

Research article

Open Access

The *Caenorhabditis* globin gene family reveals extensive nematode-specific radiation and diversification

David Hoogewijs¹, Sasha De Henau¹, Sylvia Dewilde², Luc Moens²,
Marjolein Couvreur³, Gaetan Borgonie³, Serge N Vinogradov⁴, Scott W Roy⁵
and Jacques R Vanfleteren*¹

Address: ¹Department of Biology and Center for Molecular Phylogeny and Evolution, Ghent University, B-9000 Ghent, Belgium, ²Department of Biomedical Sciences, University of Antwerp, B-2610 Antwerp, Belgium, ³Department of Biology, Nematology section, Ghent University, 9000 Ghent, Belgium, ⁴Department of Biochemistry and Molecular Biology, Wayne State University School of Medicine, Detroit, Michigan 48201, USA and ⁵National Center for Biotechnology Information, National Library of Medicine, National Institutes of Health, Bethesda, MD 20814, USA

Email: David Hoogewijs - David.Hoogewijs@UGent.be; Sasha De Henau - Sasha.DeHenau@UGent.be;
Sylvia Dewilde - Sylvia.Dewilde@ua.ac.be; Luc Moens - Luc.Moens@ua.ac.be; Marjolein Couvreur - Marjolein.Couvreur@UGent.be;
Gaetan Borgonie - Gaetan.Borgonie@UGent.be; Serge N Vinogradov - svinogra@med.wayne.edu; Scott W Roy - scottwroy@gmail.com;
Jacques R Vanfleteren* - Jacques.Vanfleteren@UGent.be

* Corresponding author

Published: 9 October 2008

Received: 3 July 2008

BMC Evolutionary Biology 2008, 8:279 doi:10.1186/1471-2148-8-279

Accepted: 9 October 2008

This article is available from: <http://www.biomedcentral.com/1471-2148/8/279>

© 2008 Hoogewijs et al; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Background: Globin isoforms with variant properties and functions have been found in the pseudocoel, body wall and cuticle of various nematode species and even in the eyespots of the insect-parasite *Mermis nigrescens*. In fact, much higher levels of complexity exist, as shown by recent whole genome analysis studies. *In silico* analysis of the genome of *Caenorhabditis elegans* revealed an unexpectedly high number of globin genes featuring a remarkable diversity in gene structure, amino acid sequence and expression profiles.

Results: In the present study we have analyzed whole genomic data from *C. briggsae*, *C. remanei*, *Pristionchus pacificus* and *Brugia malayi* and EST data from several other nematode species to study the evolutionary history of the nematode globin gene family. We find a high level of conservation of the *C. elegans* globin complement, with even distantly related nematodes harboring orthologs to many *Caenorhabditis* globins. Bayesian phylogenetic analysis resolves all nematode globins into two distinct globin classes. Analysis of the globin intron-exon structures suggests extensive loss of ancestral introns and gain of new positions in deep nematode ancestors, and mainly loss in the *Caenorhabditis* lineage. We also show that the *Caenorhabditis* globin genes are expressed in distinct, mostly non-overlapping, sets of cells and that they are all under strong purifying selection.

Conclusion: Our results enable reconstruction of the evolutionary history of the globin gene family in the nematode phylum. A duplication of an ancestral globin gene occurred before the divergence of the Platyhelminthes and the Nematoda and one of the duplicated genes radiated further in the nematode phylum before the split of the Spirurina and Rhabditina and was followed by further radiation in the lineage leading to *Caenorhabditis*. The resulting globin genes were subject to processes of subfunctionalization and diversification leading to cell-specific expression patterns. Strong purifying selection subsequently dampened further evolution and facilitated fixation of the duplicated genes in the genome.

Background

Globins are small globular proteins, usually consisting of about 140–150 amino acids that comprise eight α -helical segments (named A-H), displaying a characteristic 3-over-3 α -helical sandwich structure that encloses an iron-containing heme group. Vertebrate globin genes predominantly contain three exons separated by two introns inserted at highly conserved positions B12.2 (intron located between codon positions 2 and 3 of the 12th amino acid of globin helix B) and G7.0 (intron inserted between the codons for amino acids 6 and 7 of helix G). Organisms can express multiple globin molecules that have variant properties and functions. Vertebrates typically express haemoglobin in red blood cells, myoglobin in muscle, neuroglobin in nervous tissue [1] and cytoglobin in a variety of non-neuronal cells [2]. Invertebrate globins constitute a more heterogeneous group in terms of structure and function. They range from single-domain globins to large, multisubunit, multidomain hemoglobins and can be fused with nonglobin subunits forming chimeric proteins. Besides a conventional O₂ storage and transport function, a wealth of diverse functions has been described for invertebrate globins [3,4]. Nematodes express distinct globin isoforms in the pseudocoel, body wall and cuticle [5]. The emergence of whole genome analysis tools has revealed even higher levels of complexity. *In silico* analysis of the *C. elegans* genome identified 33 putative globin genes in this species using a robust alignment procedure based on conserved structural features of the classical globin fold. These globins feature a wide diversity in gene structure, amino acid sequence and expression profiles. Despite this remarkable variety some of them display significant sequence similarity to vertebrate myoglobin, neuroglobin and cytoglobin [6,7].

The availability of full genomic sequences of two additional *Caenorhabditis* species presents a unique opportunity to explore the evolutionary globin dynamics of these species. In this study we provide a comprehensive evolutionary analysis of the *Caenorhabditis* globin gene family and we document that globins are found in a broad range of other nematode species.

Results and discussion

Occurrence of globins in the Nematoda

We identified globin gene sequences and exon-structures from full genomic sequences of five nematodes: the *Caenorhabditis* species *C. elegans*, *C. briggsae* and *C. remanei* (Rhabditina, Rhabditidae) and the distantly related nematodes *Pristionchus pacificus* (Rhabditina, Diplogasteromorpha) and *Brugia malayi* (Spirurina, Spiruromorpha) (Additional file 1). This allows comparison of nematode globins across three levels – ortholog evolution within the *Caenorhabditis* genus, ortholog evolution across more dis-

tantly related nematodes and evolution of paralogous genes.

Ortholog conservation for *C. elegans* globins varied across levels. We identified clear orthologs of all 33 *C. elegans* genes in *C. briggsae* and *C. remanei*, but reciprocal blast searches with the TBLASTN algorithm [8] and the *C. elegans* globin amino acid sequences identified only 24 and 13 globins in the genomes of *Pristionchus pacificus* and *Brugia malayi*, respectively, reducing the number of orthologs shared by all 5 species to 10.

By screening NEMBASE [9] we found 103 parasitic globins in several other parasitic nematodes with E-values below e^{-05} . Some matched sequences of already known nematode globins from GenBank, while others lacked parts of the A-or H-helix reducing the total number of different NEMBASE EST clusters to 85 (Table 1). Thus far, we can conclude that globins are present in 31 different nematode species representing 4 nematode clades (I, III, IV and V [10,11]). In addition to the species listed in Table 1, partial globin sequences were also detected in *Heterodera schachtii*, *Litomosoides sigmodontis*, *Meloidogyne chitwoodi* and *Strongyloides ratti*.

Two separate globin classes?

Overall, we recovered over 120 new globin-like sequences from exhaustive database searches. Globin domains were extracted, aligned (Additional files 2 and 3) and subjected to phylogenetic analysis. Bayesian inference of the globin domains of all 33 *C. elegans* globins and globins from a representative set of 26 non-Rhabditid, mostly parasitic, species and globins from plants, trematodes and a sea anemone (*Nematostella vectensis*) as outgroup taxa clearly separated two classes of nematode globins. Class I globins comprises *C. elegans* ZK637.13, well known globins from other nematode species and novel related nematode globin isoforms as well as the trematode globins and is supported by 100% posterior probability. Class II globin genes consist of the remaining 32 *Caenorhabditis* globins and novel orthologs thereof, identified in parasitic nematodes (Figure 1). They form a more diverse clade that is resolved from the sea anemone globins with moderate support ($p = 0.78$). This pattern was essentially retained, though not satisfactorily supported, when more globins from distantly related eukaryotes were included in the analysis (results not shown). We included the cnidarian and trematode globins in this analysis because a recent phylogenomic analysis resolved the Cnidaria as a sister taxon to the Bilateria and the Lophotrochozoa (comprising the Platyhelminthes) as a sister taxon to the Ecdysozoa, to which the Nematoda belong [12]. Thus Bayesian inference seems to indicate that a class II ancestral globin evolved before the divergence of the Platyhelminthes and the Nematoda and radiated further in the nematode phy-

