Prefoldin 6 mediates longevity response from heat shock factor 1 to FOXO in *C. elegans*

Heehwa G. Son,^{1,12} Keunhee Seo,^{1,12} Mihwa Seo,^{1,2,3,4} Sangsoon Park,¹ Seokjin Ham,¹ Seon Woo A. An,¹ Eun-Seok Choi,^{1,10,11} Yujin Lee,¹ Haeshim Baek,¹ Eunju Kim,⁵ Youngjae Ryu,⁶ Chang Man Ha,⁶ Ao-Lin Hsu,^{5,7,8} Tae-Young Roh,^{1,9} Sung Key Jang,¹ and Seung-Jae V. Lee^{1,2}

¹Department of Life Sciences, ²School of Interdisciplinary Bioscience and Bioengineering, Pohang University of Science and Technology, Pohang, Gyeongbuk 37673, South Korea; ³Center for plant Aging Research, Institute for Basic Science, ⁴Department of New Biology, Daegu Gyeongbuk Institute of Science and Technology, Daegu 42988, South Korea; ⁵Department of Internal Medicine, Division of Geriatric and Palliative Medicine, University of Michigan, Ann Arbor, Michigan 48109, USA; ⁶Research Division, Korea Brain Research Institute, Daegu 41068, South Korea; ⁷Research Center for Healthy Aging, ⁸Institute of New Drug Development, China Medical University, Taichung 404, Taiwan; ⁹Division of Integrative Biosciences and Biotechnology, Pohang University of Science and Technology, Pohang, Gyeongbuk 37673, South Korea

Heat shock factor 1 (HSF-1) and forkhead box O (FOXO) are key transcription factors that protect cells from various stresses. In *Caenorhabditis elegans*, HSF-1 and FOXO together promote a long life span when insulin/IGF-1 signaling (IIS) is reduced. However, it remains poorly understood how HSF-1 and FOXO cooperate to confer IIS-mediated longevity. Here, we show that prefoldin 6 (PFD-6), a component of the molecular chaperone prefoldin-like complex, relays longevity response from HSF-1 to FOXO under reduced IIS. We found that PFD-6 was specifically required for reduced IIS-mediated longevity by acting in the intestine and hypodermis. We showed that HSF-1 increased the levels of PFD-6 proteins, which in turn directly bound FOXO and enhanced its transcriptional activity. Our work suggests that the prefoldin-like chaperone complex mediates longevity response from HSF-1 to FOXO to increase the life span in animals with reduced IIS.

[**Keywords:** aging; *C. elegans*; DAF-16/FOXO; HSF-1/HSF1; PFD-6/PFDN6]

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The insulin/IGF-1 signaling (IIS) pathway regulates life span in many species, including *Caenorhabditis elegans*, *Drosophila*, and mammals (Kenyon 2010; Altintas et al. 2016). In *C. elegans*, inhibition of *daf-2*, an insulin/IGF-1 receptor homolog, or its downstream phosphoinositide 3-kinase cascade components substantially increases life span (Kenyon et al. 1993; Antebi 2007; Kenyon 2010). The longevity conferred by reduced IIS is mediated by the activation of transcription factors, including the DAF-16/forkhead box O (FOXO) transcription factor, which results in the induction of its target genes (Kenyon et al. 1993; Kenyon 2010). Heat shock transcription factor 1 (HSF-1) also acts downstream from IIS to regulate longevity (Hsu et al. 2003; Morley and Morimoto 2004; Cohen

et al. 2006; Kenyon 2010; Chiang et al. 2012; Seo et al. 2013). Activation of HSF-1 induces genes encoding molecular chaperones, including HSP70 and HSP90 family members and small heat shock proteins, for the maintenance of protein homeostasis. HSF-1 and DAF-16/FOXO act together in part to promote longevity by the transactivation of common targets, most notably small heat shock proteins that contribute to longevity (Hsu et al. 2003; Murphy et al. 2003). In addition, the up-regulation of HSF-1 in neurons increases the activity of DAF-16/FOXO in other tissues in a cell-nonautonomous manner, which is crucial for extending life span (Douglas et al. 2015). Nevertheless, how HSF-1 and DAF-16/FOXO cooperate to exert their longevity response remains poorly understood.

In addition to classical heat shock proteins whose expression is regulated by HSF-1, other chaperones and

¹²These authors contributed equally to this work.

Present addresses: ¹⁰Division of Precision Medicine, Research Institute, National Cancer Center, Goyang, Gyeonggi-do 10408, South Korea; ¹¹Department of Environmental Medical Biology, Institute of Tropical Medicine, Yonsei University College of Medicine, Seoul 03722, South Korea.

Corresponding author: seungjaelee@postech.ac.kr

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chaperonins contribute to protein homeostasis. For example, the prefoldin complex functions as a molecular chaperone for various essential proteins (Boulon et al. 2010; Lundin et al. 2010). In eukaryotes, the prefoldin complex is composed of six protein subunits: two α -type subunits (prefoldin 3 [PFD-3] and PFD-5) and four β -type subunits (PFD-1, PFD-2, PFD-4, and PFD-6) (Boulon et al. 2010; Lundin et al. 2010). The prefoldin complex binds to nascent polypeptide chains of cytoskeletal proteins, actins, and tubulins to help transfer these proteins to the group II chaperonin, TRiC (TCP1 ring complex)/CCT (chaperonin-containing TCP1) for proper folding (Boulon et al. 2010; Lundin et al. 2010) in the cytoplasm. Several prefoldin subunits have other functions in addition to those as components of the prefoldin complex. Mammalian PFD-2 and PFD-6 are components of the R2TP (Rvb1, Rvb2, Tah1 [TPR-containing protein], and Pih1 [protein interacting with Hsp90])/prefoldin-like complex (Cloutier et al. 2009). The R2TP/prefoldin-like complex is required for the assembly of noncytoskeletal complexes, such as RNA polymerase II in the cytoplasm (Boulon et al. 2008, 2010; Cloutier et al. 2009). In addition, several components in prefoldin and R2TP/prefoldin-like complexes function as transcription cofactors acting in the nucleus (Millan-Zambrano and Chavez 2014). Despite the important roles of prefoldin and R2TP/prefoldin-like complex subunits in various cellular functions, their roles in aging and life span regulation remain unknown.

In the present study, we functionally characterized the life span-regulating role of *C. elegans* prefoldin-6/PFD-6, which we identified from a genome-wide *hsf-1* modifier RNAi screen. We showed that *pfd-6* was required for life span extension conferred by *daf-2* mutations. We found that *pfd-6* expression in the intestine and hypodermis was required for *daf-2(-)* longevity. We showed that PFD-6-containing R2TP/prefoldin-like complex contributed to the longevity of *daf-2* mutants. Furthermore, PFD-6 was up-regulated in *daf-2(-)* mutants. We then showed that PFD-6 up-regulated DAF-16/FOXO through physical interaction to induce a subset of DAF-16/FOXO target genes. Thus, PFD-6 appears to extend the life span in animals with reduced IIS by mediating longevity response from HSF-1 to DAF-16/FOXO.

Results

A genome-wide RNAi screen identifies genes that affect longevity conferred by reduced IIS

We sought to identify novel longevity genes that interacted with *hsf-1*. To this end, we performed a genome-wide RNAi screen using reduction-of-function *hsf-1(sy441)* mutants, which display a temperature-sensitive sterility phenotype (Hajdu-Cronin et al. 2004). We identified 17 RNAi clones that enhanced the sterility of the *hsf-1* mutants compared with wild-type animals at 20°C (Supplemental Table S1). We then tested whether any of these RNAi clones affected longevity conferred by *daf-2/insulin/IGF-1* receptor mutations, which up-regulate HSF-1 and require HSF-1 for exerting longevity (Hsu et al.

