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RESEARCH COMMUNICATION

LIN-39/Hox triggers cell division and represses EFF-1/fusogen-dependent vulval cell fusion

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General mechanisms by which Hox genes establish cell fates are known. However, a few Hox effectors mediating cell behaviors have been identified. Here we found the first effector of LIN-39/HoxD4/Dfd in *Caenorhabditis elegans*. In specific vulval precursor cells (VPCs), LIN-39 represses early and late expression of EFF-1, a membrane protein essential for cell fusion. Repression of *eff-1* is also achieved by the activity of CEH-20/Exd/Pbx, a known cofactor of Hox proteins. Unfused VPCs in *lin-39(-);eff-1(-)* double mutants fail to divide but migrate, executing vulval fates. Thus, *lin-39* is essential for inhibition of EFF-1-dependent cell fusion and stimulation of cell proliferation during vulva formation.

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Throughout animal evolution, members of the Hox family of homeodomain proteins have been responsible for establishing regional identities along the anterior-posterior axis. Much has been learned about Hox genes, in particular from their expression patterns, and about downstream genes that are targets of Hox transcriptional regulation. Although most of the downstream targets identified thus far encode for transcription factors or signaling molecules, only a very limited number of Hox target genes have been found to be effectors or genes encoding structural proteins that participate in the actual intra- and intercellular events during morphogenesis (Graba et al. 1997). A given Hox gene can function in different cell types and at multiple times during development, activating some targets and inhibiting others (Salser and Kenyon 1996; Graba et al. 1997). Part of this differential function depends on the presence or absence of proteins (e.g., EXD/PBX) that work as coactivators or corepressors along with the Hox genes (Mann and Afolter 1998). The Hox gene *lin-39*, the paralog of *deformed* (*Dfd*) from *Drosophila* and mammalian *HoxD4*, controls vulva formation as well as other developmental events in *Caenorhabditis elegans* (Clark et al. 1993;

Wang et al. 1993; Eisenmann et al. 1998; Grant et al. 2000; Liu and Fire 2000). In one case, *lin-39* was shown to act with *ceh-20*, an ortholog of the *extradenticle* (*exd*) cofactor, to regulate patterning of the postembryonic mesoderm in *C. elegans* (Liu and Fire 2000).

During vulva development, *lin-39* possibly acts at two developmental stages. At the first larval stage (L1), it is active in six of eleven ventral epidermal Pn.p cells, inhibiting them from fusing to the surrounding hypodermis (epidermis) like their sister cells (Ch'ng and Kenyon 1999). At the third larval stage (L3), *lin-39* permits three of these vulval precursor cells [VPCs; P(5–7).p], to escape the fusion fate (Malooof and Kenyon 1998; Gleason et al. 2002). Although expressed at low levels in the other three VPCs [P(3,4,8).p], *lin-39* is not sufficient for these cells to escape cell fusion (Clandinin et al. 1997; Malooof and Kenyon 1998). Only when its activity is elevated (e.g., through ectopic Wnt signaling) can these cells escape fusion and adopt vulval fates (Fig. 1; Gleason et al. 2002).

Due to its importance as a Hox/homeotic gene, and specifically as the most downstream gene in the signaling pathways involved in vulva formation, *lin-39* was intensively studied in the last decade. It was found that at the first larval stage, its expression is suppressed in the posterior body region by the chromatin regulatory factor *egl-27* (Ch'ng and Kenyon 1999) as well as by redundant activity of the *hairy* homolog *ref-1* and the Hox gene *mab-5* (Alper and Kenyon 2001). In males, the zinc-finger protein REF-2 also interacts with *lin-39* and other Hox genes to regulate cell fusion in Pn.p cells (Alper and Kenyon 2002). At L3, *lin-39* expression is elevated due to activity of the transcription factor *sem-4* (Grant et al. 2000) and members of the Wnt pathway (Eisenmann et al. 1998; Hoier et al. 2000). However, how *lin-39* regulates cell fusion and whether it has other roles in patterning vulva formation remain unknown.

