Alternative Splicing Generates a Novel Truncated $Ca_{\beta}^{2+}$ Channel in Neonatal Rat Heart*

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Ping Liao$^{1,1}$, Dejie Yu$^s$, Zhenyu Hu$^s$, Mui Cheng Liang$^s$, Jue Jin Wang$^s$, Chye Yun Yu$^s$, Gandi Ng$^s$, Tan Fong Yong$^s$, Jia Lin Soon$^s$, Yeow Leng Chua$^3$, and Tuck Wah Soong$^{1,2}$

From the $^1$National Neuroscience Institute, 11 Jalan Tan Tock Seng, Singapore 308433, $^2$Duke-NUS Graduate Medical School Singapore, Singapore 169857, $^3$Department of Physiology, Yong Loo Lin School of Medicine, National University of Singapore, Singapore 117597, and $^3$National Heart Centre Singapore, 5 Hospital Drive, Singapore 169609

**Background:** L-type $Ca_{\beta}^{2+}$ channel undergoes extensive alternative splicing, generating functionally different channels. Alternatively spliced $Ca_{\beta}^{2+}$ channels have been found to be expressed in a tissue-specific manner or under pathological conditions. To provide a more comprehensive understanding of alternative splicing in $Ca_{\beta}^{2+}$ channel, we systematically investigated the splicing patterns in the neonatal and adult rat hearts. The neonatal heart expresses a novel 104-bp exon 33L at the IVS3-4 linker that is generated by the use of an alternative acceptor site. Inclusion of exon 33L causes frameshift and C-terminal truncation. Whole-cell electrophysiological recordings of $Ca_{\beta,33L}$ channels expressed in HEK 293 cells did not detect any current. However, when co-expressed with wild type $Ca_{\beta}$ channels, $Ca_{\beta,33L}$ channels reduced the current density and altered the electrophysiological properties of the wild type $Ca_{\beta}$ channels. Interestingly, the truncated 3.5-domain $Ca_{\beta,33L}$ channels also yielded a dominant-negative effect on $Ca_{\beta,3}$ channels, but not on $Ca_{\beta,3}$ channels, suggesting that $Ca_{\beta,3}$ subunits are required for $Ca_{\beta,3}$ regulation. A biochemical study provided evidence that $Ca_{\beta,33L}$ channels enhanced protein degradation of wild type channels via the ubiquitin-proteasome system. Although the physiological significance of the $Ca_{\beta,33L}$ channels in neonatal heart is still unknown, our report demonstrates the ability of this novel truncated channel to modulate the activity of the functional $Ca_{\beta}$ channels. Moreover, the human $Ca_{\beta}$ channel also contains exon 33L that is developmentally regulated in heart. Unexpectedly, human exon 33L has a one-nucleotide insertion that allowed in-frame translation of a full $Ca_{\beta}$ channel. An electrophysiological study showed that human $Ca_{\beta,33L}$ channel is a functional channel but conducts $Ca_{\beta}^{2+}$ ions at a much lower level.

Voltage-gated calcium channels govern the depolarization-induced $Ca_{\beta}^{2+}$ entry into cardiac muscles. The channel consists of a pore-forming $\alpha_{1}$ subunit associated with $\beta$, $\alpha_{\delta}$, and/or $\gamma$ auxiliary subunits to form an oligomeric complex. In mammalian myocardium, excitation-contraction coupling is characterized by a transient increase in cytosolic $Ca_{\beta}^{2+}$. The influx of $Ca_{\beta}^{2+}$ through voltage-gated calcium channels subsequently induces $Ca_{\beta}^{2+}$ release from sarcoplasmic reticulum in a process known as $Ca_{\beta}^{2+}$-induced $Ca_{\beta}^{2+}$ release. This process is prominent in the adult heart. The contraction of neonatal hearts relies more directly on the influx of $Ca_{\beta}^{2+}$ through L-type voltage-gated calcium channels, in particular the $Ca_{\beta,1}$ channels (1, 2).

L-type $Ca_{\beta,1}$ channels are high voltage-activated channels and are expressed widely in cardiovascular and nervous systems (3). The human $Ca_{\beta,1}$ gene, $CACNA1C$, contains 55 exons, of which >19 exons are subjected to alternative splicing (4). Novel splice variants of $Ca_{\beta,1}$ channels were uncovered (5, 6) as of the first systemic screening of cardiac $Ca_{\beta,1}$ channels (7). Alternative splicing of the $Ca_{\beta,1}$ channels is coupled to the generation of splice variants with altered electrophysiological and pharmacological properties (4, 8–11). They also show tissue-specific expressions (8, 9, 12) and can be altered under pathological conditions (13, 14).

In this report we screened the major alternatively spliced loci of $Ca_{\beta,1}$ channels and identified a novel exon 33L in neonatal rat hearts that has not been reported previously. Inclusion of exon 33L generates a truncated channel $Ca_{\beta,1,33L}$ that does not conduct $Ca_{\beta}^{2+}$. However, $Ca_{\beta,1,33L}$ produced a dominant-negative effect when co-expressed with functional $Ca_{\beta,1}$ channels.

