C. elegans Ring Finger Protein RNF-113 Is Involved in Interstrand DNA Crosslink Repair and Interacts with a RAD51C Homolog

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Abstract

The Fanconi anemia (FA) pathway recognizes interstrand DNA crosslinks (ICLs) and contributes to their conversion into double-strand DNA breaks, which can be repaired by homologous recombination. Seven orthologs of the 15 proteins associated with Fanconi anemia are functionally conserved in the model organism C. elegans. Here we report that RNF-113, a ubiquitin ligase, is required for RAD-51 focus formation after inducing ICLs in C. elegans. However, the formation of foci of RPA-1 or FCD-2/FANCD2 in the FA pathway was not affected by depletion of RNF-113. Nevertheless, the RPA-1 foci formed did not disappear with time in the depleted worms, implying serious defects in ICL repair. As a result, RNF-113 depletion increased embryonic lethality after ICL treatment in wild-type worms, but it did not increase the ICL-induced lethality of rfs-1/rad51C mutants. In addition, the persistence of RPA-1 foci was suppressed in doubly-deficient rnf-113/rfs-1 worms, suggesting that there is an epistatic interaction between the two genes. These results lead us to suggest that RNF-113 and RFS-1 interact to promote the displacement of RPA-1 by RAD-51 on single-stranded DNA derived from ICLs.

Introduction

Fanconi anemia (FA) is a rare recessive disease involving bone marrow failure, developmental abnormalities including short stature and rudimentary (or absent) thumbs, and susceptibility to cancers [1,2]. The cells of FA patients have chromosomal defects including breaks, gaps, and rearrangements, and are especially hypersensitive to interstrand DNA crosslinking agents. There are 15 complementation groups of FA and the corresponding proteins have all been identified. The 15 FA-associated proteins include FANC A, B, C, D1, D2, E, F, G, I, J, L, M, O, and P, eight of which (A, B, C, E, F, G, L, and M) form a core complex. FANCM and FANCL of the complex have been shown to have well-defined functions in mammalian cells. On the other hand, it is possible that the amino acid sequences of these core complex components are not conserved to levels recognizable by homology searches. Thus, we have taken a functional approach to identifying candidate FA components.

The FANCD2 homolog, FCD-2, has been reported to be mono-ubiquitinated or associated with a ubiquitinated protein [26,27]. However, the E3 ligase has not been found in C. elegans. A ring finger protein, RNF-113, was shown to interact with FCD-2 in a high throughput yeast two-hybrid assay [36–30]. RNF-113 was predicted to be an E3 ligase and a candidate for the enzyme that ubiquitinates FCD-2 [32]. In this study, we have investigated the function of RNF-113 in response to ICL-induction using proliferating germ cells of C. elegans, and characterized its genetic relationships to FCD-2/FANCD2, RFS-1/RAD51C, and RAD-51.
Results

RNF-113 has a role in preventing DNA crosslink hypersensitivity

In a high throughput yeast two-hybrid assay a C. elegans ring finger protein, RNF-113, interacted with a number of proteins including DAF-16 (FOXO homolog) and FCD-2 (FANC-D2 homolog) [36,37]. The C. elegans protein is most closely related (35% identity in amino acid sequence) to human RNF113A (Figure S1 in File S1), a ring finger protein for which no biological function has been reported. Although C. elegans RNF-113 does not have significant sequence identity with mammalian FANC-L, its physical interaction with FCD-2 suggested a possible role for this prospective ubiquitin ligase in modifying FCD-2. Therefore, we knocked down RNF-113 (see Materials and Methods) and examined the effect on repair of interstrand DNA crosslinks (ICLs). The RNAi-treated worms without exogenously-induced DNA damages produced embryos, only 80(±7) SEM% of which hatched later (Figure 1A), suggesting a role of RNF-113 in normal embryogenesis. We then investigated the response of nf-113 knockdown worms to treatment with the crosslinking agent, TMP (4,5′-8-trimethylpsoralen), followed by exposure to UVA radiation. The photoactivated bifunctional psoralen induces DNA crosslinks, which occur almost exclusively between DNA strands, resulting in a very low level of intrastrand-crosslinks [39]. The ICL-treated L4 worms produced 33(±8)% hatched embryos after nf-113 knockdown, which was much lower than the corresponding value of 83(±2)% in wild-type worms. The yield (33±8%) of nf-113(RNAi) worms after ICL treatment was thus much lower than the yield (63±7%) obtained by simply taking into account the embryonic lethality (17±2%) induced by ICLs in wild-type worms and the death (20±7%) resulting from nf-113 knockdown. The synergism between ICL treatment and nf-113 knockdown leads to the conclusion that RNF-113 plays a critical role in repairing DNA crosslinks.