2003; Morley and Morimoto 2004). RNAi targeting each of five genes—*pfd-6*, *fat-7* (fatty acid desaturase 7) (Murphy et al. 2003), C54C8.3/acetylglucosaminyltransferase, *vps-4* (vacuolar protein sorting 4), and *ntl-2/CCR4-NOT* transcription complex subunit—decreased the life span of the *daf-2* mutants at least 10% greater than that of wild type (Fig. 1A–F); these extents are similar to the effects of RNAi targeting *hsf-1* or *daf-16* (Fig. 1A). Thus, our genetic screen identified novel *hsf-1*-interacting genes required for longevity caused by reduced IIS.

pfd-6 is specifically required for the longevity of daf-2(-) and osm-5(-) mutants

We further determined the requirement of these genes for *daf-2(-)* longevity using available mutants (Supplemental Fig. S1A–D). Strong loss-of-function alleles of *pfd-6* cause semiembryonic lethality and larval arrest phenotypes (Supplemental Fig. S2A). We therefore characterized a point mutant allele of *pfd-6(gk493446)* that causes a Thr16-to-isoleucine (T16I) change in a highly conserved PFD-6 (Supplemental Fig. S2B–E). Similar to *pfd-6* RNAi, *pfd-6(gk493446)* mutations substantially decreased the life span of *daf-2* mutants but had a little effect on that of the wild type (Supplemental Fig. S1A). In addition, *pfd-6* mutation and/or RNAi knockdown suppressed the long life span of *daf-2* mutants with or without 5-fluoro-2'-deoxyuridine (FUDR) (Fig. 2A,B; Supplemental Fig. S2F, G); FUDR is used for preventing *C. elegans* reproduction in life span assays (Park et al. 2017). Thus, *pfd-6* is required for the longevity of *daf-2(-)* mutants.

We asked whether *pfd-6* was required for the longevity of various other long-lived mutants; these include sensory-defective *osm-5* mutants, dietary-restricted *eat-2* mutants, mitochondrial respiration-impaired *isp-1* mutants, TOR downstream *rsk-1/S6* kinase mutants, HIF-1-hyperactive *vhl-1* mutants, and HSF-1-hyperactive *hsb-1* mutants. We found that *pfd-6* RNAi decreased the longevity of *osm-5* mutants (Fig. 2C) while not decreasing the life span of the other tested long-lived mutants (Fig. 2D–H; Supplemental Fig. S2H; see figure legends for discussion). Because impaired sensory neuron-mediated longevity also acts through IIS (Jeong et al. 2012), these data suggest that *pfd-6* influences life span by acting mainly through IIS.

PFD-6 in the intestine and hypodermis is required for daf-2(-) longevity

Next, we characterized transgenic animals that expressed *pfd-6::GFP* under a *pfd-6* promoter (*pfd-6p::pfd-6::GFP*). PFD-6::GFP fusion protein was localized in both the cytosol and nuclei of intestinal, hypodermal, muscle, and neuronal cells throughout the developmental stages (Fig. 3A; Supplemental Fig. S3A). The *pfd-6::GFP* transgene restored longevity in *pfd-6; daf-2* mutants (Supplemental Fig. S3B) while having a marginal effect on the wild-type life span (Supplemental Fig. S3C). We then asked in which tissues *pfd-6* was required for the longevity of *daf-2(-)* mutants using tissue-specific RNAi systems (Qadota

Figure 1. Five *hsf-1*(*sy441*) sterility enhancer RNAi clones decrease the longevity of *daf-2* mutants. (A) The graph shows percentage changes in the average life span of wild-type (X-axis) and *daf-2*(*e1370*) [*daf-2*(-); Y-axis] animals treated with each of 17 RNAi clones obtained from the primary screen. Each circle, square, or triangle indicates an average value of two different sets of life spans. A dashed line indicates an arbitrary cutoff showing RNAi clones that have 10% more life span-decreasing effects on *daf-2* mutants than on wild type. Error bars indicate the standard error of the mean (SEM) of two independent experiments. Worms were treated with RNAi only during adulthood to exclude the possibility that RNAi clones may affect the health of the animals during development. *hsf-1* RNAi and *daf-16* RNAi clones were used as positive controls. (B–F) *pfid-6* RNAi (B), *fat-7* RNAi (C), *C54C8.3* RNAi (D), *vps-4* RNAi (E), and *ntl-2* RNAi (F) substantially decreased the life spans of *daf-2* mutants while having little effect on that of wild-type animals. For *pfid-6*, we confirmed the increased sterility caused by *pfid-6* RNAi in *hsf-1* mutants using solid plates (Supplemental Fig. S1E). See Supplemental Table S2 for statistical analysis and additional repeats.

et al. 2007; Calixto et al. 2010). As expected, dsRNA of *pfid-6* had little effect on the life span of RNAi-defective *rde-1*(-) or systemic RNAi-defective *sid-1*(-) mutants (Supplemental Fig. S3D,E). We found that intestine- or hypodermis-specific *pfid-6* RNAi significantly decreased the long life span of *daf-2*(-) mutants but not that of *daf-2*(+) control animals (Fig. 3B,C). In contrast, muscle- or neuron-specific knockdown of *pfid-6* hardly affected the life span in control or *daf-2*(-) mutant backgrounds (Fig. 3D, E). Thus, together with the expression pattern of *pfid-6::GFP* (Fig. 3A), *pfid-6* in the intestine and hypodermis appears to contribute to the longevity of *daf-2* mutants.

PFD-6 as a component of the R2TP/prefoldin-like complex contributes to longevity

Mammalian PFD-6 acts as a subunit of R2TP/prefoldin-like and prefoldin complexes (Fig. 4A). By performing a large-scale yeast two-hybrid screen, we found that PFD-6 bound various proteins, including F35H10.6/UXT and PFD-5, homologs of R2TP/prefoldin-like and prefoldin complex components, respectively (Fig. 4B; Supplemental Table S7). Thus, *C. elegans* PFD-6 appears to be a subunit of R2TP/prefoldin-like and prefoldin complexes, similar to that in other species (Lundin et al. 2010; Millan-Zambrano and Chavez 2014). We then examined whether *daf-2*(-)-induced longevity was affected by RNAi targeting various components of these two complexes (Fig. 4A; Supplemental Fig. S4A,B). We found that the full longevity of *daf-2*(-) mutants was decreased by RNAi knockdown of *ruvb-1*, *ruvb-2*, or *uri-1* (Fig. 4C–E). We also found

that knockdown of *pfid-2* modestly decreased the longevity conferred by *daf-2* mutations (Fig. 4F). In addition, the life span-shortening effects of *ruvb-1* RNAi, *ruvb-2* RNAi, or *uri-1* RNAi were not additive to those of *pfid-6* mutations (Supplemental Fig. S4D–F). In contrast, RNAi clones targeting prefoldin complex-specific components showed minimal effects on longevity (Supplemental Fig. S4H–K). These data suggest that the R2TP/prefoldin-like complex, rather than the prefoldin complex, contributes to the long life span of *daf-2*(-) mutants.

daf-2 mutations increase PFD-6 levels in an HSF-1-dependent manner

We next asked whether reduced IIS affected the level of PFD-6 and found that the PFD-6::GFP level was increased by genetic inhibition of *daf-2* using fluorescence imaging and Western blot assays (Fig. 5A–E). Importantly, we showed that this increase was suppressed by *hsf-1* RNAi (Fig. 5A,B; Supplemental Fig. S5). In contrast, *daf-16* RNAi did not affect the level of PFD-6::GFP (Fig. 5A,B; Supplemental Fig. S5). Different from the PFD-6 protein level, the *pfid-6* mRNA level was not changed in *daf-2* mutants (Fig. 5F). These data suggest that inhibition of *daf-2* up-regulates PFD-6 protein levels via HSF-1.