Whereas in *C. elegans* and related nematodes, the non-vulval Pn.p cells fuse with the hypodermis (Louvvet-Vallee et al. 2002), the equivalent cells in *Pristionchus pacificus* (a nematode that has been separated from *C. elegans* for ~100 million years) undergo programmed cell death (Felix 1999; Shemer and Podbilewicz 2000). This apoptotic fate is inhibited in the VPCs by the activity of the *P. pacificus* *lin-39*, probably by repression of the apoptotic effector *ced-3*, the only candidate effector regulated by a *Dfd* paralog (Eizinger and Sommer 1997). These evolutionary differences between *C. elegans* and *P. pacificus* could be explained by differences in the regulatory sequences of the *lin-39* promoter (Grandien and Sommer 2001), or due to different effectors expressed in the VPCs in each species. To expand on the finding that *lin-39* inhibits cell fusion of vulval cells in *C. elegans*, we asked whether it does so by inhibiting a fusion effector. We recently isolated *eff-1*, a gene encoding type-I membrane proteins essential for all epithelial cell fusion events throughout embryonic and postembryonic development (Mohler et al. 2002). Here, we asked whether *eff-1* may be a cell fusion effector down-regulated by *lin-39* and other cofactors and whether *lin-39* acts only permissively to inhibit cell fusion of specific Pn.p cells or also to execute vulva formation through other effectors of cell division and cell migration.

[**Keywords:** *Caenorhabditis elegans*; cell fusion; vulva development; Hox]

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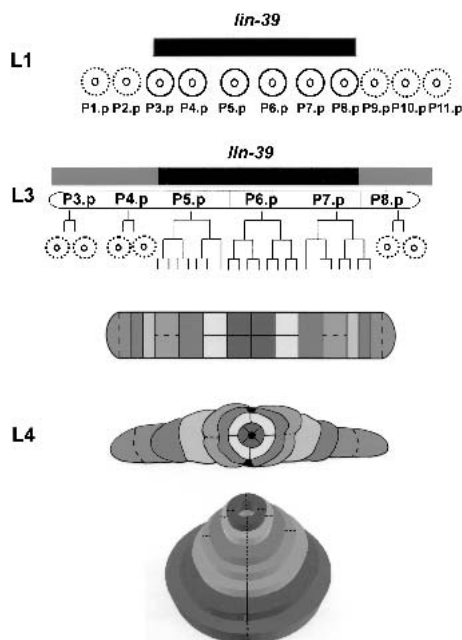


Figure 1. *lin-39* inhibits cell fusion during vulva formation. *lin-39* is expressed and active at the first larval stage (L1) in six epidermal cells (P3.p-P8.p; black circles), also known as the vulva precursor cells (VPCs). This activity (black box above cells) is necessary and sufficient to inhibit the VPCs from adopting the fate of their neighboring cells [P(1,2,9–11).p]—fusion to the surrounding syncytium hyp7 (dashed circles). At L3, the VPCs are subjected to various signaling pathways. P(3,4,8).p undergo one cycle of division, followed by fusion of their daughter cells to hyp7 (dashed circles). In about half of the cases, P3.p does not divide before cell fusion. P(5–7).p escape cell fusion and continue to divide, yielding a 22-cell vulval primordium. It was proposed that basal activity of LIN-39 in P(3,4,8).p is not sufficient for these cells to escape cell fusion, and only elevated activity of LIN-39 induced by the Ras pathway in P(5–7).p in wild-type (Malloof and Kenyon 1998) or by ectopic Wnt signaling in all VPCs in some mutants (Gleason et al. 2002), allows these cells to escape cell fusion (black and shaded boxes above cells). Migration, cell fusion (dashed lines), ring formation, and invagination of the primordial cells at L4 lead to a tube-shaped adult vulva (Sharma-Kishore et al. 1999).

Results and Discussion

eff-1 is epistatic to *lin-39*

In *C. elegans*, *lin-39* may repress VPCs from cell fusion fate in the first and third larval stages (L1 and L3), may promote vulval fates by keeping VPCs responsive to other signaling events in L3, or may combine both activities, inhibiting cell fusion and activating the response to inductive signals downstream of the RAS pathway (Clandinin et al. 1997; Malloof and Kenyon 1998; Gleason et al. 2002). In the *lin-39(n1760)* loss-of-function null mutant, all Pn.p cells have a cell fate transformation and fuse with the hypodermis at L1, developing no vulva (Clark et al. 1993; Wang et al. 1993). In contrast, in the *eff-1(hy21)* mutant, none of the Pn.p cells fuse at L1 or at L3, a protruded vulva is formed, and some animals have extra vulvae (Mohler et al. 2002).

