Research

The Family of *Caenorhabditis elegans* Tyrosine Kinase Receptors: Similarities and Differences with Mammalian Receptors

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Transmembrane receptors with tyrosine kinase activity (RTK) constitute a superfamily of proteins present in all metazoans that is associated with the control and regulation of cellular processes. They have been the focus of numerous studies and are a good subject for comparative analyses of multigene families in different species aimed at understanding metazoan evolution. The sequence of the genome of the nematode worm *Caenorhabditis elegans* is available. This offers a good opportunity to study the superfamily of nematode RTKs in its entirety and to compare it with its mammalian counterpart. We show that the *C. elegans* RTKs constitute various groups with different phylogenetic relationships with mammalian RTKs. A group of four RTKs show structural similarity with the three mammalian receptors for the vascular endothelial growth factors. Another group comprises RTKs with a short extracellular region, a feature not known in mammals; the genes encoding these RTKs are clustered on chromosome II with other gene families, including genes encoding chitinase-like proteins. Most of the *C. elegans* RTKs have no direct orthologous relationship with any mammalian RTK, providing an illustration of the importance of the separate evolution of the different phyla.

[The sequences in this paper have been submitted to GenBank under the following accession numbers: AF188748, AF188749, AF188750, and AF188751.]

Understanding the processes that have governed metazoan genome evolution is an important issue of modern biology that is largely nourished by the wealth of data generated by genome projects devoted to the analysis of selected key species. Comparison of the data obtained in several species has so far led to the general belief that the number of genes per genome was doubled after the separation of Protostomia and Deuterostomia and multiplied by three or four times in the chordate phylum after the separation from echinoderms. Such an increase in gene number could be due to tetraploidization events, as first proposed by Ohno (1970). Accordingly, it is believed that the vertebrate ancestors had single copies of genes now found in multiple copies in vertebrates due to these polyploidization events (Schughart et al. 1989; Lundin 1993; Garcia-Fernández and Holland 1994; Holland et al. 1994; Holland and Garcia-Fernández 1996; Sidow 1996; Bailey et al. 1997; Coulier et al. 1997; Spring 1997; Pébusque et al. 1998). The number of genes found in vertebrates is roughly fourfold that found in nonvertebrates, and many single copy genes in nonvertebrates have a direct correspondence (orthology) with a gene family in vertebrates. Recent findings on fish ge-

nomes have indicated that many gene families comprise even more members than in mammals, again supporting large-scale duplications as the essential driving force toward diversity (Aparicio et al. 1997; Holland 1997; Amores et al. 1998; Finnerty and Martindale 1998; Postlethwait et al. 1998; Prince et al. 1998; Wittbrodt et al. 1998). To this, individual local duplications add some complexity for some gene families in which newly duplicated paralogs are positively selected and gradually increase in number (Kirschner and Gerhart 1998). These families with maximal evolvability are encoding molecules involved in environment-oriented defense and recognition processes such as immunoglobulin, major histocompatibility antigen, and olfactory receptor gene families.

Investigating various species of triploblastic metazoans will provide information about their common or specific evolution. In particular, the availability of the nematode worm *Caenorhabditis elegans* genome sequence allows for the first time the closest possible look at a present-day metazoan genome and putative transcriptome and proteome (The *C. elegans* Sequencing Consortium 1998).

We are particularly interested in examining multigene families in *C. elegans* and comparing them with the mammalian homologs. In this paper we have studied the tyrosine kinase receptors (RTKs). RTKs are type

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I transmembrane proteins that are involved in the control and regulation of several key cellular processes during development and in adult life. They are constituted of an extracellular ligand-binding region of variable size made of various domains, a transmembrane hydrophobic domain and an intracellular region bearing a kinase domain (TK), which is sometimes split in two parts (TK1 and TK2). Based on their structure and ligand-binding specificities, several classes of RTKs have been distinguished in mammals. In three classes, the extracellular region is made solely of immunoglobulin-like domains. To date, two groups of RTKs have been described in *C. elegans*: orthologs of RTKs known in mammals (e.g., DAF-2, EGL-15, KIN-8, LET-23, and VAB-1) (Table 1) and RTKs identified only in *C. elegans* (e.g., KIN-15 and KIN-16) (Morgan and Greenwald 1993). Some RTKs are absent in *C. elegans* (e.g., RET) but present in mammals and in *Drosophila melanogaster*, and others are present in mammals (e.g., PDGFR and VEGFR) but have not yet been described in the fruit fly nor in the worm. Few of the *C. elegans* RTKs have characterized or potential ligands (Hill and Sternberg 1992; Burdine et al. 1997). Conversely, some ligands have not yet been associated with receptors (Roubin et al. 1999).

RTKs are an excellent model for studying evolution of cellular processes in metazoans: First, they represent large families of modular proteins; Second, they are major representatives of the intercellular regulatory pathways that are specific of metazoans and Third, they belong to the most widespread contingency-generating agents of eukaryotic cells that are the protein kinases (Kirschner and Gerhart 1998). Furthermore, because tyrosine kinases have been under close scrutiny for many years in various species, they are associated with a wealth of information, and most of the genes encoding these proteins are probably identi-

fied in mammals; these molecules thus provide a good starting point for comparative studies of multigene families in different species.

We describe here the organization of 21 new *C. elegans* genes that encode putative RTKs phylogenetically related to various mammalian RTK classes.

RESULTS

Identification of RTK Genes by Exploration and Annotation of the *C. elegans* Sequence

The availability of the complete genomic sequence of the nematode *C. elegans* allowed the identification of new putative RTK genes following a keyword search on the Wormpep database and a TBLASTN search on the genomic sequence with either the entire sequence or the tyrosine kinase domain sequence of receptors belonging to 13 classes of RTK described in mammals (see Table 1). The protein sequences indexed in the databases as containing a tyrosine kinase domain were checked for the presence of this domain by SMART search. Only the sequences with a canonical tyrosine kinase domain (and not with a serine/threonine type or of undetermined specificity) were used. Prior to further analyses, we made some annotations with respect to the available sequences (see Methods).