PFD-6 regulates the expression of a large subset of DAF-16/FOXO target genes in *daf-2* mutants

Because the R2TP/prefoldin-like complex can regulate transcription (Millan-Zambrano and Chavez 2014), we

Figure 2. *pfd-6* is required for the longevity of *daf-2* mutants. (A,B) *pfd-6(gk493446)* [*pfd-6(-)*] point mutation causing a T16I change with *pfd-6* RNAi partly suppressed the longevity of *daf-2(e1370)* [*daf-2(-)*] mutants with (A) or without (B) FUDR. (A) *pfd-6* RNAi during adulthood slightly decreased the short life span of *pfd-6; daf-2* double mutants but did not affect that of *pfd-6* mutants (two out of four trials). (C) *pfd-6* RNAi partially suppressed the longevity conferred by *osm-5(p813)* [*osm-5(-)*] mutations. (D–G) *pfd-6* RNAi did not decrease the longevity of *eat-2(ad1116)* [*eat-2(-)*] (D), *isp-1(qm150)* [*isp-1(-)*] (E), *rsk-1(tm1714)* [*rsk-1(-)*] (F), and *vhl-1(ok161)* [*vhl-1(-)*] (G) animals. (H) *pfd-6* mutation did not decrease the life span of *hsb-1(cg116)* [*hsb-1(-)*] mutants. Considering our data suggesting that *pfd-6* mediates the longevity effects of HSF-1, it is surprising to find that genetic inhibition of *pfd-6* did not decrease the longevity of *rsk-1* or *hsb-1* mutants, which require HSF-1 for a longer life span (Chiang et al. 2012; Seo et al. 2013). We speculate that up-regulation of HSF-1 by *rsk-1* and *hsb-1* mutations may affect *pfd-6* differently than that by *daf-2* and *osm-5* mutations, which will be interesting to examine in future studies. See Supplemental Table S3 for statistical analysis and additional repeats.

wondered whether *pfd-6* influenced global gene expression in *daf-2(-)* mutants. By performing mRNA sequencing (mRNA-seq), we identified 189 up-regulated and 162 down-regulated genes in *daf-2* mutants whose expression was dependent on *pfd-6* (fold change >1.5 and <0.67 for up-regulated and down-regulated genes, respectively; $P < 0.05$) (Fig. 6A; Supplemental Fig. S6A–D). Gene ontology (GO) analysis using these genes identified several enriched GO terms, including flavonoid biosynthetic process among up-regulated genes and centriole replication among down-regulated genes (Fig. 6B,C). We then examined which longevity factors in IIS acted together with PFD-6. Specifically, we compared the transcriptome regulated by PFD-6 in *daf-2(-)* in this study with those regulated by DAF-16/FOXO (Riedel et al. 2013; Chen et al. 2015), SKN-1/NRF (Ewald et al. 2015), and SMG-2/UPF1 (Son et al. 2017), which are crucial for longevity conferred by reduced IIS (Kenyon 2010; Blackwell et al. 2015; Son and Lee 2017). Interestingly, genes differentially regulated in *daf-2* mutants by PFD-6 displayed a highly significant overlap with those regulated by DAF-16/FOXO but not with those regulated by SKN-1/NRF or SMG-2/UPF1 (Fig. 6D–F; Supplemental Fig. S6E–K). Thus, PFD-6 appears to act together with DAF-16/FOXO to regulate a subset of gene expression in *daf-2(-)* mutants. We confirmed these results for selected genes using quantitative RT-PCR (qRT-PCR) (Fig. 6G–M; Supplemental Fig. S6L–N). We also found that the consensus sequence of the DAF-16-binding element (DBE) was significantly enriched in the 5' upstream regions of *pfd-6*-dependent up-regulated genes in *daf-2(-)* mutants (Fig. 6N; Supplemental Fig. S6O,P). Together, these data sug-

gest that PFD-6 regulates the expression of a subset of DAF-16/FOXO target genes in *daf-2(-)* mutants.

PFD-6 appears to increase the activity of DAF-16/FOXO via physical interaction

Next, we asked how PFD-6 affected the activities of DAF-16/FOXO. First, we established that knocking down *hsf-1* or *daf-16* in the intestine, hypodermis, or muscles significantly decreased the long life span of *daf-2* mutants (Fig. 7A–D; Supplemental Fig. S7A–D). Thus, consistent with published reports regarding the tissue-specific transgenic expression of DAF-16/FOXO (Libina et al. 2003; Zhang et al. 2013) and our tissue-specific *pfd-6* RNAi data (Fig. 3B,C), PFD-6 appears to act with HSF-1 and DAF-16 in common tissues, such as the intestine and hypodermis.

We next tested whether *pfd-6* affected any of several known DAF-16/FOXO activation mechanisms, nuclear translocation, increases in the protein level, and regulation of transcriptional activity via physical binding (Altintas et al. 2016). We found that the genetic inhibition of *pfd-6* did not affect the subcellular localization of DAF-16/FOXO (Fig. 7E,F) while having variable effects on the level of DAF-16/FOXO (Supplemental Fig. S7E–H). We then showed that PFD-6 bound to DAF-16/FOXO by using yeast two-hybrid and split GFP system assays (Fig. 7G,H; Supplemental Fig. S7I,J). In contrast, PFD-6 did not bind HSF-1 (Fig. 7I), and this result is consistent with our data suggesting that HSF-1 acts upstream of PFD-6 to regulate its protein level (Fig. 5A,B; Supplemental Fig. S5). Together, these data suggest that HSF-1 up-regulates

Figure 3. Intestinal and hypodermal PFD-6 contributes to the long life span of *daf-2* mutants. (A) *pfd-6p::pfd-6::GFP* was expressed in various tissues (panel a), including the intestine (panel b), the hypodermis (panel c), neurons (panel d), and muscles (panel e). Arrowheads indicate PFD-6::GFP in the intestine, hypodermis, neurons, or muscles. Bars, 50 μ m. (B,C) The longevity of *daf-2* (*e1370*) [*daf-2(-)*] mutants was significantly decreased by intestine [*rde-1(ne219); kbls7[nhx-2p::rde-1; rol-6D]*]-specific *pfd-6* RNAi (B) or hypodermis [*rde-1(ne219); kzl9[lin-26p::mIs::gfp; lin-26p::rde-1; rol-6D]*]-specific *pfd-6* RNAi (C). We noticed that life span-decreasing effects of the intestine- or the hypodermis-specific *pfd-6* RNAi were greater than the life span-decreasing effects of *pfd-6* RNAi on *rde-1(-); daf-2(-)* animals (Supplemental Fig. S3D). (D,E) Muscle [*rde-1(ne219); kzl9[lin-26p::mIs::gfp; lin-26p::rde-1; rol-6D]*]-specific (D) or neuron [*sid-1(pk3321); uls69[myo-2p::mCherry; punc-119::sid-1]*]-specific (E) *pfd-6* RNAi did not affect the

life span of *daf-2(+)* or *daf-2(-)* animals. We confirmed the efficiency of neuron- and muscle-specific *pfd-6* RNAi knockdown using fluorescence-tagged PFD-6 proteins (Supplemental Fig. S3F–I). See Supplemental Table S4 for statistical analysis and additional repeats.

PFD-6, which increases the activity of DAF-16/FOXO (likely through physical interaction) and promotes the longevity of *daf-2* mutants.

Discussion

PFD-6 mediates the cross-talk between HSF-1 and DAF-16/FOXO for longevity

HSF-1 and DAF-16/FOXO are key longevity-promoting transcription factors that act in the IIS pathway. Although the functional relationship between HSF-1 and DAF-16/FOXO for longevity regulation has been established, how HSF-1 and DAF-16/FOXO communicate with each other has remained incompletely understood. In this report, we showed that *C. elegans* PFD-6 acted as a key life span regulatory protein that mediated cross-talk between HSF-1 and DAF-16/FOXO in the IIS pathway.

Previous studies have indicated that *C. elegans* HSF-1 and DAF-16/FOXO transcription factors act both separately and together in IIS to elicit physiological responses. HSF-1 is up-regulated in worms with reduced IIS via dissociation from its inhibitory factors, such as DDL-1 (Chiang et al. 2012). Inhibition of IIS promotes the translocation of DAF-16/FOXO from the cytoplasm to the nucleus and increases DAF-16/FOXO activity via diverse regulators (Kenyon 2010; Murphy and Hu 2013). HSF-1 and DAF-16/FOXO also work in parallel in IIS by inducing common targets, including small heat shock proteins (Hsu et al. 2003; Murphy et al. 2003). For example, both DAF-16- and HSF-1-binding motifs exist in the promoter regions of genes encoding several heat shock proteins (GuhaThakurta et al. 2002; Hsu et al. 2003; Murphy et al. 2003; Xu and Kim 2012; Tepper et al. 2013). In addition, HSF-1 overexpression promotes longevity through increasing DAF-16 activity

(Hsu et al. 2003; Douglas et al. 2015). However, it remained unclear whether and how a linear process between HSF-1 and FOXO contributes to longevity responses. Here, we show that the up-regulation of PFD-6 via HSF-1 increases the transcriptional activity of DAF-16/FOXO. Therefore, PFD-6 appears to promote longevity by acting in a linear pathway from HSF-1 to DAF-16/FOXO under reduced IIS.