To test the genetic epistatic relationship between *lin-39* and *eff-1*, we followed the fates of the different VPCs in wild-type, single, and double mutants during the L1 and L3 developmental stages. Several signaling pathways

determine distinct cell fates for descendants of the different VPCs (Sharma-Kishore et al. 1999). We found that in wild-type worms, only P(3–8).p cells escaped fusion in L1; P(5–7).p cells escaped fusion fate in L3, divided, and yielded the characteristic seven vulval rings at the L4 stage (Figs. 1, 2A,E). In *lin-39(n1760)* null mutants, all of the Pn.p cells fused to the surrounding hypodermis at mid-L1 (Fig. 2B,F). A similar pattern was seen in *lin-39(n1490ts)* conditional mutants (data not shown): at the semipermissive temperature of 20°C, occasionally one Pn.p cell escaped cell fusion at L1. This cell could either fuse at the L3 stage or it could stay unfused through L3. However, even in the latter case, this cell yielded an abnormal and nonfunctional “mini-vulva.” In *eff-1(hy21)* mutants, all of the VPCs [P(3–8).p] and the

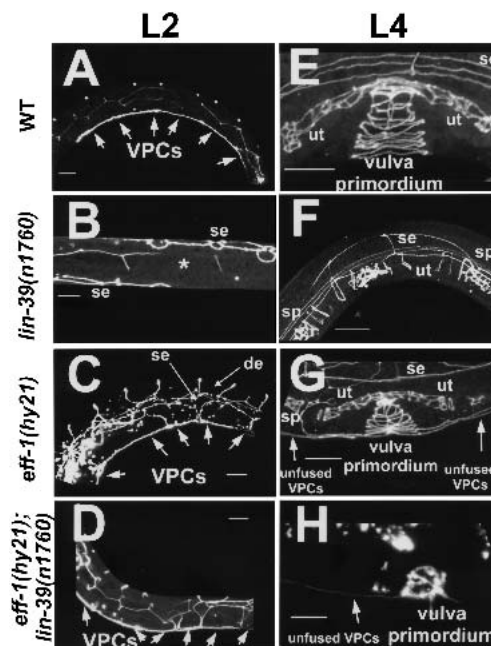


Figure 2. *eff-1* is epistatic to *lin-39* during early and late cell-fusion events. Confocal reconstructions of worms carrying an *ajm-1::GFP* construct marking the adherens junctions of epithelial cells (Shemer et al. 2000; Koepfen et al. 2001). The worms were assayed at early L2 (A–D) after the early fusion events of mid-L1 and at mid-L4 (E–H) after the late fusion events of mid-L3 (see Fig. 1 for details). (A) Wild-type worms showing the six VPCs that escape fusion to the hypodermis after they attach to each other ($n > 100$). (E) Three of these VPCs escape cell fusion to hyp7 at L3, and their 22 great-granddaughters invaginate, forming a stack of seven rings ($n > 100$). (B,F) In *lin-39(n1760)* single null mutants, no VPCs formed after all of the Pn.p cells had fused to the hypodermis (*, $n = 50$), resulting in the absence of a vulva [Vul phenotype] at late L4 (F; $n = 100$). (C) *eff-1(hy21)* mutants grown at the restrictive temperature of 25°C showing the VPCs that fail to fuse with hyp7 ($n = 50$). P3.p, P4.p, and P8.p also fail to fuse in the L3 and attach to the vulva primordium formed from the descendants of P(5–7).p. (G) The result is a stack of rings connected to a row of ectopic cells ($n = 70$). Due to fusion failure in *eff-1(hy21)*, ectopic dorsal epithelia (de) and lateral hypodermal seam cells (se) are present and migrate throughout the body of the worm. (D,H) Pn.p cells fail to fuse in *eff-1(hy21);lin-39(n1760)* double mutants at L1 (D; $n = 45$) and later at L3 (H; $n = 28$). The unfused VPCs migrate and invaginate, forming a “pseudo” vulval primordium that is incomplete and abnormal structurally, resulting in a nonfunctional vulva (cf. H and G,E). ut, uterus; sp, spermatheca. Anterior is to the left and dorsal is up, except for B, a ventral view. The fusion status of the cells was also confirmed by staining worms with the MH27 antibody (Podbilewicz and White 1994). Arrows mark unfused cells. Bar, 10 μ m.

