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Temperature-Activated Ion Channels in Neural Crest Cells Confer Maternal Fever-Associated Birth Defects

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Abstract

Birth defects of the heart and face are common and most have no known genetic cause, suggesting a role for environmental factors. Maternal fever during the first trimester is an environmental risk factor linked to these defects. Neural crest cells are precursor populations essential to the development of both at-risk tissues. Here, we report that two heat-activated transient receptor potential ion channels, TRPV1 and TRPV4, were present in neural crest cells during critical windows of heart and face development. TRPV1 antagonists protected against the development of

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Author Contributions: C.L. and E.J.B. conceived the FeRIC technology and designed and supervised the project. E.J.B. and K.P. designed and constructed all the channel constructs. C.L., I.A. and E.G. designed and constructed RF coils and RF delivery systems. C.L. and E.G. performed MRI and MRI thermometry studies. M.R.H., A.L.K., C.L., E.G., M.K. and E.J.B. designed, performed, and interpreted all avian craniofacial and heart studies. C.L. and L.X. performed cardiac MRI and image reconstructions. M.H.M., E.J.B., J.G., M.R.H., and P.B.R. designed, performed, and interpreted Ca²⁺ imaging studies. E.E.D., Z.A.K., K.E., M.R.H. and N.K. performed zebrafish studies. E.J.B. wrote the paper with input from C.L. M.R.H., A.L.K., and JG.

Data and materials availability: TRPV1^{FeRIC}, TRPV1^{WT}, TRPV4^{FeRIC}, TRPV4^{WT} and TRPV1^{FeRIC} plasmids will be made available upon request and will require a materials transfer agreement from E.J.B or C.L.

Competing Interests: C.L. and E.J.B. have filed a patent application (WO2016004281 A1 PCT/US2015/038948) relating to the use of FeRIC for cell modulation and treatments.

hyperthermia-induced defects in chick and zebrafish embryos. Conversely, treatment with chemical agonists of TRPV1 or TRPV4 replicated hyperthermia-induced birth defects in both model systems. To test whether transient TRPV channel permeability in neural crest cells was sufficient to induce these defects, we engineered iron-binding modifications to TRPV1 and TRPV4 that enabled remote and noninvasive activation of these channels in specific cellular locations and at specific developmental times in chick embryos with radiofrequency electromagnetic fields. Transient stimulation of radiofrequency-controlled TRP channels in neural crest cells replicated fever-associated defects in developing chick embryos. Our data provide a previously undescribed mechanism for congenital defects whereby hyperthermia activates ion channels that negatively affect fetal development.

Introduction

Birth defects of the heart and face are extremely common. Congenital heart defects affect about 1% of live births in the United States and often require surgical correction within the first year of life(1). Craniofacial clefts involving the lip and or the palate affect over 4,000 infants per year and typically require corrective surgery(2). Genetic associations have been identified in only 15% of heart or 30% of craniofacial defects leaving the majority of cases without a known etiology(3, 4). Environmental factors likely contribute. First trimester maternal fever is an environmental factor linked to both craniofacial and heart defects(5). Highly associated heart defects include right- and left-sided obstructive lesions (pulmonary atresia or pulmonary or aortic stenosis) and conotruncal defects (double outlet right ventricle (DORV), tetralogy of Fallot) (6–8). Hyperthermia-associated craniofacial defects include midface hypoplasia and cleft lip and or palate(5, 9). Specific modes of infections, such as respiratory compared to urinary or pelvic infections, confer differential susceptibilities leading some to speculate that infection itself is the critical event(8, 10). However, maternal hyperthermia following hot tub exposure is sufficient to confer risk of birth defects suggesting that hyperthermia itself is the principal teratogen(11). Nonetheless, the teratogenic mechanism(s) of maternal fever are unknown.

Craniofacial clefts, abnormal arch artery anatomy, aorticopulmonary septation and conotruncal heart defects can be linked through neural crest cell dysfunction(12, 13). Neural crest cells are a pluripotent migratory cell population that arises from the dorsal neural tube. Cranial and cardiac neural crest cells migrate to the head and pharyngeal arches where they contribute to the developing face and heart. The cranial neural crest cells differentiate into the bone and cartilage that form facial features including the jaw and palate(14). The cardiac neural crest cells are required for septation of the aorta and pulmonary trunk(12). In addition, they differentiate into the smooth muscle cells of the aortic arch arteries, the distal aorta and pulmonary trunk, and are required for aortic arch patterning(15). Further, neural crest cells influence early cardiac function and the development of the secondary heart field, a cardiogenic cell population in the pharynx required for proper alignment of the outflow vessels with respect to the ventricles(16–18). Therefore, fever-induced changes to neural crest cell function could provide a unifying mechanism to account for the types of birth defects observed following fever.

During our investigation, we discovered that two temperature-activated transient receptor potential (TRP) ion channels, TRPV1 and TRPV4, are present in cardiac and cranial neural crest cells during critical windows of heart and facial development. Several TRP channels are the principle components for temperature sensation within sensory neurons(19–21), although other TRP channel functions include modulating Ca^{2+} during cytoskeleton rearrangements, promoting cell polarity critical for cell migration, and regulation of cellular metabolism, suggesting highly diverse functions outside of the nervous system(22–24). We hypothesized that transient changes in the activity of temperature-activated TRP channels in neural crest cells confers susceptibility to fever-associated craniofacial and congenital heart defects. Using pharmacologic approaches to antagonize TRP activity, we protected developing chick embryos from hyperthermia-mediated birth defects. In addition, we utilized a TRP channel agonist to replicate associated birth defects in normothermic conditions. Finally, we developed a technology to remotely control transient permeability of temperature-activated TRP channels with cellular resolution using radiofrequency (RF) waves. Transient activation of these TRP channels in neural crest cells was sufficient to replicate febrile-associated birth defects.

Results

Hyperthermia induced neural crest cell-related birth defects.

We investigated the developmental effects of hyperthermia on neural crest cell-derived craniofacial and cardiovascular structures in chick embryos. The mechanisms of neural crest cell development are largely conserved between species and the chick is a well-established model to study human cardiovascular and craniofacial defects. To target critical stages in neural crest cell development, we heated Hamburger Hamilton (HH) stage 11–15 embryos to 40–41°C for 1 hour then re-incubated them at 37°C and analyzed for craniofacial and heart defects at HH36. We performed alcian blue and alizarin red stains to analyze the craniofacial features in experimental animals (Fig. 1A). Individual upper beak lengths were measured and normalized to femur lengths to account for small variations in staging (Fig. 1B). Our analysis revealed that hyperthermia exposure resulted in significant craniofacial defects including a reduction in upper beak lengths compared to normothermic control chicks with severe beak defects observed in 16% of hyperthermia exposed animals (Fig. 1C and fig. S1, A and B). Cardiovascular defects were increased in hyperthermia-exposed embryos. These defects included a significant incidence of conotruncal defects including double outlet right ventricle (DORV, Fig. 1D,E) and aortic arch patterning defects (fig. S2, A to F). All of these defects are directly linked to neural crest cell dysfunction(13). Due to the high association of outflow tract obstruction defects with maternal fevers in humans we used stereology to analyze the luminal cross-sectional areas of the aorta and pulmonary trunk using the Cavalieri probe(25). We identified a 72% and 55% reduction in luminal cross-sectional areas of the aorta and pulmonary trunk, respectively (Fig. 1F,G). The obstructive lesions were supravalvular and coincident with neural crest-derived smooth muscle vessel regions. These data demonstrate that experimentally controlled hyperthermia in chick models can produce neural crest-associated birth defects reminiscent of those observed in human infants following maternal fever.

TRPV1 channel inhibition mitigated fever-associated birth defects.

To identify potential teratogenic mechanisms of hyperthermia, we first examined the expression profile of mRNAs encoding temperature-activated vanilloid TRP channels (TRPV1–4) in neural folds containing neural crest cells in chick embryos at HH9 using RT-PCR. As a control, mRNA transcripts for all four TRPV channels were detected in whole embryo RNA from HH14 and HH22–24 embryos. However, only transcripts for *TRPV1*, *TRPV2*, and *TRPV4* were detected in dissected neural folds enriched for pre-migratory neural crest cells at HH9 (fig. S3A). TRPV2 is activated by temperatures $>52^{\circ}\text{C}$ thus making this channel less likely to be impacted by temperatures experienced during fever(20). However, TRPV1 can mediate heat-evoked Ca^{2+} currents $>40^{\circ}\text{C}$ and TRPV4 $>34^{\circ}\text{C}$ making these likely candidates for fever-induced activation(19, 26). To verify expression of *TRPV1* and *TRPV4* in migrating neural crest cells, neural tubes were electroporated with a plasmid encoding green fluorescent protein (GFP) at HH9, explanted and cultured overnight to allow for GFP⁺ neural crest cells to migrate from the neural tube (fig. S3B). RT-PCR analysis on FACS sorted GFP⁺ neural crest cells (fig. S3C,D) confirmed that both *TRPV1* and *TRPV4* transcripts were present in this population (fig. S3E).

The neural crest-related defects observed after hyperthermia exposures between HH11–15 suggested a critical developmental window of susceptibility. To determine if cranial and cardiac neural crest cells expressed *TRPV1* and or *TRPV4* during these developmental stages, we combined in situ hybridization for *TRPV4* or *TRPV1* mRNAs with immunohistochemistry for the neural crest cell marker HNK1. *TRPV1* transcripts were detected in HNK1⁺ premigratory and migrating HNK1⁺ cranial and cardiac neural crest cells between HH11 and HH13 (fig. S4, A to J). Similarly, *TRPV4* transcripts were also detected at these stages in HNK1⁺ cranial and cardiac neural crest cells (fig. S5, A to F). Thus both *TRPV1* and *TRPV4* transcripts are detected in neural crest cells during a critical developmental window of susceptibility to hyperthermia-induced defects. These *TRPV1*⁺ and *TRPV4*⁺ neural crest cells populate pharyngeal arches 1 and 2 and are the primary contributions to the bone and cartilage of craniofacial structures. Neural crest cells populating pharyngeal arches 3, 4, and 6 form the outflow septum, the smooth muscle of the great arteries, and influence aortic arch patterning. These neural crest cells are also critical to secondary heart field development and influence conotruncal alignment(18).

Next, we confirmed the responsiveness of chick TRPV 1 and TRPV4 with channel specific ligands. Treatment of chick TRPV4-expressing CHO cells with the TRPV4 agonist, GSK1016790A (GSK101), increased Ca^{2+} permeability as assessed by the genetically encoded Ca^{2+} indicator GCaMP6 (fig. S6A). Identifying an agonist for chick TRPV 1 was more challenging because it lacks the capsaicin binding site and is therefore insensitive to capsaicin and related agonists(27). However, the cysteine knot (ICK) peptide spider toxin from *P. cambridgei*, vanilloid toxin 3 (VaTx3), interacts with the outer pore of TRPV1 and is predicted to activate chick TRPV1(28). VaTx3 produced comparable changes in Ca^{2+} permeability in HEK293T cells transiently expressing either the murine or chick TRPV1 channels (fig. S6B). The TRPV1 specific antagonist SB366791 inhibited VaTx3-dependent chick TRPV1 activity (fig. S6C) suggesting the VaTx3-mediated response was TRPV1-specific.

To determine if *TRPV* transcripts detected in neural crest cells translated into functional channels, we electroporated chick neural tubes *in ovo* at HH9–10 with GCaMP6 and explanted them to culture plates to allow GCaMP6⁺ neural crest cells to migrate from the neural tubes (Fig. 2A). Explanted neural crest cells were exposed to the TRPV4 agonist GSK101, which produced a robust increase in Ca²⁺ permeability that was blocked by pretreatment with the TRPV4 antagonist RN1734 (Fig. 2B)(29). Explanted neural crest cells exposed to VaTx3 showed increased Ca²⁺ responses, demonstrating functional TRPV1 channels in neural crest cells (Fig. 2C). To confirm that neural crest cell-specific TRPV1 and TRPV4 channel activity extended beyond avian species, we analyzed primary mammalian neural crest cells for changes in intracellular Ca²⁺ following exposure to TRPV1 and TRPV4 ligands. Exposing mouse neural crest cell explants to increasing doses of GSK101 produced corresponding increases in Ca²⁺ responses (Fig. 2D), which were blocked by the TRPV4 inhibitor RN1734 (Fig. 2E). Capsaicin application induced Ca²⁺ permeability in mouse neural crest cells, an effect that was blocked with the TRPV1 inhibitor SBB366791 (Fig. 2F). These data demonstrate that TRPV 1 and TRPV4 are also present in neural crest cells during mammalian development. Exposure of primary chick neural crest cells to 40°C resulted in increased Ca²⁺ permeability and this response was reduced by pretreatment with the TRPV4 inhibitor RN1734 (Fig. 2G-I). Thus, neural crest cells responded to hyperthermia in a manner that was partially mediated through TRPV channels.

We next hypothesized that hyperthermia-induced defects in neural crest-dependent structures in our chick model could be rescued by inhibition of TRPV channels. However, treatment of HH11 embryos with the TRPV4 inhibitor RN1734 under normothermic conditions produced a significant decrease in beak length and a 20% incidence of conotruncal heart defects (fig. S7, A to I). These defects suggested that endogenous TRPV4 activity was necessary for normal craniofacial and heart development, thus precluding us from targeting TRPV4 activity to rescue hyperthermia-associated birth defects. In contrast, treatment of HH11 embryos with the TRPV1 inhibitor SB366791 treatment at normal temperatures did not produce detectable heart or facial phenotypes. Pretreatment of embryos with SB366791 before hyperthermia prevented hyperthermia-induced craniofacial defects (Fig. 3A,B) and significantly reduced supravalvular stenosis (Fig. 3C-E). The number of conotruncal defects were reduced by ~50%, although this trend did not reach statistical significance (Fig. S8, A to G). These data link TRPV channel activation with neural crest-related congenital defects in our chick hyperthermia model.