A genome-wide RNAi screen identified novel daf-2(-) longevity-regulating factors

C. elegans is an excellent model organism to identify novel functions of genes through genetic screens. A previous genome-wide RNAi screen identified genes that are required for longevity conferred by *daf-2* mutations (Samuelson et al. 2007). Among them, a high fraction of genes is involved in vesicle transport, such as *vps-16* (vacuolar protein sorting 16). Despite no direct overlap between the previous study (Samuelson et al. 2007) and the present work, our screen identified a vacuolar protein, *vps-4* (Fig. 1E), suggesting that proper vesicle transport is crucial for *daf-2(-)* longevity. It will be interesting to determine how vesicular transport regulates longevity conferred by reduced IIS in future research.

PFD-6 appears to affect longevity as a component of the R2TP/prefoldin-like complex

The prefoldin complex is composed of PFD-1 through PFD-6 and acts as a chaperone for the proper folding of many essential proteins, including actins and tubulins (Vainberg et al. 1998; Lundin et al. 2010; Millan-Zambrano and Chavez 2014). During aging, intestinal lumens deteriorate and suffer from the loss of microvilli, which

Figure 4. Prefoldin-like complex components are required for the long life span of *daf-2* mutants. (A) Schematic model of the R2TP/prefoldin-like complex and the prefoldin complex in mammals; the R2TP/prefoldin-like complex interacts with RNA polymerase II in the cytoplasm to help its assembly, and the prefoldin complex is composed of PFD-1, PFD-2, PFD-3, PFD-4, PFD-5, and PFD-6, which bind to nascent actin or tubulin polypeptides. Prefoldin complex-bound polypeptides are transferred to TRiC/CCT chaperonin for proper folding. (B) Yeast two-hybrid screening with full-length PFD-6 as bait was performed using the yeast PBN204 strain. The interaction was validated using nutritional selection (*ADE2*) for the growth assay. See Supplemental Table S7 for the description of candidate genes. (C–F) Several R2TP/prefoldin-like complex components were required for life span extension caused by reduced IIS. RNAi targeting *ruvb-1* (C), *ruvb-2* (D), *uri-1* (E), or *pfd-2* (F) significantly decreased the long life span of *daf-2(e1370)* [*daf-2(-)*] mutants. Knockdown of potential UXT homolog F35H10.6 did not affect the life span of wild-type, *daf-2(-)*, *pfd-6(-)*, or *pfd-6*; *daf-2* mutants (Supplemental Fig. S4C,G). See Supplemental Table S5 for statistical analysis and additional repeats.

contain core actin filaments and are important for cytoskeletal integrity (McGee et al. 2011). ACT-5 (actin 5), a potential target of the prefoldin complex, is required for the formation of microvilli in intestinal cells (MacQueen et al. 2005). Overexpression of a modified HSF-1 extends life span and enhances cytoskeletal integrity (Baird et al. 2014). In addition, the chaperonin activity of the TRiC/CCT complex, which works together with the prefoldin complex for cytoskeletal integrity (Vainberg et al. 1998), contributes to longevity in *C. elegans* (Noormohammadi et al. 2016). Therefore, we tested whether the genetic inhibition of *pfd-6* affected the cytoskeletal integrity in *daf-2* mutants during aging. We used filamentous actin (F-actin) reporter *Lifeact::gfp*-expressing *C. elegans* (Farooqui et al. 2012) and found that genetic inhibition of *pfd-6* did not affect the *Lifeact::GFP* expression pattern in wild-type or *daf-2* mutant worms (Supplemental Fig. S8). These data are consistent with the possibility that *pfd-6* regulates the longevity of *daf-2* mutants independently of the maintenance of cytoskeletal integrity. However, one caveat of this interpretation is that we used a hypomorphic mutant allele of *pfd-6*, as *pfd-6*-null mutations cause semiembryonic lethality and larval arrest. Thus, it will be crucial to further examine the relationship between prefoldins, aging, and cytoskeletal integrity with comprehensive genetic and biochemical tools in future studies.

The R2TP/prefoldin-like complex is expressed in the nucleus as well as cytoplasm (Millan-Zambrano and Chavez 2014), consistent with our results regarding PFD-6 expression (Fig. 3A). The R2TP/prefoldin-like complex pro-

motes assembly of RNA polymerase II in the cytoplasm (Boulon et al. 2010). Several molecular functions of the nuclear R2TP/prefoldin-like complex have also been discovered (Millan-Zambrano and Chavez 2014). *C. elegans* *uri-1* is required for DNA stability (Parusel et al. 2006). Mammalian UXT is a cofactor of the NF- κ B enhanceosome, which promotes the induction of NF- κ B target gene expression (Sun et al. 2007). In addition, a large fraction of nuclear proteins that bind the R2TP/prefoldin-like complex are transcription factors (Millan-Zambrano and Chavez 2014). A previous study also reported that PFD-6 physically interacts with DAF-16/FOXO by using mass spectrometry (Riedel et al. 2013). Consistent with this, we showed that PFD-6 bound DAF-16/FOXO in the nucleus as well as cytoplasm using an in vivo split-GFP system assay (Fig. 7H; Supplemental Fig. S7J). Thus, PFD-6 as a component of the R2TP/prefoldin-like complex is crucial for the longevity of *daf-2(-)* mutants through regulating the transcriptional activity of DAF-16/FOXO.

The tissue-specific requirement of PFD-6 is consistent with that of HSF-1 and DAF-16/FOXO

The activities of HSF-1 and DAF-16/FOXO in several tissues have been shown to contribute to the longevity of *daf-2* mutants (Libina et al. 2003; Morley and Morimoto 2004; Zhang et al. 2013; Douglas et al. 2015). Overexpression of HSF-1 in the intestine, neurons, or muscles increases life span in wild-type worms (Morley and Morimoto

Figure 5. HSF-1 increases the levels of PFD-6. (A) Mutations in *daf-2* increased PFD-6 levels, which were decreased by *hsf-1* RNAi but not by *daf-16* RNAi. Bar, 100 μ m. The following animals that expressed an extrachromosomal array of *pfd-6::GFP* were used: IJ570 (*yhEx134[pfd-6p::pfd-6::GFP; odr-1p::RFP]*) and IJ810 (*daf-2(e1370) [yhEx134[pfd-6p::pfd-6::GFP, odr-1p::RFP]*). (B) Quantification of A. $n > 15$ from two independent experiments. (C) GFP levels in integrated *pfd-6::GFP* transgenic animals (IJ1249: *yhIs74[pfd-6p::pfd-6::GFP; odr-1p::RFP]*) were examined. $n > 32$ from three independent experiments. (D) Western blot assays were performed using the IJ1249 strain. A representative blot is shown from three repeats that displayed consistent results. α -Tubulin was used as a loading control. (WB) Western blot. (E) Quantification of D. (F) mRNA levels of *pfd-6* in wild-type, *daf-2(e1370) [daf-2(-)]*, *pfd-6(gk493446, RNAi) [pfd-6(-)]*, and *pfd-6(-); daf-2(-)* animals were measured by using quantitative RT-PCR. $n = 3$. Error bars represent SEM. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$, two-tailed Student's *t*-test.

