEFLQPLYAK	LEGDSEFLQPLYAK	
	LLFSADDSEVVVGAER	LLFSADDSEVVVGAER	LLFSADDSEVVVGAER	
	VLGAEGTVFPK	VLGAEGTVFPK	VLGAEGTVFPK	None
	VFEVcVcANGR		VFEVcVcANGR	
	VLTQLVTGPK	VLTQLVTGPK	VLTQLVTGPK	
	SRPGNALYFK			
	SLVMEAPK			
<b>ζ-SG</b>	ELHLQSTEGEIFLNADSIR		ELHLQSTEGEIFLNADSIR	
	VLFSADEDEITIGA EK		VLFSADEDEITIGA EK	
	LEGISEFLLPLYVK	LEGISEFLLPLYVK	LEGISEFLLPLYVK	
	LGNLPIGSFSSSTSSNSR		LGNLPIGSFSSSTSSNSR	None
	QTVYELcVcPhGK		QTVYELcVcPhGK	
	GVQVSAAGDFK	GVQVSAAGDFK	GVQVSAAGDFK	
			STDLDIQELK VTGTEGAVFGH SVETPHIR	

**Table 5**

Blinded video analyses of abnormal movements in preweanling *Sgce<sup>m+/pGt</sup>* mice and WT littermates at P14 and P16.

Sex, Genotypes (numbers, age)	Total score <sup>a</sup> (mean ± SEM)	Number of genotype-phenotype matches <sup>b</sup>	Percent correct
M, WT (n=5, P14)	1.7 ± 1.2	4/5	80.0
M, <i>Sgce<sup>m+/pGt</sup></i> (n=5, P14)	8.3 ± 3.3 *	5/5	100.0
M, WT (n=10, P16)	18.7 ± 4.7	22/30	73.3
M, <i>Sgce<sup>m+/pGt</sup></i> (n=10, P16)	36.0 ± 5.3 *	18/30	60.0
F, WT (n=10, P16)	20.0 ± 8.1	23/30	76.7
F, <i>Sgce<sup>m+/pGt</sup></i> (n=10, P16)	39.0 ± 10.4 *	22/30	73.3

<sup>a</sup> Mean total scores for abnormal movements were generated by three blinded raters.

<sup>b</sup> Genotype-phenotype matches were declared if 2/3 of the raters scored the presence of abnormal movements in *Sgce<sup>m+/pGt</sup>* mice or if 1/3 scored abnormal movements in WT littermates.

\*  $p < 0.05$ .

