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C-terminal phosphorylation of Na_v1.5 impairs FGF13-dependent regulation of channel inactivation

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Voltage-gated Na⁺ (Na_v) channels are key regulators of myocardial excitability, and Ca²⁺/calmodulin-dependent protein kinase II (CaMKII)-dependent alterations in Na_v1.5 channel inactivation are emerging as a critical determinant of arrhythmias in heart failure. However, the global native phosphorylation pattern of Na_v1.5 subunits associated with these arrhythmogenic disorders and the associated channel regulatory defects remain unknown. Here, we undertook phosphoproteomic analyses to identify and quantify *in situ* the phosphorylation sites in the Na_v1.5 proteins purified from adult WT and failing CaMKII δ_c -overexpressing (CaMKII δ_c -Tg) mouse ventricles. Of 19 native Na_v1.5 phosphorylation sites identified, two C-terminal phosphoserines at positions 1938 and 1989 showed increased phosphorylation in the CaMKII δ_c -Tg compared with the WT ventricles. We then tested the hypothesis that phosphorylation at these two sites impairs fibroblast growth factor 13 (FGF13)-dependent regulation of Na_v1.5 channel inactivation. Whole-cell voltage-clamp analyses in HEK293 cells demonstrated that FGF13 increases Na_v1.5 channel availability and decreases late Na⁺ current, two effects that were abrogated with Na_v1.5 mutants mimicking phosphorylation at both sites. Additional co-immunoprecipitation experiments revealed that FGF13 potentiates the binding of calmodulin to Na_v1.5 and that phosphomimetic mutations at both sites decrease the interaction of FGF13 and, consequently, of calmodulin with Na_v1.5. Together, we have identified two novel native phosphoryla-

tion sites in the C terminus of Na_v1.5 that impair FGF13-dependent regulation of channel inactivation and may contribute to CaMKII δ_c -dependent arrhythmogenic disorders in failing hearts.

Voltage-gated Na⁺ (Na_v) channels are critical determinants of myocardial excitability, and defects in Na_v channel functioning or regulation in the context of inherited or acquired cardiac disease increase the risk of life-threatening arrhythmias (1). Under physiological conditions, the primary ventricular Na_v channel subunits, the Na_v1.5 channels, activate and inactivate rapidly to generate the transient peak Na⁺ current, *I*_{Na}, responsible for the depolarization phase and the propagation of action potentials. However, a small proportion of Na_v channels inactivates slowly to generate a late or persistent Na⁺ current, referred to as *I*_{NaL}, contributing markedly to determining action potential waveform, duration, and refractoriness. Among the various determinants recognized to cause Na_v channel dysfunctions and consequent increased risk of acquired arrhythmias is the activation of Ca²⁺/calmodulin-dependent protein kinase II (CaMKII)³ (2–7). A prime example of this dysregulation is heart failure in which an increased *I*_{NaL} has been linked to the activation of CaMKII (2, 4–7) and associated alteration in Na_v1.5 phosphorylation (7). This increased *I*_{NaL} in failing hearts is sometimes accompanied by a decreased Na_v channel availability caused by a hyperpolarizing shift of the voltage dependence of steady-state *I*_{Na} inactivation (2, 7). Three phosphorylation sites in the first linker loop of Na_v1.5, at positions 516 (8, 9), 571 (10–12), and 594 (8), have been suggested to play causative roles in these deleterious CaMKII-dependent mechanisms. Nonetheless, the global native phosphorylation pattern of Na_v1.5 channels associated with CaMKII activation

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This article contains supplemental Tables S1–S4 and Fig. S1.

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³ The abbreviations used are: CaMKII, Ca²⁺/calmodulin-dependent protein kinase II; CaMKII δ_c -Tg, transgenic mouse overexpressing the cytosolic isoform of the δ subunit of Ca²⁺/calmodulin-dependent protein kinase II; CTD, C-terminal domain of Na_v channels; iFGF, intracellular fibroblast growth factor; IP, immunoprecipitation; MS1, mass spectrum of precursor ions; m α Na_vPAN, anti-Na_v α subunit monoclonal antibody; Na_v α subunit, voltage-gated Na⁺ (Na_v) channel pore-forming (α) subunit; TTX, tetrodotoxin; FA, formic acid; ACN, acetonitrile; ANOVA, analysis of variance; pF, picofarad; HP, holding potential; CaM, calmodulin; m, mouse.

Phosphorylation and inactivation of cardiac Na_v1.5 channels

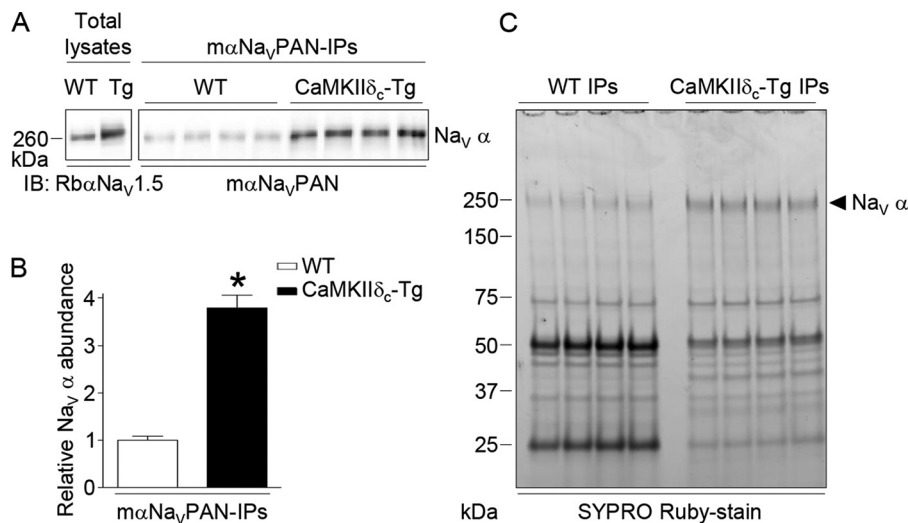


Figure 1. Immunoprecipitation of Na_v channel complexes from adult WT and CaMKIIδ_c-Tg mouse ventricles. *A*, representative Na_v α Western blots of total lysates and immunoprecipitated proteins from adult WT and CaMKIIδ_c-Tg mouse ventricles with the anti-Na_vPAN monoclonal antibody (*mαNa_vPAN-IPs*) probed with the anti-Na_v1.5 rabbit polyclonal (*RbαNa_v1.5*) and the *mαNa_vPAN* antibodies, respectively. *B*, mean ± S.E. relative Na_v α protein abundance in WT (*n* = 4) and CaMKIIδ_c-Tg (*n* = 4) IPs. *, *p* < 0.05, Mann-Whitney test. *C*, SYPRO Ruby-stained gel of *mαNa_vPAN-IPs* from WT and CaMKIIδ_c-Tg mouse ventricles. Relative abundance of proteins running at the molecular weight of Na_v α subunits is higher in CaMKIIδ_c-Tg IPs than in WT IPs.

in failing hearts as well as the molecular mechanisms associated with altered phosphorylation that underlie the defects in channel inactivation remain unknown.

The understanding of Na_v1.5 channel inactivation has received much attention over the past decade and has recently been buttressed by the generation of several crystal structures of the cytoplasmic C-terminal domain (CTD) of Na_v1.5 (as well as of other Na_v subunits) in complex with Ca²⁺-free or Ca²⁺-bound calmodulin (CaM) and/or a member of the intracellular fibroblast growth factor (iFGF) family of proteins (13–16). The Ca²⁺-binding protein CaM acts as a sensor for Ca²⁺ and regulates inactivation of Na_v channels, while underlying mechanisms still elude consensus. Depending on whether CaM is loaded with Ca²⁺, it binds different sites within and around the IQ motif in the CTD and/or the third linker loop of Na_v channels, thereby inducing conformational changes and affecting channel inactivation and the late Na⁺ current (13–19). iFGFs, which include FGF11–14, are also constitutive Na_v channel-associated proteins, binding the CTD of Na_v channels, just upstream of the IQ motif (14–16, 20–22). The most consistent effect of iFGFs is to increase the availability of Na_v channels by shifting the voltage dependence of steady-state inactivation toward depolarized potentials (15, 20–26). The central role of the Na_v1.5 CTD as well as of iFGFs and CaM in regulating the inactivation properties of Na_v1.5 channels is highlighted by the effects of several mutations observed in the arrhythmic long QT3 (LQT3) and Brugada syndromes that map to their binding interfaces (14, 27–29).

This study uses a mass spectrometry (MS)-based phosphoproteomic analysis to identify and quantify *in situ* the native phosphorylation sites in the Na_v1.5 proteins purified from failing CaMKIIδ_c-overexpressing (CaMKIIδ_c-Tg) *versus* non-failing wild-type (WT) mouse ventricles. The rationale for using this transgenic and failing mouse model is to identify the Na_v1.5 phosphorylation sites that participate in the channel inactivation defects associated with the activation of CaMKIIδ_c

in heart failure. Further biochemical and electrophysiological approaches in human embryonic kidney 293 (HEK293) cells were then used to investigate the impact of phosphorylation at two C-terminal serine residues on the interaction of FGF13 and CaM with Na_v1.5 and on the inactivation properties of Na_v1.5 channels.

Results

Purification and characterization of Na_v channel complexes from WT and CaMKIIδ_c-Tg mouse ventricles

Total lysates from four adult WT and four CaMKIIδ_c-Tg (30) mouse (m) ventricles were prepared, pooled, and used in eight distinct IPs (four WT IPs and four CaMKIIδ_c-Tg IPs) using the *mαNa_vPAN*-specific antibody. As illustrated in Fig. 1A and consistent with previous findings (7), Western blot analyses of total lysates showed greater Na_v1.5 protein expression in the CaMKIIδ_c-Tg compared with the WT ventricles. This difference in total protein expression resulted in significantly (*p* < 0.05) higher Na_v1.5 protein abundance (3.8-fold) in the *mαNa_vPAN-IPs* from the CaMKIIδ_c-Tg than from the WT ventricles (Fig. 1, A and B). Accordingly, analyses of *mαNa_vPAN-IPs* on SYPRO Ruby-stained gels revealed the presence of a band corresponding to the molecular weight of Na_v α subunits (31), the intensities of which are higher in the CaMKIIδ_c-Tg IPs than in the WT IPs (Fig. 1C).

The protein components of isolated Na_v channel complexes from WT and CaMKIIδ_c-Tg ventricles were identified by MS using three distinct mass spectrometers: an LTQ-Orbitrap XL, an LTQ-Orbitrap Elite, and a TripleTOF 5600 Plus. The Na_v1.5 protein was the most abundant protein in the *mαNa_vPAN-IPs* from both WT and CaMKIIδ_c-Tg ventricles with average numbers of total exclusive MS/MS spectra acquired increasing from 95 using the Orbitrap XL to 177 using the Orbitrap Elite and 524 using the TripleTOF (Fig. 2A). These greater sensitivities of the Orbitrap Elite and TripleTOF mass spectrometers improved

A

Protein		LTQ-Orbitrap XL	LTQ-Orbitrap Elite	TripleTOF 5600 Plus
Na _v 1.5	Average number of unique peptides (total spectra)	50 (95)	57 (177)	84 (524)
	Amino acid sequence coverage	27%	28%	32%
	Amino acid sequence coverage (excluding TD)	36%	38%	43%
Na _v 1.4	Average number of unique peptides (total spectra)	4 (5)	6 (7)	16 (30)
Na _v 1.7	Average number of unique peptides (total spectra)	-	1 (1)	4 (4)
Na _v 1.8	Average number of unique peptides (total spectra)	-	1 (1)	2 (2)
Na _v 1.3	Average number of unique peptides (total spectra)	-	-	3 (3)

B

1 MANFLLPRTGSSFRFRTR**ESLAAIEKR**MAEKQAR**CSATSQES**REGLP~~EE~~APRPQDLQASKLPDLYGN
 N-TERM
 71 PPRELIGEPLELDPPFYSTQKTFIVLNK**GKTI**FR**F**SATNALYVLSPPFHPVRRAAVK**LLVHSLFSMLIMCT**
 N-TERM IS1
 141 ILLTNCVFMAGHDPPPPWTKYVEYTFITAIYTFESLVKILARGFCLHAFTFLRD**PWNWLD**FSVIVMAYVSENI
 IS2 IS3
 211 KLGNSALRTRFRVLRALK**TI**SVIPGLK**TI**VGALIQSVK**KLAD**VMVLTVFCLSVFALIGLQLEFMGNLRHKC
 IS4 IS5
 281 VRNFTELNGTNGSVEADGIVVNSLDVYLNDPANYLLKNGTIDVLLCGNSSDAGTCPEGYRCLKAGENPDH
 351 GYTSFDSFAWAFALFLR**LMTQDC**WERLYQQ**TLR**SAGKIY**MI**FFMLVIFLGSFYLVNLI**LAVV**AMAYEEQN
 IS6
 421 QATIAETEKEEK**R**FQEAEMELKKEHEAL**TIR**GVDT**VSR**SL**EM**SPLAPVNHERRSKRR**KRLSS**C**EDGG**
 Loop I
 491 **DDRLPKS**D**EDG**PRALNQL**SL**THGL**R**TSMP**R**SS**R**GS**I**FTFRRRD**Q**G**EAD**FADDENSTAGESESHRTS
 Loop I
 561 **LLVPWPLRR**ST**Q**GQ**P**FGT**S**APGHV**LN**GKR**N**STVDCNGVVS**LL**GAGDAEATSPGSHLLRPIVLDLDRPPDT
 Loop I
 631 TTPSEEPGGQMLTPQAPCADGFEEPGAR**Q**AL**S**AVSVLTSAL**EE**LEESH**RK**CPPCWNRFAQHYLWIECC
 Loop I
 701 PLWMSIQKVK**FV**VM**DF**FAD**L**TI**MC**IVL**NT**LM**FA**LEHYNMTAE**FE**ML**Q**VGNLV**FT**G**I**FT**AE**MT**FK**II**A**
 IIS1 IIS2
 771 **LD**PPYY**F**Q**GW**NI**F**DS**I**IV**IL**SL**ME**L**GL**SR**MG**NLS**V**LR**S**FRLLRV**FK**LAK**S**W**PT**NT**L**IK**I**IGNSV**GA**L**G**
 IIS3 IIS4
 841 **N**L**RL**V**L**AI**IV**F**FA**V**VM**Q**L**FG**K**NY**S**ELRH**R**IS**DS**GL**LR**WHM**DF**H**FA**FL**I**FR**IL**CG**E**W**IE**T**M**W**DC**ME
 IIS5
 911 VSG**QS****L**CL**L**V**L**FL**VM**V**IG**N**L**V**LN**L**FL**ALL**SS**F**S**AD**N**L**T**AP**ED**GM**N**L**Q**LAL**AR**I**Q**RL**F**V**K**R**TT**W
 IIS6 Loop II
 981 **D**FC**CG**LL**R**RR**PK****K**PA**A**L**A**TH**S**Q**L**PS**C**IA**AP****R****P**PP**PE**VE**K**AP**P**ARK**ET**RF**E**ED**K**RP**G**Q**T**PG**D**TE**P**VC**V**P
 Loop II
 1051 IAVA**ES**DT**DD**QE**ED**EN**SL**GT**EE**ES**SK****Q**ES**Q**V**VS**GG**HE**PP**Q**E**PR**AW**S**Q**VS**ET**TS**SE**AE**AST**S**Q**AD**W**Q**
 Loop II
 1121 ERE**AE**PR**AP**CG**ET**PE**DS**Y**EG**ST**AD**MT**NT**AD**LE**Q**I**PD**L**GED**VK****DP**ED**CF**T**EG**CV**RR**CC**PM**CV**DT**Q**AP**
 Loop II
 1191 **G**K**V**WR**L**R**K**TC**YR**I**VE**HS**WF**ET**FI****I**FM**LL**SS**GA**L**AF**ED**I**Y**LE**ER**K**TI**K****V**LE**Y**AD**K**M**F**TY**V**F**LE**ML**LK**
 IIS1 IIS2
 1261 **W**V**AY**GF**K**K**Y**FT**NA**WC**W**LD**PL**IV**DS**LV**SL**V**ANT**L**GF**AE**MG**PI**K**SL**RT**L**R**AL**R**PL**RS**RF**EG**MR**V**V**N**AL
 IIS3 IIS4
 1331 V**G**A**I**PS**IM****V**LL**V**CL**I**F**W**L**F**IS**MG**V**N**LE**F**AG**FR**GC**IN**Q**TE**GD**L**PL**NY**TI**V**NN**K**SE**CS**FN**VT**G**EL**Y**W**TK
 IIS5
 1401 **V**K**V**N**F**DN**V**GA**Y**L**AL**L**Q**V**AT**FK**W**MD**IM**Y**AA**VD**S**R**GE**Y**EE**Q**P**Q**W**ED**N**L**MY**I**Y**F**V**V**I**I**F**GS**F**TT**L**N**L**F**IG**
 IIS6
 1471 **V**I**I**DN**F**N**Q**Q**K****KL**GG**Q**DI**FM**TE**EQ**K**Y**Y**N**AM**K****LG**SK**K**P**Q**K**P**I**PR**L**N**K**Y**Q**G**F**I**FD**IV**T**K**Q**A**FD**V**T**IM**FL
 Loop III IIS1
 1541 **I**CL**N**VM**TV**ME**TD**DD**Q**SP**E**K**V**N**L**L**AK**IN**L**LF**V**AI**FT**G**EC**IV**K**MA**L**R**H**Y**FT**NS**W**NI**F**DE**V**V**V**LL**S**IV**GT**V
 IIS2 IIS3
 1611 **L**SD**I**IQ**K**Y**FS**PT**LF**RV**IR**L**AR**I**GR**IL**LR**I**RG**AK**IR**TL**LF**AL**M**MS**L**PA**LF**NI**GL**LL**FL**VM**F**Y**S**I**F**GM**A**
 IIS4 IIS5
 1681 **N**F**AY**V**K**WE**AG**ID**DM**FN**Q**TF**AN**S**ML**CL**F**Q**IT**TS**AG**W**D**GL**LS**P**IL**NT**GP**PY**CD**PN**LS**NG**S**R**GN**CG**SP**AV
 1751 **G**IL**FF**TT**Y**II**IS**FL**IV**N**MY**IA**IL**EN**FS**V**AT**EE**ST**EL**SE**DD**FD**MF**E**Y**IE**W**FK**DP**E**AT**Q**F**IE**YL**AL**SD**F**
 IIS6 C-TERM
 1821 AD**AL**SE**PL**R**T**AK**P**Q**IS**SL**IN**MD**LP**MS**GD**R**I**HC**MD**IL**FA**FT**KR**V**L**GES**G**EM**D**AL**K**I**Q**ME**E**K**F**MA**AN**P**R**I
 C-TERM iFGF-binding site
 1891 **S**Y**E**PI**TT**LL**RR****K**HE**VS**AT**VI**Q**RA**FR**RL**L**Q**RS**V**K**H**AS**FL**FR**Q**Q**AG****S**GL**S**DE**D**AP**ER**E**GL**I**AY**MM**N**EN**F**
 IQ-motif C-TERM
 1961 **S**RR**S**GL**SS**SS**IS**TS**FF**PS**Y**DS**V**TR**AT****D**N**LP**V**R**AS**D**YS**R**SE**D**L**AD**FP**SP**DR**D**RES**IV**
 C-TERM

