Timing is Everything

PTTH Mediated DHR4 Nucleocytoplasmic Trafficking Sets the Tempo of Drosophila Steroid Production

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Timing is everything: PTTH mediated DHR4 nucleocytoplasmic trafficking sets the tempo of *Drosophila* steroid production

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During development, multicellular organisms must become sexually mature in order to reproduce. The developmental timing of this transition is controlled by pulses of steroid hormones, but how these pulses are generated have remained unclear? A recent paper shows that in *Drosophila* larvae, nucleocytoplasmic trafficking of DHR4, a nuclear receptor, in response to prothoracotropic hormone signaling, is critical for producing the correct temporal pulses of steroid hormones that coordinate the juvenile–adult transition.

**Keywords:** 20-hydroxyecdysone, DHR4, critical weight, Cyp6t3, metamorphosis

In many animals reaching a species-specific size induces a developmental switch from juvenile growth to reproductive maturity. This transition occurs when a developmental timing program is aligned with permissive checkpoint signals, such as nutritional status and photoperiod, to activate a neuroendocrine circuit that promotes steroid-induced maturation in organisms ranging from flies to humans (Dungan et al., 2006; Tennesen and Thummel, 2011).

In some insects, it is thought that attainment of a proper level of nutrient stores, known as the critical weight, plays a major role in activation of neuroendocrine signal(s) that trigger metamorphosis, a process that produces a reproductively mature adult (McBrayer et al., 2007; Mirth and Riddiford, 2007). Although the mechanisms for assessing size/weight are poorly understood (see however Callier and Nijhout, 2011), the output of achieving critical weight in *Drosophila* correlates well with production of an early low-level pulse of the steroid hormone ecdysone (E), ~8–10 h after ecysis from the second to the third instar stage. This is followed by at least two other low-level E pulses midway through the third instar stage, and finally by a much larger E peak just as pupariation commences (Warren et al., 2006; Mirth and Riddiford, 2007). The mechanisms responsible for generating the initial critical weight-associated pulse, and the importance of this and the other low-level E pulses for orchestrating the proper timing of the late major larval E pulse, that initiate metamorphosis, have remained quite vague.

In a recent paper published in PLoS Biology, Ou et al. (2011) begin to provide some insight into this issue by demonstrating that the trafficking of the nuclear receptor DHR4 between the nucleus and the cytoplasm in response to a brain-derived neuropeptide known as prothoracicotropic hormone (PTTH) controls the dynamics of the E peaks and the timing of metamorphosis.

**PTTH SIGNALING AND METAMORPHOSIS**

Previous studies in both Lepidoptera and *Drosophila* have demonstrated a key role for PTTH in stimulating E production and metamorphosis (Rybczynski, 2005; McBrayer et al., 2007). In *Drosophila*, ablating the PTTH producing neurons causes substantial developmental delay in the onset of pupariation. In contrast, over-expression of PTTH leads to the opposite phenotype, i.e., early onset or precocious pupariation (McBrayer et al., 2007; Rewitz et al., 2009b). PTTH stimulates E production by binding to its receptor Torso, a receptor tyrosine kinase, which is expressed in prothoracic gland (PG) cells, the endocrine tissue of the larva responsible for E production. Torso activates the Ras/Raf/Mek pathway resulting in the phosphorylation and targeting of phosphorylated Erk (pErk) to the nucleus. Interestingly, expression of an activated form of Ras in the PG induces precocious pupariation similar to what is seen when PTTH is expressed ubiquitously (Rewitz et al., 2009b).

