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Mutations in gld-1, a female germ cell-specific tumor suppressor gene in Caenorhabditis elegans, affect a conserved domain also found in Src-associated protein Sam68

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The gld-1 gene of Caenorhabditis elegans is a germ-line-specific tumor suppressor gene that is essential for oogenesis. We have cloned the gld-1 gene and find that it encodes two proteins that differ by 3 amino acids. The predicted proteins contain a ~170-amino-acid region that we term the GSG domain (GRP33/Sam68/GLD-1), on the basis of significant similarity between GLD-1, GRP33 from shrimp, and the Src-associated protein Sam68 from mouse (also described as GAPap62 from humans). A conserved structural motif called the KH domain is found within the larger GSG domain, suggesting a biochemical function for GLD-1 protein in binding RNA. The importance of the GSG domain to the function of gld-1 in vivo is revealed by mutations that affect 5 different conserved GSG domain residues. These include missense mutations in an absolutely conserved residue of the KH domain that eliminate the tumor suppressor function of gld-1.

[Key Words: C. elegans; germ-line development; tumor suppressor; KH domain; GSG domain]

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An important contribution to our understanding of cancer biology has been the identification of tumor suppressor genes in which loss of gene function is a factor in tumor formation. A number of tumor suppressor genes are involved in the cell cycle (for recent review, see Hartwell and Kastan 1994). These include cell cycle checkpoint genes (e.g., p53) and DNA repair genes (e.g., MSH2). Other tumor suppressor genes, possibly including WT1 (Kreidberg et al. 1993), appear to be tied to development, acting in cell type specification or developmental aspects of cell cycle control.

Two recent reports describe genetic studies of gld-1, a tumor suppressor gene that is required for normal oocyte development in the nematode Caenorhabditis elegans (Francis et al. 1995a,b). In wild-type C. elegans, diploid animals with a single X chromosome develop as males that make only sperm, whereas diploids with two X chromosomes are hermaphrodites that produce some sperm before switching to the production of oocytes (Fig. 1). gld-1 has no essential function in the male germ line or soma but is absolutely required for oogenesis in XX hermaphrodites (Francis et al. 1995a). In gld-1(null) hermaphrodites, germ cells that should develop as oocytes enter the meiotic pathway normally and progress as far as the pachytene stage of meiotic prophase. However, these cells then exit meiotic prophase, re-enter a mitotic cell cycle, and proliferate ectopically to produce a germline tumor. Neither the somatic sexual environment nor the chromosomal sex appear important for generation of the hermaphrodite tumor. Instead, tumor formation requires only that the germ-line sex determination pathway be set in the female mode that normally leads to oogenesis (Francis et al. 1995b). On the basis of these properties, gld-1 can be regarded as a tumor suppressor locus that acts to direct oocyte differentiation and meiotic prophase progression (Fig. 1, function 1).

Previous studies have indicated that in addition to its essential role in oogenesis, gld-1 is likely to participate in at least two other aspects of germline development. gld-1 appears to act to regulate mitotic proliferation negatively among premeiotic germ-line stem cells (Fig. 1, function 2), as a gld-1(null) allele can partially suppress the premeiotic proliferation defects conferred by certain alleles of the glp-1 gene (Francis et al. 1995b). In contrast to its role in oogenesis, the activity of gld-1 in premeiotic germ cells is nonessential, possibly because of genetic redundancy, and functions in both the male and hermaphrodite germ lines. The finding that gld-1 acts to regulate premeiotic proliferation negatively suggests...

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Figure 1. gld-1 (+) functions in three aspects of C. elegans germ-line development. Diagrammed is one gonad arm from a wild-type adult hermaphrodite. Germ cell nuclei with characteristic chromosomal morphology are shown. Male germ-line development is complete by the adult mode; (2) a nonessential and sex nonspecific function in negatively regulating proliferation in premeiotic germ cells; and (3) a physical location of gld-1 to a five-cosmid interval (flanked by F29D11 and ZK1014; Fig. 2B).

Genomic Southern blot analyses revealed that two gld-1 alleles are correlated with restriction fragment length polymorphisms (RFLPs). We found an insertion of 2.3 kb associated with gld-1(q343), and a small deletion associated with the psoralen-induced allele, gld-1(q485). Both RFLPs are detected by a 6.8-kb subclone (pAJ37) of cosmid C29D7 (Fig. 2C).

Gene structure

We sequenced most of pAJ37 [data not shown]. The portion of this genomic sequence corresponding to gld-1 cDNAs [see below] is shown in Figure 3. All 32 gld-1 mutations [Fig. 6, below] affect the gene as defined in Figure 3, providing strong evidence that this is the gld-1 gene.

To determine the intron/exon structure of gld-1, we isolated cDNA clones from a mixed-stage cDNA library [see Materials and methods]. One clone, judged to be nearly full length [pLAJ1], was sequenced in its entirety [Fig. 3]. The intron/exon structure of gld-1 is summarized in Figure 2D. The cDNA contains a single open reading frame of 1392 nucleotides. A 5'-untranslated region (UTR) of 16 nts is trans-spliced to SL1 (see below). The 3' UTR is 812 nts and ends with a stretch of poly(A) 10 nts downstream of the consensus polyadenylation signal AATAAA. To determine the precise 5' end of the gld-1 transcript, we isolated additional cDNA clones by 5' rapid amplification of cDNA ends (RACE)-PCR [Frohman et al. 1988]. Sequence of these clones [see Materials and methods] verified that the gld-1 transcript is trans-spliced to SL1 [Krause and Hirsh 1987].

Northern blot analysis detected a single band of 2.5 kb in both mixed-stage poly(A)-selected RNA and total RNA from various developmental stages and germ-line mutants [described below; A. Jones and T. Schedl, unpubl.]. This transcript size is consistent with that of the cDNA pLAJ1 (2.3 kb). However, when sequencing 5' RACE-PCR products, we discovered evidence for an al-

Results

Physical mapping of the gld-1 locus

gld-1 maps genetically to chromosome I between unc-13 and lin-10 [Fig. 2A; Francis et al. 1995a]. This region is represented entirely by an ordered set of yeast artificial chromosome (YAC) and cosmid clones [Coulson et al. 1988]. Through PCR deletion mapping [Barstead et al. 1991; see Materials and methods], we delimited the

that the role of gld-1 in directing oocyte development may also involve negative regulation of mitotic cell cycle factors.

