

WORMBASE – NEMATODE BIOLOGY AND GENOMES

Paul Davis, Wellcome Trust Genome Campus.

WormBase Consortium, (PI List) Lincoln Stein – OICR, Paul Sternberg – CALTECH, John Spieth - GSC, Richard Durbin – WTSI

WormBase www.wormbase.org

WormBase is the major public online database resource for the Caenorhabditis research community. The database was developed primarily for the nematode *C. elegans* but expanded to host genomes and biological data from other closely related nematode species including *C. briggsae*, *C. remanei*, *C. brenneri*, *C. japonica* and *Pristionchus pacificus*. WormBase has developed tools to mine the data held within the database and compare the hosted species. Over the years we have developed a variety of curation pipelines which often begin in a "first-pass" literature curation step. This involves a brief overview of the literature before directing it to specialised data curators who extract all relevant information. Curators focus on particular data types or experimental techniques such as gene structure changes (see the Sequence curation poster), variations, phenotypes or RNAi and their expertise in these fields make curation efficient. WormBase works with many other groups and consortiums to validate, process and integrate both large and small scale data resources. WormBase also provides data that will be of interest to the wider biomedical and bioinformatics communities allowing researchers to utilise the information and techniques offered by nematodes to study wider aspects including medicine and disease.

Automated First Pass Paper Curation

Paper and Data Type Identification:
There is a project underway to move from a manual approach to a semi automated pipeline with author input. Currently we are in a transitional phase moving from WB Curators towards Authors and Text Mining.



searched using keyword 'elegans'
Manual selection of papers

PDFs Download Automatically and stored in a database.



27 Data types extracted (2009)

Data curator Data curator Data curator Data curator

Author First-Pass Form

Please click the box next to the type of data your publication includes.

C. elegans ? Add information. *C. elegans* other than *C. elegans*? ? Add information. Nematode species other than *C. elegans*? ? Add information. Non-nematode species? ? Add information.

Gene Identification and Mapping:

Genes identified in this paper? ? Add information. Newly cloned genes? ? Add information. Newly created alleles? ? Add information. Genetic mapping data? ? Add information.

Gene Function:

Mutant, RNAi, Overexpression, or Chemical-based Phenotypes. Please specify your data type. Small-scale RNAi (less than 100 individual experiments)? ? Add information. Large-scale RNAi (greater than 100 individual experiments)? ? Add information. Overexpression phenotype? ? Add information. Chemicals? ? Add information. Mouse analysis? ? Add information. Tissue or cell site of action? ? Add information. Time of action? ? Add information. Molecular function of a gene product? ? Add information. Homology of a human disease-associated gene? ? Add information.

Interactions:

Genetic interactions? ? Add information. Functional complementation? ? Add information. Gene product interaction? ? Add information.

Regulation of Gene Expression:

New expression pattern for a gene? ? Add information. Microarray? ? Add information. Alterations in gene expression by genetic or other treatment? ? Add information. Regulatory sequence features? ? Add information. Position frequency matrix (PFM) or position weight matrix (PWM)? ? Add information.

Reagents:

C. elegans antibodies? ? Add information. Integrated transgene? ? Add information. Transgene used for tissue markers? ? Add information.

Protein Function and Structure:

Protein analysis in vivo? ? Add information. Analysis of protein domains? ? Add information. Covalent modification? ? Add information. Structural information? ? Add information. Mass spectrometry? ? Add information.

Genome Sequence Data:

Gene structure correction? ? Add information. Sequencing mutant alleles? ? Add information. New SNPs, not already in WormBase? ? Add information.

Cell Data:

Ablation data? ? Add information. Cell function? ? Add information.

In Silico Data:

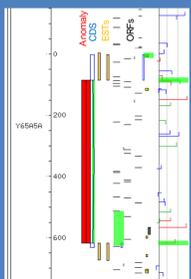
Phylogenetic analysis? ? Add information. Other bioinformatics analysis? ? Add information.

Other:

Authors will be automatically contacted once their paper is downloaded, at this point the paper will not be visible to the curators. Once a set period of time has elapsed the text mining will be conducted and work distributed within the consortium.

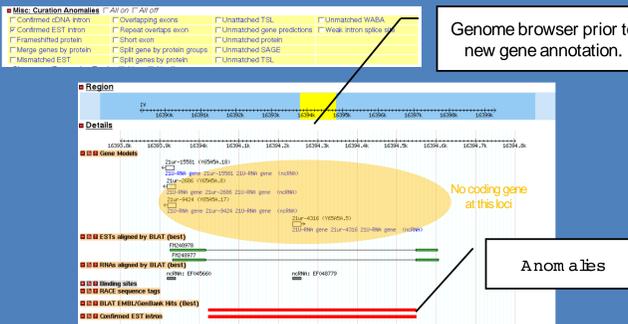
Curation Anomaly display

(see sequence curation poster for more details)



Curators use the Fmap within ACeDB to add/modify genes that go into the WormBase dataset. Curators use pre-computed anomalies to identify gene models that require attention as well as missing genes. Here we have an Fmap display with a Confirmed intron not in gene model anomaly. The curator created this two exon gene model based on the 2 EST sequences which contain a single intron.