**MATERIALS AND METHODS**

**RT-PCR and Single-cell RT-PCR**—Young adult male Wistar rats (2 months, 150–200 g) were sacrificed by CO$_2$ and subsequent cervical dislocation. Neonatal hearts were harvested from postnatal day 1 rats of either sex. This study was approved...
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and performed in accordance with the guidelines of the Institutional Animal Care and Use Committee of the University. Fetal human heart cDNA was purchased from Clontech. Adult human heart tissues were obtained from donors for heart transplantation in the National Heart Center of Singapore with the approval of the institutional review board. The work was approved by the Institutional Review Board committees of the National Heart Center of Singapore, Singapore General Hospital. Total RNA was isolated from the left ventricles. cDNA was generated from total RNA by incubating with reverse transcriptase and 18-mer oligo(dT). The PCR protocol includes: a denaturation step at 95 °C for 5 min; 35 cycles of 95 °C for 30 s, 52 °C or 55 °C for 45 s, and 72 °C for 1 min; a final extension step at 72 °C for 10 min. The primers used for the rat Ca$_{v}$.1.2 exon 33L were: 5′-GCCCTCTCCAGTGTTGGAG-3′ (forward) and 5′-TCCC- CAATCACTGCAATGATAA-3′ (reverse). The primers used for human Ca$_{v}$.1.2 exon 33L are: 5′-CCAAGACCTAGAATACTTC-ACG-3′ (forward) and 5′-CTACCCAGGGTTGTCC-ACC-3′ (reverse). The primers used for amplifying human Ca$_{v}$.1.2 exons 18 to 24 as a control were: 5′-ATGAGGATATAAACACCCCT- (forward) and 5′-CTCAGCAAGATTCTTCACAGATTG-3′ (reverse). For single-cell RT-PCR, after electrophysiological characterization by the patch clamp method, the cells were collected into a vial containing reverse transcription reaction mix for first strand cDNA synthesis using the reverse primer. Single-cell PCR protocol was similar to RT-PCR, except for a 40-cycle amplification.

Colony Screening and Restriction Enzyme Digestion—This method was described previously to identify novel splice variations (7, 14). In brief, the PCR products were cloned into pGEM-T Easy vector (Promega). After being transformed into DH10B Escherichia coli cells, each transformant was selected and grown in a single well in a 96-well plate. Colony PCR was performed with the same set of primers and conditions to identify the component of exons in each colony. Usually 96 colonies were selected for each sample. The genotype of each PCR product amplified from a colony was predicted by size and randomly confirmed by DNA sequencing. To differentiate mutually exclusive exons that are of same size, restriction enzyme digestions were used to identify the expression of individual exons. The digestion was carried out in a 30-μl reaction volume at 37 °C for 5 h. To control for complete digestion, the same amount of PCR product of a positive control DNA was subjected to similar digestion conditions.

Sequencing, Analysis, and Cloning of 33L Channels—Rat exon 33L was verified by DNA sequencing. The sequences were compared with rat genome sequence from the Human Genome Sequencing Center, Baylor College of Medicine. To characterize the functions of exon 33L containing channels, a fragment containing exon 33L was cloned into a wild type rat Ca$_{v}$.1.2 channel with AvrII and NheI sites. This wild type Ca$_{v}$.1.2 channel contains the exons predominantly expressed in rat cardiac tissues.

Generation of Polyclonal Antibody against Exon 33L Polypeptide—Rat exon 33L was cloned in-frame into pGEX-4T-1. Amino acid sequences from adjacent exons were not included. GST-fused protein was purified with glutathione-agarose (Sigma). Purified protein was used to immunize female New Zealand White rabbit once a month. Complete Freund’s adjuvant was first mixed with GST-33L for immunization, and incomplete Freund’s adjuvant was used in subsequent injections once a month. Serum collected after immunization was preabsorbed with GST protein first to remove GST antibodies, and 33L polyclonal antibody was affinity-purified from immobilized 33L protein with an IgG elution buffer (Pierce). The antibody concentration is 1 μg/μl. Serum from rabbit before immunization was used as preimmune control.

Western Blot—For protein isolation, heart tissues were homogenized in HEPES lysis buffer: 20 mmol/liter HEPES, 137 mmol/liter NaCl, 1% Triton X-100, 10% glycerol, 1.5 mmol/liter MgCl$_2$, and 1 mmol/liter EGTA supplemented with protease inhibitors (1:50 dilution, Roche Diagnostics). Homogenized samples were then centrifuged at 14,000 rpm for 15 min, and supernatants containing protein samples were collected. Protein concentration was determined using Bradford assay. 200 μg of lysates were separated by 8% SDS-PAGE and transferred overnight at 30 V onto PVDF membranes at 4 °C. Subsequently, the membrane was blocked with 1% BSA in 1× PBS + 0.1% Tween 20 for 1 h and probed with primary antibodies overnight at 4 °C. The primary antibodies used in the study include rabbit anti-exon 33L polyclonal (1:5000), rabbit anti-Ca$_{v}$.1.2 or Ca$_{v}$.1.3 (1:1000, Alomone), mouse anti-ubiquitin (1:1000, Invitrogen), and mouse anti-β-actin (1:5000, Sigma). The next day, the membrane was washed 3 times with 0.1% Tween 20 in 1× PBS. The membrane was then probed with HRP-conjugated secondary antibody for 1 h. After washing, the band was detected using Amersham Biosciences ECL Western blotting Analysis System (RPN2109, GE Healthcare).

Immunofluorescent Staining—Animals were sacrificed and perfused with saline and subsequently 4% paraformaldehyde. The heats were then collected and post-fixed with 4% paraformaldehyde for 2 h. Dehydration was subsequently carried out by immersing the hearts into 15% sucrose and subsequently 30% sucrose. Next, the rat heart was sectioned at 20 μm of thickness. After washing with 0.2% Triton X-100 phosphate-buffered saline (PBST), 100 μl of blocking serum (10% goat serum and 1% bovine serum albumin in 0.2% PBST) was added onto the sections for 1 h. The sections were then incubated with primary antibodies overnight at 4 °C. Monoclonal antibodies anticonnexin 43 (MAB 3068) was purchased from Chemicon International. On the following day, tissue sections were washed 3 times with TNT wash buffer (0.1 M Tris-HCl buffer, pH 7.5, containing 0.15 M NaCl and 0.05% Tween 20). The slides were incubated with FITC-conjugated or Texas red-labeled secondary antibodies for 1 h at room temperature. After washing three times of wash buffer, the slides were mounted with FluorSaveTM reagent (Merck). The results were visualized using laser scanning confocal microscope system (Fluoview BX61, Olympus). The negative control was done in an identical procedure except for primary antibody incubation, and no positive signal was identified.

