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### Authors

Cho, Kook-Ho  
Daubnerová, Ivana  
Park, Yoonseong  
[et al.](#)

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## Secretory Competence in a Gateway Endocrine Cell Conferred by the Nuclear Receptor $\beta$ FTZ-F1 Enables Stage-specific Ecdysone Responses Throughout Development in *Drosophila*

Kook-Ho Cho<sup>1</sup>, Ivana Daubnerová<sup>2</sup>, Yoonseong Park<sup>3</sup>, Dusan Zitnan<sup>2</sup>, and Michael E. Adams<sup>1,\*</sup>

<sup>1</sup>Departments of Entomology and Cell Biology & Neuroscience, 2103 Biological Sciences Bldg., University of California, Riverside, CA 92521 USA

<sup>2</sup>Institute of Zoology, Slovak Academy of Sciences, Dubravska cesta 9, 84506 Bratislava, Slovakia

<sup>3</sup>Department of Entomology, 123 Waters Hall, Kansas State University, Manhattan, KS 66506 USA

### Abstract

Hormone-induced changes in gene expression initiate periodic molts and metamorphosis during insect development. Successful execution of these developmental steps depends upon successive phases of rising and falling 20-hydroxyecdysone (20E) levels, leading to a cascade of nuclear receptor-driven transcriptional activity that enables stage- and tissue-specific responses to the steroid. Among the cellular processes associated with declining steroids is acquisition of secretory competence in endocrine Inka cells, the source of ecdysis triggering hormones (ETH). We show here that Inka cell secretory competence is conferred by the orphan nuclear receptor  $\beta$ FTZ-F1. Selective RNA silencing of  *$\beta$ ftz-f1* in Inka cells prevents ETH release, causing developmental arrest at all stages. Affected larvae display *buttoned-up*, the ETH-null phenotype characterized by double mouthparts, absence of ecdysis behaviors, and failure to shed the old cuticle. During the mid-prepupal period, individuals fail to translocate the air bubble, execute head eversion and elongate incipient wings and legs. Those that escape to the adult stage are defective in wing expansion and cuticle sclerotization. Failure to release ETH in  *$\beta$ ftz-f1* silenced animals is indicated by persistent ETH immunoreactivity in Inka cells. Arrested larvae are rescued by precisely-timed ETH injection or Inka cell-targeted  $\beta$ FTZ-F1 expression. Moreover, premature  *$\beta$ ftz-f1* expression in these cells also results in developmental arrest. The Inka cell therefore functions as a “gateway cell”, whose secretion of ETH serves as a key downstream physiological output enabling stage-specific responses to 20E that are required to advance through critical developmental steps. This secretory function depends on transient and precisely timed  *$\beta$ FTZ-F1* expression late in the molt as steroids decline.

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\*Corresponding author: Michael E. Adams, Dept. Cell Biology & Neuroscience, University of California, Riverside, CA 92521, michael.adams@ucr.edu.

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## Keywords

*Drosophila*; ecdysis; ecdysis triggering hormone (ETH); Inka cells;  $\beta$ FTZ-F1; RNA-silencing; secretory competence

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## Introduction

Hormones regulate developmental sequences and reproductive state through induction of stage- and tissue-specific responses. The response of a cell at any given time depends upon patterns of gene expression that confer “competence”. In insects, levels of 20-hydroxyecdysone (20E) and juvenile hormones fluctuate throughout ontogeny to determine stage-specific developmental events during embryogenesis, larval stages, and metamorphosis to the adult. Stage-dependent responses to these generalized hormonal signals are specified by nuclear receptor cascades originally recognized as temporal sequences of chromosomal puffing patterns corresponding to transcriptional activity (Ashburner et al., 1974; Becker, 1959; Yamada et al., 2000).

A key molecule in the 20E-induced signaling cascade is the “competence factor”  $\beta$ FTZ-F1, an orphan nuclear receptor whose transient expression during the hours preceding each ecdysis is induced as steroids decline to minimal levels. Early accounts showed that  $\beta$ FTZ-F1 confers cellular competence for mid-prepupal responses to 20E (Broadus et al., 1999; Yamada et al., 2000).  $\beta$ FTZ-F1 mutants exhibit a series of deficits, including failure to translocate the air bubble, perform head eversion and elongate incipient legs and wings, all hallmarks of the pre-pupal to pupal transition that occurs during pupal ecdysis. These mutants also fail to advance through larval stages: they progress to the pharate stage, characterized by formation of new cuticle, spiracles, mouthparts, and trachea, but die through failure to shed the corresponding old structures via ecdysis (Yamada et al., 2000). These observations raise the question: What cellular elements are involved in these  $\beta$ FTZ-F1-dependent responses to 20E?

In moths, transition from high to low steroid levels during the latter part of the molt is essential for ecdysis, which can be delayed by prolongation of high steroid levels (Curtis et al., 1984; Truman et al., 1983; Zitnan et al., 1999). Maintenance of high steroid levels also inhibits acquisition of secretory competence in Inka cells, the source of ecdysis triggering hormones (ETH) (Kingan and Adams, 2000; Zitnan et al., 1999). Dependence of both  $\beta$ FTZ-F1 expression and Inka cell secretory competence on declining steroid levels suggests these two events may be causally related.

We therefore tested whether  $\beta$ FTZ-F1 confers secretory competence in Inka cells through use of the GeneSwitch/UAS system (Osterwalder et al., 2001) for conditional regulation of gene expression specifically in these cells. Our findings demonstrate that Inka cell-selective silencing of  *$\beta$ ftz-f1* blocks ETH secretion, leading to failure of larval ecdyses, the prepupal-pupal transition, and adult eclosion. The timing of  $\beta$ FTZ-F1 expression in Inka cells is critically important, since its premature expression causes similar ecdysis defects. Finally, precisely timed ETH injection or targeted expression of  $\beta$ FTZ-F1 in Inka cells rescues  *$\beta$ ftz-f1* null mutants. These findings indicate that Inka cells play a crucial “gateway” role by enabling physiological outcomes initiated by the 20E-induced nuclear receptor cascade at onset of each molt.

## Materials and Methods

### Fly Stocks

The  $\beta$ FTZ-F1 null mutant ( $yw; \beta ftz-f1^{ex7}hs-\beta FTZ-F1/TM3, y^+$ - designated henceforth as  $\beta ftz-f1^{ex7}hs-\beta FTZ-F1$ ) fly line was kindly provided by Dr. Hitoshi Ueda. Additional fly lines used in this study are described below. All *Drosophila* stocks were maintained on a standard cornmeal-molasses diet at 25°C in a 12/12 hour light/dark cycle.

### Staging of animals

Larval stages were distinguished by morphology of the anterior spiracle (Ashburner, 1989). Larvae with double mouth hooks were selected and kept at 25°C until appearance of double vertical plates (dVP). Prepupae were staged by hours after puparium formation.

