

natriuretic peptides are potential therapeutic targets for heart disease (9–14).

Mice transgenic for various loci, including the 5'-flanking regions of the natriuretic peptide genes, have been used to identify the regulatory elements required for transcriptional activation either during heart development or in the diseased heart. These studies reported that the 5'-flanking regions of the natriuretic peptide genes regulated their expression during heart development (9, 10, 13); however, the 5'-flanking regions were not responsible for their specific reactivation in the diseased heart (11, 12). A recent study identified the distal enhancer elements regulating the natriuretic peptide genes in the developing heart by examining cardiac-specific transcription factor binding sites; however, these enhancer elements did not respond to heart failure (14). Therefore, the stress-responsive regulatory elements that function during heart failure have not yet been identified and are potentially located outside the 5'-flanking regions.

In this study, we aimed to identify the novel stress-responsive enhancer elements of the *Nppa* and *Nppb* genes in the failing heart. Furthermore, we established a noninvasive and quantitative live imaging assay to monitor the transcriptional activity of candidate enhancers in the failing heart. *In vivo* live imaging of the firefly luciferase reporter in a single mouse enabled us to analyze the sequential changes in enhancer activity during the progression of heart failure. Combined with a fine mapping technique using epigenetic markers, we identified a 650-bp stress-responsive enhancer that was strongly activated by cardiac hypertrophy and heart failure.

MATERIALS AND METHODS

Animals

All procedures were performed according to the U.S. National Institutes of Health (NIH) Guide for the Care and Use of Laboratory Animals (NIH publication no. 85-23, revised 1996) and were approved by the Animal Experiments Committee, Osaka University (approval no. 21-78-10).

Reagents and antibodies

Phenylephrine (PE) was purchased from Sigma-Aldrich (St. Louis, MO, USA). The anti-RNA polymerase II and anti-histone H3 trimethylated at lysine 4 (H3K4me3) antibodies used for chromatin immunoprecipitation sequencing (ChIP-seq) were kind gifts from Dr. H. Kimura (Graduate School of Frontier Biosciences, Osaka University).

Primary culture of neonatal rat cardiomyocytes

Ventricular myocytes obtained from 1- or 2-d-old Wistar rats were prepared and cultured overnight in Dulbecco's modified Eagle's medium (Sigma-Aldrich) containing 10% FBS, as described previously (15).

Comparative genomics

Genome-wide multiple alignments of the genomic sequences containing the *Nppa* and *Nppb* genes were performed using the University of California Santa Cruz (UCSC) Genome Browser (16); 8 vertebrate species were compared, including mouse (mm9, July 2007), rat (m4, Nov. 2004), human (hg18, Mar. 2006), orangutan (ponAbe2, July 2007), dog (canFam2, May 2005), horse (equCab1, Jan. 2007), opossum (monDom4, Jan. 2006), and chicken (galGal3, May 2006). We used vertebrate Multiz alignment of DNA sequences (17) to analyze the homology of DNA sequences among mouse and other species. We used the Placental Mammal Base-wise Conservation assessed by PhyloP (18) to assess the degree of mammalian conservation. Next, we identified discrete conserved fragments. The transcribed sequences within the conserved set were filtered out using known genes, spliced ESTs, and mRNA annotations obtained from the UCSC genome browser. Finally, we manually curated the data set to remove any additional false positives by visual examination of the UCSC genomic data. We defined the noncoding conserved regions (CRs) that were homologous at least in the human and mouse genomes and at least 1 kb away from the transcription start sites as the enhancer candidates.

ChIP sequencing on mouse heart tissues

Whole hearts were isolated from 8-wk-old C57BL6 mice, perfused rapidly with cold PBS, flash-frozen in liquid nitrogen, homogenized using a sterile tissue grinder, and cross-linked with 0.3% paraformaldehyde. Subsequently, chromatin isolation, sonication, and immunoprecipitation using an anti-RNA polymerase II antibody and an anti-H3K4me3 antibody were performed. The ChIP DNA and input samples were sheared by sonication, end-repaired, ligated to the sequencing adapters, and amplified. The purified ChIP DNA library samples were sequenced using the Illumina Genome Analyzer II (Illumina, Inc., San Diego, CA, USA). Unfiltered sequence reads were aligned to the mouse reference genome [U.S. National Center for Biotechnology Information (NCBI) build 37, mm9] using Bowtie. RNA polymerase II- and H3K4me3-enriched regions were identified using MACS (19) with the default parameters.

Lentiviral enhancer assay

Eleven CRs were PCR amplified from the mouse BAC clone containing the *Nppa* and *Nppb* loci (clone RP23-128E8; BACPAC Resources Center, Children's Hospital Oakland, Oakland, CA, USA); primers and probes are listed in Supplemental Table S1). The PCR fragments were subcloned into the pCR-Blunt II-TOPO vector (Invitrogen, Carlsbad, CA, USA) and recombined into a lentiviral vector encoding the firefly luciferase reporter (pGreenFire Transcriptional Reporter Lentivector; System Biosciences, Mountain View, CA, USA). The lentiviral particles were produced by transfection of 293T cells with the 3 lentiviral packaging plasmids (*i.e.*, pMDLg/pRRE, pRSV-Rev, and pMD2.VSV.G) using Lipofectamine 2000 (Invitrogen). The supernatant from 293T cells containing the lentiviral particles was collected 48 h after transfection, sterilized using a 0.45- μ m cellulose acetate filter, and concentrated by centrifugation (Peg-it Virus Precipitation Solution, System Biosciences).

Rat neonatal cardiomyocytes isolated as described above were plated in 96-well plates. The next day, the medium was replaced with a serum-free medium containing the lentiviral vector, and the cells were incubated for 12 h. Subsequently, the cardiomyocytes were exposed to 100 μ M PE for 48 h prior to the luciferase assay.