Whole-cell Patch Clamp Electrophysiology—HEK293 cells were transiently transfected with rat α$_1$ (1.25 μg), β$_{2a}$ (1.25 μg), and α$_{26}$ (1.25 μg) subunits with the calcium phosphate transfection method. The GFP-tagged β$_{2a}$ and α$_{26}$ clones were provided by Dr. Terrance Snutch (University of British Columbia).
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The rat Cav3.2 channel and Cav1.3 channel were described earlier (15, 16). I_{Ca} was recorded with a whole-cell patch clamp technique at room temperature (~25 °C) 48–72 h post-transfection. Patch electrodes were pulled using a Flaming/Brown micropipette puller (Sutter Instrument) and polished with a microforge. The external solution contained 140 mmol/liter tetraethylammonium methanesulfonate, 10 mmol/liter HEPES, and 1.8 mmol/liter CaCl_2 (the pH was adjusted to 7.4 with CsOH and osmolarity to 300–310 mossm with glucose). For I_{Ba} recording, 1.8 mM Ca^{2+} was replaced by 5 mM Ba^{2+}. The internal solution (pipette solution) contained 138 mmol/liter Cs-MeSO_3, 5 mmol/liter CsCl_2, 0.5 mmol/liter ethylene glycol tetraacetic acid, 10 mmol/liter HEPES, 1 mmol/liter MgCl_2, and 2 mg/ml MgATP, pH 7.3 (adjusted with CsOH). The osmolarity was adjusted to between 290 and 300 mossm with glucose. Under voltage clamp and using an Axopatch 200B amplifier (Axon Instruments), whole-cell currents were filtered at 1–5 kHz and sampled at 5–50 kHz. The series resistance was normally <5 megaohms after compensation. The capacity transient was compensated using an online P/4 protocol. The steady-state inactivation curves were obtained from experiments by stepping from a holding potential of −90 mV to a 30-ms normalizing pulse to 10 mV followed by a family of 15-s-long prepulses from −80 to +20 mV. A 104-ms test pulse to +10 mV was recorded finally. Each test pulse was normalized to the maximal current amplitude of the normalizing pulse. The steady-state inactivation (SSI) data were fitted with a single Boltzmann equation, \( I_{\text{relative}} = I_{\text{min}} + (I_{\text{max}} - I_{\text{min}})/(1 + \exp((V/2 - V)/k)) \), where \( I_{\text{relative}} \) is the normalized tail current, \( V/2 \) is the potential for half-inactivation, and \( k \) is the slope value. G-V curves were obtained from a tail activation protocol. The cells were activated by a 20-ms test pulse of variable voltage family from −60 to 100 mV, and tail currents were measured after repolarization to −50 mV for 10 ms. The tail currents were normalized to the peak currents before fitting with a dual Boltzmann equation, \( G/G_{\text{max}} = F_{\text{low}}/[1 + \exp((V_{1/2,\text{low}} - V)/k_{\text{low}})] + (1 - F_{\text{low}})/[1 + \exp((V_{1/2,\text{high}} - V)/k_{\text{high}})] \), where \( G \) is the tail current, and \( G_{\text{max}} \) is the peak tail current, \( F_{\text{low}} \) is the fraction of low threshold component, and \( V_{1/2,\text{low}}, V_{1/2,\text{high}}, k_{\text{low}}, \) and \( k_{\text{high}} \) are the half-activation potentials and slope factors for the low and high threshold components, and \( V_{1/2,\text{act}} \) was calculated when \( G = 0.5G_{\text{max}} \).

RESULTS

Exon 31 and 32 Are Developmentally Regulated—Ca_{1,2} channel undergoes extensive alternative splicing, which affects its tissue distribution, pharmacology, and electrophysiological properties (4, 10). Although up to 20 exons are alternatively spliced (17), only exons 31 and 32 have been reported to be developmentally regulated in the heart (18). Mutually exclusive exons 31 and 32 encode the IVS3 transmembrane segment and part of the IVS2-S3 intracellular linker, whereas another alternatively spliced exon 33 encodes the IVS3-S4 extracellular linker. There exist extensive alternative splicings within this region (7) involved in many cardiovascular diseases (6, 13, 14, 19).

FIGURE 1. A, RT-PCR across exons 30–35 could produce a fragment of 382 bp when exon 33 (33 bp) is present. Exons 31 and 32 are mutually spliced exons. Digestion at exon 32 with NsiI will generate two fragments of 68 and 314 bp, b, partial digestion with NsiI was observed in both NH and AH. Exon 32 is expressed higher in AH as more PCR product was digested. Equal amounts of DNA from both AH and NH were digested under same conditions in parallel.

RT-PCR across exons 30–35 will generate a 382-bp amplicon containing exon 33 and one of the mutually exclusive exons 31 or 32. Enzyme digestion on exon 32 by NsiI produces two smaller fragments with the sizes of 68 and 314 bp, respectively (Fig. 1A). In the presence of exon 31, only one band exists (382 bp), as no digestion occurs. Deletion of the cassette exon 33 (33 nt in length) has rarely been found in rat hearts (8). When we digested RT-PCR products from the neonatal and adult rat hearts, partial digestions were observed in both samples, indicating that neonatal and adult hearts contain a mixture of exons 31 and 32. However, the extent of digestion by NsiI was different, as more PCR products from the adult hearts were digested than from the neonatal hearts. Therefore, exon 32 expression is higher in adult rat hearts than in neonatal rat hearts (Fig. 1B). Such developmental changes of exon 31 and 32 have been reported by Diebold et al. (18), and our experiment supports the previous finding.