C

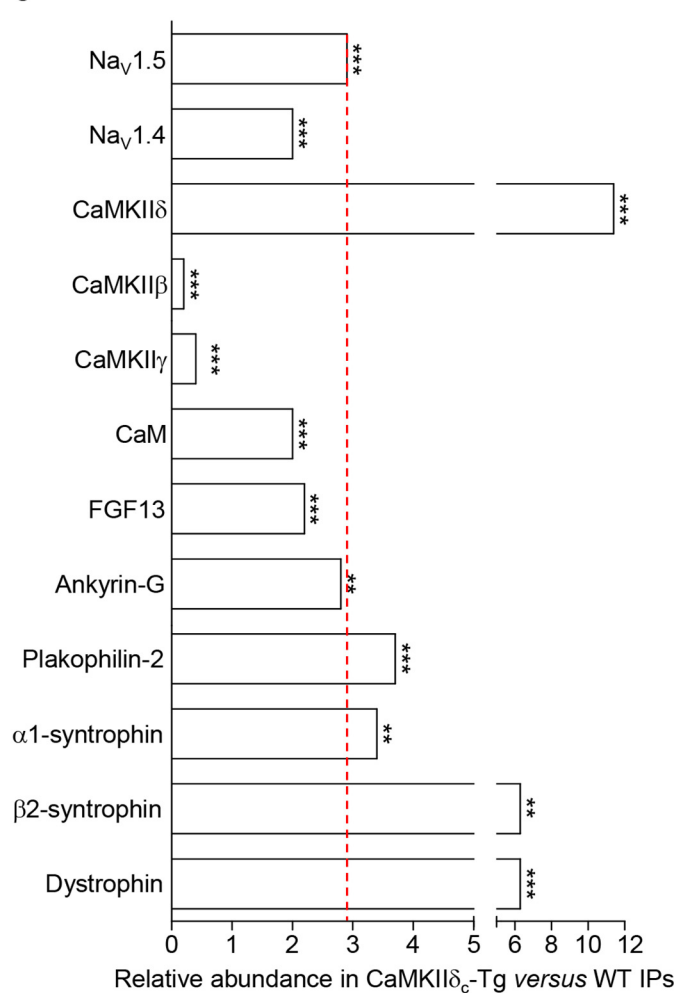


Figure 2. MS protein identification in immunoprecipitated Na_v channel complexes from WT and CaMKII_δ-Tg mouse ventricles. A, Na_v α subunits identified using the LTQ-Orbitrap XL, LTQ-Orbitrap Elite, and TripleTOF 5600 Plus mass spectrometers. The average numbers of exclusive unique peptides and total spectra for each Na_v α subunit and the percent amino acid sequence coverages obtained for Na_v1.5, including or excluding transmembrane domains (TD), are presented. In addition to Na_v1.5, which is the most abundant protein in the mαNa_vPAN-IPs, Na_v1.4 is also detected, and the greater sensitivity of the Orbitrap Elite and TripleTOF mass spectrometers allowed the identification of Na_v1.7, Na_v1.8, and Na_v1.3. B, amino acid sequence coverage obtained for the (mouse) Na_v1.5 protein (NP_001240789). Detected peptides are highlighted in yellow; identified phosphorylation sites are highlighted in blue (sites already identified in our previous MS analyses) and red (newly identified sites in the present study); transmembrane segments (S1–S6) in each domain (I–IV) are in bold and underlined in black; loops I, II, and III correspond to interdomains I and II, II and III, and III and IV, respectively; and binding sites for iFGF and calmodulin (IQ-motif) are boxed in green and orange, respectively. C, relative abundances of Na_v α subunits and previously characterized Na_v1.5 channel-associated/regulatory proteins in the CaMKII_δ-Tg IPs (n = 4) versus the WT IPs (n = 4) were calculated from the entire (Orbitrap XL) MS1 peptide data set using the DAnTE statistical software (**, p < 0.01; ***, p < 0.001).

Phosphorylation and inactivation of cardiac Na_v1.5 channels

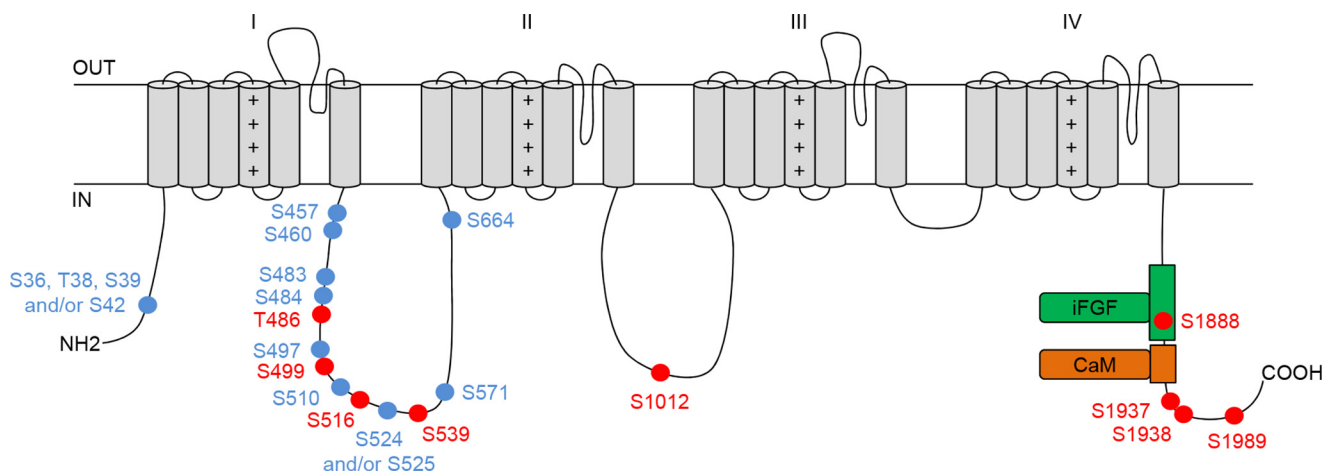


Figure 3. Localization of MS-identified *in situ* phosphorylation sites on the mouse ventricular Na_v1.5 α subunit protein. Among the 19 phosphorylation sites identified, 10 (in blue) had already been identified in our previous MS analyses and 9 (in red) are novel. Four and two phosphorylation site locations are possible at amino acids 36–42 and 524–525, respectively. The three newly identified C-terminal phosphoserines at positions 1888, 1937, and 1938 (pSer-1888, pSer-1937, and pSer-1938) are in close proximity to the binding sites for iFGF and calmodulin (IQ-motif).

the Na_v1.5 amino acid sequence coverage from 27 to 28 and 32%, respectively, and from 36 to 38 and 43% with the transmembrane domains removed. Na_v1.4 was the second most abundant Na_v α subunit with much fewer spectra acquired: 5, 7, and 30 using the Orbitrap XL, Orbitrap Elite, and TripleTOF, respectively. Most interestingly, the greater sensitivity of the TripleTOF mass spectrometer allowed the identification of three additional Na_v α subunits, Na_v1.7 (four spectra), Na_v1.3 (three spectra), and Na_v1.8 (two spectra). These experiments also led to the reliable identification of several previously identified Na_v1.5 channel-associated/regulatory proteins (32), including the δ , β , and γ subunits of CaMKII, CaM, FGF13, and ankyrin-G (Fig. 2C). The supplemental Tables 1–3 provide the complete lists of identified peptides and proteins using the Orbitrap XL, Elite, and the TripleTOF mass spectrometers, respectively.

The relative abundances of identified proteins in the α Na_v PAN-IPs from the CaMKII δ_c -Tg *versus* the WT ventricles were calculated from the MS1 peptide data (Orbitrap XL) using the DAnTE statistical analysis tool (33–35). Consistent with the biochemistry data (Fig. 1), the Na_v1.5 protein is 2.9-fold ($p < 0.001$) more represented in the CaMKII δ_c -Tg IPs than in the WT IPs (Fig. 2C). Interestingly, this quantitative MS analysis also revealed that of the 10 Na_v1.5-associated/regulatory proteins identified, the abundance ratios of CaMKII δ , β 2-syntrophin, and dystrophin in the CaMKII δ_c -Tg *versus* the WT IPs are substantially greater than the 2.9-fold abundance ratio obtained for Na_v1.5. On the contrary, the CaMKII β and γ subunits as well as CaM and FGF13 are relatively less represented compared with Na_v1.5. The abundance ratios obtained for ankyrin-G, plakophilin-2, and α 1-syntrophin are close to the Na_v1.5 abundance ratio. Taken together, these observations demonstrate that Na_v1.5 is more expressed in the CaMKII δ_c -Tg than in the WT mouse ventricles, which, as a consequence, led to greater Na_v1.5 IP yields from the CaMKII δ_c -Tg than from the WT ventricles. These analyses also suggest different relative compositions of associated/regulatory proteins in Na_v1.5 channel complexes in the CaMKII δ_c -Tg compared with the WT ventricles.

Identification of native Na_v1.5 phosphorylation sites from WT and CaMKII δ_c -Tg mouse ventricles

To determine the global native phosphorylation pattern of Na_v1.5 channels associated with the overexpression of CaMKII δ_c , a quantitative phosphoproteomic analysis of the Na_v1.5 proteins purified from the CaMKII δ_c -Tg and the WT mouse ventricles was performed. This phosphoproteomic analysis unambiguously allowed the identification of 19 native phosphorylation sites in the mouse ventricular Na_v1.5 protein (Figs. 2B and 3). Table 1 lists the phosphopeptides enabling the best phosphorylation site assignment(s), with the mass spectrometer used for the identification and the percentages of maximum intensities of site-discriminating ions, for each phosphorylation site. Representative MS/MS (and MS1) phosphopeptide spectra are presented in supplemental Fig. 1. Descriptions of all detected site-discriminating and supporting ions (calculated mass errors, confirmations of charge states) are given in supplemental Tables 4. Among the 19 Na_v1.5 phosphoserines (pSer) or phosphothreonines (pThr) identified, 10 (in blue) had already been identified in our previous phosphoproteomic analyses (31) and nine (in red) are novel (Figs. 2B and 3). These nine novel phosphorylation sites are located in the first (pThr-486, pSer-499, pSer-516, and pSer-539) and second (pSer-1012) intracellular linker loops as well as in the CTD (pSer-1888, pSer-1937, pSer-1938, and pSer-1989) of Na_v1.5. Interestingly, the three C-terminal pSer-1888, pSer-1937, and pSer-1938 are in close proximity to the binding sites for the iFGFs (14–16, 20–22) and CaM (IQ-motif) (13–19).

To determine whether one or several of these MS-identified Na_v1.5 phosphorylation sites are associated with the overexpression of CaMKII δ_c , the relative abundance of each Na_v1.5 phosphopeptide in the CaMKII δ_c -Tg *versus* the WT α Na_v PAN-IPs was determined by label-free quantification of MS1 data (36). As illustrated in Fig. 4A, and consistent with the quantification of the biochemical (Fig. 1) and the MS protein (Fig. 2C) data, the unphosphorylated Na_v1.5 peptides are 3.6-fold more represented in the CaMKII δ_c -Tg *versus* the WT IPs. Of the 86 unphosphorylated and 32 phosphorylated Na_v1.5 peptides

Table 1

Phosphorylation sites, phosphopeptides, and site-discriminating ions identified in immunoprecipitated Na_v1.5 α subunits from adult WT and/or CaMKIIδ_c-Tg mouse ventricles using MS

The site-discriminating ions observed in the high-resolution (LTQ-Orbitrap Elite or TripleTOF 5600 Plus) or low-resolution (LTQ-Orbitrap XL) MS/MS spectra of each annotated Na_v1.5 phosphopeptide support the assignment of the indicated phosphorylation site(s). The charge state as well as the percentage of maximum intensity of each observed unphosphorylated and phosphorylated site-discriminating (b and y) ion are reported in parentheses; the minus symbol indicates that the charge state could not be determined.