The third function of gld-1 is promotion of the male germ cell fate in the hermaphrodite germ line (Fig. 1, function 3). Because a reduced level of spermatogenesis is observed in XX hermaphrodites with only one functional gld-1 allele, the role of the gene in promoting spermatogenesis appears to be haploinsufficient. gld-1 is not required for sperm formation in either sex, however, and its role in promoting hermaphrodite spermatogenesis may be limited to assisting the set of genes [fog-1, fog-3, and the fem genes] that act to specify the male fate [Francis et al. 1995b]. Several classes of gld-1 gain-of-function [gf] alleles have been characterized that cause transformations in germ-line sexual fate by poisoning the germ-line sex determination process. Two classes of gf alleles feminize the XX [hermaphrodite] germ line dominantly (causing all germ cells to develop as oocytes); a third class of gf mutations feminizes both the XX and XO (male) germ lines; and a fourth class masculinizes the XX germ line (causing germ cells to develop inappropriately as sperm instead of oocytes). Thus, gld-1 can be mutated in different ways to cause gf disruptions in the process that specifies germ-line sexual fates.

The genetic properties of gld-1, including its multiple germ-line functions and classes of mutant alleles, suggest that the gene's regulation and function are likely to be complex. This report presents a molecular study of gld-1 that includes sequence analysis of the wild-type gene and 32 mutants.

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Figure 2. A summary of the mapping of the *gld-1* locus. (A) A genetic map showing the position of *gld-1* in relation to other genetic markers on chromosome I (adapted from Francis et al. 1995a). Deficiencies (ozDf5, nDf24, and nDf25) are shown with approximate end points. (B) A contiguous physical map showing the cosmids from the physical mapping project (Coulsion et al. 1988) that fall within the two PCR assays (opposing arrows) used to delimit the region containing *gld-1* (see Materials and methods). (C) A restriction map showing the position of restriction sites as well as the two RFLPs associated with *gld-1* alleles, as determined from Southern blotting and subcloning. (X)XhoI; (H)HindIII; (E)EcoRI; (P)PstI. q485 (●) is a small deletion; q343 is a 2.3-kb insertion. The double bar indicates the subclone from cosmid C29D7 (pAJ37) that was sequenced. (D) A composite summary of the gene structure as determined from cDNA sequencing. Coding regions are shown as rectangles. The gene structure represents the two transcripts as described in the text, with the alternate splice on exon 6 occurring as shown. *Trans*-splicing occurs at the 5' end of the transcript to the 22-nucleotide leader SL1 (see text). Putative initiator methionine codon (ATG), stop codon (TAA), polyadenylation signal (AATAAA), and 3'-untranslated region (3' UTR) occur as shown. Regions of homology as described in the text are shown as shaded regions superimposed on the gene structure. The GSG domain (see text) is hatched and includes the single KH motif in exon 5. (●) The CGA region (light shading; see Fig. 4) is in exon 7 as shown.

Conserved domains in the GLD-1 predicted protein

Conceptual translation of the cDNA pLAJ1 yields a 463-amino-acid protein. BLAST data base searches (Altschul et al. 1990) revealed significant sequence identity with several proteins. These proteins are shown in alignment in Figure 4A, and diagrammatically in Figure 4B. The most extensive similarity is found with GTPase-activating protein (GAP)-associated p62 (GAPap62) from humans (Wong et al. 1992), and glycine-rich protein [GRP33] from brine shrimp [Cruz-Alvarez and Pellicer 1987]. The sequence reported for GAPap62 is highly related to a 68-kD protein from mouse that is associated with Src during mitosis, and we refer to this protein as Sam68 (for Src-associated in mitosis 68 kD; see Courtneidge and Fumagalli 1994). The similarity between Sam68/GAPap62 and GRP33 has been reported previously (Wong et al. 1992). However, this original alignment is extended by >50 amino acids in the amino-terminal direction when GLD-1, Sam68, and GRP33 are aligned [Fig. 4A]. We term the ~170-amino-acid region of similarity shared by these three proteins a GSG domain [for GRP33, Sam68, and GLD-1]. Overall, the GSG domains of GLD-1 and Sam68 are 34% identical and 61% similar.

Within the GSG domain is a smaller motif proposed to be involved in RNA binding, termed the KH domain [Siomi et al. 1993a]. Both GRP33 and Sam68 have been identified as containing a KH motif [Gibson et al. 1993; Siomi et al. 1993b]. Other genes containing this motif include *FMR-1*, the gene responsible for fragile X syn-
encompassing the locus. The genomic nucleotide sequence below. Splice sites (inverted L's) exon 6 is underlined with dashes, and the al. 1993a). From the analysis of >40 potential KH motifs, a-helices in the order B-a-a-B-6. Shown in alignment in Figure 5 is the central a-a-B prediction of the KH motifs from humans (Verkerk et al. 1991), and hnRNP K, a disease in males (for _Carboxy-terminal GSG domain-associated) region. We call this short sequence the (for _Carboxy-terminal GSG domain-associated) region. We call this short sequence the

Two other proteins show significant similarity to GLD-1, Sam68, and GRP33 (see Fig. 4): A predicted protein from C. elegans, BO280.11 [Wilson et al. 1994], and ZFM1 from humans [Toda et al. 1994]. We confirmed that a gene is transcribed from the BO280.11 region by isolating and sequencing cDNA clones (see Materials and methods). Because this gene has a different intron/exon structure than predicted for BO280.11, we refer to the protein encoded by our cDNAs as BO280.1lb. Similarity between BO280.1lb and ZFM1 with the other GSG proteins does not extend as far amino-terminally, but both of these proteins contain a small region of similarity to GLD-1 that is just carboxy-terminal to the GSG domain (see Fig. 4). We call this short sequence the CAG (for Carboxy-terminal GSG domain-associated) region.

To identify conserved portions of the gld-1 transcripts and coding region, we cloned and sequenced cDNAs that include part of the coding region of the gld-1 gene from the male/female nematode species Caenorhabditis remanei [data not shown; see Materials and methods]. The gld-1 transcripts from both C. elegans and C. remanei contain identical trans-spliced SL1 leaders, and the same length [although not identical] 5' UTRs. The partial coding region [corresponding to amino acids 1-370 in C. elegans] showed 88% identity, with 99% similarity within the GSG domain and CGA region. All of the amino acids mutated in the gld-1 missense alleles described below are conserved in C. remanei.