RNA extraction and quantitative RT-PCR

The total RNA was prepared from rat cardiomyocytes, rat cardiac fibroblasts, murine hearts, and murine brains using the RNA-Bee RNA isolation reagent (Tel-Test, Friendswood, TX, USA) and then converted to cDNA using the high-capacity cDNA reverse transcription kit (Applied Biosystems, Foster City, CA, USA), according to the manufacturer's instructions. The quantitative RT-PCR was performed using the TaqMan technology and the StepOnePlus real-time PCR System (Applied Biosystems). All samples were processed in duplicate. The level of each transcript was quantified according to the threshold cycle (C_t) method using GAPDH as an internal control. Inventoried TaqMan gene expression assays were used: *Nppa*, Rn0056661, Mm01255748; *Nppb*, Rn00580641, Mm01255770; *Gapdh*, rodent GAPDH control reagent.

3C analysis

The whole hearts of the mice were isolated, perfused rapidly with cold PBS, flash-frozen in liquid nitrogen, homogenized using a sterile tissue grinder, and fixed with 1% paraformaldehyde. The cross-linked tissues utilized for 3C experiments were subjected to digestion with *Bam*HI following standard protocols (20, 21). The mouse BAC DNA containing *Nppa* and *Nppb* (clone RP23-128E8) was used as a control. The TaqMan real-time PCR was performed using probes near the restriction sites; the primers and probes are listed in Supplemental Table S2.

Transgenic mouse enhancer assay

The candidate enhancer regions were cloned into a vector encoding the minimal CMV promoter driving the luciferase gene as described above. Transgenic mouse embryos were generated by pronuclear injection into the zygotes of BDF1 mice using standard methods. Because black fur attenuates light transmission, albino mice were generated by crossing the transgenic founders to ICR albino mice.

In vivo bioluminescence imaging

Prior to *in vivo* imaging, the mice were anesthetized using isoflurane, and the black mice were shaved from the neck to the lower torso to allow the optimal visualization of fluorescence without interference from the black fur. A D-luciferin solution was injected intraperitoneally (150 mg/kg i.p.) or intravenously (75 mg/kg i.v.). The mice were imaged using an *in vivo* live imaging system (IVIS Lumina II; Caliper Life Sciences, Waltham, MA, USA). For quantification, the bioluminescence light intensity was measured at the region of interest and expressed in relative light units (RLU/min) using Living Image 4.0 (Caliper Life Sciences). To calculate the enhancer activity in the heart, we defined the ratio of heart to brain luciferase intensities as the cardiac-specific enhancer activity.

Transverse aortic constriction (TAC)

Transgenic mice aged 8 wk and weighing 20–25 g were subjected to pressure overload, as described previously (22). Briefly, the chest was entered *via* the second intercostal space at the upper left sternal border. After the arch of the aorta was isolated, a TAC was created using a 7-0 suture tied twice around a 27-gauge needle and the aortic arch, between the innominate and left common carotid arteries. After the

suture was tied, the needle was gently removed, yielding 60–80% constriction of the aorta.

PE-induced hypertrophy

Transgenic mice aged 8 wk and weighing 20–25 g were treated with PE (75 mg/kg/d) using an osmotic minipump (Alzet, Cupertino, CA, USA) to induce cardiac hypertrophy, as previously reported (23, 24).

Statistical analysis

Data are expressed as means \pm SE. The 2-tailed Student's *t* test and repeated ANOVA were used to analyze differences between the groups. Values of $P < 0.05$ were considered to represent a significant difference.

RESULTS

Identification of candidate enhancers near the *Nppa-Nppb* locus using comparative genomics and ChIP-seq

To identify potential enhancers, we performed a comparative analysis of the genomic sequences of mouse and divergent species and identified CRs that may function as common regulatory sequences (25–27). We defined CRs that were homologous at least in the human and mouse genomes and at least 1 kb away from the transcription start sites of *Nppa* and *Nppb* as the candidate enhancers. First, we analyzed the 50-kb *Nppa-Nppb* locus bounded by the binding sites of 2 CCCTC-binding factors (CTCFs), which can function as insulators (28, 29). Using a genome database (30), we identified 11 CRs, including the *Nppa* and *Nppb* introns in the 50-kb region (Fig. 1).

Next, we performed a ChIP-seq analysis on RNA polymerase II and H3K4me3 in the adult mouse heart. We analyzed the epigenetic modifications near the *Nppa* and *Nppb* genes combined with the ChIP-seq analysis using a public database of the adult mouse heart (30). We hypothesized that the normal heart would have activated epigenetic marks because *Nppa* and *Nppb* are expressed, albeit at low levels, in normal conditions. Recent genome-wide studies have determined that enhancers can be defined as DNA sequences bound by the RNA polymerase II and transcriptional coactivator protein p300, and where histone H3 monomethylated at lysine 4 (H3K4me1) accumulates instead of H3K4me3 (31–34). Among the 11 CRs identified, only CR9 coincided with the binding sites of RNA polymerase II and p300, and overlapped with the gene areas modified by H3K4me1, and filled all criteria for the enhancer (Fig. 1). In addition, H3K4me1 modifications in CR9 were only observed in the heart but not in the other organs (Fig. 1 and Supplemental Fig. S1). Therefore, we analyzed the 11 CRs, including CR9, as the most likely distal candidate enhancers for the stress-responsive regulatory regions of the natriuretic peptide genes.