P(1,2,9–11).p cells escaped fusion at L1 and failed to fuse again during the L3 stage, forming a vulva primordium of unfused vulval cells linked to unfused epidermal cells (Fig. 2C,G). If the double mutant had the phenotype of a single *eff-1*, with respect to the fusion of the Pn.p cells, it would mean that *lin-39* controls cell fusion in *C. elegans* by inhibiting the activity of *eff-1*. We found that in *lin-39(n1760);eff-1(hy21)* double mutants, all of the Pn.p cells failed to fuse both at L1 and at L3 despite the fact that *lin-39* inhibitory activity was absent (Fig. 2D,H). Similarly, the use of a *lin-39(n1490ts)* weaker allele in combination with *eff-1(hy21ts)* showed the same phenotype. Thus, during cell fusion events regulated by *lin-39* in vulva formation, *eff-1* is epistatic to *lin-39*. This is true for the VPCs at L1 where *lin-39* is active, and for P(5–7).p at L3 where there is sufficient *lin-39* activity to maintain these cells unfused. The similarity of the cell fusion phenotypes of *eff-1* single mutants and *lin-39;eff-1* double mutants suggests that *eff-1* acts downstream of *lin-39* at both early and late fusion events during vulva formation.

lin-39 represses *eff-1* expression

To rule out an alternative model in which *lin-39* inhibits cell fusion in the VPCs by repressing an unknown fusion effector that works nonredundantly with *eff-1* to promote cell fusion, we analyzed the expression of *eff-1* RNA in wild-type and *lin-39* mutant worms using a transcriptional *eff-1p::GFP* fusion construct (Mohler et al. 2002). Analysis of wild-type worms showed that *eff-1p::GFP* was expressed in the daughter cells of P3.p, P4.p, and P8.p that are committed to fusion, and was absent in the daughter cells of P(5–7).p which escape the fusion fate (Fig. 3A). In contrast, in *lin-39(n1760)* null mutants, which completely lack LIN-39 activity, all of the VPCs expressed *eff-1p::GFP* (Fig. 3B). Thus, the expression of *eff-1* RNA is regulated by LIN-39 activity, showing that *eff-1* works downstream of *lin-39* rather than in parallel. Consistent with these results, at the restrictive temperature of 25°C, *lin-39(n1490ts)* mutants expressed *eff-1p::GFP* in all of the VPCs. After shifting these conditional mutant animals to the permissive temperature, we found that GFP expression was completely dependent on LIN-39 activity. *eff-1p::GFP* was absent in P(5–7).p only in cases where remnants of *lin-39* activity were present, permitting P(5–7).p. to be further induced and to adopt vulval fates. In summary, our results show that *lin-39* inhibits epidermal fusion of the VPCs [P(3–8).p] at L1 and that of P(5–7).p at L3 by repressing *eff-1* expression (Fig. 4A).

ceh-20/exd represses *eff-1*-dependent fusion and vulva expression

A previous study showed that *lin-39* acts together with the *exd* homolog *ceh-20* in *C. elegans* to regulate the *twist* transcription factor ortholog *hlh-8* in postembryonic mesodermal cells (Liu and Fire 2000). To test whether *ceh-20* participates in regulation of cell fusion, we analyzed the fusion pattern of the VPCs in *ceh-20(ay42)* mutants. We found that all VPCs fused to hyp7 at mid-L3 in these mutant animals, except for one case in which one VPC escaped cell fusion. As a result, all of these mutants lacked a vulva (Fig. 3C). To test whether this fusion inhibition involves repression of *eff-1*, we