To see whether RNF-113 interacts genetically with FCD-2, we depleted RNF-113 in the fcd-2(tm1298) mutant. These depleted worms laid only 12 embryos, compared with 241 from the wild type, 201 from fcd-2(tm1298), and 100 from nf-113(RNAi) (Figure S2A in File S1), and the percentage of hatched embryos from the doubly deficient worms (26±12%) was much lower than from the untreated nf-113 (80±7%) worms. In fact, the number of progeny of the doubly deficient worms was too small to measure an effect of ICL on survival (Figures 1A and S2A in File S1). To investigate the reasons for this very small brood size, we examined the germ line, and found that the number of endomitotic (Emo) oocytes in the fcd-2(tm1298);nf-113(RNAi) worms was much higher than in either singly-deficient strains (Figure S2B in File S1). The ‘Emo’ phenotype was previously observed in oocytes with defective ovulation and resulted from uncontrolled DNA replication [40]. The endomitotic oocytes observed in our study contained either conglomerte chromosomes with enormous amount of DNA or greatly increased numbers of bivalent condensed chromosomes (Figure S2B in File S1).

The nuclear level of RNF-113 increases in response to ICLs

To probe the intracellular location of RNF-113, we stained the gonads of wild-type and nf-113(RNAi) worms with RNF-113 antibody (Figure 1B). The protein was present in the cytoplasm and on the periphery of the nuclei of germ cells. After ICL treatment, the amount of protein increased in both nucleus and cytoplasm, the maximum effect being seen between 9 h and 16 h after treatment i.e. the period when the germ cells were enlarged due to cell cycle arrest. The accumulation of RNF-113 decreased over time and had almost disappeared by 24 h in most of the germ cells. This corresponds to the time when cell cycling resumed based on nuclear size (Fig. 1B) and the disappearance of RAD-51 foci (data not shown). The specificity of the antibody was confirmed using RNAi knockdown of RNF-113, which effectively eliminated the anti-RNF-113 signal. The result also confirmed the efficiency of the RNAi treatment.

RNF-113 is required not for FCD-2 focus formation but for RAD-51 focus formation after ICL treatment

Since RNF-113 interacted with FCD-2 in a high throughput yeast two-hybrid assay [36–38], we examined the effect of its depletion on FCD-2 focus formation. FCD-2 (FANC-D2 homolog) appeared as nuclear foci in wild-type germ cells 18 h after ICL, as previously reported, and the FCD-2 foci formed even upon the depletion of RNF-113 (Figure 2) [26,33]. Most of the FCD-2 foci disappeared by 24 h after ICL in both of wild-type and RNF-113 depleted cells (data not shown). In mammalian cells, FANC-D2 is ubiquitinated by FANC-L, and as a result forms nuclear foci [3,5]. Therefore, the fact that FCD-2 focus formation is not affected by nf-113 knockdown suggests that FCD-2 is not a ubiquitination target of RNF-113.

Having observed that FCD-2 focus formation was not affected by RNF-113 knockdown, we examined RAD-51 focus formation in germ cell nuclei [41]. In mammalian cells, RAD51, acting downstream of FANC-D2, is recruited to DSBs resulting from incisions at ICLs and initiates repair of the DSBs via homologous recombination [42,43]. As observed previously [33], RAD-51 formed nuclear foci after ICL treatment in wild-type C. elegans germ cells, the focus signal being most prominent between 9 h and 18 h (Figures 3A and S3A in File S1). However, when RNF-113 was depleted, the number of RAD-51 foci was greatly reduced (from 5.0±0.6 (SEM) foci to 5.0±0.6 foci per nuclear focal plane), and the number did not increase significantly with time after ICL treatment (Figures 3 and S3A in File S1). This indicates that RNF-113 is needed for efficient loading of RAD-51 on DNA sites damaged by interstrand DNA crosslinking.