Figures 1 and 2 show the distribution in the *C. elegans* genome of the genes cited in the text. A total of 28 protein sequences, including the corrected T17A3.1, T17A3.8, W04G5.6A, W04G5.6B, and R09D1.12 sequences (see Methods), were thus available for subsequent analysis (Tables 1 and 2).

Phylogenetic Analysis of RTKs in *C. elegans*

A phylogenetic tree based on the alignment of the tyrosine kinase domain of RTKs was constructed; the sequences used were those identified by the *C. elegans*

Table 1. RTK Classes in Mammals and *C. elegans*

Class ^a	RTKs in mammals	Orthologs in <i>C. elegans</i>
I	EGFR, ERBB (4 known)	LET-23
II	insulin receptor, IGF1-R	DAF-2
III (5 Ig) ^b	PDGFRs, CSF1R, KIT, FLT3	none
IV (3 Ig)	FGF receptors (4 known)	EGL-15
V (7 Ig)	VEGF receptors (3 known)	discussed here
VI	MET/HGFR, RON, SEA	F11E6.8 (discussed here)
VII (2 Ig)	TRK/NGFRs (3 known), ROR	KIN-8/CAM-1
VIII (1 Ig)	ephrin receptors (14 known)	VAB-1
IX	GDNF/NTN receptor (RET)	none
X (2 Ig)	TIE1, TIE2	none
XI (2 Ig)	AXL, TYRO3	none
XII	collagen (disoidin) receptors (2 known)	F11D5.3 (discussed here)
XIII	agrin receptor (MUSK)	none

^aThe class definition is taken from Ullrich and Schlessinger (1990) and Rosnet and Birnbaum (1993) and updated. The order of classes is arbitrary.

^bWhen present in the extracellular region, the number of immunoglobulin-like domains (Ig) is indicated.

Figure 1 Distribution of RTK genes (at right of the chromosomes) and chitinase genes (at left) in the *C. elegans* genome. The genetic distances are indicated on a scale bar at left.

Sequencing Consortium (1998) and already included in ACeDB and the modifications that we made (see Methods). Thus, in addition to seven characterized nematode RTKs (i.e., DAF-2, EGL-15, LET-23, VAB-1, KIN-8, KIN-15, and KIN-16), we included 21 new putative RTKs.

As seen in Figure 3, DAF-2, EGL-15, LET-23, and VAB-1 had the expected phylogenetic relationship with their known mammalian orthologs: the insulin and IGF1 receptors, the fibroblast growth factor receptors, the receptors of the EGF receptor/ERBB family, and the ephrin receptors, respectively (Table 1). KIN-8 and the F11D5.3 putative protein shared a common ancestor with the mammalian ROR, TRK, MUSK, and discoidin receptor families. F11E6.8 shared a common ancestor with the MET/RON receptor family. The mammalian RTKs from classes III, IV, and V (Ig RTKs) grouped together and with class IX and X RTKs.

Twenty-one known or predicted *C. elegans* RTKs did not share a recent common ancestor with mammalian RTKs, and some of them formed distinct phylogenetic families (Fig. 3, shaded boxes). A grouping of 13 RTKs (designated as LERF and SERF in Fig. 3; de-

scribed below) was supported by a bootstrap value of >900/1000. Among these 13 RTKs were the KIN-15 and KIN-16 kinases (Morgan and Greenwald 1993). Four RTKs (F59F3.5, F59F3.1, T17A3.8, and T17A3.1) constituted a distinct monophyletic family (bootstrap value of >700). Four other RTKs (F09A5.2, F09G2.1, F08F1.1, and C24G6.2) grouped together (bootstrap value of >900).

Structural Features and Organization of *C. elegans* RTKs

Based on structural features, the 28 RTKs encoded by these known or predicted sequences could be divided into two groups.

In the first group, we classified the seven known or potential orthologs of mammalian RTKs (DAF-2, EGL-15, LET-23, VAB-1, KIN-8, F11D5.3, and F11E6.8). Based on a similar domain architecture and the phylogenetic analysis, F11D5.3 (Table 1) may be considered as the potential ortholog of collagen receptors DDR and TKT, RTKs with factor V-factor VIII domains in their extracellular region, also called discoidin-type

Figure 2 Details of chromosome localization of predicted *C. elegans* RTK genes described in this paper. Symbols are as follows (black arrows) RTK genes; (black arrowheads) RTK pseudogenes; (gray arrows) chitinase genes; (gray arrowheads) other genes. (A) clustering of genes encoding known and putative RTKs with a short extracellular region and chitinase-like genes on chromosome II. (B) Predicted RTK genes on chromosome III. (C) Predicted RTK and chitinase-like genes on chromosome V. (D) Predicted and known RTK genes on chromosome X.