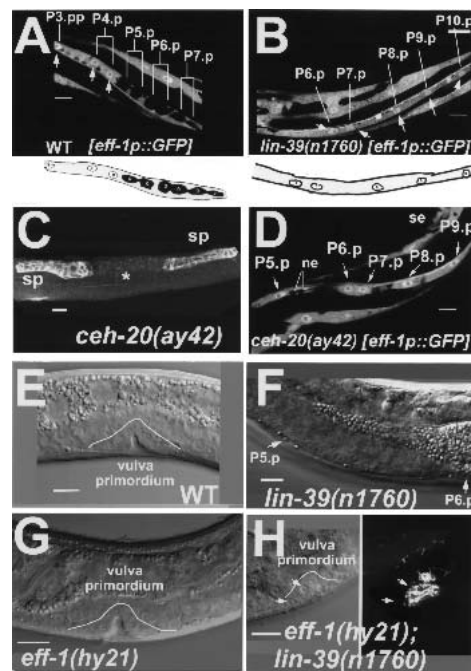


Figure 3. *lin-39* and *ceh-20* repress *eff-1* expression. (A,B) Z-serial sections from confocal reconstructions of L3 worms showing expression of *eff-1promoter::GFP* (white) in vulval primordial cells in the presence (A) and absence (B) of *lin-39* activity. (A) White arrows mark nuclei of cells that fuse with the surrounding hypodermal syncytium hyp7. All of these nuclei express *eff-1* (arrows and white circles in the diagram beneath; nucleoli are small circles within nuclei, $n = 35$). Daughter cells of P(5–7).p, which escape fusion to the hypodermis due to *lin-39* activity, do not express *eff-1*. (B) In the absence of *lin-39* activity, all of the VPCs express *eff-1* [P(6–10).p in this picture; see arrows and white nuclei in the diagram beneath, $n = 40$]. (C,D) *ceh-20* inhibits cell fusion by repressing *eff-1* expression. (C) *ceh-20* L4 mutant stained with MH27 showing that in the absence of *ceh-20*, no vulva (*) is formed as a result of VPC fusion ($n = 18$). The uterine cells are shown in the weak staining between the spermathecae (sp). (D) Expression of *eff-1promoter::GFP* (white) in vulval primordial cells in the absence of *ceh-20* activity. These cells express GFP [P(5–9).p in this picture]. Seam cells (se) and ventral neural cells (ne) are also shown ($n = 22$). (E–H) Nomarski micrographs showing vulval nuclei during organogenesis. (E) Wild-type worm at late L4. This vulva primordium is comprised of 22 nuclei (white line, invagination with 10 nuclei in this focal plane; $n > 100$). (F) *lin-39(n1760)* null mutant worm at early L4 showing only P5.p and P6.p that in the absence of *lin-39* activity fused to hyp7 ($n = 30$). (G) *eff-1(hy21)* worm at exactly the same stage as that in E (white line, invagination; $n = 60$). (H) In the absence of *lin-39*, the VPCs do not proliferate in *eff-1(hy21);lin-39(n1760)* mutants ($n = 34$). However, the cells that fail to fuse in the absence of *eff-1* activity, migrate and ultimately invaginate (white line). Right image shows a stack of four vulval rings (*ajm-1::GFP*) from the same vulva. Anterior is left and dorsal is up, except for A, C, and D, which are ventral views. All worms were grown at 25°C for complete penetrance of the *eff-1(hy21)* mutation. Bar, 10µm.

tested *eff-1p::GFP* expression in *ceh-20(ay42)* mutants. As in the case of *lin-39(-)*, in the absence of *ceh-20* activity, GFP expression was detected in all VPCs at L3 (Fig. 3D). Thus, *ceh-20* also regulates cell fusion by repressing *eff-1* expression in specific cells. Whether LIN-39 acts with CEH-20 as a protein complex and whether this regulation is direct will be tested in future experiments.