Because RAD-51 focus formation after ICL treatment was greatly diminished by RNF-113 depletion, we asked if the level of RAD-51 protein or its activation by phosphorylation was affected by RNF-113 depletion (Figure S4A in File S1). C. elegans worm extracts were separated on a 10% SDS-polyacrylamide gel, and RNAi-RAD-51 was probed by western blotting. Two bands of RAD-51 were seen in wild-type worms before and after ICL treatment. The intensity of the upper band increased after the ICL treatment, while that of the lower band decreased. This shift to the upper band is thought to be due to phosphorylation of RAD-51, as mammalian RAD51 is phosphorylated by CHK1 after DNA damage [44]. RNF-113 knockdown did not significantly reduce either the level of RAD-51 protein or affect its gel mobility shift, indicating that RNF-113 is not needed for phosphorylation of RAD-51. As a control, we observed that chk-1 RNAi partly inhibited the shift to the upper band, supporting the conclusion that the upper band is a phosphorylated form of RAD-51. We conclude that RNF-113 regulates focus formation by RAD-51 at ICLs without affecting the amount or the phosphorylation of RAD-51.

RNF-113 is not required for RAD-51 focus formation after DSB formation

In order to examine whether RNF-113 responds specifically to ICLs and not to other types of DNA damage such as DSBs...
we probed RAD-51 focus formation after irradiating worms with γ-rays. RAD-51 foci appeared clearly at 3 and 9 h after γ-irradiation in both wild-type and *rnf-113* RNAi germ cells (Figures 3B and S3B in File S1), suggesting that RNF-113 is required for repair of ICLs, but not for DSBs induced by γ-rays. In agreement with these results, RNF-113 depletion did not increase embryonic lethality after γ-irradiation (Figure S5B in File S1). Nevertheless, RNF-113 protein increased in the germ cells after the irradiation (Figure S5A in File S1), leaving open the possibility that RNF-113 plays a minor role in DSB repair downstream of RAD-51.

**Loss of RFS-1 has an epistatic relationship with RNF-113 deficiency in terms of ICL-sensitivity and RAD-51 focus formation.**

RNF-113 depletion resulted in hypersensitivity to ICL-inducing agents and greatly attenuated the resulting RAD-51 focus formation. To further characterize this effect, we examined RFS-1, a RAD51C homolog, which is required for RAD-51 focus formation.
The fact that the doubly-deficient (*p* values 0.54 and 0.55, respectively). In contrast, embryonic increase embryonic lethality either with or without ICL treatment than for the wild type (83% in the wild type (from 9.5% to 4%) gonads were immuno-stained using FCD-2 polyclonal antibody at 18 h after ICL (TMP/UVA) treatment. The scale bar is 10 μm. doi:10.1371/journal.pone.0060071.g002

**Figure 2.** Focus formation by FCD-2 after ICL treatment is not affected by RNF-113 depletion. The mitotically proliferating regions of wild-type and *rnf-113(RNAi)* gonads were immuno-stained using FCD-2 polyclonal antibody at 18 h after ICL (TMP/UVA) treatment. The scale bar is 10 μm.

formation after ICL treatment but not after ionizing-radiation [28]. For this purpose, we measured embryonic lethality in the deletion mutant *tsf-1(ok1372)* with or without depletion of RNF-113 (Figure 1A). The *tsf-1* mutant showed an embryonic survival of 49±4% in the presence of ICL-inducing agents, much lower than for the wild type (83±2%), in agreement with a previous report [28]. Depletion of RNF-113 in the *tsf-1* mutant did not increase embryonic lethality either with or without ICL treatment (*p* values 0.54 and 0.55, respectively). In contrast, embryonic survival after RNF-113 depletion in the wild-type strain was 80±7% before ICL treatment and 34±8% after the treatment. The fact that the doubly-deficient *tsf-1;rnf-113(RNAi)* strain was not significantly more sensitive to ICLs than the singly defective *rnf-113(RNAi)* strain (Student’s *t* test, *p* value = 0.12) or the *tsf-1* strain (*p* = 0.55), demonstrates that RNF-113 functions in the same pathway as RFS-1. The *tsf-1* mutation also reversed the decreased brood size of *rnf-113(RNAi)* worms seen in the absence of exogenous ICLs (Figure S2A in File S1).