Table 2. Known and Putative *C. elegans* RTK Genes and Deduced Proteins

	Wormpep	GeneBank Acc	Gene name	Length	Protein			Similarity to	Expression ¹	Chromosome
					SP	TM	TK			
S	-	AF012437	<i>daf-2</i>	1846 aa	+	1184-1204	1246-1520	Insulin receptor family	cloned	III
	-	U39761	<i>egl-15</i>	1040 aa	+	529-549	640-929	FGF receptor family	cloned	X
	-	Z35595	<i>kin-8</i>	806 aa	-	348-368	468-735	ROR receptor	cloned	II
	-	P24348	<i>let-23</i>	1323 aa	+	819-841	885-1152	EGF receptor/ERBB family	cloned	II
	-	AF040269	<i>vab-1</i>	1122 aa	-	559-579	692-984	EPH receptor family	cloned	II
	F11D5.3	AAA83276	-	791 aa	-	377-397	521-779	Discoidin receptor family	EST	X
	F11E6.8	CAB02923	-	442 aa	-	49-69	136-399	MET TK domain	EST	IV
S	-	P34891	<i>kin-15</i>	484 aa	+	46-66	140-448	Tyrosine kinase receptor	cloned	II
	-	P34892	<i>kin-16</i>	495 aa	+	61-81	140-454	Tyrosine kinase receptor	cloned	II
	C08H9.5	CAA91146	-	502 aa	-	75-95	175-461	KIN-15 and KIN-16 TK domain	NP	II
	M01B2.1	CAB05559	-	483 aa	-	27-47	133-454	KIN-15 and KIN-16 TK domain	NP	V
	R09D1.12	CAA93871	-	474 aa ²	-	63-83	160-435	KIN-15 and KIN-16 TK domain	RT-PCR	II
	R09D1.13	CAA93872	-	388 aa	+	-	80-360	KIN-15 and KIN-16 TK domain	NP	II
	W04G5.6A ⁴	CAB07686 ⁵	-	?	?	?	?	KIN-15 and KIN-16 TK domain	NP	I
	W04G5.6B ⁴	-	-	?	?	?	?	KIN-15 and KIN-16 TK domain	NP	I
	ZK938.5	CAA90147	-	498 aa	+	54-74	171-457	KIN-15 and KIN-16 TK domain	NP	II
	F59F3.1	CAA91990	-	1227 aa	+	765-785	847-1163	Ig superfamily and KIN-15 and KIN-16 TK domain	EST	X
E	F59F3.5	CAA91991	-	1199 aa	+	771-791	853-1158	Ig superfamily and KIN-15 and KIN-16 TK domain	RT-PCR	X
	T17A3.1	AAC26948	-	1083 aa ²	+	739-756	803-1072	Ig superfamily and KIN-15 and KIN-16 TK domain	RT-PCR	III
	T17A3.8	AAC26954	-	649 aa ³	-	90-110	165-407	Ig superfamily and KIN-15 and KIN-16 TK domain	NP	III
	C16D9.2	AAB18281	-	2380 aa	+	1836-1856	1865-2131	-	EST	V
O	C24G6.2 ⁶	AAC19210	-	903 aa	+	475-495	543-841	Fibronectin type III domains	EST	V
	F08F1.1	AAB71301	-	478 aa	-	40-60	166-445	-	NP	X
H	F09A5.2	CAA93646	-	894 aa	-	355-375	464-782	-	EST	X
E	F09G2.1	AAB71320	-	929 aa	-	449-469	487-771	-	EST	V
R	R151.4	AAA50850	-	276 aa	-	-	2-256	-	EST	II
S	T10H9.2	AAC19236	-	1414 aa	+	899-919	969-1234	Low-density lipoprotein receptor class A domain	EST	V
	T14E8.1	AAA82385	-	1096 aa	-	652-672	735-1003	-	EST	X

(SP) signal peptide; (TM) transmembrane domain; (TK) tyrosine kinase domain; (ND) not determined; (SERF) short extracellular region family; (LERF) long extracellular region family.

¹(Cloned) Complete cDNA sequence is found in databases; (EST) partial sequence is found in databases; (RT-PCR) we showed the existence of mRNA by RT-PCR; (NP) no proof of mRNA exist.

²Corrected after RT-PCR study.

³As predicted by GENEFINDER.

⁴Sequences derived from W04G5.6 predicted protein.

⁵Accession no. for W04G5.6.

⁶See Methods.

RTKs. The sequence of the tyrosine kinase domain of F11E6.8 (Table 1) is very similar to that of the tyrosine kinase domain of class VI RTKs; however, the predicted protein is very short, with an extracellular region of only 40 amino acids lacking a signal peptide. Search in the genomic sequence did not reveal any similarity with the extracellular region of MET family of RTKs. We considered this putative protein as a wrong GENE-FINDER prediction and a potential ortholog of class VI RTKs. No significant match and no modular architecture corresponding to RTKs from the other classes were found (Table 1).

Twenty-one predicted or characterized proteins with no extensive similarity to a particular class of mammalian RTKs constitute the second group. When their sequence and domain composition were compared, three different types could be identified (Table 2; Fig. 3).

1. One type contains two already characterized proteins, KIN-15 and KIN-16 (Morgan and Greenwald 1993), and seven other proteins. The extracellular region of KIN-15 and KIN-16 is very short (between 40 and 50 amino acids). We will hereafter refer to this family as the "short extracellular region" family ("SERF"). Such an unusual structure for a RTK has not been described in mammals. The other seven SERF proteins (C08H9.5, M01B2.1, R09D1.12, R09D1.13, W04G5.6A, W04G5.6B, and ZK938.5) have not yet been characterized in *C. elegans*. Thus, a total of nine putative RTKs belong to this family. By using the PSORT II and SMART tools, which give information on various sequence features related to protein sorting signal, we identified a putative peptide signal and a transmembrane region in the majority of the putative proteins belonging to SERF.
2. A second type is composed of four putative mol-

Figure 3 Phylogenetic analysis of known and predicted tyrosine kinase receptors of *C. elegans* based on tyrosine kinase domain alignment. Proteins constituting monophyletic families (LERF, SERF, and four other RTKs) are indicated by shaded boxes. Symbols indicate the bootstrap value [(●) >500; (▲) >700; (◆) >900; 1000 replicates were done; statistical significance is reached with a value of 700]. The sequence of the human nonreceptor type tyrosine kinase ABL1 (underlined) is used as outgroup. Human sequences are with an asterisk (*), and accession nos. are as follows: ABL1 (X16416); DDR (Q08345); EGFR (P00533); EPHA1 (M18391); FGFR1 (M63888); INSR (M32972); MET (P08581); MUSK (AF006464); PDGFRA (P16234); RET (P07949); RON (X70040); ROR1 (M97675); ROS (M34353); TIE1 (X60957); TRKA (M23102); TYRO3 (P55146); UFO (P30530); VEGFR1 (X51602). A phylogenetic tree including more sequences can be found on our web site (<http://olan.marseille.inserm.fr/u119/home.html>).

ecules (F59F3.1, F59F3.5, T17A3.1, and T17A3.8) containing an extracellular region of ~850 amino acid residues. They will be referred to as the “long extracellular region” family (“LERF”). In the protein sequence of LERF RTKs, except in T17A3.8 that contains an erroneous GENEFINDER prediction for the extracellular region (see Methods), six immunoglobulin-like domains were identified by Pfam analysis using the fragment search option (we will see later that they actually have seven immunoglobulin-like domains, but the fourth one is distinct). Their tyrosine kinase domain was found to be bipartite, with 40–50 amino acid residues separating two subdomains.