Pharmacologic activation of TRPV4 replicated heat-induced birth defects in multiple animal models.

As a complementary approach, we exposed developing chick embryos at HH11–15 to the small molecule TRPV4 agonist GSK101 under normothermic conditions to determine if pharmacologic activation of TRPV4 *in ovo* induced neural crest-related defects. Conotruncal defects were detected in 20% of GSK101-exposed embryos at HH36 (Fig. 4A,B and fig. S9, A to D). GSK101-treatment also replicated the severe supravalvular stenosis as indicated by a 50% and 44% reduction in luminal cross-sectional areas of the aorta and pulmonary trunk, respectively (Fig. 4C-E). As in the hyperthermia-exposed chicks, GSK101-treated embryos had significantly shorter beak-to-femur ratios compared to controls (Fig. 4F-H). Together,

these data provide evidence that activation of TRPV4 channel permeability during critical windows of susceptibility can produce cardiac and craniofacial lesions that resemble those associated with maternal fevers in humans.

We posited that if TRPV activation is a broadly relevant driver mechanism for neural crest-dependent congenital defects, then pharmacologic TRPV4 activation should reproduce these defects in another vertebrate. Because *TRPV4* transcripts are detected in neural crest-derived jaw structures of zebrafish during early head development(30), we targeted TRPV4 activation in developing zebrafish larvae. While zebrafish single ventricle heart anatomy limits our ability to assess conotruncal alignment defects, it offers an excellent model of craniofacial development. First, we confirmed that GSK101 activates zebrafish TRPV4 transiently expressed in CHO cells (Fig. 5A). We next used 1.4*cola1:egfp* transgenic zebrafish larvae which express *egfp* gene under the *cola1* promoter to drive expression in jaw structures(31) to monitor craniofacial development in live fish over time. We analyzed for craniofacial defects in zebrafish larvae treated at 2dpf with GSK101 or GSK1153218 (a structurally related compound which lacks TRPV4 activity, table S1). By 3dpf and 4dpf, only GSK101 exposed larvae exhibited a significant reduction in jaw length, as indicated by the decreased distance between Meckel's and ceratohyal cartilages (Fig. 5B-D). Moreover, histological examination revealed disruption of the cellular organization in the ethmoid plate suggesting defective palatogenesis (fig. S10, A and B). Consistent with the developmental timing and observed phenotypes of our chick studies, these data demonstrate the impact of aberrant TRPV channel activation on neural crest-derived craniofacial development extends beyond an avian model.

Remote activation of TRP channels in neural crest cells phenocopied fever-associated birth defects.

To test the hypothesis that congenital defects can be caused by transient activation of TRPV channels specifically within neural crest cells, we needed temporal and cell-specific control over these channels during development. Under specific conditions, modified ferritin-bound TRPV channels can be remotely controlled in vivo using radio frequency electromagnetic fields(32–34) or static magnetic fields(35). We engineered FeRIC (Fe³⁺ Redistribution to Ion Channels) modifications to TRPV1 and TRPV4 channels to direct endogenous intracellular ferritin to the cytoplasmic domains of TRPV 1 or TRPV4 channels. Specifically, we fused the ferritin-binding domain 5 (D5) of Kininogen-1 (36) with the C-terminal cytoplasmic domain of either TRPV1 (TRPV1^{FeRIC}) or TRPV4 (TRPV4^{FeRIC}) to confer responsiveness to RF (Fig. 6A,B and fig. S11, A to C, and S12, A to C). By targeting endogenous ferritin found in cells, we eliminated the need for potentially cell-toxic exogenous iron administration or co-expression of modified ferritin(33, 37).

To confirm FeRIC-mediated ferritin redistribution to the cell membrane, we investigated the subcellular localization of a ferritin heavy chain-mCherry fusion (FTH1^{mCherry}) protein in TRPV1^{WT} and TRPV1^{FeRIC}-expressing HEK293T cells. In TRPV1^{WT} cells, FTH1^{mCherry} showed a cytoplasmic distribution consistent with normal FTH1 expression. However, when FTH1^{mCherry} was co-transfected with TRPV1^{FeRIC}, FTH1^{mCherry} was redistributed to the cell membrane by 24 hours (Fig. 6C). In addition, both TRPV1^{FeRIC} and TRPV4^{FeRIC}, but

not TRPV1^{WT}, TRPV4^{WT} or FLAG constructs co-immunoprecipitated FTH1 (Fig. 6D). Next, we tested the influence of FeRIC-mediated ferritin redistribution on the bioavailability of cellular iron metabolism by measuring changes in four well-characterized iron responsive genes. Redistribution of cellular ferritin using TRPV1^{FeRIC} or TRPV4^{FeRIC} did not significantly change mRNA concentrations of *Fpn1* (which encodes ferroportin-1), *TFR1* (which encodes transferrin receptor 1), *SLC11A2* (which encodes divalent metal transporter 1), or mRNAs encoding heavy and light chains of ferritin (fig. S13, A to E), suggesting that iron bioavailability was not substantially altered.

To remotely activate FeRIC channels, we used a 175 MHz RF coil that resonates at a frequency similar to that generated by a standard 3-Tesla MRI scanner. Measurement of the current in the RF coil was 1.5A peak-to-peak, generating a mean magnetic field of 36 μ T and a specific absorption rate (SAR) of 6.8 W/kg (fig. S14, A to D). We confirmed that this magnetic field did not induce bulk temperature increases that could activate FeRIC channels by measuring the temperature of the cell media immediately before and after applying RF. The temperature fluctuation was measured at $< \pm 1^\circ\text{C}$. Because membrane-localized Fe³⁺ combined with RF could potentially be cytotoxic, we investigated cellular viability and membrane integrity. Treating TRPV1^{FeRIC}-positive HEK293T cells with 10 minutes of RF had no detectable impact on membrane integrity and did not affect cell viability at 24 hours (fig. S15, A to C). We next tested the responsiveness of TRPV1^{FeRIC} and TRPV4^{FeRIC} to RF in HEK293T cells co-expressing GCaMP6. RF stimulation of unmodified TRPV1^{WT} or TRPV4^{WT} did not alter Ca²⁺ permeability. However, channels responded to capsaicin (TRPV1, Fig. 6E) or GSK101 (TRPV4, Fig. 6F) agonists as expected. Conversely, only TRPV1^{FeRIC} or TRPV4^{FeRIC} were transiently responsive to RF stimulation and their respective ligands. In both cases, the FeRIC-dependent RF responses were abolished in the presence of channel-specific inhibitors (Fig. 6E,F). We next confirmed that remote activation of FeRIC channels required ferritin expression. Using CRISPR/Cas9, we deleted the ferritin heavy chain gene (*FTH1*) in HEK293T cells (*HEK^{FTH1-KO}*, fig. S16A-C). Deletion of *FTH1* abolished TRPV1^{FeRIC} responses to RF but did not interfere with capsaicin responses (fig. S16D,E,G). Transfection of *HEK^{FTH1-KO}* cells with human *FTH1* rescued RF responsiveness to TRPV1^{FeRIC} (fig. S16D-G).

To gain mechanistic insight into how RF activates TRPV1^{FeRIC} we generated TRPV1^{T FeRIC} which has mutations that abolish temperature activation (N628K, N652T, and Y653T) (38) without affecting chemical sensitivity (fig. S17A-C). When expressed in HEK293T cells, the FeRIC-modified triple mutant was not responsive to temperature changes as expected but retained its responsiveness to capsaicin (fig. S17D,E). In addition to the disruption of temperature-activation, these three point mutations also abolished RF-induced changes in permeability, suggesting that RF stimulation may act on TRPV1^{FeRIC} through a temperature-specific mechanism (fig. S17F).

We next sought to test whether cell-specific activation of the candidate TRPV channels phenocopied fever-associated heart defects. RF applied to chick neural crest cells expressing TRPV1^{FeRIC} increased Ca²⁺ permeability, which was blocked by the TRPV1 inhibitor SB366791 (Fig. 7A), thus confirming that TRPV1^{FeRIC} was responsive to RF in these cells. For *in ovo* experiments, TRPV1^{FeRIC} or TRPV1^{WT} plasmid DNA was bilaterally

electroporated into premigratory neural crest cells at HH9. Expression of TRPV1^{FeRIC} alone did not appreciably affect neural crest cell migration to the pharyngeal arches at HH13–14 (Fig. 7B). Twenty-four hours after electroporation, eggs were either exposed for 10 min to RF (RF^{10min}) or not (RP^{neg}). Of the TRPV1^{WT} groups, RF^{neg}TRPV1^{WT} and RF^{10min}TRPV1^{WT} did not show conotruncal defects (Fig. 7C,D). In contrast, RF^{10min}TRPV1^{FeRIC} embryos exhibited conotruncal alignment defects in 48% of the embryos (Fig. 7C,D and movie S1). These defects included overriding aorta, DORV, and tetralogy of Fallot and were accompanied by a perimembranous VSD (fig S18, A to L). In contrast, we observed only a single heart with an overriding aorta and VSD in RP^{neg}TRPV1^{FeRIC} embryos (Fig. 7D). Remotely induced conotruncal defects were rescued by pretreatment with the TRPV1 inhibitor SB366791 (Fig. 7D and fig. S18, A to L). To control for generalized RF effects, unelectroporated eggs were exposed to 10 minutes of RF and all unelectroporated RF^{10min} control eggs had structurally normal hearts (Fig. 7D).

Next, we examined the vessels for outflow obstructive lesions and found no significant differences in cross-sectional luminal areas of the outflow tracts between TRPV1^{WT} embryos with or without RF stimulation or between RP^{neg}TRPV1^{FeRIC} embryos and controls (Fig. 7E-G). However, similar to hyperthermia and GSK101-exposed embryos, RF^{10min}TRPV1^{FeRIC} embryos displayed a 61% and 37% decrease in luminal cross-sectional areas in the aorta and pulmonary trunk respectively (Fig. 7E-G). Supravalvular stenosis was also rescued by pretreatment with SB366791 (Fig. 7E-G).

We also analyzed the craniofacial structures in TRPV1^{FeRIC}-expressing embryos following remote activation in cranial neural crest cells and compared them to RP^{neg}TRPV1^{FeRIC} controls. Remote activation of TRPV1^{FeRIC} resulted in a significant reduction in upper beak length compared to control or RP^{neg}TRPV1^{FeRIC} embryos (fig. S19, A to C). These data demonstrate that transient activation of TRPV1 in cranial neural crest cells is sufficient to phenocopy craniofacial defects associated with maternal fevers.

We analyzed neural crest cell-specific remote activation of TRPV4^{FeRIC} in chick neural crest explants. RF induced transient Ca²⁺ permeability in neural crest cells expressing TRPV4^{FeRIC} that was blocked using RN1734 (Fig. 8A). Similar to FeRIC modified TRPV1, remote activation of TRPV4^{FeRIC} in neural crest cells produced a 57% incidence of conotruncal defects in the RF^{10min}TRPV4^{FeRIC} group compared to no defects in the RP^{neg}TRPV4^{FeRIC} group (Fig. 8B,C). Remote activation of TRPV4^{FeRIC} also resulted in significant supravalvular stenosis of the aorta and pulmonary trunk (Fig. 8B, D, and E). Taken together, these data indicate that transient activation of neural crest cell-specific TRPV1 or TRPV4 can replicate the types of clinically important heart defects that have been linked to maternal fevers in humans.

To confirm that FeRIC-mediated defects were not caused by RF-induced bulk temperature increases in tissues, we non-invasively mapped temperature changes throughout the egg and embryo at the beginning and end of the 10 min RF exposure using MRI thermometry(39). No substantial temperature differences were observed between RF^{10min} and RP^{neg} groups (fig. S20, A to H). Thus, RF exposure does not significantly heat the embryo to induce

congenital defects, which is also supported by the 0% incidence of cardiovascular defects in control eggs exposed to RF (Fig. 7D).

Discussion

Our study identified the presence of two temperature-activated ion channels, TRPV 1 and TRPV4, in cranial and cardiac neural crest cells during critical windows of heart and facial development. We demonstrated that experimentally controlled activation of these channels through temperature, pharmacological, or genetic approaches was sufficient to produce congenital heart and craniofacial defects similar to those associated with human fever in the first trimester. We linked TRP channel activity to hyperthermia-induced defects by showing that TRPV1 inhibition during heat exposure mitigated the majority of the congenital defects. Both TRPV4 agonist and antagonists elicited similar neural crest cell-related defects under normothermic conditions. It is not uncommon for a gain or loss of function of a signaling pathway to result in similar defects, particularly in secondary heart field defects like DORV. For example, in previous studies we have shown that too much or too little FGF signaling in the pharynx (which is mediated by neural crest cells) results in proliferation of the secondary heart field progenitors or premature differentiation of the progenitors respectively(18, 40, 41). Both result in a failure of the secondary heart cells to lengthen the outflow tract which is necessary for proper outflow tract alignment. Similarly, Noonan and Leopard Syndromes have overlapping cardiac and craniofacial phenotypes. Noonan syndrome is attributed to an activating mutation in *PTPN11* in 50% of patients whereas Leopard Syndrome is associated with an inactivating mutation in the same gene.