Identification of Exon 33L from Neonatal Rat Heart—Besides mutually exclusive exons 31, 32, and cassette exon 33, there exists an additional 66-nt extension at the 5’ end of exon 34 with different acceptor sites, producing exon 34a, an isoform of exon 34 (6). To characterize the detailed expression of exon 33 and 34a, we used the colony screening method that is able to detect exons of lower expression (8). The inclusion or exclusion of exons 33 and/or 34a could be easily identified by differences in size of the PCR products, whereas mutually exclusive exons 31 and 32 cannot, as they are of the same length. Colony PCR screening showed that the size of the majority of colonies was ~382 bp, indicating that the splice variants with exon 33 deletion or exon 34a inclusion are not predominant in neonatal and adult rat hearts (Fig. 2, A and B). DNA sequencing confirmed that the colonies with smaller size contained only exon 31 or 32 but with the exon 33 deletion. However, some colonies of a

\( ^{\text{a}} \) The abbreviations used are: SSI, steady-state inactivation; nt, nucleotide(s); NH, neonatal heart; AH, adult heart; pF, picofarad.
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To determine whether the aberrant splice variants are translated into proteins, a polyclonal antibody was generated against the 22-amino acid polypeptides encoded by exon 33L (Fig. 3A). The 22-amino acid polypeptides were expressed as a fusion protein with GST (Fig. 4A) and were injected into rabbits to produce exon 33L specific antibody α-33L. No adjacent amino acids were included for antibody production. Using α-33L, we detected by Western blot a band of 160 kDa in neonatal and adult hearts, having the same molecular mass of the truncated 33L containing Cav1.2 channel expressed in HEK 293 cells (Fig. 4B).

Therefore, neonatal rat hearts expressed higher levels of Cav1.233L channel than adult hearts. To prove the specificity of α-33L antibody, HEK cells were transfected with Cav1.233L channel or the reference channel Cav1.2WT, which does not contain exon 33L. α-33L antibody stained Cav1.233L-transfected cells but not Cav1.2WT-transfected cells (Fig. 4C). Next, we used an immunohistological method to examine the cellular expression of the Cav1.233L channel in rat hearts during development. Connexin 43 was labeled for the gap junctions between adjacent cardiac muscle cells. Again, α-33L staining showed a stronger fluorescent signal in neonatal heart than in adult heart, supporting a higher expression of Cav1.233L channel in neonatal rat heart (Fig. 4D). It should be noted that Cav1.233L did not colocalize with Connexin 43 in both neonatal and adult hearts, suggesting that the majority of Cav1.233L channels may not traffic to the plasma membrane.

Presence of Cav1.233L Channels Reduces Currents of Functional Cav1.2 Channels—The functional significance of the inclusion of exon 33L in Cav1.2 channels has not been reported. We sought to assess exon 33L functions by whole-cell patch clamp recording. Exon 33L was cloned into a reference rat cardiac-form Cav1.2 channel that contained exon 1a and 8a, with the absence of exon 9* (8). To control for transfection efficiency and the conditions of the HEK 293 cells, we performed the experiments in parallel by transfecting the HEK 293 cells on the same day and did the electrophysiological experiments alter-
nately using the same solutions on the same day. The presence of successful transfection of Cav1.233L channels was evidenced by the single cell RT-PCR. All the cells patched with green fluorescence (H92522a is tagged with GFP) contained Cav1.233L channels. The cells were maintained at a H1100290 mV holding potential before a test pulse of 0 mV for 900 ms was applied. No currents were recorded (Fig. 5A,B) for Cav1.233L channel, indicating that the presence of exon 33L generates a novel Cav1.2 channel that is impermeable to Ca2+ ions. To investigate whether the presence of the 33L channels affects the reference cardiac channels, we co-transfected the 33L channels and the reference channels into HEK cells. The exon 33L expression in neonatal heart and adult heart is 9.7% and 4.3%, respectively (Fig. 2C). Therefore, we transfected Cav1.233L channels into HEK 293 cells at 5 and 10% to the reference wild type Cav1.2; lane 4, AH; lane 5, NH. C, immunofluorescent staining of Cav1.233L- or Cav1.2wt-transfected HEK cells with polyclonal antibody α-33L (red). GFP-tagged β2a was co-transfected together with the α, subunits. D, representative immunofluorescent staining showed exon 33L expression in NH and AH. Connexin 43 (Cx43) was co-stained, and nuclei were labeled with DAPI. Scale bar: 50 μm.