Users can choose to see these anomalies/possible errors in the genome browser by selecting from these tracks.



Genome browser prior to new gene annotation.

Anom

Gene Summary Page: Concise descriptions

Curators are working to produce a manually generated concise description for all *C. elegans* genes (extending to include tier II nematodes). The aim of this is to produce an abstract like summary of the gene and its function so that WormBase users get a good understanding of the gene, with a minimum amount of effort.

Gene Summary for unc-13

Specify a gene using a gene name (unc-20), a predicted gene ID (R19A5.9), or a protein ID (CE0271.1) [unc-13]

Identification: **IDs:** unc-13 (L2/C026.0) via Person evidence: Jonathan Hodgkin **Sequence name:** ZK524.2 **WB Gene ID:** WBGene00006752

Concise Description: unc-13 encodes at least five protein isoforms that regulate neurotransmitter release by altering the conformation of syntaxin. UNC-13 proteins are required for normal physiological pumping and trapping in liquid, normally short lifespan, normally large blood sizes, and full adult body sizes. UNC-13 proteins have orthologs in vertebrates and Drosophila. UNC-13 proteins are complex, with multiple C2, prothol ester-binding, and DUF1541 domains. UNC-13 protein form is localized to most or all synapses; many of the unc-13 mutant alleles with viable phenotypes are transcript-specific, while homozygotes with an unc-13 null (deletion) allele die as paralyzed first-stage larvae. [details]

WormBook

WormBook is the online text companion to WormBase, the *C. elegans* model organism database. WormBook contains original reviews on all aspects of *C. elegans* biology and up-to-date descriptions of technical procedures used to study this animal.

WormBook Sections

- Genetics and genomics
- Developmental control
- Neurobiology and behavior
- Molecular biology
- Post-embryonic development
- Evolution and ecology
- Biochemistry
- Sex determination
- Disease models and drug discovery
- Cell biology
- The germ line
- WormMethods
- Signal transduction

Complete Chapter Listings By Section | By Publication Date

Phenotype ontology

Our Phenotype Ontology has been modified to curate nematode species other than *C. elegans* "N2" strain

Phenotype Ontology
A hierarchy-based ontology
1823 terms, 66% defined,
55% associated with a variation

Modifications required
Changes to Term Names
from "_abnormal" to "_variant".

Changes to Definitions
Use of "control animals" rather than "wild-type" or "N2" (*C. elegans* strain).

C. elegans-specific terminology, e.g., "hermaphrodite", were removed from definitions when possible.

Example:
WBPhenotype:0000037: egg_morphology_abnormal
Def: "Any deviation in the overall structure or appearance of fertilized oocytes that are deposited by adult hermaphrodites."

Changed to:
WBPhenotype:0000037: egg_morphology_variant
Def: "Any variation in the overall structure or appearance of fertilized oocytes that are laid compared to those laid by control animals." (The "abnormal" version of the term is kept as a synonym so people used to these terms will still be able to find them.)

Phenotype curation captures multiple attributes reported by authors and requires the efforts of many data curators

Phenotype increases curation accuracy and efficiency by use of ontologies and drop down lists.

COORDINATED WITH OBJECT CURATORS:
If object (allele or transgene) does not exist in the latest release of the database, an e-mail is automatically sent to the curator responsible for creating those objects.

COORDINATED WITH ONTOLOGY CURATOR:
Phenotype curators can request a term, send a suggested definition and hierarchy placement through the Phenote interface. New terms are automatically assigned to the record when they are approved.

OTHER ATTRIBUTES CAPTURED INCLUDE:
Genotype, Treatment, Nature of allele (recessive, semi-dominant, dominant), Penetrance (incomplete, low, high, complete), Maternal effect (strictly maternal, with maternal effect), Paternal effect, Temperature sensitivity, Haploinsufficiency, Allele type (amorph, hypomorph, etc.).

Phenotypes are linked to genes through allele or RNAi curation

Phenotypes reported as observed

Phenotypes reported as NOT observed

3626 / 23709* genes with alleles were annotated with phenotype data (includes NOT annotations) as of WS200

	May 2008 WS188	March 2009 WS200
Allele-phenotype connections (total # alleles)	9771	15951
Alleles Curated (total # alleles)	28% (15326)	34% (17448)
Papers curated (total papers flagged)	NA	23% (+125 unflagged papers)