**DHR4 IS A KEY MEDIATOR OF PTTH SIGNALING AND IS ESSENTIAL TO PREVENT PREMATURE METAMORPHOSIS**

A major block to a more comprehensive understanding of how the PTTH signal stimulates E production in the PG and times the onset of metamorphosis has been the lack of knowledge concerning downstream mediators of pathway activity, especially potential transcription factors that are targets for phosphorylation by Erk. Previous work on the nuclear receptor *DHR4* had shown...
that larvae carrying mutations in this gene undergo precocious
maturation, reducing the juvenile growth period and producing
small animals (King-Jones et al., 2005) similar to that caused by
hyper-activating PTTH signaling. Importantly, Ou et al. (2011)
demonstrated that the developmental timing defects exhibited by
DHR4 mutants are phenocopied by specific loss of DHR4 in the
PG cells and not by knockdown of DHR4 in the fat body, another
tissue in which DHR4 shows strong expression. These observations
led to the question of whether there might be a functional connec-
tion between PTTH signaling and DHR4 activity. As illustrated
in Figure 1, they discovered a very interesting inverse correla-
tion between PTTH mediated Erk nuclear localization and the
subcellular distribution of DHR4 in the PG cells. In the absence
of PTTH signaling, DHR4 was found primarily in the nucleus,
while constitutive PTTH signaling resulted in mainly cytoplasmic
DHR4. While it was not shown that Erk directly phosphorylates
DHR4, this seems highly likely since simple inspection of the
DHR4 primary sequence reveals the presence of several consensus
Erk phosphorylation sites.

The inverse correlation of Erk and DHR4 nuclear localization
suggests that the primary role of nuclear DHR4 in the PG is to
dampen the PTTH signal to prevent early metamorphosis. This
hypothesis was confirmed with an epistasis experiment where
over-expression of DHR4 in the PG was found to suppress the
early pupation phenotype of activated Ras expression in the PG.
Furthermore, using carefully timed heat shock pulse-driven DHR4
RNAi, the authors were able to demonstrate that the crucial time
at which DHR4 is required to prevent precocious metamorpho-
sis is around the time of the early third instar when the critical
weight and the low-level E pulses are determined. Intriguingly,
they found that in DHR4-depleted larvae, the E titer rose faster
and did not decline as in the controls leading to a step-like appear-
ance in the E titer profile instead of a discrete pulse. Taken together,
these data indicate that DHR4 is a key target of PTTH signaling
in the PG and provide the first compelling evidence that the small
early third instar E peak is essential for setting the tempo of larval
development.

Cyp6t3: A KEY ECDYSONE BIOSYNTHETIC COMPONENT AND
TARGET OF DHR4 REPRESSION?

The observation that PTTH signaling appears to upregulate E
titers by removing DHR4 from the nucleus leads to the obvious
questions of what genes DHR4 regulates and how they impact E
synthesis or degradation? To begin to address this question, Ou
et al. (2011) used a microarray approach in which they specifi-
cally isolated RNA from carefully timed wildtype Drosophila ring
glands, a tissue that contains the PG cells, and compared the
profiles to similarly timed glands in which the level of DHR4
expression was reduced by RNAi knockdown. This technically
demanding experiment was able to identify a small number of genes (∼50) that showed either enhanced or reduced expression
upon loss-of-DHR4 function. The most interesting among these,
with respect to E metabolism, were several Cyp gene family mem-
bers that code for P450 enzymes of which several are required for
either synthesis or degradation of E (Gilbert and Warren, 2005;
Rewitz et al., 2010). Intriguingly, none of the known E biosyn-
thetic P450s showed much change in expression in response to
loss-of-DHR4 function, at least not at the time points that were
examined. However, two members of the Cyp6 family, including
Cyp6t3, showed significant upregulation upon DHR4 depletion.
The authors focused their analysis on Cyp6t3 since its expres-
sion was restricted to the ring gland and showed oscillations
that correspond to the nucleocytoplasmic oscillations of DHR4.
Knockdown of Cyp6t3 in the PG was found to produce E defi-
ciency phenotypes that could be rescued by feeding larvae E or
one of several E biosynthetic precursors. These experiments tenta-
tively place Cyp6t3, along with Spook, another P450, and Shroud
a short chain dehydrogenase, at the mysterious “black box” step
that catalyzes conversion of 7-dehydrocholesterol to diketol by an
unknown mechanism. Lastly, epistasis experiments showed that