### Construction of ETH-GeneSwitch, UAS- $\beta$ FTZ-F1, and UAS- $\beta$ FTZ-F1 dsRNA fly lines

To drive conditional, Inka cell-specific expression of transgenes, we constructed a driver fly line carrying the *eth* promoter upstream of the RU486-dependent GAL4-progesterone receptor fusion protein (GeneSwitch). The *eth* promoter consists of the 362 bp sequence from -367 to -5 immediately upstream of the *eth* open reading frame (*ethup*; Park *et al.*, 2002). It was PCR amplified with primers carrying *Mlu*I and *Not*I restriction motifs for 5' and 3' ends, respectively; the forward primer was 5'-ACGCGTATACTTTGTATATTTATTATT-3' and the reverse primer was 5'-GCGGCCGCACCTGACTCCTGCTCCACAAT-3'. The amplified fragment was inserted into the *Drosophila* transformation vector pP{UAS-GeneSwitch} (Osterwalder *et al.*, 2001) (kindly supplied by Dr. Thomas Osterwalder, Yale University) by replacing adjacent 5×UAS and hsp70 sequences with the *ethup* sequence using *Mlu*I and *Not*I restriction sites. This driver line was designated "ETHGS" (ETH-GeneSwitch). Plasmid constructs used for fly transgenesis were sent to *Drosophila* Genetic Service Center in Duke University for germ-line transformation. Inverse PCR determined the transgene insertion to be in the 2<sup>nd</sup> chromosome.

For creation of UAS- $\beta$ FTZ-F1 fly lines, the entire ORF of *Drosophila*  $\beta ftz-f1$  (2450 bp) was PCR amplified from total RNA extracted from tracheal tissue. To facilitate cloning, EcoRI and KpnI adapter sequences were added to the PCR primers, whose sequences were as follows: the forward primer was 5'-CATGAATTCATGTTATTAGAAATGGATCAGC-3', and the reverse primer was 5'-TGAGGTACCTCATAAATGATTAAGTATTCCG-3'. Amplified cDNA encoding  $\beta ftz-f1$  was introduced into the pUAST vector and confirmed by nucleotide sequencing. Following germ-line transformation, the insertion was determined to be on the 2<sup>nd</sup> chromosome.

For RNA silencing of  $\beta ftz-f1$ , we created a transgenic fly line carrying a symmetrically transcribed, UAS-inducible double-stranded  $\beta ftz-f1$  sequence (Giordano *et al.*, 2002). The construct was PCR amplified using the following primers: the forward primer was 5'-GTTTCGAGCGGATAGAAATGCGTGGTG-3' and the reverse primer was 5'-AGTATTCGGTGTACGTTCTCCCGAC-3'. The fragment was cloned into pGEMT-easy, digested with EcoRI and inserted into the SympUAST vector. The resulting lines used in experiments described here carried transgenes on chromosome 3.

### Use of RU486-induced GeneSwitch for RNA silencing and rescue

RU486 (Sigma) was dissolved in absolute ethanol at 10 mg/ml concentration and kept at -20°C until use. Working dilutions of RU486 were prepared in ethanol and mixed with larval diet to achieve a final concentration of 100-200  $\mu$ g/ml RU486 and 4% ethanol. Fly lines were raised on a standard diet to the desired age (2<sup>nd</sup> or 3<sup>rd</sup> instar) and transferred to

diet containing RU486 (100-200  $\mu\text{g/ml}$ ). Larvae maintained on RU486 diet were scored for developmental defects under a dissection microscope. Trachea containing Inka cells were extirpated and processed for *in situ* hybridization with a  $\beta\text{FTZ-F1}$  probe followed by immunohistochemistry with antibodies to  $\beta\text{FTZ-F1}$  or ETH (see below).

To rescue the  $\beta\text{ftz-f1}^{\text{ex7}}\text{hs-}\beta\text{FTZ-F1}$  null mutant phenotype, we used either heat treatment or GeneSwitch-mediated expression of  $\beta\text{FTZ-F1}$  specifically in Inka cells using progeny of the following parental cross:  $yw; \text{ETHGS}/\text{ETHGS}; \beta\text{ftz-f1}^{\text{ex7}}\text{hs-}\beta\text{FTZ-F1}/\text{TM3}, y^+ \times \text{UAS-FTZ-F1}/\text{UAS-FTZ-F1}; \beta\text{ftz-f1}^{\text{ex7}}\text{hs-}\beta\text{FTZ-F1}/\text{TM3}, y^+$ . Larval progeny homozygous for  $\beta\text{ftz-f1}^{\text{ex7}}$  were recognized by the  $y^-$  marker, selected every 2 hours for heat-shock treatment (33°C for 60 minutes) during the time period 13-16 hours after egg laying (AEL), and maintained at 25°C. Surviving 1<sup>st</sup> instar larvae were collected during 42-46 hours AEL and exposed either to a second heat treatment (33°C for 60 min) or to RU486-containing diet. The number of 1<sup>st</sup> and 2<sup>nd</sup> instar larvae was determined during the time interval 54-59 hours AEL.

### Viability and developmental defects in flies with silenced and induced $\beta\text{FTZ-F1}$

Freshly ecdysed 2<sup>nd</sup> instar larvae produced from  $\text{ETHGS} \times \text{UAS-FTZ-F1}$  or  $\text{ETHGS} \times \text{UAS-}\beta\text{FTZ-F1}$  crosses were collected and transferred to RU486-containing diet (100-200  $\mu\text{g/ml}$ ). Animals were scored as developmentally arrested or surviving at 5-12 hour intervals; double mouth hook (dMH) larvae or puparia were selected for observation of ecdysis and survival. Mortality or developmental defects of flies were calculated as the percentage of arrested pharate 3<sup>rd</sup> instar larvae, pharate pupae or eclosed adults with short wings.

### ETH injections into $\beta\text{FTZ-F1}$ silenced larvae

Pharate 3<sup>rd</sup> instar larvae at the dVP stage were immobilized on double sticky tape and injected with 1-10 fmol of *Drosophila melanogaster* ETH1 (DrmETH1) dissolved in 10 nl of distilled water using a glass capillary attached to a “Nanoject” nanoliter injector (Drummond Scientific). Surviving larvae were transferred to a small agar plate and observed for ecdysis behavior. In parallel experiments, control larvae were injected with the same volume of distilled water.

### *In situ* hybridization and immunohistochemistry

For *in situ* hybridization and antibody staining, the tissues were dissected in phosphate-buffered saline (PBS) (130 mM NaCl, 7 mM  $\text{Na}_2\text{HPO}_4$ , 3 mM  $\text{NaH}_2\text{PO}_4$ , pH 7.4) and fixed in 4% paraformaldehyde (in PBS) for ~1 hour at room temperature or overnight at 4°C. *In situ* hybridization and immunohistochemical staining were performed as previously described (Kim et al., 2006b). For detection of  $\beta\text{FTZ-F1}$  expression in Inka cells, digoxigenin-labeled DNA probe was prepared by incorporation of Dig-11-dUTP (Roche Applied Science, Basel) into a single-stranded antisense DNA. Primers used for production of Drm  $\beta\text{FTZ-F1}$ -specific probe were as follows:  $\beta\text{FTZ-F1}$  sense primer 5' - GGACACCACCTCCTCACACT-3' and antisense primer 5' - GGAATTGGTTCCTCCTCCTC-3'. Negative control included *in situ* hybridization with a probe produced by the sense primer.

For immunohistochemical staining tracheae with attached Inka cells were incubated with primary antisera for 2 days at 4°C at a 1:2,000 dilution of DrmETH1 antiserum (Park et al., 2002) or 1:1,000 for Drm $\beta\text{FTZ-F1}$  antiserum, kindly supplied by Dr. Hitoshi Ueda (Yamada et al., 2000). To reveal specific binding of the primary antibodies, we used Alexa Fluor 488-labeled goat anti-rabbit IgG and Alexa Fluor 555-labeled goat anti-mouse IgG (Invitrogen). *In situ* hybridization and immunohistochemical staining were observed under confocal microscopes (Zeiss 510, or Leica TCS SPE-II).