Molecular analysis of gld-1 mutations that affect oogenesis

The gld-1 gene performs an essential function in directing oogenesis and a nonessential role in promoting spermatogenesis in the hermaphrodite [Francis et al. 1995a]. Thirty-two gld-1 mutations have been placed into six classes (A-F) on the basis of the nature of germ-line phenotypes observed in homozygotes [Francis et al. 1995a]. We determined the molecular lesions in each gld-1 allele [Table 1; Fig. 6; see Materials and methods]. Class A gld-1 alleles are null or strong loss-of-function mutations, which abolish the major function of gld-1. In XX animals homozygous for class A alleles, presumptive female germ cells never undergo oogenesis but, instead, form a tumor. As expected, many of the class A alleles
contain lesions that are likely to eliminate gene function. q485, the canonical genetic null mutation (Francis et al. 1995a), contains an 83-bp deletion that shifts the reading frame near the amino terminus of the predicted protein. In addition to another small deletion (oz127), other class A alleles contain premature stop codons (q365, q268, q495, and q93oz49), a splice site mutation (q93oz53), and missense mutations (qz89, q93oz55, q361, and oz17oz47). Of particular interest is that three of the class A missense mutations (qz89, q93oz55, and q361) affect a single residue [Gly-227 Ser or Asp] in the gld-1 KH motif. This glycine is absolutely conserved in the alignment of KH motifs [see Fig. 5].

The class B, E, and F gld-1 alleles are partial reduction-of-function mutations that block oogenesis in either early or late stages. Class B alleles show a recessive phenotype in which presumptive female germ cells arrest in pachytene of meiosis I. Most class B alleles are intragenic revertants of the class C allele q93 [see below] and contain both the q93 mutation and an additional change that is either missense [q93oz56, q93oz12, and q93oz45] or affects a splice site [q93oz52]. oz116 is the only class B allele that was not obtained as an intragenic revertant. This mutation changes the 3' splice acceptor site in the last intron, from cagAGC to caaAGC. RT–PCR analysis revealed that the adjacent AG is used as a splice acceptor at an appreciable level in oz116 RNA [see Materials and methods], resulting in a frameshift predicted to produce a carboxy-truncated protein.

One class B allele [q93oz50] has a nonsense change in codon 13 of the predicted protein. Although this might be expected to be a molecular null mutation, q93oz50 does retain residual gene activity [Francis et al. 1995a]. Two hypotheses to explain the residual q93oz50 gene activity are exon skipping during splicing [Fisher et al. 1993] and translational initiation at a downstream AUG.

The phenotypic classes E and F are each represented by a single allele. Both alleles produce abnormal oocytes and behave as partial loss-of-function mutations with respect to oogenesis [Francis et al. 1995a]. The class E
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Figure 5. An amino acid alignment of the KH domain. The amino acid sequences of several KH domains are shown. The alignment is based on several descriptions of the content of the KH domain that have placed >10 different proteins ranging from bacteria to higher eukaryotes into the KH domain family (cf. Gibson et al. 1993). The α-β architectural predictions are delimited by lines above the alignment and are from Gibson et al. (1993). Shown in alignment are the KH domains from the human proteins hnRNP K [(1,2, and 3); Siomi et al. 1993a], and FMR-1 [(1 and 2); Verkerk et al. 1991], a yeast protein HX [(1, 2, 3, and 4); Delahodde et al. 1986], GRP33 (Cruz-Alvarez and Pellicer 1987), Sam68/GAPap62 [Wong et al. 1992], ZFM1 (Toda et al. 1994), BO280.1lb (see Materials and methods), and GLD-1 (Fig. 3). Conserved hydrophobic residues are boxed, with the absolutely conserved glycine residues in black. The glycine residue circled in the GLD-1 protein is missense in three gld-I tumorous mutations (see text).

Figure 6. Molecular lesions in 32 alleles of gld-1. A summary of the position of each mutation in relation to the structure of the gene (see legend to Fig. 2D) by phenotypic class (see Table 1 and text). Blocked lines (for q127, q485, q266, and q110) indicate deletions, opposite pointing arrows indicate insertions (for q343, insertion not drawn to scale). For the intragenic revertants (e.g., q93oz50), only the new mutation is shown.

Molecular analysis of gld-1 mutations that affect hermaphrodite spermatogenesis

Aside from its essential role in directing oogenesis, gld-1 has a nonessential role in germ-line sex determination, promoting spermatogenesis in hermaphrodites. Class D gld-1 alleles form normal oocytes. However, these two alleles eliminate hermaphrodite spermatogenesis [a feminization of the germ-line (Fog) phenotype] as the result of a gf defect. In addition, class D alleles show a novel gld-1 defect; they feminize the XO male germ line, caus-
Molecular analysis of gld-1