Since the *tsf-1* mutation was epistatic to *rnf-113* knockdown in terms of the survival of embryos derived from germ cells exposed to ICL agents (Figure 1A), we tested whether the two genes also interacted in the same way with respect to RAD-51 focus formation (Figures 3A and 3C). In agreement with the report by Ward et al. [28], the number of RAD-51 foci after ICL treatment was reduced by *tsf-1(ok1372)* mutation to one fourth the level in the wild type (from 9.5±0.5 (SEM) foci to 2.3±0.5 foci per nuclear focal plane). This reduction was two-fold greater than that caused by *rnf-113* knockdown. In the doubly-deficient *tsf-1;rnf-113 (RNAi)* strain, the number of RAD-51 foci did not decrease further (Student’s *t* test, *p* = 0.30). This epistatic interaction between *rnf-113* and *tsf-1* in RAD-51 focus formation agrees well with the data on embryonic survival after ICL treatment (Figure 1A).

ICL-induced RPA-1 foci do not disappear with time in RNF-113-depleted cells

RPA-1, which is the large subunit of RPA in *C. elegans*, activates the checkpoint pathway involving ATL-1 (ATR homolog) and CHK-1 in *C. elegans* [45], and is essential for the FCD-2 focus formation [33]. RPA-1 focus formation reached a maximum by 4 h (usually between 3 h and 6 h) after ICL treatment in wild-type worms, and most of the foci had disappeared by 12 h, probably due to their replacement by RAD-51 (Figure 4). RPA-1 focus formation was normal in RNF-113-depleted cells as measured at 4 h after ICL treatment (Figure 4). This was expected, since nuclear foci of FCD-2, which acts downstream of RPA-1, formed normally (Figure 2). RPA-1 foci even formed normally in *gsf-1(ok1372)* cells, since RFS-1 is only involved in the assembly and disassembly of RAD-51 filaments, not in earlier steps. The rate of dissipation of RPA-1 foci in *tsf-1(ok1372)* cells was also very similar to that in the wild type, although the formation of RAD-51 foci and ensuing homologous recombination were defective in these cells. In contrast to the situation of wild-type and *tsf-1(ok1372)* worms, RPA-1 foci did not disappear with time in the RNF-113-depleted worms, most of them being still present 24 h after ICL treatment. This suggests that the stalled replication forks or single-stranded DNA regions to which RPA-1 binds, persist in the depleted strain due to serious defects in ICL repair. In comparison, the *tsf-1;rnf-113(RNAi)* strain was similar to the *tsf-1* mutant with respect to the dynamics of RPA-1 foci, although a minor fraction of RPA-1 foci remained 24 h after ICL treatment. These results agree well with the data on embryonic survival in Figure 1A, showing that embryonic survival after ICL was not significantly decreased by *rnf-113* depletion in the *tsf-1* mutant background.