## Results

### ***ethup*-GeneSwitch Induces Conditional, Inka Cell-specific Expression of Transgenes**

To analyze regulatory functions for  $\beta$ FTZ-F1 in secretory competence of Inka cells and ecdysis behaviors, we created a transgenic fly line capable of driving UAS-transgene expression under precise spatial and temporal control. We showed previously that the *eth* promoter sequence located immediately upstream of the *eth* open reading frame (*ethup*) drives Inka cell-specific gene expression (Park et al., 2002). We inserted this promoter sequence upstream of GeneSwitch (Osterwalder et al., 2001) and properties of the resulting ETH-GeneSwitch (ETHGS) line were checked first by crossing it to a UAS-EGFP line. No EGFP expression was observed unless progeny were fed RU486, which induced expression specifically in Inka cells within 2 hours; no EGFP fluorescence was observed in other tissues or organs (data not shown).

We performed an additional experiment, using the ETHGS driver line to induce  $\beta$ FTZ-F1 expression. Normally,  $\beta$ FTZ-F1 expression occurs during a narrow time window preceding ecdysis; i.e. in late 1<sup>st</sup> instar larvae (~45-48 hours AEL), late 2<sup>nd</sup> instar larvae (~69-72 hours AEL), and late 3<sup>rd</sup> instar larvae (~8-12 hours APF) preceding pupal ecdysis (Yamada et al., 2000). We induced premature expression of  $\beta$ FTZ-F1 by crossing the ETHGS line with a UAS-*βftz-f1* line; F1 progeny in 2<sup>nd</sup> or 3<sup>rd</sup> larval stages were fed RU486 and subjected to a double staining procedure: *in situ* hybridization was performed using a *βftz-f1* digoxigenin-labeled DNA probe, followed by immunostaining with a DrmETH antibody. Inka-cell specific expression of the  $\beta$ FTZ-F1 transcript became detectable 1-2 hours after larvae were transferred to diet containing RU486 (Fig. 1). The *in situ* hybridization reaction product depicted in Fig. 1A corresponds to the location of the Inka cell, as evidenced by the occurrence of ETH-like immunoreactivity (ETH-IR) in the same location (Fig. 1B,C arrow). These results confirm that the ETHGS driver line induces transgene expression under precise spatial and temporal control.

### **Selective Knockdown of *βftz-f1* Transcripts in Inka Cells**

We performed RNA-silencing of *βftz-f1* in Inka cells by crossing the ETHGS driver line with the UAS-*βftz-f1i* line carrying a double-stranded RNA construct; progeny were fed RU486 during the 3<sup>rd</sup> instar. Presence of  $\beta$ FTZ-F1 protein in knock-down and control flies was analyzed by immunohistochemical staining. Inka cells of prepupal flies fed RU486 were devoid of  $\beta$ FTZ-F1-IR, while strong staining for the protein was detected in nuclei of Inka cells in control flies (Fig. 2). Neighboring tracheal epidermal cells exhibited positive  $\beta$ FTZ-F1 immunostaining in both sets of flies, demonstrating Inka cell specificity of *βftz-f1* silencing.

### **$\beta$ FTZ-F1 Knockdown in Inka Cells Causes Lethal Ecdysis Deficiencies**

We examined phenotypes in all stages (larva, pupa, adult) subjected to RNA silencing of *βftz-f1*, effected by feeding RU486 to *ETHGS/UAS-βftz-f1i* flies during larval development (2<sup>nd</sup> and 3<sup>rd</sup> instars). The developmental profile of RU486-fed larvae was indistinguishable from wild type controls until they reached the double vertical plate (dVP) stage. Targeted  $\beta$ FTZ-F1 knockdown in Inka cells produced lethal ecdysis deficiency in 47% of late 2<sup>nd</sup> instar larvae (Table 1). Larvae were arrested at the dVP stage, failed to shed the old cuticle and died 1-2 days later. Arrested larvae exhibited the typical “buttoned-up” phenotype (Fig. 3A), originally described in *eth*-null mutants (Park et al., 2002), consisting of dVP, double mouth hooks (Fig. 3B), double tubular tracheae with anterior and posterior spiracles, and a new layer of cuticle for the next instar. The high percentage of lethality (47%) in late 2<sup>nd</sup> instar larvae with fully developed structures appropriate for the next larval stage indicates that targeted  $\beta$ FTZ-F1 knock-down in Inka cells resulted in failure of the ecdysis sequence.

RU486-fed larvae that escaped to the 3<sup>rd</sup> instar developed normally and formed the puparium (Fig. 3C), but the vast majority failed to advance through the pre-pupal to pupal transition. Cumulative mortality reached 97% during or shortly after unsuccessful pupation (Table 1). Most of these animals (86%) were arrested during translocation of the air bubble to the anterior region, failed to complete head eversion, and exhibited an abnormal space between the posterior abdomen and puparium (Figs. 3D, E). In freshly ecdysed control pupae, the air bubble reached the anterior end of the puparium (Fig. 3F). Some RU486-treated pharate pupae (12%) completed air bubble translocation and head eversion, but showed a space at the posterior tip of the puparium and failed to expand legs and wings (Fig. 3G). These appendages reached only one third of the body length in the puparium (Figs. 3G-I), in contrast to those of control pupae, which extended appendages to two thirds of the body length (Fig. 3J). In certain instances, head and thorax continued to develop for 1-2 days, whereas the abdomen failed to show pigmentation and retained a morphology characteristic of the early pupal stage (Fig. 3H). Two flies eclosed successfully (Fig. 3I), but these individuals failed to expand the wings and exhibited dimples on the dorsal thorax and soft abdomens (Fig. 3K), indicative of incomplete cuticle sclerotization. No other abnormalities were observed in these flies. These observations indicate that *βFTZ-F1* silencing in Inka cells leads to lethal defects at larval and pupal ecdysis and adult eclosion.

### ***βFTZ-F1* Knockdown Blocks Release of ETH from Inka Cells**

Resemblance of the *βFTZ-F1* knockdown phenotype in larvae to the ETH-null mutant, *buttoned-up* (Park et al., 2002), led us to hypothesize failure of Inka cells to acquire secretory competence, resulting in ETH deficiency. To test this possibility, Inka cells of *βftz-f1* knock-down larvae arrested at the dVP stage were subjected to immunohistochemical staining with ETH-specific antiserum. We observed strong ETH-IR in Inka cells of both control (ETHGS/UAS-EGFP) and *βFTZ-F1* knock-down larvae at the dVP stage (Fig. 4A-D). ETH-IR disappeared during the 10 min following dVP in control flies, demonstrating ETH release from Inka cells (Figs. 4A'-C'). However ETH staining persisted for hours in Inka cells of dVP-arrested *ETHGS/UAS-βftz-f1* larvae, indicating failure to release ETH (Fig. 3D'). This would explain ecdysis defects and lethal phenotypes observed in *ETHGS/UAS-βftz-f1* flies. Accumulation of ETH-IR in Inka cells of *βFTZ-F1* silenced larvae also implies that *βFTZ-F1* is not required for ETH expression, but is essential for inducing competence to release this peptide hormone.

