Title:

Essential Function of Membrane-Bound Transcription Factor MYRF in Promoting Transcription of miRNA lin-4 during C. elegans Development

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Abstract

Precise developmental timing control is essential for organism formation and function, but its mechanisms are unclear. In *C. elegans*, the microRNA *lin-4* plays a critical role in developmental timing. While *lin-4*’s downregulation of LIN-14 is well-established, the mechanisms behind *lin-4* upregulation remain unknown. We demonstrate that the membrane-associated transcription factor MYRF-1 is necessary for *lin-4* upregulation in late L1 stage. MYRF-1 is initially localized on the cell membrane, and its increased cleavage and nuclear accumulation coincide with *lin-4* expression timing. We show that MYRF-1 regulates *lin-4* expression cell-autonomously and that hyperactive MYRF-1 can prematurely drive *lin-4* expression in early L1 and embryos. The tandem *lin-4* promoter DNA recruits MYRF-1GFP to form visible loci in the nucleus, suggesting that MYRF-1 directly acts on the *lin-4* promoter. Our findings identify a crucial link in understanding developmental timing regulation and establish MYRF-1 as a key regulator of *lin-4* expression.
Introduction

It is during the embryonic stage that the basic form and pattern of organisms are mostly established. However, the majority of growth in animals occurs during post-embryonic stages[1]. Some extreme examples are seen in insects undergoing metamorphosis and vertebrate amphibians[2]. In many cases, there is often a considerable increase in body size, but profound transitions take place, such as neural circuitry remodeling throughout juvenile years and sexual maturation[3-6]. Undoubtedly, the genetic program determines how different species grow to different characteristic sizes and forms, yet such mechanisms remain largely unknown. One key question is how developmental timing is controlled.

When hatching out of their egg shells, the nematode C. elegans exhibits a miniature yet grossly similar form to adults, and demonstrates a set of behavioral abilities including sensory response, locomotion, and learning[7-9]. They go through four larval stages before becoming fertile adults, and the transition between two consecutive larval stages is marked by a molting event. Changes in the temporal pattern of development are best exemplified by stage-specific cell division and differentiation of blast cells, as well as stage-specific epidermal cuticle formation fate. Such stage-specific patterns can be skipped or reiterated in so-called heterochronic mutants, and lin-4 and lin-14 are the two that have been studied in details[10-16]. lin-4 mutants exhibit complex cell lineage defects, including reiterated larval stage 1-specific cell division in some progenitors, while lin-14 mutants exhibit precocious patterning where stage-specific events are skipped. The studies of these mutants led to the discovery of the first microRNA regulatory pathway as follows: The ubiquitously expressed nuclear factor LIN-14 promotes L1 patterns and suppresses progression to L2. The microRNA lin-4 is upregulated during late L1, and suppresses LIN-14 post-transcriptionally, consequently initiating progression to L2 pattern[17, 18]. lin-4 - LIN-14 pair not only controls the division pattern of blast cells but also controls the maturation and plasticity of the neural circuit during L1-L2 transition[19-21]; lin-4 is also involved in other diverse biological
processes[22]. The vertebrate ortholog of *lin-4*, known as *miR-125*, has been found to promote neuronal differentiation and maturation[23, 24].

What remains a mystery is the factors that trigger the expression of the *lin-4* microRNA during mid-late L1. It is known that the coding sequence of *lin-4* is embedded within an intron of a host gene, and its transcription uses its own promoter and bound Pol II complexes[13, 17, 25]. Once produced, *lin-4* can self-enhance its own transcription by directly interacting with cis-elements in the promoter region[26, 27]. It is also known that during embryonic stages, the transcription of *lin-4* is suppressed by FLYWCH transcriptional factors, but this suppression only extends until late embryogenesis[28]. What is not known is the regulation that turns on the *lin-4* transcription in late L1.

The onset of *lin-4* expression in late L1 is likely linked to the nutritional state. *C. elegans* needs to feed to initiate the post-embryonic developmental programs of the L1 stage. When newly hatched animals encounter an environment without food, they enter into a diapause state in which development is suspended, and they become more resistant to environmental stress. In the case of epidermal blast cell division, the cycling inhibitors are promoted by more activated FOXO transcription factor DAF-16 due to starvation, which suppresses the blast's division[29, 30]. During L1 diapause, *lin-4* expression is suppressed, but this suppression is largely independent of *daf-16*[30], suggesting that *lin-4* expression onset during mid-late L1 under nutrient-rich environment is not a result of simple attenuation of *daf-16* activity. It is worth noting that external food signals are insufficient to drive *lin-4* expression as *lin-4* doesn't promptly turn on when animals encounter food. Therefore, *lin-4* expression must reflect the combination of the decision of the internal nutritional state and developmental progression.

Another unique developmental event that occurs at late L1 is the synaptic rewiring of DD motor neurons[21, 31, 32]. The process enables a structural and functional switch between dendritic
and axonal domains of DDs without an obvious transition in gross neuronal morphology. We previously identified the membrane-bound transcription factor MYRF-1 as an essential positive regulator of synaptic rewiring in DDs[33]. MYRF-1 is broadly expressed, and myrf-1 loss of function results in animals being arrested at the end of L1 or during L2. Intriguingly, the MYRF-1 protein is localized on the cell membrane during early and mid-L1 but increasingly undergoes self-cleavage towards late L1 in response to unidentified signaling mechanisms[34]. The enzymatic reaction of cleavage is expected to occur after trimerization of full-length MYRF-1, which releases N-MYRF-1 in trimeric form[35, 36], but the regulatory mechanisms underlying the process are unclear. Although MYRF-1’s essential role in regulating the development of animals (including humans) is well-established[37-39], the functional targets of MYRF-1 remain elusive. In the course of studying the genetic interaction between myrf-1 and lin-14, we discovered that MYRF-1 is required for lin-4 expression. We present data demonstrating that MYRF-1 is a direct, cell-autonomous driver of lin-4 expression.

Results

MYRF-1 is required for lin-4 upregulation

The accumulation of lin-4 microRNA occurs during mid to late L1, primarily due to an upregulation of transcription in lin-4 primary RNA, as supported by multiple studies [25, 28][Figure 1A, C]. We confirmed their findings by observing a Plin-4::GFP (mals134) reporter generated by Ambros group[28], consisting of 2.4 kb of DNA sequences upstream of the mature lin-4 fused to GFP. After its on-set, lin-4 expression appears to constitutive and ubiquitous throughout the larval stages (see discussion). MYRF-1 is also expressed broadly in larvae, but unlike lin-4, MYRF-1 transcription is active in both embryos and early L1, with an increase in transcription observed towards late L1[33]. However, the activity of MYRF-1 as a transcription factor is determined by the presence of N-MYRF-1 in the nucleus[34]. Initially, full-length MYRF-1 localizes to the cell membrane during early L1, and only during mid to late L1 is the processing of MYRF-1 cleavage
increased, resulting in an elevated amount of N-MYRF-1 being released and shuttled into the nucleus (Figure 1A, B). Therefore, the nuclear accumulation of N-MYRF-1 coincides with the upregulation of \textit{lin-4} both temporally and spatially.

The mutation \textit{ju1121}(G274R) in the MYRF-1 protein not only abolishes its DNA binding capability but also negatively interferes with its close paralogue MYRF-2, resulting in \textit{myrf-1(ju1121)} displaying phenotypic resemblance to double mutants of \textit{myrf-1} and \textit{myrf-2}[33]. Animals with the \textit{myrf-1(ju1121)} mutation exhibit developmental arrest at L2 and eventually perish during L2-L3 molting. We find that \textit{lin-4} transcription reporter fails to be upregulated in \textit{myrf-1(ju1121)} at any viable stages that can be analyzed (Figure 2A, B). Similarly, another two loss of function mutants of \textit{myrf-1(syb1468)} and \textit{myrf-1(syb1491)} also shows a lack of \textit{lin-4} transcription reporter signals (Figure S1A, B). Therefore, according to the 2.4 kb promoter reporter analysis, it is evident that the upregulation of \textit{lin-4} is dependent on the presence of MYRF.

