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Alternatively spliced *Robo2* isoforms in zebrafish and rat

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Abstract *Robo2*, a member of the *robo* gene family, functions as a repulsive axon guidance receptor as well as a regulator of cell migration and tissue morphogenesis in different taxa. In this study, a novel isoform of the zebrafish *robo2* (*robo2_tv2*), which included an otherwise alternatively spliced exon (CAE), has been characterized. *Robo2_tv2* is expressed differentially in most non-neuronal tissues of adult zebrafish whereas *robo2_tv1* expression to a great extent is restricted to the brain and eye. In zebrafish, *robo2_tv2* exhibits a very-low-level basal expression starting from 1 day post fertilization until the mid-larval stages, at which time its expression increases dramatically and could be detected throughout adulthood. Our findings demonstrate that the amino acid sequence coded by CAE of the *robo2* gene is highly conserved between zebrafish and mammals, and also contains conserved motifs shared with *robo1* and *robo4* but not with *robo3*. Furthermore, we provide an account of differential transcription of the CAE homolog in various tissues of the adult rat. These results suggest that the alternatively spliced *robo2* isoforms may exhibit tissue specificity.

Keywords *Danio rerio* · *Rattus norvegicus* · *robo2* · Alternative splicing · Differential expression

Introduction

Robo family genes, which belong to the immunoglobulin superfamily of receptors, play evolutionarily conserved roles in invertebrates and vertebrates with respect to axon guidance and cell migration (Seeger et al. 1993; Yuan et al. 1999; Marillat et al. 2002; Bagri et al. 2002; Long et al. 2004; Sundaresan et al. 2004). Within this family, the vertebrate *robo1*, 2, and 3 are known to guide neuronal cell migration (Wong et al. 2002; Sundaresan et al. 2004) while *robo4* is specifically involved in angiogenesis during embryonic development (Park et al. 2003).

A common characteristic of immunoglobulin superfamily receptors is the frequent alternative splicing events they undergo (Brummendorf and Lemmon 2001). For example, alternative splicing of *robo1* in the mouse has been shown to generate two different isoforms, *robo1* and *dutt1*, which show differential expression patterns. The *dutt1* isoform is widely expressed in development and is also present in adult tissues whereas the *robo1* isoform is expressed strictly in embryonic brain, eye, and kidney tissues (Clark et al. 2002). In addition, *rig1* (*robo3*) has different isoforms, which are structurally and functionally distinct (Jen et al. 2004; Camurri et al. 2005). However, no experimentally confirmed alternative splicing event has yet been reported in the literature for *robo2* or *robo4*.

The zebrafish, a well-established model organism in the developmental genetics field, provides a multitude of opportunities to analyze the extent of alternative splicing events in a stage- and tissue-specific manner (Kimmel et al. 1995; Shen et al. 2002). Zebrafish express all four *robo* paralogs starting from embryogenesis (Lee et al. 2001; Challa et al. 2001; Bedell et al. 2005). Among these, the zebrafish *robo3* gene has been recently shown to produce two distinct variants that function in motor axon pathfinding and dorsoventral cell fate specification, respectively (Challa et al. 2005). Potential alternative splicing events in other zebrafish *robo* members remain unexplored.

In zebrafish, *robo2/astray* is responsible for the proper retinal axon guidance, anterior–posterior pathfinding, midline crossing, and fascicular retinal projection (Fricke

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et al. 2001). The predicted amino acid sequence based on the full-length *robo2* cDNA sequence (NM_131633; Lee et al. 2001) indicates that the *robo2* protein has an extracellular domain comprising five immunoglobulin domains and three fibronectin type III domains, a single transmembrane domain, and a cytoplasmic domain containing several conserved motifs. The neuronal expression of *robo2* mRNA in zebrafish is present at 12 h post-fertilization (hpf) and continues through 72 hpf while the expression of other *robo* paralogs, *robo1* and *robo3*, are diminished by that time (Lee et al. 2001). In addition to neuronal expression, zebrafish *robo* family mRNAs are present in non-neuronal tissues including pectoral fin buds during the first 3 days of development (Lee et al. 2001). However, the larval, juvenile, and adult expression pattern or function of *robo2* in non-neuronal cells of zebrafish is not well known. On the other hand, several studies have reported the expression pattern and functional contribution of *robo2* in non-neuronal tissues in chicken, rodents, and humans (Piper et al. 2000; Nagase et al. 2000; Vargesson et al. 2001; Anselmo et al. 2003; Greenberg et al. 2004; Grieshammer et al. 2004). A comprehensive study focusing on the human homolog of *robo2* (KIAA1568) expression has indicated that *robo2* is expressed in various fetal and adult tissues but is excluded from adult pancreas, heart, liver, and skeletal muscle (Nagase et al. 2000). These findings suggest that *robo2* expression may be temporally and spatially compartmentalized.