Table 1: Parasitic nematode EST clusters identified in NEMBASE

species	NEMBASE ID
<i>Ancylostoma caninum</i>	ACP00369_1
<i>Ancylostoma caninum</i>	ACP01487_2
<i>Ancylostoma caninum</i>	ACP03829_1
<i>Ascaris lumbricoides</i>	ALP00043_1
<i>Ascaris suum</i>	ASP00019_1
<i>Ascaris suum</i>	ASP00780_1
<i>Ascaris suum</i>	ASPI7423_1
<i>Ancylostoma ceylanicum</i>	AYP00272_1
<i>Ancylostoma ceylanicum</i>	AYP00544_1
<i>Ancylostoma ceylanicum</i>	AYP02881_1
<i>Ancylostoma ceylanicum</i>	AYP01816_1
<i>Ancylostoma ceylanicum</i>	AYP03578_1
<i>Ancylostoma ceylanicum</i>	AYP03942_1
<i>Dirofilaria immitis</i>	DIP00455_1
<i>Dirofilaria immitis</i>	DIP00540_1
<i>Globodera rostochiensis</i>	GRP00115_1
<i>Globodera rostochiensis</i>	GRP00048_1
<i>Haemonchus contortus</i>	HCP00202_1
<i>Haemonchus contortus</i>	HCP00202_2
<i>Haemonchus contortus</i>	HCP00202_3
<i>Haemonchus contortus</i>	HCP00202_4
<i>Haemonchus contortus</i>	HCP00202_5
<i>Haemonchus contortus</i>	HCP00208_1
<i>Haemonchus contortus</i>	HCP00208_2
<i>Haemonchus contortus</i>	HCP00333_2
<i>Haemonchus contortus</i>	HCP00759_1
<i>Haemonchus contortus</i>	HCP00759_2
<i>Haemonchus contortus</i>	HCP00759_3
<i>Haemonchus contortus</i>	HCP00770_1
<i>Haemonchus contortus</i>	HCP00770_2
<i>Haemonchus contortus</i>	HCP00770_3
<i>Haemonchus contortus</i>	HCP00786_1
<i>Haemonchus contortus</i>	HCP01314_3
<i>Haemonchus contortus</i>	HCP02815_1
<i>Haemonchus contortus</i>	HCP08501_1
<i>Haemonchus contortus</i>	HCPI3111_1
<i>Heterodera glycines</i>	HGP06223_2
<i>Heterodera glycines</i>	HGP00385_1
<i>Meloidogyne hapla</i>	MHP02564_2
<i>Meloidogyne hapla</i>	MHP04412_1
<i>Meloidogyne incognita</i>	MIP00586_2
<i>Meloidogyne javanica</i>	MJP04640_1
<i>Meloidogyne paranaensis</i>	MPP00429_1
<i>Necator americanus</i>	NAP00041_1
<i>Necator americanus</i>	NAP00088_1
<i>Nippostrongylus brasiliensis</i>	NBP00095_1
<i>Nippostrongylus brasiliensis</i>	NBP00124_1
<i>Nippostrongylus brasiliensis</i>	NBP00197_1
<i>Nippostrongylus brasiliensis</i>	NBP00328_1
<i>Ostertagia ostertagi</i>	OOP00190_1
<i>Ostertagia ostertagi</i>	OOP00190_2
<i>Ostertagia ostertagi</i>	OOP00214_1
<i>Ostertagia ostertagi</i>	OOP03092_2
<i>Ostertagia ostertagi</i>	OOP03348_2
<i>Ostertagia ostertagi</i>	OOP03513_1
<i>Onchocerca volvulus</i>	OVP00634_1
<i>Onchocerca volvulus</i>	OVP06929_1
<i>Onchocerca volvulus</i>	OVP04040_1

lum. Alternative phylogenetic methods, neighbor-joining (MEGA [13]) and maximum likelihood (RAxML [14]), were consistent with Bayesian analysis but statistical support was not significant (data not shown).

The finding that 9 parasite ESTs resolve with class II globins feeds our expectation that many more are to be detected as more nematode genome sequences will become available. The phylogenetic trees represented in Figure 2 and Figure 3 clearly illustrate that the divergence of the globin genes preceded species divergence. Also, molecular and species trees (*Brugia* separates first, followed by *Pristionchus* and next the *Caenorhabditis* cluster) agree very well for most globin genes. Of note are the very short branch lengths from the *Caenorhabditis* ancestor to the three extant species, considering that these species diverged around 100 million year ago [15] (Figure 3). These branch lengths are disproportionately short, relative to the branch lengths generated for *Brugia* and *Pristionchus* and even more so to those representing the divergence of the globin genes. A likely explanation is that the rate of evolution of the globin genes drastically slowed down in *Caenorhabditis*. Alternatively, diversity within *Caenorhabditis* could represent less of nematode history (i.e. a much more recent divergence) than previously appreciated. To distinguish between these possibilities we explored the rate of evolution of several other gene families including astacins, superoxide dismutases, glutathione S-transferases, nicotinic acetylcholine receptors, hedgehog-related proteins and ATP-binding proteins. We assembled a matrix of as many as possible five-way orthologs for the five taxa and analyzed the resulting matrix by Bayesian inference using partitioning of the data (Additional file 4). The ratios of the branch lengths from *Brugia malayi* or *Pristionchus pacificus* to the *Caenorhabditis* internal node and the average branch length of that node to the extant species suggest that the evolutionary rate is quite variable among gene families (Table 2). However, these results also suggest that four (globins, nicotinic acetylcholine receptors, astacins and hedgehog-related proteins) out of the seven multigene families tested evolved more slowly in the *Caenorhabditis* lineage.

Tissue-specific expression

We constructed gene fusions of approx 800 – 3000 bp upstream promoter and enhancer regions for all 33 globin genes to the coding region of GFP. Spatial expression patterns were observed for 32 genes. Although we have not yet unambiguously identified the specific cells expressing these genes it is clear that these globins are expressed in distinct, mostly non-overlapping sets of cells. Most globin genes were expressed in neuronal cells in the head and tail portions of the body, and the nerve cord. A minority of the globin genes were expressed in non-neuronal tissues,

Table 1: Parasitic nematode EST clusters identified in NEMBASE (Continued)

<i>Parastrongyloides trichosuri</i>	PTP03438_1
<i>Radophylus similis</i>	RSP00034_1
<i>Strongyloides stercoralis</i>	SSP00231_1
<i>Strongyloides stercoralis</i>	SSP00309_1
<i>Strongyloides stercoralis</i>	SSP02226_1
<i>Strongyloides stercoralis</i>	SSP04285_1
<i>Strongyloides stercoralis</i>	SSP04654_1
<i>Toxocara canis</i>	TCPO0537_1
<i>Teladorsagia circumcincta</i>	TDP00008_1
<i>Teladorsagia circumcincta</i>	TDP00008_2
<i>Teladorsagia circumcincta</i>	TDP00008_3
<i>Teladorsagia circumcincta</i>	TDP00008_4
<i>Teladorsagia circumcincta</i>	TDP00009_1
<i>Teladorsagia circumcincta</i>	TDP00009_2
<i>Teladorsagia circumcincta</i>	TDP00032_1
<i>Teladorsagia circumcincta</i>	TDP00084_1
<i>Teladorsagia circumcincta</i>	TDP00127_1
<i>Teladorsagia circumcincta</i>	TDP00173_1
<i>Teladorsagia circumcincta</i>	TDP01113_1
<i>Teladorsagia circumcincta</i>	TDP01113_2
<i>Trichuris muris</i>	TMP00180_1
<i>Trichuris muris</i>	TMP01615_1
<i>Trichinella spiralis</i>	TSP03467_1
<i>Trichuris vulpis</i>	TVP00077_1
<i>Trichuris vulpis</i>	TVP00688_1
<i>Xiphinema index</i>	XIP00721_1
<i>Zeldia punctata</i>	ZPP00218_1

Accession numbers correspond to those in NEMBASE.

including body wall and vulval muscle and the pharynx; only one globin gene, C26C6.7 was expressed in muscular tissue exclusively. We never observed GFP expression in the intestine or gonad (Table 3 and Figure 4).