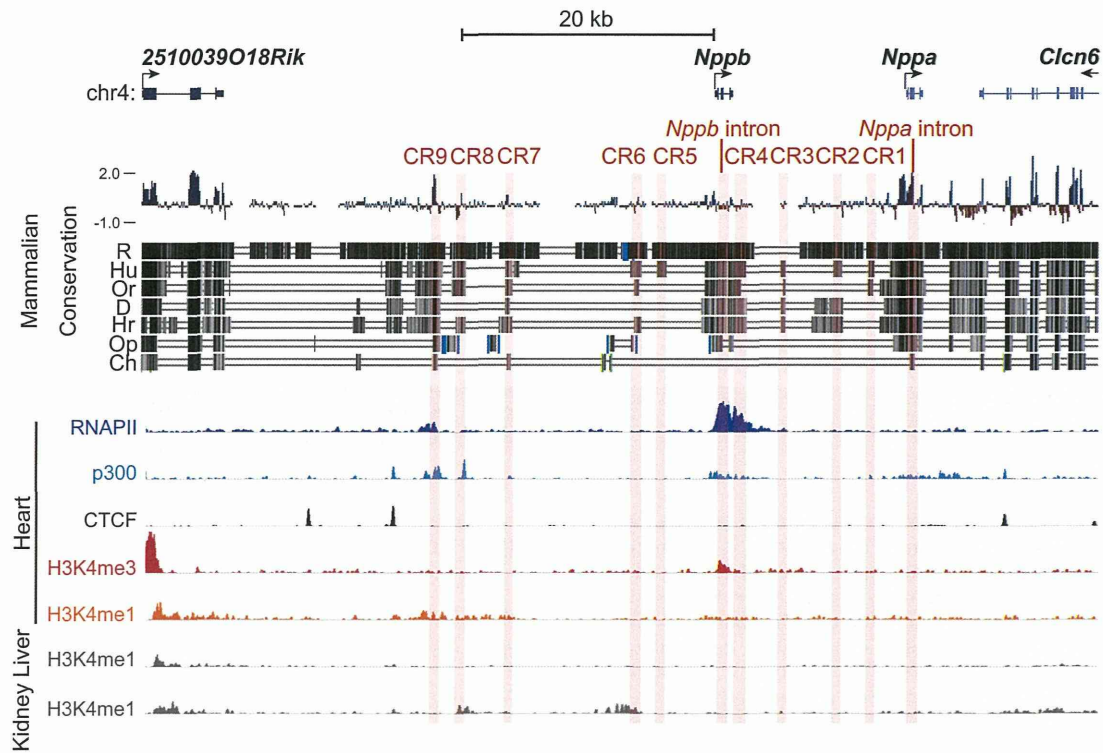


Figure 1. Mammalian evolutionarily conserved regions and ChIP-seq data surrounding the murine *Nppa* and *Nppb* loci. We used an open database on the University of California Santa Cruz (UCSC) Genome Browser to assess the degree of DNA sequence conservation around *Nppa* and *Nppb* gene loci. Blue and red vertical lines, the Placental Mammal Basewise Conservation assessed by PhyloP; black vertical lines, the vertebrate Multiz alignment of DNA sequences among mice and 7 other species (rats, humans, orangutans, dogs, horses, opossums, and chickens). We defined noncoding conserved regions (CRs) that were homologous at least in the human and mouse genomes and at least 1 kb away from the transcription start sites of *Nppa* and *Nppb* as the candidate enhancers. CRs are highlighted as light red vertical bars (CR1-9, *Nppa* intron, and *Nppb* intron). ChIP-seq data for H3K4me1, p300, and CTCF were obtained from an open database of the adult mouse heart. Some CRs coincided with the peaks for H3K4me1, RNA polymerase II, and the transcriptional coactivator protein p300. R, rat; Hu, human; Or, orangutan; D, dog; Hr, horse; Op, opossum; Ch, chicken.

Identification of a distal enhancer element responsive to an α_1 -adrenergic receptor agonist

We screened the candidate enhancers for potential stress-responsive regulatory regions. We analyzed the enhancer activity of these 11 CRs after treatment with PE, an α_1 -adrenergic receptor agonist, which mimics cardiac overload and induces *Nppa* and *Nppb* expression in cardiomyocytes (35). We confirmed that PE induced the expression of endogenous *Nppa* and *Nppb* specifically in cardiomyocytes but not in cardiac fibroblasts (Fig. 2A). Then, we introduced the 11 CRs with a minimum human cytomegalovirus (CMV) promoter and the luciferase gene into rat cardiomyocytes using a lentiviral vector system.

Among the 11 CRs tested, only CR9, which is located 22 kb upstream from the *Nppb* transcription start site and shows high mammalian conservation score in the Placental Mammal Basewise Conservation by PhyloP (Fig. 2B), reproducibly increased the PE-induced luciferase activity by ~ 5 -fold compared to the minimal CMV promoter alone (Fig. 2C). However, CR9 did not respond to PE in cardiac fibroblasts (Fig. 2C). These

results suggest that CR9 is the regulatory element that is responsive to PE specifically in cardiomyocytes.

Long-range physical interaction between the distal enhancer element and the proximal promoters of the *Nppa* and *Nppb* genes

Confirming the looping interactions between distal elements and promoters is one way to demonstrate the transcriptional regulatory activity of distal elements. We performed a 3C assay (20) to comprehensively investigate whether the genomic region containing CR9 moved closer to the *Nppa* or *Nppb* promoter in an adult murine heart treated with a continuous infusion of PE *in vivo*.

