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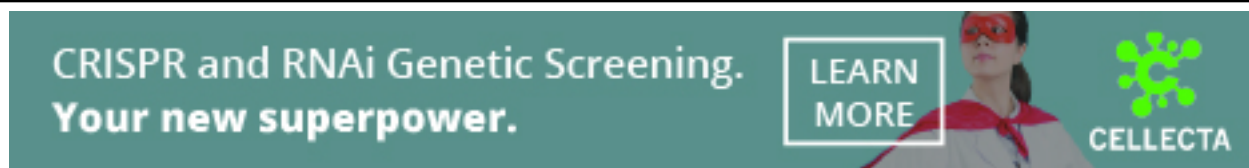
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# Toward Improving *Caenorhabditis elegans* Phenome Mapping With an ORFeome-Based RNAi Library

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The recently completed *Caenorhabditis elegans* genome sequence allows application of high-throughput (HT) approaches for phenotypic analyses using RNA interference (RNAi). As large phenotypic data sets become available, “phenoclustering” strategies can be used to begin understanding the complex molecular networks involved in development and other biological processes. The current HT-RNAi resources represent a great asset for phenotypic profiling but are limited by lack of flexibility. For instance, existing resources do not take advantage of the latest improvements in RNAi technology, such as inducible hairpin RNAi. Here we show that a *C. elegans* ORFeome resource, generated with the Gateway cloning system, can be used as a starting point to generate alternative HT-RNAi resources with enhanced flexibility. The versatility inherent to the Gateway system suggests that additional HT-RNAi libraries can now be readily generated to perform gene knockdowns under various conditions, increasing the possibilities for phenome mapping in *C. elegans*.

[Supplemental material is available online at [www.genome.org](http://www.genome.org).]

Thirty years of classical genetics in *Caenorhabditis elegans* identified and characterized ~800 genes (~4% of all genes; Brenner 1974; Jorgensen and Mango 2002). With the availability of the worm genome sequence (The *C. elegans* Sequencing Consortium 1998), it is now possible to apply genome-wide reverse genetic approaches for functional characterization. One way to uncover the functions of genes is to study their loss-of-function phenotypes after deletion (or gene knockout). Given the lack of homologous recombination in *C. elegans*, gene deletions are not easily amenable to high-throughput (HT) generation, although efforts are currently underway to achieve this goal. Instead, RNA interference (RNAi; Fire et al. 1998), which relies on the introduction of dsRNA in the worm, readily allows genome-wide loss-of-function screens by gene “knockdowns” (Fraser et al. 2000; Gonczy et al. 2000; Piano et al. 2000, 2002; Hanazawa et al. 2001; Maeda et al. 2001; Zipperlen et al. 2001; Ashrafi et al. 2003; Kamath et al. 2003; Lee et al. 2003; Pothof et al. 2003; Simmer et al. 2003; Vastenhouw et al. 2003).

An RNAi-by-feeding (Timmons and Fire 1998) library was generated by using PCR-amplified genomic DNA fragments as a template (Kamath et al. 2003). This resource targets ~78% of the 19,920 genes currently predicted in Worm Sequence 112 (WS112; WormBase, <http://www.wormbase.org>; see Methods for calculations; Kamath et al. 2003; Harris et al. 2004). This library has been successfully used for several genome-wide RNAi screens, for example, in wild-type (WT) N2 animals for systemic functional analysis (Kamath et al. 2003) and for the analysis of genes in-

involved in the regulation of body fat storage (Ashrafi et al. 2003), as well as in the hypersensitive *rrf-3* *C. elegans* strain (Simmer et al. 2003). These data represent a significant step forward in the phenotypic characterization of most *C. elegans* genes.

Ultimately, the combination of loss-of-function data obtained in numerous, different conditions could lead to the classification of worm genes by “phenotypic signatures.” Phenoclustering methods have been applied on a moderate scale in *C. elegans* (Boulton et al. 2002; Piano et al. 2002) and in *Drosophila* (Boutros et al. 2004). The *C. elegans* “phenome” map resulting from such genome-wide analyses would increase our understanding of a variety of biological processes.