Intron evolution

Analysis of the gene structures for conserved globin domains for the 10 five-way conserved orthologs revealed striking differences across evolutionary levels (Table 4). Consistent with previous findings (e.g. [16]), most intron positions (23/24) were conserved across *Caenorhabditis* species. Orthologous gene structures were more diverged between genera, but majorities of positions were shared across genera for each species (17/24 *Caenorhabditis* positions were also identified in *Pristionchus* and/or *Brugia*, 25/48 *Pristionchus pacificus* insertion positions in *Caenorhabditis* and/or *Brugia*, and 23/30 *B. malayi* positions in *Caenorhabditis* and/or *Pristionchus*). To determine the relative contribution of intron loss and gain to these patterns, we used previously published methods [17,18] to reconstruct evolution (Figure 5). Estimated numbers of gains and losses were similar (22 versus 34), in stark contrast to patterns observed within *Caenorhabditis* [19-22]. These reconstructions suggest very different histories in *Caenorhabditis* (four losses for each gain) and *P. pacificus* (60% more gains than losses).

The most striking pattern is seen at the level of divergence between paralogs. In contrast to the generally conserved intron positions of most eukaryotic globins, nematode globins exhibit a tremendous variability in intron positions (Additional file 1). Interestingly, only one *Caenorhabditis* globin (F21A3.6) displays the typical ancestral intron positions common to vertebrates and other eukaryotes (at positions B12.2 and G7.0), as do the *B. malayi* and *P. pacificus* orthologs. In addition, the *Brugia* ortholog of *C. elegans* globin gene ZK637.13, which has only one intron inserted at E3.2, has B12.2 and G7.0, whereas the *Pristionchus* ortholog contains B12.2 and G7.0 as well as E3.2. Finally, we found no strong evidence for or against any particular model of intron gain – consistent with the apparent age of the introns (most dating to early nematodes), no clear sequence similarity was observed to other genomic sequences (as expected by transposition of existing introns or intron creation by transposable element insertion) or to flanking exonic sequences (as expected by transformation of duplicated genomic sequence into a new intron). The diversity of globin intron positions in nematodes stands in stark contrast to that observed in some other lineages. For instance, in vertebrates, despite a degree of sequence divergence among paralogous globins comparable to that in nematodes, intron positions are generally conserved. Notably, intron-exon structure of nematode globins follows protein divergence; F21A3.6 is both the only gene to retain both ancestral introns, and shows the highest sequence identity to vertebrate globins (specifically, vertebrate myoglobin; data not shown).

In total, then, nematode globin intron-exon structures suggest a large amount of intron change (loss of ancestral introns and gain of new positions) in deep nematode ancestors, followed by differential loss and gain in individual subsequent lineages, with the *Caenorhabditis* lineage showing a pronounced excess of loss over gain. These data deepen the mystery of atypical nematode intron evolution (including high rates of loss and gain, frequent trans-splicing, and atypical intron splicing signals).

Mode of selective pressure

The striking diversity of nematode globins raises the question of the role of natural selection on the evolution of these genes. We determined the ratio of non-synonymous (Ka) to synonymous substitution (Ks) rates for each orthologous *briggsae/elegans* pair (Table 5). The substitution ratio (ω) is expected to be near 1 for genes under neutral selection, greater than 1 for genes under positive selection and smaller than 1 for those under negative selection. All ω -ratios ranged from 0.0070 to 0.1876 with an average of 0.0367, indicating functional constraint. Similar results were found for the orthologous *elegans/remanei* (average of 0.0359) and *briggsae/remanei* (0.0463

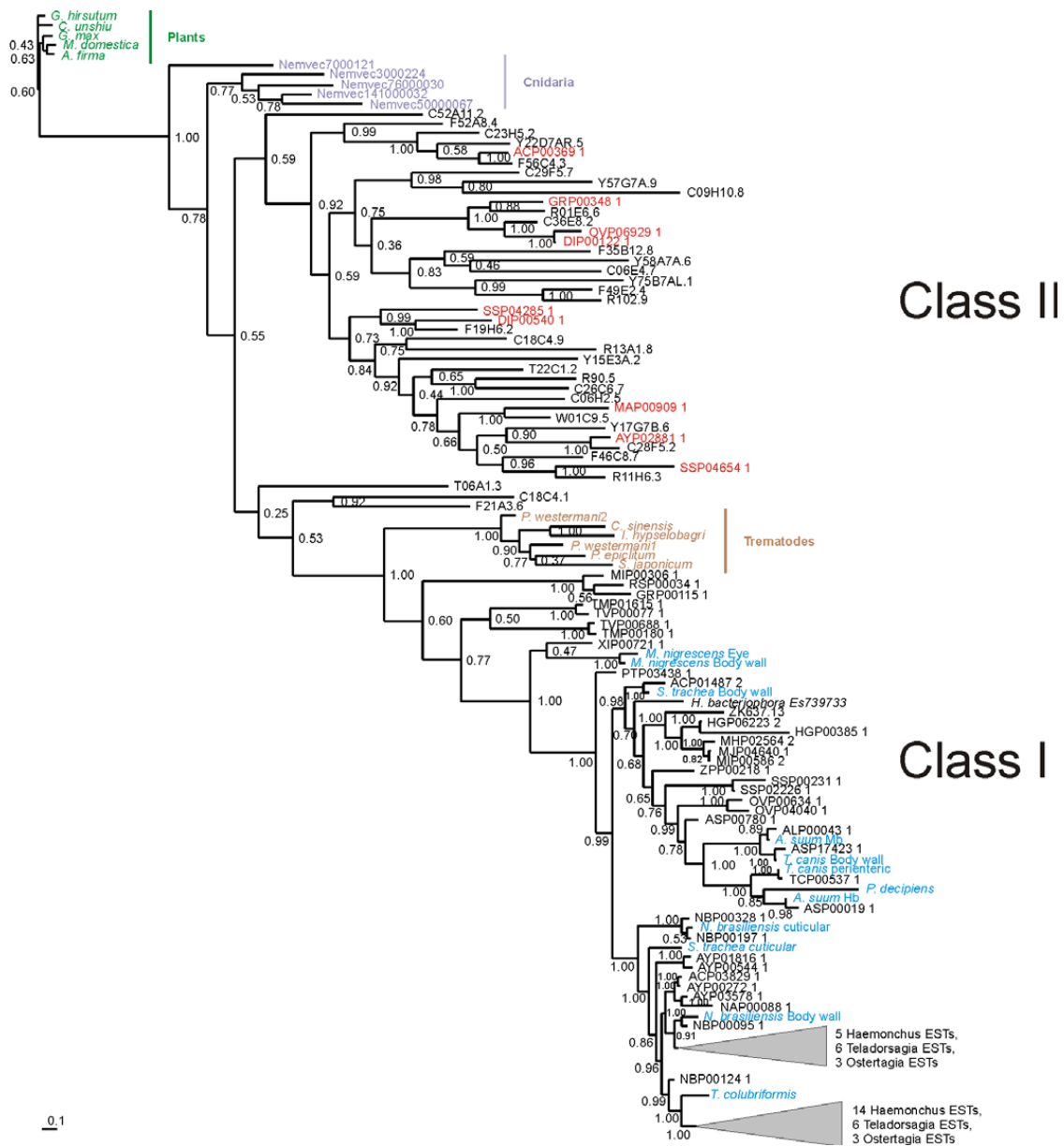


Figure 1
Bayesian phylogenetic tree of globins from *C. elegans* and from 30 other nematode, predominantly parasitic, species. Orthologs from parasitic species in clade II are marked in red. Clade I globins from parasitic nematodes that have been known for long because of their high abundance are marked in blue. The numbers at the nodes represent Bayesian posterior probabilities. All accession numbers from NEMBASE start with the initials of the species followed by P: *Ancylostoma caninum* (AC), *Ancylostoma ceylanicum* (AY), *Ascaris lumbricoides* (AL), *Ascaris suum* (AS), *Dirofilaria immitis* (DI), *Globodera rostochiensis* (GR), *Haemonchus contortus* (HC), *Heterodera glycines* (HG), *Meloidogyne chitwoodi* (MC), *Meloidogyne hapla* (MH), *Meloidogyne incognita* (MI), *Meloidogyne javanica* (MJ), *Necator americanus* (NA), *Nippostrongylus brasiliensis* (NB), *Ostertagia ostertagi* (OO), *Onchocerca volvulus* (OV), *Parastrongyloides trichosuri* (PT), *Strongyloides stercoralis* (SS), *Toxocara canis* (TC), *Teladorsagia circumcincta* (TD), *Trichuris muris* (TM), *Trichinella spiralis* (TS), *Trichuris vulpis* (TV), *Xiphinema index* (XI), *Zeldia punctata* (ZP). Plant, trematode (Platyhelminthes) and sea anemone (Cnidaria) globins were included as outgroup globins. The following plant globins (green) were included: *Gossypium hirsutum* (AAX86687), *Malus domestica* (AAP57676), *Glycine max* (AAA97887), *Alnus firma* (BAE75956), *Citrus unshiu* (AAK07675). The trematode globins (brown) were: *Schistosoma japonicum* (AAP06216), *Paramphistomum epiclitum* (AAG48877), *Paragonimus westermani* (AAX11352 and AAX11353), *Clonorchis sinensis* (AAM18464), *Isoparorchis hypselobagri* (P80722). *Nematostella vectensis* globins (purple): Nemvec141000032, Nemvec3000224, Nemvec50000067, Nemvec7000121 and Nemvec76000030.

