Cellular identity and Ca^{2+} signaling activity of the non-reproductive GnRH system in the Ciona larva

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ABSTRACT

Tunicate larvae have a non-reproductive GnRH system with multiple ligands and receptor heterodimerization enabling complex regulation. In the *Ciona* larva, one of the *gnrh* genes, *gnrh2*, is conspicuously expressed in the motor ganglion and nerve cord, which are homologous structures to the hindbrain and spinal cord, respectively, of vertebrates. The *gnrh2* gene is also expressed in the proto-placodal sensory neurons, which are the proposed homologue of vertebrate olfactory neurons. The tunicate larvae occupy a non-reproductive dispersal stage, yet the roles of their GnRH system remain elusive. In this study, we investigated neuronal types of *gnrh2*-expressing cells in the *Ciona* larva and visualized activity of these cells by fluorescence imaging using a calcium sensor protein. Some cholinergic neurons as well as dopaminergic cells express *gnrh2*, suggesting that a role of GnRH in the control of swimming behavior. By contrast, none of the *gnrh2*-expressing cells overlap with glycinergic or GABAergic neurons. A role in the motor control is also suggested by correlation between activity of some *gnrh2*-expressing cells and tail movements. Interestingly, *gnrh2*-positive ependymal cells in the nerve cord, known as a kind of glia cells, actively produced Ca²⁺ transients, suggesting that neuroendocrine signaling occurs in glia cells of the nerve cord.

Keywords: gonadotropin-releasing hormone (GnRH); ascidian; calcium imaging;
Introduction

Gonadotropin-releasing hormone (GnRH) is a key regulator of reproductive functions in vertebrates (Okubo and Nagahama 2008; Oka 2009). Non-reproductive roles of GnRH have been suggested in the nervous system and during development (Dolan et al. 2003; Albertson et al. 2008; Sherwood and Wu 2005; Wu et al. 2006; Abraham et al. 2008; Kanaho et al. 2009; Ramakrishnan et al. 2010). Compared to its reproductive roles, the non-reproductive roles of GnRH are less well understood.

Tunicates are the sister group of vertebrates (Delsuc et al. 2006; Putnam et al. 2008). A conspicuous non-reproductive GnRH system has been reported in the larva of the sessile tunicate *Ciona* (Kusakabe et al. 2012; Kamiya et al. 2014). Six GnRH peptides and four receptors are encoded by the *Ciona* genome (Adams et al. 2003; Kusakabe et al. 2003; Tello et al. 2005; Sakai et al. 2010, 2012). In the *Ciona* larva, the GnRH genes are strikingly expressed in the central nervous system (CNS) through the entire antero-posterior body axis (Kusakabe et al. 2012). Correspondingly, the GnRH receptor genes are specifically expressed in the tissues and organs located along the CNS, namely the notochord, the tail muscle, and epidermal sensory neurons (Kusakabe et al. 2012). One of the *Ciona gnrh* genes, *gnrh2*, is conspicuously expressed in the motor ganglion and nerve cord of the larva, which are homologous structures to the hindbrain and spinal cord, respectively, of vertebrates. The *gnrh2* gene is also expressed in the proto-placodal sensory neurons, which are the proposed homologue of vertebrate olfactory neurons (Abitua et al. 2015). *Ciona* GnRH has been implicated to play a pivotal role in the control of metamorphosis (Kamiya et al. 2014). Considering the complex and well-developed nature of the larval GnRH system in *Ciona*,

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**References**

- Okubo and Nagahama 2008
- Oka 2009
- Dolan et al. 2003
- Albertson et al. 2008
- Sherwood and Wu 2005
- Wu et al. 2006
- Abraham et al. 2008
- Kanaho et al. 2009
- Ramakrishnan et al. 2010
- Adams et al. 2003
- Kusakabe et al. 2003
- Tello et al. 2005
- Sakai et al. 2010, 2012
- Kusakabe et al. 2012
- Kamiya et al. 2014
- Abitua et al. 2015
- Kamiya et al. 2014
- Delsuc et al. 2006
- Putnam et al. 2008
GnRH may play diverse and important roles in developmental and physiological processes in the *Ciona* larva. To date, however, the roles of the *Ciona* GnRH system remain elusive.