It is possible that the effect of Cav1.233L channels on the reference channels is masked by the variability of the data recorded. We decided to increase the 33L channel ratio by transfecting equal amounts of both Cav1.233L channels and the reference channels into HEK 293 cells for electrophysiological recordings. The control experiments were carried out by transfecting the same amount of empty pcDNA3 vector together with the reference channels. Single cell RT-PCR was performed to confirm that both the 33L channels and the reference channels were equally expressed (Fig. 6A). The data collected from cells with unequal expression were discarded. Similar to the dominant-negative effect observed by Ebihara et al. (20) in the three-domain Cav1.2 channels, the presence of Cav1.233L channels reduced the tail current density of the reference Cav1.2 channels (Fig. 6B) in 1.8 mM Ca2+ bath solution, mimicking a physiological condition. The tail current density at +100 mV was 41.6 ± 13.2 pA/pF (n = 7) when control pcDNA3 vectors were co-transfected with the reference channels, whereas the tail current density was 11.3 ± 2.3 pA/pF (n = 9) when
Ca\textsubscript{1.2}\textsubscript{33L} channels were co-transfected, representing a 73% decrease in the presence of Ca\textsubscript{1.2}\textsubscript{33L} channels (p = 0.025, Student’s t test). Measuring the ON-gating charge (Q\textsubscript{ON}) at the reversal potential (typically +50 mV, Fig. 6C) provides a convenient measure of the number of channels at the membrane (21). Therefore, we measured the Q\textsubscript{ON} of the Ca\textsubscript{1.2}\textsubscript{33L} and pcDNA3-transfected cells. Interestingly, 33L transfection greatly reduced the Q\textsubscript{ON} as compared with pcDNA3-transfected cells: 33L Q\textsubscript{ON}, 0.8374 ± 0.2296 femtocoulombs/pF, n = 7; pcDNA3 Q\textsubscript{ON}, 2.499 ± 0.4639 femtocoulombs/pF, n = 6; p = 0.0063, Student’s t test. The reduction of 66.5% in Q\textsubscript{ON} was similar to the 73% reduction in current density.

Ca\textsubscript{1.2}\textsubscript{33L} Channels Exert a Dominant Negative Effect on Ca\textsubscript{1.3} Channels but Not on Ca\textsubscript{3.2} Channels—To examine whether Ca\textsubscript{1.2}\textsubscript{33L} channels down-regulate other voltage-gated calcium channels, we cotransfected Ca\textsubscript{1.2}\textsubscript{33L} channels with the paralogous L-type Ca\textsubscript{1.3} channels or with the T-type Ca\textsubscript{3.2} channels at a ratio of 1:1. Again, pcDNA3 vector was used as a control. One difference between Ca\textsubscript{1.3} channels and Ca\textsubscript{3.2} channels is that Ca\textsubscript{1.3} channels require the β subunit for functional expression, whereas Ca\textsubscript{3.2} channels do not need the β subunit. Current densities from I-V curves were measured in 1.8 mM Ca\textsuperscript{2+} solution. Interestingly, the dominant negative effect of Ca\textsubscript{1.2}\textsubscript{33L} channels is prominent in Ca\textsubscript{1.3}-expressed cells but not in Ca\textsubscript{3.2}-expressed cells (Fig. 7). Compared to the pcDNA3 control, the presence of Ca\textsubscript{1.2}\textsubscript{33L} channels reduced Ca\textsubscript{1.3} current densities at −10 mV from −16.16 ± 2.205 pA/pF (n = 8) to −7.952 ± 1.250 pA/pF (n = 15); p = 0.0021, Student’s t test (Fig. 7A). However, in Ca\textsubscript{3.2}-expressed cells, the current densities were similar in pcDNA3 and Ca\textsubscript{1.2}\textsubscript{33L}-transfected cells (Fig. 7B). At −20 mV, the current densities of Ca\textsubscript{1.2}\textsubscript{33L}-expressed cells were −16.93 ± 4.158
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![Diagram](https://example.com/diagram.png)

**FIGURE 8.** Effect of Ca_{v}1.2_{33L} channels on the gating properties of the reference Ca_{v}1.2 channels when transfected at 1:1 ratio. A, sample traces from the recordings for the tail currents and steady-state inactivation. B, plots of steady-state inactivation ($f_{\infty}$) and activation ($d_{\infty}$) curves of the reference cardiac Ca_{v}1.2 channels co-expressed with Ca_{v}1.2_{33L} (●) and pcDNA3 vector (○). Ca_{v}1.2_{33L} channels slightly shifts activation and SSI curves of the reference channels toward negative potential. All recordings were performed in 1.5 mM Ca^{2+} bath solution. The protocols and fittings were described under “Materials and Methods.”

pA/pF, $n = 9$, whereas those in pcDNA3-expressed cells were $-17.13 \pm 3.257$ pA/pF, $n = 7$ ($p = 0.9698$, Student’s $t$ test).

**Presence of Ca_{v}1.2_{33L} Channels Changes Electrophysiological Properties of Functional Ca_{v}1.2 Channels—**To investigate whether the electrophysiological properties of the reference cardiac channel is altered with the presence of Ca_{v}1.2_{33L} channels, we used a recording protocol for the SSI and a tail protocol for the activation properties, which were described previously (8, 9). Interestingly, the presence of Ca_{v}1.2_{33L} channels negatively shifted both the SSI and the activation properties of the reference channels (Fig. 8). The $V_{1/2}$ of the SSI was hyperpolarized-shifted for $-3.54$ mV ($p = 0.0174$, Student’s $t$ test), $V_{1/2SSIPcDNA}$ ($-22.96 \pm 0.86$ mV ($n = 6$)) versus $V_{1/2SSIC33L}$ ($-26.5 \pm 0.9$ mV ($n = 6$)), whereas the $V_{1/2}$ of the activation curve was hyperpolarize-shifted for $-5.73$ mV ($p = 0.0036$, Student’s $t$ test), $V_{1/2ActPcDNA}$ ($18.43 \pm 1.05$ mV ($n = 9$)) versus $V_{1/2ActC33L}$ ($12.7 \pm 1.28$ mV ($n = 7$)). Therefore, the calcium window current of the reference cardiac channel was shifted to a more negative potential in the presence of Ca_{v}1.2_{33L} channels.