### **ETH Injection Rescues Developmental Arrest in *βFTZ-F1* Knockdown Larvae**

If ETH deficiency is responsible for developmental arrest in *ETHGS/UAS-βftz-f1* larvae, injection of synthetic peptide should be an effective rescue strategy. DrmETH1 was chosen because of its efficiency in rescue of ETH-null mutants (Park et al., 2002). First instar larvae (*ETHGS/UAS-βftz-f1*) were transferred to a diet containing RU486 and injected when they reached the late 2<sup>nd</sup> instar stage. DrmETH1 (10 fmol) was injected 30 min after appearance of double vertical plates (dVP). The number of rescued animals was compared with that of water-injected control larvae. Most *ETHGS/UAS-βftz-f1* larvae arrested at the dVP stage performed the ecdysis sequence following DrmETH1 injection (19 out of 20; Table 2). Considering that the percentage of lethality was 47% in *βftz-f1* knock-down flies (see Table 1), injection of DrmETH1 rescued 89% of knock-down flies. This result supports the hypothesis that the failure of ecdysis in *ETHGS/UAS-βftz-f1* is caused by suppression of ETH release.

### **Targeted Ablation of Inka Cells Phenocopies *βFTZ-F1* silencing**

We have shown that *βftz-f1* silencing in Inka cells leads to failure of ETH release, resulting in a pattern of ecdysis defects similar to those observed previously in ETH-null mutants

(Park et al., 2002). We sought to confirm that lack of ETH leads to ecdysis deficiencies in later developmental stages by comparing phenotypes of  $\beta$ FTZ-F1 silenced flies (*ETHGS/UAS- $\beta$ ftz-f1*) with those of Inka cell-ablated flies (*ETHGS/UAS-reaper-hid*). Progeny of the latter cross, which were fed RU486 during the second instar, showed no DrmETH1-IR in the tracheal system, consistent with ablation of Inka cells (data not shown). At the time of ecdysis from 2<sup>nd</sup> to 3<sup>rd</sup> instar, these larvae showed phenotypes identical to those observed in the *ETHGS/UAS- $\beta$ ftz-f1*. Affected larvae exhibited the *buttoned-up* phenotype and lethal defects at larval ecdysis (98%; Table 3). Likewise, all 3<sup>rd</sup> instar larvae fed RU486 showed severe defects at pupal ecdysis (Table 3), indicated by incomplete translocation of the air bubble, failure of head eversion and reduced extension of pupal wing pads and legs.

### Proper Timing of $\beta$ FTZ-F1 Expression in Inka cells is Essential for Larval Ecdysis, Pupation, and Adult Development

*$\beta$ FTZ-F1* expression is temporally restricted to a short period prior to each larval ecdysis and pupation, coinciding with decline of the ecdysteroid peak to a low level (Yamada et al., 2000). To assess the importance of stage-specific  *$\beta$ FTZ-F1* expression in Inka cells, we analyzed the effect of its premature and prolonged expression on larval development and ecdysis in *ETHGS/UAS- $\beta$ FTZ-F1* flies.  *$\beta$ FTZ-F1* expression in Inka cells was activated by raising newly eclosed 2<sup>nd</sup> instar larvae on diet containing RU486. Premature  *$\beta$ FTZ-F1* expression in Inka cells did not affect feeding and larval growth, but resulted in severe ecdysis defects. Almost one-third of these larvae (average 30%) failed to execute ecdysis and exhibited the same morphological “*buttoned-up*” phenotype as observed in  *$\beta$ FTZ-F1* -silenced animals shown in Table 1: double mouth hooks, double vertical plates, and new cuticle, spiracles and tracheae for the next larval instar (Table 4). Survivors continued to receive RU486 in the diet throughout the 3<sup>rd</sup> instar. All larvae in this group showed normal growth and puparium formation, but most (96%) failed to progress through pupal ecdysis and died (Table 4). Phenotypes included failure of air bubble translocation, head eversion and elongation of wings and legs. Three survivors eclosed to the adult stage, but failed to expand wings as observed in  *$\beta$ FTZ-F1*-silenced flies (Table 4). These results show that premature expression of  *$\beta$ FTZ-F1* in Inka cells disrupts ecdysis with lethal consequences at all stages (larval, pupal, adult).

### Targeted $\beta$ FTZ-F1 Expression in Inka Cells Rescues *$\beta$ ftz-f1* Deletion Mutants

Homozygous  *$\beta$ ftz-f1<sup>ex7</sup>* progeny are embryonic lethal, but can be rescued by temporally restricted heat treatment, resulting in ectopic expression of  *$\beta$ FTZ-F1* and progression to the first instar (Yamada et al., 2000). These rescued larvae are arrested at ecdysis to the 2<sup>nd</sup> instar, exhibiting the *buttoned-up* phenotype unless rescued by a second heat treatment. While  *$\beta$ FTZ-F1* is expressed widely in *Drosophila* tissues (Lam and Thummel, 2000), we have shown here that similar lethal phenotypes are caused by Inka cell-specific knockdown in *ETHGS/UAS- $\beta$ ftz-f1* flies. This suggests that failure of  *$\beta$ FTZ-F1* expression in Inka cells may be responsible for the most severe defects in  *$\beta$ ftz-f1* mutants. To test this hypothesis, we asked whether  *$\beta$ ftz-f1<sup>ex7</sup>* mutants could be rescued through temporally restricted, Inka cell-selective  *$\beta$ FTZ-F1* expression using GeneSwitch.

We compared viability of  *$\beta$ ftz-f1<sup>ex7</sup>* null flies treated by heat shock (hs), which induces  *$\beta$ FTZ-F1* expression in all tissues versus Inka cell-selective  *$\beta$ FTZ-F1* expression via activation of GeneSwitch. To accomplish this, we collected homozygote progeny (*yw*; *ETHGS/UAS-FTZ-F1*;  *$\beta$ ftz-f1<sup>ex7</sup>hs- $\beta$ FTZ-F1*/ *$\beta$ ftz-f1<sup>ex7</sup>hs- $\beta$ FTZ-F1*) obtained from the following cross: *yw*; *ETHGS/ETHGS*;  *$\beta$ ftz-f1<sup>ex7</sup>hs- $\beta$ FTZ-F1/TM3*, *y<sup>+</sup>* × *yw*; *UAS-FTZ-F1/UAS-FTZ-F1*;  *$\beta$ ftz-f1<sup>ex7</sup>hs- $\beta$ FTZ-F1/TM3*, *y<sup>+</sup>*, using the *y<sup>-</sup>* marker for selection. First instar larvae were rescued through the embryonic stage by heat treatment 13-16 hours after egg laying (AEL). Surviving 1<sup>st</sup> instar larvae were transferred during the time interval 42-46



hours AEL into a new dish containing food with RU486 (100-200  $\mu\text{g/ml}$ ) to elicit  $\beta\text{FTZ-F1}$  expression only in Inka cells. In parallel experiments, larvae with the same genotype were placed into a dish containing standard diet and heat treated during 42-46 hours AEL, which resulted in organism-wide expression of  $\beta\text{FTZ-F1}$ . The number of 1<sup>st</sup> and 2<sup>nd</sup> instar larvae was counted during 54-60 hours AEL. We found that RU486 treatment rescued  $\sim 77\%$  to the 2<sup>nd</sup> instar, similar to 80% rescue of larvae by the ectopic expression of  $\beta\text{FTZ-F1}$  following heat treatment (Table 5). These results show that restricted expression of  $\beta\text{FTZ-F1}$  in Inka cells is sufficient for rescue of the  $\beta\text{ftz-f1}^{ex7}$  null mutant.