The ligation frequencies were quantified by TaqMan real-time PCR using specific primers and probes and were compared to the ligation frequency of noncross-linked *Bam*HI-digested BAC DNA containing the *Nppa*-*Nppb* locus. We observed that CR9 interacts with both the *Nppa* and *Nppb* promoter regions at a higher frequency relative to other gene areas (Fig. 2D); furthermore, PE treatment strengthened these interac-

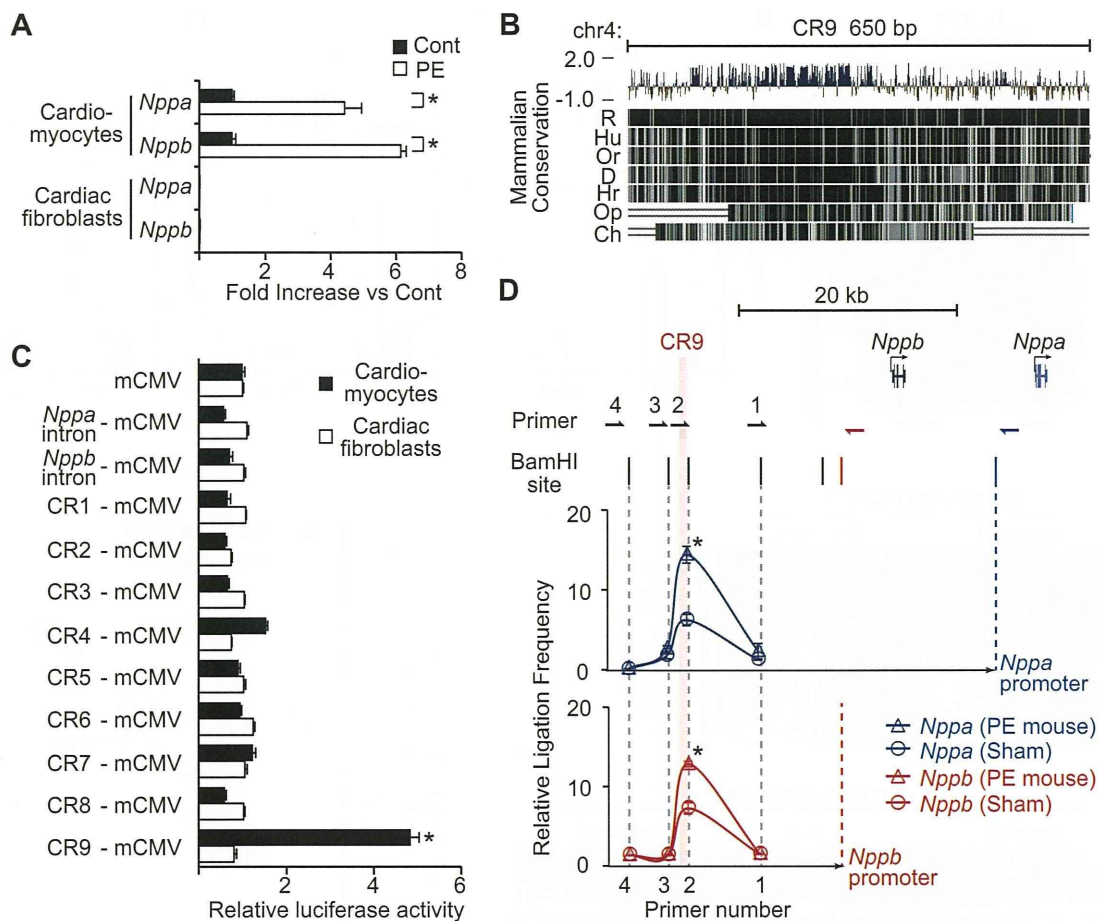


Figure 2. Identification of a distal enhancer element that is responsive to an α_1 -adrenergic receptor agonist. *A*) Relative transcript levels of *Nppa* and *Nppb* in rat neonatal cardiomyocytes and cardiac fibroblasts 48 h after treatment with PE (100 μ M). Values are means \pm SE ($n=3$ cultures). * $P < 0.01$ vs. control; t test. *B*) CR9 is a highly conserved genomic region in vertebrates. *C*) Relative luciferase reporter activities of CRs in rat neonatal cardiomyocytes and cardiac fibroblasts 48 h after treatment with PE (100 μ M). PE-induced luciferase activity driven by the mCMV promoter was defined as 1. Values are means \pm SE ($n=5$ cultures). * $P < 0.001$ vs. mCMV alone; t test. *D*) *In vivo* 3C analysis of the murine *Nppa* and *Nppb* loci, showing relative ligation frequencies of each primer to the *Nppa* promoter (blue triangle, mouse with PE treatment; blue circle, mouse without PE) and the *Nppb* promoter (red triangle, mouse with PE treatment; red circle, mouse without PE). Vertical bars and arrows show the positions of *Bam*HI sites and primers. Data were normalized to the amplification value of a *Bam*HI-digested and religated BAC clone, which included the *Nppa* and *Nppb* loci (means \pm SE; $n=2$ hearts). R, rat; Hu, human; Or, orangutan; D, dog; Hr, horse; Op, opossum; Ch, chicken. * $P < 0.05$ vs. control; t test.

tions (Fig. 2D). These results suggest that there is a close proximity between the distal genomic region containing CR9 and the proximal promoters of the *Nppa* and *Nppb* genes in the PE-induced hypertrophic heart.

Establishment of an *in vivo* live imaging system for gene expression in a murine model of heart disease

We confirmed the activity of the newly identified enhancer CR9 in the heart *in vivo*. The conventional histological evaluation of LacZ reporter expression in the heart only provides data at a single time point; therefore, this method cannot be employed for kinetic assessments or time course analyses of reporter expression in a live heart.

To overcome this difficulty, we established a nonin-

vasive and quantitative live imaging system that allowed real-time monitoring of the firefly luciferase reporter. We generated 3 transgenic mouse lines (Tg-line1, Tg-line2, and Tg-line3) in which the CR9 enhancer element and a minimal CMV promoter driving the luciferase reporter gene were introduced into the germline. The live-imaging system detected luciferase expression in the heart, brain, and intestine of the Tg-line1 (Fig. 3A), in the heart, salivary glands, and skin of the Tg-line2 (Supplemental Fig. S2A), and in the heart of the Tg-line3 (Supplemental Fig. S2E).