Ca_{v}1.2_{33L} Channels Down-regulate Functional Ca_{v}1.2 Channels by Enhancing Protein Degradation via the Ubiquitin-Proteasome System—To investigate the mechanism for the dominant negative effect of Ca_{v}1.2_{33L}, the channels were first co-transfected with the reference channels at 1:1 ratio. A Western blot showed that the total protein level (membrane and cytosol) of the reference Ca_{v}1.2 channels was reduced by the presence of Ca_{v}1.2_{33L} (Fig. 9A). Treatment with 3 μM MG132, a proteasome inhibitor, successfully increased the expression of the reference Ca_{v}1.2 channels, suggesting that Ca_{v}1.2_{33L} might enhance proteasomal degradation. A similar down-regulation effect of Ca_{v}1.2_{33L} was observed on Ca_{v}1.3 channels (Fig. 9B) but with a smaller reduction. Next, we examined protein ubiquitination using immunoprecipitation and subsequent Western blot. Interestingly, transfection with Ca_{v}1.2_{33L} greatly enhanced wild type Ca_{v}1.2 channel ubiquitination. The ubiquitinated Ca_{v}1.2 channels were further increased with the treatment of 3 μM MG132 (Fig. 9C). Again, Ca_{v}1.3 channel ubiquitination was also increased with Ca_{v}1.2_{33L} co-expression (Fig. 9D).

**Human Exon 33L-containing Channel Conducts Ca^{2+} Ions—** When we compared the exon 33L sequences of mouse, rat, and human (Fig. 10A), we found that the acceptor for exon 33L is preserved across the three species, indicating that exon 33L may exist in all these species. The mouse and the rat shared exactly the same sequences of exon 33L, whereas the human owns a different sequence. Interestingly, the human genomic sequence had an additional base within exon 33L compared to mouse or rat. Therefore, no frameshift was generated (Fig. 10B), and the human Ca_{v}1.2_{33L} channel may conduct Ca^{2+} ions. To prove this, we first performed RT-PCR on human heart samples using primers specific for exon 33L. The sequences of the primers are described under “Materials and Methods.” A band with a size of 502 bp was found in the left and right ven-
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FIGURE 10. Human Cav1.233L channels conduct Ca2+ ions and are developmentally regulated in heart. A, alignment of the exon 33L DNA sequences from Rattus norvegicus, Mus musculus, and Homo sapiens. The canonical -ag- acceptors (indicated in the gray box) is preserved in all three species. The mouse and the rat share the same exon 33L sequence. Human exon 33L is predicted to contain an additional nucleotide. All the differences in DNA sequences are highlighted by boxes. B, alignment of the amino acid sequences of exon 33L predicted from R. norvegicus, M. musculus, and H. sapiens. The differences in amino acid sequences are labeled by boxes. No stop codon is generated by the inclusion of human exon 33L. C, using human exon 33L-specific primers, a band of 502 bp was found in human adult left ventricle (LV), adult right ventricle (RV), and fetal left ventricle (FH). M, marker. DNA sequencing confirmed the sequence to be human exon 33L. PCR using primers across exon 18–exon 24 was illustrated as the loading control. As a water control (–), cDNA was replaced by water to exclude possible contaminations. D, human exon 33L was cloned into a Cav1.2 reference channel. I-V curves were obtained in a bath solution containing 1.8 mM Ca2+. In HEK cells, human Cav1.233L channels indeed conduct Ca2+ ions with a smaller current density. E, conductance was calculated as $G = I/(V_m - E_{Ca})$ and normalized by the maximum conductance value and fit with Boltzmann equation, where $E_{Ca}$ is reverse potential.

Truncated Na+ and Ca2+ channels have been found in brain and muscles (22, 23). In rabbit heart aberrant splicing in the II-III loop of Cav1.2 channels produces two truncated channels that contain only domains I and II (23). The truncation is caused by the deletion of exons 17 and 18 or the deletion of exon 19, which results in a frameshift leading to premature termination of translation. In contrary, alternative splicing of exon 33L...
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of the Ca\(_{\text{v}1.2}\) channels we report here yielded 2 significant findings. 1) The functional modification is species-specific in that rat exon 33L results in a truncated channel that likely competes for β subunit binding to wild type channels. However, the human exon 33L has a single nucleotide insertion resulting in a longer functional full-length channel that, however, produced reduced current density. 2) The truncated rat Ca\(_{\text{v}1.2\beta 33L}\) variant not only competes for β subunit binding to expose the wild type Ca\(_{\text{v}1.2}\) channel to presumably ERAD degradation (24), but it also similarly down-regulated the expression of another β subunit requiring L-type Ca\(_{\text{v}1.3}\) channel but not the T-type Ca\(_{\text{v}3.2}\) channel.

The presence of rat exon 33L shifts the coding frame and leads to a deletion of the last three transmembrane segments of the domain IV and the downstream C terminus. Therefore, the predicted topography of this rat Ca\(_{\text{v}1.2\beta 33L}\) channels is larger than the two domain channels reported previously (23).

Rat Ca\(_{\text{v}1.2\beta 33L}\) channels do not conduct Ca\(^{2+}\) ions, indicating that the last three transmembrane segments within domain IV are critical in forming a complete functional channel. The pore of the channel is likely to be disrupted in particular with the deletion of IVS4 containing positively charged amino acids. Another result from this study is that the presence of Ca\(_{\text{v}1.2\beta 33L}\) channels could down-regulate the current flux through the reference wild type channels. The inhibitory effect is not prominent when Ca\(_{\text{v}1.2\beta 23L}\) channels are expressed at 5% or 10%, possibly due to data variability when recording small changes in current density. However, when Ca\(_{\text{v}1.2\beta 33L}\) was expressed in equal amount to the reference channels, the dominant negative effect is clear. We found a 73% current density reduction, largely in correlation with a 66.5% decrease in gating current, indicating that the current density reduction is mainly due to a decrease of the functional Ca\(_{\text{v}1.2}\) channels on the cell membrane. The slight difference between the changes in current density and gating current (~6.5%) suggests that a small number of channels on the surface membrane may not conduct Ca\(^{2+}\) ions. This small population of channels is likely the truncated Ca\(_{\text{v}1.2\beta 23L}\) channels. Immunostaining of Ca\(_{\text{v}1.2\beta 33L}\) channels in rat neonatal and adult hearts suggests that the majority of Ca\(_{\text{v}1.2\beta 33L}\) channels were not expressed on the plasma membrane. Therefore, only very few Ca\(_{\text{v}1.2\beta 33L}\) channels may traffic to the membrane, contributing to a small portion of the gating currents.