As a control experiment, homozygous embryos from  $yw; \beta\text{FTZ-F1}^{ex7}hs-\beta\text{FTZ-F1}/TM3, y^+$  parents were first rescued by heat shock during 13-16 hours AEL. Surviving 1<sup>st</sup> instar larvae were exposed again either to heat shock or RU486 during the developmental interval 42-46 hours AEL. The number of 1<sup>st</sup> or 2<sup>nd</sup> instar larvae was tabulated during 54-59 hours AEL. The proportion of larvae rescued by heat shock (79%) is similar to that obtained in the above described experiment, whereas those fed RU486 failed to undergo ecdysis and died as expected (Table 5).

## Discussion

### Lethal Phenotypes of $\beta\text{FTZ-F1}$ Deficiency are Induced by Blocking ETH Signaling

We have shown that GeneSwitch-mediated  $\beta\text{ftz-f1}$  silencing selectively in Inka cells prevents stage-specific responses to 20E in larvae, mid-prepupae, and adults that were observed previously for the  $\beta\text{ftz-f1}^{ex7}$  null mutant (Yamada et al., 2000). Larval phenotypes include duplicate mouthparts, spiracles, and cuticular layers, all of which are ecdysis defects characteristic of ETH deficiency. Identical phenotypes result from Inka cell ablation (this work) and *eth* excision (Park et al., 2002).  $\beta\text{ftz-f1}$  knockdown in Inka cells is further characterized by persistent ETH-IR, suggesting failure to release the peptide. Rescue of these phenotypes by ETH injection lends credence to this view.

We also performed genetic rescue of the  $\beta\text{ftz-f1}^{ex7}$  null mutant using two different methods. It was shown previously that the embryonic lethal genotype  $\beta\text{ftz-f1}^{ex7}hs-\beta\text{FTZ-F1}$  could be rescued to the first instar by heat treatment. Although these survivors fail to advance to the 2<sup>nd</sup> instar, a second heat treatment rescues a majority ( $\sim 70\%$ ) of them (Yamada et al., 2000). By incorporating ETH-GeneSwitch and  $\beta\text{FTZ-F1}$  into this fly line, we were able to compare rescue efficiencies of organism-wide  $\beta\text{FTZ-F1}$  expression by heat treatment with Inka cell-specific  $\beta\text{FTZ-F1}$  expression. Our results indicate that larval mortality at the end of the first instar can be rescued equally well by heat treatment (80%) and Inka cell specific expression of  $\beta\text{ftz-f1}$  (77%). These results provide further evidence that the major defects arising from the  $\beta\text{ftz-f1}^{ex7}$  null mutant are ecdysis related and stem from ETH deficiency.

Inka cell-specific  $\beta\text{ftz-f1}$  silencing also phenocopies the  $\beta\text{ftz-f1}^{ex7}$  null mutation at the mid-prepupal to pupal transition, which involves air bubble translocation (pre-ecdysis), head eversion (ecdysis) and elongation of incipient wings and legs (post-ecdysis). Despite normal muscle morphology,  $\beta\text{ftz-f1}$  mutants fail to perform abdominal contractions necessary to generate internal hydrostatic pressure for the prepupal-pupal transition. These observations can be explained by block of ETH release required for activation of motor circuitry driving abdominal musculature at the appropriate developmental time (Fortier et al., 2003). The entire ecdysis sequence is triggered by ETH via a downstream receptor-mediated signaling cascade of central peptidergic neurons that express ETH receptors (ETHR) (Kim et al., 2006b; 2006a; Park et al., 2002). ETHR neurons are organized as sequentially recruited peptidergic ensembles that release kinin, diuretic hormones, FMRFamides, eclosion hormone (EH), crustacean cardioactive peptide (CCAP), myoinhibitory peptide (MIP) and bursicon (Kim et al., 2006b; 2006a). These ensembles recruit successive steps in the ecdysis

sequence; their disruption by cell-specific ablation causes defects similar to those observed in *βftz-f1*-null mutants and in flies in which *βftz-f1* is silenced selectively in Inka cells (Clark, 2004; Kim et al., 2006b; Loveall and Deitcher, 2010; McNabb et al., 1997; J. H. Park, 2003; Peabody et al., 2008). For example, targeted ablation of the CCAP/MIPs/bursicon ensemble causes failure of head eversion and incomplete wing and leg extension, most likely due to insufficient blood pressure and cuticle extensibility normally conferred by co-release of these peptides (Kim et al., 2006b; J. H. Park, 2003). Further studies with pburs null mutants showed that this gene is mainly responsible for these phenotypes (Lahr et al., 2012). In the case of Inka cell specific *βftz-f1* knock-down flies, most animals show the same phenotypes as described above and die during pupal ecdysis, while a small number of surviving adults fail to expand the wings. These defects are most likely explained by failure to recruit CCAP/MIP/bursicon neurons, normally activated by ETH (Kim et al., 2006b).

It is important to note that the phenotype associated with *βftz-f1* loss of function mutants could be explained partially by defects in remodeling of motor neurons innervating larval muscles during metamorphosis. Proper pupal ecdysis requires activation of EcR-B1 expression in muscles controlled by the *ftz-f1/Hr39* nuclear receptor pathway. Production of TGF-β/BMP ligands during muscle degeneration provides a signal that initiates retraction of motor neurons (Boulanger et al., 2012). However this cannot explain lethal phenotypes observed in *ETHGS/UAS-βftz-f1i* knockdown prepupae deficient in βFTZ-F1 protein only in Inka cells. Equally severe developmental defects clearly demonstrate the importance of Inka cell peptides in activation of neuronal networks controlling key processes in pupation.

Similarity of phenotypes resulting from deficiencies in βFTZ-F1 and ETH indicates they are signals in a common pathway initiated by 20E at the beginning of the molt and culminating with the ETH-driven behavioral cascade that terminates molting through completion of the ecdysis sequence.

### Correct Timing of βFTZ-F1 Expression and ETH Release is Critical for Normal Development

Previous experiments with *βftz-f1<sup>ex7</sup>hs-βftz-f1* null mutant flies demonstrated that developmental arrest could be rescued by heat-induced βFTZ-F1 expression, but only at developmental times corresponding to normal expression of βFTZ-F1; premature heat treatment was ineffective for rescue (Yamada et al., 2000). Likewise, our results show that, while Inka cell-specific βFTZ-F1 expression at the appropriate time rescues *βftz-f1<sup>ex7</sup>* null mutants, premature expression of βFTZ-F1 in Inka cells leads to high mortality. Such mortality is likely due to premature ETH release. We have shown previously that premature ETH injection fails to rescue ETH-null mutants (Park et al., 2002). We also showed in the moth, *Manduca sexta* that premature injection of ETH into pharate larvae is lethal (Zitnanová et al., 2001). Taken together, these observations suggest that temporal pattern of βFTZ-F1 expression is critically important in determining the correct timing of ETH release.

### βFTZ-F1 Confers Secretory Competence in Inka Cells

In *Manduca sexta*, release of ETH from Inka cells requires decline of 20E to low levels (0.1 μg/ml; 210 nM) and new gene expression in the hours preceding ecdysis (Kingan and Adams, 2000). Prior to this time, Inka cells have not yet acquired competence to release ETH in response to eclosion hormone. Among the genes newly expressed during this period is *βftz-f1* (Hiruma and Riddiford, 2001).