In addition to the vulval ectodermal cells, *eff-1* is essential for cell fusion in many epithelial and mesodermal

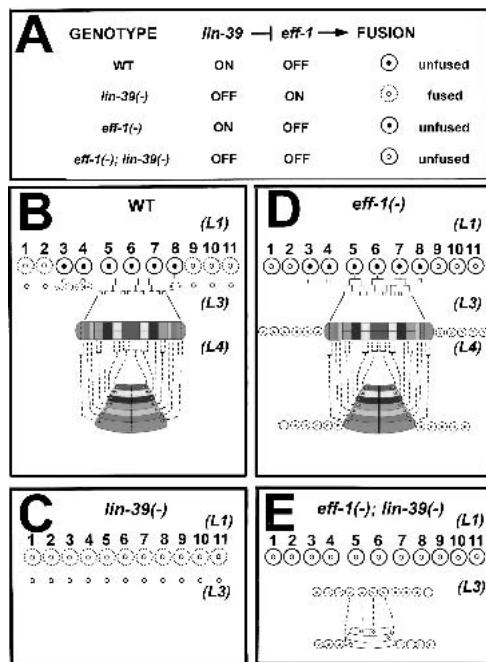


Figure 4. A model of *lin-39* activity during vulva formation. (A) *lin-39* inhibits cell fusion in the VPCs by repressing the fusogen *eff-1*. This model of interactions among *lin-39*, *eff-1*, and cell fusion predicts the fusion pattern of the VPCs in wild-type, single, and double mutants. In all diagrams, dashed lines represent fusing cells, and solid lines represent nonfusing cells. Cells that express LIN-39 have black nuclei. (B–E) Structural fates of the vulval cells in different genotypes. (B) In wild-type worms, *lin-39* acts in P(3–8).p at L1 and in P(5–7).p at L3, preventing these cells from fusion. The result is a 22-cell primordium that forms the seven vulval rings. (C) In *lin-39(-)* single mutants, *eff-1* is not repressed and at L1, all cells fuse and contribute their nuclei to the surrounding hypodermis. (D) In *eff-1(-)* single mutants, none of the cells are able to fuse with the hypodermis, resulting in ectopic cells that migrate along with the vulva precursors. (E) In *eff-1(-); lin-39(-)* double mutants, despite the lack of the inhibitory activity of *lin-39*, the Pn.p cells are not able to fuse at L1 and at L3 in the absence of *eff-1* activity. In the absence of *lin-39*, the cells fail to proliferate. They do succeed in forming rings (three in this example), but these rings are structurally abnormal and nonfunctional.

cells in which *lin-39* is not expressed or active (Shemer and Podbilewicz 2000). Other transcription factors that are known to regulate fusion in these cells might also work by regulating *eff-1* expression (Witze and Rothman 2002). Future work analyzing the relationships between *eff-1* and these regulators (e.g., the ELT-5/6 GATA transcription factors in the seam cells) should give a comprehensive view on how cell fusion is controlled throughout development in *C. elegans*.

lin-39 is required for proliferation of vulval cells

In wild-type worms the unfused VPCs divide, migrate, form rings, and invaginate to form the adult vulva (Fig. 4B; Sharma-Kishore et al. 1999). It has been proposed, using temperature-sensitive *lin-39* mutants and *heat-shock promoter::lin-39* transgenic worms, that the activity of *lin-39* is essential to keep the VPCs responsive to other inductive signaling events (Clandinin et al. 1997; Maloof and Kenyon 1998). Because cell fusion could not be bypassed in those experiments, the fusion of