However, several technical points need to be resolved before a comprehensive phenome map can be generated for *C. elegans*. First, such a map relies upon the generation of RNAi data for all genes. Today, not all worm genes are yet targeted by an RNAi clone. Second, RNAi screens should be performed in different environmental conditions or various genetic backgrounds, applying different procedures. For example, experiments leading to the loss-of-function of two genes, simultaneously, can be carried out in the worm (Simmer et al. 2003; Tewari et al. 2004). Furthermore, RNAi can be applied by different methods of delivery, such as injection (Fire et al. 1998), feeding (Timmons and Fire 1998), or soaking (Tabara et al. 1998), at different times during the development of the worm such as different larval stages, and with different DNA templates for the synthesis of dsRNA, such as genomic DNA fragments, cDNAs, or ORFs. Finally, a new RNAi technology, referred to as inducible hairpin RNAi (Tavernarakis et al. 2000; N.M. Johnson, C.A. Behm, and S.C. Trowell, unpubl.), opens new possibilities for targeting genes. Indeed, genes can be knocked-down at specific times during the development and specific cell types such as neurons can now be targeted. Other improved technologies for loss-of-function assays will likely be de-

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veloped soon. To accommodate these numerous applications, RNAi resources should thus be generated by using cloning tools that are as flexible as possible (Fig. 1).

In the infancy of the genomic era (Collins et al. 2003), resources need to be dynamic and flexible. Ideally, a single, comprehensive set of cloned ORFs, that is, an “ORFeome” (Walhout et al. 2000b; Reboul et al. 2003; Rual et al. 2004), should be usable for various HT applications. An ORFeome resource should be updated (ORFs corrected, removed, or added) regularly as new genome annotations become available. An ORFeome should be cloned in a format that allows facile transfer of the ORFs to any desired vector as new technologies become available (Fig. 1). Recently developed recombinational cloning technologies (Liu et al. 1998; Hartley et al. 2000; Walhout et al. 2000a; Zhang et al. 2002) allow such versatile cloning approaches at HT (Reboul et al. 2003; Dricot et al. 2004).

Upon the release of the *C. elegans* genome sequence (The *C. elegans* Sequencing Consortium 1998), a genome-wide approach was initiated to clone all predicted ORFs of the worm ORFeome (Walhout et al. 2000b; Reboul et al. 2001, 2003; Vaglio et al. 2003; WorfDB, <http://worfdb.dfci.harvard.edu>) using the Gateway recombinational cloning system (Hartley et al. 2000; Walhout et al. 2000a). Version 1.1 of the *C. elegans* ORFeome resource contains 11,942 cloned ORFs (Reboul et al. 2003) and constitutes a platform for proteomic and functional genomic approaches (Reboul et al. 2003; Rual et al. 2004). The *C. elegans* ORFeome resource is dynamic, with new ORFs added or corrected continuously (Lamesch et al. 2004). Phenome mapping, as well as other “omic” approaches, should benefit from the flexibility of this resource.

By using the *C. elegans* ORFeome v1.1 resource (Reboul et al. 2003), we began generating a companion resource for RNAi screening. Herein we describe this ORFeome-RNAi v1.1 library, as well as the results from using this library for a genome-wide RNAi-by-feeding screen of WT *C. elegans* hermaphrodites. We conclude that the use of an ORFeome approach is feasible for the generation of an alternative RNAi library, and we suggest that this cloning strategy could be applied to other functional genomic approaches.

## RESULTS

### Generation of the ORFeome-RNAi v1.1 Library

The *C. elegans* ORFeome v1.1 library contains 11,942 ORFs cloned as Gateway Entry clones, comprising 10,623 ORFs cloned “in frame” plus 1319 ORFs cloned “out of frame” (Reboul et al. 2003). ORFs were cloned out of frame because of mispredictions of their ATG or STOP codons. Only the in-frame ORFs can be

used for protein expression, but both sets of clones can be used for RNAi. HT-recombinational cloning protocols were used to transfer ORFs from the pDONR201 Entry plasmid into the pL4440-dest-RNAi Destination vector by using the LR Gateway reaction (see Methods). In all, 11,804 RNAi clones were generated (Supplemental Table A). These clones were archived as glycerol stocks of transformed *Escherichia coli* strain HT115(DE3) used in RNAi-by-feeding and as miniprep plasmid DNAs (Timmons and Fire 1998). These DNA preps can be used as templates for in vitro dsRNA synthesis before RNAi application by soaking or injection. The ORFeome-RNAi v1.1 library is available to the community upon request (Open Biosystems, <http://www.openbiosystems.com>; MRC geneservice, <http://www.rfcgr.mrc.ac.uk/geneservice>; WorfDB, <http://worfdb.dfci.harvard.edu>).