Our primary aim was to test the presence of alternatively spliced forms of *robo2* and, if any such forms exist, to characterize their developmental and tissue-specific expression pattern. We also investigated whether these splicing events were conserved between fish and mammals using the rat as a model system. Our findings indicated the presence of a highly conserved alternative exon (herein named CAE) located in the cytoplasmic portion of the *robo2* mRNA. Reverse transcriptase-polymerase chain reaction (RT-PCR) results showed that CAE was more likely to be spliced out in neuronal than in non-neuronal tissues of both organisms. Furthermore, CAE was highly conserved across the paralogs *robo1*, *robo2*, and *robo4* but was not incorporated into the *robo3* gene structure. These findings implicated CAE in the production of multiple *robo2* proteins which may perform different functions in neuronal and non-neuronal tissues.

Materials and methods

Fish husbandry

Zebrafish (*Danio rerio*), purchased from pet-shops or provided by the University of Oregon at Eugene, OR, USA (AB strain), were kept and raised using standard methods. The fish were maintained on a cycle of a 14-h day and a 10-h night at 28.5°C. Before they were killed, the animals were anesthetized with an overdose of tricaine solution

(0.12%). Adult fish were dissected using sterile equipment to obtain different tissues. The embryos were collected in the morning following the setup and maintained in plates filled with system water. Embryos, larvae, and juvenile fish were staged according to hours post-fertilization (hpf), days post-fertilization (dpf), and relevant morphological criteria (Kimmel et al. 1995).

Animals

Nine-week-old, 200 to 250-g male or female Sprague–Dawley rats were used. They were housed under controlled environmental condition (22°C) in the animal holding facility of Bilkent University, Turkey on a 12-h light and 12-h dark cycle. All of the animals received care according to the criteria outlined in the “Guide for Care and Use of Laboratory Animals” prepared by the National Academy of Science and this study protocol complied with Bilkent University’s guidelines on humane care and use of laboratory animals.

Comparison of zebrafish *robo2* sequences in NCBI and Ensembl databases

A complete *robo2* genomic sequence is not currently available. However, the Ensembl database contains a predicted transcript (Ensdart00000014877; <http://www.ensembl.org>; WTSI Zv5; Scaffold791) which partially corresponds to zebrafish *robo2* (NM_131633). Several NCBI zebrafish ESTs, which are aligned primarily to the 5' and 3' ends of the *robo2* cDNA sequence (NM_131633), could also be detected (CK686381.1, AL920855.1, CN173823.1, CN318984.1, and AI437295.1). The exon/intron structure of Ensdart00000014877 in the genomic assembly was analyzed in further detail by comparing it with the corresponding Genscan prediction. The most 3' end of Genscan00000032249 (corresponding to the zebrafish chromosome 15: 40,471,855 to 40,472,040 bp; www.ensembl.org, v37) exhibited a potential alternative exon which was further pursued in the present study (S1a in “Appendix”).

Rat *robo2* sequences in NCBI and Ensembl databases

A full length rat *robo2* mRNA sequence was computationally predicted by NCBI (XM_213677). In the Ensembl database, several computationally predicted isoforms have been reported (S1b in “Appendix”); these differed in their length and their usage of alternative exons including the rat homolog of CAE (Ensmot00000044621, Ensmot00000043725, Ensmot00000042437, Ensmot00000050397, Ensmot00000047577, Ensmot00000046648, Q9QZ13_RAT; Chromosome 11).