In *Drosophila*, a similar pattern of steroid decline and βFTZ-F1 expression occurs at the end of each developmental stage prior to ecdysis, namely during the hours 45-47 hours AEL (first instar), 69-71 hours AEL (second instar), and 8-12 hours after pupariation (Yamada et

al., 2000). The null mutant  $\beta ftz-f1^{ex7}hs-\beta FTZ-F1$  is arrested at the first ecdysis, but can be rescued by heat treatment to the second instar. Rescued animals become arrested at the transition from 2<sup>nd</sup> to 3<sup>rd</sup> instar, but can be rescued again by heat treatment. Surviving animals fail to pass through the pre-pupal to pupal transition unless heat-treated again. These treatments induce  $\beta FTZ-F1$  expression in all tissues that normally express this gene. Quite striking are the results presented here, showing that the  $\beta ftz-f1^{ex7}hs-\beta FTZ-F1$  null mutant line is rescued by Inka cell-specific expression of  $\beta FTZ-F1$ . Although not all of the cellular phenotypes (e.g., salivary gland histolysis) associated the  $\beta FTZ-F1$  knockout line may be rescued by Inka cell expression alone, it is nevertheless noteworthy that such restricted expression of  $\beta FTZ-F1$  is sufficient to advance development to the next stage.

We conclude that a major physiological outcome downstream of transient, precisely timed expression of  $\beta FTZ-F1$  is ETH secretion by Inka cells. Based on our findings,  $\beta FTZ-F1$  appears to be both necessary and sufficient to confer secretory competence in Inka cells at stage-specific times to advance development.

### 20E-induced Signaling Cascades in the Inka Cell

Although the precise sequence of molecular events leading to Inka cell secretory competence following  $\beta FTZ-F1$  expression is unknown, previous findings shed some light on signal transduction in this endocrine cell. We propose a model combining evidence gained from studies on moths and flies (see graphical abstract). The surge of ecdysteroid levels that initiates changes in gene expression, including induction of the ETH gene via EcR and CRC (Gauthier and Hewes, 2006; Gauthier et al., 2012; Zitnan et al., 1999). High ecdysteroid levels induce expression of the gene *eth* (downward curved arrow) and inhibit expression of *βftz-f1* (downward inhibitory bar); only after 20E levels fall to low levels (straight downward arrow) does *βftz-f1* expression occur, allowing for acquisition of secretory competence. We have shown in this paper that silencing of *βftz-f1* prevents ETH release, leading to ecdysis defects larvae, pre-pupae, and adults. Injection of ETH or targeted expression of *βftz-f1* in Inka cells rescues these phenotypes.

Studies in *Manduca sexta* and *Bombyx mori* showed that Inka cells release ETH in response to corazonin and EH. Corazonin released from Ia<sub>1</sub> neurons in the brain induces an initial, low level secretion of ETH, which acts back on the CNS to elicit the release of EH from VM neurons (Ewer et al., 1997; Kim et al., 2004; Kingan et al., 1997). EH induces elevation of cGMP in Inka cells, followed by massive release and complete depletion ETH. Pharmacological manipulations of Inka cells using various kinase and phosphatase inhibitors revealed Ca<sup>2+</sup> and cGMP-mediated pathways are likely to be involved in EH-induced secretion of ETH in *M. sexta* (Kingan, 2001). Identification of membrane bound guanylyl cyclases as EH receptors confirmed a cGMP-mediated transduction pathway for this neuropeptide also in flies (Chang et al., 2009). Elevation of cGMP in Inka cells at ecdysis thus provides evidence in support of EH-induced ETH release in *Drosophila* (Clark, 2004). Further support for neuropeptide regulated release of ETH comes from detection of corazonin and EH receptor expression in Inka cells of moths and flies (Chang et al., 2009; Kim et al., 2004; Zitnan and Adams, 2013). However, EH alone appears not to be essential for ETH secretion in *Drosophila*, since EH knockout flies complete ecdysis even in the absence of this neuropeptide and cGMP elevation in Inka cells. In these flies ETH release may be induced by corazonin. Thus far, injections of corazonin or EH in *Drosophila* failed to induce premature ETH secretion and initiation of the ecdysis sequence (Clark, 2004). It is possible that this failure is related to a very brief window of neuropeptide receptor expression induced by  $\beta FTZ-F1$ . We speculate that brief appearance of receptors for corazonin and EH shortly before ecdysis initiation may be crucial for acquisition of Inka cell secretory competence mediated by  $\beta FTZ-F1$ .

## Conclusions

Our findings provide precise physiological outcomes resulting from 20E-induced  $\beta$ FTZ-F1 expression in insect development and help to explain various lethal phenotypes associated with  $\beta$ FTZ-F1-null mutants in different developmental stages (Lam and Thummel, 2000; Yamada et al., 2000). Furthermore, they implicate the Inka cell as a key cellular “gateway” for stage-specific outcomes induced by 20E. Further investigation of the precise actions of  $\beta$ FTZ-F1 in Inka cell signal transduction will hopefully reveal molecular mechanisms underlying secretory competence in endocrine cells.

## Acknowledgments

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## Abbreviations

<b>ETH</b>	ecdysis triggering hormone
<b>ETH-IR</b>	ecdysis triggering hormone-like immunoreactivity
<b>EH</b>	eclosion hormone
<b>DrmETH</b>	<i>Drosophila melanogaster</i> ETH
<b>ETHGS</b>	ETH-GeneSwitch

### Highlights

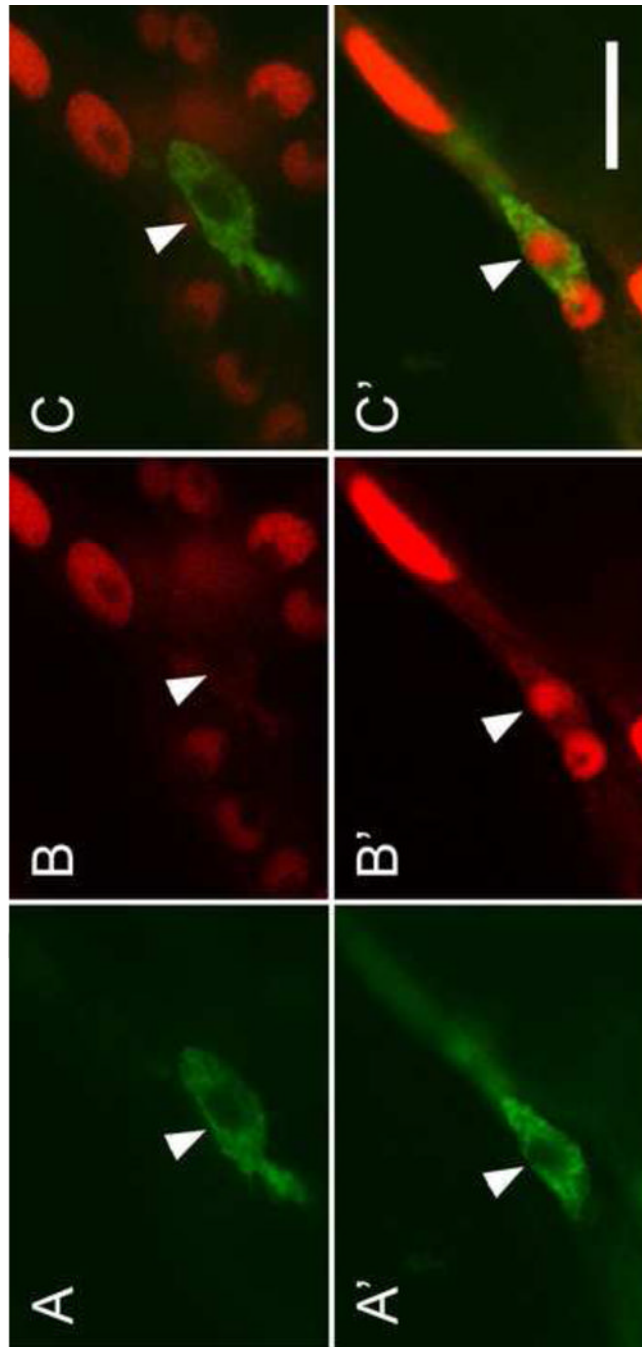
- Silencing of *βftz-f1* in Inka cells blocks ETH release and arrests development.
- Developmental arrest rescued by ETH injection.
- *βftz-f1* null mutants rescued by Inka cell-specific expression of *βftz-f1*.
- The Inka cell provides a “gateway” endocrine output of ecdysone signaling.