To confirm the crucial role of MYRF in the upregulation of \textit{lin-4} transcription during late L1 stage, we investigated how \textit{myrf} influences \textit{lin-4} transcription by employing a reporter system with a nuclearly localized mScarlet protein, endogenously inserted at the \textit{lin-4} locus (\textit{umn84}), wherein the reporter open reading frame replaced the primary RNA sequence of \textit{lin-4}. Our observations revealed that mScarlet signals were not detected in embryos and early L1 larvae (Figure 2C, Figure 5A). However, these signals were significantly upregulated during late L1 stage and exhibited even stronger intensity in early L2 stage. Notably, in \textit{myrf-1(ju1121)} mutants, the mScarlet signals were largely absent throughout the body during late L1 stage (14h), except for an interesting presence in 8-9 nuclei in the posterior bulb of the pharynx, with intensity comparable to that observed in the wild-type counterparts (Figure 2C, D, Figure S1C). The overall absence of \textit{lin-4} upregulation persisted in \textit{myrf-1(ju1121)} mutants during early L2 stage (21h), except for the aforementioned subset of pharyngeal nuclei where the mScarlet signals increased, although the ascent was much weaker compared to the wild-type controls.
We carried out qPCR analysis using probes specifically targeting lin-4 microRNA to examine endogenous lin-4 expression in wild type and myrf-1(ju1121) mutants, and observed a significant reduction in the levels of mature lin-4 microRNA in myrf-1(ju1121) mutants (Figure 2E). This decrease of mature lin-4 microRNA has been further confirmed through microRNA sequencing analysis (Figure 7A). These findings, combined with the analysis of endogenous lin-4 reporter, provide compelling evidence supporting the critical role of MYRF in the upregulation of lin-4 during the late L1 stage.

To investigate if MYRF-1 is continuously required for lin-4 transcription after its initial upregulation, we used the auxin-inducible degradation (AID) system[41]. We combined the ubiquitously expressed F-box protein TIR1 with degron-tagged MYRF-1 to acutely deplete MYRF-1 protein. The degradation was induced by treating animals at L2, L3, and L4 stages with an NAA solution, an auxin analog. Our results showed a significant reduction in the lin-4 transcription reporter signals in animals of all tested stages within 10 hours' post-treatment (Figure S2). This demonstrates that MYRF-1 is necessary for lin-4 expression throughout the larval stages.

**Sustained high level of LIN-14 protein in myrf-1 mutants**

The LIN-14 protein is typically present in embryos and early L1 but is downregulated as development progresses. Loss of the microRNA lin-4 leads to sustained high levels of LIN-14 protein throughout larval development[11]. To investigate how myrf-1 mutations may affect LIN-14, we examined the signals of endogenously tagged LIN-14::GFP in myrf-1(ju1121) mutants[40]. The LIN-14::GFP signal remains bright and shows no signs of decreasing at late L1 and beyond in myrf-1(ju1121) mutants (Figure 3A, B), consistent with the low levels of lin-4 microRNA observed in the mutants.
MYRF-1 regulates *lin-4* expression cell-autonomously

Although MYRF is essential for larval development, there is limited understanding of how MYRF specifically affects tissues and cells during development. Identifying *lin-4* as a potential transcriptional target of MYRF-1, we aimed to investigate how MYRF-1 may regulate *lin-4* transcription. The primary question is whether MYRF-1 promotes *lin-4* expression in target cells or whether MYRF-1 acts to modify unidentified systemic signals that subsequently lead to upregulation of *lin-4* in terminal cells.

We conducted tissue-specific rescue experiments by expressing MYRF-1 under the *myo-3* promoter, which is specific to body wall muscles, in *myrf-1(ju1121)* mutants. We observed a significant induction of *lin-4* transcription in body wall muscles but not in other tissues (Figure 4A). We also used the epidermis-specific promoter of *dpy-7*, a collagen, for another MYRF-1 rescue experiment, which resulted in the appearance of *lin-4* transcription reporter signals only in the epidermis. Interestingly, the reporter GFP signals were not detected in the seam cells, a group of specialized epidermal cells embedded in the large syncytium epidermal hpy7 cell. Seam cells are derived from the epidermal lineage and also express the *dpy-7* gene; however, this experiment used a short promoter of *dpy-7*, which is not activated in seam cells, thus excluding the induction of *lin-4* in these cells. These results demonstrate that MYRF regulates *lin-4* transcription autonomously within specific cells.

One possible explanation for the deficiency of *lin-4* upregulation in *myrf-1(ju1121)* mutants is that it may be due to overall developmental arrest in L2. To investigate this possibility, we generated a conditional allele of *myrf-1loxP*(ybq98) using CRISPR-Cas9 editing. We then obtained an epidermis-specific Cre-expressing transgene and combined it with the *myrf-1loxP* allele in animals (Figure 4B, C). Our analysis showed that the *lin-4* transcription reporter signals in the epidermis are lost in animals with the combined Cre transgene and *myrf-1loxP* allele, while the signals were
still present in all surrounding tissues. These results provide further evidence that MYRF-1 promotes \textit{lin-4} upregulation cell autonomously and suggest that MYRF-1 acts directly in terminal tissues to regulate cell development.

\textbf{Hyperactive form of MYRF-1 drives premature expression of \textit{lin-4}}

The next question was whether MYRF-1 is sufficient to drive the upregulation of \textit{lin-4}. Because of the intricate nature of regulated MYRF-1 cleavage, overexpressing full-length MYRF-1 by transgene does not enhance its transcriptional activity effectively, as excess MYRF-1 is unable to traffic to the cell membrane or undergo adequate cleavage\cite{33}. Overexpressing N-MYRF-1 alone is also insufficient as it unlikely forms trimers efficiently. In another study, we have identified elements of the cleavage mechanism, which we will report in another manuscript. Using this information, we created a truncated form of MYRF-1 (deleting 601-650) that is expected to bypass the need for cell-membrane trafficking and circumvent the developmental signals that control cleavage. We expected that overexpressing this modified MYRF-1 would produce abundant N-MYRF-1 in trimeric form and enhance its endogenous function.

In this experiment, we used the endogenously mScarlet-tagged line to report the transcription activity of \textit{lin-4 (umn84)}. As mentioned earlier, mScarlet expression is not detected in embryos and early L1 larvae (Figure 2C, Figure 5A). We find that overexpressing the hyperactive MYRF-1 (Δ601-650) driven by the ubiquitous promoter \textit{rpl-28} is detrimental to larval development, as all F1 progeny with the transgene arrest at early larval stages. Strikingly, expressing this hyperactive MYRF-1 variant causes upregulation of \textit{lin-4} transcription reporter in both embryos and early L1 larvae, indicating that MYRF-1 alone is sufficient to activate \textit{lin-4} transcription (Figure 5B). Given that \textit{lin-4} transcription is suppressed by FLYWCH during embryogenesis and by distinct yet unidentified mechanisms during early L1, MYRF-1 may play a predominant role in promoting \textit{lin-4} transcription.
In our previous study, we characterized a mutant of *myrf-1^700*(syb1313) that exhibited precocious synaptic remodeling and M-cell division during mid-late L1, albeit in a discordant manner\[34\]. Our analysis of MYRF-1^700 localization led us to infer that MYRF-1(syb1313) functions as a constitutively active protein in specific cell types, but weakly and inconsistently. In the current study, we investigated the expression of the *lin-4* transcription reporter in *myrf-1*(syb1313) and observed a clear upregulation of the reporter in the mutants, especially in neurons, at 6 hours, while it was undetectable in wild-type animals at this stage (Figure 5C, D). It is worth noting that at the time (15h) when *lin-4* is typically upregulated in wild-type animals, the reporter GFP expression is significantly absent in the pharynx and epidermis of *myrf-1*(syb1313) mutants. However, there is sustained, higher-than-wild type expression of the reporter GFP in neurons of these mutants. Considering the precocious phenotype observed in DD neurons in *myrf-1*(syb1313), these data support that hyperactive MYRF-1 promotes the premature transcription of *lin-4*.

We conducted further analysis to investigate the impact of *myrf-1^700*(syb1313) on the endogenously tagged mScarlet *lin-4* transcription reporter. In the mutants, we observed premature upregulation of mScarlet signals in a subset of nuclei, likely neurons based on their position and nucleus size, at 6 hours (Figure 5E, F). As mutant animals progressed towards late L1 (14h), the signal intensity significantly increased and remained consistently higher compared to the wild type controls. From this stage on, while the mScarlet signals in many wild type individuals were still in the process of upregulation, the signal intensity in *myrf-1^700*(syb1313) mutants did not exhibit a comparable sustained increase. This observation aligns with our assessment of *myrf-1^700*(syb1313) mutants, indicating an inconsistent precocity and developmental progression deficiency. Nonetheless, these results strongly support the notion that hyperactive MYRF can precociously activate endogenous *lin-4* transcription.
It is worth noting that in \textit{myrf}-1\textsuperscript{v700}(\textit{syb1313}) mutants, the mScarlet signals exhibit strong upregulation in a few nuclei located in the posterior bulb of the pharynx at 14 hours, in comparison to the wild type (Figure 5E, F). These cells appear to belong to the same subset, as determined by their position and pattern, where the mScarlet signals have not experienced a significant decrease as observed in many other neighboring cells when MYRF function is lost (Figure 2C, D). This result indicates that hyperactive MYRF indeed enhances \textit{lin}-4 transcription in this specific subset of pharyngeal cells. However, it is noticeable that at 6 hours, the mScarlet signals in these nuclei of \textit{myrf}-1\textsuperscript{v700}(\textit{syb1313}) mutants are not as bright as nearby neurons, suggesting that MYRF-1\textsuperscript{v700} does not activate \textit{lin}-4 expression effectively in these pharyngeal nuclei at early L1.