To identify the organs in which CR9 functioned as a stress-responsive enhancer, we examined the luciferase reporter expression in each organ by quantitative PCR. Continuous infusion of PE increased the blood pressure and resulted in cardiac hypertrophy (24, 36). The

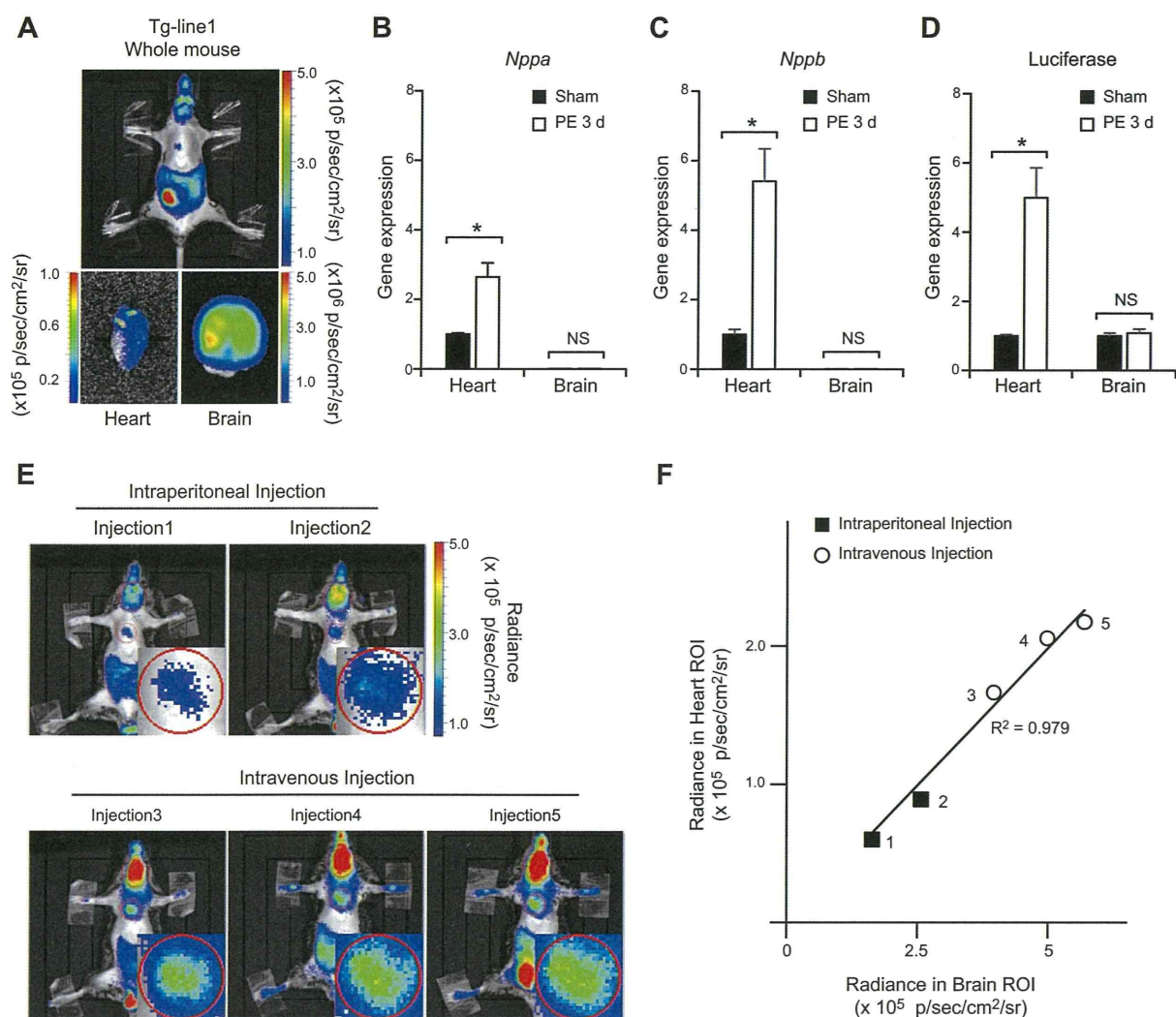


Figure 3. Establishment of an *in vivo* live imaging system for enhancer activity. *A*) Chemiluminescence imaging of CR9 in a mouse of Tg-line1. Top panel: result from whole-animal *in vivo* live imaging. Bottom panels: chemiluminescence images of the heart and brain in the same mouse. *B*, *C*) Relative transcript levels of *Nppa* and *Nppb* in the ventricular myocardium and brain of CR9 Tg-line1 mice treated with continuous infusion of PE for 3 d. Average transcript level in the ventricular myocardium of preinfused mice was defined as 1 (means \pm SE; $n=5$ hearts). $*P < 0.01$ vs. sham-infused mice; *t* test. *D*) Relative transcript levels of luciferase reporter in the ventricular myocardium and brain of the CR9 Tg-line1 mice continuously infused with PE for 3 d. Average transcript level in the ventricular myocardium and brain of preinfused mice was defined as 1. (means \pm SE; $n=5$ hearts). $*P < 0.01$ vs. sham-infused mice; *t* test. *E*) Comparison of the chemiluminescence intensities obtained using different luciferin injection methods in a Tg-line1 mouse; injections 1 and 2, intraperitoneal injections (top panels), injections 3, 4, and 5, intravenous injections (bottom panels). Injections were performed ≥ 4 h apart to eliminate the effect of the previous injection. Inset in each panel shows a magnified image of the heart. *F*) Scatterplots of the chemiluminescence intensities in the heart and brain. Plots indicate the independent experiments shown in each panel in *E*. There is a linear relationship between the expression in the heart and the brain, $R^2 = 0.979$.