In this study, we investigated neuronal types of *gnrh2*-expressing cells in the *Ciona* larva and visualized activity of these cells by fluorescence imaging using a calcium sensor protein. Some cholinergic motor neurons as well as unique cholinergic cells along the nerve cord express *gnrh2*, suggesting that a role of GnRH in the control of swimming behavior. By contrast, none of the *gnrh2*-expressing cells overlap with glycinergic or GABAergic neurons. A role in the motor control is also suggested by simultaneous activation of some *gnrh2*-expressing cells with tail movements. Interestingly, *gnrh2*-positive ependymal cells in the nerve cord, known as a kind of glia cells, produced Ca\(^{2+}\) transients, suggesting that active signaling, presumably involving GnRH, occurs in the nerve cord.

**Results**

*Gnrh2 is expressed in proto placode-derived sensory neurons and caudal glial ependymal cells*

The 4.3-kb upstream region of *gnrh2* connected with a fluorescence reporter can recapitulate the expression patterns of *gnrh2* (Kusakabe et al. 2012) (Fig. 1). This upstream region was used to express mCherry and G-CaMP8 in cells expressing *gnrh2* (Figs. 1,2 and 3). Cell types were identified by double fluorescent staining of larvae with cell type-specific markers.

First, we examined whether *gnrh2*-expressing cells include glutamatergic neurons. In the *Ciona* larva, glutamate is a major neurotransmitter in the peripheral sensory neurons and photoreceptor cells (Horie et al. 2008). Some interneurons in the
posterior brain is also glutamatergic (Horie et al. 2008). As previously reported (Abitua et al. 2015), proto-placode derived gnrh2-expressing epidermal neurons (aATENs) are glutamatergic (Fig. 2A). aATENs are the only glutamatergic neurons that express gnrh2. None of the glutamatergic neurons express gnrh2 in the CNS.

Then we examined whether any of GABAergic/glycinergic neurons express gnrh2 using vesicular inhibitory amino acid transporter (VIAAT) as a marker. None of the VIAAT-positive cells were overlapped with the reporter expression under the control of the gnrh2 cis-regulatory region (Fig. 2C and 2D). In the anterior tail region, there are two pairs of VIAAT-positive neurons called ACINs, which align with glial ependymal cells in the lateral wall of the anterior nerve cord (Horie et al. 2010; Nishitsuji et al. 2012). The ACINs do not express gnrh2 whereas the lateral ependymal cells express gnrh2 (Fig. 2D). The gnrh2 expression in the lateral wall ependymal cells of the nerve cord is consistent with the in situ hybridization data previously reported (Kusakabe et al. 2012).

Cellular retinaldehyde binding protein (CRALBP) is specifically localized in the glial ependymal cells in the brain vesicle and the motor ganglion (Tsuda et al. 2003; Kusakabe et al. 2009). CRALBP-positive cells were never overlapped with gnrh2-expressing cells (Fig. 2B). Thus, in contrast to the conspicuous gnrh2 expression in the ependymal cells of the nerve cord, gnrh2 is not expressed in the ependymal cells in the brain vesicle and the motor ganglion.

**Some cholinergic and dopaminergic neurons express gnrh2**

Acetylcholine is a major neurotransmitter at the neuromuscular junctions of the Ciona larva (Yoshida et al. 2004; Horie et al. 2010). Cholinergic neurons were visualized by a
fluorescence protein expressed under the control of cis-regulatory region of the vacht gene (Yoshida et al. 2004). Gnrh2-expressing neurons were shown to be cholinergic both in the brain vesicle and the motor ganglion (Fig. 3A).

The caudal part of the CNS (nerve cord) mainly consists of non-neuronal ependymal cells (Katz 1983). The nerve cord also contains two types of neurons: ACINs and bilateral pairs of cholinergic caudal neurons (Horie et al. 2010). Some of these cholinergic caudal neurons seem to express gnrh2 (Fig. 3B).