2004; Douglas et al. 2015). In addition, a transgene that expresses a dominant-negative HSF-1 in neurons or body wall muscles shortens the long life span of *age-1*/PI3 kinase mutants (Morley and Morimoto 2004). Transgenic expression of *daf-16*/FOXO in the intestine, neurons, or hypodermis also partially restores the longevity of *daf-2(-)* mutants (Libina et al. 2003; Zhang et al. 2013). Here, we performed life span assays with tissue-specific RNAi targeting *hsf-1* and *daf-16* and found that knockdown of *hsf-1* or *daf-16* in the intestine, hypodermis, or muscles significantly decreased longevity in *daf-2(-)* mutants. Overall, these findings are in part consistent with our data using PFD-6, whose activities in the intestine and hypodermis were crucial for the longevity of *daf-2* mutants. Moreover, we found that PFD-6 bound DAF-16/FOXO in vivo. Therefore, a life span-regulating pathway from HSF-1 to DAF-16/FOXO through PFD-6 appears to act together at least in these two tissues: the intestine and hypodermis. It will be important to determine whether the communication among HSF-1, PFD-6, and DAF-16/FOXO is strictly tissue-autonomous or has tissue-nonautonomous features.

HSF-1 may up-regulate HSP-70 and HSP-90, which in turn stabilize PFD-6 with reduced IIS

How does up-regulated HSF-1 increase PFD-6 protein levels in *daf-2* mutants? In flies and mammals, the R2TP/prefoldin-like complex functions with chaperones HSP70 and HSP90 (Boulon et al. 2010; Benbahouche Nel et al.

2014), both of which are targets of HSF-1 (Vihervaara and Sistonen 2014; Dai 2018). Consistently, we found that both *C. elegans* HSP-90 and HSP-70 bound PFD-6 (Supplemental Fig. S9). Based on these results, we speculate that HSP-70 and HSP-90, which are up-regulated by HSF-1 in *daf-2* mutants, bind and stabilize PFD-6 to contribute to longevity.

The R2TP/prefoldin-like complex may play roles in aging or age-related diseases through heat shock factors and FOXO transcription factors

The physiological roles of heat shock factors and FOXOs in aging and age-related diseases are relatively well established. As master transcription factors for the induction of cytosolic chaperones, heat shock factors are crucial for modulating proteostasis-related and age-associated diseases, including Alzheimer's and Parkinson's diseases (Gomez-Pastor et al. 2018). Multiple studies with human populations have indicated that polymorphisms in FOXO3A are linked to longevity (Kenyon 2010; Tazearslan et al. 2012; Webb and Brunet 2014; Martins et al. 2016). Therefore, it will not be surprising to find that heat shock factors and FOXO transcription factors act together to influence aging and age-associated diseases in mammals, consistent with discoveries made using *C. elegans*. Prefoldin and R2TP/prefoldin-like complex components, including PFD-6, are evolutionarily conserved from archaea to humans. Although the role of the

Figure 6. PFD-6 up-regulates a large subset of DAF-16 target genes in *daf-2* mutants. (A) Scatter plots show the comparison of gene expression between wild-type and *daf-2* (*e1370*) [*daf-2(-)*] or wild-type and *pfd-6* (*gk493446*, RNAi); *daf-2* (*e1370*) [*pfd-6(-); daf-2(-)*] animals. Black dots indicate gene expression in *daf-2* mutants compared with wild type, whose levels were significantly increased (fold change > 1.50 ; $P < 0.05$; left) or decreased (fold change < 0.67 ; $P < 0.05$; right). Red dots indicate expression levels in *pfd-6*; *daf-2* mutants compared with wild type. See Supplemental Table S8 for the list of up-regulated or down-regulated genes in *daf-2* mutants compared with wild-type or *pfd-6*; *daf-2* double mutants. (B,C) Overrepresented GO terms of up-regulated (B) or down-regulated (C) genes that are shown in A. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$. (D–F) Venn diagrams indicate overlapping genes whose expression was increased by *daf-2* mutations in a *pfd-6*- and *daf-16*-dependent (Riedel et al. 2013) (D), *pfd-6*- and *skn-1*-dependent (Ewald et al. 2015) (E), or *pfd-6*- and *smg-2*-dependent (Son et al. 2017) (F) manner. Fold change > 1.5 . $P < 0.05$. (RF) Representation factor. P -values for the overlaps of Venn diagrams were calculated using the hypergeometric probability test. (G–M) qRT-PCR results showing mRNA levels of DAF-16 target genes *F12A10.1* (G), *F26G1.2* (H), *C06G3.3* (I), *C02F12.5* (J), *gpd-4* (K), *sod-3* (L), and *D1086.1* (M). $n > 4$. Error bars represent SEM. (*) $P < 0.05$; (**) $P < 0.01$; (***) $P < 0.001$, two-tailed Student's *t*-test. (N) Motif analysis was performed by using 800-base-pair (bp) upstream sequences of *pfd-6*-dependent genes that were up-regulated in *daf-2* mutants. Fold change > 1.5 . $P < 0.05$. See Supplemental Figure S6, O and P, for other enriched motifs.

recently discovered R2TP/prefoldin-like complex in diseases remains unexplored, prefoldin complex components are implicated in protecting against Huntington's and Alzheimer's diseases (Sorgjerd et al. 2013; Tashiro et al. 2013). Therefore, it will be interesting to test whether PFD-6 or the R2TP/prefoldin-like complex plays roles in age-related diseases or longevity in mammals through working with heat shock factors or FOXOs.

Materials and methods

Strains

The following strains were used in this study.

Some strains were provided by the *Caenorhabditis* Genetics Center, which is funded by the National Institutes of Health Office of Research Infrastructure Programs (P40 OD010440). The strains were N2 wild type, CF1041: *daf-2* (*e1370*) III, CF2495: *hsf-1* (*sy441*) I, CF1085: *daf-16* (*mu86*) I, *daf-2* (*e1370*) III, IJ462: *pfd-6* (*gk493446*) I, IJ485: *pfd-6* (*gk493446*) I, *daf-2* (*e1370*) III, IJ547: *C54C8.3* (*gk896531*) I, IJ548: *C54C8.3* (*gk896531*) I, *daf-2* (*e1370*)

III, IJ479: *ntl-2* (*gk390728*) II, IJ480: *ntl-2* (*gk390728*) II, *daf-2* (*e1370*) III, IJ373: *fat-7* (*wa36*) V, IJ389: *daf-2* (*e1370*) III, *fat-7* (*wa36*) V, EQ150: *hsb-1* (*cg116*) IV, IJ1099: *hsb-1* (*cg116*) IV, IJ1105: *pfd-6* (*gk493446*) I, *hsb-1* (*cg116*) IV, CF1908: *eat-2* (*ad1116*) II, CF2172: *isp-1* (*qm150*) IV, IJ109: *rsks-1* (*tm1714*) III, CF2553: *osm-5* (*p813*) X, IJ7: *vhl-1* (*ok161*) X, *pfd-6* (*tm3759*) I/hT2 [*bli-4* (*e937*) *let-3* (*q782*) *qIs48* [*Pmyo-2::gfp*; *Ppes-10::gfp*; *Pges-1::gfp*]], *pfd-6* (*tm3510*) I/hT2, IJ943: *daf-2* (*gk493446*) I, *daf-2* (*e1370*) III, *yhEx133* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ944: *pfd-6* (*gk493446*) I, *daf-2* (*e1370*) III, *yhEx134* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ568: *yhEx132* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ569: *yhEx133* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ570: *yhEx134* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ810: *daf-2* (*e1370*) III, *yhEx134* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ1249: *yhIs74* [*pfd-6p::pfd-6::gfp*; *odr-1p::rfp*], IJ1562: *yhEx420* [*vha-6p::hsp-70::spGFP1-10*; *odr-1p::rfp*], *yhEx406* [*vha-6p::pfd-6::spGFP11*; *unc-122p::rfp*], IJ1579: *yhEx424* [*vha-6p::hsp-90::spGFP1-10*; *odr-1p::rfp*], *yhEx406* [*vha-6p::pfd-6::spGFP11*; *unc-122p::rfp*], IJ1732: *yhEx413* [*vha-6p::spGFP1-10*; *odr-1p::rfp*], *yhEx406* [*vha-6p::pfd-6::spGFP11*; *unc-122p::rfp*], IJ1532: *yhEx405* [*vha-6p::pfd-6::spGFP1-10*; *odr-1p::rfp*], *yhEx406* [*vha-6p::pfd-6::spGFP11*; *unc-122p::rfp*], IJ1058: *daf-16* (*mu86*) I, *mulIs112* [*daf-16p::gfp::daf-16cDNA*; *odr-1p::rfp*], IJ1230: *pfd-6* (*gk493446*) *daf-16* (*mu86*) I, *mulIs112* [*daf-16p::gfp::*