The strong induction of mScarlet signals in these pharyngeal nuclei at late L1 and onwards in \textit{myrf}-1\textsuperscript{v700}(\textit{syb1313}) mutants contrasts with the lack of \textit{maIs134} signals in the pharynx of the mutants. One possible explanation for this contrasting expression pattern is the presence of additional pharynx-enhancing elements located outside the 2.4 kb promoter region of \textit{lin}-4 used in the \textit{maIs134} transgene. Alternatively, it should be considered that the threshold for transcription factor activation required to drive the expression of the endogenous reporter (mScarlet) versus the multicopy DNA array reporter (\textit{maIs134}) is likely different. Therefore, resolving this discrepancy would require further investigation.

\textbf{MYRF-1 interacts with \textit{lin}-4 promoter directly}

We next investigated whether MYRF-1 physically binds to \textit{lin}-4 regulatory sequences to regulate \textit{lin}-4 transcription. Gel mobility shift assays are commonly used to test the direct interaction between protein and DNA fragments. A series of MYRF-1 variants were expressed in HEK cells by transfection, and cell lysis was tested for their binding with 498 bp DNA of the \textit{lin}-4 promoter. None of the \textit{C. elegans} MYRF-1 variants including full-length and N-MYRF-1\textsuperscript{1-482} display significant binding with 498 bp \textit{lin}-4 promoter. This negative result is difficult to interpret.
considering when *C. elegans* MYRF-1 is expressed HEK cell lines, they do not assemble into trimer nor be cleaved (while mammalian MYRF does) for unknown reasons[33], which may render them incompetent to bind target DNA.

While analyzing *Plin-4-GFP(mals134)*, we observed that the multicopy transgene of *lin-4* promoter DNA caused unexpected developmental abnormalities in animals. Previously, we used combined alleles, *myrf-1(ju1121)* along with a rescuing multicopy transgene *myrf-1loxP(ybqIs112)* for the conditional ablation of *myrf-1*[33]. In our current study, we introduced the *Pdpj-7-Cre* transgene into the combined-conditional *myrf-1* alleles (*ju1121-ybqIs112*) to examine *mals134* expression in the epidermis where *myrf-1* is to be ablated. We observed a certain percentage of dauers emerging on plates even when the food *E. Coli* OP50 had not been exhausted (Figure 6A, B; Figure S3). Dauer refers to the alternative L3 stage that animals develop into when they encounter unfavorable living conditions such as low food abundance and high temperature. Because wild type *C. elegans* would never become dauer before the food runs out, this type of dauer formation is only associated with certain types of mutants, termed as the constitutive dauer formation (Daf-C). With intrigue, we initially thought that the dauer is due to the loss of *myrf-1* in the epidermis; however, the *mals134* transgene animals themselves produce a low percentage of Daf-C, although the combinatorial effect of the conditional *myrf-1* alleles (*ju1121-ybqIs112*) and epidermal-Cre remains to be characterized. We also noticed that animals carrying *mals134* invariably develop slower than wild-type animals, even though they show no obvious defect in becoming adults and in fertility (Figure 6B-F). The developmental delay starts from L2 and onwards based on their appearance and body length measurement. The L2 animals of *mals134* are typically darker, longer, fatter than normal L2 animals and resemble pre-dauer (L2d) animals in gross morphology (Figure 6F). We interpret the Daf-C phenotype in *mals134* with the reasoning that the transgene containing tandem array of *lin-4* promoter DNA sequestered a significant amount of the MYRF-1 protein, causing MYRF-1 to drop to such a low level that impedes normal development. A salient phenotype of *myrf-1* mutants is developmental arrest
after all. If the hypothesis was true, expressing more MYRF-1 in mals134 animals might suppress the Daf-C phenotype. In the rescuing transgene myrf-1LoxP(ybqls112) MYRF-1 protein is expected to increase slightly more than its endogenous level. This notion is indeed consistent with the observation that ybqls112 transgene completely suppressed the slow-growth phenotype in mals134, as well as the Daf-C (Figure 6A-D), suggesting that the transgene of lin-4 promoter DNA negatively interferes with the normal function of MYRF-1 in development.

These observations prompted us to investigate further whether the transgene array of lin-4 promoter DNA effectively binds a significant amount of MYRF-1 protein. To test this, we injected the lin-4 promoter (498 bp or 2.4 kb) reporter plasmid into myrf-1GFP(ybq14), and remarkably, we detected some puncta of intensified GFP signals in a subset of nuclei in animals carrying transgene (Figure 6G-I; Figure S4). This suggests that the MYRF-1GFP protein is condensed in discrete nuclear locations, a phenomenon that we never observed in wild type myrf-1GFP(ybq14) animals. To determine the specificity of their interaction, we designed a plasmid of lin-4 promoter DNA without the sequence encoding RFP or GFP, and a second plasmid containing 7x TetO sequence to be bound by TetR. We generated a line of ubiquitously expressed TetR DNA binding domain fused with RFP (TetR::RFP) as a single copy transgene (ybqS1233) to ensure consistent expression of TetR::RFP at a moderate level (Figure 6G, H). The formation of the TetO tandem DNA array indeed led to the formation of TetR::RFP puncta, demonstrating the effectiveness of the method. The presence of RFP puncta can then mark the location of DNA arrays in the nucleus. The formation of the TetO DNA array alone was insufficient to sequester MYRF-1GFP, while only the addition of the lin-4 promoter DNA into the tandem array caused the emergence of the MYRF-1GFP puncta, indicating the specificity of the lin-4 promoter-MYRF-1 interaction. Therefore, the collection of serendipitous observations and the nuclear loci co-labelling in vivo supports the direct regulator role of MYRF-1 in driving lin-4 transcription.
**MYRF-1 regulates a selective subset of microRNAs during L1-L2 transition**

We wanted to investigate the extent to which MYRF-1’s transcriptional activity might affect the landscape of microRNA expression in genome, as microRNAs often have stronger effects when present in combination with homologous microRNAs or in synergy with other microRNA families[22]. We performed microRNA-targeted sequencing analysis on *myrf-1(ju1121)* mutants of late L1 (16 hours) compared to controls. The sequencing analysis showed that a small subset of microRNA species was differentially expressed between *myrf-1(ju1121)* and control animals (P<0.05) (Figure 7A). We performed a phylogenetic analysis to analyze the relationships between these microRNA species (Figure 7B). Remarkably, 6 out of the 7 up-regulated microRNAs are clustered on one phylogenetic branch, which is separate from the other two. Among the differentially expressed, *lin-4* was the most decreased gene, consistent with our present analysis on *lin-4*. We then selected several candidates of relative abundance and examined their expression using transcriptional reporters. We generated single copy insertion transgenes carrying 2 kb upstream sequences of the candidate microRNA and eventually obtained lines for *mir-48, mir-73, and mir-230* showing consistent GFP signals (Figure 7C, D). The *mir-48* reporter (*ybqSi206*) was decreased, while *mir-73* (*ybqSi208*) and *mir-230* (*ybqSi209*) were increased in *myrf-1(ju1121)* at early L2, thus confirming the microRNA sequencing results. Notably, all three microRNAs were increased from L1 to L2 by transcriptional reporter; among the three, *mir-48* reporter is hardly detected in L1 and shows the most dramatic upregulation when animals transition to L2. The loss of *myrf-1* significantly decreased *mir-48*’s upregulation, but did not completely block it as it did with *lin-4*. *mir-48* belongs to the *let-7* microRNA family, and two other family members, *mir-84* and *mir-241*, were also downregulated in *myrf-1(ju1121)*, while *let-7*, the founding member, itself was upregulated in the mutants. Among the microRNAs upregulated in the *myrf-1* mutants, *mir-42* belongs to the *mir-35* family, which is essential for embryogenesis, and *mir-71* has been extensively studied for its upregulation in adults and regulatory role in aging. At this stage, it is unclear how these candidate microRNAs may be involved in regulating post-
embryonic development, but our results suggest an intricate genetic circuit in which MYRF-1’s activity may either enhance or dampen the expression of select microRNA species.