Phosphorylation site	MS machine	Phosphopeptide sequence	m/z (charge)	b ion	Phospho b ion	y ion	Phospho y ion
pS36, pT38, pS39 and/or pS42	Elite	35-GSATSQESREGLPEEEAPRPQLDLQASK	773.365 (+4)	(--)	b12 (-, 21.7)	y16 (+2, 72.4) y16 (+3, 24.3)	(--)
pS457 + pS460	TTOF	452-GVDTV(pS)RS(pS)LEMSPLAPVTNHER	661.305 (+4)	b5 (-, 1.1)	b6 (-, 2.7) b7 (+1, 3.0) b8 (-, 0.4) b11 (-, 3.2)	y12 (+1, 4.2) y13 (+2, 1.1)	y18 (-, 0.4) y20 (+3, 0.4)
pS460	Elite	452-GVDTVSR(pS)LEMSPLAPVTNHER	854.739 (+3)	b8 (+1, 2.3)	b11 (+1, 11.9)	y12 (+1, 3.0)	(--)
pS483 + pS484	TTOF	481-RL(pS)(pS)GTEDGGDDR(L)PK	621.596 (+3)	b2 (-, 37.6)	b3 (-, 2.1) b6 (+1, 2.0)	y12 (+1, 3.0)	y13 (-, 1.9)
pS483 or pS484 + pT486	TTOF	480-KRLSSGTEDGGDDR(L)PK	664.294 (+3)	b3 (-, 16.7)	b6 (-, 19.5) b9 (+1, 7.6)	y10 (+1, 10.2)	y11 (-, 6.0)
pS483 + pS484 + pT486	TTOF	481-RL(pS)(pS)G(pT)EDGGDDR(L)PK	648.251 (+3)	b2 (-, 98.9)	(--)	y9 (+1, 83.8)	(--)
pS484	TTOF	482-LS(pS)GTEDGGDDR(L)PK	542.906 (+3)	b2 (+, 20.2)	b3 (-, 9.2)	y12 (+1, 40.9)	y13 (+2, 22.3)
pS497	TTOF	482-LSSGTEDGGDDR(L)PK(pS)DSEDGPR	618.266 (+4)	b12 (-, 2.4)	(--)	y7 (+1, 18.6)	y8 (-, 6.1) y12 (+2, 5.1)
pS499	XL	497-SD(pS)EDGPR	471.665 (+2)	b2 (-, 2.1)	b5 (-, 7.8)	y4 (-, 19.6)	y6 (-, 16.6)
pS510	Elite	505-ALNQL(pS)LTHGLSR	745.380 (+2)	b4 (+1, 10.5)	b9 (+2, 13.8)	y7 (+1, 29.4)	y8 (-, 5.4)
pS510	Elite	505-ALNQL(pS)LTHGLSR	497.256 (+3)	b3 (-, 5.5)	b9 (+2, 11.3)	y7 (+1, 12.4)	y10 (-, 6.8) y11 (+2, 5.0)
pS516	TTOF	505-ALNQLSLTHGL(pS)R	497.260 (+3)	b8 (-, 36.7)	(--)	(--)	y3 (-, 6.3) y4 (-, 2.8)
pS524 and/or pS525	Elite	524-SSRGSIFTR	619.290 (+2)	(--)	(--)	y7 (-, 28.3)	(--)
pS539	TTOF	536-DQG(pS)EADFADDENSTAGESESH	845.312 (+3)	b2 (+1, 29.9) b3 (-, 6.6)	b7 (+1, 10.4)	y17 (-, 29.8) y18 (+2, 9.0)	(--)
pS571	TTOF	569-RP(pS)TQQQPGFGTSAPGHV(L)NGK	569.029 (+4)	b2 (+1, 63.5)	b4 (+1, 2.0)	y15 (-, 100) y19 (-, 8.2)	y20 (-, 1.8)
pS664	XL	662-AL(pS)AVSVLTSALEEELEESHK	783.731 (+3)	(--)	b3 (-, 0.8) b5 (-, 2.0)	y17 (-, 27.0)	y19 (-, 34.4)
pS1012	TTOF	993-KPAALATHSQLPSC(+57)IAAPR(pS)PPPPEVEK	758.148 (+4)	b18 (-, 14.2)	(--)	y8 (-, 100)	y11 (+1, 4.6)
pS1888	TTOF	1884-AANP(pS)KISYEPITTLR	647.994 (+3)	(--)	(--)	y10 (+1, 12.6) y11 (+2, 2.1)	(--)
pS1937 and/or pS1938	TTOF	1933-Q(-17)QAGSSGLSDE(D)APER	855.334 (+2)	b4 (+1, 23.8)	b7 (-, 8.1)	y10 (+1, 37.1)	(--)
pS1937 + pS1938	TTOF	1934-QAG(pS)(pS)GLSDE(D)APER	560.211 (+3)	(--)	(--)	y10 (+1, 7.6)	y11 (-, 3.0)
pS1989	TTOF	1987-AT(pS)DNLPVR	526.742 (+2)	b2 (-, 2.9)	b5 (-, 4.8)	y6 (-, 15.4)	y7 (-, 19.7)

(118 peptides total), only the three phosphopeptides AT(pS)DNLPVR, RL(pS)(pS)GTEDGGDDR, and AL(pS)AVSVLTSALEEELEESHK (marked with a † in Fig. 4A), exhibiting phosphorylation(s) on serines 1989 (pSer-1989), 483 and 484 (pSer-483 and pSer-484), and 664 (pSer-664), respectively, present fold change ratios (8.96-, 7.13-, and 0.58-fold) significantly different from the median ratio (Tukey whisker analysis, Fig. 4B). Nonetheless, the other phosphopeptides assigning pSer-483 and/or pSer-484 do not show any significant abundance changes compared with the median ratio (Fig. 4, A and B); and the abundance change observed for the phosphopeptide identifying pSer-664 is not significantly different between the mαNa_vPAN-IPs from the CaMKIIδ_c-Tg and the WT ventricles (Fig. 4A). In addition, the phosphopeptide Q(-17.03)QAGSS-

GLSDE(D)APER, assigning pSer-1937 and/or pSer-1938 (Table 1), is present in the CaMKIIδ_c-Tg IPs (*n* = 3/4) and absent in the WT IPs (*n* = 0/4) (Fig. 4C). Relative abundances of all the other Na_v1.5 phosphopeptides are comparable with the median (or mean) relative abundance found for the unphosphorylated Na_v1.5 peptides (Fig. 4, A and B).

Amino acid sequence alignment of the flanking regions of pSer-1937/38 and pSer-1989 showed a good conservation of Ser-1938 and Ser-1989 across orthologous sequences (Ser-1937 is not well-conserved) and, interestingly, revealed that Ser-1989 conforms to the consensus CaMKII phosphorylation site (RXX(S/T) Fig. 4D (37)). Taken together, these phosphoproteomic analyses identified nine novel native Na_v1.5 phosphorylation sites, of which two conserved serines in the CTD, at

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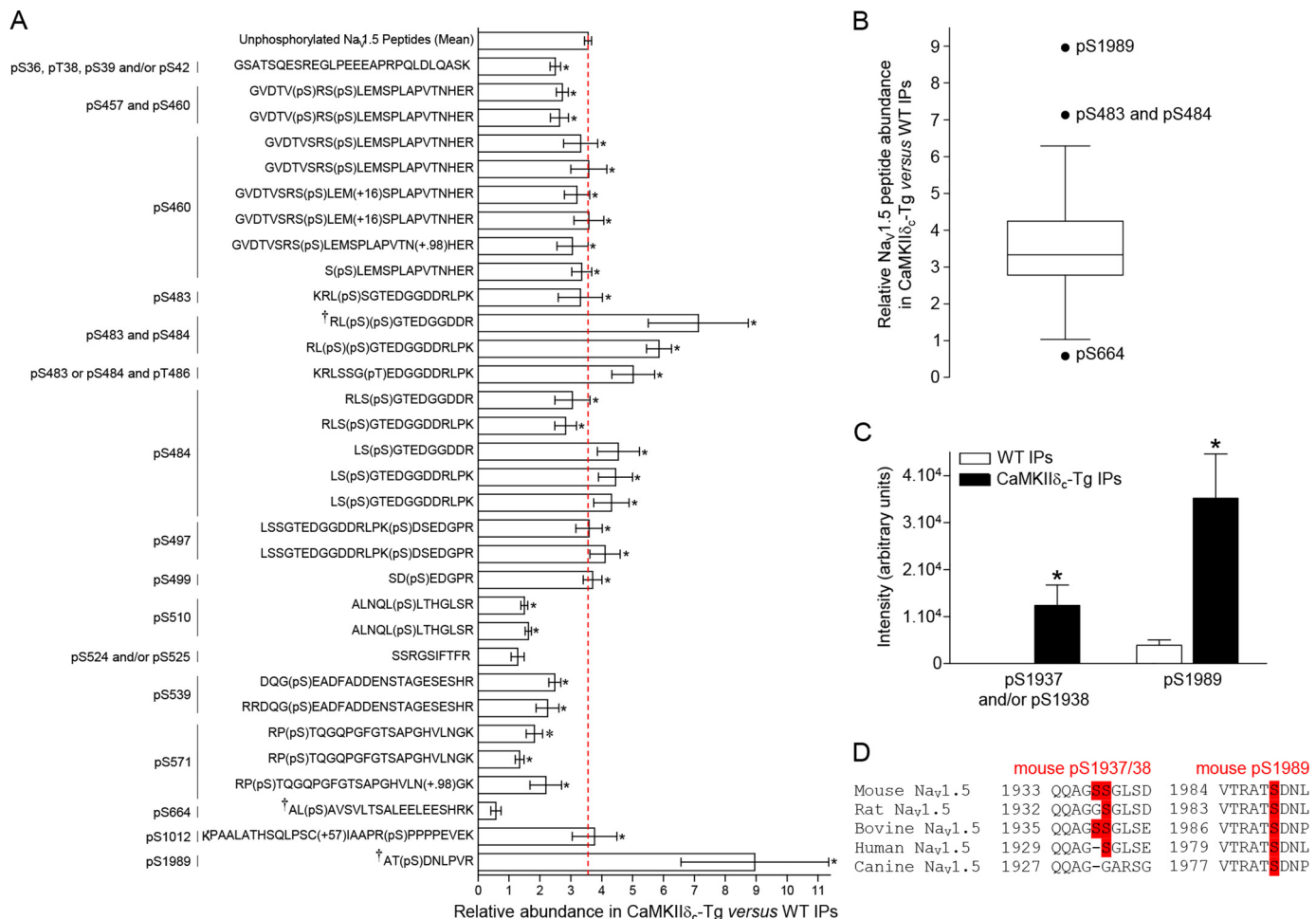


Figure 4. Quantification analysis of Na_v1.5 phosphorylation sites in the CaMKIIδ_c-Tg versus the WT mαNa_vPAN-IPs. A, relative abundances of 32 Na_v1.5 phosphopeptides allowing assignments of the listed phosphorylation sites, in the CaMKIIδ_c-Tg (*n* = 4) versus the WT (*n* = 4) IPs, were calculated using label-free quantification of the (Orbitrap XL) MS1 data. The mean ± S.E. relative abundance of unphosphorylated Na_v1.5 peptides in the CaMKIIδ_c-Tg versus the WT IPs was calculated from 86 unphosphorylated Na_v1.5 peptides (minimum Scaffold peptide probability scores of 95%). Consistent with the biochemistry data (Fig. 1) and the DANte protein statistical analysis (Fig. 2C), the unphosphorylated Na_v1.5 peptides are 3.6-fold more represented in the CaMKIIδ_c-Tg IPs than in the WT IPs (red dashed line). The relative abundances of individual Na_v1.5 phosphopeptides are significantly (*, *p* < 0.05, Mann-Whitney test) different in the CaMKIIδ_c-Tg compared with the WT IPs. B, Tukey whisker analysis of Na_v1.5 peptide relative abundance in CaMKIIδ_c-Tg versus WT IPs. Of the 118 (86 unphosphorylated and 32 phosphorylated) Na_v1.5 peptides, only the phosphopeptides AT(pS)DNLPVR, RL(pS)(pS)GTEDGGDDDR, and AL(pS)AVSVLTSALEEESHK (marked with † in A), exhibiting phosphorylation(s) on serines 1989 (pSer-1989), 483 and 484 (pSer-483 and pSer-484), and 664 (pSer-664), respectively, present fold change ratios (8.96-, 7.13-, and 0.58-fold) significantly different from the median ratio. C, mean ± S.E. intensities of phosphopeptides Q(-17.03)QAGSGLSDEDAPER, assigning pSer-1937 and/or pSer-1938 (absence in WT IPs and presence in CaMKIIδ_c-Tg IPs), and AT(pS)DNLPVR, assigning pSer-1989 (8.96-fold change ratio), in WT (*n* = 4) and CaMKIIδ_c-Tg (*n* = 4) IPs. *, *p* < 0.05, Mann-Whitney test. D, conservation of the two C-terminal Na_v1.5 serines 1938 and 1989 across orthologs.

positions 1938 and 1989, show increased phosphorylation in the CaMKIIδ_c-Tg compared with the WT ventricles.

Phosphorylation at serines 1933 and 1984 impairs FGF13-dependent regulation of Na_v1.5 channel inactivation

To determine the impact of phosphorylation at these two C-terminal (mouse) serines 1938 and 1989 on the gating properties of Na_v1.5 channels, the orthologous human serine to glutamate (Na_v1.5-S1933E/S1984E, Na_v1.5-EE) or serine to alanine (Na_v1.5-S1933A/S1984A, Na_v1.5-AA) double Na_v1.5 phosphomutants were generated and analyzed by whole-cell voltage-clamp analyses in transiently transfected HEK293 cells. Because pSer-1938 and pSer-1989 are located in close proximity to the binding site for the iFGFs (14–16, 20–22), we tested the hypothesis that phosphorylation at these two sites disrupts the interaction of the iFGFs with Na_v1.5 and the associated

iFGFs-dependent regulation of Na_v1.5 channel function. To this purpose, initial experiments were aimed at exploring the effects of FGF13 on the current density and biophysical properties of heterologously expressed Na_v1.5 channels. The isoform 2 of FGF13 (FGF13–2) was chosen as the iFGF in these experiments as it is the isoform detected in the mαNa_vPAN-IPs. As illustrated in Fig. 5, these whole-cell voltage-clamp analyses demonstrated that FGF13 significantly decreases the peak Na⁺ current (*I*_{Na}) density (*p* < 0.05, Fig. 5, A and B) and shifts the voltage dependence of steady-state current inactivation toward depolarized potentials (*p* < 0.01, Fig. 5D, see detailed densities, properties, and statistics in Table 2). In contrast, no significant differences in the voltage dependence of activation (Fig. 5C) or the kinetics of activation, inactivation, and recovery from inactivation were observed upon FGF13 co-expression (Table 2). To determine the effect of Na_v1.5 phosphorylation at