In our model, the timing of the hyperthermia and or TRP channel activation in neural crest cells coincided with their migration into the pharyngeal arches. While the immediate consequences of transient TRP channel activation in neural crest cells will be the focus of future studies, many studies have already shown that increased Ca^{2+} signaling impacts cell migration, survival and proliferation, all of which are linked to neural crest-associated heart and facial defects. Disruption of neural crest cell polarity is linked to conotruncal and craniofacial defects(42, 43). Increased maternal homocysteine is associated with neural crest-associated defects and homocysteine enhances cardiac neural crest cell attachment and alters migration in vitro through a Ca^{2+} -dependent mechanism(44, 45). Further, increased Ca^{2+} homeostasis in neural crest cells is linked to human diseases including fetal alcohol syndrome and Timothy syndrome, both of which have neural crest-related craniofacial and or conotruncal defects(46, 47). Together, these studies suggest that intracellular Ca^{2+} homeostasis is critical for normal neural crest-dependent development.

The study of environmentally-induced transient alterations in genetically normal TRP channel activation has not been described in the context of congenital birth defects. In part, this may be due to the lack of widely available technology enabling precise temporal control of targeted cell-specific activation. Purely pharmacologic approaches lack cell specificity and can be influenced by bioavailability through placental interfaces and maternal or fetal metabolism. Genetically modified channels with constitutive activity, or those that lack activity, can be targeted to cell-specific populations but cannot facilitate transient changes needed to study the impact of passing environmental factors such as fever. Technologies

using synthetic designer ligands such as DREADD receptor approaches can provide cell specificity and some temporal resolution(48, 49). However, in our study we needed much shorter restrictions on activation times to mimic the temperature spikes such as those that occur in febrile states. Once synthetic compounds used to activate DREADD receptors are injected into the egg, we would not be able to remove the compounds and therefore their effects could last hours to days depending on metabolism within the chick embryo. Our approach using RF offered the ability to precisely control the presence or absence of channel stimulus remotely. Here, we developed a simplified technology, FeRIC, that utilized the cell's endogenous iron stores to convert the modified channel into a remotely controlled channel with high temporal resolution and cellular specificity.

The physical mechanisms of ferritin-dependent electromagnetic control over ion channels is under debate (50, 51). The functionality of FeRIC channels is in line with those of similar technologies independently developed by other laboratories(33, 35, 52). Here, we showed that FeRIC channels depended on the presence of endogenous intracellular ferritin. When the *FTH1* gene was deleted *in vitro*, TRPV1^{FeRIC} was unresponsive to RF. Reintroduction of *FTH1* in these cells rescued the remote capabilities of TRPV1^{FeRIC}. In another experiment, we showed that the introduction of temperature-sensitive amino acid substitutions in the outer pore region of TRPV1 also abolished RF-dependent channel activity. These data are suggestive but not conclusive evidence for a temperature-dependent mechanism. However, one important theoretical consideration suggests that RF waves at the power level reported in the literature are not sufficient to induce significant heating of ferritin(50). Thus, whether FeRIC-modified channels are activated by RF by a heat-dependent mechanism will require further measurements, such as nanoscale-resolution mapping of iron distribution and temperature change in unexposed and RF-exposed cells expressing these channels. Should this evidence become available, we anticipate that FeRIC modifications could be expanded to study other temperature-sensitive channels, both in the context of development as well as in relation to other spatiotemporally sensitive disease mechanisms.

Lastly, maternal infections with associated fevers in the first trimester are linked to clinically important craniofacial and cardiac defects. Craniofacial defects range from midface hypoplasia to clefts and cardiac defects include obstructive lesions and conotruncal defects. Hyperthermia has been proposed as a direct teratogen; however, the mechanisms linking temperature to birth defects are unknown(53, 54). It is critical to distinguish between infection-based teratogenicity and simple hyperthermia because the latter is a modifiable risk factor. Acetaminophen is drug commonly used by pregnant women and is safe and protects against some fever-associated birth defects(8, 55, 56). However, new concerns have been raised over its repetitive use in pregnancy and the risk for behavioral problems in childhood(57). Our data provide a rational molecular mechanism for hyperthermia-induced teratogenicity mediated through transient gating of temperature-activated ion channels in neural crest cells. Because hyperthermia is a modifiable risk factor in pregnant women, these fever-associated birth defects may be preventable through public awareness and judicious use of anti-pyretics in febrile pregnant women. We hope that these observations will stimulate detailed clinical studies with a view to implement new policies in prenatal care and allow pregnant women to fully weigh the risks and benefits of antipyretic therapy during pregnancy.

Materials and Methods:

Animals

All animal experiments were approved by the Duke University Animal Care and Use Committee, and were performed in accordance with the institutional and NIH ethical guidelines. Fertilized Ross Hubert chick eggs (*Gallus gallus domesticus*, Mountaire Farms, Siler City, NC) were incubated for 1–10 days at 37°C and 70% humidity. Embryos were staged according to Hamburger and Hamilton(58). Adult zebrafish (*Danio rerio*) and the *Tg-1.4coll1a1:egfp* line were maintained at 28°C on a 14/10 h light and dark cycle. Embryos were raised at 28°C and staged according to hours post fertilization (hpf) and morphology. Embryos were growth in E3 medium (NaCl, 5mM; KCl, 0.17mM; CaCl₂, 0.33mM; MgSO₄, 0.33mM, 0.00001% Methylene Blue). All embryos (chick or zebrafish) were stage matched and randomized to the specific treatment groups described below. The investigator was not blinded to the various treatment groups.

Cell Lines

Where indicated, the human embryonic kidney cell line (HEK293T, Clontech Cat#632180) or Chinese Hamster Ovary cell line (CHO, ATCC CCL-61) were used. Both cell lines have tested negative for Mycoplasma contamination by Clontech or ATCC respectively. The identity of HEK293T cell line was verified by STR analysis at Clontech. The identity of CHO cell lines was confirmed at the ATCC using an Isoenzyme (Interspecies) assay for hamster.

Hyperthermia experiments

Chick embryos were incubated at 37°C until the desired age (HH 9–15). The eggs were then placed in a 40–42°C incubator. Internal egg temperature was monitored every 15 min using a digital thermometer probe. It generally took 45–60 min for the eggs to reach the targeted temperature. Once the target temperature was achieved, the eggs were incubated for another 60 min. Following hyperthermia, embryos were re-incubated at 37°C and allowed to develop until day 10 (HH36). The embryos were harvested and gross congenital defects were documented. All hearts and heads were further processed and analyzed for structural defects (see below). Hyperthermic exposures of 40–41°C at HH1–13 produced the most consistent defects. For the hyperthermia rescue experiments embryos were either pretreated with 10µM SB366791 or 0.1% DMSO in PBS, incubated at 41°C for 1hr as described above (see below for details of drug delivery), returned to a 37°C incubator and harvested at HH36 for analysis of heart and craniofacial defects.

Pharmacological treatments of chick embryos

Eggs were windowed at HH1–14 and treated with 20µL of varying concentrations of TRPV channel agonists or antagonist (see below) to determine an effective dose range. Controls included untreated embryos and DMSO-vehicle controls. The eggs were sealed with tape, incubated until HH36 and analyzed for heart and craniofacial defects. All drug stocks were made in DMSO. Stocks were diluted in PBS such that the DMSO concentration was never higher than 0.1% and all treatments were delivered in 20µL volume to the embryo. We

determined the effective GSK101 (TRPV4 agonist) dose to be 20 μ L of a 10–50 μ M solution per egg (~3.3nM-16.7nM final concentration in an average 60mL egg volume). The incidence and severity of defects increased at the higher doses. Lower doses did not induce defects and no concentration tested caused lethality. Eggs dosed with the 10–100 μ M RN1734 (1.2nM-12nM final concentration) caused significant incidence of heart and craniofacial defects. For the hyperthermia and TRPV1^{FeRIC} rescue experiments embryos were treated with 10 μ M SB366791 (1.0 nM final concentration).

TRPV RT-PCR

Total RNA was extracted from HH14, HH22, or dissected HH9 neural folds. cDNA was used in subsequent PCR reactions using the following published primer sets: Chick *TRPV1* (F) GTCCTGCATAGACACATGT Chick *TRPV1* (R) GCACAAAATACTGTATCCC (59); Chick *TRPV2*(F) CCCTTGAGATCACCTTACC Chick *TRPV2*(R) CTTCCCAGTCTTTGCATCT (60); Chick *TRPV3*(F) CCCCTCAATTCACCTCCTGC Chick *TRPV3*(R) GGAAAGGCATTCACCACCA (60); Chick *TRPV4*(F) TTCAAGGATTGGGCATACG Chick *TRPV4*(R) ATTAACCCTCACTAAAGGGCAACTTCCAGATGTGTTTG (59); Chick *SLUG*(F) GCCTGCCTTCAAATGCCACGCTCCTTCCTGG Chick *SLUG*(R) GGCTGCTGCGTAGCACACTGAGTCATGCAGTC (61).

For the flow sorted chick neural crest explant studies, a plasmid encoding GFP was electroporated at HH9 into the neural tube. The electroporated embryos were removed and the neural tubes were enzymatically dissociated from the surrounding tissue. Specifically, the cranial and cardiac neural crest region extending from the midbrain to the third somite was placed in a solution containing 0.5 μ g/ml dispase and 1 μ g/ml collagenase for 1–2min. The embryos were then placed in a stop/wash solution of DMEM with 20% fetal bovine serum (FBS) and the surrounding tissues were dissected away from the neural tube using fine needles. The isolated neural tubes were cultured on fibronectin-coated plates overnight at 37°C in DMEM with 20% FBS supplemented with 2% chick embryo extract. 24 hours after plating, the neural tubes were mechanically removed with fine forceps. The remaining neural crest cells were treated with 0.25% trypsin for 2 minutes at 37°C, collected, and washed in DMEM with 20% FBS. The cells were sorted in the Duke Flow Cytometry Core Facility on a MoFlo XDP (Beckman Coulter Life Sciences).

In situ hybridization and immunohistochemistry

Whole mount in situ hybridization was carried out using a standard protocol with DIG-labeled probes(62). Previously published probes were used for *TRPV1* and *TRPV4* in situ hybridizations(59). PCR amplified probe templates were cloned and sequenced at the Duke Cancer Biology Sequencing Facility. Plasmids containing probe sequences were used to generate RNA probes from the T7 or SP6 promoters in vitro. After visualization of DIG, the embryos were embedded in paraffin, sectioned and photographed to document the cellular localization of *TRPV1* and *TRPV4* transcripts. To determine co-localization of *TRPV1* and *TRPV4* transcripts with neural crest cells, the coverslips were soaked in PBS to remove from the sectioned in situ hybridizations. Tissue sections were stained with mouse IgM HNK1

antibody (ATCC) overnight at 4°C and developed with 3,3'-Diaminobenzidine tetrahydrochloride. All images were acquired on a Leica DMRA2 compound microscope.

Plasmids

To generate the TRPV1^{wt} construct, full length murine wild-type TRPV1 was PCR amplified from cDNA generated from spinal cord tissue from C57/B16 mice. 5' SpeI sites and 3' NotI sites were introduced into their respective locations outside of the open reading frame using PCR. This product was subcloned into SpeI and NotI sites within the multiple cloning site of the PLVX-IRES-mCherry vector to generate TRPV1^{wt} (Clontech, for detailed map see fig. S12). To generate the TRPV1^{FeRIC} construct, PCR primers were designed to eliminate the 3' stop site in wild-type TRPV1 and introduce a novel 3' XbaI site. PCR primers introducing a 5' XbaI site and a 3' NotI site and a stop codon were used to amplify human Kininogen1 domain 5 (FeRIC) from whole blood DNA. This FeRIC fragment was subcloned into the XbaI NotI sites within the PLVX-IRES-mCherry vector (for detailed map see fig. S12). All completed constructs were sequenced verified by the Duke Cancer Biology Sequencing Facility and analyzed using MacVector 13.0.

To generate the TRPV1^{T FeRIC} construct, a synthesized cassette of a 798 bp region was designed that included a native TRPV1 PsHA1 restriction site, introduced the code for triple mutant amino acids N628K, N652T, and Y653T, and removed the native stop codon. We added a novel XbaI restriction site for ligation in frame with FeRIC sequence. Synthesis of the cassette was done by BioBasic Inc. Original TRPV1^{FeRIC} construct was digested by psHA1 and XbaI, which removed the 3' terminal end of TRPV1 ending at the FeRIC start. The psHA1-XbaI mutant cassette was then ligated into the corresponding sites in the TRPV1^{FeRIC} construct. Complete construct with these 3 mutations was sequenced verified by the Duke Cancer Biology Sequencing Facility and analyzed using MacVector 13.0.

To generate the TRPV4^{WT} construct, full-length rat *TRPV4* cDNA was a generous gift from Dr. Robert Lefkowitz (Duke University). SpeI and NotI restriction sites were introduced using PCR as above. The full-length wild-type *TRPV4* was subcloned into PLVX-IRES-mCherry vector to generate TRPV4^{WT} (Clontech, for detailed map see fig. S13). To generate the TRPV4^{FeRIC} construct, PCR primers were designed to eliminate the 3' stop site in wild-type *TRPV4* and introduce a 3' NotI site. PCR primers introducing a 5' NotI site and a 3' BamHI site and a stop codon were used to amplify human *Kininogen1 domain 5* (FeRIC). This FeRIC fragment was subcloned into the XbaI BamHI sites within the PLVX-IRES-mCherry vector containing *TRPV4* (for detailed map see fig. S13). All completed constructs were sequenced verified by the Duke Cancer Biology Sequencing Facility and analyzed using MacVector 13.0.