Another neurotransmitter that controls swimming of Ciona tadpoles is dopamine (Razy-Krajka et al. 2012). Dopaminergic neurons are present in the brain vesicle (Moret et al. 2005; Razy-Krajka et al. 2012; Horie et al. 2018). Dopaminergic neurons were labeled with mCherry expressed under the control of the cis-regulatory region of the dopaminergic cell-specific gene fer2 (Razy-Krajka et al. 2012; Horie et al. 2018; previously described as Ptf1a, see Gyoja & Satoh 2013 for the orthologous families of the bHLH transcription factors). Double fluorescence imaging of dopaminergic neurons and gnrh2-expressing cells revealed that some of dopaminergic neurons express gnrh2 (Fig. 3C).

**Active Ca$^{2+}$ transients in aATENs and gnrh2-expressing cells of the posterior CNS**

G-CaMP8 was used to monitor temporal changes in intracellular Ca$^{2+}$ in gnrh2-expressing cells. Active Ca$^{2+}$ transients were observed in aATENs, the motor ganglion, and the caudal nerve cord (Figs. 4-6; Movies S1-3). The larva contains two aATENs, each has a sensory cilium (Abitua et al. 2015, Fig. 4A,B). Both aATENs showed Ca$^{2+}$ transients but their activity was not synchronized (Fig. 4C; Movie S1).

Cholinergic gnrh2-expressing neurons at the posterior part
of the motor ganglion exhibited active Ca\textsuperscript{2+} transients (Figs. 5 and 6). Periodic Ca\textsuperscript{2+} transients of a gnrh2-expressing neuron in the motor ganglion were observed at 19 hours after fertilization (arrowheads in Fig. 6B). In the tail region, Ca\textsuperscript{2+} transients were observed though the entire length of the nerve cord (Figs. 5 & 6). Both cholinergic neurons and ependymal cells showed Ca\textsuperscript{2+} transients in the tail nerve cord. For example, a narrow cell indicated by an arrow in Fig. 5A is presumably cholinergic neurons. Many of cells showing Ca\textsuperscript{2+} transients were block-shaped cells, characteristics of caudal ependymal cells (Fig. 6C).

Ca\textsuperscript{2+} transients were not synchronized between cells, including neurons and ependymal cells, located at different sites of the larva (Figs. 5 and 6). However, simultaneous activation of cells at different sites were occasionally observed as indicated by an arrowhead in Fig. 5B and Fig. 6B, suggesting the presence of a neural circuit connecting gnrh2-expressing cells at different sites.

**Correlation between tail movements and Ca\textsuperscript{2+} transients in the motor ganglion and the anterior nerve cord**

A neural circuit in the motor ganglion and the anterior nerve cord is thought to control muscle contraction of the tail (Nishino et al. 2010; Horie et al. 2010; Kusakabe 2017). Because active Ca\textsuperscript{2+} transients in gnrh2-expressing cells were observed in these regions, we examined temporal correlation between tail movement and the activity of gnrh2-expressing cells. Ca\textsuperscript{2+} transients were frequently observed when the tail stopped movement, whereas the Ca\textsuperscript{2+} transients were seemingly suppressed during the term when the tail was moving (Fig. 7). This finding suggests possible involvement of gnrh2-expressing cells in the control of swimming behavior.


**Discussion**

In this study, we identified cell types of *gnrh2*-expressing cells and visualized their activity in the *Ciona* larva. Previously, the cells expressing GnRH-encoding genes had been only partially identified in *Ciona*. The caudal ependymal cells and the aATEN epidermal neurons were reported to express *gnrh2* (Kusakabe et al. 2012; Abitua et al. 2015). We confirmed these findings and further identified CNS neurons expressing *gnrh2*.

