Dimerization-driven degradation of *C. elegans* and human E proteins

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E proteins are conserved regulators of growth and development. We show that the *Caenorhabditis elegans* E-protein helix–loop–helix-2 (HLH-2) functions as a homodimer in directing development and function of the anchor cell (AC) of the gonad, the critical organizer of uterine and vulval development. Our structure–function analysis of HLH-2 indicates that dimerization drives its degradation in other uterine cells (ventral uterine precursor cells [VUs]) that initially have potential to be the AC. We also provide evidence that this mode of dimerization-driven down-regulation can target other basic HLH (bHLH) dimers as well. Remarkably, human E proteins can functionally substitute for *C. elegans* HLH-2 in regulating AC development and also display dimerization-dependent degradation in VUs. Our results suggest that dimerization-driven regulation of bHLH protein stability may be a conserved mechanism for differential regulation in specific cell contexts.

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The E subfamily of basic helix–loop–helix (bHLH) proteins encompasses essential regulators of cell growth and differentiation in mammalian development and tissue homeostasis. Dysregulated expression or activity of E proteins has been associated with leukemias, lymphomas, and autoimmune diseases (for review, see Belle and Zhuang 2014). Thus, understanding the mechanisms that govern E-protein activity and stability are highly relevant to human development and disease.

E proteins bind DNA as obligate dimers, forming either homodimers or heterodimers with “class II” bHLH proteins (Murre et al. 1994; Massari and Murre 2000). Dimer composition provides DNA-binding specificity for the transcriptional activation of target genes (Chien et al. 1996; Grove et al. 2009). *Drosophila* and mammalian E proteins are widely expressed, and, in most developmental contexts that have been studied, E proteins function in heterodimers with class II bHLH partners (Murre et al. 1994; Grove et al. 2009). Expression of these partners is precisely controlled spatially and temporally to achieve tissue-appropriate E-protein activity and target gene expression.

There are also dimerization-based mechanisms that regulate E-protein activity and bHLH stability. The best characterized is the negative regulatory mechanism involving expression of *Drosophila* Emc or mammalian Id, which contain an HLH domain but no DNA-binding domain: Emc/Id forms a heterodimer with E that cannot bind DNA, thereby acting as dominant-negative to reduce E activity (Benezra et al. 1990; Ellis et al. 1990). Id regulates E in diverse developmental and physiological processes, and how it regulates E continues to be investigated (Ling et al. 2014; Miyazaki et al. 2015). In addition, in some contexts, E proteins may stabilize their dimerization partners (Viñals et al. 2004; Roark et al. 2012). In this study, we made the novel observation that dimer formation can instead promote bHLH protein instability. We started with studies of HLH-2, the sole *Caenorhabditis elegans* E protein, and then explored the behavior of other *C. elegans* bHLH proteins and human E2A isoforms.

HLH-2 is central to the proper specification and differentiation of the anchor cell (AC) of the hermaphrodite gonad. The AC is a unique, specialized cell in the ventral uterus that serves as a signaling nexus for uterine and vulval patterning. Four cells initially have the potential to be the AC; only one AC is specified, and the other three become ventral uterine precursor cells (VUs) (Kimble 1981; Seydoux and Greenwald 1989; Seydoux et al. 1990). *hlh-2* plays multiple and sequential roles in the specification and differentiation of the AC (Karp and Greenwald 2003, 2004; Hwang and Sternberg 2004). *hlh-2* endows prospective uterine cells with the potential to be the AC, is essential to resolve which cell becomes the AC in a LIN-12/Notch-mediated interaction, and promotes differentiation and function of the AC by directly activating transcription of target genes that organize uterine and vulval development.

Here we show that HLH-2 functions as a homodimer for all of these roles. Surprisingly, we found that while the HLH-2 homodimer is active and required for HLH-2 function in the AC, HLH-2 homodimerization also drives its post-translational degradation in VUs, thereby spatially restricting its activity to the AC. By analyzing ecotopically expressed class II proteins in the VUs, we show that stability of some class II proteins may also be regulated in a dimerization-dependent manner. Finally, we show that human E proteins can functionally substitute for *C. elegans* HLH-2 and that their stability is also regulated in a dimerization-dependent manner, suggesting that a similar mechanism may be conserved in humans.
Results and Discussion

HLH-2 is down-regulated post-translationally in the presumptive VUs

Specification of the AC is a critical event early in gonadogenesis [Fig. 1A]. Four cells of the ventral uterine region of the somatic gonad primordium, which we term "α" and "β" cells, initially have the potential to be the AC. The two β cells lose the potential to be the AC quickly and always become VUs; the two α cells interact with each other via LIN-12/Notch, engaging feedback mechanisms that amplify initial differences between them; the α cell with relatively lower activity becomes the AC, and the other becomes a VU [Seydoux and Greenwald 1989; Seydoux et al. 1990; Greenwald 2012].