3. A third type was represented by eight predictions that are very heterogeneous in their extracellular region (Table 2). Some predictions, such as F08F1.1, have a very short extracellular region lacking a signal peptide and may represent a wrong GENEFINDER prediction. Some others (F09A5.2, F09G2.1, R151.4, and T14E8.1) have a large extracellular region but do not contain any domain described to date. Two other protein predictions have in their extracellular region one (C24G6.2) or three (C16D9.2) fibronectin type III domains. The T10H9.2 protein prediction contains a low-density lipoprotein-a (LDL-a) domain, never described in mammalian RTKs.

C. elegans Putative RTKs with LERF Resemble Mammalian VEGFRs

The sequences of the extracellular region of the proteins with the LERF were aligned with the different classes of RTK containing five or seven immunoglobulin-

like domains. The highest percentage of identity was obtained in the alignment with the extracellular regions of class V RTKs. These Ig RTKs are the three VEGFRs (Mustonen and Alitalo 1995; Fournier et al. 1997); their ligands are the vascular endothelial growth factors (Fig. 4). A remarkable conservation of the cysteines involved in the immunoglobulin-like disulfide bonds of the VEGFRs is seen in this alignment (not shown). Moreover, a region highly similar to the related fourth immunoglobulin-like domains of VEGFRs, which lacks a disulfide bond, is also seen in an equivalent location in the putative proteins of the LERF family. Some positions of this immunoglobulin-like domain containing aromatic amino acids are conserved among *C. elegans* and mammals. The immunoglobulin-like domains with disulfide bonds in the VEGFRs are structurally constituted by seven β -sheets and belong to the C2-type domain (Bork et al. 1994). Like in the mammalian VEGFRs, all immunoglobulin-like domains except the fourth one are made of seven β -sheets in the predicted secondary structure of the *C. elegans* LERF RTKs.

RTKs are characterized by the presence of a highly hydrophobic transmembrane domain made of an α -helix. In the LERF RTKs, similar regions with highly hydrophobic α -helices were identified upstream of the tyrosine kinase domain in all members and at equivalent positions (Table 2). Like the kinase domain of mammalian class III, IV, and V RTKs, the catalytic domain of the predicted LERF proteins is split by a kinase insert. Its size is 40–50 amino acids, different from the

kinase insert of class IV RTKs (15 amino acids) and class III and V RTKs (70 amino acids).

Genomic Distribution of the Genes Encoding the New Predicted RTKs

The four genes encoding the LERF RTKs are located in tandem on either chromosome III or X (Fig. 2B, D). T17A3.1 and T17A3.8, the two genes from chromosome III (Fig. 2B), are oriented in an opposite direction, 6.5 kb apart. On chromosome X, F59F3.1 and F59F3.5 are in the same transcriptional orientation; no putative gene has been evidenced in the 4.5-kb region separating these genes by GENEFINDER (Fig. 2D).

The genes encoding the SERF RTKs are mostly found on chromosome II in two clusters separated by 0.5 Mb (Fig. 2A). One cluster, mapping at the approximate genetic position 0.99 and covering the M176 and R09D1 cosmids, includes the *kin-15*, *kin-16*, R09D1.13, and R09D1.12 genes, whereas the second cluster, at genetic map position 1.47 and covering the ZK938 and C08H9 cosmids, includes the ZK938.5, C08H9.5, and C08H9.8 genes. Three genes of this type are not located on chromosome II: W04G5.6A and W04G5.6B, located on chromosome I (Fig. 1), and M01B2.1, located on chromosome V (Fig. 2C).

A major feature of the RTK genes located in two clusters on chromosome II is their tandem organization. In the first cluster, the *kin-15* and *kin-16* coding sequences are separated by 529 bp and are transcribed unidirectionally (Fig. 2A). Another pair of genes, R09D1.13 and R09D1.12, are also transcribed in the same direction but are 7.8 kb apart. In the second cluster, the ZK938.5 and C08H9.8 RTK genes are also organized in tandem. Similarly, but on chromosome I, the W04G5.6A and W04G5.6B genes are in the same transcriptional orientation and separated by only 0.7 kb.

Figure 4 (Left) A schematic representation of the vascular endothelial growth factor receptors (class V RTKs) and the four *C. elegans* RTKs of the long extracellular region type; (Ig) Immunoglobulin-like domain (cysteine residues are shown), (TM) transmembrane domain, (TK) tyrosine kinase domain. (Right) The percentages of similar amino acids shared by these RTKs.

Most of the *C. elegans* RTK Sequences Are Transcribed

To determine roughly how many of the predicted RTK genes correspond to expressed genes, we looked for the presence of ESTs in the DNA Data Bank of Japan. Seventeen genes have corresponding ESTs or complete cDNAs in this database (Table 2). We looked by means of reverse transcription–polymerase chain reaction (RT–PCR) for expression of the RTK genes of the short and long ex-

tracellular region families that do not have a corresponding EST or cDNA in the database. In addition to F59F3.1, for which ESTs exist in the database, we found that two other LERF genes (F59F3.5 and T17A3.1) and one SERF gene (R09D1.12) are expressed. Thus, a minimum of 20 RTK-encoding genes from a total of 28 putative genes have transcribed RNA.

Other Multigenic Families Are Located on Chromosome II

The two clusters on chromosome II containing genes predicted to encode RTKs also contain a large number of genes encoding chitinases (on cosmids M176, R09D1, and T19H5—genetic map position 0.99—and cosmids ZK938 and C08H9—genetic position 1.47) (Fig. 2A); guanylate cyclases (predictions AH6.1, R134.1, and R134.2 at approximate genetic position

1.01), olfactory receptors (from *sra-1* to *sra-9* on the AH6 cosmid, also at approximate genetic position 1.01), and cytochrome P450 molecules (one member on cosmid ZK1320 at approximate genetic position 1.13 and eight members on cosmid T10B9 at position 1.41) are also present in this area. A tyrosine kinase–chitinase-like genes association was also found on chromosome V. One gene coding for a SERF RTK (M01B2.1) and two genes coding for chitinase-like proteins (K08F9.3 and M01B2.6) are located very closely, in a region of chromosome V covered by 0.5 genetic map units.