Table 1. Molecular lesions of gld-1 alleles

<table>
<thead>
<tr>
<th>Allele</th>
<th>Phenotypic class</th>
<th>Nucleotide change</th>
<th>Predicted result of molecular lesion</th>
</tr>
</thead>
<tbody>
<tr>
<td>q485</td>
<td>A1 (tum)</td>
<td>deletion of 481–563.</td>
<td>shift in reading frame, multiple stop codons</td>
</tr>
<tr>
<td>oz127</td>
<td>A2 (tum)</td>
<td>deletion of 261–270.</td>
<td>shift in reading frame, multiple stop codons</td>
</tr>
<tr>
<td>q93oz53</td>
<td>A2 (tum)</td>
<td>g 654 a</td>
<td>3’ splice acceptor changed, exon 3</td>
</tr>
<tr>
<td>q268, q495</td>
<td>A2 (tum)</td>
<td>g 1202 a</td>
<td>Gly-227 Ser</td>
</tr>
<tr>
<td>q93oz49</td>
<td>A2 (tum)</td>
<td>c 1275 t</td>
<td>Gly-227 Asp</td>
</tr>
<tr>
<td>oz17oz47</td>
<td>A2 (tum)</td>
<td>g 1374 a</td>
<td>Gln-238 Stop (uag)</td>
</tr>
<tr>
<td>q3oz50</td>
<td>B (undiff.)</td>
<td>c 206 t</td>
<td>Trp-267 Stop (uag)</td>
</tr>
<tr>
<td>q3oz56</td>
<td>B (undiff.)</td>
<td>g 907 a</td>
<td>Gln-292 Stop (uag)</td>
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<tr>
<td>q3oz12</td>
<td>B (undiff.)</td>
<td>c 958 t</td>
<td>Asp-310 Asn</td>
</tr>
<tr>
<td>q3oz52</td>
<td>B (undiff.)</td>
<td>g 1007 a</td>
<td>Gly-200 Glu</td>
</tr>
<tr>
<td>q3oz45</td>
<td>B (undiff.)</td>
<td>g 1281 a</td>
<td>Pro-217 Leu</td>
</tr>
<tr>
<td>oz116</td>
<td>B (undiff.)</td>
<td>g 1617 a</td>
<td>Ala-294 Thr</td>
</tr>
<tr>
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<tr>
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<td>C2 (Mog)</td>
<td>g 1101 a</td>
<td>Gly-250 Arg</td>
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<tr>
<td>oz10</td>
<td>C3 (Mog)</td>
<td>1. c 1449 t</td>
<td>1. Pro-335 Ser</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2. deletion of</td>
<td>2. deletion within the 3’ UTR</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1911–2426</td>
<td></td>
</tr>
<tr>
<td>q126, oz142</td>
<td>D (Fog)</td>
<td>g 1369 a</td>
<td>Gly-308 Glu</td>
</tr>
<tr>
<td>q266</td>
<td>E [Fog/abn. oocyte]</td>
<td>deletion of 1412–1438</td>
<td>amino acids 322–331 deleted, Ile-323 Met</td>
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<tr>
<td>q343</td>
<td>F [abn. oocyte]</td>
<td>insertion after 2164</td>
<td>insertion of Tc2 in the 3’ UTR</td>
</tr>
</tbody>
</table>

Molecular lesions were determined as described in the text and Materials and methods. The numbering of the altered nucleotides is given in Fig. 3. Amino acids affected are numbered according to the predicted protein from the major transcript, represented by the cDNA pLA1 [see text]. Phenotypic class designations are defined in Francis et al. [1995a]. Both class A1 and A2 alleles have a recessive tumorous germ-line phenotype (tum), where presumptive female germ cells exit pachytene and return to mitotic proliferation. Class A1 is null for all aspects of gld-1 function, whereas A2 alleles display gf poisoning activity that results in feminization of the homomorphodite germ line (see Discussion). Class B alleles show a recessive phenotype in which presumptive female germ cells do not differentiate [undiff.] and are arrested in pachytene. Class C alleles all result in a recessive masculinization of the germ-line (Mog) phenotype, homozygous homomorphodites produce a vast excess of sperm and either never switch to oogenesis (C1 and C2) or eventually switch in older adults (C3). Class C1 alleles show a stronger dominant Mog phenotype than class C2 alleles, and class C2 alleles display an undifferentiated phenotype at 15°C similar to class B. The Mog phenotype for all class C alleles is the result of gf poisoning activity. In addition, class C1 and C2 alleles have partial loss-of-function defects in directing oocyte development. Class D alleles show a feminization of the germ-line (Fog) phenotype in both homomorphodites and males; germ cells that would normally develop as sperm instead develop as oocytes. The class E allele shows a recessive abnormal oocyte [abn. oocyte] phenotype, where female germ cells fail to complete oocyte differentiation but undergo meiotic prophase development normally. In addition, the class E allele has a homomorphodite-specific Fog phenotype. For both classes D and E, the Fog phenotypes are the result of gf defects. The class F allele also displays an abnormal oocyte phenotype but does not have a Fog phenotype. Some class A2 and B alleles are intragenic revertants of the class C1 Mog mutations [e.g., q93oz50], both the original and a new mutation were found, but only the new mutation is presented. Missense alleles of gld-1 that affect residues conserved among Sam68, GRP33, ZFM1, and GLD-1 (see Fig. 4), are shown with the wild-type amino acid in boldface type.

ing males to make oocytes [Francis et al. 1995a]. The two class D mutations (q126 and oz142) contain identical nucleotide changes resulting in Gly-308 Glu. This mutation alters one of the few amino acids not conserved between GLD-1 and BO280.11b in the CGA region (see Fig. 4A). The class E allele q266 also alters amino acids in the CGA region [see above] and, like many of the gld-1 alleles, has a gf effect on homomorphodite spermatogenesis that results in a Fog phenotype. Unlike the class D alleles, however, q266 does not affect XO males overtly.

In contrast, class C alleles are rare gf mutations of gld-1 that increase homomorphodite sperm production [a masculinization of the germ-line (Mog) phenotype]. On the basis of distinct genetic and phenotypic properties [Francis et al. 1995a] the 11 gld-1(Mog) alleles have been divided into three subclasses (C1, C2, and C3). Homozygous C1 and C2 mutants make excess sperm and fail to switch to oogenesis. Genetic tests indicate that these alleles, while having a gf defect for male sex determination (in the homomorphodite), are partial loss of function for the major role of gld-1 in directing oogenesis. All class C1 mutations (q93, q62, oz17, oz34, and oz35) result in Gly-248 Arg, and all class C2 mutations (oz30, oz16, oz29, oz33, and oz70) result in Gly-250 Arg. Thus, missense mutations in two absolutely conserved glycine residues in the GSG domain (see Fig. 4A) are responsible for the fully penetrant Mog phenotype [all of the Mog alleles were rare independent isolates that arose at a fre-
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Figure 7. gld-I RNA expression in the germ line. A Northern blot probed with the cDNA clone pLAJ1 is shown. The first four lanes contain ~5 µg of total RNA from predominantly young adult-stage animals, with the animals in the first three lanes grown at 25°C (restrictive temperature for these mutations). gld-I RNA is easily detectable in total RNA on overnight exposure and is presumably a relatively abundant RNA. The last lane contains ~0.5 µg of poly(A)+-selected RNA from a mixed developmental stage population of wild-type animals (N2). An act-l-specific probe was used as a control for loading [a gift from L. Schriefer, Washington University, St. Louis, MO).