Although hlh-2(0) mutants are embryonic-lethal, reducing hlh-2 activity post-embryonically by RNAi revealed its essential roles in AC specification and function [Karp and Greenwald 2003, 2004; Hwang and Sternberg 2004; Schindler and Sherwood 2011]. Previous work showed that hlh-2 is transcribed in the four α and β cells and continues to be transcribed in the specified AC and VUs but that endogenous HLH-2 protein is detected by antibody staining only in the AC [Karp and Greenwald 2003]. These observations suggested that a post-transcriptional mechanism results in differential HLH-2 accumulation in early ventral uterine cells.

Since hlh-2 is an essential gene, manipulations that alter its expression level or activity might have deleterious pleiotropic effects. To facilitate analysis of HLH-2 regulation, we identified "hlh-2prox," an element from the hlh-2 5' flanking region [Supplemental Fig. S1] that is necessary and sufficient to promote transcription only in the four α and β cells of the developing ventral uterus [Fig. 1B–D] with no detectable expression elsewhere. When hlh-2prox drives expression of GFP alone, the level of fluorescence is initially similar in the α and β cells; as the primordium forms and the cells adopt their fates, GFP remains evident in the presumptive AC and the three VUs, although the VVs sometimes appear dimmer than the AC and αVU (Fig. 1D). In contrast, when hlh-2prox drives expression of GFP fused to HLH-2, GFP-HLH-2 fluorescence is initially observed in all four α and β cells, but as the primordium forms and the cell fates are specified, fluorescence is evident only in the presumptive AC and not in the VUs [Fig. 1C,D]. Thus, GFP-HLH-2(+) has the same pattern of accumulation as endogenous HLH-2 [or HLH-2 marked with a small HA tag] [Karp and Greenwald 2003], indicating that the GFP moiety does not affect HLH-2 stability.

These results establish that the regulation of HLH-2 stability is a post-translational process, since hlh-2prox::GFP-HLH-2 lacks other potential hlh-2 regulatory sequences (UTRs and introns). Furthermore, GFP-HLH-2 is stabilized in VUs when proteasome activity is reduced by RNAi, consistent with degradation of HLH-2 in VUs [Fig. 1E].

HLH-2 functions as a homodimer for all aspects of early ventral uterine development

Heterodimerization partners are involved in all developmental events requiring hlh-2 in *C. elegans* described to date, and heterodimerization is the basis for regulating E-protein function in many cell contexts in *C. elegans, Drosophila,* and mammals. Thus, a simple hypothesis is that differential expression of a heterodimerization partner in the AC or the VUs is responsible for differential stability of HLH-2. However, our expression analysis and optimized functional assays indicated that no other bHLH gene is involved in any aspect of early ventral uterine development [Supplemental Figs. S2–S5], suggesting that HLH-2 functions as a homodimer in ventral uterine development and that differential stability of HLH-2 is not regulated by a bHLH heterodimer partner.

Two additional tests support the inference that HLH-2 functions as a homodimer. First, we analyzed the
sequence of E boxes present in known transcriptional targets of HLH-2 in the AC. In vitro studies have shown that HLH-2 homodimers bind to CAGGTG-type E boxes but not CATATG E boxes, which bind heterodimers in vitro and mediate heterodimer function in vivo [Krause et al. 1997; Harfe et al. 1998; Zhang et al. 1999; Hwang and Sternberg 2004; Hwang et al. 2007; Zhao et al. 2007; Grove et al. 2009]. Thus, if HLH-2 functions as a homodimer in the AC, its known direct targets should have evolutionarily conserved CAGGTG E boxes. Indeed, we found that all four validated targets of HLH-2 in the AC have highly conserved CAGGTG E boxes [Supplemental Fig. S6]. When we altered the two CAGGTG E boxes of the AC-specific Enhancer of lin-3 [ACEL], the best characterized direct target of HLH-2 in the AC [Hwang and Sternberg 2004], to heterodimer-type CATATG E boxes, expression in the AC was lost [Fig. 2B], indicating that the homodimer-binding sequence is important for function in the AC.