RNF-113 has ubiquitin ligase activity in vitro

To understand the molecular function of RNF-113, we measured its enzymatic activity by expressing it in *E. coli* with an N-terminal 6×HIS tag and purifying the tagged protein on a Ni-NTA column. The purified protein was reacted with E2 (UbcH5c) in the presence of E1, ubiquitin (HA-tagged), and ATP. The reaction products were separated on an 8–16% SDS-polyacrylamide gel and analyzed by western blotting using HA antibody. When all the components for ubiquitination, namely HA-ubiquitin, E1, E2, RNF-113, and ATP were present, two ubiquitinated protein bands of about 100 kDa and 89 kDa were formed (Figure 5A). The bands were absent if ATP, E1 or E2 was omitted. To characterize the two ubiquitinated proteins, the reaction products of Figure 5A were separated on another 8–16% SDS-PAGE and probed with HIS antigen. Only the lanes containing RNF-113 exhibited two His-tagged proteins of about 90 kDa and 70 kDa (Figure 5B, which were also observed after protein-staining with Ponceau S (data not shown). Although the reason why two HIS-tagged polypeptides were produced is not clear (GST-tagged RNF-113 used for antibody production was also expressed as two polypeptides), both polypeptides were confirmed to be RNF-113 by MALDI-TOF mass spectrometric analysis (data not shown). To show that the ubiquitinated proteins (Figure 5A) are derived from the two HIS-tagged proteins, we analyzed the ubiquitinated proteins by two-dimensional electrophoresis involving isoelectric focusing followed by SDS-PAGE (Figure S6 File S1). The longer polypeptide having the expected isoelectric point is thought to be the full-length 6×HIS::RNF-113, whereas the shorter one with a higher isoelectric point is probably a cleaved product. The results (Figures 5 and S6 in File S1) are most consistent with the argument that the two ubiquitinated bands (Figure 5A) correspond to mono-ubiquitinated forms of the two 6×HIS::RNF-113 polypeptides (Figure 5B).

**Discussion**

In this study, we have shown that the *C. elegans* ring finger protein, RNF-113, does not affect FCD-2 (FANCd2 ortholog) focus formation, thus excluding its possible role as a ubiquitin ligase for FCD-2 (Figure 2). However, the protein has been shown to be a regulator of RAD-51 focus formation in response to interstrand DNA crosslinks (ICLs) (Figures 3A and 3C). The purified protein had E3 ligase activity, adding mono-ubiquitin to itself in the presence of E1, E2, and ATP (Figure 5B). How this E3
activity is related to its role as a regulator of RAD-51 focus formation after ICL treatment is unclear. One clue to the molecular function of RNF-113 is its epistatic interaction with RFS-1, a RAD51C homolog. RFS-1 is required for the full effects of RNF-113 depletion on embryonic lethality and brood size in untreated worms, and for persistence of RPA-1 foci in the germ cells after ICL treatment (Figures 1A, S2A in File S1, and 4). The mammalian homolog of RFS-1, RAD51C, is thought to play a role at more than one stage of homologous recombination, including RAD51 filament formation and the subsequent Holliday junction formation [15,16,46]. Nevertheless, *C. elegans* RFS-1 is more important for RAD-51 focus formation after ICL formation than after treatment with ionizing radiation [28]. Likewise, the

**Figure 3. RNF-113 depletion attenuates RAD-51 focus formation at ICLs in wild-type worms, but not in rfs-1 mutants.** (A) The mitotically proliferating regions of gonads from wild-type, *rnf-113*(RNAi), *rfs-1(ok1372)*, and *rfs-1(ok1372);rnf-113*(RNAi) worms are shown after staining for RAD-51 at 16 h following TMP/UVA treatment. (B) The immuno-staining in (A) was repeated at 3 h after IR (ionizing radiation, 75 Gy) treatment instead of ICL treatment. The scale bars are 10 μm. (C) A focal plane having a maximum number of RAD-51 foci was chosen for each nucleus of germ cells (n = 150 for wild-type and *rnf-113*(RNAi); n = 30 for *rfs-1(ok1372)* and *rfs-1(ok1372);rnf-113*(RNAi)) 16 h after ICL treatment as in (A), and the average numbers of RAD-51 foci per nuclear focal plane are plotted. The error bars are SEM.

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The role of mammalian RAD51C may be more critical in ICL repair than in DSB repair [1].