Table 2: Evolutionary distance of seven multigene families of *Pristionchus pacificus* and *Brugia malayi* compared to *Caenorhabditis*, inferred from branch length ratios (Additional file 4). *glb* globin; *gst* glutathione S-transferase; *ast* astacin, *sod* superoxide dismutase, *acr* nicotinic acetylcholine receptors; *wrt* hedgehog-related (warthog) proteins; *abc* ATP-binding proteins.

	<i>Glb</i>	<i>gst</i>	<i>sod</i>	<i>ast</i>	<i>wrt</i>	<i>acr</i>	<i>abc</i>
<i>B. malayi</i>	14.10526	7.09375	7.439024	12.15217	10.34615	17.60357	4.163636
<i>P. pacificus</i>	10.86842	5.59375	4.182927	10.65217	11.15385	16.96071	4.272727

on average) pairs. No substantial differences were obtained when this analysis was applied on the globin domains only (results not shown). All *C. elegans* globin genes are expressed [7]. Although pseudogenes have been found in *C. elegans* with ω -ratios down to 0.4 (Ian Hope, personal communication), the combination of these find-

ings refutes the possibility that some of them would be pseudogenes.

We also asked whether the large N- and C-terminal extensions of many *C. elegans* globins are required for their specific function. To this end we searched for positive selection on individual amino acids, because positive selection is thought to act only on specific residues in a protein that is under purifying selection [23]. No positive selection was detected using the maximum likelihood procedure of Thomas et al. [24,25], even in globin Y75B7AL.1, which is a chimeric polypeptide composed of a C-terminal globin domain and an N-terminal domain that has all characteristics of a G-coupled sensor. This domain contains 7 transmembrane helices and this structure is a candidate target for positive selection [25]. The strong purifying selection acting on the globin genes is consistent with the slowing of evolution of the exon-structures in the genus *Caenorhabditis*.

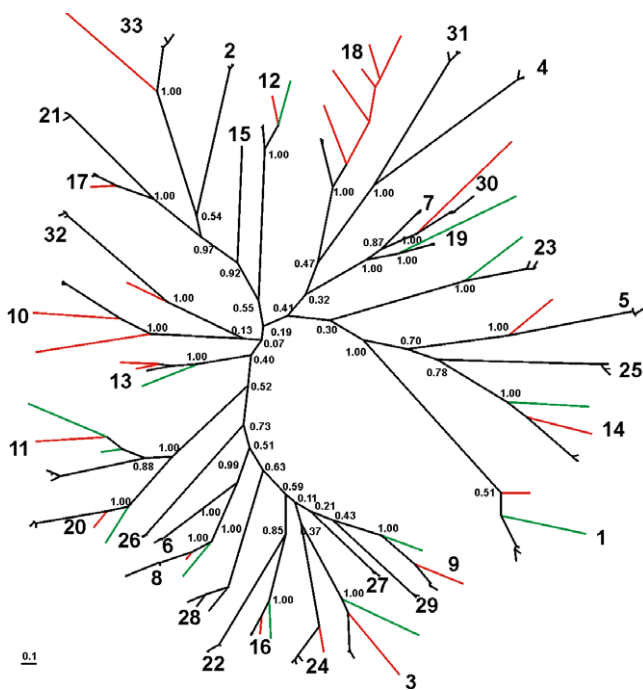


Figure 2
Unrooted bayesian phylogenetic tree of all *C. elegans*, *C. briggsae*, *C. remanei*, *Brugia malayi* (labeled in green) and *Pristionchus pacificus* (labeled in red) globins. The numbers at the nodes represent Bayesian posterior probabilities. The numbers at the branches are consistent with globin nomenclature in WormBase: ZK637.13 (1), C06E4.7 (2), C06H2.5 (3), C09H10.8 (4), C18C4.1 (5), C18C4.9 (6), C23H5.2 (7), C26C6.7 (8), C28F5.2 (9), C29F5.7 (10), C36E8.2 (11), C52A11.2 (12), F19H6.2 (13), F21A3.6 (14), F35B12.8 (15), F46C8.7 (16), F49E2.4 (17), F52A8.4 (18), F56C4.3 (19), R01E6.6 (20), R102.9 (21), R11H6.3 (22), R13A1.8 (23), R90.5 (24), T06A1.3 (25), T22C1.2 (26), W01C9.5 (27), Y15E3A.2 (28), Y17G7B.6 (29), Y22D7AR.5 (30), Y57G7A.9 (31), Y58A7A.6 (32), Y75B7AL.1 (33).

Conclusion

Our study unveils an unexpected complexity of the globin family in nematodes. *Caenorhabditis* species contain a very large number of globin genes, and even distantly related nematode species harbor orthologs to many of them. Our analysis provides some evidence for a number of gene duplication events giving rise to a class of globin genes that is likely unique to the nematode phylum. It remains to be seen whether this class will persist as more genomes will become available.

It is generally accepted that gene duplication played a major role in the evolution of eukaryotic genomes, particularly in the origin of multi-gene families [26]. Duplication events that gave rise to the radiation of nematode globin genes most likely occurred too long ago to leave behind obvious evidence of adaptive evolution. The dispersion of the globin genes over all six chromosomes of *C. elegans* is consistent with this idea. Globins R102.9 and F49E2.4 are exceptional in showing strong sequence conservation (Figure 2) and sharing 2 intron positions (Additional file 1). However, they are located on different chromosomes, arguing against a very recent tandem duplication event. Similarly, both *C. briggsae* orthologs are found on different chromosomes.