Figure 8. Analysis of the relative abundance of the two alternatively spliced products. The radioactive products of RT–PCR are shown (see text; Materials and methods), separated on a denaturing polyacrylamide gel and autoradiographed. Three PCR cycle points within the linear range of amplification were taken for each RNA sample tested, and quantitation of each band was performed. The ratio of the smaller product to the larger product calculated for each PCR cycle point is shown. The plasmid control reactions were included to demonstrate that known ratios of input DNA gave an equivalent output ratio in the assay. The control reactions were for 20 cycles of amplification, starting with 1 amole of DNA containing different ratios of two cDNA plasmid clones representing the two different splice products.

Tissue specificity of gld-I RNA

We analyzed gld-I RNA levels in wild type [Fig. 7]. A Northern blot of mixed-stage poly(A)+ RNA from wild type (N2) shows a single band of ~2.5 kb when probed with the cDNA clone pLAJ1; no additional bands are seen on extended exposures. In Northern blots of RNA from developmentally staged populations, a 2.5-kb band is also visible in the L3 larval stage and increases in abundance up to the adult stage (A. Jones, unpubl.). This single band presumably represents the two gld-I transcripts identified by cDNA sequence and RT–PCR analysis.

To examine the tissue specificity of gld-I RNA, total RNA was analyzed from adult populations of several mutants that affect the germ line. glp-4(bn2) is a conditional mutant that makes very few germ cells at the restrictive temperature [Beanan and Strome 1992]. The absence of gld-I transcript in total RNA from glp-4(bn2) adults [Fig. 7] suggests that the production of gld-I RNA is dependent on, and perhaps limited to, the germ line. The 2.5-kb gld-I RNA band is detected in total RNA from animals with germ lines undergoing both male [fem-3(q20gf); Barton et al. 1987] and female development [fem-1(hc17); Hodgkin 1986], consistent with the sex-nonspecific role of gld-I in premeiotic proliferation [Francis et al. 1995b].

Analysis of the two gld-I transcripts

Alternative splicing of gld-I RNA produces transcripts whose predicted protein products differ by 3 amino acids. The alternative splice product of gld-I is predicted to add Leu-Leu-Lys within the GSG domain [see Fig. 4A]. Because these changes could add functionality or change the specificity of RNA binding [e.g., Gorlach et al. 1994] or possibly eliminate function [e.g., Liang et al. 1991], we wanted to address whether the alternative splice was regulated developmentally or sex specific. To do this we undertook radioactive RT–PCR using RNA samples in which gld-I RNA is present at detectable levels on Northern blots. In all developmental stages tested, as well as in RNA samples from adult hermaphrodite animals that were making only sperm [fem-3(q20gf)] or only oocytes [fem-1(hc17)], the smaller RT–PCR product is present at roughly two to three times the amount of the larger product [Fig. 8]. Thus, we found no evidence that
Figure 9. A comparison of the two splice sites for exon 6 in C. elegans and C. remanei. Genomic sequence for C. elegans corresponds to nucleotides 1155–1175 (see Fig. 3). Bars match identical nucleotides, and arrows indicate the two splice sites that are used. Translations of the coding regions are shown above and below the respective sequences, and both the nucleotides and amino acids are italicized for the longer transcript.

Discussion

gld-1 has a tumor suppressor function in vivo

The gld-1 gene functions as a female germ cell-specific tumor suppressor. XX animals homozygous for the genetic null allele gld-1(q485) exhibit a phenotype in which germ cells fail to undergo oogenesis and proliferate ectopically, forming a germ-line tumor (Francis et al. 1995a). The results of the molecular analysis of gld-1(q485) are consistent with the genetic interpretation of this allele as null for the locus: q485 contains an amino-terminal frameshifting deletion and fails to make a gene product (A. Jones and T. Schedl, unpubl.).

GLD-1 contains evolutionarily conserved domains

The predicted GLD-1 protein has extensive similarity over a third of its length with two proteins: Sam68 and GRP33. We call this region, comprising ~170 amino acids, a GSG domain [see Fig. 4]. GRP33 is a putative hnRNP from brine shrimp (Cruz-Alvarez and Pellicer 1987) but has not been studied extensively since its identification. The mouse phosphoprotein Sam68 associates with Src in a cell cycle-dependent manner (Fumagalli et al. 1994; Taylor and Shalloway 1994) and is thought to be nearly identical to the sequence reported for GAP-associated p62 from humans [see Courtneidge and Fumagalli (1994) for clarification of nomenclature]. Sam68 contains several regions with the potential to interact with the Src homology regions SH2 and SH3 [Weng et al. 1994; Richard et al. 1995]; however, these regions of Sam68 lie outside the GSG domain and are not found in GLD-1 or GRP33.

The importance of the GSG domain to the in vivo function of GLD-1 is revealed by 16 independent gld-1 missense mutations that occur throughout this region of the protein, many in absolutely conserved residues (see Table 1; Fig. 4). These mutations affect gld-1 function in various ways, resulting in one of three phenotypic classes of gld-1 mutations: Missense mutations in Gly-227 abolish the essential function of gld-1 in directing oogenesis and suppressing tumor formation [phenotypic class A2]; mutations in either Gly-248 or Gly-250 result in a gf defect that leads to the masculinization of the hermaphrodite germ line [Mog phenotype: classes C1 and C2]; and mutation in any one of four residues in the GSG domain can suppress class C1 mutations intragenically to produce a loss-of-function phenotype [either mutant class A2 or B].

Another C. elegans gene, BO280.11b, encodes part of the GSG domain [see Fig. 4B]. Within this partial GSG domain, GLD-1 is more similar to Sam68 (41 amino acids identical) than it is to BO280.11b (35 amino acids identical). However, a second region of GLD-1 and BO280.11b is highly conserved [68% identical over 22 amino acids]. This small region, which we refer to as the CGA region, is just carboxy-terminal to the GSG domain and apparently not found in Sam68 and GRP33. This region, as well as a partial GSG domain, is also found in ZF1, a gene that is tightly linked to multiple endocrine neoplasia type 1 [MEN1] in humans (Toda et al. 1994). Like the GSG domain, the CGA region is important for GLD-1 function, as several gld-1 mutations affect it (refer to Table 1; and Fig. 4): An intragenic mutation in Glu-310 suppresses the class C1 mutant defect, with the resulting double mutant having a tumorous phenotype [phenotypic class A2]; mutation of Gly-308 results in a gf defect leading to a feminization of the germ-line (Fog) phenotype in both the hermaphrodite and male but does not disrupt oocyte differentiation [class D]; finally, deletion of the last 4 amino acids of the CGA region results in a defect in oogenesis as well as a gf defect that results in a hermaphrodite-specific Fog phenotype [class E].