Full-length chick *TRPV4* was synthesized and sequence verified by Biobasic Inc. The full-length construct was subcloned into the PLVX expression vector using novel SpeI (5') and NotI (3') sites introduced outside of the open reading frame. Full-length zebrafish *TRPV4* was cloned from pooled zebrafish larval cDNA generated from embryos at 24–96hpf using the following 5' primer containing a novel SpeI site and the 3' primer containing a novel xbaI site: *zTRPV4*(F)

CTATTTCCGGTGAATTCCTCGAGACTAGTCTGGCCATGACAGAGTCCTTGTCTG

and *zTRPV4*® CGGGATCCGCGGCCGCTCTAGATTAGCTTTCAGACTTGAGTCGG. All constructs were sequenced verified by the Duke Cancer Biology Sequencing Facility and analyzed using MacVector 13.0.

To generate the FTH1 constructs, the full-length human *FTH1* was cloned using RT-PCR and cDNA generated from FeCl₂ stimulated HEK293T cells. 5'-GCCGCCATGACGACCGCGT and 3'-CCGAGGCTTAGCTTTCATT primers flank the entire open reading frame. The entire open reading frame was cloned into pcDNA 3.1. For FTH1mCherry fusion construct, PCR was used to abolish the stop codon and subclone *FTH1* into in the multiple cloning site of pLVX-mCherry-N1 vector (Clontech) in frame with mCherry. Sequences were verified by the Duke Cancer Biology Sequencing Facility and analyzed using MacVector 13.0.

CRISPR Cas9 FTH1 deletion

Ferritin heavy chain (*FTH1*) was deleted in HEK293T cells using commercially available double nickase plasmids (Santa Cruz Biotechnology Inc.). This system uses two 20nt gRNA sequences targeting exon 2 of the human *FTH1* gene. Briefly, cells were transfected with plasmids encoding the gRNA sequences, Cas9, and the puromycin resistance gene. 24 hours later, transfected cells were selected in DMEM media containing 5mg/ml puromycin for 5 days. Surviving cells were harvested and washed and serially diluted into a 96-well plate. Wells containing single cells were expanded and screened by RT-PCR for full length *FTH1* mRNA using primers listed above. Full-length *FTH1* PCR products were cloned and the expected 147bp sequence deletion was confirmed in exon 2 of *FTH1*. 88% of the individual clones generated were FTH1 negative by RT-PCR and western blot. HEK^{FTH1KO} clone 2 was expanded and used for subsequent TRPV1^{FeRIC} RF analysis in this study.

RF coil

The RF emitting coil consisted of a double loop wire with a loop diameter of approximately 4 cm. The coil was connected in parallel with tuning capacitors forming an LC circuit. The circuit was tuned to a resonance frequency at approximately 175 MHz. RF signal was generated by a broadband (0–400 MHz) signal generator (Boonton Electronics, NJ, Model 102A) and amplified using a 50-W linear RF amplifier (Electronic Navigation Industries, TX, Model 550L). Electric current across the coil was measured using a current monitor (Pearson Electronics, CA, Model 2877). The peak-to-peak current was measured to be 1.5 A. During experiments, the coil sits roughly 1.5 cm above the cells or chick embryo.

RF simulation

The electromagnetic fields emitted by the RF coil were computed by cylindrical finite-difference time-domain simulation in Matlab© (Mathworks, MA, version R2014a) as described by Dib and Weller(63). Perfectly-matched-layer boundary conditions were implemented as described by Schneider(64) and the simulation resolution was 0.5 cm in the z direction, 0.25 cm in the radial direction and $\pi/28$ radians in the ϕ direction. The computed magnetic field was verified by comparison with the analytical expression for the magnetic field along the axis of a static current loop. The electromagnetic fields were simulated for a coil loaded with a dish of 0.5-cm thickness and 2-cm radius. The dish electrical conductivity

was 0.5 S/m, similar to human tissue (65). The AC input was 1.5 A peak-to-peak at 175 MHz.

Bilateral electroporations and RF delivery to eggs

For the cranial and cardiac neural crest cell electroporation studies, eggs were incubated to HH9–11. The eggs were windowed and the vitelline membrane was torn. One or two drops of Ringers solution was applied so that the embryo fell away from the vitelline membrane. Platinum electrodes were placed at the cranial end on either side of the embryo. TRPV1^{WT}, TRPV1^{FeRIC}, TRPV4^{WT}, or TRPV4^{FeRIC} constructs were injected into the neural tube using a glass micropipette. DNA concentration was 4µg/µl. Four 50ms square wave pulses were immediately applied at 20 V each. A drop of Ringers was again applied to the embryo, and injection and electroporation were repeated for the other side of the neural tube. 24 hours after electroporation, expression of the plasmid was confirmed by visualizing mCherry expression. Then the egg was placed in a 37°C chamber, the RF coil was placed on the egg surrounding the embryo and 10 minutes of RF stimulation (see above) was delivered. Control (RF^{Neg}) eggs were placed in the same 37°C chamber for 10 minutes without RF stimulation. For the TRPV1^{FeRIC} rescue experiments, TRPV1^{FeRIC}-expressing embryos were pretreated with 10µM SB366791 for 1hr prior to RF exposure.

Primary neural crest cell cultures

A mixture of GCaMP6(66) (obtained from Addgene#40753) was electroporated alone or in combination with the TRPV constructs described above into chick neural tubes at HH9. The electroporated embryos were removed from the egg and the neural tubes were enzymatically dissociated from the surrounding tissue. Specifically, the cranial and cardiac neural crest region extending from the midbrain to the third somite was placed in a solution containing 0.5µg/ml dispase and 1µg/ml collagenase for 1–2min. The embryos were then placed in a stop/wash solution of DMEM with 20% fetal bovine serum (FBS) and the surrounding tissues were dissected away from the neural tube using fine needles. The isolated neural tubes were cultured on fibronectin-coated plates overnight at 37°C in DMEM with 20% FBS supplemented with 2% chick embryo extract and Ca²⁺ imaging was performed as described below. Mouse neural crest cell explants were generated as above using E8.5 embryos from CD1 mice. Mouse neural tubes were cultured at 37°C in DMEM with 20% FBS.

Ca²⁺ imaging

HEK293T or CHO cells were plated on collagen-coated glass bottom 35 mm dishes and cultured in 10% FBS and 1% PenStrep in DMEM. After 24 hours, cells were co-transfected using the Lipofectamine LTX Plus reagent with GCaMP6 and either TRPV1^{WT}, TRPV1^{FeRIC}, TRPV4^{WT}, or TRPV4^{FeRIC}. For Ca²⁺ imaging of chick neural crest cells, neural crest cells were electroporated, isolated and cultured as described above. Twenty-four hours after electroporation, cytosolic Ca²⁺ were monitored by fluorescence imaging of cells positive for GCaMP6⁺ and TRPV channels (mCherry⁺) using a 20x objective on an inverted Olympus IX50 microscope equipped with a camera (Q-Imaging-Retiga 1300i) controlled by Metamorph software 7.8 (Molecular Devices). Images were captured at 1 frame per sec. Unless stated otherwise, all experiments were carried out at room temperature. Neural crest cell recordings were carried out in Leibovitz's L-15 Medium (ThermoFisher) supplemented

with Ca^{2+} to a final concentration of 2mM. For HEK293T cells, the imaging buffer solution contained 140mM NaCl, 2.8mM KCl, 1mM MgCl_2 , 2mM CaCl_2 , 10mM glucose, and 10mM HEPES at pH 7.4. RF was delivered for 10 min using a custom built RF emitting coil designed to fit the 35mm tissue culture dish as described above. After RF, cells were then exposed to TRP channel agonists, 1 μM capsaicin or 1 μM GSK1016790A. In a second series of experiments, HEK293T or neural crest cells were exposed to TRPV1 or TRPV4 channel inhibitors (10 μM SB366791 or 10 μM RN1734) for 10 min and then imaged for changes in intracellular Ca^{2+} with or without RF exposure. Images were analyzed using Metamorph software; cytosolic regions of interest (ROIs) were placed over those cells which co-expressed GCaMP6 and TRPV channels. GCaMP6 fluorescence intensity was measured for each image of the time-lapse acquisition (650 sec). The data were fit with a double exponential decay time and corrected for photobleaching using the Microcal Origin 7 software (OriginLab, MA). Responses are presented as F/F_0 , where F_0 is the resting fluorescence averaged over 60 s before the start of stimulation and F is the change in fluorescence over resting values. For each experimental condition, 50–150 cells were analyzed and each experiment was conducted at least 3 times. Plots in the individual figures are representative tracings.

To image Ca^{2+} changes in mouse neural crest cells, neural tubes from E8.5 embryos were dissected and plated as described above. 8–24 hours after culture, cells were washed with imaging buffer solution and incubated with 5 μM fluo-4 AM (Molecular Probes) for 40 min at room temperature. Fluo-4 loaded cells were imaged using the imaging system described above. Fluo-4 was excited using a blue filter and green emission was collected at 1 frame per second. Functional TRPV1 channels were analyzed by exposing the explants to 1 μM capsaicin 8–18 hours after plating. 10 μM SB366791 was used to block capsaicin-mediated Ca^{2+} responses. For assessment of TRPV4 channels, at 8–24 hours neural crest cells explants were exposed to GSK1016790A (0.1 – 1000 nM). The TRPV4 inhibitor RN1734 (10 μM) was used to inhibit the GSK1016790A mediated response. Data were analyzed off-line with MetaMorph Image Analysis software. Changes in cytosolic Ca^{2+} were estimated as the relative increase of fluorescence intensity (F) from baseline fluorescence (F_0).

Temperature Activation Ca^{2+} Imaging Assays

Temperature-sensitivity of TRPV channels was examined at a hyperthermic (40°C) temperature in neural crest cells or HEK293T cells transfected with GCaMP6 and either TRPV1^{FeRIC} or TRPV1^{TFeRIC}, using Lipofectamine LTX Plus reagent. Twenty-four later, transfected cells were identified by expression of both GCaMP6 and TRPV1 channels (mCherry⁺). Primary GCaMP6⁺ chick neural crest cells were also generated as described above to examine the thermosensitivity of endogenous TRPV channels. For Ca^{2+} imaging, neural crest cells or HEK293T cells were placed on a recording chamber in the stage of Nikon-eclipse Ti-s inverted microscope. Cells were epi-illuminated using a blue filter and continuously perfused with the imaging buffer solution through a gravity-driven perfusion system (~2mL per min). Drugs were applied using same gravity system. The temperature was controlled with a CL-100 temperature controller (Warner Instruments) and a SC-20 dual In-Line Heater/Cooler (Harvard Apparatus). During experiments, temperature was monitored with a thermistor (TA-29, Warner Instruments) within the recording chamber.

Heat stimulus was delivered stepwise to increase the temperature from 25°C to 40°C over a 3-minute period. After removal of the heat stimulus, cells were exposed to TRPV channel agonists such as pH 5 or 1 μ M GSK1016790A. Neural crest cells were exposed to TRPV4 channel inhibitor 10 μ M RN1734 for 10 min and then imaged for changes in intracellular Ca²⁺ with heat or GSK1016790A exposure. GCaMP6 green emission was collected from cells at 1 frame per second. Images were analyzed offline using Nikon Element software; ROIs were placed over those cells which co-expressed GCaMP6 and TRPV channels, or over the GCaMP6 positive neural crest cells for endogenous TRPV channel analysis. GCaMP6 fluorescence intensity was measured for each image of the time-lapse acquisition. Data were analyzed using the Microcal Origin 7 software (OriginLab, MA). Responses are presented as F/F_0 and plots correspond to average \pm SEM.

Membrane integrity and cell viability assays

Membrane integrity assays were conducted in HEK293T cells *in vitro*. Membrane integrity was assayed using LIVE/DEAD Fixable Green Dead Cell Stain Kit (Life Technologies) following manufacturer's instructions. In this assay, the green dye is excluded from cells with intact membranes. However, cells with disrupted membranes, the dye accumulates in the cell and reacts with cytoplasmic proteins. Briefly, mCherry or TRPV1^{FeRIC} expressing HEK293T cells were exposed to either no RF, RF for 10 minutes, or 10 minutes at 55°C. After 5 minutes, cells were stained with manufacturer's green fluorescent amine reactive dye then fixed with 1% paraformaldehyde. Cells were analyzed using a 488nm line of an argon-ion laser and green fluorescence was detected in the green channel of the flow cytometer (530/30 nm).

Cell viability was measured with the CellTiter 96 Non-Radioactive Cell Proliferation Assay (MTT, Promega). HEK293T cells (40–50% confluent) transfected with TRPV 1^{FeRIC} or mCherry were exposed to RF for 10 minutes and cultured at 37°C for 30 hours before performing the assay according to the manufacturer's instructions. RF-exposed cells from each group was compared to transfected cells without RF exposure.