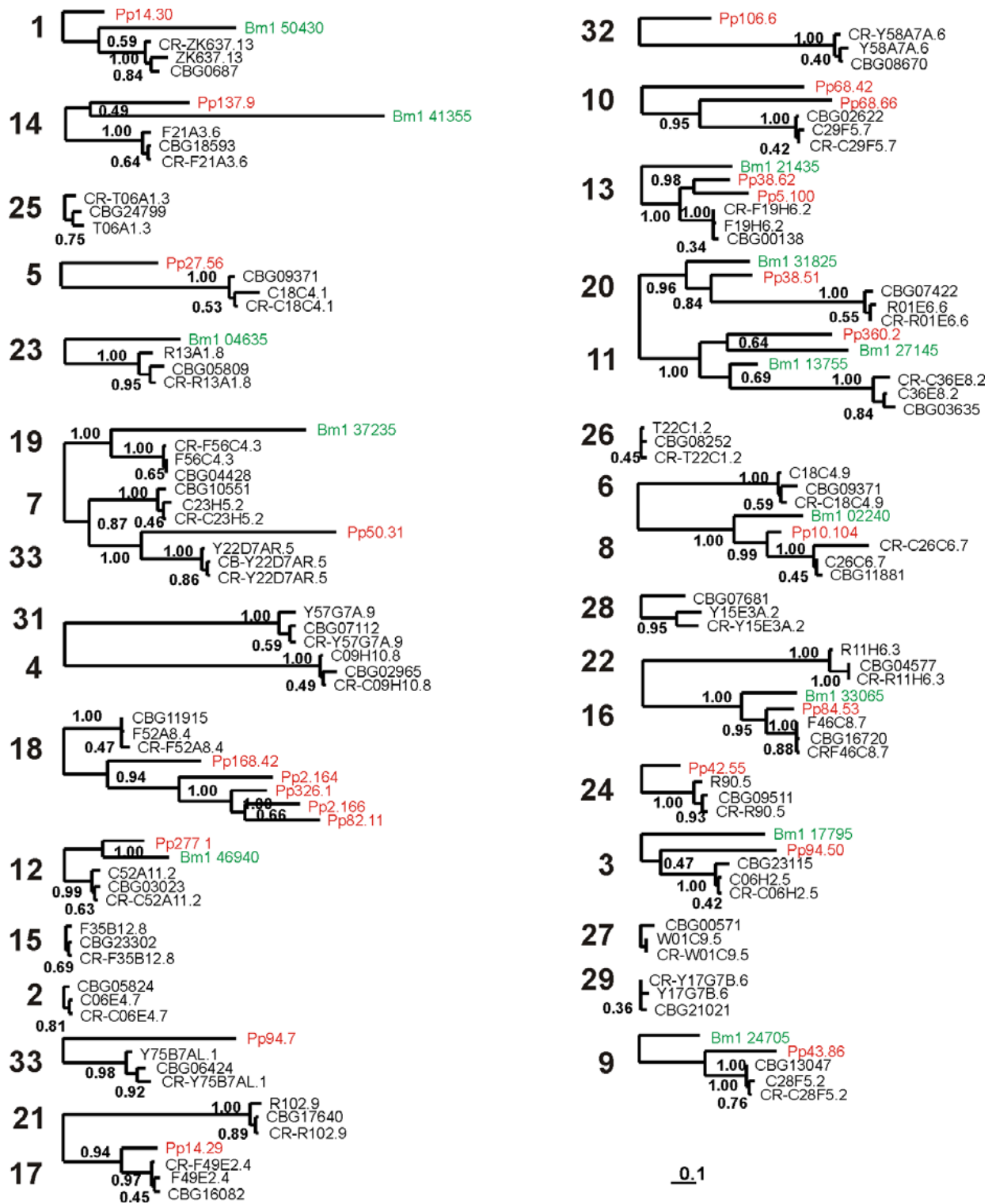


Figure 3
Detailed view of Figure 2, phylogenetic relationship of all orthologous groups of *C. elegans*, *C. briggsae*, *C. remanei*, *Brugia malayi* (labeled in green) and *Pristionchus pacificus* (labeled in red) globins. Due to the complex WashU nomenclature ID, *C. remanei* globins are referred to as *C. elegans* homologue preceded by 'CR-'. The numbers at the nodes represent Bayesian posterior probabilities.

Table 3: Overview of globin expression patterns

Gene	Expression pattern
C06E4.7	Head neurons
C06H2.5	Head and tail neurons, nerve cord
C09H10.8	Head and tail neurons, nerve cord
C18C4.1	Pharynx (Corpus, isthmus), head neurons, nerve cord
C18C4.9	Head and tail neurons, nerve cord
C23H5.2	Head and tail neurons
C26C6.7	Body wall muscle
C28F5.2	Head and tail neurons, nerve cord, vulva neurons
C29F5.7	Head and tail neurons, nerve cord
C36E8.2	Pharynx (Corpus, terminal bulb), head and tail neurons, nerve cord
C52A11.2	Head and tail neurons
F19H6.2	Head neurons, nerve cord, vulva neuron
F21A3.6	Head neurons, vulva neurons, vulval muscle
F35B12.8	No observable expression
F46C8.7	Head neurons, tail neuron
F49E2.4	Head and tail neurons, nerve cord
F52A8.4	Head and tail neurons, nerve cord
F56C4.3	Head neurons and tail neuron
R01E6.6	Stomato-intestinal muscle, depressor muscle, body wall muscle, head neurons, vulva neurons, nerve cord
R102.9	Pharynx (Corpus), head neurons, nerve cord
R11H6.3	Head and tail neurons, nerve cord
R13A1.8	Head neurons, nerve cord
R90.5	Head and tail neurons, nerve cord
T06A1.3	Head and tail neurons, nerve cord
T22C1.2	Head mesodermal cell, stomato-intestinal muscle
W01C9.5	Head and tail neurons, nerve cord
Y15E3A.2	Head and tail neurons, nerve cord
Y17G7B.6	Head and tail neurons, nerve cord
Y22D7AR.5	Head and tail neurons, nerve cord
Y57G7A.9	Head neurons
Y58A7A.6	Head and tail neurons, nerve cord
Y75B7AL.1	Head and tail neurons, nerve cord
ZK637.13	Head muscle/hypodermis, tail muscle/hypodermis, head, tail and vulva neurons, Nerve cord

Why would tiny animals like *Caenorhabditis* species need up to 33 different globins? We have shown that they are all expressed and subject to strong purifying selection. Extensive gene duplication appears to be typical for *Caenorhabditis*. Other surprisingly large gene families include astacins [27], insulins [28], chemoreceptors [20,21] and orphan nuclear receptors HNF4 [29]. Sub-functionalization is frequently invoked to explain the retention of duplicate genes, and differences in gene expression patterns of duplicate genes are generally advanced in support of this hypothesis [30-33]. It is likely that this mechanism was the driving force for the expansion of the *Caenorhabditis* globin family. We have demonstrated that these genes are expressed in distinct subsets of cells and that they are subject to strong purifying selection, in line with this hypothesis. Further support is provided by differential expression of subsets of globin genes in the dauer stage and upon oxygen deprivation [7]. Evidence is mounting that behavioral responses of *C. elegans* to attracting or repelling chemicals including oxygen and CO₂ are generated by gene activities that are deployed in different combinations of neurons [34-37]. Individual

worm neurons seem to have attained very high specialization which is in keeping with their expressing unique or small subsets of globin isoforms.

To date the precise function of none of these globins is known. It seems unlikely that they would all be required for simple oxygen transportation or storage purposes. Globins that are upregulated when oxygen supply is low might serve this function [7]. Other functions are also plausible. It has been shown that *C. elegans* senses molecular oxygen through the heme domain of a guanylate cyclase homolog (GCY-35) and reacts rapidly to changing oxygen levels with aerotaxis responses [37,38]. We expect that some globins may play similar roles in distinct sensory neurons. One likely candidate is globin Y75B7AL.1 which has all characteristics of a G-coupled sensor. Still other functions are worth consideration. Globin T22C1.2 oxidizes instantly to the ferric form in the presence of oxygen and is therefore not capable of reversible oxygen binding (unpublished results). We speculate that this globin may participate in redox reactions with an as yet unidentified reaction partner. Alternatively, peroxidase activity