Second, we tested the ability of HLH-2[C329S] [HLH (m1)] to rescue abnormalities associated with ventral uterine defects of the hypomorphic allele hlh-2(bx108). In human E2A, the corresponding cysteine, C573, stabilizes the E homodimer through a disulfide bond between the two monomers but is not required to stabilize heterodimers in vitro [Benezra 1994; Markus and Benezra 1999]. Although HLH-2[+1] efficiently rescues hlh-2(bx108), HLH-2[C329S] does not [Fig. 2C], consistent with HLH-2 function as a homodimer in this cell context.

Dimerization of HLH-2 leads to its degradation in VUs

To identify features that mediate differential stability of HLH-2 in the AC and VUs, we used hlh-2prox to express GFP-tagged mutant forms in α and β cells and assayed the effect of the mutations on the pattern of GFP-HLH-2 accumulation [Fig. 3A,B] and function [Supplemental Fig. S8].

Our initial analysis indicated that the bHLH domain is necessary and sufficient for differential stability [Fig. 3A]. The N-terminal region [Nterm], which contains two adjacent predicted protein degradation (PEST) sequences [Fig. 1C; Supplemental Material; Rogers et al. 1986], destabilized GFP, as it resulted in GFP-HLH-2[Nterm] degradation in the AC as well as the VUs, whereas deletion of the PEST sequences from GFP-Nterm [GFP-Nterm[ΔP]] led to stability in all four cells. These results indicate that the PEST sequences promote general turnover when taken out of the HLH-2 structural context but do not regulate the differential stability of HLH-2. When the PEST sequence was deleted from otherwise intact HLH-2 [GFP-HLH-2[ΔP]], or the entire Nterm region was deleted [HLH-2[ΔN]], the wild-type pattern was observed, suggesting that the bHLH domain contains the information needed for proper ventral uterine patterning, i.e., stability in the AC and instability in the VUs. In Supplemental Figure S9, we further support this inference by showing that GFP-HLH-2[ΔP] is properly regulated and retains function in the background of hlh-2(tm1768), an in-frame deletion of the PEST sequences in the endogenous gene.

We then independently mutated either the basic region, which mediates DNA binding, or the bHLH region, which mediates dimerization. Residues that mediate DNA binding of the E47 homodimer and residues that mediate dimerization are well characterized both in vitro and by crystal structure [Voronova and Baltimore 1990; Ellenberger et al. 1990]; these residues are conserved in HLH-2 [Supplemental Fig. S10]. As expected for mutations that abrogate such functions, these mutant forms did not rescue hlh-2(bx108) uterine defects [Supplemental Fig. S8]. Mutation or deletion of residues required for E47 DNA binding had a relatively weak effect on HLH-2 accumulation in VUs, with low expressivity and variable penetrance; in contrast, mutations in conserved residues required for E47 dimerization led to strong HLH-2 accumulation in VUs, with high penetrance and high expressivity [Fig. 3B,D; Supplemental Fig. S9]. Mutations that prevent DNA binding do not prevent dimerization, but dimerization is required for DNA binding [Voronova and Baltimore 1990]. Therefore, the strong stabilization observed for the dimerization mutants but not for the basic region mutants suggests that the ability to dimerize per se, rather than a secondary effect on DNA binding, is the important determinant of stability.

The inference that dimerization is important to direct down-regulation of HLH-2 in VUs was further supported by analysis of two additional mutant forms. First, in addition to the HLH region, human E-protein dimerization also requires “domain C” [Goldfarb et al. 1998], which is conserved in HLH-2 [Supplemental Fig. S10]. Deleting the region encompassing this conserved domain also stabilized HLH-2 in VUs and eliminated its rescuing activity [Fig. 3B; Supplemental Fig. S8]. Second, GFP-HLH-2[C329S] [HLH[Δm1]], which should result in unstable E-protein homodimers without preventing dimer formation per se [Benezra 1994; Markus and Benezra 1999], also is stabilized in VUs [Fig. 3B].