Immunoprecipitation

N-terminally FLAG tagged wild-type and FeRIC modified TRPV1 and TRPV4 were generated by PCR. HEK293T cells in a 6-well plate were transfected with 3 μ g of plasmid DNA using Lipofectamine PLUS reagent (Invitrogen). 8 hours after transfection, culture media was exchanged with fresh media containing 100 μ M FeCl₂ for endogenous ferritin induction. 18 hours later, protein was extracted and immunoprecipitation experiments were performed as described(67) with the following modification: CHAPS lysis buffer contained 5 mM CHAPS, 50 mM Tris (pH 7.4), 150 mM NaCl, 1 mM CaCl, 5% sucrose, 0.5% Triton X-100, protease inhibitor cocktail (Roche). Anti-FLAG M2 affinity gel (Sigma A2220) was used to immunoprecipitate FLAG tagged TRPV1 or TRPV4 constructs following the manufacturer's instructions. Samples were probed by western blot using antibodies against DDDDK (1:2,000, Abcam, ab1170) or FTH1 (1:1,000, Cell Signaling #3998).

Heart analysis

Hearts from the various treatment groups were harvested at day 10, photographed, fixed overnight in 10% formalin, paraffin embedded, serially sectioned and stained with hematoxylin and eosin and assessed for defects as previously described(41). Briefly, the hearts was scored as DORV if the aorta or the aortic portion of the outflow was not wedged between the tricuspid and mitral valve and shifted to the right in relation to the pulmonary trunk. All hearts categorized as DORV had a concomitant VSD. Hearts were classified as overriding aorta if the outflow VSD was positioned such that the aortic vestibule was in communication with the pulmonary infundibulum and the aorta was overriding the septal defect. In embryos with abnormal persistence of arch artery vessels, only vessels with patent lumens were scored. Outflow obstructions lesions were qualitatively assessed compared to stage-matched controls. Quantitative measurements were carried out as described below (see Stereology). Hearts from embryos with ectopia cordis were excluded form analysis because the etiology of the alignment defects cannot be distinguished between the experimental treatment or the open chest wall.

Stereology

Hematoxylin and eosin staining was performed on serial 10- μm sections through the entire heart from the ventricular apex through the aortic arch as previously described (41). The aortic and pulmonary valves were identified. For the aorta, we began with the last section containing coronary artery and moving superiorly, the lumens in 3–4 adjacent tissue sections were analyzed. For the pulmonary trunk, we identified the last section with pulmonary valve tissue attached to the vessel wall and analyzed 3–4 tissue sections above that. Analysis was conducted in bright field using the Cavalieri probe in Stereo Investigator 11 (MBF Biosciences, VT) on a Zeiss AxioImager2 microscope. Analysis parameters include: tissue cut thickness, 10 μm ; analysis objective, 10x; probe grid size, 50 μm x 50 μm ; grid orientation, randomized. The results are presented as the average luminal area in mm^2 for each vessel. The average coefficient of error (*CE*, Gundersen $m+1$) is a standard statistical value that is used extensively in the stereological studies. The definition of the *CE* is defined as the standard error of the mean of repeated estimates divided by the mean. An average *CE* was reported for each of the experiments within the individual figures and was less than 0.05.

MRI thermometry

MRI thermometry was used to measure internal heating in chicken eggs due to RF. This technique uses the temperature-dependent chemical shift of water to measure internal temperature changes during the time between two MRI scans of a stationary object(39). TRPV1^{FeRIC} was electroporated into the neural tube of chick embryos at HH9. Twenty-four hours later, eggs were imaged using a 7.0 T small-bore animal MRI scanner controlled by Agilent software version vnmr4 (Santa Clara, CA). A multi-echo gradient echo sequence was used to measure frequency maps. The sequence parameters were: field of view (FOV) = 7 \times 7 cm, matrix = 128 \times 128, repetition time (TR) = 200 ms, first echo time (TE1) = 5 ms, second echo time (TE2) = 10 ms, flip angle = 10°, slice thickness = 2 mm. Each scan took 20 s and was repeated every 10 minutes during 140 minutes, alternating 10 minute periods of RF and no RF. The first two echoes of each scan were used to create 14 maps that

measured changes in temperature during periods of RF and 14 maps that measured temperature changes without RF.

Anatomical MRI and heart visualization

Prior to MRI, embryo specimens were immersion fixed in 2.5 mM ProHance (Gadoteridol, Bracco Diagnostics Inc., Princeton, NJ) dissolved in 10% formalin. ProHance, a gadolinium contrast agent, was used to decrease the T_1 (longitudinal relaxation time) and to improve SNR(68). Images were acquired on a 9.4T system (400 MHz vertical bore Oxford superconducting magnet). The system consists of an 89-mm vertical bore magnet controlled by an Agilent VnmrJ 4.0 console (Agilent Technologies, Santa Clara, CA). The specimens were firmly secured in an acrylic specimen cartridge and immersed in Fomblin (Ausimont USA, Inc., Thorofare, NJ) to limit susceptibility artifacts at the tissue boundary. The cartridge was placed in a copper solenoid RF resonator (14-mm diameter, 21-mm length). A set of 3D multi-echo gradient echo images were acquired (10 echoes, $TE_1/\text{spacing}/TE_{10} = 3.9/7.6/71.9$ ms) with a field of view of $22 \times 11 \times 11$ mm³ and a matrix size of $768 \times 384 \times 384$, resulting in isotropic resolution of $28.6 \times 28.6 \times 28.6$ μm^3 . Final images were produced using a weighted sum of the individual echo images(69, 70).

One specimen was rotated to a position such that all four chambers of the heart were visible on a cross section. This position served as a reference. The coronal plane of this reference space was orthogonal to the atrial septum. Images were registered to this space using rigid body transformation, correlation cost function, and *sinc* interpolation (FMRIB Software Library, <http://www.fmrib.ox.ac.uk/fsl>). The heart wall, chambers, and vessels were segmented using seeded region growing in Avizo software (Visualization Sciences Group, Burlington, MA). The individual chambers (right atrium, right ventricle, left atrium, left ventricle) and vessels (pulmonary artery, aorta) were further segmented manually at the respective valves.

qPCR for cellular Fe²⁺ homeostasis

HEK293T cells were transfected with control mCherry, TRPV1^{WT}, TRPV1^{FeRIC}, TRPV4^{WT}, or TRPV4^{FeRIC}. Seventy-two hours after transfection, total RNA was extracted and cDNA was prepared using High Capacity cDNA RT (Applied Biosciences). qPCR was performed using SYBR green (IQ SYBR green Supermix; Biorad) on a Biorad C1000 Touch Thermal Cycler. mRNA expression was calculated using the C_t method and normalized to the relative expression of β -actin. mRNA levels for mCherry were adjusted to 1. Primers used have been previously published(71). Primer sequences are: *Fpn(F)* TTGCAGGAGTCATTGCTGCTA *Fpn(R)* TGGAGTTCTGCACACCATTGAT; *Fth(F)* CCATCAACCGCCAGATCAAC *Fth(R)* GCCACATCATCTCGGTCAAA; *Ftl(F)* GGGCCTCCTACACCTACCTC *Ftl(R)* CTCCTGGGTTTTACCCATT; *Slc11a2+1RE(F)* TGTTTGATTGCATTGGGTCTG *Slc11a2+1RE(R)* CGCTCAGCAGGACTTTTCGAG; *Tf1 ex 2,3(F)* TCAAGCCAGATCAGCATTCTC *Tf1 ex 2,3(R)* AGCCAGTTTCATCTCCACATG.

Avian craniofacial analysis

To visualize the development of the skeletal structures, day 10–11 chicks from the various treatment groups were fixed in 95% ethanol and processed for alcian blue and alizarin red staining as described by(72). Once cleared in glycerol, the length of the jaw structures (in mm) was measured using a dissecting microscope fitted with an ocular micrometer. The upper beak measurement extended from the quadratojugal to the tip of the upper beak. To ensure comparable staging across treatment groups, the upper beak length for each chick were normalized to individual femur lengths.

Zebrafish pharmacological treatment and imaging

We treated zebrafish larvae at 48 hours post fertilization (hpf) to assess the impact of TRPV4 stimulation on cranial neural crest cell convergent and extension. Zebrafish embryos were generated by natural pairwise matings of *-1.4coll1a1:egfp* transgenic adults (AB background)(31). Embryos were collected 30 minutes after spawning, stored at 28°C in embryo media (0.3g/L NaCl, 75mg/L CaSO₄, 37.5mg/L NaHCO₃, 0.003% methylene blue), and distributed randomly into separate petri dishes (50 embryos per dish). At 48hpf, embryos were treated with either 20uM GSK1016790A (GSK101) in 0.1% DMSO, 0.1% DMSO (vehicle control), or 20uM GSK1153218A, a compound that is structurally similar to GSK101 but lacks detectible activity (Supplemental Table. 1). We obtained ventral images of cartilaginous craniofacial structures at 3 and 4 days post fertilization (dpf) using the Vertebrate Automated Screening Technology (VAST; software version 1.2.2.8.) platform (Union Biometrica) as described(73). Embryos were collected after imaging and returned to their petri dish with fresh experimental media and stored at 28°C until the 4dpf imaging time point. Craniofacial defects were quantified by measuring the distance between Meckel's cartilage and the ceratohyal using ImageJ. All experiments were repeated at least 3 times.

Statistical analysis

All experiments unless otherwise noted were repeated a minimum of 3 times. Differences in categorical data sets (percentage of animals with congenital defects) were analyzed using Fisher's exact testing. Differences in continuous data sets were analyzed using a one-way ANOVA followed by Bonferroni post-hoc testing or unpaired t testing, where appropriate. Data are expressed as mean \pm SEM where applicable. A *P* value less than 0.05 was considered a statistically significant difference.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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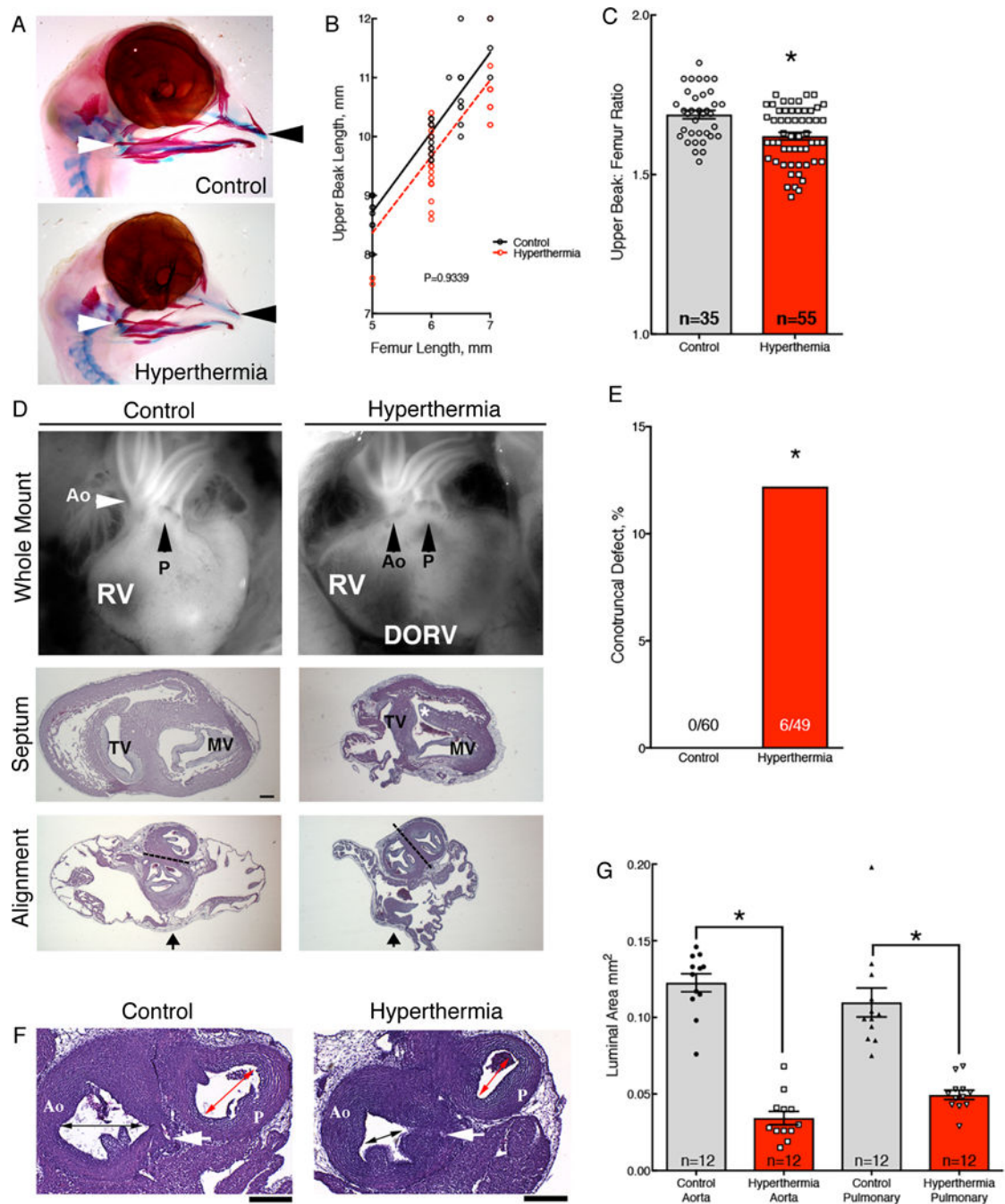


Fig. 1. Hyperthermia-induced congenital defects in neural crest-dependent tissues.