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**FIGURE 1 | Prothoracicotropic hormone/DHR4 generate cyclic steroid pulses that drive directional developmental transitions.** A model illustrating the mechanisms generating 8–16 h ultradian steroid oscillations. (A) Each cycle is initiated by the release of PTTH from terminals of brain neurons that activates the Ras pathway in the PG. This results in the phosphorylation and nuclear localization of pERK and translocation of DHR4 to the cytoplasm, which derepresses ecdysone biosynthesis. (B) In the presence of PTTH, the DHR4 repressor is displaced from promoters, including Cyp6t3 increasing its expression. During periods without PTTH signaling, nuclear DHR4 represses gene expression, although DHR4 may act as an activator in some cases. pErk, phospho-ERK; PG, prothoracic gland; PRE, PTTH response element; PTTH, prothoracicotropic hormone. Torsos: the transmembrane PTTH receptor in the PG. Dashed arrows indicate inactive mechanisms.
upregulation of Cyp6t3 contributes to the developmental timing defect of DHR4 loss since knockdown of Cyp6t3 in a DHR4 mutant background suppressed the precocious pupation phenotype.

Despite these compelling findings suggesting a key role for Cyp6t3 as a new E biosynthetic component, there are some puzzling aspects. First, Cyp6t3 shows no strong conservation among insects, which is unusual, compared to the hitherto characterized E biosynthetic enzymes. For Cyp6t3 there are numerous related Cyp family members in different insect species, however, they are not clear orthologs as can be found for all the other key Cyp E biosynthetic enzymes. Further de novo synthesis of E is required for embryogenesis and oogenesis. While the expression of Cyp6t3 in ovaries and embryos was not reported, data available through the Flybase does not support much expression in these tissues, although it may be very low as is observed in the PG. A full analysis of the null mutant phenotype will be necessary to address a possible role in embryogenesis and either clonal or RNAi knockdown in follicle cells to address a role in ovaries. Even if cyp6t3 is not required for E biosynthesis in either of these tissues, this might once again simply reflect a division of labor with a related Cyp6 family member that is expressed during embryogenesis and in the ovary. In fact, this same situation occurs for Spook and Spookier, two essential ecdysteroidogenic P450s which, interestingly enough, are also implicated at the black box step. In this case, spook is expressed in embryos and ovaries and its function is dispensable in larvae, while spookier is expressed in the larval PG and its function is not required in embryos and presumably the ovary as well. However, in this example Spook and Spookier are clear paralogs; no similarly related as well. However, in this example Spook and Spookier are clear paralogs; no similarly related orthologs as can be found for all the other key Cyp E biosynthetic enzymes. Further de novo synthesis of E is required for embryogenesis and oogenesis. While the expression of Cyp6t3 in ovaries and embryos was not reported, data available through the Flybase does not support much expression in these tissues, although it may be very low as is observed in the PG. 

The work described by Ou et al. (2011) undoubtedly demonstrates that DHR4 is an important component of the network regulating the timing of the steroid cues that drive the gene circuit underlying time-directional development in Drosophila. Like all good breakthroughs however, the reported experiments raise as many interesting questions as they answer. For example, what aspect of the altered E titer profile in DHR4 mutants is key to accelerating development? Is it the more rapid kinetics of its production, its ability in integrate the total E signal over time. There is precedence for this simply as the result of gland growth, and the larvae were able to ally, the large late-larval E peak would not occur in the absence of PTTH. However, the answer might be surprising if, for example, there was a steady increase in E in the double RNAi larvae, perhaps simply as the result of gland growth, and the larvae were able to integrate the total E signal over time. There is precedence for this ability in Manduca, where a prolonged continuous small E dose can have the same effect as a larger pulsed dose (Nijhout, 1976). Signal integration of a steady E ramp up may enable the larvae to still initiate metamorphosis with close to normal timing. The results by Ou et al. (2011) provide important insights into regulation of the steroid pulse generator that, from an evolutionary perspective, seems to be a key part of the neuroendocrine system coordinating the juvenile–adult transition in animals.

WHAT’S NEXT?
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