The gld-1 GSG domain may be involved in RNA binding

A potential biochemical function for the GSG domain is suggested by a general in vitro RNA-binding activity for Sam68 (Wong et al. 1992; Taylor and Shalloway 1994; Wang et al. 1995). This ability to bind RNA may be attributable in part to the presence of a single KH motif within the larger region of similarity (see Fig. 4B). A number of other proteins that contain the KH motif have been implicated in RNA metabolism (Gibson et al. 1993). In most cases, the KH motif is present in multiple copies (Gibson et al. 1993; Siomi et al. 1993b). GSG domain proteins, however, contain a single KH motif. Furthermore, the single KH motifs of GSG domain proteins
are more similar to each other than to any other KH motif [see Fig. 4]. The high degree of similarity between GSG domain proteins, as well as a comparison of GSG domain proteins to other KH motif-containing proteins in overall structure, provides additional support for the assertion that the GSG domain proteins represent a specialized family of KH motif-containing proteins (Gibson et al. 1993). Conserved regions of the GSG domain outside of the KH motif may themselves be critical for modulating an RNA-binding activity of the KH motif [i.e., affecting the binding affinity or specificity]. Alternatively, these regions may be involved in contacts with other proteins.

Potential RNA targets for binding by KH motif-containing proteins have not been well characterized. Specific RNA targets for Sam68 binding have not been reported. FMR-1 has been demonstrated to bind its own transcript but has also been reported to bind <4% of all mRNA from human brain, implying a general nonspecific RNA-binding ability [Ashley et al. 1993]. However, a number of hnRNPs, originally thought to be nonspecific RNA-binding proteins, have been found to bind to specific targets with high affinity [e.g., Dreyfuss et al. 1993; Burd and Dreyfuss 1994; Gorlach et al. 1994]. In addition, in vitro experiments with hnRNP K suggest that the KH motif may also mediate DNA binding [Takimoto et al. 1993]. GLD-1 protein appears to be absent from the nucleus as determined by antibody staining [A. Jones and T. Schedl, in prep.]. Therefore, if GLD-1 is an RNA-binding protein, it most likely functions in the cytoplasm to control translation or mRNA stability.

Mutations in the KH motif of GLD-1 eliminate the major in vivo function of GLD-1 in directing oogenesis. Two different substitutions of an absolutely conserved glycine in the KH motif of gld-1 [see Fig. 5] result in the formation of germ-line tumors that are similar to those found in the genetic null allele q485 [Francis et al. 1995a]. Thus, missense mutations in the KH motif of GLD-1 result in loss of the GLD-1 tumor suppressor function in vivo.

gld-1 mutant protein may poison other gene products

Many gld-1 alleles behave in a dominant manner with respect to the nonessential role of gld-1 in promoting hermaphrodite spermatogenesis. The presence or absence of sperm in a hermaphrodite is easily scored, and the effect of various gld-1 alleles on this function was quantified [Francis et al. 1995a]. The analysis revealed a weak haploinsufficient Fog phenotype for large deletions in relation to the KH motif, the class C1 and C2 Mog mutants behave as partial loss-of-function mutations with respect to oogenesis [Francis et al. 1995b]. The molecular lesions in the class C1 and C2 Mog mutants occur in amino acids absolutely conserved among the GSG domain proteins [see Fig. 4A], adjacent to the KH motif. Given the position of these mutations in relation to the KH motif, the class C1 and C2 Mog mutations might act by failing to regulate the activity of the KH motif properly. Consistent with this hypothesis is the finding that the allele q93o255, which corresponds to the cis double mutant of the Mog allele q93 [class C1] and the tumorous allele oz89 [class A2], is phenotypi-
cally indistinguishable from the oz89 single mutant, even in its gf feminizing defect [Francis et al. 1995a]. Thus, the KH motif missense mutation appears to suppress the q93 missense mutation completely. For this reason, the gf activity of class C1 alleles must in some way act through a functional KH motif.

**GLD-1 as a regulator of germ-line development**

GLD-1 might function by regulating mRNA stability or translation. By functioning in this manner, GLD-1 could control aspects of germ-line development by acting as a specific regulator of a few key RNAs or a more general regulator of many RNAs. Germ-line translational control in *C. elegans* has been shown to be an important mode of regulating proliferation ([gld-1]; Evans et al. 1994), male development ([fem-3]; Ahringer and Kimble 1991), and female development ([tra-2]; Goodwin et al. 1993). Although these genes could be regulated ostensibly by GLD-1, none stand out as likely targets. *gld-1* tumors still form in the absence of *gld-1* product [Francis et al. 1995b], indicating that tumor formation is not caused by a failure to regulate *gld-1* activity; epistasis analysis revealed that *gld-1* acts downstream of *tra-2* [Francis et al. 1995b]; and masculinization in the *gld-1* class C1 Mog allele may not result from increased *fem-3* activity [Ellis and Kimble 1995].

If GLD-1 functions as a translational regulator, how might it act as a female germ-line tumor suppressor? The *mos* proto-oncogene provides an example of a cell cycle control factor that is regulated translationally during meiotic prophase. In *Xenopus* oocytes, the *mos* serine-threonine protein kinase is needed for activation of maturation promoting factor in meiosis I and for meiosis II arrest [Yew et al. 1993]. Although *mos* mRNA synthesis begins early in oogenesis, it is only translated in mature oocytes after the stimulus for meiotic maturation [Sagata et al. 1988]. Furthermore, inappropriate expression of *mos* can result in tumor formation [Yew et al. 1993]. In *gld-1* null mutants, mRNAs encoding factors that are synthesized for use late in oocyte development and are capable of promoting mitosis [such as *mos*] might be translated inappropriately. Alternatively, loss of *gld-1* activity might result in the inappropriate translation of mitotic factors packaged in the oocyte for embryonic cell divisions. Such ectopic mitotic activity early in oocyte development might then result in an exit from meiotic prophase, a return to mitotic proliferation, and subsequent tumor formation.