Discussion

Our study reveals that the nuclear accumulation of released N-terminal MYRF-1 increases from mid L1 to late L1, coinciding with the upregulation of lin-4 expression. We demonstrated that MYRF-1 is essential for the upregulation of lin-4 in terminal tissues in L1 and also required for lin-4 expression throughout the larval stages (Figure 7E). Our data support the direct, predominant role of MYRF-1 in driving lin-4 transcription.

While lin-4 expression appears to be constitutive after late L1 by stable GFP reporter, the analysis with higher temporal resolution showed that it exhibits oscillation in each larval stage[42, 43]. The periodic activation of lin-4 transcription is likely driven by poorly understood oscillation gene regulatory circuits, some of which include C. elegans orthologs of human circadian clock genes[44-49]. Recent work has begun to reveal the regulation mechanisms underlying oscillating lin-4 transcription[43, 49, 50]. BLMP-1/PRDM1 functions as a nutrient-stimulated priming factor to decompact the lin-4 locus and boost the amplitude of lin-4 oscillation[43]. It is worth noting that the expression of BLMP-1 itself does not oscillate. However, two other oscillating nuclear hormone receptors, NHR-85/Rev-Erb and NHR-23/ROR, promote lin-4 transcription, ensuring its precise oscillation timing[50]. These two receptors are the closest nematode orthologs of human circadian transcription factors Rev-Erb and ROR, respectively. The expression of NHR-85 is downregulated post-transcriptionally by LIN-42/PERIOD, another ortholog of the human circadian factor[50]. LIN-42/PERIOD peaks in each cycle, but lags behind NHR-85 and NHR-23. Although the interactions between BLMP-1, NHR-85, and NHR-23 constitute an attractive model for explaining the oscillation of lin-4 expression, it remains unclear to what extent the upregulation of lin-4 expression, particularly during late L1, relies on these factors. For instance, there is a lack of
description regarding the impact of BLMP-1, NHR-85, and NHR-23 mutants on endogenous *lin-4* expression. Exploring how MYRF-1 interacts with the three components of *lin-4* oscillation control—BLMP-1, the NHR-85 and NHR-23 pair, and LIN-42—would be an interesting direction for further study.

Similar to the orthologs of circadian genes, *myrf-1* mRNA exhibits oscillation in each larval stage as part of an oscillation scheme consisting of over 3000 genes oscillating at specific phases in each cycle[44, 45]. These gene expression patterns, including *lin-4*, are halted during acute food removal/starvation, indicating that cycle progression is regulated by nutritional state[43, 51]. Our report includes the partial dauer-constitutive phenotype caused by the interaction between the *lin-4* promoter DNA and MYRF-1. Although we do not fully understand the phenomenon, it clearly points to a link between MYRF-1 activity and nutrient state. Investigating how MYRF-1 responds to nutrient signals, what transcriptional outcome it produces, and its position in the regulatory circuit of oscillation will be important topics for future investigation.

**Materials and Methods**

**Animals**

Wild-type *C. elegans* were Bristol N2 strain. Strains were cultured on NGM plates using standard procedures (Brenner, 1974). Unless noted, animals were cultured at 20°C for assays requiring specific developmental stages. Animals analyzed in this paper were hermaphrodite

**Naming of the Alleles**

All alleles generated in Y.B.Q. lab are designated as "ybq" alleles, and all strains, as "BLW" strains. "Ex" denotes transgene alleles of exchromosomal array. "Is" denotes integrated transgene. "Si" denotes single-copy integrated transgene. "syb" alleles (in "PHX" strains) are generated by genomic editing using CRISPR-Cas9 technique. "syb" alleles were designed by Y.B.Q. and produced by SunyBiotech (Fuzhou, China).
myrf-1 alleles by CRISPR-Cas9 editing

For the following described alleles generated by CRISPR-Cas9 editing, Cas9 and gRNA were expressed from plasmids. The positive clones were identified using PCR screening to test singled F1 resulted from microinjection. GFP::myrf-1(LoxP) (ybq98) alleles was generated in the background of strain BLW889 [GFP::myrf-1::3xFlag(ybq14)]. Two artificial introns are inserted into the third exon of myrf-1 gene and each intron carries one LoxP site.

gRNA target: sgRNA1: TCAAGTCGGCTTCTCTTTACGTGG
sgRNA2: TACGTGGCATCTCCAAAAACAGGG

ybq14(background): …GGAATGCCAAGCCCTGTCCCCGGAGATGCCACGTACGTAAG - (insertion point)

AGAAGGCGACTTGACACCCCCGTGAAACGGCAAGAAATCGCTCCAAAGCTTTGCTGGTTATG
ACGGATTTCCAGATGAGAATTACAGTCAGCAACAGGCAATCAG - (insertion point) -
ATTCTCAAAGTTCTCAAGAAAGAAGTGAGTCCACTGTATGACATCAACGCTCAACCGCTAC
AACAACTTCAA...

ybq98: …GGAATGCCAAGCCCTGTCCCCGGAGATGCCACGTACGTAAG - (gtatgtttcgaatgatactaataATAACTTCGTATAGCATACATTATACGAAGTTATAcataacatacataacatatcag)

AGAAGGCGACTTGACACCCCCGTGAAACGGCAAGAAATCGCTCCAAAGCTTTGCTGGTTATG
ACGGATTTCCAGATGAGAATTACAGTCAGCAACAGGCAATCAG - (gtatgtttcgaatgatactaataATAACTTCGTATAGCATACATTATACGAAGTTATAcataacatacataacatatcag)

ATTCTCAAAGTTCTCAAGAAAGAAGTGAGTCCACTGTATGACATCAACGCTCAACCGCTAC
AACAACTTCAA...

GFP::Degron::myrf-1::3xFlag(ybq133) was generated in the background of BLW889 [GFP::myrf-1::3xFlag(ybq14)]. Degron is inserted after the last amino acid of GFP (Lys).

sgRNA: CAATCAACCTACAAACACCCTGG
ybq14(background): …GGGATTACACATGGCATGGATGAACTATACAAA - (insertion point) –
GCAGTCAATCAACCTACAAACACCTGGCTCAACTCAA…
ybq133: …GGGATTACATGGCATGGATGAACTATACAAA – DEGRON
GCAGTCAATCAACCTACAAACACCC

**Generation of transgene alleles**

Pmyo-3-MYRF-1 transgene: The vector pQA1094 [Pmyo-3-myrf-1] was injected to BLW1424 [myrf-1(ju1121)/mIn1 II; lin-4p::GFP+unc-119(mals134)] at concentration of 0.5 ng/µl. The resulting strain is BLW1579 [myrf-1(ju1121)/mIn1 II; lin-4p::GFP+unc-119(mals134); Pmyo-3-myrf-1(ybqEx746)]

Pdpy-7-MYRF-1 transgene: The vector pQA1511 [Pdpy-7-gfp::myrf-1] injected to N2 at concentration of 0.5 ng/µl to generate BLW1562 [Pdpy-7-gfp::myrf-1(ybqEx721)]. ybqEx721 was crossed into VT1072 [unc-119(ed3) III; lin-4p::GFP+unc-119(mals134)] to generate BLW1578 [lin-4p::GFP+unc-119(mals134); Pdpy-7-gfp::myrf-1(ybqEx721)]. BLW1578 was crossed with BLW1424 [myrf-1(ju1121)/mIn1 II; lin-4p::GFP+unc-119(mals134)] to generate BLW1580 [myrf-1(ju1121)/mIn1 II; lin-4p::GFP+unc-119(mals134); Pdpy-7-gfp::myrf-1(ybqEx721)].

Overexpress GFP::MYRF-1 (delete 601-650): The vector pQA1922[Prpl-28-GFP::myrf-1(delete 601-650)] were injected into CGC177[lin-4(umn84[lin-4p::SL1::EGL-13NLS::lox2272::mScarlet-l::cMycNLS::Lox511::let-858-3’UTR::lox2722])/mIn1[dpy-10(e128) umnIs33] at 10 ng/µl.