In view of the known roles of RFS-1/RAD51C, and also the effects of RNF-113 on the dissipation of ICL-induced RPA-1 foci, RNF-113 appears to replace, probably indirectly, RPA-1 at DNA replication forks stalled by ICLs (Figure 6A). Stalled replication forks are predominantly processed by DNA incision to yield one-ended DSBs, which are then repaired by homologous recombination [47]. We propose that RFS-1 binds to single-stranded DNA (ssDNA) derived from one-ended DSBs by end-resection, like mammalian RAD51C, which forms nuclear foci concomitantly with RAD51 and RPA at DSBs [48]; thereafter RNF-113 ubiquitinates an unknown protein ‘X’ that could be RFS-1 or a subunit of RPA, at which point RPA-1 dissociates from the DNA (Figure 6A). Thereafter, RAD-51 can be loaded onto the ssDNA, and replication fork recovery is initiated via strand invasion during homologous recombination. In the absence of RNF-113 the nuclear protein ‘X’ cannot be ubiquitinated; hence RPA-1 remains associated with the ssDNA, and replication fork recovery is frozen at that step (Figure 6B). Although we have discussed the roles of RNF-113 in relation to DNA damage repair, we noted that RNF-113 is important for embryonic survival in the absence of exogenous DNA insults. Likewise, homozygotes of a deletion allele \textit{mf-113(ok1401)} have been reported in WormBase (http://www.wormbase.org/) to arrest at mid-larval stages. Therefore, RNF-113 appears to be essential for larval growth as well as for embryogenesis.

Figure 4. RNF-113 depletion does not affect the formation of RPA-1 foci after ICL treatment, but greatly retards the dissipation of RPA-1 foci. The mitotically proliferating regions of gonads from wild-type, \textit{mf-113(RNAi)}, \textit{rfs-1(ok1372)}, and \textit{rfs-1(ok1372);mf-113(RNAi)} worms are shown after immuno-staining for RPA-1 at 4 h, 12 h, and 24 h following TMP/UVA treatment. The scale bar is 10 μm.

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Figure 5. RNF-113 has E3 ubiquitin ligase activity in vitro. Recombinant 6xHis::RNF-113 was purified from \textit{E. coli} lysates using Ni-NTA agarose and incubated with E1, HA-ubiquitin, and UbcH5c as E2 enzyme. The reaction products were electrophoresed in two separate 6–18% SDS-polyacrylamide gels and probed with (A) HA and (B) HIS antibodies, respectively.

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the tumorous germ line proliferation of glp-1(ez64) [49]. Knockdown of a number of proteins participating in pre-mRNA splicing induced the same phenotype, supporting a function of RNF-113 in pre-mRNA splicing. Additional support for a role of RNF-113 in pre-mRNA splicing comes from the fact that its homolog CWC24p in S. cerevisiae is involved in splicing of pre-U3 snoRNA, therefore affecting pre-rRNA processing [50]. CWC24p has also been identified as a component of Cef1p complexes involved in pre-mRNA splicing, which also contain PRP19 [51,52]. It is intriguing that S. cerevisiae mutated in PRP19/PSO4 is hypersensitive to DNA damage, as is C. elegans deficient in RNF-113 [53]. Given the role of RNF-113 in promoting tumorous germ cell proliferation like other splicing factors, and the activity of its homolog in S. cerevisiae on pre-U3 snoRNA splicing, it is very possible that RNF-113 participates in pre-rRNA splicing. What we observed after knocking down RNF-113 expression could theoretically have been an indirect effect of inhibiting the transcription of ICL repair genes. However, the embryonic lethality and persistent RPA-1 foci due to rnf-113 depletion were not induced in the absence of RFS-1, thus supporting a direct link between the two proteins.

In summary, our study has identified a novel ubiquitin ligase RNF-113 in cellular response to interstrand DNA crosslinks in C. elegans. The protein promotes RAD-51 focus formation and shows genetic interactions with the RAD51C ortholog, RFS-1. It will be important to identify target proteins ubiquitinated by RNF-113 and to find out whether RFS-1 or a subunit of RPA complex is one of the targets in vivo, to fully elucidate the biological functions of RNF-113.

Materials and Methods

C. elegans Strains

Bristol N2 C. elegans worms were cultured at 20°C on agar containing nematode growth medium seeded with E. coli OP50-1 cells. Nematode mutations used were fbd-2(tm1298) and tsf-1(ok1372). The mutation fbd-2(tm1298) was generated by the National Bioresource Project (Japan) and previously out-crossed 6 times with wild-type N2 males [27]. The tsf-1(ok1372) mutant was obtained from the Caenorhabditis Genetics Center and out-crossed to the wild-type N2 strain 3 times.