Figure 7. PFD-6 binds and up-regulates DAF-16/FOXO in *daf-2(-)* mutants. (A–D) Intestine-specific (A,B) or hypodermis-specific (C,D) *hsf-1* or *daf-16* RNAi decreased the longevity of *daf-2(e1370)* [*daf-2(-)*] mutants. See Supplemental Table S6 for statistical analysis and additional repeats. (E) *daf-2* RNAi-induced nuclear localization of DAF-16::GFP was not affected by *pfd-6* mutations. (F) Quantification of E. *n* > 18 from three independent experiments. Bars, 100 μ m. (G) Yeast two-hybrid assays showed the direct interaction between PFD-6 and DAF-16. (H) GFP N-terminal domain-fused DAF-16 isoform a [DAF-16::spGFPN] bound GFP C-terminal domain-fused PFD-6 (PFD-6::spGFPN). See Supplemental Figure S7 for experimental results using *daf-2* RNAi-treated animals. (I) HSF-1::spGFPN did not bind PFD-6::spGFPN. Triangles indicate *odr-1p::RFP*, a coinjection marker for DAF-16::spGFPN and HSF-1::spGFPN. Asterisks indicate *unc-122p::RFP*, a coinjection marker for PFD-6::spGFPN.

daf-16cDNA; *odr-1p::rfp*], TJ356: *zIs356[daf-16p::daf-16a/b::gfp; rol-6]*, IJ1125: *pfd-6(gk493446)* I; *zIs356[daf-16p::daf-16::gfp; rol-6D]*, IJ1530: *yhEx402[vha-6p::daf-16a::spGFP1-10; odr-1p::rfp]*; *yhEx406[vha-6p::pfd-6::spGFP11; unc-122p::rfp]*, IJ1533: *yhEx411[vha-6p::hsf-1::spGFP1-10; odr-1p::rfp]*; *yhEx406[vha-6p::pfd-6::spGFP11; unc-122p::rfp]*, IJ1061: *yhEx240[rpl-28p::lifeact::egfp; odr-1p::rfp]*, IJ1063: *daf-2(e1370)* III; *yhEx240[rpl-28p::lifeact::egfp; odr-1p::rfp]*, WM27: *rde-1(ne219)* V, NR222: *rde-1(ne219)* V; *kzIs9[lin-26p::nls::gfp; lin-26p::rde-1; rol-6D]*, NR350: *rde-1(ne219)* V; *kzIs20[hhlh-1p::rde-1; sur-5p::nls::gfp]*, VP303: *rde-1(ne219)* V; *kzIs7[nhx-2p::rde-1; rol-6D]*, IJ411: *daf-2(e1370)* III; *rde-1(ne219)* V, IJ415: *daf-2(e1370)* III; *rde-1(ne219)* V; *kzIs9[lin-26p::nls::gfp; lin-26p::rde-1; rol-6D]*, IJ416: *daf-2(e1370)* III; *rde-1(ne219)* V; *kzIs20[hhlh-1p::rde-1; sur-5p::nls::gfp]*, IJ417: *daf-2(e1370)* III; *rde-1(ne219)* V; *kzIs7[nhx-2p::rde-1; rol-6D]*, NL3321: *sid-1(pk3321)* V, IJ807: *daf-2(e1370)* III; *sid-1(pk3321)* V, TU3401: *sid-1(pk3321)* V; *uIs69[myo-2p::mCherry; unc-119p::sid-1]*, IJ898: *daf-2(e1370)* III; *sid-1(pk3321)* V; *uIs69[myo-2p::mCherry; unc-119p::sid-1]*; *sid-1(pk3321)* V, IJ1862: *uIs69[myo-2p::mCherry; unc-119p::sid-1]*; *yhEx507[unc-119p::pfd-6::gfp; ofm-1p::rfp]*, and IJ1863: *rde-1(ne219)* V; *kzIs20[pDM#715(hhlh-1p::rde-1) + pTG95(sur-5p::nls::GFP)]*; *yhEx508[hhlh-1p::pfd-6::mch; ofm-1p::gfp]*.

A genome-wide RNAi screen

A genome-wide RNAi screen was conducted as described previously with modifications (Lee et al. 2010). *Escherichia coli* (HT115) bacteria expressing dsRNA targeting *C. elegans* genes were cultured in 96-well plates for 16 h at 37°C. The plates were then duplicated into 200 μ L of LB that contained 50 μ g/mL ampicillin (Usb Corp.) and cultured for 16 h at 37°C. dsRNA expression was induced by the addition of 4 mM isopropyl β -D-thiogalactoside (IPTG) (Gold Biotechnology) for 2 h at 37°C. After pelleting *E. coli* by centrifugation at 2000 rpm for 10 min to remove LB liquid, the

bacteria were resuspended in 100 μ L of NG liquid containing 50 μ g/mL ampicillin and 4 mM IPTG. Synchronized L1 larvae of N2 and *hsf-1(sy441)* animals were prepared by performing bleaching (Stiernagle 2006) and subsequent overnight starvation in liquid M9 buffer. Approximately 15–20 N2 or *hsf-1(sy441)* L1 larvae were deposited into each well of 96-well plates containing dsRNA-expressing *E. coli*. After 3 d of incubation at 20°C, the sterility of N2 and *hsf-1(sy441)* was monitored by examining the existence of larval progeny. RNAi clones that enhanced the sterility of *hsf-1(sy441)* mutants but did not cause sterility in N2 animals were scored as positive hits. The positive hits from the initial RNAi screen were confirmed by repeating four times using liquid medium. To test the sterility on solid plates, gravid N2 and *hsf-1(sy441)* worms were placed on control RNAi-seeded plates. After 1 d, one L1 worm was transferred to a new plate seeded with control RNAi or *pfd-6* RNAi bacteria, and a total of 50 individual L1 worms was transferred onto 50 plates seeded with respective RNAi bacteria. The number of plates that contained egg-laying adults was then measured after 4 d of culture.

Life span assays

Life span assays were performed as described previously with modifications (Son et al. 2017). Worms were maintained at 20°C for at least two generations before life span assays, which were performed at 20°C except for *rsks-1(tm1714)* mutants. The *rsks-1* mutant life span assay was performed as described previously at 22.5°C (Seo et al. 2013). To synchronize the other worm strains, gravid adults were allowed to lay eggs for 12 h on RNAi bacteria-seeded plates. The synchronized progeny were allowed to reach L4 stage, and ~25–35 worms were transferred to plates seeded with gene-specific RNAi bacteria. After 1 d, worms were transferred onto new RNAi plates treated with 5 μ M FUDR (Sigma). Worms were transferred to new RNAi plates every 1 or 2 d until they stopped laying eggs for the life span assay without

FUDR. During life span experiments, worms that ruptured, burrowed, crawled off the plates, or had internal hatching of progeny were censored but were included for statistical analysis, which was performed by using OASIS 2 [online application of survival analysis; <http://sbi.postech.ac.kr/oasis2>] (Yang et al. 2011; Han et al. 2016).

Sequence alignment for PFD-6

Amino acid sequence alignments of PFD-6 among species were conducted by using ClustalW2 format (McWilliam et al. 2013; <http://www.ebi.ac.uk/Tools/msa/clustalw2>) and ClustalX (Jordan and Piel 2008; <http://www.clustal.org/clustal2>).

Schematic domain structure of PFD-6

The exon–intron pattern of genomic *pfid-6* was made by using Exon–Intron Graphic Maker (<http://wormweb.org/exonintron>).