(A) Alcian blue and alizarin red stained craniofacial features of HH36 heads for control and hyperthermia-exposed chicks. Upper beak measurement extended from the quadratojugal (white arrowhead) to the tip of the upper beak (black arrowhead). (B) Upper beak length was normalized to femur lengths for control (black) and hyperthermia-exposed (red) chicks. (C) Graph of upper beak length to femur length ratios for control and hyperthermia chicks. (D) Whole mount and histological sections of HH36 hearts from control (normothermia) and hyperthermia-exposed embryos. The white and black arrows in the whole mount images

highlight the alignment of the aorta (Ao) and pulmonary trunk (P) in control heart compared to the hyperthermia-exposed heart. Histological sections of the whole mount hearts at the level of the ventricular septum and the outflow vessel semilunar valves. Dashed lines indicate the plane of outflow tract septation. Sections through the hyperthermia-exposed DORV heart with a perimembranous VSD (*) and a rightward shift of the aorta in relation to the pulmonary trunk (dashed line). Gross and histological analysis of heart anatomy was performed in 60 normothermia hearts and 49 hyperthermia-exposed hearts. Tricuspid valve (TV), mitral valve (MV). **(E)** Percentage of conotruncal defects in control (0%, n=60 hearts) and hyperthermia groups (12%, n=6/49 hearts). **(F)** Representative histological sections through the aorta and pulmonary trunk used to compare the luminal areas (arrows) of normothermic and hyperthermia-exposed hearts distal to the valves at the level of the left coronary artery (white arrowhead). Bar = 200 μ m. **(G)** Average Cavalieri probe estimates of the luminal cross-sectional areas of the aorta and pulmonary trunk immediately distal to the respective semilunar valves. Significance was determined using unpaired *t*-test(C,G) or Fishers exact test (E). *P<0.0001. The number of biological replicates is indicated by n in the graphs in (C, E, and G).

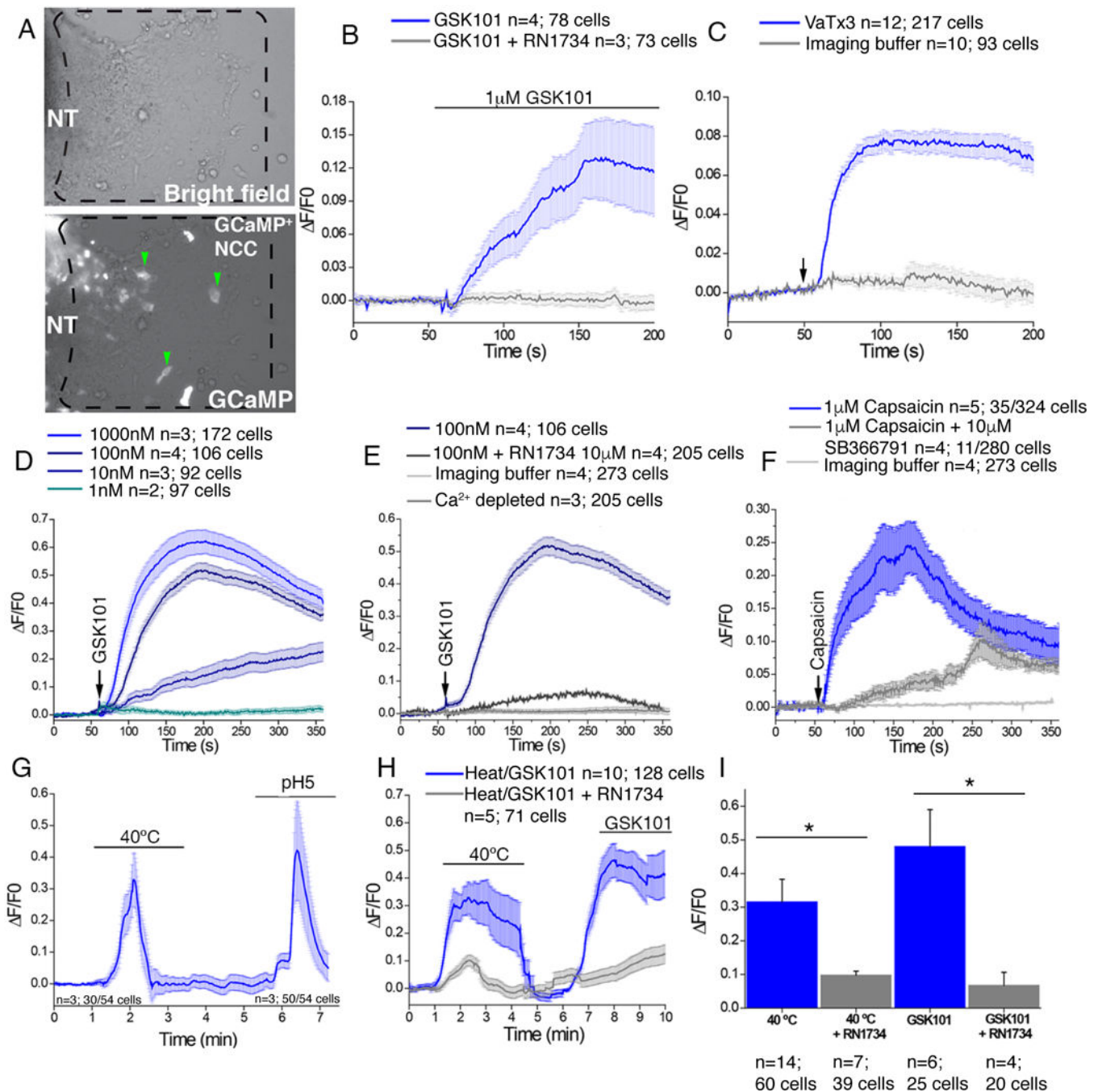


Fig. 2. Temperature-activated TRPV channels in avian and mammalian neural crest cells. (A) Images of chick primary explanted neural tubes (NT) expressing GCaMP6 after 24 hours in culture. Dashed box region shows migrating neural crest cells used in subsequent analyses (green arrowheads). (B) Change in GCaMP6 fluorescence in chick neural crest cells following exposure to 1 μ M GSK101 (blue) or 1 μ M GSK101 combined with 10 μ M RN1734 (gray line). (C) GCaMP6 fluorescence in chick neural crest cells following exposure to 1 μ M VaTx3 (arrow, blue line) or buffer alone (gray line). (D) Change in Fluo4 fluorescence in mouse primary explanted neural crest cells from E8.5 embryos in response

to the indicated doses of GSK101. **(E)** Fluo4 fluorescence in mouse primary explanted neural crest cells from E8.5 embryos in response to GSK101 (arrow, blue line) or to GSK101 and 10 μ M RN1734 (gray line). **(F)** Changes in Fluo4 fluorescence in mouse neural crest cells following exposure to 1 μ M capsaicin (arrow, blue line) or to capsaicin and SB366791 (gray line). **(G)** Representative GCaMP6 fluorescence in chick primary neural crest cells following exposure to 40°C imaging buffer followed by imaging buffer pH 5. **(H)** Representative GCaMP6 fluorescence in chick primary neural crest cells following exposure to 40°C imaging buffer or GSK101 (blue) or in the presence of RN1734 inhibitor (gray line). **(I)** Averages of more than 4 separate experiments as in (H) analyzing the number of cells/condition as indicated. Significance was determined using unpaired *t*-test. **P*<0.05. The number of biological replicates is indicated by *n* and/or the number of cells analyzed is indicated in the graphs in (B), (C), (D), (E), (F), (H), and (I).

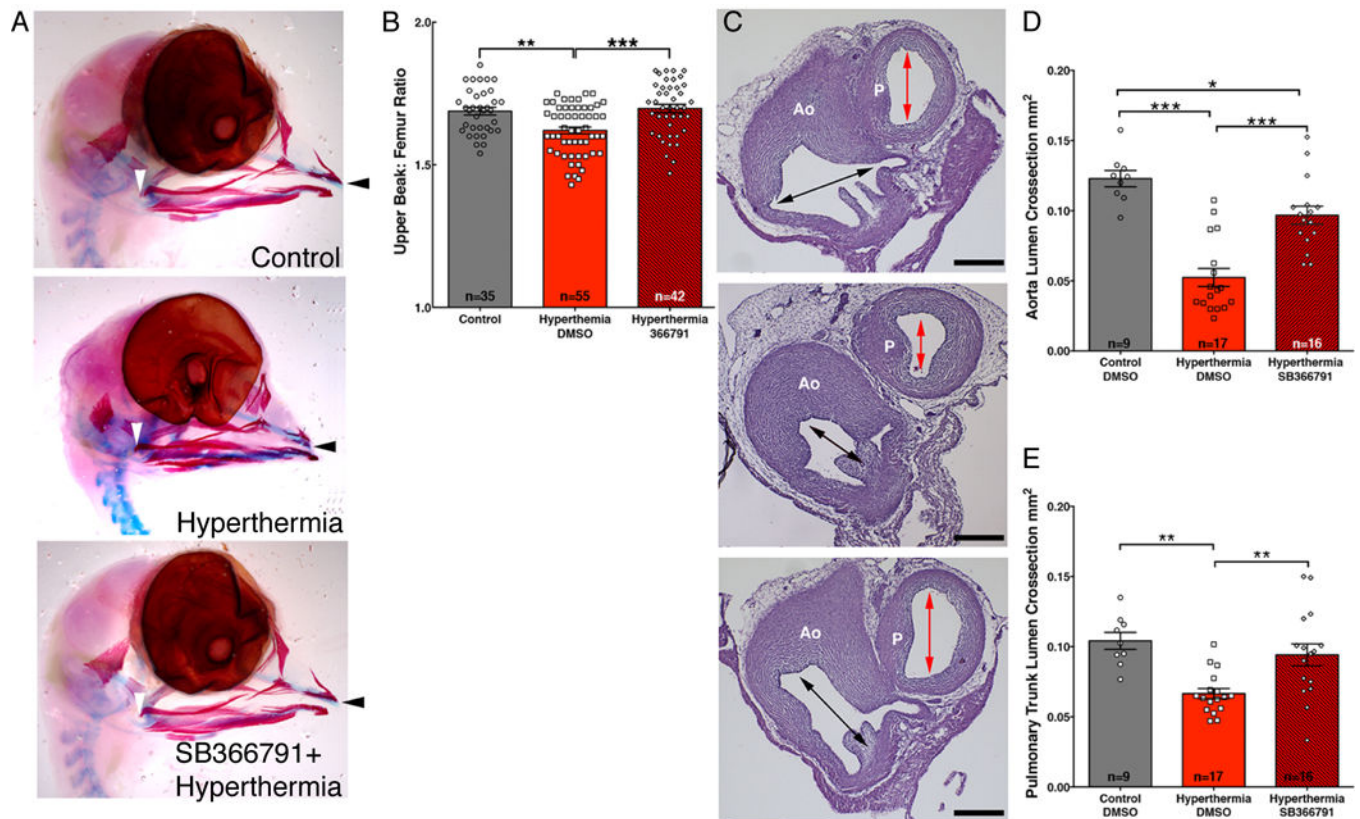


Fig. 3. TRPV1 inhibition rescues hyperthermia-induced congenital defects.

(A) Alcian blue and alizarin red stains of HH36 heads from hyperthermia-exposed and hyperthermia-exposed chicks following pretreatment with the TRPV1 inhibitor SB366791. Upper beak measurement extended from the quadratojugal (white arrowhead) to the tip of the upper beak (black arrowhead). (B) Comparison of upper beak length to femur length ratios in control, hyperthermia, or hyperthermia + SB366791 chicks. (C) Histological sections through the Aorta (Ao) and pulmonary trunk (P) comparing the luminal areas (arrows) of a control (top), hyperthermia (middle), and hyperthermia-exposed hearts with SB366791 pretreatment (lower) distal to the semilunar valves at the level of the left coronary artery (white arrowhead). Bar = 200 μ m. (D-E) Cavalieri probe estimates of the luminal cross-sectional area of the aorta (D) or pulmonary trunk (E) immediately above the aortic valve. Significance was determined by one-way ANOVA followed by Bonferroni's multiple comparisons test. * $P < 0.05$, ** $P < 0.004$, *** $P < 0.0001$. The number of biological replicates is indicated by n in the graphs in (B), (D), and (E).