**GLD-1, Sam68, and mitosis**

*gld-1* has demonstrated a role in vivo for regulating mitotic activity negatively during pachytene of meiotic prophase [Francis et al. 1995a] and a nonessential, probably redundant role in regulating proliferation negatively in pachytene germ cells [Francis et al. 1995b]. Biochemical experiments show that the mammalian protein Sam68 interacts with Src in a cell cycle-specific manner [Fumagalli et al. 1994; Taylor and Shaloway 1994] and is phosphorylated on tyrosine during mitosis [Fumagalli et al. 1994]. Given that both Sam68 and GLD-1 contain GSG domains, it is tempting to speculate that both molecules might be involved in the regulation of mitosis in similar ways. However, the region of Sam68 that binds to Src shows no similarity to GLD-1. Interestingly, tyrosine phosphorylation of the carboxy terminus of Sam68 may regulate its RNA-binding activity [Wang et al. 1995]. Regions outside of the GSG domain could regulate GLD-1 activity in an analogous manner. The similarity between GLD-1 and Sam68 within the GSG domain suggests that if these two proteins do have a regulatory role in mitosis in common, targets for interaction with the GSG domain (either proteins or RNAs) might be conserved across phyla as well.

**Materials and methods**

**Physical mapping of the gld-1 gene**

Genetic mapping placed the *gld-1* locus in relation to three multilocus deletions, *nDf24*, *nDf25*, and *ozDf5* [Fig. 2A; Francis et al. 1995a]. Using sequence information from various clones spanning this region, PCR assays were developed and used on individual homozygotes from each deletion [Barstead et al. 1991]. PCR primer sets were derived from sequence of the cosmids *MO2B2* and *ZK1014* [primers *MO2B2* (a and b) and *ZK1014* (a and b); Y. Kozono, pers. comm.] and sequence of the LRF gene, which resides on cosmid F29D11 (primers F29D11 a and b; Yochem and Greenwald 1993). These assays placed *gld-1* between the cosmid clones F29D11 and ZK1014 on the physical map, and individual cosmids in this interval were used as probes on Southern blots to identify *gld-1* RFLPs [Fig. 2B; Coulson et al. 1988].

Transgenic experiments provide results that are consistent with identification of the *gld-1* gene. Six independent extra-chromosomal array-containing lines were generated by microinjecting pAJ37 [wild-type *gld-1* at 10 μg/ml] and the dominant rolling marker plasmid pRF4 [100 μg/ml; Mello et al. 1991]. All stable lines produced variable dominant phenotypes similar to *gld-1* loss-of-function (ectopic proliferation, pachytene arrest, abnormal oocytes, and feminization of the germ line) in a wild-type background, and all failed to rescue the tumorous phenotype after crossing into *gld-1(q485)* [A. Jones unpubl.]. Similarly, wild-type transgenes of another germ-line locus, *gfp-1*, can produce a variable dominant *gfp-1* loss-of-function germ-line phenotype in a wild-type background [A. Fire, pers. comm.]. The basis of these results is not understood presently.

**Nucleic acid preparation and analysis**

Standard methods for manipulation of DNA and RNA were used [Ausubel et al. 1991]. Sequence was determined for the majority of the 6.8-kb subclone pAJ37 by methods similar to those described in Wilson et al. [1994]. RNA was prepared from developmentally staged nematode populations as described [Meyer and Casson 1986; Goetinck and Waterston 1994].

**Isolation and analysis of cDNA clones**

Clones (1 × 10^6^) from a ZAP cDNA library [a gift from R. Barstead, Oklahoma Medical Research Foundation, Oklahoma City] were screened using a 1.0-kb EcoRI–HindIII fragment as a probe [see Fig. 2C]. Five clones were characterized by restriction
analysis and sequencing. Three appear to be identical and near full length, with the 5' end containing the last five nucleotides of the C. elegans trans-spliced leader SL1 [Krause and Hirsh 1987]. One of these clones, pLA1, was sequenced entirely.

Analysis of the 5' end of the gld-1 transcript was performed using the 5' RACE system (GIBCO/BRL) according to the manufacturer's instructions. Starting material was 1 μg of mixed-stage total RNA, with primer Afg29a used to prime reverse transcription. In amplification reactions with the supplied anchor primer, both primers Afg28 and Afg23 gave single-band products as assayed on agarose gels. Amplification products were cloned, and the 5' ends of 28 independent cDNA clones were sequenced, and 5 terminated at different points within the coding region. To look for additional differentially spliced products (other than the alternate splice described in Results), equivalent amounts of RNA from each developmental stage were combined and reverse transcribed with random primers as described below for analysis of the two known spliced forms. PCR analysis of this sample using numerous combinations of 5'-1-specific primers did not provide evidence for additional alternative splicing of the gld-1 transcript (data not shown).

Characterization of BO280.11b

A BLAST search with gld-1 identified similarity to the product of the predicted gene BO280.11 on the C. elegans cosmid BO280 (GenBank accession no. U10438). This sequence was used to make primers for 5'-RACE analysis as described above for gld-1. Reverse transcription was primed with Algort32, and the cDNA was amplified with Algort30 and anchor primer. A predominant band of ~600 bp was cloned. Four independent clones were sequenced, none of which contained a trans-spliced leader sequence. However, two clones with inserts of ~600 bp had identical 5' ends. The sequence of these clones revealed two important differences from the predicted splicing pattern. First, the 5'-RACE clones do not contain the first five predicted exons of BO280.11, which code for a potential tyrosine phosphatase. Second, some of the remaining five exons use different splice sites than predicted. The resulting gene, which we refer to as BO280.11b, has the following structure [numbered according to cosmid BO280]: exon 1 (34089-33988); exon 2 (33936-33851); exon 3 (33798-33474); exon 4 (33425-33328); exon 5 (33281-33049). We believe the BO280.11b cDNAs represent an independent translational unit, as exon 1 contains stop codons in each reading frame before the putative initiator methionine codon starting at 34027. We have no information about the 3' end of BO280.11b.