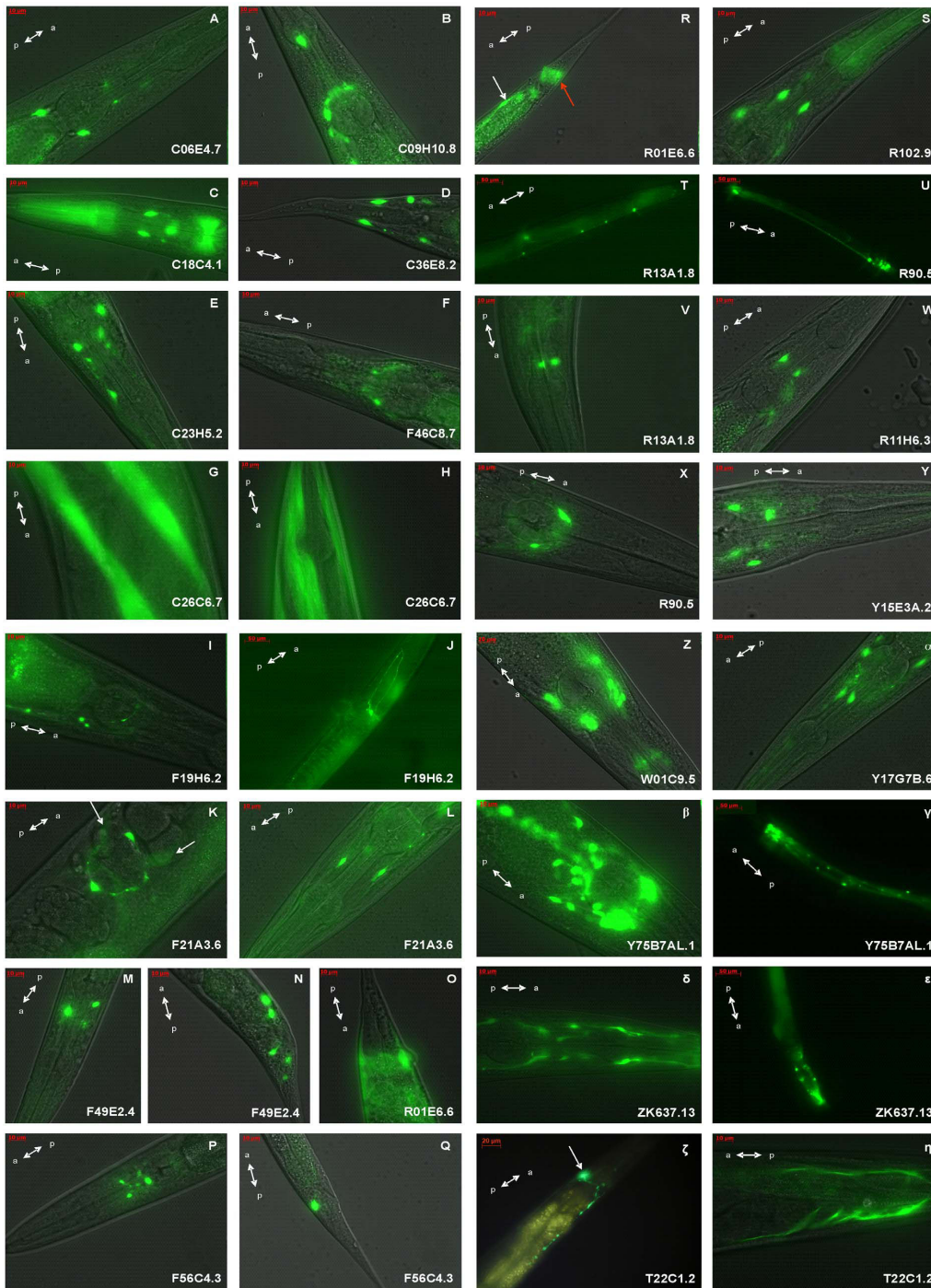


Figure 4

Expression patterns from a selected set of globin genes. A, anterior body part; P, posterior body part. **(A)** C06E4.7, **(B)** C09H10.8, **(C)** C18C4.1, **(D)** C36E8.2, **(E)** C23H5.2, **(F)** F46C8.7, **(G)** C26C6.7, **(H)** C26C6.7, **(I)** F19H6.2, **(J)** F19H6.2, **(K)** F21A3.6 white arrows denote vulval muscle, **(L)** F21A3.6, **(M)** F49E2.4, **(N)** F49E2.4, **(O)** R01E6.6, **(P)** F56C4.3, **(Q)** F56C4.3, **(R)** R01E6.6, white arrow denotes stomato-intestinal muscle, red arrow denotes anal depressor muscle, **(S)** R102.9, **(T)** R13A1.8, **(U)** R90.5, **(V)** R13A1.8, **(W)** R11H6.3, **(X)** R90.5, **(Y)** Y15E3A.2, **(Z)** W01C9.5, **(α)** Y17G7B.6, **(β)** Y75B7AL.1, **(γ)** Y75B7AL.1, **(δ)** ZK637.13, **(ε)** ZK637.13, **(ζ)** T22C1.2, white arrow denotes head mesodermal cell, **(η)** T22C1.2, stomato-intestinal muscle.

Table 4: Overview of intron insertion positions in the globin domain for all five-way conserved orthologs

Globin	<i>C. elegans</i>	<i>C. briggsae</i>	<i>C. remanei</i>	<i>B. malayi</i>	<i>P. pacificus</i>
C06H2.5	A2.2 G15.0	A2.2 G15.0	A2.2 G15.0	A2.2 B11.2 EF11.0 H16.0	A15.0 B11.0 E9.1 EF11.0 G15.0
C26C6.7	C4.0 GH4.2	C4.0 GH4.2	C4.0 GH4.2	E18.0 FGI.0	B1.0 C4.0 E18.0 G2.1 GH4.2
C28F5.2	E13.2 FG5.0	E13.2 FG5.0	E13.2 FG5.0	C7.0 E13.2 FG5.0	AB9.1 B1.0 C7.0 E13.2 F8.2 H1.0
C36E8.2	E6.0 EF14.2 H22.2	E6.0 EF14.2 H22.2	EF14.2 H22.2	E2.0 H8.2	E2.0 EF11.0 H21.1
C52A11.2	E18.0 FG2.2 GH2.0	E18.0 FG2.2 GH2.0	E18.0 FG2.2 GH2.0	A14.0 C7.0 E18.0 GH6.0	A14.0 C7.0 E18.0 F7.0 GH6.0 H10.2
F19H6.2	AB2.0 E10.1 H12.0	AB2.0 E10.1 H12.0	AB2.0 E10.1 H12.0	AB2.0 F3.0 H12.0	A3.2 D6.2 F3.0 G15.0 H12.0
F21A3.6	B12.2 E5.2 G7.0	B12.2 E5.2 G7.0	B12.2 E5.2 G7.0	B12.2 G7.0	B12.2 E17.0 G7.0
F46C8.7	B9.0 F5.1	B9.0 F5.1	B9.0 F5.1	B9.0 E10.0 F5.1 H19.0	A7.0 B9.0 CD4.2 E10.0 F5.1 GH2.1
R01E6.6	B9.0 E16.0 H1.0	B9.0 E16.0 H1.0	B9.0 E16.0 H1.0	C6.2 E16.0 H1.0	B9.0 E16.0 FG2.1 H1.0
ZK637.13	E3.2	E3.2	E3.2	B12.2 EF2.1 G7.0	B12.2 E3.2 EF7.0 G7.0 H15.1

which is a latent but inherent property of globins might have evolved to become the ultimate function of other *C. elegans* globins. Finally, we cannot exclude a potential role as an alternative oxidase during anaerobiosis, when levels of oxygen drop below saturation of cytochrome oxidase, as has been well documented for plant cells [39,40]. Future research shall provide more answers.

Methods

Sequence database searches

The BLAST algorithm [8] was employed to search the sequence databases WormBase (Release WS182), TIGR (*Brugia malayi*) and <http://www.pristionchus.org> (Assembly Freeze 1). NEMBASE 3, containing EST clusters from 37 different partial parasitic nematode genomes [9], was searched for globin motifs (PF00041, PS01033 and SSF46458). Additional EST sequences were searched from EMBL-EBI parasite EST, and the NCBI parasite EST databases using the BLAST algorithm [8] with the *Caenorhabditis* globins as query sequences and cut off E-values of e^{-05} . In cases where the identification of a putative globin was uncertain, searches employing FUGUE [41] were used to determine whether the borderline sequence should be accepted as a globin.

Additional sequences of known nematode globins were obtained from GenBank: *Trichostrongylus colubriformis* (AAA30102), *Nippostrongylus brasiliensis* cuticular globin (P51536), *Nippostrongylus brasiliensis* body wall globin (P51535), *Toxocara canis* body wall globin (AAL56428), *Toxocara canis* perienteric globin (AAL56430), *Ascaris suum* myoglobin (AAA64695), *Ascaris suum* hemoglobin

(AAA29374), *Mermis nigrescens* eye globin (AAF34874), *Mermis nigrescens* body wall globin (AAF35435), *Syngamus trachea* cuticular globin (AAL56426), *Syngamus trachea* body wall globin (AAL56427), *Pseudoterranova decipiens* (P26914). All encoded globin sequences were aligned manually as described previously [42,43].

Globin gene expression analysis

The putative promoter and enhancer sequences upstream of the predicted start codon of each globin gene were extracted from the UCSC Genome Browser database using a repeat masker function [44]. PCR primers were designed to amplify about 3 kb of N2 wild-type genomic DNA, this size was reduced as needed to exclude any upstream gene located at a shorter distance. Promotor-GFP fusion constructs were generated as described by [45]. pRF4 *rol-6*(*su1006*) plasmid and promoter-reporter fusion construct were co-injected at 100 ng/ μ l and 50 ng/ μ l, respectively, into the gonads of young adult hermaphrodites. Rolling F1 worms were transferred to fresh plates and rolling F2 progeny were examined for fluorescence using an Axiovert 200 M (Zeiss) fluorescence microscope. At least 2 independent transgenic lines were examined for each globin gene. These lines generally displayed very similar GFP expression patterns. Primer sequences are available upon request.