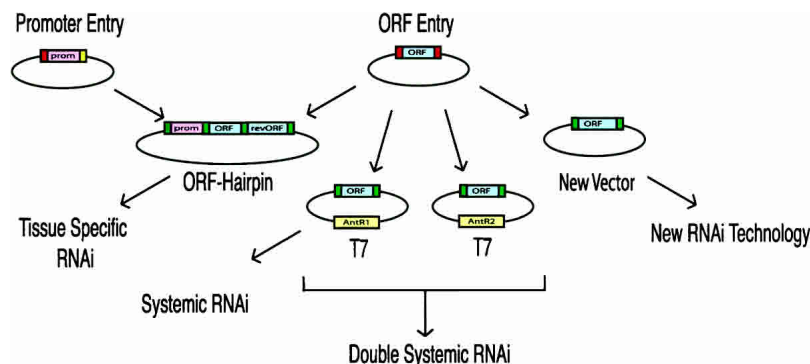
Bioinformatic analysis revealed that the ORFeome-RNAi v1.1 library contains 11,511 RNAi clones, each expected to target a single gene (see Methods; Supplemental Table A). Several of these RNAi clones target the same genes, so the RNAi library in theory can knockdown the expression of 10,953 (~55%) of the 19,920 unique protein-encoding genes predicted in WS112 (Supplemental Table B). Of these 10,953 genes, 485 are targeted by two or more RNAi clones. There are 1736 genes, not previously targeted by other existing RNAi-by-feeding libraries, that now have a clone available in the ORFeome-RNAi v1.1 library (Supplemental Table B). This represents an increase of ~9% in the total number of genes that can be targeted by RNAi-by-feeding. In combination with the previous RNAi-by-feeding library (Kamath et al. 2003) a total of 17,201 predicted worm genes (~86%) can now be targeted by an RNAi-by-feeding clone.

### Genome-Wide Phenotypic Analysis Using the ORFeome-RNAi v1.1 Library

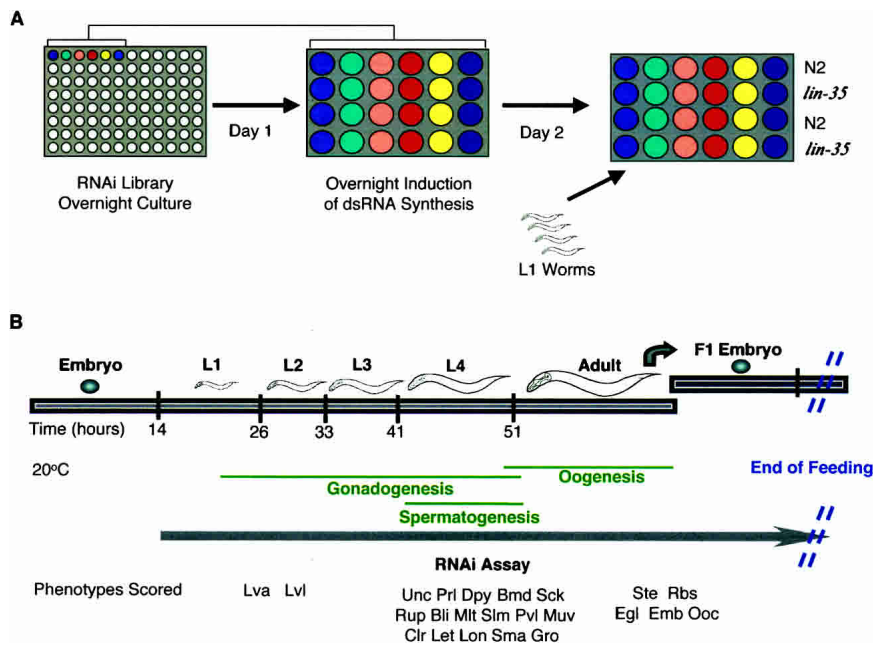
We first used the ORFeome-RNAi v1.1 library for a synthetic lethal screen by conducting a genome-wide RNAi screen simultaneously in WT N2 *C. elegans* hermaphrodites and in a *lin-35/Rb* mutant background (see Methods; Fig. 2). The results in the N2 animals are presented here, and the results of the *lin-35/Rb* synthetic phenotype screen will be reported elsewhere (J. Ceron, J.-F. Rual, M. Vidal, and S. van den Heuvel, unpubl.). Worms were subjected to RNAi-by-feeding at the first larval stage (L1). We identified a variety of phenotypic defects, including lethality and abnormal body morphology, occurring during the embryonic, larval, and adult stages.