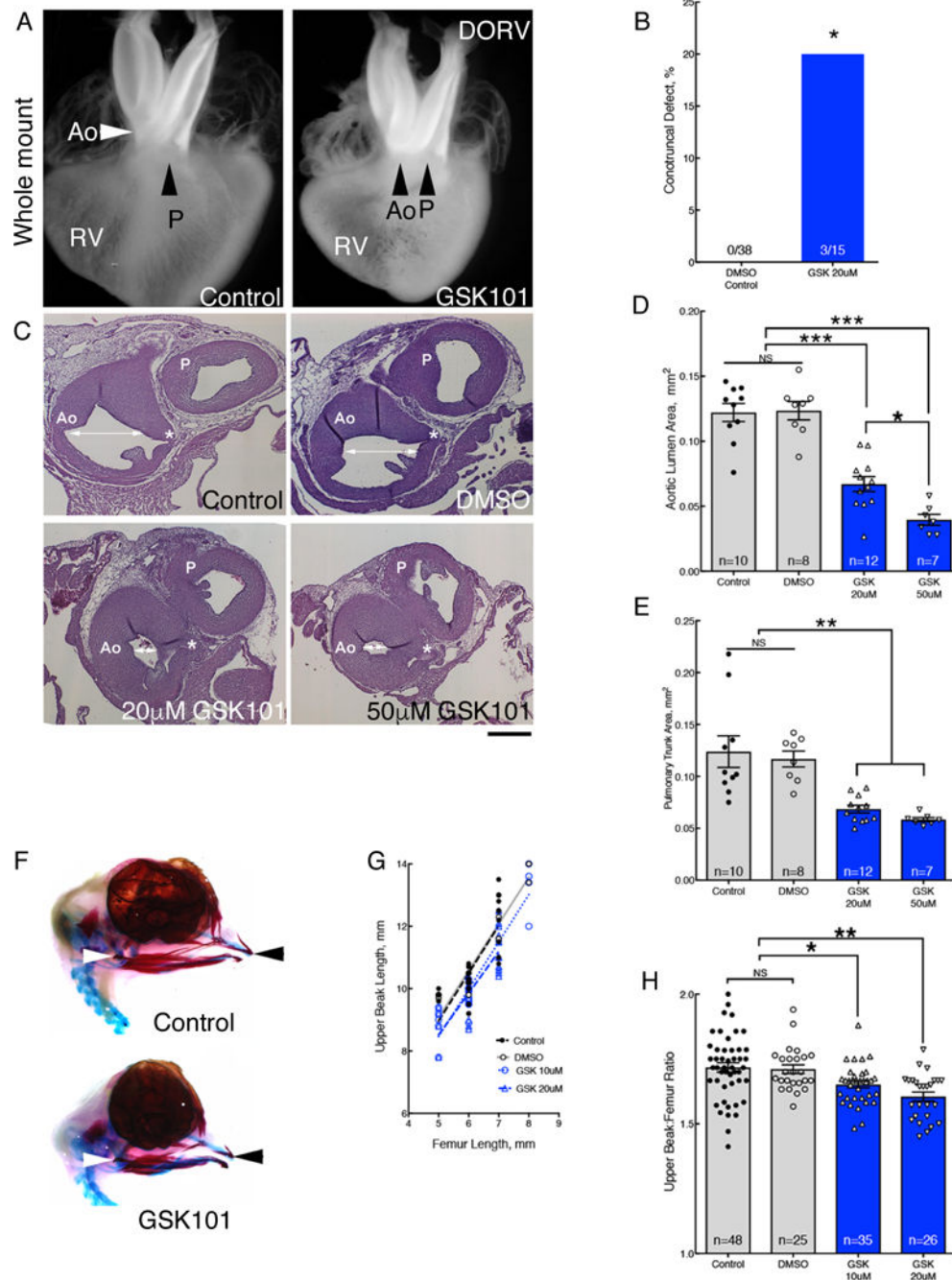


Fig. 4. Ligand activation of TRPV4 replicates hyperthermia induced birth defects.

(A) Whole mounts of hearts from control and GSK101 treated-embryos. GSK-treated heart showed DORV orientation of the aorta (Ao) and pulmonary trunk (P). See fig. S9 for histological sections. Whole mount and histological analysis of heart anatomy was performed in 38 hearts from DMSO-treated embryos and 15 GSK101-treated embryos. (B) Percentage of hearts with histologically confirmed conotruncal defects in GSK101 (GSK)-treated embryos compared to DMSO controls. (C) Panel of histological sections through the aorta (Ao) and pulmonary trunk (P) comparing the luminal areas (white arrows) at the level

of the left coronary artery (*) in control, DMSO, and 20 μ M or 50 μ M GSK101 treatment. Bar = 200 μ m. **(D,E)** Treatment with GSK101 reduced aortic luminal areas (D, average coefficient of error $(CE)=0.03$) and pulmonary trunk luminal areas (E, average $CE=0.03$). **(F)** Alcian blue and alizarin red stains of DMSO control (top) and 20 μ M GSK101 treated embryos (bottom) at HH36. **(G)** Normalization of upper beak length to femur lengths. **(H)** Graph of upper beak to femur ratios in untreated control, DMSO-treated control and 2 GSK101-treatment groups. Significance was determined using Fisher exact test (B) or one-way ANOVA followed by Bonferroni's multiple comparisons test (D, E, H). * $P<0.03$, ** $P<0.004$, *** $P<0.0001$. The number of biological replicates is indicated by n in the graphs in (D), (E), and (H).

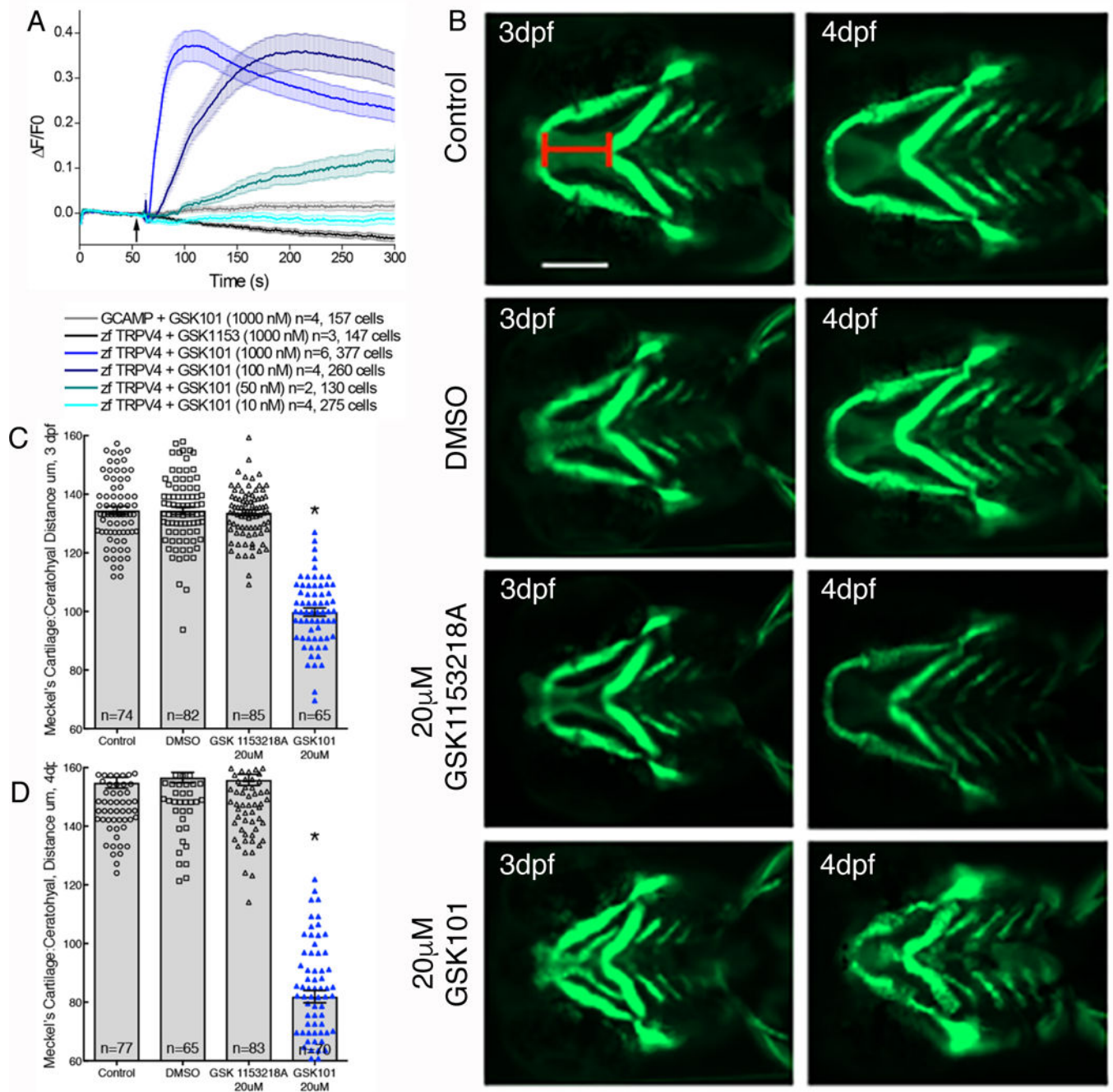


Fig. 5. TRPV4 activation disrupts jaw extension in zebrafish larvae.

(A) The activity of GSK101 at the indicated concentrations and of GSK1153218 (black tracing), a structurally related compound that lacks activity on mammalian TRPV4 (Table S1), in cloned zebrafish TRPV4 in CHO cells using GCaMP6 to assess Ca^{2+} permeability. (B) Representative images of *1.4cola1:egfp* transgenic zebrafish larvae treated with 20μM GSK101, vehicle control, or GSK1153218A. Replicate batches were imaged live at 3 and 4 dpf. Scale bar approximately 130μm. (C,D) Quantification of the distance between Meckel's cartilage and the ceratohyal was measured (red line in B). Significance was determined

using one-way ANOVA followed by Bonferroni's multiple comparisons test. * $P < 0.0001$, GSK101-treated group compared to all other groups). The number of biological replicates is indicated by n and/or the number of cells analyzed is indicated in the graphs in (A), (C), and (D).

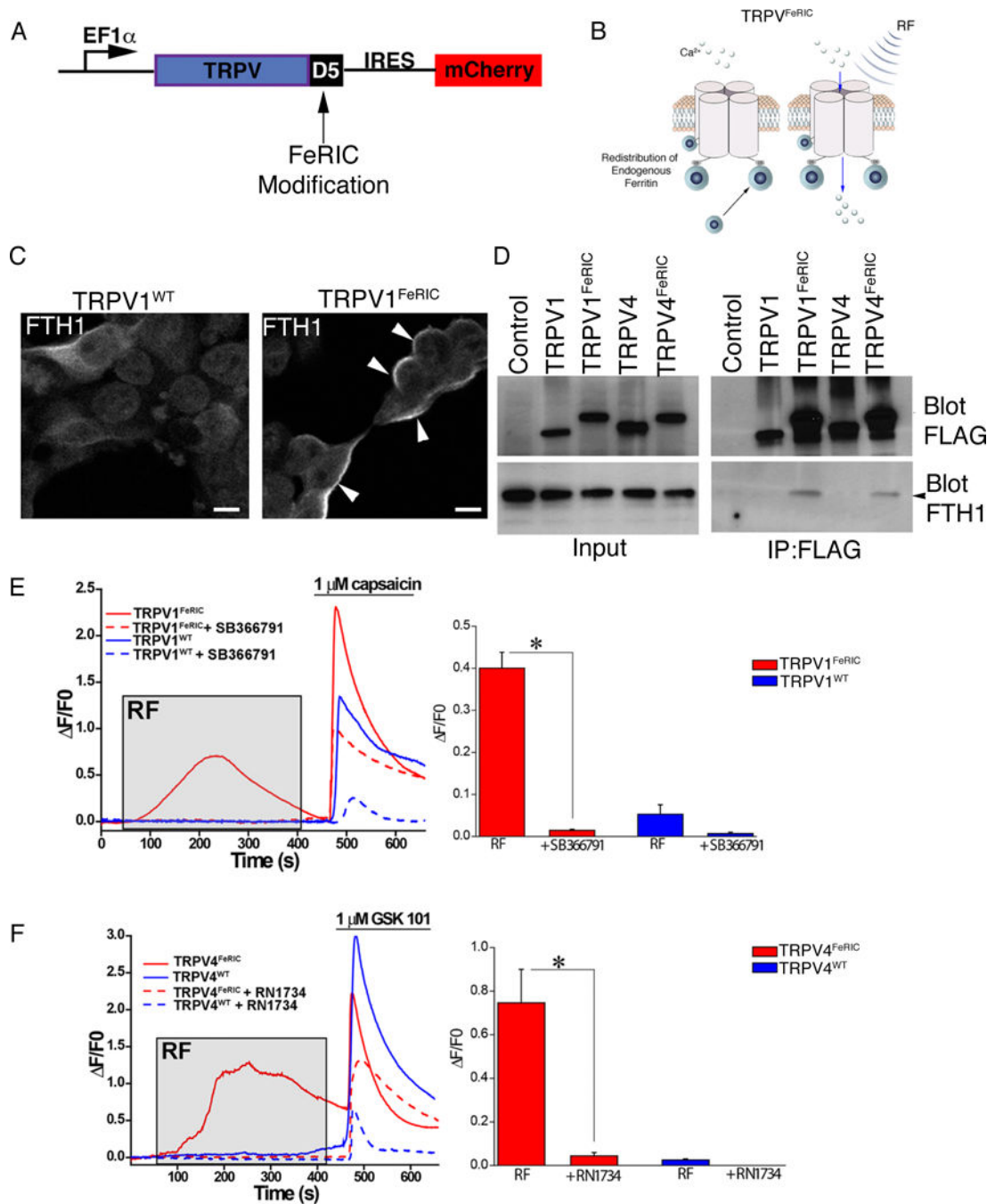


Fig. 6. Development of remotely controlled TRPV^{FeRIC} channels.

(A) TRPV channels were tagged with domain 5 of kininogen 1 (D5) and were cloned into the PLVX vector with an internal ribosomal entry site (IRES) for mCherry (fig. S12 and S13). (B) FeRIC channels were designed to recruit endogenous cellular ferritin to the modified TRPV channel at the cell membrane. (C) Cytoplasmic distribution of ferritin heavy chain fused with mCherry (FTH1^{mCherry}) in HEK293T cells expressing TRPV1^{WT} and membrane redistribution of FTH1^{mCherry} (white arrowheads) in TRPV1^{FeRIC}-expressing cells. Images were representative of 2 independent experiments. (D) Representative immunoprecipitation

western blot of HEK293T cells expressing FLAG tagged TRPV1^{WT}, TRPV1^{FeRIC}, TRPV4^{WT}, or TRPV4^{FeRIC}. The blot was probed for FLAG and FTH1. 4 independent experiments were conducted using TRPV1^{FeRIC} and 3 independent experiments were conducted using TRPV4^{FeRIC}. **(E)** GCaMP6 fluorescence in TRPV1^{WT}-expressing (blue) or TRPV1^{FeRIC}-expressing (red) HEK293T cells following RF (gray box) then 1 μ M capsaicin (bar). Bar graphs are DF/F0 averages of 4 experiments with 50–100 cells/group analyzed. **(F)** GCaMP6 fluorescence in TRPV4^{WT}-expressing (blue) or TRPV4^{FeRIC}-expressing (red) HEK293T cells following RF (gray box) then 1 μ M GSK101 (bar). Bar graphs are F/F0 averages of 5 experiments with 106–123 cells/group analyzed. Significance was determined using unpaired *t*-test. *P<0.05.