Plasmid construction and generation of transgenic worms

The 832-base-pair (bp) region of the *pfid-6* promoter and *pfid-6* genomic DNA were amplified and cloned into pPD95.75 by using the In-Fusion cloning kit (CloneTech). Plasmids for split GFP assays were generated as described previously (Jeong et al. 2017). The intestine-specific promoter region of *vha-6* (893 bp) was inserted into pPD95.75 using the In-Fusion cloning kit. The GFP N-terminal (GFP1-10) or GFP C-terminal (GFP11) region was then inserted into a linearized *vha-6* promoter-containing pPD95.75 vector using the In-Fusion cloning kit. *daf-16* isoform a (1526 bp), *hsf-1* cDNA (2015 bp), *pfid-2* cDNA (425 bp), *hsp-90* cDNA (2108 bp), or *hsp-70* (C12C8.1; 2069 bp) genomic DNA (gDNA) was then inserted into the linearized *vha-6p::GFPN* vector using the In-Fusion cloning kit. *pfid-6* cDNA (380 bp) was inserted into the linearized *vha-6p::GFPC* vector using the In-Fusion kit. To generate a neuron-specific *pfid-6* expression plasmid, a promoter region of *unc-119* (1964 bp) and genomic *pfid-6* DNA (608 bp) were inserted into pPD95.75 with the In-Fusion cloning kit. For muscle-specific *pfid-6* expression, an *hlh-1* promoter (2000 bp), genomic *pfid-6* DNA (608 bp), and mCherry (711 bp) sequences were cloned into pPD95.75 using the In-Fusion cloning kit. For *rpl-28p::lifeact::eGFP*, the *rpl-28* promoter was inserted into pPD95.75, and the *GFP* region of pPD95.75 was substituted by *lifeact::eGFP* obtained from PCR amplification using PTK92-*lifeact*-GFP as a template (Addgene). The *F35H10.6/UXT* RNAi construct was made by the insertion of the *F35H10.6/UXT* (596 bp)-coding sequence from ATG into L4440 using the In-Fusion cloning kit. For generation of transgenic animals, 25 µg/µL DNA construct was injected into the gonad of day 1 adult worms with 75 µg/µL coinjection marker, *odr-1p::xfp*, *unc-122p::xfp*, or pRF4 [*rol-6(su1006)*], except for 6.25 µg/µL *rpl-28p::lifeact::eGFP*.

Yeast two-hybrid screening

Yeast two-hybrid screen was performed as described previously with modifications (Kim et al. 2008) by Panbionet (<http://panbionet.com>). The full length of *pfid-6* cDNA (381 bp) was amplified by PCR. The PCR product was cloned into pGBKL vector, which contains the DNA-binding domain (BD) of GAL4. Screening was conducted with cotransformation of GAL4 DNA-BD-fused PFD-6 and a *C. elegans* cDNA activation domain (AD) library in pPC86 vector. Yeast strain PBN204 (Panbionet) was used, and two different reporter genes—URA3 and ADE2—were used as selection markers. Yeast transformants of the *pfid-6* bait and the *C. elegans* cDNA AD library were spread on selection medium lacking leucine and tryptophan (SD–LW). From 8.2×10^6

colonies, 640 colonies grew on medium lacking leucine, tryptophan, and uracil (SD–LWU). Among them, 147 colonies grew on medium lacking leucine, tryptophan, and adenine (SD–LWA). In order to confirm the interaction, the prey genes of 147 candidates were amplified by PCR or transformation of *E. coli* and retransformed into PBN204 with the PFD-6 bait plasmid. Subsequently, 75 clones were confirmed as real positives.

Construction of plasmids for a yeast two-hybrid assay

cDNAs used in this assay were amplified by PCR using a *C. elegans* cDNA library as a template. The 381-bp region of *pfid-6* and 1593 bp of *daf-16* were amplified and cloned into pGBKT7 or pGADT7 (Addgene) by using the In-Fusion cloning system. Transformant yeasts containing GAL-4 DNA-BD-fused PFD-6 and GAL-4 AD-fused DAF-16 were spread on a master plate (SD–LW). After 3 d, colonies were replicated on selective plates (SD–LWA). Four micrograms of DNA were used. Because Gal4 AD-DAF-16 transformed yeast displayed growth defects, the interaction between Gal4 DNA-BD-DAF-16 and AD-PFD-6 was tested for yeast two-hybrid assay.

Imaging transgenic animals

Microscope imaging was performed as described previously with some modifications (Artan et al. 2016). Animals were prepared at mixed stages and transferred onto a 2% agarose pad on glass with 100 mM sodium azide (Daejung). The images of worms were captured by using an AxioCam HRc CCD digital camera (Zeiss Corporation) with a Zeiss Axio Scope A1 compound microscope (Zeiss Corporation). ImageJ (Java-based image processing program; National Institutes of Health, <http://rsbweb.nih.gov/ij/>) was used for the quantification of fluorescent images. The double-RNAi method for Supplemental Figure S5 was used as described previously with some modifications (Rea et al. 2007). HT115 bacteria for control RNAi, *daf-2* RNAi, *hsf-1* RNAi, or *daf-16* RNAi were cultured in LB medium to optical density (OD₅₉₀) 0.9~1.0. Bacteria expressing *daf-2* dsRNA were mixed at a 1:1 ratio with those expressing control dsRNA, *hsf-1* dsRNA, or *daf-16* dsRNA. Mixed or single dsRNA-expressing bacteria (100 µL) were seeded onto 50 µg/mL ampicillin-containing NGM plates. IPTG (1 mM) was added and incubated overnight at room temperature. Transgenic worms for the fluorescent imaging (Supplemental Fig. S8) were prepared with the same method that was used for the RNAi-mediated life span assays. The confocal images in Figure 3A were acquired by using Nikon confocal microscopy (A1Rsi) at the Brain Research Core Facilities of the Korea Brain Research Institute (KBRI).

mRNA extraction and quantitative qRT-PCR assays

mRNA extraction and qRT-PCR were performed as described previously with some modifications (Hwang et al. 2015). N2, *pfid-6(gk493446)*, *daf-2(e1370)*, and *pfid-6(gk493446); daf-2(e1370)* animals were grown on control RNAi-seeded plates until L4 stage and transferred to control RNAi-seeded or *pfid-6* RNAi-seeded plates containing FUDR. Synchronized day 1 adult worms were harvested using M9 buffer after two washes and frozen in liquid nitrogen. Total RNA was extracted using RNAiso plus (Takara), and cDNA libraries were obtained from reverse transcription (Promega) with random primers. qPCR was conducted using a StepOne real-time PCR system (Applied Biosystems) with SYBR Green master mix (Applied Biosystems). Two technical repeats were averaged for one biological sample and analyzed

for all data sets. The transcripts of samples were normalized to *ama-1* mRNA levels.