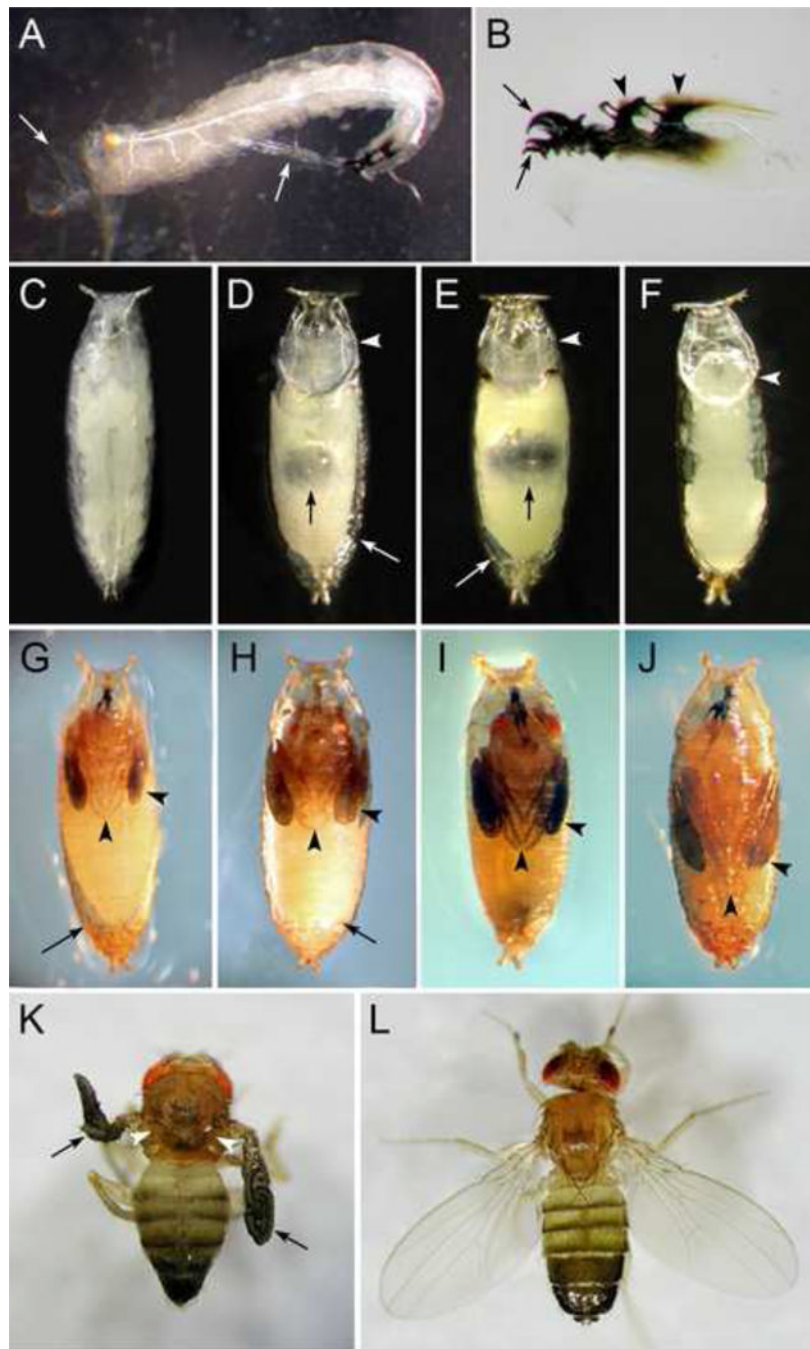
**Fig. 1. Inka cell-specific expression of  $\beta FTZ-F1$  in  $ETHGS/UAS-\beta FTZ-F1$  flies**

Targeted  $\beta FTZ-F1$  expression in Inka cells (arrows) was induced by feeding RU486 to  $ETHGS/UAS-\beta FTZ-F1$  larvae. (A) Tracheae were dissected from feeding 3<sup>rd</sup> instar larvae 2 hours later and processed for *in situ* hybridization with a  $\beta FTZ-F1$  DNA probe. (B) Specific  $\beta FTZ-F1$  expression in Inka cells was confirmed by subsequent immunostaining with an antiserum to DrmETH. (C) Merged A and B. Scale bar, 30  $\mu m$ .





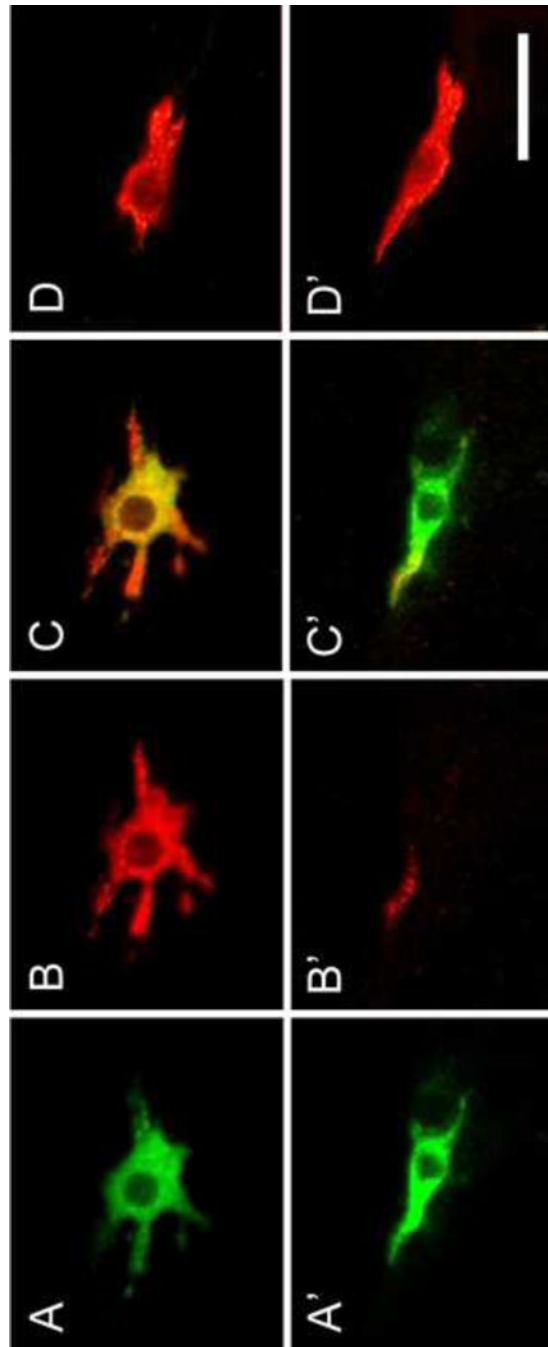
**Fig. 2.  $\beta$ FTZ-F1 expression is reduced selectively in Inka cells of ETHGS/UAS- $\beta$ FTZ-F1i flies**  
 Third instar larvae were reared on diet containing RU486. Tracheae were dissected 8 hours after puparium formation and processed for immunostaining with antisera against ETH and  $\beta$ FTZ-F1. (A) Inka cell displaying ETH-IR in the cytoplasm (green; arrowheads), but not in the nucleus. (B) Lack of  $\beta$ FTZ-F1-IR in nuclei of Inka cells in RNA-silenced flies. (C) A and B merged. (A'-C') Inka cells of control animals reared on standard diet showed ETH-IR in the cytoplasm (A') and  $\beta$ FTZ-F1-IR in the nucleus (B'). (C') Merged A' and B'. Scale bar, 30  $\mu$ m.