The results from this phenotypic analysis of 10,953 WS112 predicted genes are described in Table 1 and in Supplemental Table B. We observed phenotypes different from WT for 1066 WS112 predicted genes, representing ~10% of the total ORFs tested. This percentage is similar to that obtained in previous genome-wide RNAi-by-feeding screens (Kamath et al. 2003; Simmer et al. 2003).

We observed an RNAi phenotype for 220 WS112 predicted genes for which no phenotype was previously reported in WormBase (Supplemental Table B). For 79 of these 220 genes, no RNAi assay was listed in WormBase. The other 141 genes, targeted in previous RNAi screens, had showed a WT phenotype before. Altogether, our results increase the total number of genes associated with any RNAi phenotype by ~7%, from 3259 to 3479. Finally, an additional 1145 genes for which no RNAi assay was reported in WormBase did not show any detectable phenotype and were scored as WT in our screen.



**Figure 1** Generating RNAi resources from flexible Gateway ORFeome and promoterome collections. AntR1 and AntR2 refer to antibiotic resistance markers number 1 and 2.



**Figure 2** Overview of the RNAi screening procedure. (A) Overnight cultures of 96 bacterial clones are inoculated in sixteen 24-wells plates (six clones per plate and four wells per clone). After overnight induction of the dsRNA synthesis, three to 10 worms synchronized at the L1-stage (N2 or *lin-35* strain) were deposited into the wells. (B) RNAi experiments were performed at 20°C. We observed a wide range of phenotypes across development.

A global analysis of the data verifies observations made previously regarding the distributions of viable postembryonic phenotypes (Vpep) and nonviable phenotypes (Nonv) ascribed to various genes located on either the autosomes or the X chromosome (Piano et al. 2000; Maeda et al. 2001; Kamath et al. 2003). In particular, there appears to be a low incidence of genes located on the X chromosome that are associated with any phenotype (Table 1).

Gene inactivation by RNAi-by-feeding is often incomplete and may be too limited to cause fully penetrant loss-of-function phenotypes, thereby generating false-negative results. For example, RNAi is less effective in neurons (Tavernarakis et al. 2000), and genes encoding proteins with long half-lives are more resistant to RNAi (Hannon 2002). Previous genome-wide RNAi screens had various rates of false negatives (Kamath et al. 2003; Simmer et al. 2003), based on comparison of the RNAi results to the loss-of-function phenotypes caused by genetic mutations. Nearly 45% of the genes with a reported mutant phenotype did not show any phenotype in each of these previous studies. More precisely, rates of ~25% and ~55% of false negatives were observed for Nonv mutant phenotypes and for Vpep mutant phenotypes, respectively. Matching our RNAi data to the phenotypes described for 269 genetic mutants (resembling the control set of genetic mutants used in Kamath et al. [2003] and Simmer et al. [2003]), we observed an overall false-negative rate of ~59% (~43% for Nonv mutant phenotypes and ~83% for Vpep mutant phenotypes; Supplemental Table C). This somewhat higher frequency of false negatives can be explained mostly by our HT RNAi protocol, in which animals were followed only for one generation and not transferred to a second RNAi plate (see Discussion).

Low rates of false positives were previously described in genome-wide RNAi-by-feeding screen (Kamath et al. 2003; Simmer et al. 2003). In these studies, a false positive was scored when the RNAi phenotype was lethal, whereas the reported genetic mutant

shows a nonlethal phenotype. Published false-positive rates vary from ~0% to 0.4% (Kamath et al. 2003; Simmer et al. 2003). By applying the same strategy to a similar set of 103 loci, we counted four Nonv phenotypes in our RNAi experiment (Supplemental Table D). After sequence verification, we found one obvious cross-contamination. The remaining three clones have the correct sequence. Two of the three corresponding genes are reported as Nonv by RNAi in WormBase. Overall, our rate of false-positive results is low and in part can be due to cross-contamination.

## Reproducibility

The ORFeome-RNAi v1.1 library contains 430 pairs in which both clones target the same gene and should give rise to the same phenotype. These clones were processed independently and therefore constitute a blind control for the reproducibility of scoring. In 78% (337/430), we obtained a WT phenotype for both clones of the pair. In 10% (45/430), we observed a phenotype for both clones. Thus, the results are in agreement regarding the scoring of WT versus phenotype for ~89% of the pairs of clones (382/430; Supplemental Table E). In ~11% (48/430), there is disagreement. Variability in the results from repeated RNAi-by-feed-

ing assays was observed previously (Simmer et al. 2003). Variability ranges from ~10% to 30% between experiments and is likely due to lack of reproducibility from one RNAi assay to another (Simmer et al. 2003). To verify this phenomenon, we performed a second RNAi assay for the 48 pairs of RNAi clones in disagreement. In the second assay, we observed similar phenotypes for both clones ~52% of the time (25/48; Supplemental Table E). Thirteen pairs of clones now gave rise to a WT phenotype for both clones. Ten pairs of clones were still not in agreement; their correct identities were verified by DNA sequence analysis.