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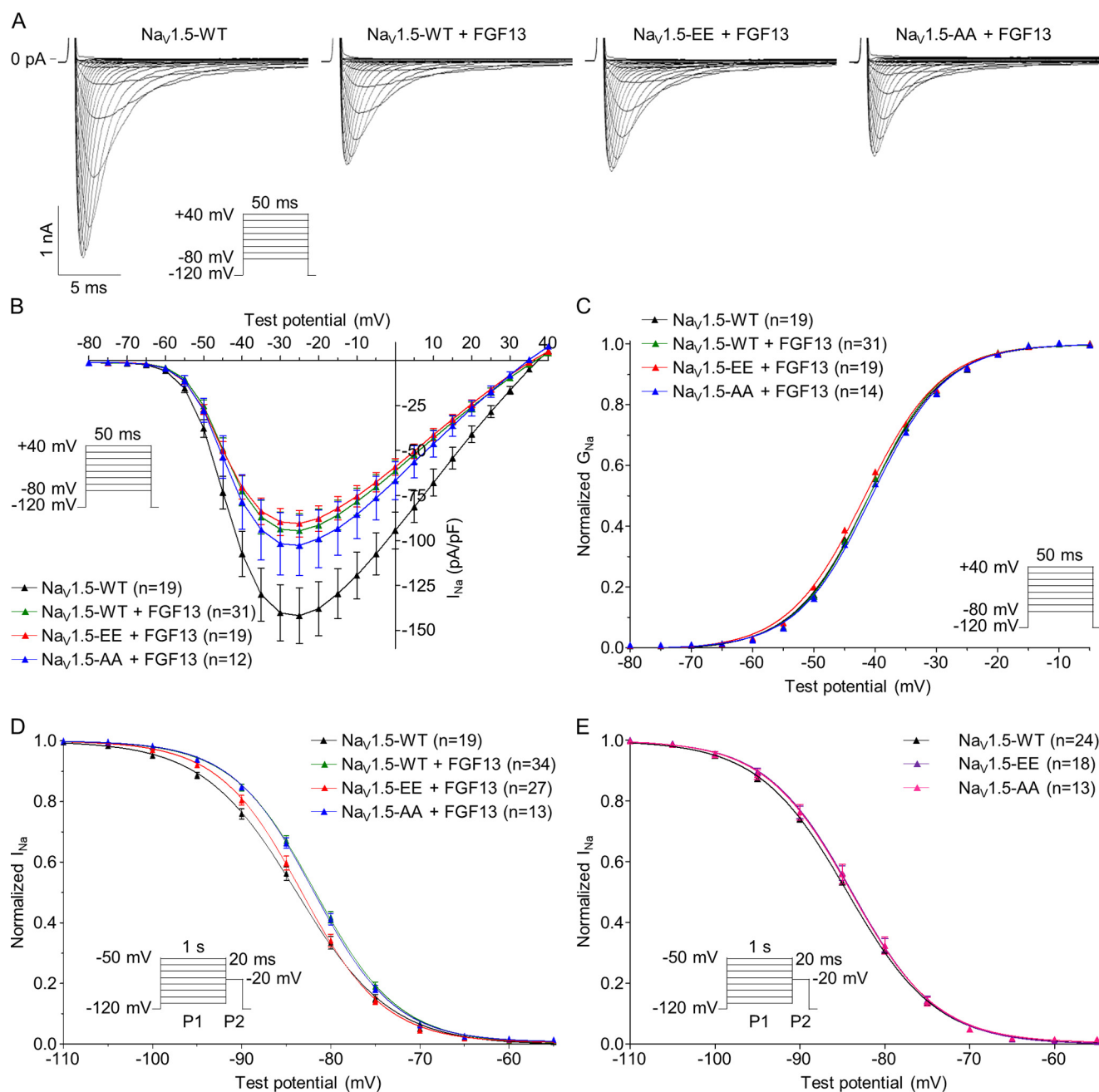


Figure 5. Phosphorylation at serines 1933 and 1984 disrupts the FGF13-dependent increase in steady-state $\text{Na}_v1.5$ channel availability. *A*, representative whole-cell voltage-gated Na^+ currents recorded from transiently transfected HEK293 cells. Currents were obtained 48 h following transfection of HEK293 cells with $\text{Na}_v1.5$ -WT (black), $\text{Na}_v1.5$ -WT + FGF13 (green), $\text{Na}_v1.5$ -S1933E/S1984E + FGF13 ($\text{Na}_v1.5$ -EE + FGF13, red), $\text{Na}_v1.5$ -S1933A/S1984A + FGF13 ($\text{Na}_v1.5$ -AA + FGF13, blue), $\text{Na}_v1.5$ -S1933E/S1984E ($\text{Na}_v1.5$ -EE, purple), and $\text{Na}_v1.5$ -S1933A/S1984A ($\text{Na}_v1.5$ -AA, pink) using the protocols illustrated in each panel. *B*, mean \pm S.E. peak Na^+ current (I_{Na}) densities are plotted as a function of test potential. *C*, voltage dependence of current activation. Mean \pm S.E. normalized conductances (G_{Na}) are plotted as a function of test potential and fitted using a Boltzmann equation. *D*, voltage dependence of steady-state current inactivation. Mean \pm S.E. normalized current amplitudes are plotted as a function of prepulse potential and fitted using a Boltzmann equation. FGF13 significantly ($p < 0.01$ versus $\text{Na}_v1.5$ -WT, one-way ANOVA followed by the Dunnett's post hoc test) shifts the voltage dependence of $\text{Na}_v1.5$ channel inactivation toward depolarized potentials, an effect reversed with the $\text{Na}_v1.5$ -EE phosphomutant ($p < 0.01$ versus $\text{Na}_v1.5$ -WT + FGF13, one-way ANOVA followed by the Dunnett's post hoc test). *E*, no significant changes in voltage dependence of steady-state current inactivation were observed between $\text{Na}_v1.5$ -WT, $\text{Na}_v1.5$ -EE, and $\text{Na}_v1.5$ -AA in the absence of FGF13. Detailed densities, properties, and statistics are provided in Table 2.

serines 1933 and 1984 on these FGF13-dependent regulations, these analyses were repeated from cells co-expressing $\text{Na}_v1.5$ -EE or $\text{Na}_v1.5$ -AA and FGF13. These recordings revealed that the voltage dependence of steady-state inactivation in cells co-expressing $\text{Na}_v1.5$ -EE and FGF13 is significantly ($p < 0.01$) shifted toward hyperpolarized potentials compared with cells co-expressing $\text{Na}_v1.5$ -WT and FGF13 and similar to cells

expressing $\text{Na}_v1.5$ -WT alone (Fig. 5*D* and Table 2). In contrast, the $\text{Na}_v1.5$ -AA phosphomutant co-expressed with FGF13 showed voltage dependence of inactivation properties similar to those recorded from cells co-expressing $\text{Na}_v1.5$ -WT and FGF13. No changes in peak I_{Na} density, voltage dependence of activation, or kinetics of activation, inactivation, and recovery from inactivation were observed with either the $\text{Na}_v1.5$ -EE or

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Table 2

Voltage-gated Na⁺ current densities and properties in transiently transfected HEK293 cells

The peak Na⁺ current density (I_{Na}), time to peak, and time course of inactivation properties presented were determined from analyses of records obtained on depolarizations to -20 mV (HP = -120 mV). All values are means ± S.E. The number of cells analyzed is provided in parentheses. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$ versus Na_v1.5-WT; #, $p < 0.01$ versus Na_v1.5-WT + FGF13; one-way ANOVA followed by the Dunnett's post-hoc test.

	Na _v 1.5-WT	Na _v 1.5-WT + FGF13	Na _v 1.5-EE + FGF13	Na _v 1.5-AA + FGF13	Na _v 1.5-S1933E + FGF13	Na _v 1.5-S1933A + FGF13	Na _v 1.5-S1984E + FGF13	Na _v 1.5-S1984A + FGF13	
I_{Na} (pA/pF)	-137.9 ± 14.9 (19)	-91.5 ± 8.9 (31)*	-87.8 ± 7.1 (19)*	-99.0 ± 15.9 (12)	-107.8 ± 10.1 (26)	-97.6 ± 17.1 (6)*	-97.6 ± 8.8 (16)	-83.9 ± 10.8 (8)*	
Time to peak (ms)	0.76 ± 0.02 (19)	0.82 ± 0.02 (31)	0.79 ± 0.02 (19)	0.78 ± 0.03 (14)	0.77 ± 0.01 (26)	0.72 ± 0.02 (6)	0.76 ± 0.02 (16)	0.74 ± 0.02 (8)	
Time course of inactivation	τ_{fast} (ms)	0.81 ± 0.03 (19)	0.84 ± 0.02 (31)	0.88 ± 0.02 (19)	0.85 ± 0.03 (14)	0.86 ± 0.02 (26)	0.73 ± 0.01 (6)	0.86 ± 0.02 (16)	0.79 ± 0.02 (8)
	τ_{slow} (ms)	5.3 ± 0.3 (19)	5.1 ± 0.3 (31)	6.1 ± 0.3 (19)	4.1 ± 0.3 (14)	5.1 ± 0.3 (26)	4.0 ± 0.4 (6)	5.9 ± 0.6 (16)	4.7 ± 0.3 (8)
	A_{fast}/A_{slow}	17.3 ± 1.7 (19)	15.7 ± 1.7 (31)	14.4 ± 2.2 (19)	15.4 ± 1.2 (14)	21.4 ± 2.9 (26)	18.3 ± 5.5 (6)	20.5 ± 4.9 (16)	18.3 ± 4.8 (8)
Voltage-dependence of activation	$V_{1/2}$ (mV)	-41.1 ± 0.4 (19)	-41.0 ± 0.4 (31)	-41.7 ± 0.4 (19)	-40.6 ± 0.7 (14)	-39.9 ± 0.3 (26)	-41.9 ± 0.5 (6)	-40.7 ± 0.5 (16)	-40.9 ± 0.6 (8)
	k (mV)	6.1 ± 0.1 (19)	5.9 ± 0.1 (31)	6.2 ± 0.1 (19)	5.9 ± 0.1 (14)	6.0 ± 0.1 (26)	5.9 ± 0.2 (6)	6.1 ± 0.1 (16)	5.9 ± 0.2 (8)
Voltage-dependence of inactivation	$V_{1/2}$ (mV)	-83.8 ± 0.5 (19)	-81.8 ± 0.4 (34)**	-83.3 ± 0.4 (27)##	-81.9 ± 0.4 (13)*	-81.0 ± 0.3 (24)***	-83.3 ± 0.6 (10)	-82.9 ± 0.4 (15)	-82.0 ± 0.3 (10)
	k (mV)	5.1 ± 0.11 (19)	4.5 ± 0.04 (34)***	4.4 ± 0.04 (27)***	4.5 ± 0.06 (13)***	4.5 ± 0.05 (24)***	4.5 ± 0.08 (10)***	4.5 ± 0.05 (15)***	4.5 ± 0.07 (10)***
Recovery from inactivation	τ_{rec} (ms)	6.9 ± 0.5 (13)	6.0 ± 0.3 (26)	6.8 ± 0.3 (18)	5.4 ± 0.3 (12)	5.3 ± 0.2 (21)	6.1 ± 0.4 (8)	6.3 ± 0.3 (11)	5.9 ± 0.4 (10)

Na_v1.5-AA phosphomutants compared with the Na_v1.5-WT co-expressed with FGF13 (Fig. 5, A–C, and Table 2). In addition, no changes in the voltage dependences of inactivation were observed between Na_v1.5-WT ($V_{1/2} = -84.5 \pm 0.6$ mV, $n = 24$), Na_v1.5-EE ($V_{1/2} = -83.9 \pm 0.5$ mV, $n = 18$), and Na_v1.5-AA ($V_{1/2} = -84.1 \pm 0.7$ mV, $n = 13$) in the absence of FGF13 (Fig. 5E), suggesting that FGF13 plays a pivotal role in mediating this phosphorylation-dependent effect. Finally, mimicking or abolishing phosphorylation at only one of the two phosphorylation sites (in cells co-expressing Na_v1.5-S1933E, Na_v1.5-S1933A, Na_v1.5-S1984E, or Na_v1.5-S1984A with FGF13) did not show any significant changes on peak I_{Na} density or channel biophysical properties compared with cells co-expressing Na_v1.5-WT and FGF13 (Table 2).

Additional voltage-clamp experiments were designed to examine the effects of FGF13 and of Na_v1.5 phosphomutants on the late Na⁺ current. These analyses showed that the TTX-sensitive late Na⁺ current (I_{NaL}) density is significantly ($p < 0.001$) smaller in cells co-expressing Na_v1.5 and FGF13 than in cells expressing Na_v1.5 alone (Fig. 6, A and B, and Table 3). Interestingly, this effect was significantly ($p < 0.05$) abrogated in cells co-expressing Na_v1.5-EE and FGF13 (Fig. 6, A and B, and Table 3). However, no such abrogation of the FGF13 effect was observed with the Na_v1.5-AA phosphomutant (Fig. 6, C and D, and Table 3). In addition, similar to the biophysical properties, no significant effects on I_{NaL} could be detected in the absence of FGF13 ($I_{NaL} = -0.32 \pm 0.04$ pA/pF, $n = 27$ for Na_v1.5-WT; $I_{NaL} = -0.46 \pm 0.07$ pA/pF, $n = 28$ for Na_v1.5-EE; and $I_{NaL} = -0.25 \pm 0.05$ pA/pF, $n = 15$ for Na_v1.5-AA, see Fig. 6E) or with the simple phosphomutants (Table 3). Taken together, these results suggest that simultaneous phosphorylation at serines 1933 and 1984 on Na_v1.5 impairs the effects of

FGF13 on channel inactivation properties, which results in decreased channel availability and increased I_{NaL} .

Phosphorylation at serines 1933 and 1984 decreases the interaction of FGF13 and CaM with Na_v1.5

To explore the hypothesis that phosphorylation at serines 1933 and 1984 in Na_v1.5 impairs the regulation of channel inactivation mediated by FGF13 by altering the binding of FGF13 and/or CaM to the channel, co-immunoprecipitation experiments were completed using the same experimental paradigm as above. As illustrated in Fig. 7A, FGF13 and (endogenous) CaM co-immunoprecipitate with Na_v1.5, whether in the WT, Na_v1.5-AA, or Na_v1.5-EE forms, in HEK293 cells. Interestingly, however, the relative abundances of FGF13 (Fig. 7B) and CaM (Fig. 7C) are significantly ($p < 0.01$ and $p < 0.001$, respectively) lower in the Na_v1.5 immunoprecipitates from cells expressing the Na_v1.5-EE phosphomutant compared with cells expressing the WT channels. No changes were observed with the Na_v1.5-AA phosphomutant. Parallel negative controls of co-immunoprecipitations obtained from cells transiently transfected with FGF13 or Na_v1.5-WT alone confirmed the specificity of detected signals. To determine whether phosphorylation at only one site was sufficient to decrease the interaction of FGF13 or CaM with the channel, co-immunoprecipitations were performed with the single alanine or glutamate phosphomutant channels. Consistent with the electrophysiological findings, no significant changes in binding affinity were observed with any of the single phosphomutant channels (Fig. 7, B and C).