Pairwise estimates of K_A and K_S

After manual alignment of orthologous *briggsae/elegans* globin pairs, alignments of corresponding coding sequences were used to calculate the ratios (ω) of non-

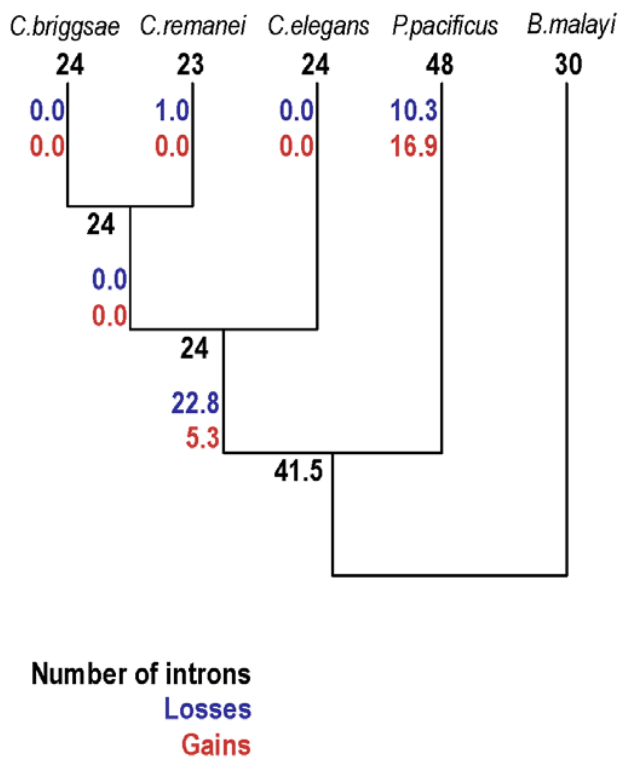


Figure 5
Estimates of intron losses and gains.

synonymous (K_A) to synonymous substitutions (K_S). We used the Yang and Nielsen [46] maximum likelihood method implemented in the YN00 program of the PAML package version 3.15 [47] since the Nei and Gojobori [48] method was not applicable in some cases. Similar results were generated in cases where both methods were applicable.

Phylogenetic analysis

Bayesian inference trees were obtained employing MrBayes version 3.1.2 [49]; four chains were run simultaneously for 4×10^6 generations and trees were sampled every 100 generations generating a total of 40000 trees. The final average standard deviations of split frequencies were stationary and in the range of about 0.016. Posterior probabilities were estimated on the final 30000 trees. The appropriate model of amino acid sequence evolution (JTT model [50]) was selected by ProtTest [51] using the Akaike Information Criterion (AIC).

Authors' contributions

DH and JRV conceived and designed the study; DH and SDH carried out data collection and genomic analysis and generated fusion constructs; DH and MC performed micro-injection experiments; DH and SWR performed data analysis; SD, LM, GB and SNV provided additional input in data analysis; DH, SWR and JRV wrote the man-

Table 5: Overview of K_A/K_S (ω)-ratios for *C. briggsae* – *C. elegans* orthologs

globin	K_A/K_S
C06E4.7	0.0171
C06H2.5	0.0184
C09H10.8	0.0115
C18C4.1	0.0248
C18C4.9	0.0295
C23H5.2	0.0376
C26C6.7	0.0223
C28F5.2	0.0199
C29F5.7	0.0172
C36E8.2	0.0519
C52A11.2	0.0400
F19H6.2	0.0096
F21A3.6	0.0535
F35B12.8	0.0070
F46C8.7	0.0260
F49E2.4	0.0178
F52A8.4	0.0318
F56C4.3	0.0274
R01E6.6	0.0473
R102.9	0.0344
R11H6.3	0.0251
R13A1.8	0.0372
R90.5	0.0292
T06A1.3	0.0381
T22C1.2	0.0506
W01C9.5	0.0153
Y15E3A.2	0.1876
Y17G7B.6	0.0173
Y22D7AR.5	0.0287
Y57G7A.9	0.0274
Y58A7A.6	0.0093
Y75B7AL.1	0.1102
ZK637.13	0.0849

uscript. All authors read and approved the final manuscript.

Additional material

Additional file 1

Overview of intron insertion positions in *Caenorhabditis*, *Brugia malayi* and *Pristionchus pacificus* globins. Identical *Caenorhabditis*-*Pristionchus* intron positions are marked in red, identical *Brugia*-*Pristionchus* intron positions in blue and identical *Brugia*-*Caenorhabditis* intron positions in green. Phase 0 introns separate two consecutive codons (annotated by the number of the amino acid residue, a dot and number 0). Phase 1 and phase 2 introns are inserted following the first or second base of a codon, respectively (annotated by the number of the amino acid, a dot and number 1 or 2, respectively). Introns inserted in the N-terminal extensions are referred to as NA counting from amino acid 1, HC for those inserted in the C-terminal extension starting with the first amino acid after the H-helix.

Click here for file

[<http://www.biomedcentral.com/content/supplementary/1471-2148-8-279-S1.doc>]

Additional file 2*Manual alignment of nematode globins.*

Click here for file

[http://www.biomedcentral.com/content/supplementary/1471-2148-8-279-S2.doc]

Additional file 3*Similarity matrix of all 33 C. elegans globins based on the alignment provided in Additional file 2.*

Click here for file

[http://www.biomedcentral.com/content/supplementary/1471-2148-8-279-S3.xls]

Additional file 4*Unrooted bayesian trees based on matrices of five-way orthologs from C. elegans, C. briggsae, C. remanei, Pristionchus pacificus and Brugia malayi. glb globin; gst glutathione S-transferase; ast astacin, sod superoxide dismutase, acr nicotinic acetylcholine receptors; wrt hedgehog-related (warthog) proteins; abc ATP-binding proteins.*

Click here for file

[http://www.biomedcentral.com/content/supplementary/1471-2148-8-279-S4.jpeg]

Acknowledgements

We thank Yves Van de Peer for critical reading of the manuscript and Guy Haegeman for use of the fluorescence microscope. This work was supported by grants from the Fund for Scientific Research Flanders (G.0331.04). SD is a postdoctoral fellow of the Fund for Scientific Research Flanders (FWO).