The most prominent example for repetitive duplications of transcribed sequences in this region is the family of chitinases. Among the 40 chitinase genes or predicted chitinase-like genes (Table 3), 25 are located on chromosome II along with SERF genes. Three other

Table 3. Chitinase Family in *C. elegans*

Wormpep	GenBank acc. no.	Length (aa)	Similarity with proteins of chitinase family	Chromosome	EST
C04F6.3	Q11174	617	glycosyl hydrolase family 18	X	+
C08B6.4	CAA96587	447	chitinase class IA	V	–
C08H9.4	CAA91151	465	glycosyl hydrolase family 18	II	–
C08H9.6	CAA91154	394	glycosyl hydrolase family 18	II	–
C08H9.7	CAA91150	447	glycosyl hydrolase family 18	II	–
C08H9.10	CAA91147	457	glycosyl hydrolase family 18	II	–
C08H9.11	CAA91148	459	glycosyl hydrolase family 18	II	–
C08H9.12	CAA91149	460	glycosyl hydrolase family 18	II	–
C08H9.13	CAA91152	459	glycosyl hydrolase family 18	II	–
C08H9.14	CAA91153	339	glycosyl hydrolase family 18	II	–
C45E5.2	AAC24288	469	glycosyl hydrolase family 18	IV	–
C45E5.3	AAC24286	266	glycosyl hydrolase family 18	IV	–
C51E3.8	CAB01643	272	chitinase class IA	V	–
F07G11.9 ^a	AAB65302	2025	glycosyl hydrolase family 18	V	–
F10G2.5 ^a	AAB04826	1614	<i>Trichoderma harzianum</i> endochitinase (family 18)	V	–
F15A4.8	CAB02942	755	glycosyl hydrolase family 18	II	–
K08F9.3	CAB03188	328	glycosyl hydrolase family 18	V	–
M176.8	CAB01653	400	glycosyl hydrolase family 18	II	–
M01B2.6	CAB05568	479	glycosyl hydrolase family 18	V	–
R09D1.1	CAA93861	432	glycosyl hydrolase family 18	II	–
R09D1.2	CAA93862	440	glycosyl hydrolase family 18	II	–
R09D1.3	CAA93863	435	glycosyl hydrolase family 18	II	–
R09D1.4	CAA93864	384	glycosyl hydrolase family 18	II	–
R09D1.5	CAA93865	444	glycosyl hydrolase family 18	II	–
R09D1.6	CAA93866	444	glycosyl hydrolase family 18	II	–
R09D1.7	CAA93867	369	glycosyl hydrolase family 18	II	–
R09D1.8	CAA93868	404	glycosyl hydrolase family 18	II	–
R09D1.9	CAA93869	216	glycosyl hydrolase family 18	II	+
R09D1.9a	CAA93873	185	probable pseudogene (family 18)	II	–
R09D1.10	CAA93870	380	glycosyl hydrolase family 18	II	–
R09D1.11	CAA93874	432	glycosyl hydrolase family 18	II	–
R10D12.15	CAB03255	453	chitinase class IA	V	–
T01C4.1	AAB09177	1484	glycosyl hydrolase family 18	V	–
T05H4.7	AAB66013	511	chitinase class IA	V	–
T10D4.3	AAC68716	410	family 18 and cartilage glycoprotein YKL39	II	–
T13H5.3	CAA91419	633	glycosyl hydrolase family 18	II	+
T19H5.1	CAA87665	764	glycosyl hydrolase family 18	II	–
T19H5.2	CAA87663	358	glycosyl hydrolase family 18	II	–
T19H5.3	CAA87664	407	glycosyl hydrolase family 18	II	–
ZK938.6	CAA90145	522	glycosyl hydrolase family 18	II	–

^aTwo different GENEFINDER predictions for the same genomic region.

chitinase-like genes, T10D4.3, F15A4.8, and T13H5.3, map on chromosome II but far away from the tyrosine kinase–chitinase-like clusters, at position –20.28,

0.24, and 16.97, respectively (Fig. 1). The 12 other chitinase-like genes are located elsewhere in the genome: two on chromosome IV (C45E5.2 and C45E5.3 at the approximate genetic map position 23.98), nine on chromosome V (T05H4.7 located at the approximate genetic map position –1.11; T01C4.1 at 0.62; F07G11.9 and F10G2.5, representing two different predictions for the same genomic region, between 0.63 and 0.64; C08B6.4 and C51E3.8 at 3.18–3.19; R10D12.15 at 10.47; K08F9.3 at 20.47; and M01B2.6 at 21.11), and one on chromosome X (C04F6.3) (Fig. 1).

Based on both amino acid sequence differences and tridimensional structure (Henrissat 1990; Davies and Henrissat 1995; Henrissat and Romeu 1995), chitinases constitute two families of glycosyl hydrolases: family 18 (which contains chitinases from viruses, bacteria, fungi, animals, and classes III and V from plants), and family 19 (chitinase classes I, II, and IV from plants). Sequence analysis of the 40 chitinase or chitinase-like coding genes showed that they encode two types of proteins: a very large number of predicted proteins similar to chitinases of class II (family 18) and four predictions (K08F9.3, T05H4.7, R10D12.15, C08B6.4) similar to chitinase class IA. We conducted a phylogenetic analysis of these different chitinases. As shown in Figure 5, the chitinase-like proteins encoded by genes located in clusters on chromosome II shared a recent common ancestor (chromosome II, shaded boxes). The chitinase genes from class IA located on chromosome V were also grouped in one branch.