Additional co-immunoprecipitation experiments were then designed to determine whether FGF13 and CaM influence each other for the binding to the channel. Interestingly, these experiments revealed that co-expression of FGF13 significantly ($p <$

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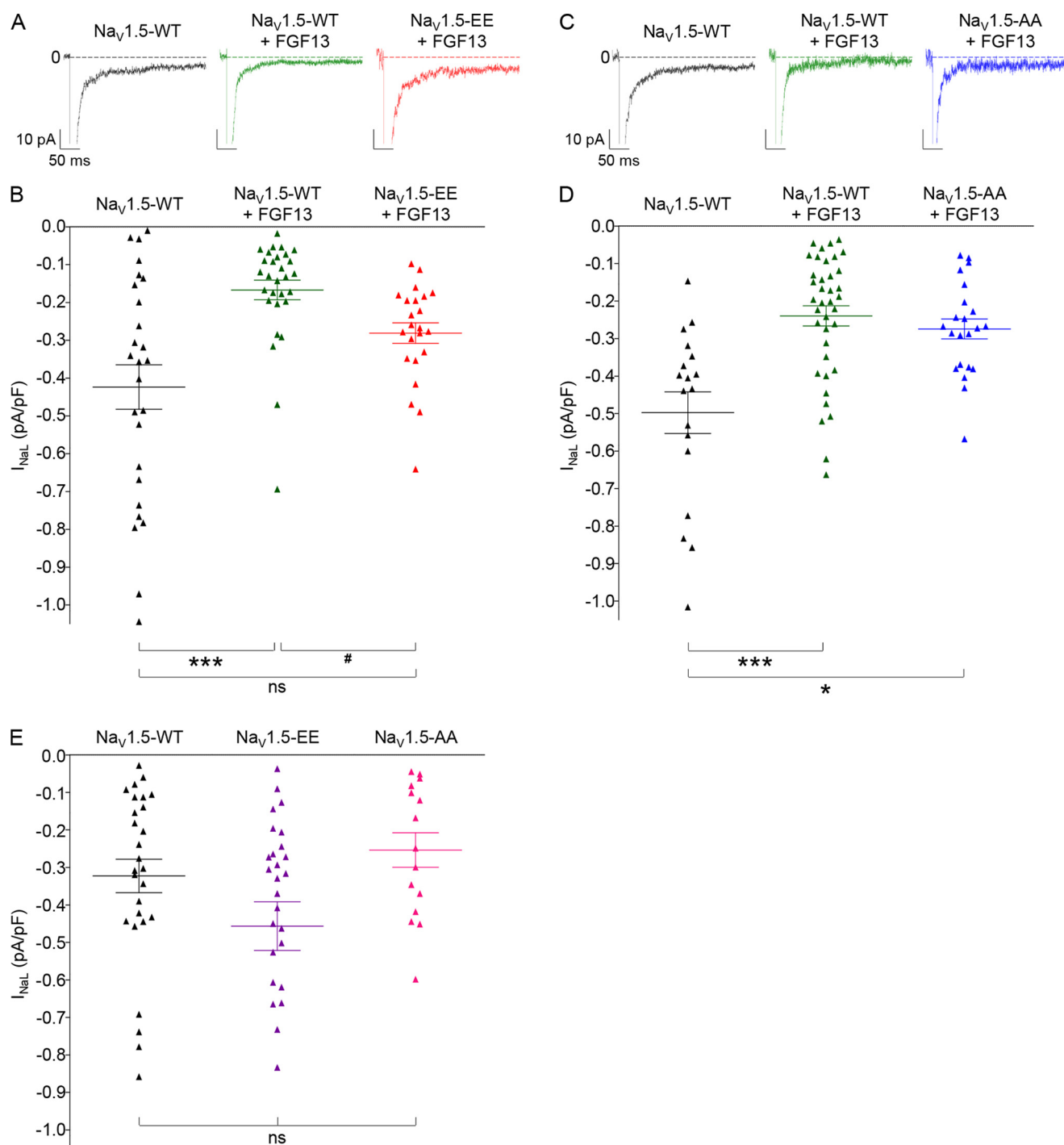


Figure 6. Phosphorylation at serines 1933 and 1984 disrupts the FGF13-dependent decrease in I_{NaL} . TTX-sensitive late Na^+ current (I_{NaL}) recordings were obtained 48 h after transfection of HEK293 cells in three different data sets. The first data set (A and B) was obtained from cells expressing $\text{Na}_v1.5\text{-WT}$ (black), $\text{Na}_v1.5\text{-WT} + \text{FGF13}$ (green), and $\text{Na}_v1.5\text{-S1933E/1984E} + \text{FGF13}$ ($\text{Na}_v1.5\text{-EE} + \text{FGF13}$, red); the second data set (C and D) from cells expressing $\text{Na}_v1.5\text{-WT}$ (black), $\text{Na}_v1.5\text{-WT} + \text{FGF13}$ (green), and $\text{Na}_v1.5\text{-S1933A/S1984A} + \text{FGF13}$ ($\text{Na}_v1.5\text{-AA} + \text{FGF13}$, blue); and the third data set (E) from cells expressing $\text{Na}_v1.5\text{-WT}$ (black), $\text{Na}_v1.5\text{-S1933E/S1984E}$ ($\text{Na}_v1.5\text{-EE}$, purple), and $\text{Na}_v1.5\text{-S1933A/S1984A}$ ($\text{Na}_v1.5\text{-AA}$, pink). A and C, representative TTX-sensitive I_{NaL} recordings evoked by prolonged depolarizations (350 ms at -20 mV) from a holding potential of -120 mV. The scale bars indicate a current amplitude of 10 pA, which corresponds to a current density of 0.8 pA/pF and time (50 ms). B, D, and E, distributions and mean \pm S.E. TTX-sensitive late Na^+ current (I_{NaL}) densities. *, $p < 0.05$; ***, $p < 0.001$ versus $\text{Na}_v1.5\text{-WT}$; #, $p < 0.05$ versus $\text{Na}_v1.5\text{-WT} + \text{FGF13}$; ns, non-significant; Kruskal-Wallis one-way ANOVA followed by the Dunn's post hoc test. Detailed densities and statistics are provided in Table 3.

0.01) increases the binding of CaM to $\text{Na}_v1.5$ (Fig. 7, D and E). Conversely, overexpression of CaM did not influence the binding of FGF13 to the channel (Fig. 7, D and E). Additionally, no changes in the binding of CaM to $\text{Na}_v1.5\text{-EE}$ or $\text{Na}_v1.5\text{-AA}$, compared with $\text{Na}_v1.5\text{-WT}$, were observed in the absence of

FGF13 (Fig. 7F). Together, these biochemical analyses suggest that FGF13 potentiates the binding of CaM to $\text{Na}_v1.5$ and that phosphorylation at both serines 1933 and 1984 decreases the interaction of FGF13 and, consequently, of CaM with the channel.

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Table 3

Late Na⁺ current densities in transiently transfected HEK293 cells

The TTX-sensitive late Na⁺ current (*I*_{NaL}) densities were measured at -20 mV (HP = -120 mV). All values are means ± S.E. The number of cells analyzed is provided in parentheses. *, *p* < 0.05; ***, *p* < 0.001 versus Na_v1.5-WT; #, *p* < 0.05 versus Na_v1.5-WT + FGF13; Kruskal-Wallis one-way ANOVA followed by the Dunn's post-hoc test.

	Na _v 1.5-WT	Na _v 1.5-WT + FGF13	Na _v 1.5-EE + FGF13	Na _v 1.5-S1933E + FGF13	Na _v 1.5-S1984E + FGF13
<i>I</i> _{NaL} (pA/pF)	-0.42 ± 0.06 (26)	-0.17 ± 0.03 (29)***	-0.28 ± 0.03 (23)#	-0.26 ± 0.04 (20)	-0.25 ± 0.03 (21)
	Na _v 1.5-WT	Na _v 1.5-WT + FGF13	Na _v 1.5-AA + FGF13	Na _v 1.5-S1933A + FGF13	Na _v 1.5-S1984A + FGF13
<i>I</i> _{NaL} (pA/pF)	-0.50 ± 0.06 (18)	-0.24 ± 0.03 (38)***	-0.27 ± 0.03 (22)*	-0.32 ± 0.06 (13)	-0.39 ± 0.06 (13)

Discussion

The results presented here provide two novel phosphorylation maps of native mouse Na_v1.5 channel subunits purified from control WT and failing CaMKIIδ_c-overexpressing ventricles, and they delineate two novel C-terminal phosphoserines, at positions 1938 and 1989, that are up-regulated in the failing CaMKIIδ_c-overexpressing ventricles. Mechanistic analyses in HEK293 cells revealed that mimicking phosphorylation at both sites (in the human sequence) impairs the regulation of Na_v1.5 channels by FGF13, resulting in decreased channel availability and increased late Na⁺ current. Co-immunoprecipitation experiments demonstrated that FGF13 potentiates the binding of CaM to Na_v1.5 and that FGF13 and, consequently, CaM bindings are decreased when phosphorylation at both sites is mimicked. Overall, these results provide evidence for a novel phosphorylation-dependent mechanism that acts at the level of the Na_v1.5 channel macromolecular complex through regulation of specific protein/protein interactions.

CaMKIIδ_c-dependent phosphorylation map of native mouse Na_v1.5 channels

The present phosphoproteomic analysis confidently identified a total of 19 native phosphorylation sites in the Na_v1.5 channel proteins purified from mouse ventricles, of which nine are novel. Consistent with our previous MS analysis (31), and with another phosphoproteomic analysis of human Na_v1.5 channels purified from HEK293 cells (9), the great majority (13 of 19) of identified phosphorylation sites are located in the first intracellular linker loop of the channel, suggesting critical roles for this region in mediating phosphorylation-dependent regulation of cardiac Na_v1.5 channels. However, none of these 13 phosphorylation sites identified in the first linker loop, including phosphoserine 571 which was reported to be CaMKIIδ_c-dependent (11, 12), appeared to be regulated in the CaMKIIδ_c-Tg ventricles compared with the WT ventricles. Note that no relative quantification could be obtained for the low abundance phosphopeptides assigning phosphoserines 516 and 1888, and that the region surrounding threonine 594 was not covered (Fig. 2B), precluding possible detection and quantification of phosphothreonine 594. The only two/three phosphorylation sites showing significant and consistent abundance changes in the CaMKIIδ_c-Tg compared with the WT ventricles are located

in the CTD of Na_v1.5, at position(s) 1937 and/or 1938 and at position 1989. Although localization(s) of phosphorylation could not be discriminated between serines 1937 and 1938 in the singly phosphorylated peptide allowing quantification (presence in the CaMKIIδ_c-Tg IPs, and absence in the WT IPs), we focused our attention on serine 1938 (human serine 1933) because serine 1937 is not well-conserved across species and is notably absent in human. In addition, the human Na_v1.5 phosphoserine 1933 has previously been shown to be CaMKIIδ_c-dependent in *in vitro* phosphoproteomic analyses (9), underscoring the potential involvement of CaMKIIδ_c in phosphorylating this site. Finally, it is also important to note here that the relative quantification of the doubly phosphorylated peptide supporting phosphorylation at both serines 1937 and 1938 (Table 1) could not be obtained because of low abundance, yet this peptide was only detected in the CaMKIIδ_c-Tg IPs. The number of phosphorylation sites identified here in association with the overexpression of CaMKIIδ_c in mouse ventricles (two phosphoserines) could seem little compared with the results of the previous phosphoproteomic study of human Na_v1.5 channels in which 23 phosphorylation sites, of the 34 identified, were found to be phosphorylated *in vitro* by CaMKIIδ_c. These seemingly distinct findings might reflect the fact that different experimental approaches were employed. Indeed, although phosphorylation sites were identified from native, *in situ* phosphorylated Na_v1.5 channels in this study, previous MS analysis was performed from heterologously-expressed Na_v1.5 channels that were immunoaffinity-purified and subsequently subjected to *in vitro* phosphorylation by recombinant CaMKIIδ_c. It is also stressed here that although the present MS analysis was performed from ventricles in which CaMKIIδ_c is overexpressed, we cannot exclude the involvement of other kinases/phosphatases whose activity may also be changed. Consistent with the direct implication of CaMKIIδ_c, however, the *in vitro* phosphoproteomic analyses of human Na_v1.5 demonstrated that human phosphoserine 1933 is CaMKIIδ_c-dependent (9). Likewise, it is interesting to note that mouse phosphoserine 1989 (and human phosphoserine 1984) is located in a well-conserved consensus CaMKII phosphorylation site (37). Together, these MS analyses from mouse ventricles therefore suggest that the two Na_v1.5 phosphoserines at positions 1938 and 1989 are associated with the overexpression of CaMKIIδ_c.

Phosphorylation and inactivation of cardiac Na_v1.5 channels

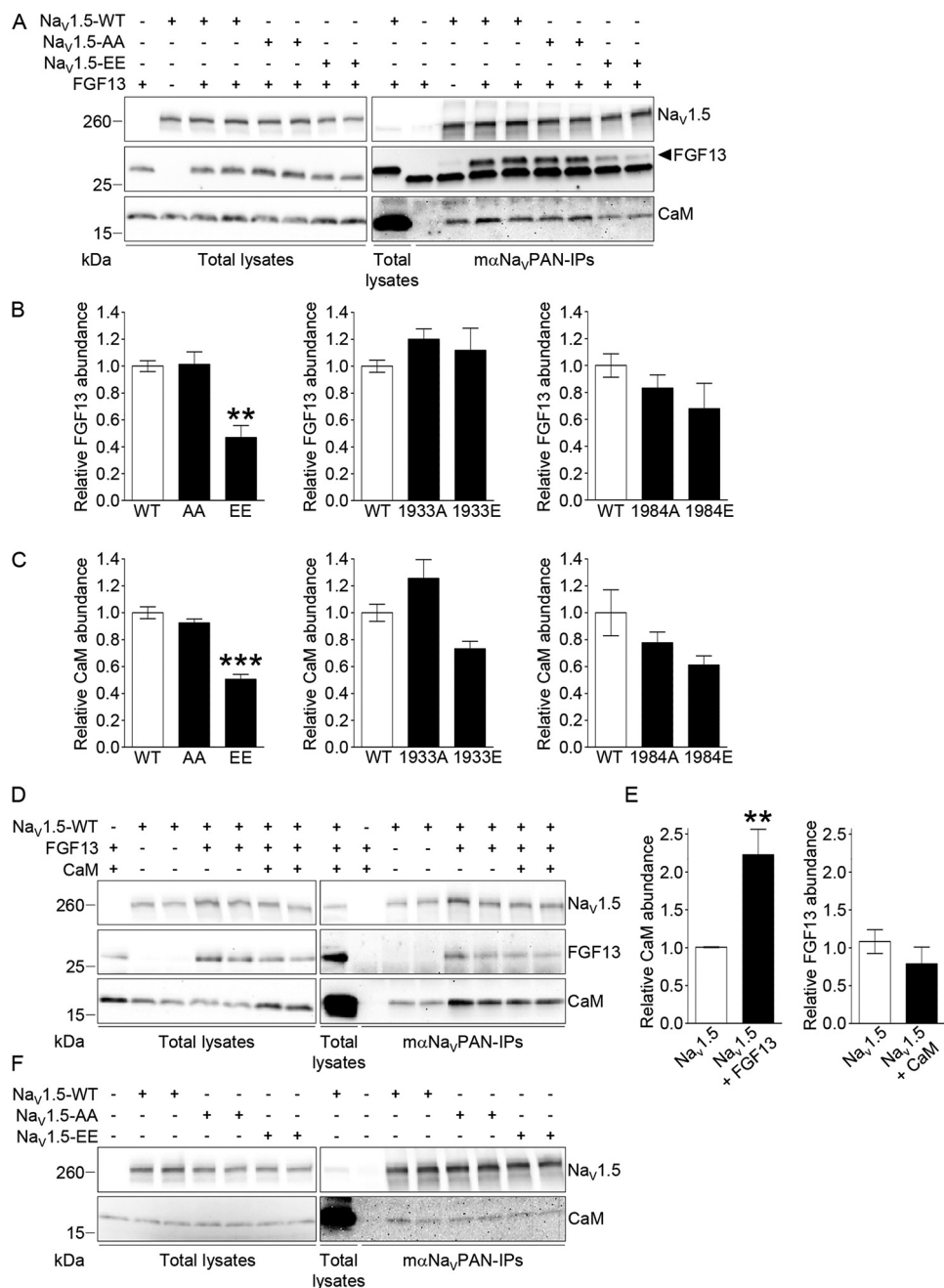


Figure 7. Phosphorylation at serines 1933 and 1984 decreases the interaction of FGF13 and consequently of CaM with Na_v1.5. Forty eight hours following transfection of HEK293 cells with Na_v1.5-WT, Na_v1.5-S1933A/S1984A (Na_v1.5-AA), Na_v1.5-S1933E/S1984E (Na_v1.5-EE), Na_v1.5-S1933A (1933A), Na_v1.5-S1933E (1933E), Na_v1.5-S1984A (1984A), Na_v1.5-S1984E (1984E), FGF13, and/or CaM, cell lysates were prepared and used for IPs with the α Na_vPAN antibody. *A*, *D*, and *F*, representative Western blots of the lysates (*left panel*) and the immunoprecipitates (*right panel*) with the monoclonal anti-Na_vPAN, anti-FGF13, and/or anti-CaM antibodies. Relative mean \pm S.E. FGF13 (*B*) and CaM (*C*) abundances in α Na_vPAN-IPs from cells expressing Na_v1.5-WT (WT, *n* = 8), Na_v1.5-AA (AA, *n* = 8), and Na_v1.5-EE (EE, *n* = 8); Na_v1.5-WT (WT, *n* = 5 and 4, respectively), 1933A (*n* = 6 and 4, respectively), and 1933E (*n* = 6 and 4, respectively); and Na_v1.5-WT (WT, *n* = 6), 1984A (*n* = 6), and 1984E (*n* = 6). *E*, relative mean \pm S.E. CaM abundances in α Na_vPAN-IPs from cells expressing Na_v1.5-WT (*n* = 8) and Na_v1.5-WT + FGF13 (*n* = 12), and FGF13 abundances in α Na_vPAN-IPs from cells expressing Na_v1.5-WT + FGF13 (*n* = 4) and Na_v1.5-WT + FGF13 + CaM (*n* = 4). **, *p* < 0.01; ***, *p* < 0.001 versus Na_v1.5-WT, Kruskal-Wallis one-way ANOVA followed by the Dunn's post hoc test (*B* and *C*), or Mann-Whitney test (*E*). FGF13 and CaM abundances in each IP were first normalized to immunoprecipitated Na_v1.5 and then expressed relative to FGF13 or CaM abundances in IPs from control conditions.