expression of endogenous *Nppa* and *Nppb* mRNA increased 3 d after the PE infusion began (Fig. 3*B*, *C* and Supplemental Fig. S2*B*, *C*, *F*, *G*). Concomitantly, the quantitative PCR analysis of the CR9 luciferase mRNA expression showed enhanced expression in the ventricular myocardium 3 d after the PE infusion (Fig. 3*D* and Supplemental Fig. S2*D*, *H*). On the other hand, in the brain and the salivary glands where neither *Nppa* nor *Nppb* is highly expressed, the CR9-driven luciferase mRNA expression did not respond to PE (Fig. 3*B–D* and Supplemental Fig. S2*B–D*, *F–H*). Therefore, the

patterns of PE-induced luciferase expression suggest that CR9 is almost exclusively active in the heart. Because the integration sites were random in these three lines, the patterns of luciferase expression depend on CR9 or other enhancers near the integrated sites. The expression of luciferase in the brain of Tg-line1 and salivary glands of Tg-line2, both of which express neither *Nppa* nor *Nppb*, might be driven by other enhancers near the integrated sites.

To evaluate the accuracy and reproducibility of this method, we measured the luminescence in the heart of

a mouse from Tg-line1. In this transgenic line, the brain, intestine, and testis expressed the reporter protein due to positional effects of the insertion site and most likely not due to CR9 activity. Because the luciferase mRNA expression in the brain remained unchanged after PE treatment (Fig. 3D), we used the reporter activity in the brain as a control. The absolute luminescence values of the heart were affected by the injection method and the amount of luciferase substrate injected (Fig. 3E). However, using brain luminescence as a control, we successfully eliminated the signal variations caused by these differences. The ratio of the luminescence in the heart and brain remained constant within each mouse, independent of the injection method (Fig. 3F). Therefore, we defined the ratio of heart to brain luciferase intensities as the cardiac-specific enhancer activity.

Distal enhancer element was activated in the murine model of heart failure

To examine whether the CR9 enhancer was also responsible for gene expression in other pathological conditions, we subjected Tg-line1 mice to heart failure induced by TAC and compared them with sham-surgery mice. This model mimics the heart condition of patients with hypertension who suffer a continuous pressure overload on the heart. The pressure overload by TAC caused potent cardiac hypertrophy at 2 wk postsurgery and reduced cardiac contractility at 3 wk postsurgery (Fig. 4A, B), as previously reported (22). The endogenous *Nppa* and *Nppb* expression increased severalfold in the ventricular myocardium 3 wk after the TAC surgery (Fig. 4C). The heart to brain luciferase intensity ratio also increased severalfold 3 wk following the TAC surgery (Fig. 4D, E and Supplemental Fig. S3). However, the heart to brain luciferase intensity ratio of sham-surgery mice did not change after the surgery (Fig. 4D, E and Supplemental Fig. S3; 3 wk after TAC surgery: 5.7 ± 1.3 fold; 3 wk after sham surgery: 1.0 ± 0.2 fold; $P < 0.001$, repeated ANOVA). These results suggest that CR9 increases transcriptional activity during mechanical pressure overload-induced hypertrophy and subsequent heart failure.

DISCUSSION

Here, we focused on the stress-responsive regulatory elements of *Nppa* and *Nppb* in heart failure. By screening the evolutionarily conserved and epigenetically modified regions around the *Nppa* and *Nppb* gene loci, we identified a 650-bp transcriptional enhancer that was responsive to an α_1 -adrenergic receptor agonist *in vitro*. Furthermore, *in vivo* 3C analysis revealed that this distal enhancer directly interacted with the 5'-flanking regions of both *Nppa* and *Nppb*. Using *in vivo* live imaging of luciferase reporter gene expression, we observed that this 650-bp enhancer caused cardiac-specific activation of reporter gene expression during

the progression of pressure overload-induced heart failure. Notably, this is the first study to provide a time series analysis for monitoring enhancer activity under pathological conditions in an individual live mouse.

Although numerous approaches have been used to explore the stress-responsive regulatory elements driving gene transcription during heart failure (11, 12, 14), these elements have not yet been identified due to the technical difficulty involved. To detect the elements that are responsive to pathological conditions such as heart failure, it is essential to confirm the activity of the responsive element using a beating heart that remains connected to the systemic cardiovascular system. Therefore, it would be beneficial to establish transgenic mouse lines carrying a reporter plasmid to assess the responsive elements driving the expression of specific genes. However, the creation of multiple stable adult mouse lines to identify these elements is time-consuming.

In this study, we utilized two improved methods for reporter analysis and successfully identified a novel potent enhancer.

First, by performing an enhancer analysis using a lentiviral vector, we accurately identified candidate enhancers in cardiomyocytes and subsequently generated transgenic reporter mice. Previous promoter analyses used electroporation or lipofection to transfect cultured cardiomyocytes with plasmids (37, 38), but the transfection efficiency of these methods in primary cardiomyocytes is too low to accurately measure reporter activity during the stress response. In this study, greater than 90% transduction efficiency of cardiomyocytes was achieved using a lentiviral vector, which enabled us to accurately identify a specific enhancer fragment. Using this method, we efficiently minimized the number of reporter plasmids to be subsequently integrated into the mouse genome to screen for potential enhancers.