In sum, the analysis of mutant forms indicates that the HLH-2 homodimer, rather than a specific sequence in the bHLH domain, is the key determinant of differential stability in the AC and VU. In addition, since dimerization...
not normally found in a and β cells and examined their stability [Fig. 4A, Supplemental Fig. S11]. All ectopically expressed bHLH proteins were stable in the AC and displayed different degrees of susceptibility to down-regulation. HLH-1/MyoD, HLH-6/ASCL4, and HLH-8/Twist were mostly stable in VUs. Strikingly, LIN-32/ATOH1 and NGN-1/Ngn3 were efficiently down-regulated in all three VUs, consistent with susceptibility to down-regulation comparable with that of HLH-2.

We further investigated LIN-32, the best-characterized dimerization partner of HLH-2 in C. elegans (Portman and Emmons 2000). We found that the bHLH region was sufficient and dimerization was necessary for down-regulation in VUs [Fig. 4A], indicating that the mechanism that degrades HLH-2 homodimers can regulate at least some bHLH heterodimers. We could not identify a specific shared sequence or structural feature in the monomers of the proteins that were more efficiently down-regulated or in the proteins that were not (Supplemental Material), suggesting that a general structural feature of bHLH dimers may be recognized by the dimerization-dependent mechanism, consistent with a dimerization-driven regulatory process.

**Human E proteins can functionally substitute for hlh-2 and display dimerization-dependent degradation in VUs**

The human E2A locus [also called TCF3] encodes two different isoforms: E47 and E12 (Massari and Murre 2000). Both can form homodimers and can form heterodimers with one another or with class II bHLH proteins [Murre et al. 1994]. E47 readily forms homodimers; E12 does so less well due to an inhibitory domain in E12 that is not present in E47 or HLH-2 [Sun and Baltimore 1991; Benezra 1994].

We expressed GFP-tagged forms of E47 and E12 in the four α and β cells and observed that, like HLH-2, both E47 and E12 stably accumulate in the AC but not in the VUs [Fig. 5A, C; Supplemental Fig. S12]. In addition, E47 and E12 each efficiently rescues defects associated with hlh-2(bx108) and hlh-2(RNAi) [Fig. 5B, Supplemental Figs. S8, S13]. Finally, we tested the dimerization dependence of E47 stability and function by making two different mutations that disrupt dimerization in vitro.
Concluding remarks

Our analysis indicates that HLH-2 acts as a homodimer to mediate its known roles in early ventral uterine development—competence, the LIN-12/Notch-mediated AC/ VU decision, and AC differentiation and function—even though these discrete events occur within the same cells in a short temporal window. There have been few roles identified for E homodimers, most notably the “classic” role in early B-lymphocyte development (Murre 2005). Indeed, study of E dimerization partners and HLH-only regulatory proteins (e.g., Id in mammals) has driven much research on the roles of post-translational regulation of bHLH activity in development. Our finding of E homodimers as the regulator of a critical signaling nexus in C. elegans organogenesis suggests that additional roles for E homodimers in mammalian development beyond the role defined in B lymphocytes are awaiting discovery.

The remarkable conservation of function and regulation of human E2A and HLH-2 revealed in our C. elegans assays suggests that the down-regulation mechanism that operates in C. elegans VUs may be conserved in humans as well. Furthermore, given our finding that the mechanism can operate on heterodimers as well as homodimers, there are a large number of possible cell contexts in which it may operate. We therefore suggest that dimerization-driven degradation of bHLH proteins will be a more common regulatory mechanism than is currently recognized.

Materials and methods

C. elegans mutants and transgenes

Strain names and full genotypes for all figures are listed in Supplemental Tables S1 and S2. Conditions used for C. elegans genetic analysis, RNAi, and generation of transgenes are described in the Supplemental Material and Supplemental Table S3.

Plasmid constructions and sequence analysis

Construction of hlh-2::cDNA constructs and methods used to analyze E-box and bHLH sequences are described in the Supplemental Material and Supplemental Figures S6 and S10.

Scoring GFP-bHLH protein stability and function

GFP-bHLH protein stability was assessed in multiple independent transgenic lines based on fluorescence visible at a 400-msec exposure time. GFP-HLH-2 and GFP-E2A protein function was assessed in rescue assays of the hypomorphic hlh-2 allele bx108 and of hlh-2(RNAi). Details of scoring and assays are described in the Supplemental Material and Supplemental Figures S8, S9, and S11–S13.

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