FGF13-dependent regulation of Na_v1.5 channel inactivation

Because our MS findings converged on the possible involvement of Na_v1.5 C-terminal phosphorylation sites in mediating CaMKII δ_c -dependent channel regulation and that the CTD of Na_v channels is well-recognized for its role in regulating channel inactivation through the binding of iFGFs (15, 20–26) and

CaM (13–19), we first characterized the roles of FGF13 in regulating Na_v1.5 channels in HEK293 cells. Our findings are in accordance with previous data demonstrating a key role for FGF13 in increasing steady-state Na_v1.5 channel availability, although the amplitude of the presently observed effect (\sim 2 mV) is smaller than previously reported effects of FGF13 or

Phosphorylation and inactivation of cardiac Na_v1.5 channels

other iFGFs (~5–10 mV) (15, 20–26). Interestingly, our data also provide a novel role for FGF13 in decreasing the late Na⁺ current in HEK293 cells. The role of FGF13, and in particular of FGF13-2, in regulating the late Na⁺ current had only been investigated once in tsA201 cells, and it was shown to have no effect (26). These different findings perhaps are due to differences in the heterologous expression systems used or are due to the fact that no Na_vβ subunits were included in the previous study compared with this study in which Na_v1.5 channels were co-expressed with FGF13–2 and Na_vβ1. Consistent with this negative effect of FGF13 on the late Na⁺ current, the conditional knock-out of FGF13 in murine hearts revealed longer action potential durations (25), which could well be caused by an increased late Na⁺ current. The answer to this question must await further analysis of the late Na⁺ current from this or other FGF13 animal models. The observed decrease in peak Na⁺ current density upon FGF13 co-expression was somewhat surprising in light of previous findings in native cardiomyocytes demonstrating a decreased Na_v1.5 channel cell-surface expression upon FGF13 knockdown (21) or a reduced peak Na⁺ current density in ventricular myocytes isolated from FGF13 conditional knock-out mice (25). This lack of consistency is nonetheless reminiscent of previous investigations of iFGF-dependent regulation of Na_v channels from Pitt and co-workers, who reported the same disparate effects on peak Na⁺ current density in native (increase) (21, 25) and heterologous (decrease) (26) cells, and it may reflect differences in channel cellular and molecular environment. In addition, note that no differences in peak Na⁺ current densities were observed in ventricular myocytes isolated from FGF13 knock-out and WT mice (24). In summary, these results suggest that FGF13 primarily regulates the inactivation properties of Na_v1.5 channels from both the closed state (increased availability) and the open state (decreased late Na⁺ current).

Impairment of FGF13-dependent Na_v1.5 channel inactivation by phosphorylation

Determination of the impact of phosphorylation at serines 1933 and 1984 in the regulation of Na_v1.5 channels first revealed that the binding of FGF13 and CaM to Na_v1.5 is reduced with the double-glutamate phosphomutant compared with the WT channels, suggesting that phosphorylation at these serines disables the interaction of FGF13 and CaM with the channel. Most importantly, this reduced interaction, as assessed by co-immunoprecipitation, correlates with an impairment of functional modulation. Indeed, all the effects of FGF13 on Na_v1.5-generated currents, but those on the peak density were nearly completely abolished with the double-glutamate phosphomutant, therefore resulting in decreased channel availability and increased late Na⁺ current. Together, these analyses show that mimicking phosphorylation at serines 1933 and 1984, which according to these and previous findings (9) are not phosphorylated at baseline in HEK293 cells, impairs FGF13-dependent regulation of Na_v1.5 channel inactivation. Of note, these functional effects associated with the decreased FGF13 and CaM bindings are similar to those previously observed with several Na_v1.5 mutants in which the binding of CaM to the Na_v1.5 CTD is disrupted (17, 19, 29). Mimicking

phosphorylation at only one site, however, did not lead to any consistent alterations in the interaction of FGF13 or CaM with the channel nor in the functional regulation by FGF13, suggesting that phosphorylation at both sites is necessary to disable FGF13/CaM bindings and associated dysregulation of channel inactivation. Consistent with these findings in HEK293 cells, our MS analyses also revealed different relative compositions of associated/regulatory proteins in Na_v1.5 channel complexes in the CaMKIIδ_c-Tg ventricles compared with the WT ventricles. Of particular interest, indeed, the FGF13 and CaM relative abundances in the CaMKIIδ_c-Tg IPs compared with the WT IP, are lower (2-fold abundance ratios) than the 3-fold abundance ratios obtained for Na_v1.5 or ankyrin-G. Together, therefore, these analyses suggest that phosphorylation at serines 1933 and 1984 impairs the binding of FGF13 and CaM to Na_v1.5 and abrogates FGF13-dependent regulation of channel inactivation.

FGF13 potentiates the binding of CaM to Na_v1.5, and phosphorylation-dependent disruption of FGF13 binding to Na_v1.5 results in reduced CaM binding

To further delineate the relationship between FGF13 and CaM for the binding to Na_v1.5 and subsequent phosphorylation-dependent regulation, the same electrophysiological and biochemical analyses were performed in the absence of FGF13. Interestingly, neither the channel availability nor the late Na⁺ current or the CaM/Na_v1.5 interaction were altered in the absence of FGF13, highlighting the primary role of FGF13 in this phosphorylation-dependent regulation. These findings also suggest that the decreased CaM binding to the double-glutamate phosphomutant channel is subsequent to the decreased FGF13 binding. In this respect, further co-immunoprecipitation experiments were performed to examine the relationship between FGF13 and CaM for the binding to Na_v1.5. Remarkably, these analyses revealed that FGF13 substantially increases the binding of CaM to the channel, demonstrating for the first time that the binding of CaM to Na_v1.5 is potentiated by the binding of FGF13. Inversely, no changes in FGF13 binding were observed upon CaM overexpression, suggesting that CaM does not influence the binding of FGF13. Whether phosphorylation or FGF13 influence directly the Ca²⁺/CaM dependence of Na_v1.5 channel inactivation remains to be investigated. Interestingly, these novel findings are evocative of recent observations from the Pitt group (29) demonstrating that the increased late Na⁺ current associated with several LQT3 syndrome mutations within the Na_v1.5 CTD correlates with a decreased binding of apoCaM to the Na_v1.5 CTD. In line with these findings, the recently reported LQT3-causing mutation at p.H1849R in the Na_v1.5 CTD is also associated with an impairment of iFGF binding and iFGF-mediated regulation of Na_v1.5 channels (28). Together with these two previous studies, our results herein support the overall hypothesis that the binding of FGF13 and/or CaM to the Na_v1.5 CTD is pivotal in the regulation of channel inactivation, as unveiled by C-terminal phosphorylation and disease mutations within the Na_v1.5 CTD, and that impairment of iFGF and/or CaM modulation may constitute a common mechanism to inherited (CTD-mutated LQT3 patients) and acquired (CaMKIIδ_c-associated) arrhythmias.

Potential broader implication of phosphorylation-dependent regulation of Na_v1.5 channels in heart failure

Evidence suggests that some of the arrhythmogenic abnormalities associated with heart failure are triggered by the activation of CaMKII (2, 30) and a subsequent CaMKII-dependent increase in I_{NaL} (4–7). By identifying two novel Na_v1.5 phosphorylation sites up-regulated in CaMKII δ_c -Tg ventricles, which are failing (30), and by demonstrating roles for these sites in altering Na_v1.5 channel inactivation, our results provide evidence for a model in which increased CaMKII δ_c -dependent Na_v1.5 phosphorylation at serines 1933 and 1984 may provide an arrhythmic substrate in heart failure. Nonetheless, it is unclear whether the functional consequences of impaired FGF13 and CaM binding to Na_v1.5 by phosphorylation, as observed here in HEK293 cells, fully match up with the Na_v channel defects associated with heart failure. Indeed, although the increased late Na⁺ current is a consistent finding in failing cardiomyocytes from both human and animal models (2, 4–7, 38), evidence for a decreased Na_v channel availability has only been observed, to our knowledge, in the CaMKII δ_c -Tg mice (7) and in a failing dog model induced by left-bundle branch ablation and right atrial pacing (2). Besides, the activation of CaMKII has consistently been associated with an increased I_{NaL} and a decreased Na_v1.5 channel availability (3, 7, 8, 11, 12). Overall, the molecular mechanisms elucidated here may be relevant in conditions of increased CaMKII δ_c expression and/or activity and may partly contribute to the arrhythmias associated with heart failure.

Experimental procedures

Animals were handled in accordance with the guidelines from Directive 2010/63/EU of the European Parliament on the protection of animals used for scientific purposes. Experimental protocols were approved by the local animal care and use committee (Comité d’Ethique pour l’Expérimentation Animale des Pays de la Loire, authorization CEEA.2010.9). Generation and characterization of the CaMKII δ_c -Tg mouse line have been described previously (30). Cardiac tissues for *in vitro* experiments were harvested after euthanasia of the mice by cervical dislocation.

Immunoprecipitations of Na_v channel complexes

Flash-frozen ventricles from four 13-month-old CaMKII δ_c -Tg and four age- and sex-matched WT mice were homogenized as described previously (31) in ice-cold lysis buffer containing 20 mM HEPES (pH 7.4), 150 mM NaCl, 0.5% amidosulfobetaine (Sigma), 1× complete protease inhibitor mixture tablet (Roche Applied Science), 1 mM phenylmethylsulfonyl fluoride (PMSF, Interchim), 0.7 μg/ml pepstatin A (Thermo Fisher Scientific), and 1× Halt phosphatase inhibitor mixture (Thermo Fisher Scientific). After a 15-min rotation at 4 °C, 8 mg of the pooled WT or CaMKII δ_c -Tg ventricular soluble protein fractions were pre-cleared with 200 μl of protein G-magnetic beads (Pierce) for 1 h and subsequently used for IPs with 48 μg of an anti-Na_vPAN mouse monoclonal antibody (mαNa_vPAN, Sigma, S8809), raised against the SP19 epitope (39) located in the third intracellular linker loop and common to all Na_v α subunits. Prior to the IP, antibodies were cross-linked to 200 μl of pro-

tein G-magnetic beads using 20 mM dimethyl pimelimidate (Thermo Fisher Scientific) (40). Protein samples and antibody-coupled beads were mixed for 2 h at 4 °C. Magnetic beads were then collected and washed rapidly four times with ice-cold lysis buffer, and isolated protein complexes were eluted from the beads in 2% Rapigest (41) (Waters), 8 M urea (Sigma), 100 mM Tris (pH 8.5) at 37 °C for 30 min.

For co-immunoprecipitations of heterologously expressed proteins, HEK293 cells were washed twice with PBS and lysed in lysis buffer (as above) 48 h after transfection. Soluble protein fractions were collected and incubated with 12.5 μl of mαNa_vPAN-coupled magnetic beads (as above). After a 2-h incubation at 4 °C, beads were washed four times in lysis buffer, and protein complexes were eluted with 1× SDS sample buffer at 60 °C for 5 min.

Gel electrophoreses and Western blot analyses

Ten percent of the immunoprecipitated mouse ventricular Na_v channel protein complexes were fractionated on one-dimensional polyacrylamide gels and analyzed using either SYPRO Ruby (Life Technologies, Inc.) staining or Western blotting using the mαNa_vPAN antibody (1:2000, Sigma, S8809) as described previously (31). The total ventricular lysates were blotted with a rabbit polyclonal anti-Na_v1.5 antibody (1:2000, RbαNa_v1.5, Alomone, ASC-005). Western blot analyses of protein eluates from co-immunoprecipitations of heterologously expressed proteins were performed using the following primary antibodies: mouse monoclonal anti-Na_vPAN (1:2000, mαNa_vPAN, Sigma, S8809); mouse monoclonal anti-FGF13 (1:300, clone N91/27, NeuroMab Facility, University of California Davis, NINDS/NIMH, National Institutes of Health); and rabbit monoclonal anti-CaM (1:300, Abcam, EP799Y). Bound antibodies were detected using horseradish peroxidase-conjugated goat anti-rabbit or anti-mouse secondary antibodies (Santa Cruz Biotechnology), and protein signals were visualized using the SuperSignal West Dura Extended Duration Substrate (Pierce). The intensities of co-immunoprecipitated FGF13 or CaM bands were normalized to the intensities of immunoprecipitated Na_v1.5 bands from the same IP sample, and FGF13 or CaM abundances in immunoprecipitations from experimental conditions are expressed relative to abundances in immunoprecipitations from control conditions.

In-solution endoprotease digestions

Samples for mass spectrometry were prepared as described previously (31). Briefly, eluted proteins were precipitated using the 2D protein clean-up kit (GE Healthcare). The resulting pellets were dissolved in 8 M urea, 100 mM Tris (pH 8.5), reduced with 5 mM tris(2-carboxyethyl)phosphine (pH 8.0) for 30 min at room temperature, and alkylated with 10 mM iodoacetamide (Bio-Rad) for 30 min at room temperature. Samples were then digested with 1 μg of endoproteinase Lys-C (Roche Applied Science) overnight at 37 °C and subsequently with 4 μg of trypsin (Sigma) overnight at 37 °C. Peptides were acidified with formic acid to a final concentration of 1%, extracted with NuTip porous graphite carbon wedge tips (Glygen), and eluted with aqueous acetonitrile (ACN, 60%) containing formic acid (FA, 0.1%). The extracted peptides were dried, dissolved in aqueous

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ACN/FA (1%/1%), stored at -80°C , and subsequently analyzed using one-dimensional liquid chromatography-tandem mass spectrometric experiments (LC-MS/MS).

Mass spectrometric analyses

Peptide mixtures were analyzed using nano-LC-MS on three high-resolution hybrid mass spectrometers, a linear quadrupole ion trap Orbitrap XL (LTQ-Orbitrap XL) (31), an LTQ-Orbitrap Elite (42) (both from Thermo Fisher Scientific), and a TripleTOF[®] 5600 Plus (SCIEX) (43). The chromatograph was a 2D Plus (Eksigent) LC with a Nanoflex module and AS2 autosampler in dual cHiPLC columns (ChromXP C₁₈ 200 $\mu\text{m} \times 15\text{ cm}$; particle size 3 μm , 120 \AA) configuration. The mobile phases were 1% FA in water (A) and 1% FA in ACN (B). The liquid chromatographs were interfaced to the mass spectrometers through a nanospray source (PicoView PV550, New Objective). The samples were loaded in a volume of 5–10 μl at a flow rate of 1.5 $\mu\text{l}/\text{min}$ followed by organic gradient elution of peptides (800 nl/min). The LC conditions for analyses on the LTQ-Orbitrap XL and LTQ-Orbitrap Elite were performed after equilibrating the columns in 98% solvent A and 2% solvent B followed by 2% B, 0–5 min; 2–25% B, 5–110 min; 25–80% B, 110–170 min; 80–2% B, 170–175 min; and isocratic elution at 2% B, 175–190 min. The survey scans (m/z 350–2000) (MS1) were acquired at high resolution (60,000 at m/z 400) in the Orbitrap XL, and the MS2 spectra were acquired in the linear ion trap at low resolution, both in profile mode. The maximum injection times for the MS1 scans in the Orbitrap and the LTQ were 500 and 200 ms, respectively. The automatic gain control targets for the Orbitrap and the LTQ were 5×10^5 and 3×10^4 , respectively, with maximum injection times of 100 ms for the MS2 scans. The MS1 scans were followed by three MS2 events in the linear ion trap with collision activation in the ion trap (parent threshold 1000, isolation width 2.0 Da, normalized collision energy 30%, activation Q 0.250, and activation time 30 ms). Dynamic exclusion was enabled ($-0.20/+1.0$ Da) for 90 s after MS2 acquisitions. A repeat count of 1, a repeat duration of 45 s, and a maximal exclusion list size of 500 were used. The following ion source parameters were used: capillary temperature 200°C , source voltage 3.0 kV, source current 100 μA , capillary voltage 33 V, and tube lens 120 V. Data were acquired using XCalibur, version 2.2 SP1 (Thermo Fisher Scientific). For data acquisition on the LTQ-Orbitrap Elite, the following modifications were employed. The survey scans ($m/z = 300$ –1650) were acquired at a resolution of 120,000 with a target value of $1\text{e}6$ ions. The HCD MS2 acquisitions were performed for the top 15 most intense ions at a resolution of 15,000, a target value of 40,000 and a low mass setting of 120 m/z . For collision-induced dissociation, a collision energy of 40% was used with an activation time of 100 ms. Dynamic exclusion was enabled (30 s) without MS2 acquisition of +1 charged parent ions. The data were acquired using Xcalibur, version 2.0.7 (Thermo Fisher Scientific).