Tandem DNA array of Plin-4(2412 bp) DNA: The vector pQA1881 [Plin-4(2412bp) -unc-54-3’UTR] was injected into BLW1827 [gfp::myrf-1(ybq14); glo-1(zu391)] at 50 ng/µl.

Tandem DNA array of Plin-4(498bp) DNA: The vector pQA1880 [Plin-4(498bp) -unc-54-3’UTR] was injected into BLW1827 [gfp::myrf-1(ybq14); glo-1(zu391)] at 50 ng/µl.

Tandem DNA array of 7xTetO: The vector pQA1961 [7xTetO] was injected into BLW2170 [gfp::myrf-1(ybq14)]; glo-1(zu391); Prpl-28-TetR-DBD::TagRFP(ybqSi233)] at 50 ng/µl.

Tandem DNA array of Plin-4 DNA+7xTetO: The vector pQA1960 [Plin-4(2412bp) -unc-54-3’UTR -7xTetO] were injected into BLW2170 [gfp::myrf-1(ybq14)]; glo-1(zu391); Prpl-28-TetR-DBD::TagRFP(ybqSi233)] at 50 ng/µl.
Pmir-48-gfp single copy transgene: The pQA1861 [Pmir-48-GFP miniMos_vector] at 50 ng/µl and standard components of miniMos system (see below) was injected into N2 animals to make single copy transgene Pmir-48-GFP(ybqSi206).

Pmir-73-gfp single copy transgene: The pQA1863 [Pmir-73-GFP miniMos_vector] at 50 ng/µl and standard components of miniMos system (see below) was injected into N2 animals to make single copy transgene Pmir-73-GFP(ybqSi208).

Pmir-230-GFP single copy transgene: The pQA1864 [Pmir-230-GFP miniMos_vector] at 50 ng/µl and standard components of miniMos system (see below) was injected into N2 animals to make single copy transgene Pmir-230-GFP(ybqSi209).

TetR-TagRFP single copy transgene: The pQA1896 [Prpl-28-TetR-DBD::TagRFP miniMos_vector] at 50 ng/µl and standard components of miniMos system (see below) was injected into N2 animals to make single copy transgene Prpl-28-TetR::TagRFP (ybqSi233).

Single-copy integrated transgene allele

The procedure for generating single-copy integrated transgenes using miniMos technology in this study followed the protocol established by Frøkjær-Jensen et al. (2014). The injection mixture contained 10 ng/µl pGH8 [Prab-3::mCherry], 2.5 ng/µl pCFJ90 [Pmyo-2::mCherry], 50 ng/µl pCFJ601 [Peft-3-Mos1-transposase], and 50 ng/µl miniMos plasmid harboring the target gene. The mixture was injected into the gonads of N2 animals. After injection, the nematodes were transferred onto NGM medium and incubated at 25°C for approximately 48 hours. Subsequently, 500 µl of 25 mg/mL G418 solution was added to each plate to screen for nematodes carrying the target transgene. After 7-10 days, healthy nematodes without mCherry co-markers were selected from the medium where all the food was consumed. Candidate single-copy integrators were grown on G418-containing plates and analyzed for target protein expression. Homozygous nematodes carrying a single-copy transgene were identified by their 100% resistance to G418 toxicity and expression of the target protein.
**Taqman real time PCR analysis for miRNA lin-4**

Samples for Taqman Real-Time PCR Assays were prepared from wild type N2 and BLW252 [myrf-1(ju1121)/mln1] strains at three stages: Early L1 (0h), Late L1 (16h), and Early L2 (21h). The worms were screened for size and fluorescence marker of the desired transgene using Biosorter. myrf-1(ju1121) mutants lack the pharyngeal GFP, which serves as a transgene marker of balancer mln1, as a criterion in sorting process. More than 2,000 worms were collected for each sample after the Biosorter process.

RNA extraction was carried out using RNAiso Plus (TaKaRa, Dalian, China) following the manufacturer’s instructions with some modifications. Briefly, 1 ml of RNAiso Plus was added to each sample, which was homogenized using a pre-cooled grinder. The homogenization buffer was then transferred to a 1.5 ml microcentrifuge tube and allowed to stand at room temperature (15 – 30°C) for 5 – 10 min. The supernatant was discarded, and 200 µl of chloroform was added and shaken for 30 s until the mixture turned milky. The mixture was then centrifuged at 12,000g at 4°C for 15 min, and the supernatant was transferred to a new microcentrifuge tube. To the supernatant, 0.5 ml of isoamylol was added and left to stand for 10 min at room temperature, then centrifuged at 12,000g at 4°C for 60 min. After discarding the supernatant, the pellet was washed with 75% ethanol at 12,000g, 4°C for 60 min, and then dissolved in 20 µl of DEPC-treated water.

The concentration of miRNA was determined using Nanodrop, and its quality was assessed through electrophoresis. Each experiment was performed using at least three replicates.

The TAKARA RNA reverse transcription kit (6110A) was used to perform RNA reverse transcription following the manufacturer’s instructions. The concentration of the resulting product was measured using Nanodrop, and its quality was evaluated through electrophoresis.

The Taqman real-time PCR was conducted according to the Taqman small RNA assay protocol. Initially, the concentration of each sample was diluted to below 10 ng/µl, followed by reverse transcription PCR using the Taqman microRNA RT kit (4366596). Subsequently, the Taqman real-time PCR mixture was prepared using Taqman small RNA assay (20X) with lin-4a (21nt) (ID: 000258) and sn2323 (ID: 001760) as the internal control. Each sample was tested using at least
three technical replicates and three biological replicates. The Taqman real-time PCR was carried out using Applied Biosystems QuantStudio 7 Flex Real-Time PCR System (Thermofisher), and all the procedures were followed as per the instructions. To determine the relative changes of each sample in each assay, the lin-4 Ct of technical replicates was averaged and then subtracted by the average sn2323 Ct to obtain the $\triangle$CT value for each sample. Within each biological replicate and assay, the difference between the $\triangle$CT of each sample and the $\triangle$CT of wild type at Early L1 was calculated to obtain $\triangle\triangle$CT. The 2 to the power of $\triangle\triangle$CT was then calculated to obtain the fold change in each sample compared to the wild type at Early L1. The fold changes were plotted in Figure 1.

**miRNA sequencing and bioinformatics analysis**

BLW53 [Phlh-8-GFP(ayIs6)] and BLW1555 [myrf-1(ju1121)/ mln1; Phlh-8-GFP(ayIs6)] strains were utilized for sample preparation. To isolate myrf-1 mutants, late L1 stage worms were selected based on size and lack of pharyngeal GFP (a balancer mln1 marker) using Biosorter. Over 10,000 worms were collected for each sample for further experimentation. Total RNA was extracted and assessed for quality and quantity using Nanodrop and Agilent 2100, respectively. Samples with a total quantity of 3-5 µg and high quality (RIN > 10) were selected for miRNA library construction. The miRNA libraries were established and sequenced by Novogene Technology Co., Ltd. (Beijing, China).

In this study, the C. elegans Assembly WBcel235/ce11 was used as the reference genome for bioinformatic analysis of the sequencing data. The reference genome and gene annotation database were obtained from the EMSEMBL database and used for downstream analysis. The sequencing data was initially assessed for quality using the software FASTQ, followed by alignment of each miRNA sequencing dataset independently to the reference genome using HiSAT2. Transcriptome from each RNA-sequencing dataset was then extracted using the StringTie software. Differential miRNA cluster analysis was performed to compare miRNA expression of control and myrf-1(ju1121) mutant using DESeq2. Data normalization was done using the TPM method, and the profiling differences between control and myrf-1(ju1121) mutant
were compared using "Fold Change" (FC). A P-value was assigned to each gene and adjusted using the Benjamini and Hochberg method. Genes with P_adj < 0.05 were considered differentially expressed.

**Microscopic analyses and quantification**

Live animals were anesthetized using 0.5% 1-Phenoxy-2-propanol in ddH₂O and mounted on 3% Agar gel pad. The animals were examined under x20, x40, or x60 oil objective on OLYMPUS BX63 microscope. The wide-field DIC or fluorescence images (single plane images or Z-focal stacks) were taken by a sCMOS camera (PrimΣ Photometrics camera (model 2) mounted on OLYMPUS BX63, which is driven by CellSens Dimension software (Olympus). Images of live animals were also acquired on Zeiss LSM880 with Airyscan. The thickness of the optical slices was typically 0.8 µm.