The mechanism for the inhibitory effect of Ca\(_{\text{v}1.2\beta 33L}\) channels could be explained in that Ca\(_{\text{v}1.2\beta 33L}\) channels compete with the reference channels, presumably for β subunit for expression on the plasma membrane. It is well known that the β subunit is required to target the functional L-type Ca\(^{2+}\) channels to the cell surface (25), and the binding requires a conserved motif in the domain I-II cytoplasmic loop of the Ca\(_{\text{v}1.2}\) channels (26). As the I-II loop is well preserved in Ca\(_{\text{v}1.2\beta 33L}\) channels, the expression of this truncated channel may scavenge β subunits and, therefore, down-regulate the surface expression of the reference channels. This mechanism is supported by a drastic reduction of gating currents caused by Ca\(_{\text{v}1.2\beta 33L}\) channels. Another possible mechanism is that the truncated channels may prevent the correct folding of wild type channels and inhibit channel synthesis or stability. This hypothesis was shown by Dolphin and co-workers (27) in two domain truncated Ca\(_{\text{v}2.2}\) channels, in which the dominant negative effect is independent on β subunit. Therefore, such modulation is generally channel-specific, seldom affecting other channels with a different protein structure. Our results from experiments performed on Ca\(_{\text{v}1.3}\) and Ca\(_{\text{v}3.2}\) channels suggest that the second mechanism may not explain the effect Ca\(_{\text{v}1.2\beta 33L}\) has on wild type Ca\(_{\text{v}1.2}\) and Ca\(_{\text{v}1.3}\) channels. The dominant negative effect of Ca\(_{\text{v}1.2\beta 33L}\) channels was exerted as well on Ca\(_{\text{v}1.3}\) channels, which require β subunits, but not on Ca\(_{\text{v}3.2}\) channels, which do not need β subunits, suggesting that β subunits are critical for Ca\(_{\text{v}1.2\beta 33L}\) suppression. The binding of β subunits has been demonstrated by the Zamponi and co-workers (24) to increase membrane expression of Ca\(_{\text{v}1.2}\) channels by preventing endoplasmic reticulum-associated protein degradation. Without β subunits, Ca\(_{\text{v}1.2}\) channels undergo robust ubiquitination. Therefore, if Ca\(_{\text{v}1.2\beta 33L}\) channels compete with wild type Ca\(_{\text{v}1.2}\) channels for β subunits, wild type channels without β subunit association will be degraded. Our biochemical results support this hypothesis. In the presence of Ca\(_{\text{v}1.2\beta 33L}\) channels, wild type Ca\(_{\text{v}1.2}\) channel ubiquitination was enhanced for degradation, and thus total protein level was reduced.

However, we cannot exclude the mechanism proposed by Dolphin completely. If the number of intracellular β subunits is fixed, the expression of Ca\(_{\text{v}1.2\beta 33L}\) channels should correlate with the inhibitory effect on the reference channels, which is an ~50% decrease when the Ca\(_{\text{v}1.2\beta 33L}\) channel and the reference channel are expressed at a 1:1 ratio. However, the current density is reduced by 73%, more than the predicted 50%, indicating the presence of the second mechanism. Furthermore, the Ca\(_{\text{v}1.2\beta 33L}\) channel suppressed the current density in Ca\(_{\text{v}1.3}\) channels from −16.16 pA/µF to −7.952 pA/µF, representing an ~50% reduction, strongly correlating with the molar ratio of Ca\(_{\text{v}1.2\beta 33L}\) channels to Ca\(_{\text{v}1.3}\) channels, which is 1:1. As Ca\(_{\text{v}1.3}\) channels have a different structure from Ca\(_{\text{v}1.2}\) channels, the second mechanism is unlikely to apply to Ca\(_{\text{v}1.3}\) channels. Therefore, we would hypothesize that the inhibitory effect of Ca\(_{\text{v}1.2\beta 33L}\) channels on Ca\(_{\text{v}1.3}\) channels is solely via competing for β subunits.

Ca\(_{\text{v}1.2\beta 33L}\) channels are also different from another three-domain Ca\(_{\text{v}1.2}\) channel that was reported earlier (20). In that truncated channel, the N terminus and the first domain as well as the entire I-II loop are deleted. Thus, there will be no competition for β subunits from this three-domain channel, which could explain why the inhibitory effect was not found on other channels. Interestingly, the three-domain Ca\(_{\text{v}1.2}\) channel exhibits a more prominent dominant-negative effect on wild type channels compared with the Ca\(_{\text{v}1.2\beta 33L}\) channel. As the C terminus is well preserved in the three-domain Ca\(_{\text{v}1.2}\) channel and the rest of the channel is the same as in the Ca\(_{\text{v}1.2\beta 33L}\) channel, the potent inhibitory effect is likely due to the intact C terminus. Zamponi and co-workers (24) showed that the endoplasmic reticulum retention motif of Ca\(_{\text{v}1.2}\) channel locates at the C terminus and likely interferes with the β subunits and the I-II linker sterically or allosterically in a properly folded channel. The three-domain Ca\(_{\text{v}1.2}\) channel with a complete C ter-
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Our results further showed that the presence of Ca\textsubscript{v}1.2\textsubscript{33L} channels could alter the electrophysiological properties of the reference cardiac Ca\textsubscript{v}1.2 channels. The negative shifting of SSI and activation curves produced a calcium window current that peaks at more hyperpolarized membrane potential. The exact mechanism of such alterations is not fully understood. In a previous study on Ca\textsubscript{v}1.1 channel (\alpha1S), the two-domain (I and II)-containing hemichannels were expressed on the cell membrane, and the gating characteristics were modified significantly, presumably by interactions with domains III and IV (28). Thus, it is possible that the interaction between the intact I and II domains of 33L channels with the reference Ca\textsubscript{v}1.2 channels may change the electrophysiological properties of those reference channels. The deletion of the C terminus in 33L channels may also cause the electrophysiological property changes in reference channels. The C terminus of Ca\textsubscript{v}1.2 channels is known to regulate Ca\textsuperscript{2+}-dependent and voltage-dependent channel inactivation (29). Therefore, the interaction with the truncated 33L channels may alter the regulatory functions of C terminus in the reference channel. Additional experiments are needed to elucidate the regulatory role of the 33L channels on functional Ca\textsubscript{v}1.2 channels.