RNA interference

Wild-type N2 or mutant worms were fed E. coli strain HT115(DE3) that had been transformed with the L4440 feeding vector (control), or the same vector carrying an appropriate cDNA insert. The full length cDNA of rnf-113 was amplified from a C. elegans cDNA pool using primers: 5'-GGATCCATGGATCTC-TTCCGAAAAC and 5'-AAGCTTTCAATCTTTTTCAGCAT (restriction sites underlined). Knockdown of chk-1 was performed using the feeding vector described in our previous work [54]. Since rnf-113 is co-transcribed at its 3'-end with hpl-2, which is a heterochromatin protein HP1 homolog [55], we checked whether hpl-2 mRNA expression was affected by the knockdown of rnf-113. The level of hpl-2 mRNA was not affected by the RNAi; hence the hypersensitivity of rnf-113(RNAi) worms is not only due to loss of RNF-113, not to any effects on HPL-2 (Figure S4B in File S1). The fact that rnf-113(RNAi) worms were not hypersensitive to γ-rays (Figures 3B and S5B in File S1), unlike a hpl-2 mutant that showed a great hypersensitivity to the radiation [55], also supports that hpl-2 expression was not significantly affected by the knockdown.

Embryonic lethality after ICL treatment

Synchronized L4 worms were transferred to NGM plates that had been seeded with E. coli HT115(DE3) cells in the presence of 1 mM IPTG. The E. coli strain HT115(DE3) contained either the L4440 control vector or the vector with a cDNA insert. When worms reached the late L4 stage at 20°C, they were soaked in 1× PTW containing 25 μg/ml TMP (4,5,8-trimethylpsoralen, Sigma) for 30 min, and irradiated with 200 J/m² of ultraviolet light (365 nm). Ten of the treated worms were individually placed on the nematode growth medium seeded with E. coli OP50-1 cells. Nematode mutations used were fcd-2(tm1298) and hpl-2(ok1372). The mutation fbd-2(tm1298) was generated by the National Bioresource Project (Japan) and previously out-crossed 6 times with wild-type N2 males [27]. The tsf-1(ok1372) mutant was obtained from the Caenorhabditis Genetics Center and out-crossed to the wild-type N2 strain 3 times.
an NGM plate with E. coli cells and allowed to lay eggs for 24 h. Hatched eggs were scored 20 h later to calculate hatching rate. All experiments were done in triplicate.

**Antibody preparation**

The full length cDNA of RNF-113 was amplified from a cDNA pool that had been prepared by reverse transcription of total RNA from wild-type C. elegans. Primers were 5'-GGATCCATGGATCTCTTTCCAGAAAAC and 5'-CTCGAGTCGAATCTTTTTCA-GCATCAGT, and the amplified DNA fragment was inserted between the BamHI and XhoI sites on pGEXKT-1 and transformed into BL21(DE3) cells (Yeasent Biotech). The recombinant protein was overexpressed by incubation with 1 mM IPTG for 4 h at 37°C, and used to generate antibodies in rats. The polyclonal antibody against RNF-113 was isolated from the serum of the immunized rats by affinity chromatography as follows. The GST::RNF-113 protein was blotted onto a nitrocellulose membrane, and the membrane was incubated with the antiserum. The antibody was stripped off the membrane in 100 mM glycine (pH 2.5), and the resulting antibody solution was neutralized and concentrated to be used for immunostaining.

**Immunostaining**

C. elegans gonads were ejected from the body with a scalpel, and fixed in 4% paraformaldehyde followed by 100% methanol. They were then incubated with RAD-51(1:50 dilution), RPA-1(1:1,000 dilution), fixed in 4% paraformaldehyde followed by 100% methanol. They were then incubated with RAD-51(1:50 dilution), RPA-1(1:1,000 dilution), and observed with a fluorescence microscope (DMR HC, Leica).

**Western blot analysis**

About 600 wild-type worms at the L4 or young adult stage were transferred to an NGM plate with E. coli cells and allowed to lay eggs for 24 h. Hatched eggs were scored 20 h later to calculate hatching rate. All experiments were done in triplicate.