**Table 6**Genes significantly up- and down-regulated in *Sgce*<sup>m+/pGt</sup> mice

Affymetrix Cluster ID	Gene Symbol	Gene Name	Fold Change	1.25 & p	0.05)
17400862	<i>Hmgcs2</i>	3-hydroxy-3-methylglutaryl-Coenzyme A synthase 2	1.250		
17522876	<i>Eomes</i>	Eomesodermin homolog (Xenopus laevis)	1.250		
17347355	<i>Eif2ak2</i>	Eukaryotic translation initiation factor 2-alpha kinase 2	1.255		
17436237	<i>Fosl2</i>	FOS-like antigen 2	1.255		
17301615	<i>Gm21685</i>	Predicted gene, 21685	1.256		
17259078	<i>Rnf213</i>	Ring finger protein 213	1.256		
17222819	<i>9330175M20Rik</i>	RIKEN cDNA 9330175M20 gene	1.262		
17300591	<i>Irf9</i>	Interferon regulatory factor 9	1.264		
17473666	<i>Vmn1r67</i>	Vomer nasal 1 receptor 67	1.266		
17435577	<i>LOC102638085</i>		1.268		
17234423	<i>Derl3</i>	Der1-like domain family, member 3	1.274		
17403237	<i>Gbp3</i>	Guanylate binding protein 3	1.275		
17411147	<i>Ifi44</i>	Interferon-induced protein 44	1.284		
17547813	<i>Gm19845</i>	Predicted gene, 19845	1.289		
17350009	<i>Gm24690</i>	Predicted gene, 24690	1.290		
17487489	<i>Pvr</i>	Poliovirus receptor	1.293		
17516383	<i>Snord14e</i>	Small nucleolar RNA, C/D box 35A	1.303		
17538425	<i>Snora35</i>	Small nucleolar RNA, H/ACA box 35	1.305		
17292632	<i>n-R5s54</i>	RNA, 5S Ribosomal Pseudogene 212	1.307		
17320684	<i>Gm24647</i>	Predicted gene, 24647	1.318		
17261033	<i>Gm22753</i>	Predicted gene, 22753	1.327		
17481504	<i>Olfir713</i>	Olfactory receptor 713	1.341		
17226622	<i>Cd55</i>	CD55 antigen	1.344		
17490622	<i>Snord35a</i>	Small Nucleolar RNA, C/D Box 35A	1.359		
17455903	<i>Asb4</i>	Ankyrin repeat and SOCS box-containing 4	1.360		
17290072	<i>Gm25985</i>	Predicted gene, 25985	1.373		
17290072	<i>Traj13</i>	T Cell Receptor Alpha Joining 13	1.389		
17361435	<i>Npas4</i>	Neuronal PAS domain protein 4	1.394		
17315178	<i>Nr4a1</i>	Nuclear receptor subfamily 4, group A, member 1	1.404		
17335467	<i>Cdkn1a</i>	Cyclin-dependent kinase inhibitor 1A (P21)	1.410		
17464654	<i>Pdk4</i>	Pyruvate dehydrogenase kinase, isoenzyme 4	1.426		
17342642	<i>Dusp1</i>	Dual specificity phosphatase 1	1.453		
17385374	<i>Nr4a2</i>	Nuclear receptor subfamily 4, group A, member 2	1.494		
17358658	<i>Gm23426</i>	Predicted gene, 23426	1.518		
17277387	<i>Fos</i>	FBJ osteosarcoma oncogene	1.698		
17413945	<i>Nr4a3</i>	Nuclear receptor subfamily 4, group A, member 3	1.718		
17308165	<i>Gm21464</i>	Predicted gene, 21464	1.856		
17252875	<i>1700016P03Rik</i>	RIKEN cDNA 1700016P03 gene	2.046		
17464588	<i>Sgce</i>	Sarcoglycan, epsilon	-2.098		



Affymetrix Cluster ID	Gene Symbol	Gene Name	Fold Change 1.25 & p 0.05)
17546762	<i>Mid1</i>	Midline 1	-1.883
17532694	<i>Pisd-ps3</i>	Phosphatidylserine decarboxylase, pseudogene 3	-1.805
17464803	<i>Ica1</i>	Islet cell autoantigen 1	-1.566
17546212	<i>G530011O06Rik</i>	RIKEN cDNA G530011O06 gene	-1.564
17403844	<i>Gm24373</i>	Predicted gene, 24373	-1.551
17270615	<i>Gm22743</i>	Predicted gene, 22743	-1.478
17491505	<i>Gm24966</i>	Predicted gene, 24966	-1.457
17308963	<i>Gm23926</i>	Predicted gene, 23926	-1.398
17278781	<i>Gm23347</i>	Predicted gene, 23347	-1.349
17455971	<i>Mios</i>	Missing oocyte, meiosis regulator, homolog (Drosophila)	-1.334
17491503	<i>Gm25499</i>	Predicted gene, 25499	-1.333
17456204	<i>Capza2</i>	Capping protein (actin filament) muscle Z-line, alpha 2	-1.333
17539189	<i>Gm23806</i>	Predicted gene, 23806	-1.330
17364365	<i>Gm24930</i>	Predicted gene, 24930	-1.327
17491527	<i>Gm25870</i>	Predicted gene, 25870	-1.318
17356739	<i>Mir194-2</i>	microRNA 194-2	-1.297
17283683	<i>Gm24334</i>	Predicted gene, 24334	-1.288
17274184	<i>Socs2</i>	Suppressor of cytokine signaling 2	-1.288
17432577	<i>Gm436</i>	Predicted gene 436	-1.287
17399769	<i>S100a13</i>	S100 calcium binding protein A13	-1.282
17491642	<i>Snord107</i>	Small Nucleolar RNA, C/D Box 107	-1.281
17363082	<i>Olfir1437</i>	Olfactory receptor 1437	-1.278
17440618	<i>Mir701</i>	microRNA 701	-1.278
17478956	<i>Gm7551</i>	Heterogeneous nuclear ribonucleoprotein A3 pseudogene	-1.277
17460861	<i>Slc41a3</i>	Solute carrier family 41, member 3	-1.273
17278840	<i>Mir376a</i>	microRNA 376a	-1.271
17288946	<i>Mir9-2</i>	microRNA 9-2	-1.261
17491507	<i>Gm22496</i>	Predicted gene, 22496	-1.259
17491515	<i>Gm25121</i>	Predicted gene, 25121	-1.256
17284609	<i>Ighv1-62-3</i>	Immunoglobulin heavy variable 1-62-3	-1.254