The LC-MS analysis on the TripleTOF[®] 5600 Plus mass spectrometer was obtained using the following gradient elution program: 0 time, 98% A, 2% B; 5 min, 2% A, 98% B; 415 min, 65% A, 35% B; and 440 min, 20% A, 80% B. Initial chromatographic conditions were restored in 5 min and maintained for 20 min.

The ion spray voltage was set at 2.9 kV, the curtain gas at 10 p.s.i., the nebulizer gas at 14 p.s.i., and the interface heater temperature set at 175°C . The MS1 data were acquired at a resolution of 25,000 with a scan range of 400–1200 m/z in 250 ms. The top 50 product ions were selected for MS2 acquisition with a dwell time of 100 ms. Four time bins were summed for each scan at a frequency of 15.4 kHz (through monitoring of the 40 GHz multichannel TDC detector with four-anode/channel detection). A rolling collision energy (CE) was applied to all precursor ions for collision-induced dissociation using the following equation: $\text{CE} = \text{slope} \cdot m/z + \text{intercept}$, where the slope for all charge states above +2 is 0.0625 and the intercept is -3 , -5 , and -6 for +2, +3, and +4, respectively.

MS data processing and analyses

The LC-MS raw files from the LTQ-Orbitrap XL and Elite were processed using MASCOT Distiller (version 2.3.02, Matrix Science) with settings previously described (31). The TripleTOF data were processed using the SCIEX MS Data Converter (version 1.3, SCIEX), converting the raw data files (*.wiff) to mgf files. The resulting MS2 centroid files were used for database searching with MASCOT (version 2.3.02) against the UNIPROT mouse protein database (downloaded on May 2, 2011, with 72,510 entries for the XL data, and on December 20, 2011, with 77,200 entries for the Elite and the TripleTOF data) using the following parameters: trypsin as the enzyme, MS tolerances of 50 (XL), 10 (Elite), and 25 (TripleTOF) ppm; MS/MS tolerances of 0.8 (XL), 0.05 (Elite), and 0.1 (TripleTOF) Da, with a fixed carbamidomethylation of Cys residues and variable modifications being oxidation (Met), pyro-glutamination (Gln), and phosphorylation (Ser, Thr, and/or Tyr), a maximal number of missed cleavages of 4, and +1, +2, +3, and +4 charge states. Scaffold (versions 3.1.4.1 (XL), 4.4.3 (Elite) and 3.6.4 (TripleTOF), Proteome software) was used to validate MS2-based peptide and protein identifications with the Peptide (44) and Protein (45) Prophet algorithms using thresholds of 50 and 95%, respectively. The supplemental Tables 1–3 provide the complete lists of identified peptides and proteins using the LTQ-Orbitrap XL, Elite, and the TripleTOF mass spectrometers, respectively.

Phosphopeptide spectra were manually interpreted by comparing the observed mass values from the spectrum in XCalibur (LTQ-Orbitrap XL and Elite data) or PeakView (TripleTOF data) with the theoretical parent and fragment $\text{Na}_v1.5$ ion mass values from MS-Product. Annotations of MS2 spectra were first automated using java-based software, which matches observed m/z values with theoretical fragment masses from MS-Product, and definitive annotations were subsequently obtained by manual verification and interpretation. Mass accuracy tolerances of 20 ppm or 0.5 Da were used as guidelines for high- and low-resolution spectral annotations, respectively, and only those ions with mass errors within these ranges were included to determine residue coverage and location of phosphorylation site(s). Additionally, for spectral annotation of high-resolution MS2 data, charge states of observed parent and fragment ions were determined, and only precursor, b- and y-ions with confirmed charge states (*i.e.* with at least the presence of the ^{13}C isotope peak) were used. The phosphorylation

site assignments were based on the presence or absence of the unphosphorylated and phosphorylated b- and y-ions flanking the site(s) of phosphorylation, ions that we call site-discriminating ions throughout this study. When site-discriminating ions were not all detected, assignments of phosphorylation sites were narrowed down to one, two (for pSer-524 and/or pSer-525), or four (for pSer-36, pThr-38, pSer-39, and/or pSer-42) possibility(ies) by elimination. In addition to mass accuracy and charge state, the relative percentage of maximum intensity of site-discriminating ions or of any supporting ions in each analyzed spectrum was also considered for spectral annotation. Representative MS1 and MS2 spectra used for each of the phosphorylation site assignments, mass errors of parent ions (in ppm), and Mascot Ion scores for each phosphopeptide are presented in [supplemental Fig. 1](#). The definition of all observed site-discriminating ions, as well as the calculated mass errors and charge state confirmations for all supporting b- and y-ions (as well as for, when detected, the loss of phosphoric acid peaks) are summarized in [Table 1](#) and [supplemental Tables 4](#).

The label-free quantitative MS analysis of protein abundance in the IPs was performed from the entire (LTQ-Orbitrap XL) MS1 peptide data set using the DAnTE (Data Analysis Tool Extension) software (33–35). To perform label-free quantitative analyses of mass spectra of precursor ion (MS1 peptides) data, the LTQ-Orbitrap XL MS1 and MS2 data from quadruplicate analyses of α Na_vPAN-IPs from WT and CaMKII δ_c -Tg mouse ventricles were imported into Rosetta ElucidatorTM (version 3.3, Rosetta Biosoftware) for retention time and *m/z* alignment of the peptide ion chromatograms using previously described parameters (36). Peak intensities of MS1 peptide features were quantified, and normalization of intensities across samples was performed using the average signal intensities obtained in each sample. Ion chromatograms and isotopic distributions of aligned Na_v1.5 peptides were all visually inspected.

Plasmids

The simple and double Na_v1.5 phosphomutant constructs were generated by mutating the serines 1933 and/or 1984 to alanine(s) or glutamate(s) by site-directed mutagenesis of the pCI-Na_v1.5 plasmid containing the human Na_v1.5 hH1C cDNA (46) (NCBI Reference Sequence NM_000335) using the QuikChange II XL site-directed mutagenesis kit (Agilent Technologies) with the following primers (with mutations underlined): Na_v1.5-S1933A-F, 5'-cagcaggcgggcccggcctctccga, and Na_v1.5-S1933A-R, 5'-tcggagaggccggcggcccgcctgctg; Na_v1.5-S1933E-F, 5'-cgctcagcaggcggcgaaggcctctccgaagag, and Na_v1.5-S1933E-R 5'-ctcttcggagaggccttcggcggcctgctgac; Na_v1.5-S1984A-F, 5'-tgtcactagagccaccgcccataacctccaggtg, and Na_v1.5-S1984A-R 5'-cacctggaggttatcggcgggtgctctagtgac; Na_v1.5-S1984E-F 5'-agtgtcactagagccaccgaagataacctccaggtg, and Na_v1.5-S1984E-R 5'-ccgacactggaggttatctcgggtgctctagtgacact. The mutated constructs were then digested with restriction endonucleases to excise the mutated fragments, which were then subcloned into the original pCI-Na_v1.5 plasmid. The human Na_vβ1 (NM_001037, a gift from A. L. George) and rat CaM (NM_017326, a gift from I. Deschenes) cDNAs were subcloned

into pRc/CMV and pcDNA3, respectively. The human transcript variant 2 of FGF13 (FGF13–2, NM_001139500), subcloned into pCMV6-XL5, was purchased from Amsbio. All constructs were sequenced to ensure that no unintentional mutations were introduced.

Culture and transient transfections

HEK293 cells were maintained in Dulbecco's modified Eagle's medium (DMEM, Life Technologies, Inc.) supplemented with 10% fetal bovine serum, 100 units/ml penicillin, and 100 μg/ml streptomycin, in 37 °C, 5% CO₂, 95% air incubator. Cells were transiently transfected at 70–80% confluence in 35-mm dishes with 0.9 μg of the WT or phosphomutant Na_v1.5 plasmid, 0.45 μg of the Na_vβ1 plasmid, with or without 0.45 μg of the FGF13 plasmid using 2 μl of Lipofectamine 2000 (Life Technologies, Inc.) following the manufacturer's instructions. For patch-clamp recordings, transfections also contained 0.2 μg of the pEGFP plasmid (enhanced green fluorescent protein plasmid, Clontech), so that the enhanced GFP expression serves as a marker of transfection. The absolute amounts of the various constructs were calculated, and the empty pcDNA3.1 plasmid (Life Technologies, Inc.) was used as a filler plasmid to keep the total DNA constant at 2 μg in each transfection.

Electrophysiological recordings

Whole-cell Na_v currents were recorded at room temperature from transiently transfected HEK293 cells using an Axopatch 200A amplifier (Axon Instruments) 48 h after transfection. Voltage-clamp protocols were applied using the pClamp 10.2 software package (Axon Instruments) interfaced to the electrophysiological equipment using a Digidata 1440A digitizer (Axon Instruments). Current signals were filtered at 10 kHz prior to digitization at 50 kHz and storage. Patch-clamp pipettes were pulled from borosilicate glass (outer diameter is 1.5 mm and inner diameter is 0.86 mm, Sutter Instrument) using a P-97 micropipette puller (Sutter Instrument), coated with wax, and fire-polished to a resistance between 1.5 and 2.5 megohms when filled with internal solution. The internal solution contained (in mM): NaCl 5, CsCl 105, HEPES 10, glucose 5, EGTA 10, CaCl₂ 8.7 (1 μM free [Ca²⁺], calculated with MaxChelator) and Mg-ATP 5 (1 mM free [Mg²⁺], MaxChelator) (pH 7.2 with CsOH, ~300 mosM). The external (low Na⁺) solution contained (in mM): NaCl 25, CsCl 94, tetraethylammonium chloride 25, HEPES 10, glucose 5, CaCl₂ 1, MgCl₂ 2 (pH 7.4 with CsOH, ~300 mosM). All the chemicals were purchased from Sigma. After establishing the whole-cell configuration, 3 min were allowed to ensure stabilization of the voltage dependence of inactivation properties, at which time 25-ms voltage steps to ±10 mV from a holding potential (HP) of –70 mV were applied to allow measurement of whole-cell membrane capacitance, input, and series resistances. Only cells with access resistance <7 megohms were used, and input resistances were typically >5 giga-ohms. After compensation of series resistance (80%), the membrane was held at a HP of –120 mV, and the voltage-clamp protocols were carried out as indicated below. Leak currents were always <200 pA at the HP (–120 mV) and corrected

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off line. Cells exhibiting peak current amplitudes <500 or >5000 pA were excluded from analyses.

Data were compiled and analyzed using ClampFit 10.2 (Axon Instruments), Microsoft Excel, and Prism (GraphPad Software). Whole-cell membrane capacitance (C_m) was determined by analyzing the decays of capacitive transients elicited by the 25-ms voltage steps to ± 10 mV from the HP (−70 mV) prior to compensation. C_m was calculated by dividing the integrated capacitive transients by the voltage. Input resistance was calculated from the steady-state currents elicited by the same ±10-mV steps (from the HP). Series resistance was calculated by dividing the decay time constants of the capacitive transients (fitted with a single exponential) by the C_m. To determine the current-voltage relationships, currents were elicited by 50-ms depolarizing pulses from −80 to +40 mV in 5-mV increments from a HP of −120 mV (5-s inter-pulse duration). Peak current amplitudes at each voltage step were defined as the maximal current amplitudes. Current amplitudes were corrected by leak currents (calculated at each voltage step from the leak current measured at the HP) and normalized to the C_m. To analyze the voltage dependence of activation properties, current amplitudes at each voltage step were transformed to conductances (G), and conductance-voltage relationships were fitted with the Boltzmann equation $G = G_{\max}/(1 + \exp(-(V_m - V_{1/2})/k))$, in which V_{1/2} is the membrane potential for half-activation, and k is the slope factor. The time course of inactivation of macroscopic current was fitted with the double exponential function $I(t) = A_{\text{slow}} \times \exp(-t/\tau_{\text{slow}}) + A_{\text{fast}} \times \exp(-t/\tau_{\text{fast}}) + A_0$, where A_{slow} and A_{fast} are the amplitudes of the slow and fast inactivating current components, respectively, and τ_{slow} and τ_{fast} are the decay time constants of A_{slow} and A_{fast}, respectively. A standard two-pulse protocol was used to generate the voltage dependence of steady-state inactivation curves; from a HP of −120 mV, cells were stepped to 1-s conditioning potentials varying from −120 to −50 mV (prepulse) in 5-mV increments, followed by 20-ms test pulses to −20 mV (5-s interpulse duration). Current amplitudes measured at each test pulse were normalized to the maximal current amplitude (I_{max}), and the steady-state inactivation curves were fitted with the Boltzmann equation $I = I_{\max}/(1 + \exp((V_m - V_{1/2})/k))$, in which V_{1/2} is the membrane potential for half-inactivation, and k is the slope factor. Recovery from inactivation was assessed using a three-pulse protocol (5-s interpulse duration) with a test pulse (at −20 mV) at variable times (from 1 to 200 ms at −120 mV) after a 1-s conditioning pulse to −20 mV (from a HP of −120 mV). The time course of recovery from inactivation was analyzed by fitting the current amplitudes measured at each test pulse normalized to the current amplitudes measured during each conditioning pulse with the single exponential function $I(t) = A \times (1 - \exp(-t/\tau_{\text{rec}}))$.

In experiments aimed at recording the TTX-sensitive late Na⁺ current, cells were bathed in external (full Na⁺) solution containing (in mM) the following: NaCl 140, CsCl 5, HEPES 10, glucose 5, CaCl₂ 1, MgCl₂ 2 (pH 7.4 with CsOH, ~300 mosM). Repetitive 350-ms test pulses to −20 mV from a HP of −120 mV (5-s interpulse duration) were applied to cells superfused locally with external (full Na⁺) solutions supplemented with 20 mM mannitol, in the absence and in the presence of 30 μM TTX

(Tocris Bioscience). Only cells exhibiting peak current amplitudes of >4000 pA were used (those with peak current of <4000 pA did not show measurable late Na⁺ current), and cells with difference in leak current amplitudes before and after TTX application of >2 pA at −20 mV (calculated from leak currents at −120 mV) were excluded from analyses. TTX-sensitive currents from individual cells were determined by off-line digital subtraction of average leak-subtracted currents obtained from five recordings in the absence and in the presence of TTX after achieving steady state. The amplitude of TTX-sensitive late Na⁺ current was defined as the steady-state current amplitude (A₀) obtained by fitting the inactivation decay of macroscopic TTX-sensitive current with the double exponential function $I(t) = A_{\text{slow}} \times \exp(-t/\tau_{\text{slow}}) + A_{\text{fast}} \times \exp(-t/\tau_{\text{fast}}) + A_0$. For each cell, the TTX-sensitive late Na⁺ current amplitude was normalized to the C_m, and expressed as TTX-sensitive late Na⁺ current density (in pA/pF).