In the brain vesicle, dopaminergic neurons and a limited number of cholinergic neurons express *gnrh2*. Pharmacological and behavioral analyses have suggested that dopaminergic cells modulate the light-off-induced swimming behavior of *Ciona* larvae (Razy-Krajka et al. 2012). The role of cholinergic neurons in the *Ciona* brain vesicle has not been elucidated. Current observation reveals previously unknown heterogeneity of cholinergic neurons in the brain vesicle and would provide a clue to investigate roles of these neurons.

Cholinergic neurons in the motor ganglion have been implicated in the regulation of tail muscle contraction (Horie et al. 2010; Nishino et al. 2010, 2011; Kusakabe 2017). Here we show that one subtype of cholinergic neurons in the motor ganglion expresses *gnrh2* (Fig. 3A). These neurons posteriorly send axons but it is unclear whether they are motor neurons directly innervating muscle cells or interneurons connecting to other CNS neurons in the caudal nerve cord. In the nerve cord, another class of cholinergic cells also expresses *gnrh2* (Fig. 3B). A potential role of cholinergic/GnRH neurons in the motor ganglion and the nerve cord may be the control of swimming behavior. These neurons may also play a role in metamorphosis because GnRH has been known to be involved in the regulation of metamorphosis (Kamiya et al. 2018).
Calcium imaging has been applied to studies of *Ciona* development (Hackley et al. 2013; Abdul-Wajid et al. 2015; Akahoshi et al. 2017). These previous studies focused on Ca\(^{2+}\) transients in embryos but not in mature larvae. The present study for the first time reported spatio-temporal patterns of Ca\(^{2+}\) transients in mature larvae of *Ciona*. Our observation includes four novel findings: i) active Ca\(^{2+}\) transients in the proto-placode-derived aATENs, ii) periodic spikes in the motor ganglion of young larvae, iii) correlation between Ca\(^{2+}\) transients and the tail movements, and iv) active Ca\(^{2+}\) transients in ependymal cells of the nerve cord.

The proto-placode-derived aATENs share morphological and molecular properties with vertebrate olfactory neurons and thought to be chemosensory cells (Abitua et al. 2015). However, olfactory receptors have not been identified in *Ciona* and chemical cues that stimulate aATENs are not known. Calcium imaging with *gnrh2>G-CaMP8* could help us search for chemical cues that triggers activation of aATENs in future studies.

Periodic Ca\(^{2+}\) transients observed in the motor ganglion of young larvae are reminiscent of spontaneous rhythmic activities observed in developing nervous systems of vertebrates (Gu et al. 1994; Wong et al. 1995; Feller et al. 1996; Zhou 1999; Kerr et al. 2005; Chang and Spitzer 2009). These periodic neuronal activities are thought to be important for the development of neural circuits in the central nervous system and the retina (Zhou 1999; Spitzer et al. 2000; Feller 2006). Similar rhythmic oscillation of Ca\(^{2+}\) transients were reported in the developing motor ganglion of the *Ciona* embryo (Akahoshi et al. 2017). By contrast, we observed rhythmic Ca\(^{2+}\) transients only in young larvae. Swimming behavior of *Ciona* larvae reveals ontogenic changes and their photo-
responsiveness appears a few hours after hatching (Nakagawa et al. 1999; Tsuda et al. 2003; Zega et al. 2006). Thus, the spontaneous rhythmic Ca\(^{2+}\) transients may play an important role in the neural circuit development of *Ciona* larvae.

We observed correlation between the tail movements and Ca\(^{2+}\) transients in the motor ganglion and the nerve cord. This suggests that *gnrh2*-expressing cells are involved in the control of swimming locomotion. Ca\(^{2+}\) transients appeared when the tail stops movement and Ca\(^{2+}\) signal was low when the tail was moving (Fig. 7B). In other words, the tail movement precedes the Ca\(^{2+}\) spike. This pattern suggests that these *gnrh2*-expressing cells are not motor neurons. In fact, majority of the cells expressing *gnrh2* in the nerve cord are ependymal cells and we observed Ca\(^{2+}\) transients correlating with tail movements in ependymal cells. An intriguing possibility is that Ca\(^{2+}\) spikes are induced in ependymal cells by muscle contraction or motor axon excitation. If so, the ependymal cells may monitor activity of muscle or motor neurons. It has been reported that various types of glia cells exhibit Ca\(^{2+}\) transients in response to neuronal activities and regulate neuronal functions in vertebrates (Reist and Smith 1992; Newman and Zahs 1997; Robitaille 1998; Haydon 2001). The ependymal cells of *Ciona* larva may have similar regulatory roles, suggesting a deep evolutionary conservation of glia function between tunicates and vertebrates. Given the simplicity of its nervous system, the *Ciona* larva could serve as a unique model for the study of glia-neuron interaction.