Interestingly, of the 10 pairs of clones in disagreement in the second RNAi assay, six showed large differences in size of the cloned ORF. For these six pairs, the longer ORF clone was the one giving rise to the observed phenotype. These numbers are not statistically significant but suggest that effectiveness of RNAi-by-feeding may be somewhat dependent on the length of the clone, the longer clones being more effective in RNAi. We went on to perform RNAi assay by injection for three of these six pairs of clones and obtained a phenotype in all six (three by two) RNAi-by-injection assays. Thus, the output of an RNAi assay depends on the method of delivery of dsRNA.

## DISCUSSION

The generation of complete resources of cloned ORFs represents an important and challenging step in the development of proteomics and functional genomics (Reboul et al. 2003; Rual et al. 2004). Within months after the generation of the *C. elegans* ORFeome v1.1 (Reboul et al. 2003), we transferred >11,000 ORFs into the pL4440-dest RNAi vector, generating a new RNAi resource. The technology that allowed this genome-wide cloning effort is now being used for the production of additional resources, such as Activation Domain and DNA Binding Domain fusion collections for yeast two-hybrid screens.

The *C. elegans* ORFeome-RNAi v1.1 library contains 11,511

**Table 1.** Summary of RNAi Data

Genome annotation	Genome					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	10,953	16,757	16,757	86.2%		
Phenotypes	1066	1722	1722	10.3%		
Nonviable phenotypes	928	1170	1170	7.0%		
Viable phenotypes only	138	552	552	3.2%		
Emb	342	929	929	5.5%		
Rbs/Ste/Stp	602	474	474	2.8%		
	<b>Chromosome I</b>					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	1670	2445	2445	ND <sup>b</sup>	1979	2978
Phenotypes	272	334	334	13.7%	202	354
Nonviable phenotypes	239	240	240	10.3%	182	240
Viable phenotypes only	33	81	81	3.3%	20	114
Emb	88	226	226	9.2%	60	204
Rbs/Ste/Stp	149	98	98	4.0%	122	76
	<b>Chromosome II</b>					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	1670	2445	2445	ND <sup>b</sup>	1533	2132
Phenotypes	272	334	334	11.9%	232	395
Nonviable phenotypes	239	240	240	8.1%	197	298
Viable phenotypes only	33	81	81	3.9%	35	97
Emb	88	226	226	6.9%	70	220
Rbs/Ste/Stp	149	98	98	2.5%	139	150
	<b>Chromosome III</b>					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	1670	2445	2445	ND <sup>b</sup>	1333	2132
Phenotypes	272	334	334	11.9%	232	395
Nonviable phenotypes	239	240	240	8.1%	197	298
Viable phenotypes only	33	81	81	3.9%	35	97
Emb	88	226	226	6.9%	70	220
Rbs/Ste/Stp	149	98	98	2.5%	139	150
	<b>Chromosome IV</b>					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	1848	2693	2693	ND <sup>b</sup>	1433	2357
Phenotypes	166	293	293	10.9%	49	131
Nonviable phenotypes	143	187	187	6.9%	41	47
Viable phenotypes only	23	106	106	3.9%	8	84
Emb	55	143	143	5.3%	16	28
Rbs/Ste/Stp	93	72	72	2.7%	22	15
	<b>Chromosome V</b>					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	1848	2693	2693	ND <sup>b</sup>	1433	2357
Phenotypes	166	293	293	10.9%	49	131
Nonviable phenotypes	143	187	187	6.9%	41	47
Viable phenotypes only	23	106	106	3.9%	8	84
Emb	55	143	143	5.3%	16	28
Rbs/Ste/Stp	93	72	72	2.7%	22	15
	<b>Chromosome VI</b>					
	ORFeome-RNAi v1.1		Kamath et al. (2003)		WS61 <sup>b</sup>	
	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>	WS112 <sup>a</sup>	WS61 <sup>b</sup>
Genes targeted	1848	2693	2693	ND <sup>b</sup>	1433	2357
Phenotypes	166	293	293	10.9%	49	131
Nonviable phenotypes	143	187	187	6.9%	41	47
Viable phenotypes only	23	106	106	3.9%	8	84
Emb	55	143	143	5.3%	16	28
Rbs/Ste/Stp	93	72	72	2.7%	22	15

<sup>a</sup>The total number of predicted genes in WS112 is 19,920; 2850 on chromosome I; 3457 on chromosome II; 2639 on chromosome III; 3242 on chromosome IV; 4970 on chromosome V; and 2762 on chromosome X.