Second, by sequentially measuring the enhancer activity in a single live mouse, we collected robust data to assess enhancer activity in the heart *in vivo*. LacZ is not a suitable reporter for this purpose because LacZ activity can only be assessed after animal euthanization. Therefore, we overcame this limitation using the luciferase reporter plasmid. Recent advances in high-sensitivity luminescence imaging have made it possible to evaluate enhancer-driven luciferase activity without operating on the mice. Therefore, we sequentially assessed reporter activity and hemodynamic changes in the same mouse throughout the time course of the development of heart failure. These data were highly reproducible and enabled us to identify an enhancer element that was activated by cardiac overload. Because this method can be applied to any organ, the *in vivo* luciferase reporter assay may be used for assessing the *in vivo* enhancer or promoter activities responsible for clinically important diseases. The noninvasive nature of this method also enabled us to simultaneously assess the hemodynamic and metabolic parameters *in vivo* along with reporter activity. Specifically, the Tg-line1

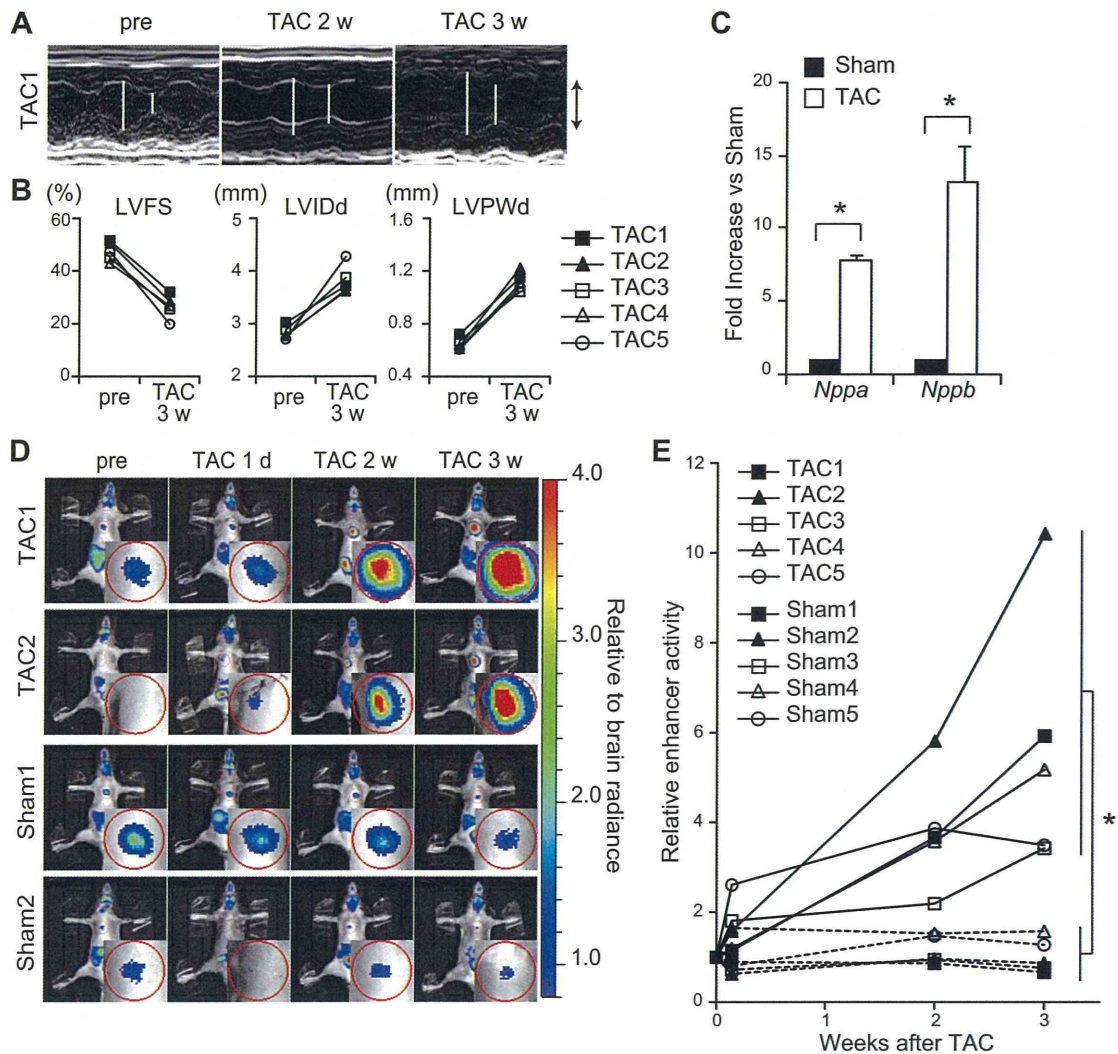


Figure 4. Distal enhancer element is reactivated in the murine model of heart failure. *A*) Representative M-mode echocardiograms in a mouse of Tg-line1 (TAC1) before and after TAC. Open bars indicate maximal left ventricular internal dimension in diastole (LVIDd) and maximal left ventricular internal dimension in systole (LVIDs). Up and down arrows represent 3 mm. *B*) Echocardiographic changes in left ventricular fractional shortening (LVFS), LVIDd, and left ventricular posterior wall thickness in diastole (LVPWd) in 5 mice of Tg-line1 (TAC1-5) before and after TAC. *C*) Relative *Nppa* and *Nppb* transcript levels in the ventricular myocardium 3 wk after the TAC procedure (means \pm SE; $n=3$ hearts). * $P < 0.05$ vs. sham-surgery mice; *t* test. *D*) Sequential *in vivo* live imaging of 4 representative Tg-line1 mice before and after TAC or sham surgery at each time point. Top 2 and bottom 2 panels represent sequential imaging data of TAC and sham-surgery mice, respectively. Sequential imaging of the 6 other surgically treated mice is shown in Supplemental Fig. S3. Insets in images show magnified images of the heart. Color scale depends on the ratio relative to brain intensity. *E*) Cardiac-specific enhancer activity plots of 10 Tg-line1 mice (TAC1, TAC2, and Sham1, Sham2, shown in *D*) and TAC3-5 and Sham3-5 shown in Supplemental Fig. S3). Heart to brain luciferase intensity ratio represents the cardiac-specific enhancer activity; enhancer activity in presurgery mice was defined as 1. 3 wk after TAC surgery: 5.7 ± 1.3 fold; 3 wk after sham surgery: 1.0 ± 0.2 fold; means \pm SE; $n = 5$. * $P < 0.001$, repeated ANOVA.

mice enabled us to accurately quantify the expression level of the natriuretic peptides. These mice are useful tools for repeatedly assessing the degree of heart failure to screen various cardiovascular drugs.