Figure 5 Phylogenetic analysis of chitinase-like proteins of *C. elegans*. Sequences analyzed in this tree are identical to those listed in Table 3 except for eight sequences that show an incomplete chitinase domain and have not been used in the phylogenetic analysis. Chromosomal localizations of *C. elegans* chitinase genes are indicated at right; the labeled shaded box indicate the two clusters localized on chromosome II (see Fig. 1). Symbols for bootstrap values are as in Fig. 3. The sequence X78325 (underlined) of *Nicotiana tabacum* is used as outgroup. Species abbreviations used for the phylogenetic tree are as follows: (Aea) *Aedes aegypti* (yellow fever mosquito); (Aga) *Anopheles gambiae* (african malaria mosquito); (Avi) *Acanthocheilonema viteae* (animal filarial nematode); (Bmo) *Bombyx mori* (silkworm); (Bta) *Bos taurus*; (Bma) *Brugia malayi* (agent of Brugian lymphatic filariasis); (Bpa) *Brugia pahangi* (filarial nematode of domestic cats); (Csp) *Chelonus sp.* (braconid wasp); (Cte) *Chironomus tentans* (nonbiting midges); (Dme) *Drosophila melanogaster* (fruit fly); (Hcu) *Hyphantria cunea* (fall webworm); (Hsa) *Homo sapiens* (human); (Mse) *Manduca sexta* (tobacco hornworm); (Mmu) *Mus musculus* (mouse); (Nta) *Nicotiana tabacum* (common tobacco); (Ovo) *Onchocerca volvulus* (agent of river blindness); (Pco) *Phaedon cochleariae* (mustard beetle); (Pja) *Panaeus japonicus* (Kuruma prawn); (Wba) *Wuchereria bancrofti* (agent of lymphatic filariasis).

DISCUSSION

Identification of Novel RTKs in *C. elegans*

A combined search of the *C. el-*

egans genome conducted along key words and through alignment (BLAST) allowed the identification of 28 putative protein-coding sequences with similarity to RTKs. Seven of these sequences were known RTKs (DAF-2, EGL-15, KIN-8, KIN-15, KIN-16, LET-23, and VAB-1) and 21 were new putative RTKs. Among the 28 RTKs, a first group of seven proteins is represented by the *C. elegans* orthologs of mammalian RTKs (Table 1); two protein predictions (F11D5.3 and F11E6.8) identified in this study represent candidate orthologs of mammalian discoidin receptors and receptors of the MET family, respectively.

Among a second group of 21 putative proteins, eight are very heterogeneous in their extracellular region; they lack similarity with any extracellular domain described to date or contain domains never described in mammalian RTKs, such as an LDL-a domain (Table 2). A subgroup of 13 putative RTKs was subdivided into two types according to the structure of their non-tyrosine kinase portion: Nine molecules containing a SERF were of the first type, and four putative RTKs containing a LERF were of the second type. Together they constitute a separate subgroup that does not share a recent common ancestor with any particular mammalian RTK class. It cannot be excluded that this is due to a rapid evolution of the genes, as is thought to be the case for two-thirds of protein-coding genes in *C. elegans* (Mushegian et al. 1998; Roberston 1998), even if the tree topology does not support it.

SERF RTKs have only 20–80 amino acids in their extracellular region. The previously described (Morgan and Greenwald 1993) KIN-15 and KIN-16 molecules, as well as seven other members, belong to this group. The function of KIN-15, KIN-16, and R09D1.12, for which the presence of messenger RNA was found, is unknown.

None of the putative LERF proteins has been described in *C. elegans*. They resemble the three mammalian VEGFRs. Like the VEGFRs, they contain an extracellular region composed of seven type C Ig domains (the fourth being a pseudo domain because it lacks the typical cysteine residues), a transmembrane domain, and an intracellular region composed of two tyrosine kinase subdomains separated by a kinase insert. From the phylogenetic tree built on the alignment of the tyrosine kinase domains, it seems that the two RTK types (LERF and SERF) described here have evolved very closely (Fig. 3). The loss of most of the extracellular domain from a seven immunoglobulin-like ancestor may have been the mechanism of apparition of the SERF RTKs. Further local duplications in *cis* or *trans* would have allowed the growth of this family up to nine members. The limited length of the branches and a high bootstrap value found in the phylogenetic analysis suggest that some of these duplications could be recent (C08H9.5 and ZK938.5 genes; see Fig. 3).

RTKs Resembling Mammalian VEGFRs Exist in *C. elegans*

Up to now, VEGFRs have only been described in vertebrates. Such molecules have not yet been recognized in Mollusca and Arthropoda; however, a genome from these phyla has not yet been completely sequenced.

In mammals, the three VEGFRs described to date participate in the development and architectural organization of both the circulatory and blood systems during embryogenesis and are involved in the regulation of cell permeability (Mustonen and Alitalo 1995). The existence of four molecules in the worm, which does not possess per se a circulatory system, is intriguing but may be a simple example of how the same molecules can be adapted to different roles in different species while performing the same basic processes. Chemotaxis, cell migrations, and pervasive growth are important processes of angiogenesis. The LERF receptors may be similarly involved in these processes in the worm, even if the cells in which they are expressed have a different differentiation status. It is also possible that the identification of the function of potential VEGFRs in *C. elegans* may point to new functions for this class of RTKs in vertebrates. We did not find any *C. elegans* sequence with significant similarity with any of the six known mammalian VEGFs. The identification of potential ligands of the RTKs with a LERF may provide some clues about their function and possible related function in mammals.

Possible Scheme of Evolution of RTKs in the *C. elegans* Lineage

Because of the remarkable conservation of domain architecture between *C. elegans* LERF RTKs and mammalian VEGFRs, convergent evolution may not be the cause of *C. elegans* VEGFR-like existence, although it cannot be ruled out. Based on the phylogenetic analysis and chromosome localization, we propose that the four LERF RTKs derive from a common ancestor through a series of *cis*, then *trans* and finally, *cis* duplications. They do not group with the mammalian VEGFRs. This is in favor of an independent expansion of these families after the separation from the last common ancestor of Protostomia and Deuterostomia. The *egl-15* gene, encoding an FGFR, is, like F59F3.1 and F59F3.5, located on an overlapping cosmid on chromosome X, at position 15.79 (Fig. 2D). This suggests that the original linkage group that contained duplicated Ig RTK genes may have been maintained in the nematode lineage, whereas it has been disrupted in mammals where genes encoding class IV, on the one hand, and class III and VI RTKs, on the other hand, are separated (Rosnet and Birnbaum 1993; Pébusque et al. 1998).