<sup>b</sup>The total number of predicted genes in WS61 is 19,427; the number of predicted genes for each chromosome could not be found; ND means not determined.

RNAi clones, each expected to target a single gene (Supplemental Table A), and includes 1736 *C. elegans* genes not targeted previously by any RNAi-by-feeding clones (Supplemental Table B). We used the *C. elegans* ORFeome-RNAi v1.1 library to perform a genome-wide RNAi screen in WT *C. elegans* hermaphrodites (Fig. 2). We observed a phenotype for ~10% of the WS112 genes studied (Supplemental Table A; Table 1). Our RNAi results increase by 220 the number of genes for which an RNAi phenotype has been described. We confirmed phenotypes for numerous genes.

As previously observed (Simmer et al. 2003), we found that the RNAi effect varies from one RNAi-by-feeding screen to another. This lack of reproducibility is likely due to the inability to control the amount of dsRNA animals are exposed to, resulting in a high rate of false negatives. We also suggest that the RNAi effect could depend on the method of dsRNA delivery and on the size of the RNAi clone, with longer clones generally more effective, as described previously for RNAi-by-injection (Parrish et al. 2000). As for most large-scale approaches, results obtained from genome-wide RNAi screens should be interpreted carefully, especially negative results. Several issues need to be addressed before a comprehensive phenome map of *C. elegans* can be generated, for example, by generating genome-wide RNAi data in different conditions and by developing new RNAi protocols. In this regard, ORFeome-based RNAi resources should help improve phenome mapping by taking advantage of the versatility of recombinational cloning to apply new RNAi technologies to genome scale (Fig. 1).

Cloned ORFs versus cloned genomic DNA fragments, as template DNA could improve the efficiency of RNAi. Cloned ORFs are free of introns and thus provide more template for siRNAs. Moreover, a study based on the RNAi results obtained with the genomic RNAi library (Kamath et al. 2003) showed a lower incidence of RNAi phenotype among loci with a small portion of coding sequence and large introns. This observation suggested that genomic DNA as RNAi template reduces RNAi efficiency (Cutter et al. 2003). Nevertheless, our proportion of hits (Table 1) and rate of false-negative results are comparable to previous results (Kamath et al. 2003; Simmer et al. 2003). Furthermore, an analysis using the current version of the genome annotation demonstrates that the fraction of coding gene sequence exerts a weaker negative effect than initially reported (A.D. Cutter, pers. comm.). Overall, it seems that the presence of introns in the DNA template does not have a dramatic negative impact on the efficiency of RNAi.

In our screen, animals were exposed to RNAi-by-feeding during the L1 stage (Fig. 2), as opposed to the L4 stage in previous screens (Kamath et al. 2003; Simmer et al. 2003). Consequently, we observed phenotypes linked to sterility (sterile progeny, and reduced brood size) for a larger number of genes (~5.5% of those analyzed) than reported before (~2.8%; Table 1; Kamath et al. 2003). On the other hand, Kamath et al. (2003) observed an embryonic lethal phenotype (Emb) for ~5.5% of assayed genes, whereas we observed 3.1%. This difference might arise because many genes are essential for *both* the development of the adult germ line and the embryos. For these genes, scoring Emb versus sterile phenotypes could depend on the time of initiation of the RNAi assay, larval stage L1 versus L4.

Several factors are likely to contribute to a reduced frequency of phenotypes as well as to phenotypes that differ from those listed in WormBase for a given gene. First, we used a different RNAi protocol, and the criteria for scoring phenotypes were slightly more stringent than those used previously. To increase the throughput of the screening, we initiated the screening on L1 stage worms, we followed the animals for one generation and did not transfer them to a second RNAi plate. Phenotypes occurring late in the screen were difficult to score because

food was limited. Subtle phenotypes may have been overlooked because we were scoring for visually obvious phenotypes.

Despite the higher rate of false negatives, we observe a comparable rate, ~10%, of hits for all genes targeted (Table 1). This is probably due to higher representation in the ORFeome v1.1 resource of genes that can give rise to a phenotype. Highly expressed genes that are more represented in the ORFeome1.1 resource (Reboul et al. 2003) have more chance to be essential (Cutter et al. 2003).