Isolation of gld-1 from C. remanei

To obtain sequence from the C. remanei gld-1 gene, we used the primer pair Afg16 and Alg20 in low annealing temperature PCR reactions on C. remanei genomic DNA. Using the sequence of the amplification product, we designed the C. remanei-specific primers AfgRg15, AfgRg16, and AfgRg16. 5'-RACE cloning and sequencing, as described above, was performed on C. remanei RNA using AfgRg16 and AfgRg15. Several clones that contained a trans-spliced leader identical to the C. elegans trans-spliced leader SL1 [Krause and Hirsh 1987] were sequenced completely. Additional sequence was obtained from the RT-PCR product of primers AfgRg16 and AfgR11.

The genomic sequence between exons 5 and 6 in C. remanei was determined from the amplification product of primers AfgRg16 and AfgR8 on genomic DNA (sequence around the splice sites is shown in Fig. 9). On the basis of this sequence, we constructed the primer AfgR11, which contains the last 12 nucleotides of exon 5 and the first 6 nucleotides of the alternate splice. AfgR11, in combination with AlgR8, AlgRc9, or AfgR11 gave single-band products in PCR on reverse-transcribed C. remanei RNA (data not shown), providing evidence that this splice site is used in C. remanei.

Analysis of gld-1(oz116) splicing

For the RT-PCR analysis of gld-1(oz116), RNA was isolated from the strain ndP4; unc-13(e51) gld-1(oz116). Primer AfgR20 was used to prime reverse transcription of 1 μg of total RNA. PCR was performed using primers AfgR19 and AfgR2, and the products were cloned and sequenced. Twenty-six independent clones were sequenced; 18 had the predicted wild-type sequence through the splice site junction at the final exon, and 8 had a 2-bp deletion relative to wild-type cDNAs, suggesting that these clones represent aberrant splice products that use an AG acceptor 2 nucleotides downstream of the normal acceptor. The wild-type clones may either represent RNA from the wild-type copy of gld-1 on ndP4 and/or correct splicing of mutant oz116 RNA similar to that seen by Arnaud et al. (1993).

RT-PCR analysis of gld-1 RNA

Approximately 5 μg of each RNA sample was reverse transcribed with 200 units of Superscript RT (GIBCO/BRL) at 42°C for 30 min with 15 μg of random hexamer primers (GIBCO/BRL). One-tenth of each reaction was PCR amplified directly using primers Alg20 [32p end-labeled] and AfgRc9. The amplification cycle profile was 94°C for 30 sec, 64°C for 30 sec, and 72°C for 45 sec, ending with 7 min at 72°C after the final cycle. Separate reactions were done at different numbers of cycles to determine the linear range of amplification for each sample. Amplification products were separated on denaturing polyacrylamide gels (5%, 7 m urea). Quantitation was performed using a Phosphorimager (Molecular Dynamics). To show that both spliced products would amplify at the same efficiency, experiments were first done on cDNA plasmid clones of known concentration [Fig. 8, data not shown]. Because of the sensitivity of the assay, a no-DNA control was always included and all RNA samples were tested for contamination by performing the reaction on RNA that had not been reverse transcribed.

Allele sequencing

Five adult animals homozygous for each allele of gld-1 were processed essentially as described in Williams et al. (1992). PCR was performed on these samples using one of three primer pairs to generate DNA for sequencing (AfgR9/Afg23, Afg21/Afg23, or Afg17/Afg37). Of each product, 25–50 ng was sequenced using [32p] end-labeled primers in a cycle sequencing reaction [Craik 1991]. The genomic DNA containing the gld-1 coding sequence was sequenced completely for all alleles with the exception of Mog alleles oz20, oz33, q62, oz34, oz16, and oz70, most of which were only sequenced with Alg21 and confirmed on the opposite strand with Alg18. In addition, six alleles (q93, q126, q268, q93oz50, oz10, and oz35) were sequenced through the 810-bp 3' UTR. For q343, ~200 bp of sequence from each end of the 2.3-kb insertion was obtained using the primers Alg28 and Alg32. A TA target site duplication occurred at the site of insertion, as has been described previously for Tc2 (Ruvolo et al. 1992). All mutations were confirmed on the opposite DNA strand.
Oligonucleotide primers used

The following oligonucleotides were used for this work, with the corresponding genomic sequence [numbered according to the sequence in Fig. 3] in parentheses; descending numbers indicate that the primer is on the opposite strand: AJgf9 (7-27); AJgf14 (521-540); AJgr14 (550-539); AJgf16 (678-695); AJgr17 (813-796); AJgf18 (939-958); AJgr20 (1089-1068); AJgf20 (1068-1088); AJgr21 (1182-1164); AJcr8 (1209-1193); AJgf21 (1206-1223); AJgr22 (1301-1316, 1366-1364); AJRcc9 (1377-1362, 1316-1314); AJgf23 (1442-1424); AJgf24 (1501-1520); AJcr11 (1634-1618, 1574-1571); AJgf25a (1589-1586); AJgf27 (1814-1831); AJgf28 (1831-1814); AJgf29a (2055-2034); AJgf30 (2112-2131); AJgr32 (2319-2301); AJgf33 (2375-2394); AJcr19 (2478-2459); AJgr20 (2530-2516); AJgr37 (2798-2781). Additional oligonucleotides used were MO2B2a GACAAGTCAAAGTCATTC, AJRGrl5 GCGACGTGCA, F29D1lb CAGAATGAGCAAGACGATCAGTC, GTCATGAGTGGATTG, F29D1la CATGGAGGTGAATGT- MO2B2a GCTCAGACCGCAGTTTCAG, MO2B2b GAGCACCG, AJgotr30 GCGAATTCGGATCCATCTTTTGTACATCCTTT- GCGGGATCCTCGAGTCGTTCTTTGGGACATAG, AJgotr32 GAATTCGGATCCTACACGAACACGACCG, AJRgfl6 GC-

The GSG domain and the CGA region of the GLD-1 protein are conserved between GLD-1 and the predicted gene product (T.A. Ebersole, Q. Chen, M.J. Justice, and K. Artzt, pers. comm.). All eight of the amino acids in which missense mutations are found are conserved between GLD-1 and the predicted quaking protein.

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