Statistical analyses

Results are expressed as means ± S.E. Data were first tested for normality using the D'Agostino and Pearson normality test. Depending on the results of normality tests, statistical analyses were then performed using the Mann-Whitney nonparametric test, the one-way ANOVA followed by the Dunnett's post hoc test, or the Kruskal-Wallis one-way ANOVA followed by the Dunn's post hoc test as indicated in the figures and tables. All of these analyses, the Tukey whisker analysis and the plots, were performed using Prism (GraphPad software).

Author contributions—C. M. designed the study and wrote the paper. C. M., M. R. M., C. F. L., J. M. N., R. R. T., and L. S. M. designed, performed, and/or analyzed the mass spectrometry experiments shown in Figs. 1–4 and Table 1. S. B., F. C. C., M. L., G. L., F. C., and C. M. designed, performed, and/or analyzed the experiments shown in Figs. 5–7 and Tables 2 and 3. J. H. B. and L. S. M. provided the CaMKIIδc-Tg mice. All authors reviewed the results and approved the final version of the manuscript.

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References

1. Remme, C. A., and Bezzina, C. R. (2010) Sodium channel (dys)function and cardiac arrhythmias. *Cardiovasc. Ther.* **28**, 287–294
2. Aiba, T., Barth, A. S., Hesketh, G. G., Hashambhoy, Y. L., Chakir, K., Tunin, R. S., Greenstein, J. L., Winslow, R. L., Kass, D. A., and Tomaselli, G. F. (2013) Cardiac resynchronization therapy improves altered Na channel gating in canine model of dyssynchronous heart failure. *Circ. Arrhythm. Electrophysiol.* **6**, 546–554
3. Dybkova, N., Wagner, S., Backs, J., Hund, T. J., Mohler, P. J., Sowa, T., Nikolaev, V. O., and Maier, L. S. (2014) Tubulin polymerization disrupts cardiac β-adrenergic regulation of late I_{Na}. *Cardiovasc. Res.* **103**, 168–177
4. Maltsev, V. A., Reznikov, V., Undrovinas, N. A., Sabbah, H. N., and Undrovinas, A. (2008) Modulation of late sodium current by Ca²⁺, calmodulin, and CaMKII in normal and failing dog cardiomyocytes: similarities and differences. *Am. J. Physiol. Heart Circ. Physiol.* **294**, H1597–H1608
5. Maltsev, V. A., and Undrovinas, A. (2008) Late sodium current in failing heart: friend or foe? *Prog. Biophys. Mol. Biol.* **96**, 421–451

6. Toischer, K., Hartmann, N., Wagner, S., Fischer, T. H., Herting, J., Danner, B. C., Sag, C. M., Hund, T. J., Mohler, P. J., Belardinelli, L., Hasenfuss, G., Maier, L. S., and Sossalla, S. (2013) Role of late sodium current as a potential arrhythmogenic mechanism in the progression of pressure-induced heart disease. *J. Mol. Cell. Cardiol.* **61**, 111–122
7. Wagner, S., Dybkova, N., Rasenack, E. C., Jacobshagen, C., Fabritz, L., Kirchhof, P., Maier, S. K., Zhang, T., Hasenfuss, G., Brown, J. H., Bers, D. M., and Maier, L. S. (2006) Ca²⁺/calmodulin-dependent protein kinase II regulates cardiac Na⁺ channels. *J. Clin. Invest.* **116**, 3127–3138
8. Ashpole, N. M., Herren, A. W., Ginsburg, K. S., Brogan, J. D., Johnson, D. E., Cummins, T. R., Bers, D. M., and Hudmon, A. (2012) Ca²⁺/calmodulin-dependent protein kinase II (CaMKII) regulates cardiac sodium channel NaV1.5 gating by multiple phosphorylation sites. *J. Biol. Chem.* **287**, 19856–19869
9. Herren, A. W., Weber, D. M., Rigor, R. R., Margulies, K. B., Phinney, B. S., and Bers, D. M. (2015) CaMKII phosphorylation of Na(V)1.5: novel *in vitro* sites identified by mass spectrometry and reduced S516 phosphorylation in human heart failure. *J. Proteome Res.* **14**, 2298–2311
10. Glynn, P., Musa, H., Wu, X., Unudurthi, S. D., Little, S., Qian, L., Wright, P. J., Radwanski, P. B., Gyorke, S., Mohler, P. J., and Hund, T. J. (2015) Voltage-gated sodium channel phosphorylation at Ser571 regulates late current, arrhythmia, and cardiac function *in vivo*. *Circulation* **132**, 567–577
11. Hund, T. J., Koval, O. M., Li, J., Wright, P. J., Qian, L., Snyder, J. S., Gudmundsson, H., Kline, C. F., Davidson, N. P., Cardona, N., Rasband, M. N., Anderson, M. E., and Mohler, P. J. (2010) A β (IV)-spectrin/CaMKII signaling complex is essential for membrane excitability in mice. *J. Clin. Invest.* **120**, 3508–3519
12. Koval, O. M., Snyder, J. S., Wolf, R. M., Pavlovicz, R. E., Glynn, P., Curran, J., Leymaster, N. D., Dun, W., Wright, P. J., Cardona, N., Qian, L., Mitchell, C. C., Boyden, P. A., Binkley, P. F., Li, C., *et al.* (2012) Ca²⁺/calmodulin-dependent protein kinase II-based regulation of voltage-gated Na⁺ channel in cardiac disease. *Circulation* **126**, 2084–2094
13. Chagot, B., and Chazin, W. J. (2011) Solution NMR structure of apocalmodulin in complex with the IQ motif of human cardiac sodium channel NaV1.5. *J. Mol. Biol.* **406**, 106–119
14. Gabelli, S. B., Boto, A., Kuhns, V. H., Bianchet, M. A., Farinelli, F., Aripirala, S., Yoder, J., Jakoncic, J., Tomaselli, G. F., and Amzel, L. M. (2014) Regulation of the NaV1.5 cytoplasmic domain by calmodulin. *Nat. Commun.* **5**, 5126
15. Wang, C., Chung, B. C., Yan, H., Lee, S. Y., and Pitt, G. S. (2012) Crystal structure of the ternary complex of a NaV C-terminal domain, a fibroblast growth factor homologous factor, and calmodulin. *Structure* **20**, 1167–1176
16. Wang, C., Chung, B. C., Yan, H., Wang, H. G., Lee, S. Y., and Pitt, G. S. (2014) Structural analyses of Ca²⁺/CaM interaction with NaV channel C-termini reveal mechanisms of calcium-dependent regulation. *Nat. Commun.* **5**, 4896
17. Kim, J., Ghosh, S., Liu, H., Tateyama, M., Kass, R. S., and Pitt, G. S. (2004) Calmodulin mediates Ca²⁺ sensitivity of sodium channels. *J. Biol. Chem.* **279**, 45004–45012
18. Sarhan, M. F., Tung, C. C., Van Petegem, F., and Ahern, C. A. (2012) Crystallographic basis for calcium regulation of sodium channels. *Proc. Natl. Acad. Sci. U.S.A.* **109**, 3558–3563
19. Shah, V. N., Wingo, T. L., Weiss, K. L., Williams, C. K., Balsler, J. R., and Chazin, W. J. (2006) Calcium-dependent regulation of the voltage-gated sodium channel hH1: intrinsic and extrinsic sensors use a common molecular switch. *Proc. Natl. Acad. Sci. U.S.A.* **103**, 3592–3597
20. Goetz, R., Dover, K., Laezza, F., Shtraizent, N., Huang, X., Tchetchik, D., Eliseenkova, A. V., Xu, C. F., Neubert, T. A., Ornitz, D. M., Goldfarb, M., and Mohammadi, M. (2009) Crystal structure of a fibroblast growth factor homologous factor (FHF) defines a conserved surface on FHFs for binding and modulation of voltage-gated sodium channels. *J. Biol. Chem.* **284**, 17883–17896
21. Wang, C., Hennessey, J. A., Kirkton, R. D., Wang, C., Graham, V., Purnam, R. S., Rosenberg, P. B., Bursac, N., and Pitt, G. S. (2011) Fibroblast growth factor homologous factor 13 regulates Na⁺ channels and conduction velocity in murine hearts. *Circ. Res.* **109**, 775–782
22. Wang, C., Wang, C., Hoch, E. G., and Pitt, G. S. (2011) Identification of novel interaction sites that determine specificity between fibroblast growth factor homologous factors and voltage-gated sodium channels. *J. Biol. Chem.* **286**, 24253–24263
23. Bosch, M. K., Carrasquillo, Y., Ransdell, J. L., Kanakamedala, A., Ornitz, D. M., and Nerbonne, J. M. (2015) Intracellular FGF14 (iFGF14) is required for spontaneous and evoked firing in cerebellar Purkinje neurons and for motor coordination and balance. *J. Neurosci.* **35**, 6752–6769
24. Park, D. S., Shekhar, A., Marra, C., Lin, X., Vasquez, C., Solinas, S., Kelley, K., Morley, G., Goldfarb, M., and Fishman, G. I. (2016) Fhf2 gene deletion causes temperature-sensitive cardiac conduction failure. *Nat. Commun.* **7**, 12966
25. Wang, X., Tang, H., Wei, E. Q., Wang, Z., Yang, J., Yang, R., Wang, S., Zhang, Y., Pitt, G. S., Zhang, H., and Wang, C. (2017) Conditional knock-out of Fgf13 in murine hearts increases arrhythmia susceptibility and reveals novel ion channel modulatory roles. *J. Mol. Cell. Cardiol.* **104**, 63–74
26. Yang, J., Wang, Z., Sinden, D. S., Wang, X., Shan, B., Yu, X., Zhang, H., Pitt, G. S., and Wang, C. (2016) FGF13 modulates the gating properties of the cardiac sodium channel Nav1.5 in an isoform-specific manner. *Channels* **10**, 410–420
27. Gabelli, S. B., Yoder, J. B., Tomaselli, G. F., and Amzel, L. M. (2016) Calmodulin and Ca²⁺ control of voltage gated Na⁺ channels. *Channels* **10**, 45–54
28. Musa, H., Kline, C. F., Sturm, A. C., Murphy, N., Adelman, S., Wang, C., Yan, H., Johnson, B. L., Csepe, T. A., Kilic, A., Higgins, R. S., Janssen, P. M., Fedorov, V. V., Weiss, R., Salazar, C., *et al.* (2015) SCN5A variant that blocks fibroblast growth factor homologous factor regulation causes human arrhythmia. *Proc. Natl. Acad. Sci. U.S.A.* **112**, 12528–12533
29. Yan, H., Wang, C., Marx, S. O., and Pitt, G. S. (2017) Calmodulin limits pathogenic Na⁺ channel persistent current. *J. Gen. Physiol.* **149**, 277–293
30. Zhang, T., Maier, L. S., Dalton, N. D., Miyamoto, S., Ross, J., Jr., Bers, D. M., and Brown, J. H. (2003) The δ C isoform of CaMKII is activated in cardiac hypertrophy and induces dilated cardiomyopathy and heart failure. *Circ. Res.* **92**, 912–919
31. Marionneau, C., Lichti, C. F., Lindenbaum, P., Charpentier, F., Nerbonne, J. M., Townsend, R. R., and Mèrot, J. (2012) Mass spectrometry-based identification of native cardiac Nav1.5 channel α subunit phosphorylation sites. *J. Proteome Res.* **11**, 5994–6007
32. Abriel, H., Rougier, J. S., and Jalife, J. (2015) Ion channel macromolecular complexes in cardiomyocytes: roles in sudden cardiac death. *Circ. Res.* **116**, 1971–1988
33. Karpievitch, Y., Stanley, J., Taverner, T., Huang, J., Adkins, J. N., Ansong, C., Heffron, F., Metz, T. O., Qian, W. J., Yoon, H., Smith, R. D., and Dabney, A. R. (2009) A statistical framework for protein quantitation in bottom-up MS-based proteomics. *Bioinformatics* **25**, 2028–2034
34. Polpitiya, A. D., Qian, W. J., Jaitly, N., Petyuk, V. A., Adkins, J. N., Camp, D. G., 2nd, Anderson, G. A., and Smith, R. D. (2008) DAnTE: a statistical tool for quantitative analysis of -omics data. *Bioinformatics* **24**, 1556–1558
35. Taverner, T., Karpievitch, Y. V., Polpitiya, A. D., Brown, J. N., Dabney, A. R., Anderson, G. A., and Smith, R. D. (2012) DanteR: an extensible R-based tool for quantitative analysis of -omics data. *Bioinformatics* **28**, 2404–2406
36. Neubert, H., Bonnert, T. P., Rumpel, K., Hunt, B. T., Henle, E. S., and James, I. T. (2008) Label-free detection of differential protein expression by LC/MALDI mass spectrometry. *J. Proteome Res.* **7**, 2270–2279
37. Songyang, Z., Lu, K. P., Kwon, Y. T., Tsai, L. H., Filhol, O., Cochet, C., Brickey, D. A., Soderling, T. R., Bartleson, C., Graves, D. J., DeMaggio, A. J., Hoekstra, M. F., Blenis, J., Hunter, T., and Cantley, L. C. (1996) A structural basis for substrate specificities of protein Ser/Thr kinases: primary sequence preference of casein kinases I and II, NIMA, phosphorylase kinase, calmodulin-dependent kinase II, CDK5, and Erk1. *Mol. Cell. Biol.* **16**, 6486–6493
38. Valdivia, C. R., Chu, W. W., Pu, J., Foell, J. D., Haworth, R. A., Wolff, M. R., Kamp, T. J., and Makielski, J. C. (2005) Increased late sodium current in myocytes from a canine heart failure model and from failing human heart. *J. Mol. Cell. Cardiol.* **38**, 475–483

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39. Vassilev, P. M., Scheuer, T., and Catterall, W. A. (1988) Identification of an intracellular peptide segment involved in sodium channel inactivation. *Science* **241**, 1658–1661
40. Schneider, C., Newman, R. A., Sutherland, D. R., Asser, U., and Greaves, M. F. (1982) A one-step purification of membrane proteins using a high efficiency immunomatrix. *J. Biol. Chem.* **257**, 10766–10769
41. Yu, Y. Q., Gilar, M., and Gebler, J. C. (2004) A complete peptide mapping of membrane proteins: a novel surfactant aiding the enzymatic digestion of bacteriorhodopsin. *Rapid Commun. Mass Spectrom.* **18**, 711–715
42. Michalski, A., Damoc, E., Lange, O., Denisov, E., Nolting, D., Muller, M., Viner, R., Schwartz, J., Remes, P., Belford, M., Duniach, J. J., Cox, J., Horning, S., Mann, M., and Makarov, A. (2012) Ultra high-resolution linear ion trap Orbitrap mass spectrometer (Orbitrap Elite) facilitates top down LC MS/MS and versatile peptide fragmentation modes. *Mol. Cell. Proteomics* **11**, O111-013698
43. Andrews, G. L., Simons, B. L., Young, J. B., Hawkrige, A. M., and Mudiman, D. C. (2011) Performance characteristics of a new hybrid quadrupole time-of-flight tandem mass spectrometer (TripleTOF 5600). *Anal. Chem.* **83**, 5442–5446
44. Keller, A., Nesvizhskii, A. I., Kolker, E., and Aebersold, R. (2002) Empirical statistical model to estimate the accuracy of peptide identifications made by MS/MS and database search. *Anal. Chem.* **74**, 5383–5392
45. Nesvizhskii, A. I., Keller, A., Kolker, E., and Aebersold, R. (2003) A statistical model for identifying proteins by tandem mass spectrometry. *Anal. Chem.* **75**, 4646–4658
46. Makielski, J. C., Ye, B., Valdivia, C. R., Pagel, M. D., Pu, J., Tester, D. J., and Ackerman, M. J. (2003) A ubiquitous splice variant and a common polymorphism affect heterologous expression of recombinant human SCN5A heart sodium channels. *Circ. Res.* **93**, 821–828