**Two-dimensional gel electrophoresis**

The reaction products of an in vitro ubiquitination (50 μl) involving 6× His::RNF-113, HA-ubiquitin, E1, and E2, were concentrated using Vivaspin (Satorius, Germany) in the buffer (9 M Urea, 4% CHAPS). After adding the rehydration buffer (IPG buffer, 5 mM DTT, bromophenol blue), the reaction products were applied to an IPG strip (pH 4–7, 7 cm, GE Healthcare). After 16 h of rehydration at room temperature, the IPG strip was placed into the Ettan IPG phor 3 system (GE Healthcare) to perform isoelectric focusing. The applied voltage was increased gradually from 100 V to 3500 V for 9.6 h. After isoelectric focusing, the strip was equilibrated with the buffer (50 mM Tris-Cl, pH 8.8, 6 M urea, 20% glycerol, 2% SDS, 2.5% acrylamide, 0.34% tributylphosphine) and placed on an 0–16% SDS-PAGE. After second-dimensional gel electrophoresis, proteins were transferred to a nitrocellulose membrane and detected using HIS or HA antibody (Sigma, 1:5000).

**Supporting Information**

**File S1** Figure S1. Comparison of amino acid sequences of Ring finger protein 113 homologs in Homo sapiens and C. elegans. Both proteins have a zinc finger domain (yellow) and a ring finger domain (red), and are identical in 35% of their amino acids. Alignment was carried out with Vector NTI Advance (10.0.1) software. Figure S2. Differential effects of RNF-113 depletion on brood size and oocyte chromosomal abnormalities in wild type, fsl-2, and rfs-1 backgrounds. (A) Knockdown of rnf-113 was performed by feeding RNAi from the young adult stage (P0 generation) of wild-type, fsl-2(tm21), and rfs-1(ok1372) worms. The total numbers of eggs laid during the first 3 days of F1 adults were determined. (B) The gonads of F1 adult worms (n = 20) were dissected and stained with DAPI, to count endomitotic (Emo) oocytes. The error bars are SEM. The scale bar is 10 μm. Figure S3. Effects of RNF-113 depletion on RAD-51 focus formation with time lapse after ICL treatment. (A) The mitotically proliferating regions of gonads from wild-type and rnf-113(RNAi) worms are shown after staining for RAD-51 at 9 and 18 h following TMP/UVA treatment. (B) The immunostaining in (A) was repeated at 3 and 9 h after IR (ionizing radiation, 75 Gy) treatment instead of ICL treatment. The scale bars are 10 μm. Figure S4. Knockdown of m-113 expression does not affect the level of RAD-51 protein or hpl-2 transcripts. (A) Western blot of extracts of wild type, rnf-113(RNAi), and chk-1(RNAi) worms at the adult stage using antibodies to RAD-51 and β-tubulin. The upper band of RAD-51 is thought to be its phosphorylated form, pRAD-51. (B) hpl-2 expression relative to that of γ-tubulin. Mixed stages of wild type and rnf-113(RNAi) worms were harvested 18 h after ICL treatment and total RNA isolated. Briefly, 2 μg of total RNA was used to synthesize a strand of cDNA using oligo(dT) primer and AMV reverse transcriptase (Intron, Korea). The resulting cDNA was amplified using iQ SYBR Green Supermix (Bio-Rad) in a real time PCR instrument (CFX96 Touch, Bio-Rad). cDNA amplification was analyzed with CFX Manager Software. The primers were 5'-GGGAGATTGTTAGAGGGGA and 5'-CTGCTTGGCTTCTCAGTGA for hpl-2, and 5'-AAGATCTTAATGTGTATCCAGGC and 5'-CTGTAACCTCTTGTC-
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Author Contributions

Conceived and designed the experiments: HL HSK. Performed the experiments: HL AA MSP. Analyzed the data: HL HSK. Contributed reagents/materials/analysis tools: HSK. Wrote the paper: HSL AR HSK.


detected using HA antibody, deprobed, and then reprobed with HIS antibody. (PDF)

References