In conclusion, the present study reveals dynamic Ca\(^{2+}\) transients of *gnrh2*-expressing cells at various sites in the *Ciona* larva. Our findings suggest a connection between activity of *gnrh2*-expressing cells and tail movements of the larva. An important yet unsolved question is whether GnRH2 is involved in these processes.
Future studies should address developmental and physiological roles of \textit{gnrh2}-expressing cells and GnRH peptides based on the findings of this study.

\textbf{Methods}

\textbf{Animals and embryos}

Mature adults of \textit{Ciona intestinalis} Type A (also called \textit{Ciona robusta}) were provided by the Maizuru Fisheries Research Station of Kyoto University and by the Misaki Marine Biological Station of the University of Tokyo through the National Bio-Resource Project (NBRP) of MEXT, Japan, and maintained in indoor tanks of artificial seawater (ASW) (Marine Art BR, Tomita Pharmaceutical Co., Ltd., Tokushima, Japan) at 18°C. The adults were also collected from the pond on the Fukae campus of Kobe University, Kobe, Japan and from the fishing harbor in Murotsu, Hyogo, Japan. Eggs and sperm were obtained surgically from the gonoducts, and eggs were fertilized \textit{in vitro}. After insemination, embryos were raised in ASW containing 50 \(\mu\)g/ml streptomycin sulfate (S6501, Sigma-Aldrich, St. Louis, MO, USA) at 18°C.

\textbf{Preparation of reporter constructs and electroporation}

Construction of the \textit{vglut>kaede} was described previously (Horie et al., 2011, Razy-Krajka et al. 2012). The \textit{vacht>cfp} plasmid was made by inserting the 3.8-kb upstream region of \textit{Ciona vacht} (Yoshida et al. 2004) into the \textit{SalI/BamHI} site of pSP-CFP (Horie et al. 2011). The 2.4-kb upstream region of \textit{Ciona fer2} (Gene ID KH.L116.39) was previously cloned in the pSP-CFP vector (Razy-Krajka et al. 2012). The reporter sequence was replaced with a DNA fragment coding for mCherry to generate \textit{fer2>mcherry}
using NotI/EcoRI sites. The *gnrh2*-*kaede* and *gnrh2*-*mcherry* plasmids were made by inserting the 4.3-kb upstream region of *Ciona gnrh2* (Kusakabe et al. 2012) into the XhoI/NotI sites of the pSP-Kaede vector and pSP-mCherry vector, respectively (Hozumi et al. 2010). The *gnrh2* upstream region was also used to generate the *gnrh2*-*g-camp8* construct. The Kaede coding sequence of pSP-Kaede was replaced with a DNA fragment coding for G-CaMP8 (Ohkura et al. 2012) using NotI/EcoRI sites. The *gnrh2* upstream region was amplified from the *gnrh2*-*kaede* plasmid using a pair of nucleotide primers (5'-GAATCGGCCAACGCGGGATCCAGGAGCAGACGTCATAAGTA-3' and 5'-TGACGCGGCCGCTGTTACGTTATCTCTCTTAGAAG-3'), digested with BamHI and NotI, and then inserted into the BamHI/NotI sites upstream of the G-CaMP8 in the pSP vector. Plasmid DNA constructs were electroporated into *Ciona* fertilized eggs as described by Corbo et al. (1997).

**Immunofluorescent staining**

Immunofluorescent staining was carried out according to the method described by Nishitsuji et al. (2012). Fluorescent images were obtained by using a laser scanning confocal microscope (FV1200 IX83; Olympus, Tokyo, Japan).