Orthologous CAE in genomic assembly databases

The human, rat, chimpanzee, dog, and chicken homologs of the zebrafish *robo2* protein sequences were extracted from public databases (GenBank, <http://www.ncbi.nlm.nih.gov>; Ensembl, <http://www.ensembl.org>). Orthologs of CAE mapped, in general, to Genscan predicted putative exonic sites near the 3' end of *robo2* [human: Chr. 3: 77,764,315:77,764,497 bp, (Genscan00000045332); chimpanzee: Chr. 3: 79,987,335:79,987,517 bp, (Genscan00000073054); dog: Chr. 31: 12416150: 12416332 bp, (Genscan00000000765)] for all species examined, except chicken and rat. For chicken, Ensembl provided Ensgalt00000025020, which included the 3' end of *robo2*. Although this transcript did not contain CAE, its corresponding Genscan sequence [Chr. 1: 90,811,644: 90,811,826 bp, (Genscan00000007903)] incorporated a sequence highly orthologous to zebrafish CAE. In rats, *robo2* CAE was already annotated as transcribed based on computational prediction algorithms, although this was not shown experimentally (S1b in "Appendix"). There has been no EST evidence for CAE for any of the aforementioned organisms including the rat.

Primer design

To detect whether the conserved alternative exon (CAE) was expressed in a developmental and/or tissue-specific manner in zebrafish, primers have been designed against various exons in Ensdart00000014877, NM_131633, and the genomic region corresponding to CAE (Chr. 15: 40,471,855:40,472,040; S1a in "Appendix"; Table 1; Fig. 3). In addition, rat-specific primers were designed to amplify the rat homolog of zebrafish CAE and the flanking sequences in the neighboring exons (S1b in "Appendix"; Table 1; Fig. 6). Degenerate primers targeting the conserved regions of the CAE and the flanking sequences (MAMF–MAMRO primer pair) also were used particularly for sequencing purposes (Table 1).

Total RNA extraction

Dissected fish and rat tissues and embryos were kept in the RNAlater solution (Qiagen, USA, Cat # 7020) or in liquid nitrogen before performing RNA isolations. All material and solutions were treated with diethyl pyrocarbonate to prevent RNase contamination. RNA extraction was performed from zebrafish embryos of various dpfs in an ascending order from embryonic to juvenile stages. RNA

Table 1 *Robo2*, *mipep*, and *cyclophilin* primer sequences and conditions used for RT-PCR for zebrafish and rat cDNAs amplification and sequencing

Name	Sequence (5'-3')
NCBI1F	CTCGGCAGCCTGCAGATCAA
NCBI2F	AAGACTTCCGCGATGTGATG
E21F	CCTTAGTGTCTGACATCGAG
E22R	GGACGAGGTCTATGAGGAA
AER	ATCATCTTCTGAAGCCGAGC
RATF	CCCCTCAGAGCACTAGACCA
RATRI	TCCATCCAGCCTAAAACCAG
RATRO	TCCCTCCGATGAGGTAACAC
MAMF	GAGCACT(G/A)GACCAGAC(T/A)CC
MAMRO	T(G/T)GGTCCATCCG(C/T)CCTC
MIPEPF	ACATCCTCAGAGAATTGCAG
MIPEPR	GGTGCTGGATCTCGATGTT
CYCF	GGGAAGGTGAAAGAAGGCAT
CYCR	GAGAGCAGAGATTACAGGGT

Primer pair	Denaturation	Cycle number	Denaturation	Annealing	Extension	Extension
NCBI1F–E22R	93°C, 2 min	40	93°C, 10 s	56°C, 30 s	72°C, 3 min	72°C, 10 min
NCBI1F–AER	93°C, 2 min	40	93°C, 10 s	55°C, 30 s	72°C, 4 min	72°C, 10 min
NCBI2F–E22R	93°C, 2 min	40	93°C, 10 s	56°C, 30 s	72°C, 3 min	72°C, 10 min
NCBI2F–AER	93°C, 2 min	40	93°C, 10 s	55°C, 30 s	72°C, 4 min	72°C, 10 min
E21F–E22R	95°C, 5 min	30	95°C, 30 s	58°C, 30 s	72°C, 30 s	72°C, 10 min
E21F–AER	95°C, 5 min	30	95°C, 30 s	58°C, 30 s	72°C, 30 s	72°C, 10 min
RATF–RATRI	95°C, 5 min	30	94°C, 30 s	55°C, 30 s	72°C, 30 s	72°C, 10 min
RATRF–RATRO	95°C, 5 min	30	94°C, 30 s	58°C, 30 s	72°C, 30 s	72°C, 10 min
MAMF–MAMRO	95°C, 5 min	35	95°C, 30 s	55°C, 30 s	72°C, 30 s	72°C, 10 min
MIPEPF–MIPEPR	95°C, 5 min	30/35	95°C, 30 s	58°C, 30 s	72°C, 30 s	72°C, 10 min
CYCF–CYCR	95°C, 5 min	25	95°C, 30 s	58°C, 30 s	72°C, 30 s	72°C, 10 min