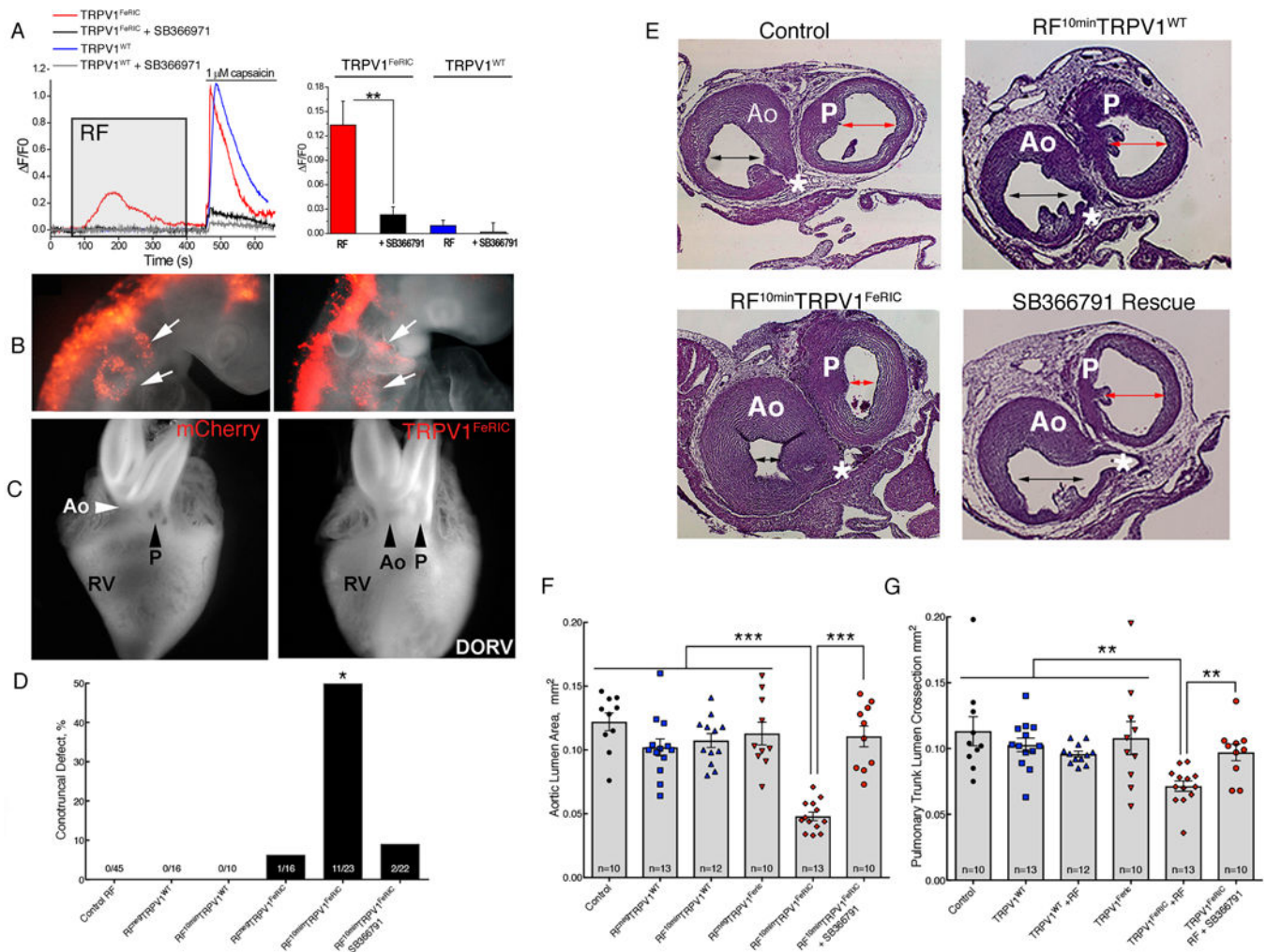


Fig. 7. Fever-associated heart defects following remote activation of TRPV1 in neural crest cells. (A) GCaMP6 fluorescence in primary chick neural crest cells electroporated with TRPV1^{WT} (blue lines) or TRPV1^{FeRIC} (red lines) in the absence or presence of the TRPV1 inhibitor SB366971 (gray and black lines) following RF (gray box) then 1 μ M capsaicin (bar). Bar graph shows cumulative responses representing 3–4 separate experiments with 22–138 cells/group analyzed. (B) mCherry⁺ neural crest streams migrating to the pharyngeal arches (white arrows) in HH14 control or TRPV1^{FeRIC} electroporated embryos. 6 independent experiments. (C) Whole-mounted hearts from RF^{Neg}TRPV1^{FeRIC} embryo and RF^{10min}TRPV1^{FeRIC} embryo with arrows noting the position of the aorta (Ao) and pulmonary trunk (P) with respect to the right ventricle (RV). 5 independent experiments. See Supplemental Movie 1 for an MRI reconstruction of a RF^{10min}TRPV1^{FeRIC}-induced DORV heart defect in chick. (D) Percent of embryos with histologically confirmed conotruncal heart defects within indicated groups at HH36. (E) Representative histological sections through the aorta and pulmonary trunk at the level of the coronary artery (*) in Control RF, RF^{10min}TRPV1^{WT}, RF^{10min}TRPV1^{FeRIC}, and RF^{10min}TRPV1^{FeRIC} with SB366971 pretreatment. The double arrowheads highlight the luminal areas of the aorta and pulmonary trunk estimated using the Cavalieri probe. Bar = 200 μ m. (F) Graph of the

Cavalieri probe estimates of cross sectional areas through the aorta at the level of the coronary artery in the indicated treatment groups ($CE=0.03$). (G) Graph of the Cavalieri probe estimates of the crosssectional areas through the pulmonary trunk distal to the semilunar valve in the indicated treatment groups ($CE=0.03$). Significance was determined using unpaired *t*-test (A), Fisher exact test (D) or one-way ANOVA followed by Bonferroni's multiple comparisons test (F,G). * $P<0.02$, ** $P<0.005$, *** $P<0.0001$. The number of biological replicates is indicated by n in the graphs in (F) and (G).

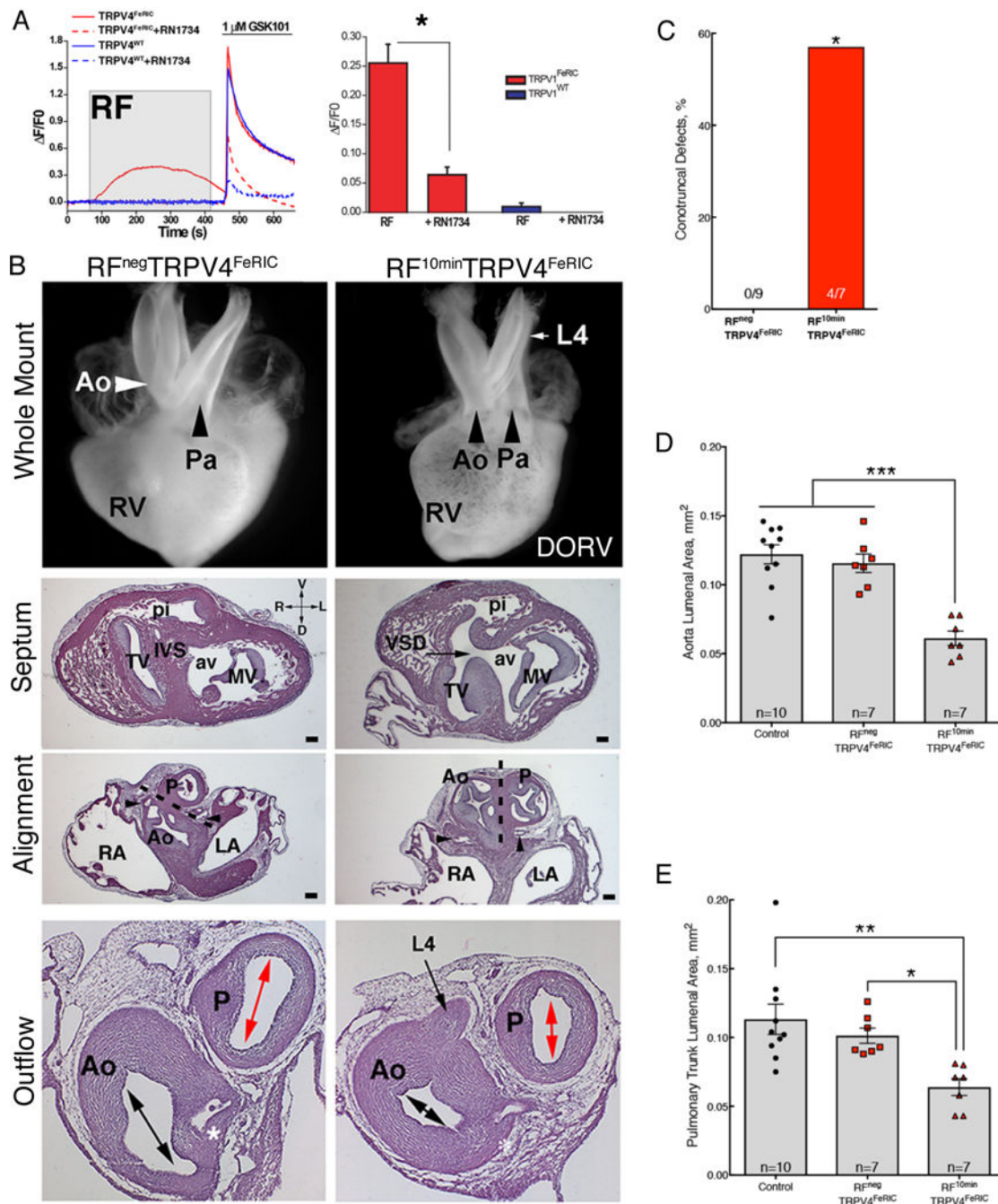


Fig. 8. Fever-associated heart defects following remote activation of TRPV4 in neural crest cells. (A) GCaMP6 fluorescence in chick neural crest cells electroporated with TRPV4^{WT} (blue lines) and TRPV4^{FeRIC} (red lines) following RF (gray box) then GSK101 (bar). TRPV4 inhibitor RN1734 inhibits response (dashed lines). Bar graph shows cumulative responses representing 4 separate experiments with 45–49 cells/group analyzed. (B) Whole mount and histological sections of HH63 hearts from RF^{Neg}TRPV4^{FeRIC} embryo and RF^{10min}TRPV4^{FeRIC} embryo with DORV and a persistent L4 arch artery. The white and black arrows in the whole mount images highlight the alignment of the aorta (Ao) and

pulmonary trunk (P) in the $RF^{NegTRPV4^{FeRIC}}$ and $RF^{10minTRPV4^{FeRIC}}$ hearts. Histological sections of the whole mount in $RF^{negTRPV4^{FeRIC}}$ and $RF^{10minTRPV4^{FeRIC}}$ hearts (above) at the level of the ventricular septum (IVS), at the level of the semilunar valves of the aorta (coronary arteries, black arrowheads) and pulmonary trunk and more distally through the smooth muscle walls of the aorta and pulmonary trunk at the level of the left coronary artery (*). The dashed line indicates the plane of outflow tract septation. Black arrows indicate a VSD and persistent L4 arch artery in the section through the $RF^{10minTRPV4^{FeRIC}}$ heart. Double headed arrows indicate the luminal cross sectional areas of the aorta and pulmonary trunk measured in the Cavalieri estimates in (D-E). Bar = 200 μ m. 3 separate experiments were performed and a total of 9 $RF^{NegTRPV4^{FeRIC}}$ and 7 $RF^{10minTRPV4^{FeRIC}}$ hearts were analyzed. Tricuspid valve (TV), mitral valve (MV), aortic vestibule (av), pulmonary infundibulum (pi) right atrium, (RA), and left atrium (LA). (C) Percentage of histologically confirmed conotruncal defects in $RF^{NegTRPV4^{FeRIC}}$ embryos compared to $RF^{10minTRPV4^{FeRIC}}$ embryos. (D) Graph of the Cavalieri probe estimates of cross sectional areas through the aorta at the level of the coronary artery in the indicated treatment groups ($CE=0.05$). (E) Graph of the Cavalieri probe estimates of the cross-sectional areas through the pulmonary trunk distal to the semilunar valve in the indicated treatment groups. ($CE=0.05$). Significance was determined using unpaired *t*-test (A), Fisher exact test (C) or one-way ANOVA followed by Bonferroni's multiple comparisons test (D,E). * $P<0.02$, ** $P<0.005$, *** $P<0.0001$. The number of biological replicates is indicated by n in the graphs in (C, D and E).