To quantify the patterns of GFP::MYRF-1, we acquired images of stage-synchronized animals using wide-field microscopy as described above. The same parameters were used, including the power of excitation light, an identical objective, exposure duration, and screen display setting. The acquired images were examined, and the patterns of GFP::MYRF were categorized based on the consistency of the signals observed at specific subcellular locations throughout the animal body. A "weak or unclear signal" means that either no clear signals are detected or there are some weak signals that are inconsistent throughout the animal body. We carried out three independent rounds of culture, imaging, and scoring, and the data were pooled and presented in percentage column graphs.

To quantify the general Plin-4-GFP(mals134) transcriptional reporter fluorescence intensity, we opened each acquired image stack in ImageJ and selected a single slice with a focused pharynx for further analysis. A square with sides 20 pixels (3.25 µm) long was drawn to the pharynx isthmus. The intensity of the region of interest (ROI) was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

To quantify the pharyngeal Plin-4-GFP(mals134) transcriptional reporter fluorescence intensity in myrf-1(syb1491) and myrf-1(syb1468) mutants, we opened each acquired image stack in ImageJ
and selected a single slice with a focused pharynx bulb for further analysis. The posterior pharynx bulb was selected as the ROI. The intensity of the ROI was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

To quantify the neuronal Plin-4-GFP(mals134) transcriptional reporter fluorescence intensity in myrf-1(syb1491) and myrf-1(syb1468) mutants, we opened each acquired image stack in ImageJ and selected a single slice with a focused head nerve ring for further analysis. The dorsal nerve ring area was selected as the ROI. The intensity of the ROI was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

To quantify the whole body lin-4p::nls::mScarlet(umn84) transcriptional reporter fluorescence intensity, the z-stack was maximally projected to produce a single image in ImageJ. The whole animal area was selected as ROI. The intensity of the ROI was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

To quantify the pharyngeal lin-4p::nls::mScarlet(umn84) transcriptional reporter fluorescence intensity in myrf-1(ju1121) and myrf-1(syb1313) mutants, we opened each acquired image stack in ImageJ and selected a single slice with a focused pharynx bulb for further analysis. Two posterior pharynx bulb nuclei were selected as the ROI. The intensity of the ROI was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

To quantify the neuronal lin-4p::nls::mScarlet(umn84) transcriptional reporter fluorescence intensity in myrf-1(ju1121) and myrf-1(syb1313) mutants, we opened each acquired image stack in ImageJ and selected a single slice with a focused nerve ring for further analysis. Several neuron nuclei were selected as the ROI. The intensity of the ROI was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

To quantify the LIN-14::GFP fluorescence intensity, we opened each acquired image stack in ImageJ and selected a single slice with a focused head neurons for further analysis. The head region was selected as the ROI. The intensity of the ROI was measured and presented as Mean ± SEM (t-test, *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).
To generate the line plot of TetR::TagRFP and GFP::MYRF-1, we opened each acquired image stack from Zeiss Airyscan in ImageJ and selected a single slice with a focused TetR::TagRFP spot for further analysis. We drew a 2-pixel (~0.1 μm)-thick, 12 μm-long ROI line to cross the center of one bright TetR::TagRFP spot. The middle point of the ROI line was positioned at the center of the spot. We ran the "plot profile" program located in the Analyze menu and recorded plot data in the RFP and GFP channels, respectively. The final graph was generated in GraphPad Prism 8. Each plot line represented the intensity distribution across a single TetR::TagRFP spot from an independent cell.

To generate a multiple line plot of GFP::MYRF-1, each image stack acquired from Zeiss Airyscan was opened in ImageJ, and a single slice with a focused GFP::MYRF-1 spot was selected. When no obviously bright GFP::MYRF-1 spot was found in the set of image slices, we chose a slice with clearly focused GFP::MYRF-1 signals in nucleus. A 2-pixel (~0.1μm)-thick, 10μm-long ROI line was drawn to cross the center of the GFP::MYRF-1 punctum (or brightest nucleus signal as we could fine). We then ran the program "plot profile", located in the Analyze menu, to record the plot data in the GFP channel. The final graph was generated in GraphPad Prism 8. The maximum intensity position in the ROI line was defined as the "0" position in the X axis. Measurement points within two microns centered at the "0" position were graphed. Each plot line represented the intensity distribution across a single GFP::MYRF-1 spot from an independent cell.

To generate a line plot of epidermal Plin-4-gfp in epidermal knock-out MYRF-1 animals, each acquired image stack was opened in ImageJ and a single slice with a focused seam cell was selected for further analysis. A 2-pixel (~0.1μm) thick, 12μm-long ROI line was drawn to cross a seam cell in a ventral-to-dorsal direction. The middle point of the ROI line was positioned at the center of the seam cell. The data was recorded by the "plot profile" program located in the Analyze menu. The final graph was generated using GraphPad Prism 8. Each plot line represented the intensity distribution across a single seam cell from an independent animal.

For quantification of miRNA transcriptional reporter fluorescence intensity, each image stack was opened in ImageJ and a single slice with focused head was selected for further analysis. A circle
ROI with a diameter of 10 pixels (1.625 μm) was drawn in the head region. For the mir-48 and mir-73 reporter, a circle was drawn on the pharynx bulb. For the mir-230 reporter, a circle was drawn on the first seam cell in the head. The intensity of the ROI was measured and presented as Mean ± SEM (t test. *, p<0.05; **, p<0.01; ***, p<0.001; ****, p<0.0001).

**Assay of dauer formation with 1% SDS**

To assay dauer formation, approximately 30 gravid adults (aged ~60 hours) were transferred to a fresh 3.5cm NGM plate where they laid eggs for four hours. Afterward, all adults were removed, and the synchronized eggs were maintained at 25°C for 50 hours. To analyze the ratio of dauer formation, animals were washed from the NGM plate and treated with 1% SDS for 20 minutes. Surviving animals were counted and considered as dauer, while dead animals were counted and considered as non-dauer.

**Assay of L2d formation by imaging**

To assay L2d formation, approximately 30 gravid adults (aged ~60 hours) were transferred to a fresh 3.5cm NGM plate where they laid eggs for four hours. Afterward, all adults were removed, and the synchronized eggs were maintained at 20°C for 48 hours. To analyze the morphology of animals, DIC images of animals were acquired on an OLYMPUS BX63 microscope using a x10 objective. Animals exhibiting a leaner morphology and more condensed intestinal granules were counted and considered as L2d.

**Assay of dauer formation by imaging**

To assay dauer formation, synchronized fresh-hatched L1 were seeded at ~200 per 3.5cm NGM plate and cultured at 20°C for 70 hours. To analyze the morphology of animals, DIC images of animals were acquired on an OLYMPUS BX63 microscope using a x10 objective. Animals exhibiting a leaner and darker morphology were counted and considered as dauer.

**Quantification of animal length**

To quantify animal length and analyze their growth, DIC images of animals were acquired on an OLYMPUS BX63 microscope using a x10 objective. The lengths of animals in the images were
measured using the polyline tool in OLYMPUS imaging software, and the data were further analyzed in GraphPad Prism.

**Auxin treatment**

To perform auxin treatment, worms were transferred to OP50-seeded NGM plates containing ~4 mM K-NAA (1-Naphthaleneacetic acid potassium salt). Briefly, fresh OP50-seeded 3.5cm NGM plates with 3mL NGM agar and a 25mM K-NAA solution in ddH2O were prepared. Next, 500 μL of the 25mM K-NAA solution was added to the NGM plates and the lid was kept on until the solution dried. For all K-NAA treatment experiments, 500 μL of ddH2O was used as a control.

**Phylogenetic analysis of differentially expressed miRNA genes**

Phylogenetic analyses were performed using the MEGA 11.0 program. First, all miRNA sequences were imported into MEGA 11.0 and aligned using ClustalX to search for conserved bases and motifs. Pairwise distance was then estimated using the default parameters, except for the bootstrap replication number parameter, which was set to 1000, and the substitution model parameter, which was set to P-distance. The phylogenetic tree for our specific miRNAs was generated using the Neighbor-Joining method and default parameters, with branches corresponding to partitions reproduced in less than 0.1 bootstrap replicates filtered out.

**Acknowledgments**

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References


Figure 1

A

MYRF-1<sup>grp</sup>(ybr14)

P(1n-4-GFP(mals134)

<table>
<thead>
<tr>
<th>Time</th>
<th>MYRF-1&lt;sup&gt;grp&lt;/sup&gt;</th>
<th>P(1n-4-GFP(mals134)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early-L1</td>
<td>Nucleus</td>
<td>Cell membrane</td>
</tr>
<tr>
<td>Late-L1</td>
<td>Nucleus</td>
<td>Cell membrane</td>
</tr>
<tr>
<td>Early-L2</td>
<td>Nucleus</td>
<td>Cell membrane</td>
</tr>
</tbody>
</table>

B

MYRF-1<sup>grp</sup> localization

- Nucleus
- Cell membrane
- Nucleus and Cell membrane

C

P(1n-4-GFP intensity

- Nucleus
- Cell membrane
- Cell membrane and Nucleus
Figure 1. The nuclear accumulation of N-MYRF-1 coincides with the upregulation of lin-4 in developmental timing.