was extracted based on the instructions of the manufacturer (Qiagen Rneasy Mini Kit, USA, Cat# 74124) from five embryos, one to five larvae, or one juvenile fish per dpf, and from multiple adult tissues of two or more zebrafish (i.e., eye, brain, liver, muscle, heart, caudal fin, pectoral fin, dorsal fin, spleen, gills, digestive tract, and ovaries). Total RNA from different rat tissues (brain, liver, kidney, lung, spleen, heart, skeletal muscle, testes, and ovary) was isolated by using TriPure Isolation Reagent (Roche Diagnostics, USA, Cat # 11667157001) according to the instructions of the manufacturer.

RT-PCR analysis and sequencing of *robo2* products

Equal amounts of total RNA from zebrafish embryo, larvae, juveniles, and various tissues of adult fish and rat were converted into the first strand cDNA according to the manufacturer's protocols (Fermentas, Canada). PCR reactions were performed using the appropriate primer pairs for each cDNA sample (Techne, fitGENE2D, Cambridge, England). Each reaction contained a total of 25- μ l reaction volume that included 1 μ g cDNA, 10 pmol reverse and forward primers, 0.5 ml dNTP (0.2 mM, each), 1.5 mM MgCl₂, 2.5 μ l 10 \times PCR buffer, and 1 unit Taq DNA polymerase (Fermentas, Canada). PCR conditions for each primer pair are indicated in Table 1. Size-matched amplicons with or without CAE from zebrafish and rat (for primer pairs, see Table 1) were sequenced (Iontek Sequencing Facility, Istanbul, Turkey; ABI PRISM 310 Genetic Analyzer, Applied Biosystems, USA) upon gel

extraction and purification (QIAquick Gel Extraction Kit, Cat # 28704).

Results and discussion

Identification and conservation of the putative alternative exon

Transcript and genomic information from the Ensembl database (Ensdart00000014877 and ENSDARG00000014891 in Chr.15: 40,369,682–40,475,421; WTSI Zv5) were used to identify possible alternatively spliced exons not present in the current *robo2* cDNA sequence but predicted by Genscan (see "Materials and methods" for details). We previously had observed that primers targeted to a region spanning 3,955–4,126 bp (E21F–E22R; Table 1) of *robo2* mRNA sequence (NM_131633) resulted in amplification of two different products in zebrafish, suggestive of an alternative splicing event in *robo2*. A 186-bp sequence, herein called CAE that corresponds to a region in chromosome 15 (from 40,471,855 to 40,472,040 bp and in between 21st and 22nd exons of Ensdart00000014877), has accordingly been identified. Distant homology searches for the presence of a CAE homolog in the human, chimpanzee, dog, rat, and chicken genomic sequences suggested that CAE was likely to be expressed, although previously no ESTs with CAE had been reported (see "Materials and methods" for details). Unlike in vertebrates, no homologous sequence to CAE could be identified for invertebrates.