The *C. elegans* ORFeome v1.1 and ORFeome-RNAi v1.1 libraries represent ~55% of the predicted worm ORFeome. Importantly, these collections should be considered as dynamic resources. New Gateway Entry clones, generated in the ongoing ORFeome project (Lamesch et al. 2004), will be subsequently transferred into the RNAi vector in the near future. If a new improved RNAi vector is made available to the community, the Gateway cloning system will allow quick transfer of all cloned ORFs into it. For example, a hairpin RNAi method (Tavernarakis et al. 2000) was developed by using the ORFeome v1.1 resource (Reboul et al. 2003) to generate hairpin RNAi constructs in the pWormgate system (N.M. Johnson, C.A. Behm, and S.C. Trowell, unpubl.). In addition, such hairpin constructs could be fused to any specific promoter cloned by MultiSite Gateway (Fig. 1; Dupuy et al. 2004).

The complete ORFeome-RNAi v1.1 library (Supplemental Table A) is available upon request from Open Biosystems (<http://www.openbiosystems.com>) and MRC geneservice (<http://www.rfcgr.mrc.ac.uk/geneservice>; WorfdB, <http://worfdb.dfci.harvard.edu>). The data generated in the WT RNAi screen (Supplemental Table B) will be available in WormBase shortly (<http://www.wormbase.org>; Harris et al. 2004) and in RNAiDB (<http://nematoda.bio.nyu.edu>; Gunsalus et al. 2004). Integrated with other functional genomic data, these RNAi resources should contribute to a deeper understanding of the biology of *C. elegans*.

In summary, we demonstrated the value of an ORFeome-based approach for the generation of alternative HT-RNAi resources and ultimately, for the improvement of phenome mapping in *C. elegans*.

## METHODS

### Generation of the ORFeome-RNAi v1.1 Library

ORFs were transferred into pL4440-dest-RNAi, a Gateway-compatible RNAi vector adapted from the original pL4440-RNAi vector (Timmons and Fire 1998). Gateway recombinational cloning reactions were performed as described by the manufacturer (Invitrogen) with minor changes. The pL4440-dest-RNAi vector was digested with EcoRI and NcoI before the LR reaction, which significantly improved (>10-fold) the efficiency of the LR reaction. LR reaction plates were incubated overnight at 25°C. By using 96 well plates and HT-liquid handling systems, up to 1500 LR reactions were carried out by one person per day.

It is not possible to transform LR products directly into the RNAi-by-feeding bacterial strain HT115 (DE3), an RNase III-deficient strain of *E. coli*. Indeed, HT115 (DE3) bacteria are resistant to the *ccdB* toxic gene used in the Gateway cloning system. Therefore, LR products were first transformed into *E. coli* DH5 $\alpha$  strain. Bacterial transformations were performed with 100  $\mu$ g/mL ampicillin selection in liquid cultures. Plasmid DNA minipreps were prepared in a 96-well format for each pool of transformants by using a Qiagen 9600 Robot. DNA preparations were subsequently transformed into the RNAi-by-feeding bacterial strain HT115 (DE3). Out of the 11,942 ORFs cloned as Entry clones in the *C. elegans* ORFeome v1.1 resource, we obtained HT115 (DE3) transformants for 11,804 ORFs (Supplemental Table A).

All the Entry clones used in the LR reaction were verified by DNA sequencing analysis (Reboul et al. 2003). We also verified

the identity of 96 RNAi clones picked at random from the library by sequencing. The universal primers used for PCR amplification and sequencing are pL4440-dest-RNAi-FOR (5'GTTTCCAGT CACGACGTT3') and pL4440-dest-RNAi-REV (5'TGGATAACCG TATTACCGCC3'). We observed the expected sequences in 96% of the time. Three of the clones contained the wrong ORFs, and one clone did not contain any insert. We conclude that a tiny portion of the phenotypes may have been attributed to the wrong genes due to cross-contamination. The identity of the 220 RNAi clones corresponding to genes for which no phenotype was previously reported in WormBase has been verified by DNA sequence analysis.

## Bioinformatic Analysis

For the *C. elegans* genome sequence and related genome annotation, we used version WS112 available at WormBase (<http://www.wormbase.org>; Harris et al. 2004). There are 19,920 protein encoding predicted genes in WS112 (Supplemental Table B).