A. Nuclear localization of GFP::MYRF-1 is increased in late L1, coinciding with the upregulation of Plin-4-GFP. GFP, endogenously inserted at MYRF-1 Ala171, labels both full-length MYRF-1 and post-cleaved N-MYRF-1. Plin-4-GFP(mals134) is a transcriptional reporter of lin-4, carrying a 2.4 kb sequence upstream of the lin-4 gene that drives GFP. While GFP::MYRF-1 is initially localized at the cell membrane in early-mid L1 (6 post-hatch hours), it becomes enriched in the nucleus towards late L1 (15 post-hatch hours). Plin-4-GFP is barely detected in early L1 but is upregulated in late L1. The insert shows a zoomed-in view of the framed area, covering part of the pharynx.

B. Quantification of animals showing a particular pattern of GFP::MYRF-1 (as shown in (A)) at various stages. The number of animals analyzed is indicated on each bar.

C. The fluorescence intensity of the lin-4 transcriptional reporter (as shown in (A)) was quantified and presented as mean ± SEM (t-test, ****p < 0.0001); n > 20, where n is the number of nematodes analyzed.
Figure 2

(A) 

Plin-4-GFP(mals134)

<table>
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<tr>
<th></th>
<th>wt</th>
<th>myrf-1(u1121)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early L1</td>
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<tr>
<td>(0h)</td>
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<tr>
<td>Late L1</td>
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<tr>
<td>(16h)</td>
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<tr>
<td>Early L2</td>
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<tr>
<td>(21h)</td>
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</table>

(B) 

Plin-4-gfp intensity

(C) 

Lin-4p:nls:mScarlet(mnr84)[m984]

<table>
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<tr>
<th></th>
<th>wt</th>
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(D) 

Lin-4p:nls:mScarlet intensity

(E) 

lin-4 microRNA quantity

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<td>Early L2</td>
<td></td>
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<tr>
<td>(21h)</td>
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Fold change (in wt vs myrf-1(u1121))

lin-4 microRNA quantity

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<tbody>
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<td>Early L1</td>
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<td></td>
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<tr>
<td>(0h)</td>
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<tr>
<td>(16h)</td>
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<tr>
<td>Early L2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>(21h)</td>
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</table>

Fluorescence intensity

in pharyngeal cells

in head neurons

whole body
Figure 2. MYRF is required for lin-4 upregulation during late L1

A. Plin-4-GFP is not upregulated in myrf-1(ju1121). The expression of the lin-4 transcriptional reporter in wild-type and myrf-1(ju1121) animals was examined at the early L1 (0h), late L1 (16h), and early L2 stages (21h).

B. The fluorescence intensity of the lin-4 transcriptional reporter (as shown in (A)) was quantified and presented as mean ± SEM (t-test, ****p < 0.0001). More than 20 animals were analyzed.

C. The endogenous lin-4 transcriptional reporter lin-4p::nls::mScarlet(umn84) is not upregulated in myrf-1(ju1121) except for a few pharyngeal nuclei. The expression of the lin-4p::nls::mScarlet in wild-type and myrf-1(ju1121) animals was examined at the early-mid L1 (6h), late L1 (14h), and early L2 stages (21h).

D. The fluorescence intensity of lin-4p::nls::mScarlet(umn84) (as shown in (C)) was quantified and presented as mean ± SEM (t-test, ****p < 0.0001). The "whole body" refers to the region of interest (ROI), which encompasses the entire body of the animal. The ROI of "in head neurons" comprises multiple head neuron nuclei found in each animal. For "in pharyngeal cells", the ROI includes the 8-9 pharyngeal nuclei that exhibit bright mScarlet signals. More than 15 animals were analyzed.

E. The abundance of mature lin-4 miRNAs in wild-type and myrf-1(ju1121) animals was examined at the early L1 (0h), late L1 (16h), and early L2 stages (21h) using qPCR analysis with probes specifically detecting lin-4 microRNA. Each data point represents relative fold change of each sample comparing to the wild type Early L1 sample within one set of experiment. The data represent three biological replicates. Statistics used t-test. *p < 0.05, ***p < 0.001.
Figure 3

A

LIN-14^{GFPI(cc2841)}

wt  

myrf-1(ju1121)  

Early L1  
(0h)

Late L1  
(16h)

Early L2  
(21h)

B

Phenotypic Intensity

LIN-14^{GFPI} intensity

Early L1  Late L1  Early L2

Control  

myrf-1  
(ju1121)
Figure 3. There is a sustained high level of LIN-14 protein in myrf-1 mutants.

A. Expression of LIN-14::GFP(cc2841) in wild-type and myrf-1(ju1121) at the early L1 (0h), late L1 (16h), and early L2 (21h) stages. GFP was endogenously tagged at the LIN-14 C-terminus. LIN-14::GFP is bright in early L1 and downregulated in late L1. LIN-14::GFP is not affected by myrf-1(ju1121) at early L1 but significantly brighter than wild-type control at late L1 and L2.

B: The fluorescence intensity for LIN-14::GFP (as shown in (G)) was measured and presented as mean ± SEM (t-test, ns: not significant, p > 0.05; ***: p < 0.001) from the analysis of more than 20 nematodes. Each data point represents the mean intensity of the head region in an individual animal. The head region was selected due to its low autofluorescence background.
Figure 4

A

\(\text{wt}\)

\(\text{myr-1}(\text{ju1121})\)

\(\text{Pmyo-3-myrf-1}\)

\(\text{Pdpy-7-myrf-1}\)

\(\text{Plin-4-GFP(mals134)}\)

B

\(\text{myr-1}(\text{ybq98})\)

\(\text{Control}\)

\(\text{Pdpy-7-Cre(mls1028)}\)

\(\text{Plin-4-GFP(mals134)}\)

C

Fluorescence intensity vs. Distance across one seam cell (μm)
Figure 4. MYRF-1 is sufficient to drive lin-4 expression in a cell-autonomous manner.

A. Genetic rescue of MYRF-1 in myrf-1(ju1121) using tissue-specific promoters. Plin-4-GFP(mals134) signals are observed specifically in body wall muscles and epidermis (asterisk) of myrf-1(ju1121) carrying transgene Pmyo-3-myrf-1 and Pdpy-7-myrf-1, respectively, while no detectable Plin-4-GFP is observed in L2 of myrf-1(ju1121).

B. Tissue-specific ablation of myrf-1 in the epidermis. myrf-1loxP(ybq98) combined with Pdpy-7-NLS::Cre(tmls1028) caused loss or drastic decrease of Plin-4-GFP(mals134) in the epidermis, while signals were detected in other tissues. Representative images of L2 (24h) animals are shown.

C. Fluorescence intensity measurements (as shown in (B)) are displayed for a ROI bar drawn transversely and centered at one seam cell. Individual lines represent signals from one animal.
Figure 5. Hyperactive MYRF-1 drives premature expression of lin-4.

A. The reporter of lin-4 transcription, labeled by endogenously inserted nls::mScarlet (umn84), which also produces a loss-of-function allele of lin-4. The fluorescence was not observed in embryos or early L1, but in late L1, confirming the previous reports.

B. Overexpression of a hyperactive MYRF-1 mutant, GFP::MYRF-1(delete 601-650) caused premature lin-4 transcription in embryos and early L1, labeled by lin-4p::nls::mScarlet.

C. The expression of Plin-4-GFP(mals134) in wild type and myrf-1(syb1313) mutants. At 6 hours, Plin-4-GFP expression is elevated in the neurons of myrf-1(syb1313) mutants but undetectable in wild type. By late L1 (15h), Plin-4-GFP is upregulated in multiple tissues in wild type. Although GFP expression is sustained in neurons (arrow) of the mutants, it is significantly weak or absent in the pharynx (asterisk) of the mutants.

D. The fluorescence intensity of the lin-4 transcriptional reporter (as displayed in (C)) was measured and presented as mean ± SEM (t-test, *p < 0.05, **p < 0.01, ***p < 0.001). Each data point represents the mean intensity of the head neurons or pharynx region in individual animal, which were imaged using confocal microscopy.