The integration of activities from multiple enhancers could confer specificity and robustness to transcriptional regulation (1). Warren *et al.* (14) identified the *Nppa* enhancer in the embryonic heart by examining Nkx2-5 binding regions around the *Nppa* locus, but the

enhancer did not respond to heart failure. This enhancer does not overlap with CR9 and might regulate *Nppa* expression only during the embryonic stage (14). On the other hand, Horsthuis *et al.* (11) showed that the regulatory region from -27 to $+58$ kb relative to the transcription start site of *Nppa* was sufficient for *Nppa* gene expression in the failing heart, similar to CR9. However, because this 85-kb regulatory region does not include CR9, *Nppa* may have multiple enhanc-

ers that regulate its expression during heart failure. Furthermore, the length of the 85-kb region poses a challenge for understanding its specific biological role.

This is the first study to provide a time course imaging analysis of enhancer activity using an individual live diseased mouse model. Using this new method, we identified a novel heart enhancer. This method can be widely used for identifying enhancers that regulate transcriptional activity only under pathological conditions. FJ

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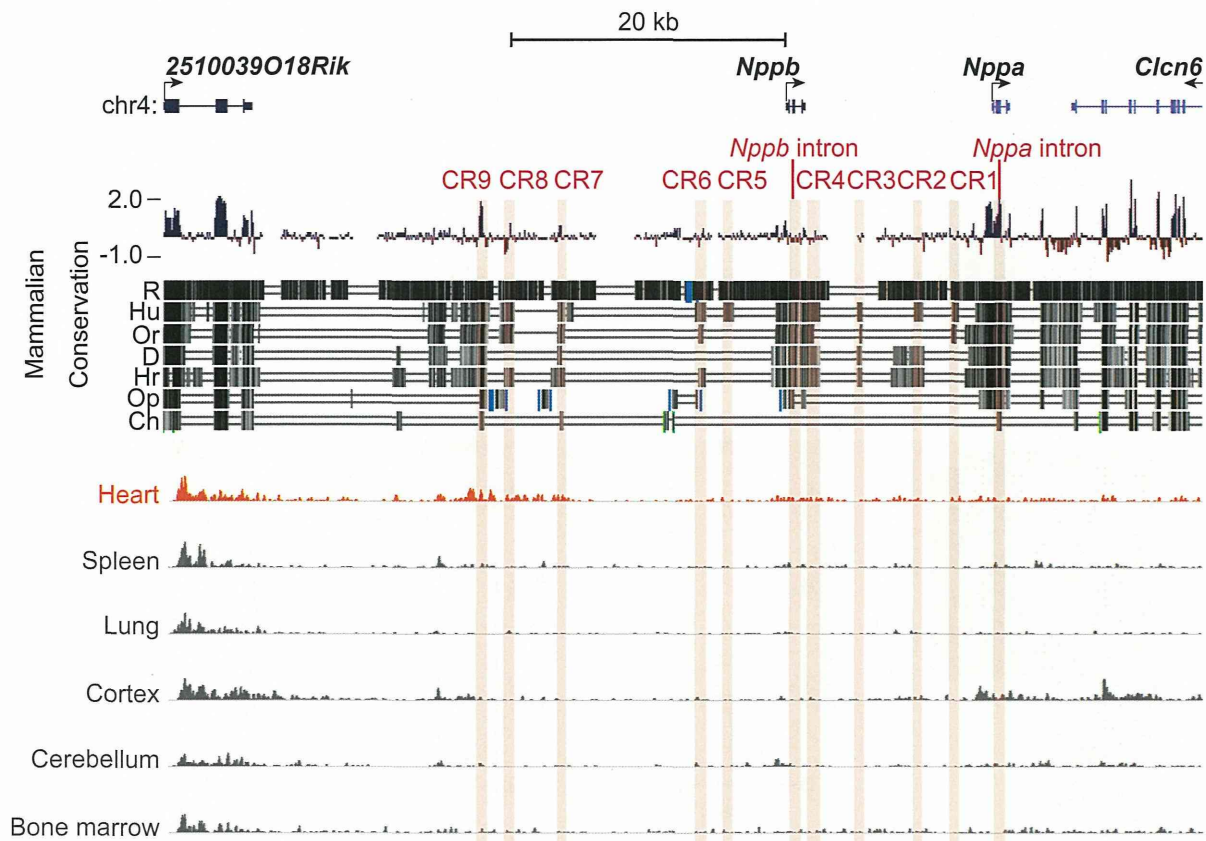
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Matsuoka_Supplemental Figure S1



Supplemental Figure S1. CR9 overlapped with the gene region that is modified by H3K4me1 in a cardiac-specific manner. Mammalian evolutionarily conserved regions (CRs) and ChIP-seq data for H3K4me1 surrounding the murine *Nppa* and *Nppb* loci. Non-coding conserved regions are highlighted as light red vertical bars (CR1-9, *Nppa* intron, and *Nppb* intron). ChIP-seq data for H3K4me1 in various organs was obtained from an open database of the adult mouse heart. R, Rat; Hu, Human; Or, Orangutan; D, Dog; Hr, Horse; Op, Opossum; Ch, Chicken.