To visualize localization of cell-type specific proteins, a mouse antiserum against *Ciona* VIAAT (Horie et al. 2010) or a rabbit antiserum against *Ciona* CRALBP (Tsuda et al. 2003) was diluted 1:1000 in 10% goat serum in T-PBS (0.1% Triton X-100 in PBS) and used as the primary antibody. The secondary antibody was an Alexa Fluor 594–conjugated anti-mouse IgG (A11005; Thermo Fisher Scientific) or an Alexa Fluor 594–conjugated anti-rabbit IgG (A11012; Thermo Fisher Scientific).

Primary antibodies used to visualize localization of
fluorescent reporter proteins are rabbit anti-Kaede polyclonal (PM012; Medical & Biological Laboratories Co., Ltd., Nagoya, Japan; for Kaede), rabbit anti-GFP polyclonal (A11122; Thermo Fisher Scientific; for G-CaMP8 and CFP), rat anti-RFP monoclonal (5F8; ChromoTek GmbH, Martinsried, Germany; for mCherry), and rat anti-GFP monoclonal (GF090R; Nacalai Tesque, Kyoto, Japan; for G-CaMP8 double stained with anti-CRALBP) antibodies. All the primary antibodies are diluted 1000-fold as described above. Secondary antibodies were an Alexa Fluor 488−conjugated anti-rabbit IgG (A11008; Thermo Fisher Scientific) for G-CaMP8 and CFP, an Alexa Fluor 488−conjugated anti-rat IgG (A11006; Thermo Fisher Scientific) for G-CaMP8, and an Alexa Fluor 594−conjugated anti-rat IgG (A11007; Thermo Fisher Scientific) for mCherry.

**In vivo Ca**\(^{2+}\) **imaging**

Electroporated *Ciona* larvae expressing the G-CaMP8 transgene was placed in ASW on a 35-mm glass based dish (coverslip diameter 12 mm, #3931-035; Iwaki, Japan). For imaging, a microscope (IX81, Olympus, Tokyo, Japan) equipped with an EMCCD camera (EVOlVE512, Photometrics, USA) was used. Images were taken with a 50-ms exposure time per 1 second and 1×1 binning. For each animal, 300 images were taken in 6 minutes. Changes in intracellular calcium concentrations were measured as the changes of green fluorescence of G-CaMP8. MetaMorph image analysis software system (Molecular Devices) was used for the analysis of imaging. Image processing was also performed with ImageJ (U. S. National Institutes of Health, Bethesda, Maryland, USA, https://imagej.nih.gov/ij/).

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Figure Legends

Figure 1. Expression patterns of gnrh2 in the Ciona larva. (A) Schematic diagram showing the central nervous system (CNS) of the larva. (B) Localization of gnrh2 mRNA visualized by in situ hybridization. (C) Immunofluorescent localization of G-CaMP8 expressed under the control of the cis-regulatory region of gnrh2. Scale bar, 200 µm.

Figure 2. Immunohistochemical identification of types of cells expressing gnrh2 in the Ciona larva. (A) Glutamatergic neurons and gnrh2-expressing cells were labeled with Kaede (green) and mCherry (magenta), respectively. The proto placode-derived sensory neurons (arrows) were shown to express gnrh2. (B) Cellular retinal aldehyde binding protein (CRALBP)-positive cells (magenta) were not overlapped with gnrh2-expressing cells (green). Arrows indicate gnrh2-expressing cells in the brain vesicle. (C,D) GABAergic/glycinergic neurons were visualized by immunostaining with anti-VIAAT antibody (magenta). Arrows in (C) indicate GABAergic/glycinergic neurons in the motor ganglion. Arrows in (D) indicate VIAAT-positive ACINs. Scale bars, 30 µm.