To determine which WS112 genes are targeted by the ORFeome-RNAi v1.1 clones, we first retrieved the sequences from the primer pairs used in the ORFeome project (Reboul et al. 2003; Vaglio et al. 2003) and used these sequences to perform BLAST analysis against version WS112 of the worm genome. For each RNAi clone, the resulting position was then queried for an overlap of >100 nucleotides with the transcript(s) (UTRs plus exonic sequences) of one or more predicted WS112 genes.

With this strategy, we determined that there are 153 RNAi clones that target more than one WS112 predicted gene (Supplemental Table A). Furthermore, there are 140 RNAi clones that do not target any WS112 predicted genes (Supplemental Table A). These results are somewhat surprising because the cloned ORFs were PCR amplified from a worm cDNA library (Reboul et al. 2003). The information related to these cloned ORFs should be corrected in forthcoming releases from WormBase. The results obtained with these 293 RNAi clones were removed from our analysis. We found 10,953 WS112 genes targeted by the collection of 11,511 remaining RNAi clones. Out of the 10,953 genes, 485 were targeted by more than one RNAi clone. Given the low rate of false-positive results and the high rate of false-negative results in RNAi screens (see Results; Kamath et al. 2003; Simmer et al. 2003), all the phenotypes different from WT were scored by default for those 485 genes in the Supplemental Tables B, C, and D.

The data related to the previous RNAi library (Kamath et al. 2003) were retrieved from WormBase WS112 (WormBase, <http://www.wormbase.org>; Harris et al. 2004). In WS112, there are 16,537 RNAi clones that target 15,465 WS112 genes. However, 16,757 RNAi clones were described as targeting 16,757 WS61 genes (Kamath et al. 2003). The discrepancy is probably caused by altered gene predictions between WS61 and WS112, and missing data for 220 RNAi clones in WormBase WS112.

## RNAi Screening

We performed the RNAi screen simultaneously in WT N2 *C. elegans* hermaphrodites and in a *lin-35/Rb* mutant background (Fig. 2). Standard methods were used for culturing worms on nematode growth medium (NGM; Brenner 1974). Screening was performed in 24-well format plates containing NGM supplemented with 50 µg/mL ampicillin, 12.5 µg/mL tetracycline, and 1.43 mg/mL IPTG. Unseeded feeding plates were kept in the dark for less than a week at 4°C. Before bacterial inoculation, the feeding plates were dried for 2 to 3 h in a laminar airflow hood or in a 37°C incubator. RNAi bacterial clones were grown in 96-well format for 24 h at 37°C in 600 µL LB with 50 µg/mL carbenicillin and 12.5 µg/mL tetracycline. Each well of the 24-well format plates was inoculated with 115 µL of bacterial culture by using an extendable automated multichannel pipette. The expression of dsRNA was induced overnight in the presence of IPTG (1.43 mg/mL), at room temperature and in the dark. After dsRNA induction, three to 10 worms synchronized in the L1-stage were deposited into the wells (Fig. 2).

Most of the previous RNAi studies were done by initiating

RNAi on young adults (Fraser et al. 2000; Gonczy et al. 2000; Piano et al. 2000, 2002; Maeda et al. 2001; Zipperlen et al. 2001; Kamath et al. 2003; Simmer et al. 2003). Instead, we fed worms at the L1 larval stage (Fig. 2). This procedure necessitates fewer worm manipulations and thus allows a higher throughput. A disadvantage of this method is food limitation in the F1 generation. This might explain the high rate of false-negative results in our assay, particularly for Vpep phenotypes.

Phenotypes were scored visually twice on days 4 and 5 of the 20°C culture. We looked for 18 different Vpep phenotypes: Lva (larval arrest), Unc (uncoordinated), Prl (paralyzed), Dpy (dumpy), Bmd (body morphology defect), Sck (sick), Bli (blister), Mlt (molting defect), Slm (slim), Him (high incidence of male), Pvl (protruding vulva), Muv (multivulva), Clr (clear), Lon (long), Sma (small), Gro (growth defect), Egl (egg laying defect), and Soc (social). We looked for eight different Nonv phenotypes: Ste (sterile), Rbs (reduced brood size), Let (Lethal), Emb (embryonic lethal), Ooc (oocytes), Stp (sterile progeny), Rup (ruptured), and Lvl (larval lethal). When the phenotype occurs in the F1 generation, F1 is added before the phenotype name (e.g., F1 Lva). Emb was defined as >20% dead embryos. Phenotypes observed with partial penetrance are indicated with a "p", for example, pEmb for partially penetrant embryonic lethality. A partial phenotype is scored when the phenotype occurs in <90% of the P<sub>0</sub> or F1 progeny.

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