The nine genes coding for the SERF RTKs are clustered on chromosomes I, II (in close proximity of chiti-

nase-like genes), and V. From the relationships of these genes and their genome distribution, we propose a scenario of evolution implying large-scale duplications and local duplications. We hypothesize that a couple of tyrosine kinase–chitinase genes were represented in an “ancestral” linkage group that represented the precursor of a tyrosine kinase–chitinase-like cluster located on chromosome II. A second precursor cluster was generated in the close proximity of the first. Successive local duplications in *cis* allowed the expansion of tyrosine kinase and chitinase-like genes. This separate evolution of each cluster on chromosome II by the means of an earlier duplication of an ancestral linkage group followed by local amplification of genes is suggested by the topology of the trees of chitinase-like and tyrosine kinase genes. The phylogenetic analysis of the chitinases-like protein sequences revealed that the molecules located on each of the two chromosome II clusters form a separate branch and that both branches are grouped separately from the other chitinase genes of different species. Among the chitinase-like genes located on the other chromosomes, only M01B2.6 (on the same cosmid as M01B2.1 SERF RTK gene) branches with the genes of one chromosome II cluster supporting the hypothesis of a partial *trans* duplication.

For chitinase-like proteins as for kinases, some duplication events could be recent, suggesting that the concerted expansion process may be ongoing. The presence of genes encoding olfactory receptors in the same clusters is interesting. Robertson (1998) has previously suggested an ongoing process of rapid evolution of olfactory receptor genes with extensive gene duplications, movement, and diversification. We propose below two nonexclusive types of possible explanation for this expansion.

Chitinases are glycosyl hydrolases that catalyze the degradation of chitin, a polymer of *N*-acetylglucosamine. In plants and insects, they are an important inductive host defense mechanism against fungi. It could be that chitinase-like proteins from *C. elegans* are also implicated in defense against parasites. In this case, the expansion of the chitinase family would be dependent on a positive selection mechanism, and passive “hitchhiking” by this mechanism may explain the combined expansion of RTK genes. However, some predictions for chitinase-like sequences yield proteins that lack the chitin-binding domain or the catalytic domain, and, at present, their role remains unknown.

A large region containing multiple copies of RcC9, RcD1, Rc35, and Rc123 DNA repetitive elements is located between the two clusters of tyrosine kinase–chitinase genes from chromosome II. These elements, showing an internal sequence organization resembling that of minisatellite sequences described in mammals, have a tendency to cluster at some positions on the

chromosomes of *C. elegans* (Naclerio et al. 1992). The suprapericentric patterns, which are different in the closely related species of *C. elegans*, *Caenorhabditis briggsae*, and *Rhabditis maupasi*, are constant between arrays with different chromosomal locations suggesting an active mechanism of homogenization (crossing-over, gene conversion) (Naclerio et al. 1992; La Volpe 1994). Because in human cell lines, minisatellite sequences promote homologous recombination (Wahls et al. 1990), an increased rate of crossing-over promoted by the presence of repeated sequences, associated with possible slippage in the homologous pairing due to lower complexity of the region containing the repeated sequences, might have resulted in the progressive increase of copies number of genes surrounding the region. An example of gene expansion associated with a particular type of complex satellite structures has been shown in the human genome (Eichler et al. 1998).

Conclusions

The identification and comparison of orthologs of human proteins in the nematode may yield interesting clues as to the respective functions of proteins in the two species. Furthermore, the particular organization of genes in *C. elegans*, such as kinase and chitinase genes, may provide clues as to the mechanisms that have driven their typical, rapid evolution by duplication and, thus, mechanisms that influence evolution. These may be associated with positive selection as the resulting proteins could be involved in defense mechanisms or with instability of a genomic region due to repetitive elements. Both mechanisms may have worked together because evolutionary pressure may have favored expansion over contraction of the region.

In metazoan evolution, each lineage, having evolved independently over long periods of time, shows some extent of specificity with regard to genome organization (Ruddle 1997). Independent and specific evolution may be frequent. A good example is that of *HOX* genes in teleosts (Aparicio et al. 1997; Amores et al. 1998). We show here an example of independent evolution in a multigene family of RTKs in the *C. elegans* lineage. Examples in *C. elegans* will certainly be numerous, and several have been reported already (Robertson 1998; Jansen et al. 1999; Sluder et al. 1999). This shows that this type of evolution concerns various types of genes and various types of species. In *C. elegans*, the apparently rapid processes of genetic duplications and movements within the genome may greatly favor this independent evolution.

METHODS

Database Searches

To identify putative RTKs in *C. elegans*, two approaches were

used. First, a search in the last release (no. 16) of Wormpep database using different combinations of the key words =el-gans=, =tyrosine=, =kinase=, and =receptor= was done by means of the Entrez search tool (<http://www.ncbi.nlm.nih.gov/Entrez/>). We thus identified the GENEFINDER predictions indexed as putative RTK or only as tyrosine kinases. Second, a *C. elegans*-specific TBLASTN search (http://www.sanger.ac.uk/Projects/C_elegans/blast_server.shtml) using tyrosine kinase domain sequences was used for the identification of some genomic regions coding for such a domain and missed by GENEFINDER.

The sequences recovered by the two approaches larger than 400 amino acids were further analyzed for motif or domain composition using Pfam (<http://www.sanger.ac.uk/Software/Pfam/>) and SMART (<http://coot.embl-heidelberg.de/SMART/>) tools. This eliminated some serine/threonine kinases or undetermined proteins that were recovered from the initial searches. Then, to eliminate the potential nonreceptor type tyrosine kinases, a search in the PSORT II program (<http://psort.nibb.ac.jp>) was done. Two sequences (CO8H9.8 and T01G5.1) lacking a consensus ATP-binding site were discarded. For the C24G6.2 gene, there are two alternative predictions of putative transcripts, C24G6.2A (coding for a protein with a large extracellular region with fibronectin type III domains) and G24G6.2B (coding for a protein with a short extracellular region lacking any described domain).