Figure 3. Some cholinergic and dopaminergic neurons express gnrh2. (A,B) Cholinergic neurons and gnrh2-expressing cells were labeled with CFP (green) and mCherry (magenta), respectively. Arrows indicate cells co-expressed both markers. (C) Dopaminergic cells and gnrh2-expressing cells were labeled with mCherry (magenta) and G-CaMP8 (green), respectively. Some dopaminergic cells were also labeled with G-CaMP8 (arrows). Scale bars, 30 µm.

Figure 4. Calcium imaging of gnrh2-expressing chemo...
neurons. Fluorescence images of *Ciona* larvae expressing G-CaMP8 in the chemosensory apical trunk epidermal neurons (aATENs). (A) The putative sensory cilia of each aATEN was labeled with G-CaMP8 fluorescence (dotted circle). (B,C) An example of a larva showing dynamic Ca\(^{2+}\) transients in the pair of aATENs (*arrowheads*) at 19 hpf (hours post fertilization). (C) Representative images of the larva recorded at the time (s) indicated. The serial images of the larva shown in (B,C) is shown in **Movie S1**.

**Figure 5.** Interconnection between *gnrh2*-expressing cells in the larval central nervous system. (A) Fluorescence image of a larva at 19.5 hpf, showing G-CaMP8 fluorescence in the motor ganglion and the anterior nerve cord. (B) The graph shows temporal patterns of fluorescence intensity at the three sites indicated by circles in (A). Colors of lines correspond to the sites indicated by circles in respective colors. The Ca\(^{2+}\) transients occurred independently of each other, but sometimes occurred at the same time (*arrowheads*). (C) Representative images of the larva recorded at the time (s) indicated. The serial images of the larva shown in (A) is shown in **Movie S2**.

**Figure 6.** Periodic oscillation of Ca\(^{2+}\) transients in a *gnrh2*-expressing cell in the motor ganglion. (A) Fluorescence image of a larva at 20 hpf, showing G-CaMP8 fluorescence in the motor ganglion and the anterior nerve cord. (B) The graph shows temporal patterns of fluorescence intensity at the four sites indicated by circles in (A). Colors of lines correspond to the sites indicated by circles in respective colors. Ca\(^{2+}\) spikes were periodically observed at regular intervals in the cell indicated by the magenta circle in (A) (*arrowheads*). (C) Representative images of the larva recorded at the time (s) indicated. The serial images of
the larva shown in (A) is shown in Movie S3.

**Figure 7.** Correlation between Ca\(^{2+}\) transients in *gnrh2*-expressing cells and the tail movement. (A) Fluorescence image of a larva at 19.5 hpf, showing G-CaMP8 fluorescence in the anterior nerve cord. (B) The graph shows temporal patterns of fluorescence intensity at the four sites encircled by colored lines. Colors of lines in the graph correspond to the sites encircled by lines in respective colors. Gray vertical lines indicate the period when the tail was moving. Ca\(^{2+}\) transients generally occurred when the tail movement stopped. (C) Representative images of the larva recorded at the time (s) indicated. The serial images of the larva shown in (A) is shown in Movie S4.

**Movie S1.** Serial fluorescence images showing Ca\(^{2+}\) transients in the larva shown in Figure 4.

**Movie S2.** Serial fluorescence images showing Ca\(^{2+}\) transients in the larva shown in Figure 5.

**Movie S3.** Serial fluorescence images showing Ca\(^{2+}\) transients in the larva shown in Figure 6.

**Movie S4.** Serial fluorescence images showing Ca\(^{2+}\) transients in the larva shown in Figure 7.
Figure 1

Figure A: Diagram of a larval cyclopiform comprising sensory brain vesicle, brain vesicle, nerve cord, motor ganglion, neck, ocellus, otolith, sensory vesicle, posterior brain, adhesive organ, and notochord.

Figures B and C: Micrographs showing gnrh2 mRNA expression and gnrh2>G-CaMP8 expression.
Figure 2

A

VGLUT>Kaede

gnrh2>mCherry

B

gnrh2>G-CaMP8

CRALBP

C

gnrh2>G-CaMP8

anti-VIAAT

D

gnrh2>G-CaMP8

anti-VIAAT
Figure 5
Figure 6
Figure 7

A

B

C

Figure 7