E. The expression of lin-4p::nls::mScarlet(umn84) in wild type and myrf-1(syb1313) mutants. At 6 hours, mScarlet expression is elevated in certain neurons of myrf-1(syb1313) mutants but undetectable in wild type. By late L1 (14h), mScarlet is upregulated in multiple tissues in both wild type and myrf-1(syb1313) mutants. The mutants exhibit stronger mScarlet signals than wild type.

F. The fluorescence intensity of lin-4p::nls::mScarlet (as displayed in (E)) was measured and presented as mean ± SEM (t-test, *p < 0.05, **p < 0.01, ***p < 0.001). The ROI for "whole body" encompasses the entire body of the animal. The ROI of "in head neurons" comprises multiple head neuron nuclei in each animal. The ROI for "in pharyngeal cells" includes the 8-9 pharyngeal nuclei that exhibit strong mScarlet signals. Each data point represents the mean intensity of the ROI in individual animal.
Figure 6. *lin-4* promoter DNA recruits MYRF-1 protein *in vivo.*

**A.** A subset of animals carrying the *maIs134* transgene constitutive dauer formation even in the absence of food depletion on culture plates. Dauer formation was assessed by treating the animals with 1% SDS for 20 minutes. MYRF-1 overexpression (*ybqIs112*) suppresses the constitutive dauer formation in *maIs134*. Animals were cultured at 20°C for 70 hours starting from freshly laid eggs. The number of animals analyzed is indicated on each bar.

**B.** Morphological assessment shows that a subpopulation of animals carrying the *maIs134* transgene become dauer larvae, which exhibit a lean body and darkened intestine (Figure S3). MYRF-1 overexpression (*ybqIs112*) suppresses the constitutive dauer formation in *maIs134*. Animals were cultured at 20°C for 70 hours starting from young L1. The number of animals analyzed is indicated on each bar.

**C.** The development of *maIs134* is delayed compared to wild-type animals. with the majority of *maIs134* animals exhibiting pre-dauer-like characteristics while most of the wild-type animals become L4. While most of the wild-type animals become L4, the majority of *maIs134* animals are pre-dauer-like. Animals were cultured at 20°C for 48 hours starting from freshly laid eggs.

**D.** Representative images of animals from experiments in C.

**E.** Measurements of body length of wild type and *maIs134* animals show a growth delay in *maIs134* starting from L2. The mean body length of analyzed animals at a series of time points is shown on the graph, with the mean ± S.D. indicated. The number of animals analyzed for each data point is more than 20.

**F.** Representative images of animals from experiments in E. At 26h *maIs134* animals are thinner than wild type and have dark intestinal granules, which are characteristic of pre-dauer (L2d).

**G.** A tandem DNA array containing *lin-4* promoter (2.4 kb) DNA causes puncta of GFP::MYRF-1(*ybq14*) in the nucleus. As a control, a 7xTetO sequence-containing DNA array causes the puncta formation of TetR::tagRFP(*ybqSi233*), while it doesn’t cause the aggregation of GFP::MYRF-1. Only the addition of *lin-4* promoter DNA causes the formation of GFP::MYRF-1 puncta.
H. Representative images of animal cells carrying transgenes described in G, but in a high magnification view.

I. Line plots of signal intensity measurements along the bar ROI drawn across one red punctum in images, examples of which were shown in H. The bar ROI is centered at the fluorescent spot and examples of bar ROI were shown in H. Each individual panel represents signals from one cell.
Figure 7

A

B

C

D

E
Figure 7. MYRF-1 regulates a selective subset of microRNAs during L1-L2 transition.

A. The hierarchical clustering diagram shows differential miRNA expression levels in wild-type and \textit{myrf-1(ju1121)} animals at L2 stage. The color scale represents the log2 of the fold change, with red indicating highly expressed miRNAs and blue indicating low-expressed miRNAs. The numbers indicate the mean TPM of the specific miRNA from three replicates. Significance was determined at P<0.05.

B. The phylogenetic analysis shows the relationship between differentially expressed miRNA genes in \textit{myrf-1(ju1121)}. The three branches are color-coded.

C. The expression of transcriptional reporters, \textit{Pmir-48-gfp(ybqSi206)}, \textit{Pmir-73-gfp(ybqSi208)}, and \textit{Pmir-230-gfp(ybqSi209)} in wild-type and \textit{myrf-1(ju1121)} animals at early-mid L1 (6h) and middle L2 (24h) is shown.

D. The fluorescence intensity of transcriptional reporters shown in (C) is quantified and presented as mean ± SEM (t-test: ns, not significant, *p<0.05, ****p<0.0001); n >10, where n is the number of animals analyzed.

E. An illustration for how MYRF is processed and promotes \textit{lin-4} expression during development. At early L1, MYRF is mainly localized to the cell membrane in a PAN-1 dependent manner. However, during late L1, N-MYRF is released from the membrane through the catalytic activity of the ICE (Intramolecular Chaperone of Endosialidase) domain following its trimerization. Subsequently, N-MYRF translocates to the nucleus to enhance the transcription of \textit{lin-4}.
Figure S1. *Plin-4-GFP* expression in *myrf-1* loss of function mutants.

**A.** *Plin-4-GFP*(maIs134) expression was examined in wild-type, *myrf-1*(syb1491), and *myrf-1*(syb1468) animals at the L2 stage (24h). *Plin-4-GFP* was not detected in *myrf-1* mutants.

**B.** The fluorescence intensity of *Plin-4-GFP* was quantified and presented as mean ± SEM (t-test; ****p<0.0001); n >30, where n is the number of animals analyzed.

**C.** An individual of *myrf-1*(ju1121) mutant, showing DIC image and *lin-4p::nls::mScarlet*(umn84) fluorescence. mScarlet is expressed in 8-9 pharyngeal nuclei.
Figure S2

A

myrf-1<sup>Dpy25</sup> (ybq133); Peft-3-TIR(wrdS23); Plin-4-GFP(mals134)

L2 12h

L2+mock  L3+mock  L4+mock  L2+NAA  L3+NAA  L4+NAA

B

Plin-4-GFP intensity

Fluorescence intensity

NAA  Mock  L2  L3  L4

Figure S2. MYRF-1 is required for lin-4 transcription in all larval stages.

A. Larval stages L2, L3, and L4 animals were treated with or without 4 mM K-NAA for 12 h. The lin-4 transcription reporter was down-regulated in the drug-treated animals compared to the pre-treated and mock-treated animals.

B. The fluorescence intensity of lin-4 transcription reporter was quantified and presented as mean ± SEM (t-test; ****p<0.0001); n >30, where n is the number of animals analyzed.
Figure S3. Constitutive dauer phenotype observed in a subpopulation of *Plin-4-GFP(mals134)* animals.

Representative images of animals from the experiments in Figure 4B, which were cultured at 20°C for 70 hours, starting from synchronized young L1 larvae. A subset of animals carrying the *mals134* transgene exhibited a constitutive dauer phenotype even when food was not depleted on the culture plate.
Figure S4

A

Transgene co-marker alone

Transgene co-marker + Plin-4 DNA (498 bp)

B

Transgene co-marker alone

Transgene co-marker + Plin-4 DNA (2412 bp)

C

Transgene co-marker alone

Transgene co-marker + Plin-4 DNA (2412 bp)

D

MYRF-1<sup>off</sup> intensity

Fluorescence intensity

Distance across one MYRF-1<sup>off</sup> spot (µm)
Figure S4. *lin-4* promoter DNA causes aggregation of GFP::MYRF-1(ybq14) in the nucleus.

**A.** Tandem DNA arrays containing *lin-4* promoter (498 bp) DNA caused aggregation of GFP::MYRF-1(ybq14) in the nucleus (arrow), while a control co-marker (*Pmyo-2-mCherry*) containing DNA array did not cause the aggregation of GFP::MYRF-1.

**B.** Tandem DNA arrays containing *lin-4* promoter (2.4 kb) DNA caused aggregation of GFP::MYRF-1(ybq14) in the nucleus (arrow), while a control co-marker (*Pmyo-2-mCherry*) containing DNA array did not cause the aggregation of GFP::MYRF-1.

**C.** Representative images of animal cells carrying transgenes described in B, but in a high magnification view.

**D.** Line plots of signal intensity measurement along the bar ROI drawn across the GFP spots as illustrated in representative images in C. The bar ROI is centered at one fluorescent punctum. Each individual line represents signals from one representative cell. Examples of bar ROI were drawn on the images in C.