Sequences of primers used for qRT-PCR

The primers used for qRT-PCR were *ama-1*-Forward (TGGAAC TCTGGAGTCACACC), *ama-1*-Reverse (CATCCTCCTTCATT GAACGG), *pfid-6*-Forward (GAAACGGCTTGAGTTTATCG), *pfid-6*-Reverse (CACCTGTCTCGCTGTTCCG), *pfid-1*-Forward (CTGAAGCTAAACAGTCAAAAGAG), *pfid-1*-Reverse (CGTC TACTTTGAATCAGCTCG), *pfid-2*-Forward (GCAGAATAATA TCGCTAATTTGAC), *pfid-2*-Reverse (GTGAGCAGACGAATG TTGTGTG), *pfid-3*-Forward (CAGAAGCCCGAGAAGGTATC), *pfid-3*-Reverse (CTCGTCGACCACTTTTGAAC), *pfid-4*-Forward (CAAAACAAACCCAGTAAAAGCAG), *pfid-4*-Reverse (GCAGCTTCCAATTCTGCATGG), *pfid-5*-Forward (GGAACA TATTACAAAGCAGGTTG), *pfid-5*-Reverse (GTGGCGAGTTG AGACTGAAC), *ruvb-1*-Forward (CCTAAAATAAGAAGTGA TATCCATTC), *ruvb-1*-Reverse (GGAATGAGATAAGTTGTA GAAGAAG), *ruvb-2*-Forward (CAAGTTGATAAGGCTTCTG GACG), *ruvb-2*-Reverse (GTCTCTTTTGAATTTCTCCATCT GG), *F35H10.6/UXT*-Forward (GAGTTTTTATGAATATTGA AGTGAGAG), *F35H10.6/UXT*-Reverse (CTGTCCATCTTCCG ATCAGATG), *uri-1*-Forward (GTTGGAAGCTCCTATTTTGG CG), *uri-1*-Reverse (CTGTATGAATGCTGCATTCTCCG), *F12 A10.1*-Forward (ATGAATTCGCAAACCTACTTTTGG), *F12A 10.1*-Reverse (GGCGTTTTTCACGAACATTTATATTG), *F26 G1.2*-Forward (GATGTGGTTTCTTATACAGAAGACGG), *F26 G1.2*-Reverse (GTGGAGCTCCAGTAATCGTACGG), *C06G 3.3*-Forward (GAAACCTGATCAAAGCCAACACG), *C06G3.3*-Reverse (GAAATGGTATATTAATGGAAGCCAC), *C02F12.5*-Forward (CTTGAATCTTGTAACCCAGCTGAC), *C02F12.5*-Reverse (CTCCCATCACACACACCGTTC), *grd-4*-Forward (GCT GTAGCTGTTCAAGGCC), *grd-4*-Reverse (CATCACCAAAAT AGCTCAACGTTTC), *sod-3*-Forward (CTATCTTCTGGACCAA CTTGG), *sod-3*-Reverse (GCAAGTTATCCAGGGAACCG), *D1 086.1*-Forward (GGTATTTTCAGTGACAAATCAAGAATG), *D1 086.1*-Reverse (CGTGGAACTCTCCACACGCC), *nlp-35*-Forward (CAAGTTCTCGCACCAAGATTCCGA), *nlp-35*-Reverse (GGT TTTGGAGCAAATGTTGGAGG), *M02D8.6*-Forward (GCTTC TGTATATATCAGCAAG), *M02D8.6*-Reverse (CAAGCATCATC TCGAAGAAC), *nas-5*-Forward (GTGGTCAATGGACGGGG TC), and *nas-5*-Reverse (CTTTCCCGTATTGGTGAACATA AAC).

mRNA library preparation and mRNA-seq

mRNA extraction for RNA-seq was performed as described previously with some modifications (Lee et al. 2015). N2, *pfid-6* (*gk493446*), *daf-2(e1370)*, and *pfid-6(gk493446); daf-2(e1370)* were synchronized until the worms reached L4 stage on control RNAi plates and transferred to control or *pfid-6* RNAi plates containing FUDR. After 1 d, worms were harvested with M9, washed to remove bacteria, and frozen in liquid nitrogen. Total RNAs were isolated from frozen worms using RNAiso plus (Takara) and ethanol precipitation. The paired-end mRNA-seq library was prepared and sequenced by a HiSeq 2500 platform at Chunlab (Chunlab, Inc.). A total of three biological repeats of samples was analyzed for all conditions. RNA-seq experiments for two biological repeats were performed first, and the last set was added later to increase the robustness of the data.

mRNA-seq analysis

RNA-seq data analysis was performed as described previously (Pertea et al. 2016). Reads were aligned to the *C. elegans* genome

cell and analyzed by HISAT2 software (version 2.0.5) (Pertea et al. 2016) with default parameters. Transcript assembly and quantification were performed by using StringTie (version 1.3.3) (Pertea et al. 2015) and Ballgown (version 2.0.0) (Frazee et al. 2015). Differentially expressed genes were analyzed by limma (version 3.24.15) (Phipson et al. 2016) with count values. Batch effects among biological repeats were removed with lmFit, contrasts.fit, and eBayes functions in limma (version 3.24.15) (Law et al. 2014). *P*-values were calculated by using limma (version 3.24.15) (Phipson et al. 2016) with R version 3.1.3 (<http://www.r-project.org>). Scatter plot (Riedel et al. 2013), Venn diagram, and heat map analyses (Ewald et al. 2015) were performed as described previously. Heat maps were generated by clustering expression profiles using Cluster 3.0 (Eisen et al. 1998) and visualized by Java Treeview (Saldanha 2004). Venn diagrams were made by using Venn diagram plotter (<http://omics.pnl.gov/software/venn-diagram-plotter>), and *P*-values of the overlaps between two different sets of genes were calculated by using hypergeometric probability test (http://nemates.org/MA/progs/overlap_stats.html).

GO biological process

GO analysis for *pfid-6*-dependent up-regulated genes in *daf-2(-)* animals was performed by using DAVID (Huang da et al. 2009; <https://david.ncifcrf.gov/home.jsp>). The enriched GO terms were selected by using the criterion of *P*-value < 0.05.

Analysis of DNA motifs

DNA motif analysis was performed as described previously with modifications (Seo et al. 2015). Enriched DNA sites located from 800 bp upstream of *pfid-6*-dependent up-regulated genes and down-regulated genes in *daf-2* mutants were searched by using RSAT (RNA sequence analysis tools; <http://www.rsat.eu>). DNA motifs were predicted by using a default parameter, oligomer lengths of 6 and 7, pseudo-frequency of <0.01, and flanking residue of 3.

Subcellular localization of DAF-16::GFP

Microscope imaging for GFP subcellular localization was performed as described previously with modifications (Seo et al. 2015). Gravid adult worms were transferred to RNAi bacteria-seeded plates and allowed to lay eggs. After 2 or 3 d, day 1 adult progeny were mounted on 2% agar pads with 2 mM levamisole (tetramisole) (Sigma). The number of worms with nuclear-localized DAF-16::GFP in the anterior intestinal cells was counted. ImageJ was used for the quantification of fluorescent images.

Coimmunoprecipitation and Western blot assays using cultured mammalian cells

Coimmunoprecipitation was performed as described previously with some modifications (Seo et al. 2015). HA-tagged HSP-90 and Flag-tagged PFD-6 were expressed using HEK293T cells. The cells were maintained with DMEM (HyClone) supplemented with penicillin/streptomycin (HyClone) and 10% (v/v) FBS (HyClone) in a 5.0% (v/v) CO₂-provided incubator at 37°C. Transfection was performed with Lipofectamine LTX and Plus reagent (Invitrogen). Three micrograms of plasmids expressing Flag-HSP-90 and/or HA-PFD-6 was used for transfection. HEK293T cell lysates were collected using PBS (Welgene; 137 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, 2 mM KH₂PO₄ at pH 7.4) and then centrifuged at 800g for 1 min. The supernatant was removed, and the

pellet was resolved with a lysis buffer (20 mM Tris-HCl at pH 7.4, 10 mM KCl, 10 mM MgCl₂, 2 mM EDTA, 10% glycerol, 1% Triton X-100, 2.5 mM β -glycerophosphate, 1 mM NaF, 1 mM DTT, 1 mM PMSF, protease inhibitor mixture [Biovision]). The cell lysates were sonicated (power output 30 W), and 210 mM NaCl was added. The lysates were incubated for 1 h at 4°C on an end-over-end rotator and subsequently centrifuged at 13,000g for 20 min at 4°C. A total of 2 mg of protein was incubated for 1 h~overnight with Flag antibody (Sigma). Protein G agarose was added (GE Healthcare) and then incubated for 1 h at 4°C on an end-over-end rotator. The mixture was washed three times with a wash buffer (20 mM Tris-HCl at pH 7.4, 500 mM NaCl, 10 mM MgCl₂, 2 mM EDTA, 1% Triton X-100) with centrifugation at 800g for 1 min. The mixture was denatured by using 2× SDS sample buffer (Bio-Rad) at 95°C, and proteins were detected by using standard Western blot assays. Primary antibodies against HA (1:2000; Sigma) and Flag (1:5000; Sigma) were used with anti-rabbit (1:5000; Thermo) and anti-mouse (1:5000; Thermo) secondary antibodies, respectively.

Western blot assays with *C. elegans* samples

Western blot assay using *C. elegans* was performed as described previously (Jeong et al. 2017).

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Prefoldin 6 mediates longevity response from heat shock factor 1 to FOXO in *C. elegans*

Heehwa G. Son, Keunhee Seo, Mihwa Seo, et al.

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