the VPCs alone, in the absence of *lin-39* activity, could explain lack of vulval induction (Fig. 4C). The exact role of *lin-39* during late stages of vulva development is thus unclear; *lin-39* may have additional effectors in the VPCs and may be responsible for one or more of three processes: maintaining inhibition of cell fusion, promoting proliferation of P(5–7).p, and/or inducing formation of vulval rings. To prevent cell fusion of the VPCs to the hypodermis, we used *eff-1(hy21)* at 25°C at the L3 stage (Figs. 3G, 4D) and investigated the behavior of these cells and whether they were able to form vulval rings in *lin-39(-); eff-1(-)* and *lin-39(ts); eff-1(-)*. We found that although the VPCs escaped fusion in the double mutants, they failed to proliferate (Figs. 3H, 4E). These results show that (1) cells that fail to fuse because the cell fusion effector is inactive in *eff-1(-)* need additional cues to divide, and (2) *lin-39* may control a cell division effector in the VPCs that is required for them to divide. In the double mutant, the unfused VPCs may be blocked in proliferation because a putative cell division effector of *lin-39* is not receiving positive input from this Hox gene. In wild-type worms, remnants of *lin-39* from previous stages may be sufficient to induce one cycle of division in P(3,4,8).p before fusion to epidermis, whereas in the *lin-39(-); eff-1(-)* mutants, residual *lin-39* activity is absent, resulting in a block of cell division. Thus, in wild-type animals, *lin-39* may activate an unidentified effector of cell division in the VPCs, and *lin-39* complete loss-of-function causes cell cycle arrest. Alternatively, in the absence of *lin-39* activity, unfused VPCs may undergo a cell fate transformation from vulval precursor cells to differentiated vulval fates equivalent to the great-granddaughters of P(5–7).p cells.

Unfused stem cells (VPCs) differentiate without dividing in *lin-39(-); eff-1(-)* mutants

We next investigated the behavior of nonproliferative unfused cells. During the first hours of the L4 stage, the primordial cells normally form rings and invaginate. In *lin-39(-); eff-1(-)* animals, unfused and undivided VPCs did not show any morphological changes. Surprisingly however, at mid-L4 the VPCs succeeded to form one to four vulval rings or half-rings that stacked to form a pyramidal invagination (Figs. 2H, 3H, 4E). Thus, *lin-39* is not necessary to promote cell migration and ring formation during vulva development. These results imply that there are no cell migration effectors of *lin-39* in the process of vulva formation. To further examine the role of unfused and undivided VPCs in vulva morphogenesis, we examined the formation of vulval structures and found that the vulval rings were defective compared to wild-type and *eff-1* animals (Fig. 3, cf. H and E,G), and these vulvae were nonfunctional. Actually, these structures resembled the structures of pseudovulval rings seen in *let-60/ras(gf)* mutants that also result from a relatively small number of precursors (Shemer et al. 2000), showing that in *C. elegans*, a 6-cell vulval primordium is not sufficient to form a functional organ. In conclusion, our findings show that vulval/epidermal stem cells in *C. elegans* continue a differentiation program regardless of cell division. We demonstrated that these cells in developing worms continue their cell fate determination and execution programs even when the cell-division cycle is blocked, as occurs in other organisms (Harris and Hartenstein 1991; Amthor et al. 1998).

In summary, *lin-39* activity is essential for two sequential stages in the VPCs: inhibition of *eff-1*-dependent cell fusion to the epidermis, and activation of cell-division cycle mediated by an unidentified Hox effector of proliferation. In addition, it appears that *lin-39* does not control cell migration effectors essential for ring formation and stacking during invagination. Because other Hox genes in *Drosophila*, vertebrates, and *C. elegans* also regulate multiple cell behaviors such as migration, fusion, proliferation, differentiation, and morphogenesis (Salser and Kenyon 1996), it is possible that some of the unidentified effectors are evolutionarily conserved. Identification of these effectors may facilitate the understanding of mechanisms that regulate stem cell division, fusion, and differentiation in mammals (Ying et al. 2002).

Materials and methods

General methods and strains

Worms were handled as described (Brenner 1974). All experiments were performed at 20°C unless otherwise indicated. Wild-type animals were *C. elegans* Bristol N2 strain. The following genes and alleles were used: (LGII) *eff-1(hy21)* (Mohler et al. 2002); (LGIII) *lin-39(n1760)*, *lin-39(n1490ts)*, *ceh-20(ay42)*, *dpy-17(e164)*, *unc-32(e189)*, *unc-36(e251)*; (LGIV) *him-8(e1489)* (Riddle et al. 1997). SU93 *jcls1[ajm-1::GFP, pRF4[rol-6(su1006)]]IV* (Koeppen et al. 2001) served as control in all experiments testing strains on a *jcls1* background.

FC50 *zzEx10[p]E3, pRF4* contains *pE3*, an *eff-1p::GFP* transcriptional construct comprised of a 7.5-kb promoter sequence of *eff-1* fused to the GFP vector pPD95.75 (gift from A. Fire; Mohler et al. 2002).