References

- Burmester T, Weich B, Reinhardt S, Hankeln T: **A vertebrate globin expressed in the brain.** *Nature* 2000, **407(6803)**:520-523.
- Burmester T, Ebner B, Weich B, Hankeln T: **Cytoglobin: a novel globin type ubiquitously expressed in vertebrate tissues.** *Mol Biol Evol* 2002, **19(4)**:416-421.
- Weber RE, Vinogradov SN: **Nonvertebrate hemoglobins: functions and molecular adaptations.** *Physiol Rev* 2001, **81(2)**:569-628.
- Vinogradov SN, Moens L: **Diversity of globin function: Enzymatic, transport, storage and sensing.** *J Biol Chem* 2008, **283**:8773-8777.
- Blaxter ML: **Nemoglobins: divergent nematode globins.** *Parasitol Today* 1993, **9(10)**:353-360.
- Hoogewijs D, Geuens E, Dewilde S, Moens L, Vierstraete A, Vinogradov S, Vanfleteren J: **Genome-wide analysis of the globin gene family of C. elegans.** *IUBMB Life* 2004, **56(11-12)**:697-702.
- Hoogewijs D, Geuens E, Dewilde S, Vierstraete A, Moens L, Vinogradov S, Vanfleteren JR: **Wide diversity in structure and expression profiles among members of the Caenorhabditis elegans globin protein family.** *BMC Genomics* 2007, **8**:356.
- Altschul SF, Madden TL, Schaffer AA, Zhang J, Zhang Z, Miller W, Lipman DJ: **Gapped BLAST and PSI-BLAST: a new generation of protein database search programs.** *Nucleic Acids Res* 1997, **25(17)**:3389-3402.
- Parkinson J, Whitton C, Schmid R, Thomson M, Blaxter M: **NEMBASE: a resource for parasitic nematode ESTs.** *Nucleic Acids Res* 2004:D427-430.
- Blaxter ML, De Ley P, Garey JR, Liu LX, Scheldeman P, Vierstraete A, Vanfleteren JR, Mackey LY, Dorris M, Frisse LM, Vida JT, Thomas WK: **A molecular evolutionary framework for the phylum Nematoda.** *Nature* 1998, **392(6671)**:71-75.
- Meldal BH, Debenham NJ, De Ley P, De Ley IT, Vanfleteren JR, Vierstraete AR, Bert W, Borgonie G, Moens T, Tyler PA, Austen MC, Blaxter ML, Rogers AD, Lambshhead PJ: **An improved molecular phylogeny of the Nematoda with special emphasis on marine taxa.** *Mol Phylogenet Evol* 2007, **42(3)**:622-636.
- Dunn CW, Hejnal A, Matus DQ, Pang K, Browne WE, Smith SA, Seaver E, Rouse GW, Obst M, Edgecombe GD, Sorensen MV, Hadcock SH, Schmidt-Rhaesa A, Okusu A, Kristensen RM, Wheeler WC, Martindale MQ, Giribet G: **Broad phylogenomic sampling improves resolution of the animal tree of life.** *Nature* 2008, **452(7188)**:745-749.
- Tamura K, Dudley J, Nei M, Kumar S: **MEGA4: Molecular Evolutionary Genetics Analysis (MEGA) software version 4.0.** *Mol Biol Evol* 2007, **24(8)**:1596-1599.
- Stamatakis A, Hoover P, Rougemont J: **A Rapid Bootstrap Algorithm for the RAxML Web-Servers.** *Syst Biol* 2008 in press.
- Stein LD, Bao Z, Blasiar D, Blumenthal T, Brent MR, Chen N, Chinwalla A, Clarke L, Clee C, Coghlan A, Coulson A, D'Eustachio P, Fitch DH, Fulton LA, Fulton RE, Griffiths-Jones S, Harris TW, Hillier LW, Kamath R, Kuwabara PE, Mardis ER, Marra MA, Miner TL, Minx P, Mullikin JC, Plumb RW, Rogers J, Schein JE, Sohrmann M, Spieth J, Stajich JE, Wei C, Willey D, Wilson RK, Durbin R, Waterston RH: **The genome sequence of Caenorhabditis briggsae: a platform for comparative genomics.** *PLoS Biol* 2003, **1(2)**:E45.
- Kent WJ, Zahler AM: **Conservation, regulation, synteny, and introns in a large-scale C. briggsae-C. elegans genomic alignment.** *Genome Res* 2000, **10(8)**:1115-1125.
- Roy SV, Gilbert W: **Rates of intron loss and gain: implications for early eukaryotic evolution.** *Proc Natl Acad Sci USA* 2005, **102(16)**:5773-5778.
- Roy SV, Gilbert W: **Complex early genes.** *Proc Natl Acad Sci USA* 2005, **102(6)**:1986-1991.
- Kiontke K, Gavin NP, Raynes Y, Roehrig C, Piano F, Fitch DH: **Caenorhabditis phylogeny predicts convergence of hermaphroditism and extensive intron loss.** *Proc Natl Acad Sci USA* 2004, **101(24)**:9003-9008.
- Robertson HM: **Two large families of chemoreceptor genes in the nematodes Caenorhabditis elegans and Caenorhabditis briggsae reveal extensive gene duplication, diversification, movement, and intron loss.** *Genome Res* 1998, **8(5)**:449-463.
- Robertson HM: **The large srh family of chemoreceptor genes in Caenorhabditis nematodes reveals processes of genome evolution involving large duplications and deletions and intron gains and losses.** *Genome Res* 2000, **10(2)**:192-203.
- Roy SV, Penny D: **Smoke without fire: most reported cases of intron gain in nematodes instead reflect intron losses.** *Mol Biol Evol* 2006, **23(12)**:2259-2262.
- Nielsen R, Yang Z: **Likelihood models for detecting positively selected amino acid sites and applications to the HIV-1 envelope gene.** *Genetics* 1998, **148(3)**:929-936.
- Thomas JH: **Adaptive evolution in two large families of ubiquitin-ligase adapters in nematodes and plants.** *Genome Res* 2006, **16(8)**:1017-1030.
- Thomas JH, Kelley JL, Robertson HM, Ly K, Swanson WJ: **Adaptive evolution in the SRZ chemoreceptor families of Caenorhabditis elegans and Caenorhabditis briggsae.** *Proc Natl Acad Sci USA* 2005, **102(12)**:4476-4481.
- Lespinet O, Wolf YI, Koonin EV, Aravind L: **The role of lineage-specific gene family expansion in the evolution of eukaryotes.** *Genome Res* 2002, **12(7)**:1048-1059.
- Mohrlen F, Hutter H, Zwilling R: **The astacin protein family in Caenorhabditis elegans.** *Eur J Biochem* 2003, **270(24)**:4909-4920.
- Pierce SB, Costa M, Wisotzkey R, Devadhar S, Homburger SA, Buchman AR, Ferguson KC, Heller J, Platt DM, Pasquelli AA, Liu LX, Doberstein SK, Ruvkun G: **Regulation of DAF-2 receptor signaling by human insulin and ins-1, a member of the unusually large and diverse C. elegans insulin gene family.** *Genes Dev* 2001, **15(6)**:672-686.
- Robinson-Rechavi M, Maina CV, Gissendanner CR, Laudet V, Sluder A: **Explosive lineage-specific expansion of the orphan nuclear receptor HNF4 in nematodes.** *J Mol Evol* 2005, **60(5)**:577-586.
- Cusack BP, Wolfe KH: **When gene marriages don't work out: divorce by subfunctionalization.** *Trends Genet* 2007, **23(6)**:270-272.
- Freilich S, Massingham T, Blanc E, Goldovsky L, Thornton JM: **Relating tissue specialization to the differentiation of expression of singleton and duplicate mouse proteins.** *Genome Biol* 2006, **7(10)**:R89.

32. Fuchs C, Luckhardt A, Gerlach F, Burmester T, Hankeln T: **Duplicated cytoglobin genes in teleost fishes.** *Biochem Biophys Res Commun* 2005, **337(1)**:216-223.
33. Lynch M, Force A: **The probability of duplicate gene preservation by subfunctionalization.** *Genetics* 2000, **154(1)**:459-473.
34. Chang AJ, Bargmann CI: **Hypoxia and the HIF-1 transcriptional pathway reorganize a neuronal circuit for oxygen-dependent behavior in *Caenorhabditis elegans*.** *Proc Natl Acad Sci USA* 2008, **105(20)**:7321-7326.
35. Hallem EA, Sternberg PW: **Acute carbon dioxide avoidance in *Caenorhabditis elegans*.** *Proc Natl Acad Sci USA* 2008, **105(23)**:8038-8043.
36. Bretscher AJ, Busch KE, de Bono M: **A carbon dioxide avoidance behavior is integrated with responses to ambient oxygen and food in *Caenorhabditis elegans*.** *Proc Natl Acad Sci USA* 2008, **105(23)**:8044-8049.
37. Cheung BH, Cohen M, Rogers C, Albayram O, de Bono M: **Experience-dependent modulation of *C. elegans* behavior by ambient oxygen.** *Curr Biol* 2005, **15(10)**:905-917.
38. Gray JM, Karow DS, Lu H, Chang AJ, Chang JS, Ellis RE, Marletta MA, Bargmann CI: **Oxygen sensation and social feeding mediated by a *C. elegans* guanylate cyclase homologue.** *Nature* 2004, **430(6997)**:317-322.
39. Igamberdiev AU, Hill RD: **Nitrate, NO and haemoglobin in plant adaptation to hypoxia: an alternative to classic fermentation pathways.** *J Exp Bot* 2004, **55(408)**:2473-2482.
40. Igamberdiev AU, Hill RD: **Plant Mitochondrial Function During Anaerobiosis.** *Ann Bot (Lond)* 2008.
41. **FUGUE** [<http://tardis.nibio.go.jp/fugue/>]
42. Vinogradov SN, Hoogewijs D, Bailly X, Arredondo-Peter R, Guertin M, Gough J, Dewilde S, Moens L, Vanfleteren JR: **Three globin lineages belonging to two structural classes in genomes from the three kingdoms of life.** *Proc Natl Acad Sci USA* 2005, **102(32)**:11385-11389.
43. Vinogradov SN, Hoogewijs D, Bailly X, Arredondo-Peter R, Gough J, Dewilde S, Moens L, Vanfleteren JR: **A phylogenomic profile of globins.** *BMC Evol Biol* 2006, **6**:31.
44. **UCSC Genome Browser** [<http://genome.ucsc.edu/>]
45. Hobert O: **PCR fusion-based approach to create reporter gene constructs for expression analysis in transgenic *C. elegans*.** *Biotechniques* 2002, **32(4)**:728-730.
46. Yang Z, Nielsen R: **Estimating synonymous and nonsynonymous substitution rates under realistic evolutionary models.** *Mol Biol Evol* 2000, **17(1)**:32-43.
47. Yang Z: **PAML: a program package for phylogenetic analysis by maximum likelihood.** *Comput Appl Biosci* 1997, **13(5)**:555-556.
48. Nei M, Gojobori T: **Simple methods for estimating the numbers of synonymous and nonsynonymous nucleotide substitutions.** *Mol Biol Evol* 1986, **3(5)**:418-426.
49. Huelsenbeck JP, Ronquist F: **MRBAYES: Bayesian inference of phylogenetic trees.** *Bioinformatics* 2001, **17(8)**:754-755.
50. Jones DT, Taylor WR, Thornton JM: **The rapid generation of mutation data matrices from protein sequences.** *Comput Appl Biosci* 1992, **8(3)**:275-282.
51. Abascal F, Zardoya R, Posada D: **ProtTest: selection of best-fit models of protein evolution.** *Bioinformatics* 2005, **21(9)**:2104-2105.

Publish with **BioMed Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours — you keep the copyright

Submit your manuscript here:
http://www.biomedcentral.com/info/publishing_adv.asp