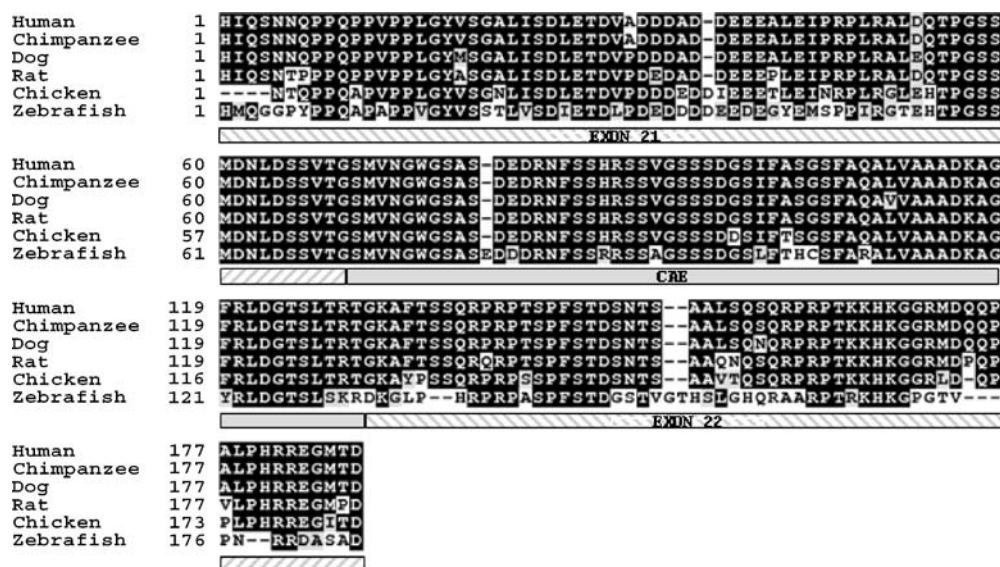


Fig. 1 Multiple alignment of the *robo2* protein sequences for the human, rat, chimpanzee, dog, chicken, and zebrafish was performed using ClustalW 1.8 in BCM Search Launcher (<http://www.searchlauncher.bcm.tmc.edu/>) and then visualized using BOX-SHADE 3.1 (http://www.ch.embnet.org/software/BOX_form.html). The conserved alternative exon has been manually translated and incorporated into the protein sequence when not included in the original NCBI database sequences. The accession numbers and

location of the amino acid used at the first position of the alignment are given for each species in parentheses: human *robo2* (NP_002933; 1,186), rat *robo2* (XP_213677; 1,313), chimpanzee *robo2* (XP_516591; 1,121), dog *robo2* (XP_544815; 1,186), chicken *robo2* (ENSGALP00000024974; 1,073), and zebrafish *robo2* (NP_571708; 1,242). The location of CAE and those of the exons 21 and 22 are depicted using shaded boxes

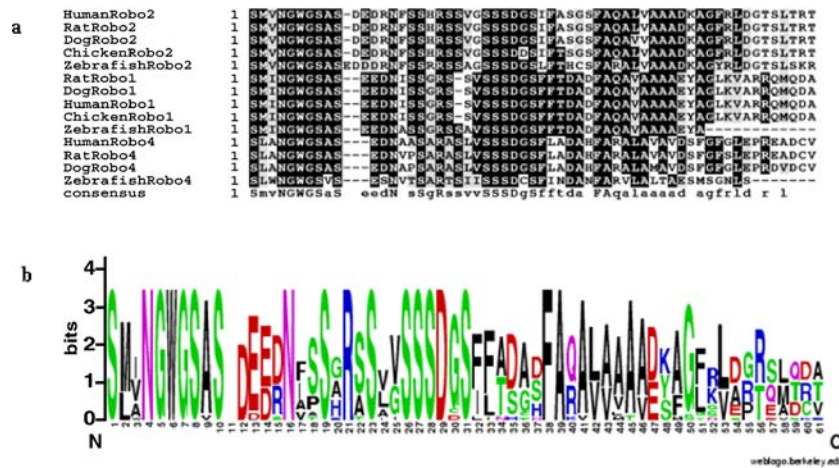


Fig. 2 **a** Multiple alignment of the protein sequences of robo proteins from different species. Human: robo1 (NP_002932, residues 1369–1426), robo4 (NP_061928, residues 872–929); rat: robo1 (NP_071524; 1,369–1,426), robo2 (XP_213677; 1,382–1,441), robo4 (NP_852040; 890–947); dog: robo1 (XP_544814.2; 1,420–1,477), robo4 (XP_546424.2, 912–969); zebrafish: robo1 (NP_571556; 1,388–1,434), robo4 (AAQ10890; 1,014–1,064); and chicken: robo1 (XP_416673; 1,690–1,747). The first position shown

in the alignment corresponds to the initial residue given in the parentheses above. Robo2 CAE sequences are as previously described in Fig. 1. The *last* row of the multiple alignment refers to a consensus sequence. No protein sequence with high similarity to the consensus motif could be identified for chicken robo4 in the genome; thus, the chicken robo4 sequence is not aligned. **b** CAE motif highly conserved among robo paralogs