The truncated Ca\textsubscript{v}1.2\textsubscript{33L} channel may not have a major physiological impact on neonatal heart function, as the expression level is only at 9.7% and even lower in adult hearts. However, during the search for more altered exonal expression in neonatal hearts, at least one more site was identified in the IIIS2 region involving exons 21 and 22 in forming an aberrant channel (data not shown). This aberrant channel again loses the ability for Ca\textsuperscript{2+} influx but exhibits a dominant-negative effect on functional channels. Therefore, we believe that aberrant channels in neonatal hearts may have some functions during cardiac development. More experiments with gene manipulation can help to identify the physiology of these aberrant channels.

When we searched human genomic sequence of Ca\textsubscript{v}1.2 channel, we identified a similar acceptor site for human exon 33L. Unexpectedly, human exon 33L sequence contained an additional nucleotide and was unlikely to generate a stop codon that is found in rat exon 33L. We successfully cloned human exon 33L into a reference Ca\textsubscript{v}1.2 channel, and electrophysiology studies on these channels eventually showed that they are functional in conducting Ca\textsuperscript{2+} ions, albeit with a smaller current density. At 0 mV, the current density of human Ca\textsubscript{v}1.2\textsubscript{33L} channels is only ~40% that in wild type channels. The mechanism for the smaller current remains unknown. Exon 33L codes for the extracellular loop between IVS3 and IVS4, which has been shown to affect channel activation property (7). It was proposed that a longer IVS3-S4 loop shifts the activation potential to a more positive potential. This trend was previously shown for alternatively spliced exon 33, which is of 33 bp in length. Deletion of exon 33 in smooth muscles shifts the activation and inactivation properties of Ca\textsubscript{v}1.2 channels to a more negative potential. Such structural variations are believed to contribute to the electrophysiological differences between cardiac and smooth muscle Ca\textsubscript{v}1.2 channels (8). With the inclusion of human exon 33L, which encodes an additional 24 amino acids, the loop between IVS3 and IVS4 is much longer and may change the channel properties as well. Indeed, when we compared the activation properties of the human Ca\textsubscript{v}1.2\textsubscript{33L} channel with the wild type channel, the \( V_{1/2} \) of the human Ca\textsubscript{v}1.2\textsubscript{33L} channel is significantly more positive than that of wild type channel.

Interestingly, human exon 33L is also developmentally regulated in heart, with a similar trend as seen in rat exon 33L, suggesting that the mechanism for regulating exon 33L expression during cardiac development is conserved in human and rodents. In the brain, two exons, 9* and 33, of Ca\textsubscript{v}1.2 channel are developmentally regulated (30). Fox proteins 1 and 2 regulate the expression of the two exons during embryonic mouse brain development. In this study, the longer exon 33L is formed by adopting an alternate acceptor site. As there is no intron between exon 33L and the downstream exon 33, the splicing mechanism of 33L is likely regulated by Fox proteins as well. The increase of Fox proteins during embryonic development causes more inclusion of exon 33 in cerebral Ca\textsubscript{v}1.2 channel (30). Therefore, the lower level of Fox proteins in fetal rats may affect the selection of the acceptor sites for exon 33 or exon 33L. Further experiments in changing the Fox binding elements locating within the intron after exon 33 will help to elucidate the underlying mechanism.

To the best of our knowledge, only one report showed an alteration of the structure of IVS3 segment (encoded by alternatively spliced exons 31/32) of Ca\textsubscript{v}1.2 channels during cardiac development (18). Interestingly, subsequent studies revealed a reappearance of the fetal Ca\textsubscript{v}1.2 channel in hearts of rat myocardial infarction and human heart failure (13, 19), indicating an alteration of Ca\textsubscript{v}1.2 channel in response to cardiac malfunctions.

In conclusion, we report here for the first time that the neonatal rat heart expresses a higher level of an aberrant truncated Ca\textsubscript{v}1.2\textsubscript{33L} channel. Albeit not permeable to Ca\textsuperscript{2+} ions, Ca\textsubscript{v}1.2\textsubscript{33L} channels affect the functional reference channels, mainly via competing for \( \beta \) subunits. Due to a genomic difference, human Ca\textsubscript{v}1.2\textsubscript{33L} channels are complete and functional channels that conduct Ca\textsuperscript{2+} ions. Importantly, human exon 33L is also developmentally regulated in heart. Neonatal Ca\textsubscript{v}1.2 splice variants have been shown to re-emerge in chronic heart diseases (13, 19). Therefore, it is important to understand whether these neonatal channels appear and function in cardiac diseases.

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