The information for cosmids, genes, and gene products (Wormpep database) of *C. elegans* is available through the ACeDB database (http://www.sanger.ac.uk/Projects/C_elegans/webace_front_end.shtml).

Searches for expressed sequence were done in the DNA Data Bank of Japan maintained by Yuji Kohara (<http://www.ddbj.nig.ac.jp/>).

Sequence Analysis and Alignment

Sequence similarity searches were done using the BLASTP and TBLASTN algorithms (Altschul et al. 1990, 1997). The accession numbers for sequences in mammals and *C. elegans* are listed in the legend to Figure 3 and in Tables 2 and 3, respectively. The protein sequences were aligned using the CLUSTAL W (Thompson et al. 1994) program followed by manual edition. The protein sequence analysis and the alignments were confirmed by searches with the PSORT II (<http://psort.nibb.ac.jp>), Pfam, and SMART tools. Alignments were done with the entire protein sequences or after partitioning into distinct domains.

Annotations

Annotation 1

The T17A3.1 predicted protein contains a truncated tyrosine kinase domain (with the TK1 domain only). A region presenting a significant similarity to both kinase insert (KI) and TK2 domain, referred to as F40G9.13 gene by GENEFINDER, was identified downstream of the putative stop codon of the predicted T17A3.1, in the neighboring F40G9 cosmid (see Fig. 2B). We considered T17A3.1 as an erroneous prediction and corrected the protein sequence by the addition of the missing KI and TK2 region identified on cosmid F40G9. A short region presenting a weak similarity with the extracellular region of F59F3.1 was also identified on cosmid F40G9 but in an opposite orientation to the T17A3.1 transcript. We considered this region lacking a GENEFINDER prediction as a pseudogene.

Annotation 2

In the region of the T17A3.8 transcript, we identified two short sequences with high similarities to the one encoding the putative extracellular region of F59F3.1, which were not predicted by GENEFINDER. We corrected the deduced protein sequence of T17A3.8 by including these two sequences. We identified by BLAST search on the genomic sequences an exon not predicted by GENEFINDER coding for a region downstream of the ATP-binding site. On the same cosmid, the gene prediction T17A3.10, which is oriented in the same direction as T17A3.8 (see Fig. 2B), has a similarity with the region contained between amino acid positions 450 and 850 of putative RTKs F59F3.1 and F59F3.5. It was considered as a probable pseudogene because we were unable to identify upstream and downstream sequences with similarity to the remaining of an RTK molecule.

Annotation 3

When we used the Pfam module recognition tool, we found that one of the genes, namely W04G5.6, contained sequences encoding two tyrosine kinase domains that, when analyzed by TBLASTN, happened to be very similar. We considered these two regions as belonging to two different transcripts transcribed in the same direction that were missed by the GENEFINDER prediction. They will hereafter be referred to as W04G5.6A and W04G5.6B (see Fig. 1).

Annotation 4

Sequence analysis of the cDNA fragment amplified by RT-PCR using R09D1.12 primer pair showed that exon four of the R09D1.12 gene is 75 bp longer in its 3' end than predicted. The corrected protein sequence, by addition of 25 amino acids in the tyrosine kinase domain, was used for phylogenetic analysis.

Phylogenetic Analyses

Phylogenetic trees were inferred using neighbor-joining algorithms (Saitou and Nei 1987) of the CLUSTAL W phylogenetic package (Thompson et al. 1994) to determine the evolutionary relationships among sequences. Default parameters of CLUSTAL W were used. The results were analyzed using the bootstrap method (1000 replicates) to provide confidence levels for the tree topology (Felsenstein 1985). The construction of trees was done with TREEVIEW (Page 1996).

For the phylogenetic analysis of RTKs, we used only the tyrosine kinase domain of proteins; due to the length variability of the kinase insert region between different proteins, these were eliminated. The large gaps were also eliminated. The tyrosine kinase domain was defined as follows: For vertebrate tyrosine kinases, the sequences used were those defined by the Pfam program; and for *C. elegans* kinases, the domain was identified by SMART and aligned with the alignment existing for the vertebrate sequences in Pfam. This alignment can be found on our Web site (<http://olan.marseille.inserm.fr/u119/home.html>). The tree was rooted using the kinase domain of human ABL1, a cytoplasmic tyrosine kinase.

The chitinase domains of well-characterized chitinases and of chitinase-like proteins from different species of mammals, insects, and worms were used for the phylogenetic analysis of chitinases-like proteins. A plant chitinase was used for rooting the tree.

Expression Analyses

RT reactions were performed from 2 µg of total RNA extracted from mixed-staged worm preparations by lithium chloride precipitation (MacLeod et al. 1981), using random hexanucleotides and SuperScript II Reverse Transcriptase (GIBCO BRL) according to manufacturer's instructions. The equivalent of 200 ng of reverse transcribed RNA and 50 ng of total DNA as control were used for PCR detection of messengers using the following primers for genes encoding RTK: R09D1.12F, 5'-CCCAAGATTAACCTCGATCGATCG-3'; R09D1.12R, 5'-TCCTTCAATGAATCCGAAACG-3'; F59F3.5F, 5'-TACAGCAAAGTGGTCCAGG-3'; F59F3.5R, 5'-CTGTCAATTTGGCTCATCG-3'; T17A3.1F, 5'-CAACGGATTATCGGATTCAG-3'; and T17A3.1R, 5'-TTGAGCTGCATATCC-TTTCCTA-3'.

The following conditions were used for PCR amplification: initial denaturation at 95°C for 5 min, 35 cycles of 30-sec denaturation at 95°C, 30-sec annealing at 55°C (for R09D1.12 and T17A3.1) or 52°C (for F59F3.5), 1-min extension at 72°C, and a final extension at 72°C for 10 min. The fragments were gel-purified, cloned using pGEM-T Easy kit (Promega), and sequenced at Génome Express (Grenoble, France) using an automated sequencer (Applied Biosystems 373).

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NOTE ADDED IN PROOF

The KIN-8 receptor tyrosine kinase has been renamed CAM-1 (Forrester et al. 1999. *Nature* **400**: 881–885).

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