**Table 7**

Validation of whole-genome gene expression with relative quantitative RT-PCR.

Gene symbol	Protein	Function	Fold Change ( <i>p</i> value) *	
			Microarray	QRT-PCR
<i>Nr4a3</i>	Nuclear receptor subfamily 4 group A member 3	Transcriptional activator	1.72 ± 0.17 (9.0E-4)	1.93 ± 0.15 (6.0E-5)
<i>Nr4a2</i>	Nuclear receptor subfamily 4 group A member 2	Transcriptional activator	1.49 ± 0.14 (4.0E-3)	1.29 ± 0.10 (0.010)
<i>Dusp1</i>	Dual specificity protein phosphatase 1	Regulation of cellular proliferation	1.45 ± 0.14 (3.0E-4)	2.17 ± 0.12 (8.0e-7)
<i>Cdkn1a</i>	Cyclin-dependent kinase inhibitor 1	Regulator of cell cycle progression	1.41 ± 0.16 (4.0E-3)	1.50 ± 0.12 (1.0E-3)
<i>Eomes</i>	Eomesodermin	Transcriptional activator	1.25 ± 0.06 (0.035)	1.63 ± 0.17 (1.0E-4)
<i>Spry4</i>	Sprouty homolog 4	Inhibitor of mitogen-activated protein kinase (MAPK) signaling pathway	1.24 ± 0.09 (0.027)	1.21 ± 0.08 (0.018)
<i>Slc41a3</i>	Solute carrier family 41 member 3	Cation transmembrane transporter activity	-1.27 ± 0.09 (0.045)	-1.42 ± 0.07 (3.0E-4)
<i>S100a13</i>	S100 calcium-binding protein A13	Calcium ion binding and lipid binding	-1.28 ± 0.04 (1.7E-3)	-1.52 ± 0.05 (1.0E-4)
<i>Socs2</i>	Suppressor of cytokine signaling 2	Suppressor of cytokine signaling (SOCS)	-1.29 ± 0.09 (0.016)	-1.29 ± 0.04 (4.0E-4)
<i>Capza2</i>	F-actin-capping protein subunit alpha-2	Regulator of the actin filaments	-1.33 ± 0.05 (6.0E-5)	-1.18 ± 0.16 (0.131)
<i>Mios</i>	WD repeat-containing protein mio	Inhibitor of the target of rapamycin complex I	-1.33 ± 0.04 (2.0E-4)	-1.22 ± 0.11 (0.049)
<i>Ica1</i>	Islet cell autoantigen 1	Membrane protein on Golgi complex and immature secretory granules	-1.56 ± 0.04 (6.0E-6)	-1.86 ± 0.04 (3.0E-6)
<i>Pomt2</i>	Protein-O-mannosyltransferase 2	Modification of the protein alpha-dystroglycan	-1.01 ± 0.02 (0.678)	1.01 ± 0.07 (0.886)
<i>Sgca</i>	Sarcoglycan, alpha	Component of the dystrophin-glycoprotein complex (DGC)	1.04 ± 0.02 (0.216)	1.05 ± 0.03 (0.459)
<i>Sgcb</i>	Sarcoglycan, beta	Component of DGC	-1.06 ± 0.02 (0.209)	-1.05 ± 0.05 (0.308)

\* Means ± SEM (*p* value)