**Fig. 3. Targeted  $\beta$ FTZ-F1 silencing selectively in Inka cells leads to ecdysis failure throughout development**

(A) Ecdysis failure and “buttoned-up” phenotype in late 2<sup>nd</sup> instar larva; ecdysis failure is indicated by partially shed old cuticle (arrows) and death. (B) Failure to shed old mouthparts indicated by double mouth hooks (arrows) and double vertical plates (arrowheads). (C) Surviving larvae show normal pupariation indistinguishable from controls. (D, E) Pupal ecdysis failure, indicated by incomplete translocation of the air-bubble (black arrows), lack of head eversion (arrowheads) and gaps between pupal abdomen and puparium (white arrows). (F) A freshly pupated wild-type (*W1118*) pupa with completed head eversion (white arrowhead). (G-I) Arrested development after head eversion. (G) Developmental

defects, indicated by lack of pigmentation, failure to elongate wings and legs (arrowheads), and a gap between the abdomen and puparium (arrow). (H) Successful development and pigmentation of the head and thorax, while wings and legs are short (arrowheads) and the abdomen exhibits persistent early prepupal phenotype (arrow). (I) Successful development and pigmentation, but failure to elongate wings and legs (arrowheads). (J) Normal development and elongated appendages (arrowheads) in a wild-type (*W1118*) pharate adult. (K) A few emerged adults with silenced *βFTZ-F1* in Inka cells fail to expand their wings (arrows) and contain dimples on the dorsal thorax (arrowheads). (L) Normally developed control adult (*W1118*).



**Fig. 4. Persistent ETH-IR in Inka cells of *ETHGS/UAS- $\beta$ ftz-*flt** flies**

*ETHGS* flies were crossed with UAS-EGFP line for Inka cell visualization at dVP larval stage (A-C) and just after ecdysis (A'-C'). A, A'; anti-GFP staining (green, Alexa 488). B, B'; The same tracheae were stained with ETH antiserum to detect ETH levels before and after ecdysis (red, Alexa 555); C, C; merged images. Note that ETH was depleted after ecdysis (B'). (D, D') In  *$\beta$ FTZ-F1* knock-down flies, ETH staining showed no difference between Inka cells in dVP stage (D) and arrested larvae 3 hr after appearance of dVP (D'; red, Alexa 555). Scale bar, 25  $\mu$ m.

**Table 1**Lethality and developmental defects resulting from *βFTZ-F1* silencing in Inka cells.

Fly lines	n	Accumulated lethality		Short wings/normal
		Pharate 3 <sup>rd</sup> instar	Pharate pupae	Adults
<b>RU486 +</b>				
ETHGS × UAS-βFTZi	68	32 (47)	34 (97)	2/0
ETHGS × UAS- <i>reaper-hid</i>	53	52 (98)	1 (100)	0/0
ETHGS × UAS-EGFP	60	0	0	0/60
<b>RU 486 -</b>				
ETHGS × UAS-βFTZi	50	0	0	0/50
ETHGS × UAS- <i>reaper-hid</i>	45	1	0	0/44
W1118	60	0	0	0/60

\* Numbers in parentheses correspond to percent mortality at each stage.

**Table 2**

Rescue of larvae with targeted  $\beta$ FTZ-F1 knockdown in Inka cells by DrmETH1 injection into late 2<sup>nd</sup> instar larvae.

Fly lines	n	Ecdysis
ETHGS $\times$ UAS- $\beta$ FTZi (100 fmol DrmETH1)	20	19
ETHGS $\times$ UAS- $\beta$ FTZi (dist. water)	20	9

**Table 3**

Lethality and developmental defects of flies with ablated Inka cells.

Fly lines	<i>n</i>	Accumulated lethality		Short wings/normal
		Pharate 3 <sup>rd</sup> instar	Pharate pupae	
<b>RU486 +</b>				
ETHGS × UAS- <i>reaper-hid</i>	57 (2 <sup>nd</sup> )	56 (98)	1 (100)	0/0
ETHGS × UAS- <i>reaper-hid</i>	64 (3 <sup>rd</sup> )	-	64 (100)	0/0
<b>RU 486 -</b>				
ETHGS × UAS- <i>reaper-hid</i>	42 (2 <sup>nd</sup> )	1	0	0/41
ETHGS × UAS- <i>reaper-hid</i>	38 (3 <sup>rd</sup> )	-	0	0/3

\* Numbers in parentheses correspond to percent mortality at each stage.

**Table 4**Lethality and developmental defects caused by premature expression of  $\beta FTZ-F1$  in Inka cells.

Fly lines	n	Accumulated lethality		Short wings/normal
		Pharate 3 <sup>rd</sup> instar	Pharate pupae	
<b>RU486 +</b>				
ETHGS × UAS- $\beta FTZ$	50	13 (26%)	35 (95%)	2/0
ETHGS × UAS- $\beta FTZ$	42	14 (33%)	27 (97%)	1/0
ETHGS × UAS-EGFP	54	1	2	0/51
<b>RU 486 -</b>				
ETHGS × UAS- $\beta FTZ$	47	0	0	0/47
W1118	64	0	0	0/64



Table 5

Rescue of the  $\beta ftz-f1^{ex7}hs-\beta FTZ-F1$  null mutant larvae by targeted  $\beta FTZ-F1$  expression in Inka cells.

Total number of larvae used after hs of embryos						
	1 <sup>st</sup> instar	1 <sup>st</sup> instar	2 <sup>nd</sup> instar	rescued		
<i>ETHGS/UAS-βFTZ-F1; ftz-f1<sup>ex7</sup>hs-βFTZ-F1</i>						
hs	59	hs	11	48	81%	
hs	67	hs	14	53	79%	
hs	56	RU486	11	45	80%	
hs	59	RU486	15	44	75%	
hs	90	RU486	22	68	76%	
<i>ftz-f1<sup>ex7</sup>hs-βFTZ-F1</i>						
hs	87	hs	18	69	79%	
hs	75	RU486	75	0	0%	

Embryos were rescued by heat shock (hs) during 13-16 hours AEL. Surviving 1<sup>st</sup> instar larvae were heat treated or fed RU486 treated during 42-46 hours AEL. The number of 1<sup>st</sup> or 2<sup>nd</sup> instar larvae was counted during 54-60 hours AEL.