To analyze single and double mutants with or without *eff-1p::GFP* expression, the following strains were generated: BP92 [*eff-1(hy21)II; unc-36(e251)*, *lin-39(n1490ts)III*]; *jcls1 IV*], BP93 [*unc-36(e251)*, *lin-39(n1490)III*]; *jcls1 IV*], BP94 [*eff-1(hy21)II; lin-39(n1760)III*]; *jcls1 IV*], BP95 [*lin-39(n1760) III*]; *jcls1 IV*], BP121 [*lin-39(n1760)III*]; *zzEx10*], BP123 [*eff-1(hy21) II*]; *unc-36(e251)*, *lin-39(n1490) III*]; *zzEx10*], BP124 [*unc-36(e251)*, *lin-39(n1490) III*]; *zzEx10*], and BP126 [*ceh-20(ay42)*, *unc-36(e251)III*]; *zzEx10*].

BP92 was constructed by mating *jcls1* males with *unc-36(e251)*, *lin-39(n1490)III* hermaphrodites at 15°C (10% of the hermaphrodites have normal vulva at 15°C) and using outcross males for mating with *eff-1(hy21)* hermaphrodites. Rol F1s of this mating were isolated, and F2s were screened for Unc;Vul;Rol worms. These worms were further isolated and F3 were screened for Eff;Unc;Vul;Rol that showed temperature-sensitivity regarding the Vul phenotype. BP93 was constructed during construction of BP92, only Unc;Vul;Rol non-Eff worms were isolated from F3 progeny. BP94 was constructed by mating *jcls1* males with BP76 [*eff-1(hy21)II;jcls1 IV*] and using outcross males for mating with MT4009 [*lin-39(n1760)/dpy-17(e164);unc-32(e189)III*] hermaphrodites. After isolation of Rol F1s, F2s were screened for Rol;Vul worms, and the progeny of these worms (F3) were screened for Eff;Vul;Rol mutants. BP95 was constructed during construction of BP94, only Vul;Rol non-Eff worms were isolated from F3 progeny. BP121 was constructed by mating *him-8(e1489)IV* males with FC50 [*zzEx10(p)E3, pRF4*] hermaphrodites and using outcross males for mating with MT4009 [*lin-39(n1760)/dpy-17(e164);unc-32(e189)III*] hermaphrodites. Rol F1s that express *eff-1p::GFP* were isolated, and Rol F2s were cloned. F3s were screened for Vul *eff-1p::GFP* expressing mutants. BP123 was constructed by mating FC50 [*zzEx10(p)E3, pRF4*] males with BP92 [*eff-1(hy21)II;unc-36(e251);lin-39(n1490ts)III;jcls1*] hermaphrodites at 15°C. Rol non-Unc F1s were transferred to the restrictive temperature of 20°C, and Eff F2s that express *eff-1p::GFP* in the epithelia were isolated. From these worms, F3 were screened for Eff;Unc;Vul worms that express *eff-1p::GFP*. BP124 was constructed during construction of BP123, only Vul F2s expressing *eff-1p::GFP* were isolated and screened for the absence of Eff progeny. BP126 was constructed by mating FC50 [*zzEx10(p)E3, pRF4*] males with NH2296 [*ceh-20(ay42) unc-36(e251)/sma-3(e491) unc-36(e251) III*] hermaphrodites. Rol non-Unc F1s were isolated, and F2 were scored for Unc;Vul worms that express *eff-1p::GFP*.

Immunofluorescence and microscopy

To image larvae, GFP-expressing worms were anesthetized with 5mM NaN₃ or 0.01% levamisole or fixed briefly in 2% paraformaldehyde. To check *ajm-1::GFP* expression, we fixed and stained worms using the mouse monoclonal antibody MH27 (1:300 dilution) that recognizes AJM-1 within the adherens junctions (zonula adherens) of epithelial cells (Francis and Waterston 1991; Shemer et al. 2000). Worms were staged and visualized as described (Shemer et al. 2000). To follow proliferation of cells, worms were analyzed by Nomarski optics.

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