Multiple alignment of robo2 orthologous sequences indicated that the protein sequence of zebrafish CAE was highly similar to that of human (80%), chimpanzee (80%), rat (80%), dog (78%), and chicken (80%) CAE (Fig. 1), whereas the residues from 5' and 3' neighboring exons were not as highly conserved (5' exon: 62% to human, 62% to chimpanzee, 62% to rat, 63% to dog, and 68% to chicken; 3' exon: 43% to human, 43% to chimpanzee, 33% to rat, 43% to dog, and 41% to chicken). CAE is predicted to yield an in-frame insertion without stop codons (Fig. 1). It corresponds to a region between the second and the third cytoplasmic domains (between the 1,311th and 1,312th residues of NM_131633) of the zebrafish robo2 protein sequence.

The high sequence homology and preserved reading frame between the orthologous sequences of CAE suggest that this sequence is functionally important and conserved. It also is possible that CAE could be incorporated in multiple robo2 isoforms other than the ones specified in this study. Robo2 as a guidance molecule might be crucial for the regulation of cell–cell and cell–extracellular matrix interactions in adult life.

CAE also was conserved between paralogs robo1 and robo4, among different vertebrate taxa (Fig. 2a). Robo1, robo2, and robo4 proteins all possessed a highly conserved motif (Fig. 2b) exhibiting greater sequence similarity within the members of a particular paralog (e.g., a paralog-specific submotif: SMINGWGSAS for robo2, SMINGWGSAS for robo1, and SLABGWGSAS for robo4). On the other hand, the conserved motif shown in Fig. 2b was not present in robo3 in any of the organisms.

One evolutionarily plausible hypothesis to explain the differential incorporation of CAE into the functional protein structure of robo family members could involve the exclusion of CAE in robo3 and tissue-specific inclusion

of CAE in robo2. On the other hand, robo1 and robo4 could contain CAE either constitutively or alternatively, although no EST without CAE has been reported yet. It is possible that the incorporation of the CAE into robo1 and robo4 protein structure might be crucial for their function, whereas its exclusion could be dispensable or only condition-specifically advantageous, thus leading to functional divergence in other robo paralogs. The alternative usage of CAE in robo paralogs accordingly warrants future functional studies.

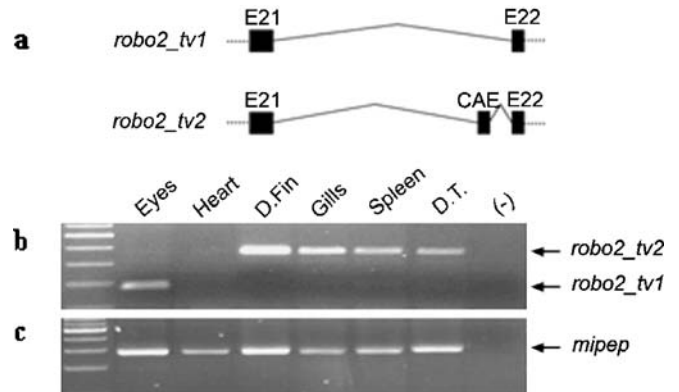


Fig. 3 Genomic representation and differential expression of robo2 alternative isoforms in zebrafish adult tissues. **a** Genomic representation of *robo2* isoforms with or without the CAE. **b** RT-PCR results with primers E21F–E22R (Table 1) on various tissues of zebrafish adults are shown. The expected size of the amplicon is 172 bp when the alternative exon, CAE, is spliced out while amplicon size increases to 358 bp when CAE is included. The *top* band of the 100 bp DNA Ladder Plus corresponds to 600 bp. **c** The RT-PCR of the same tissues also was performed for amplification of *mipep* as a house-keeping gene (expected size, 318 bp) using MIPEP F-R primers

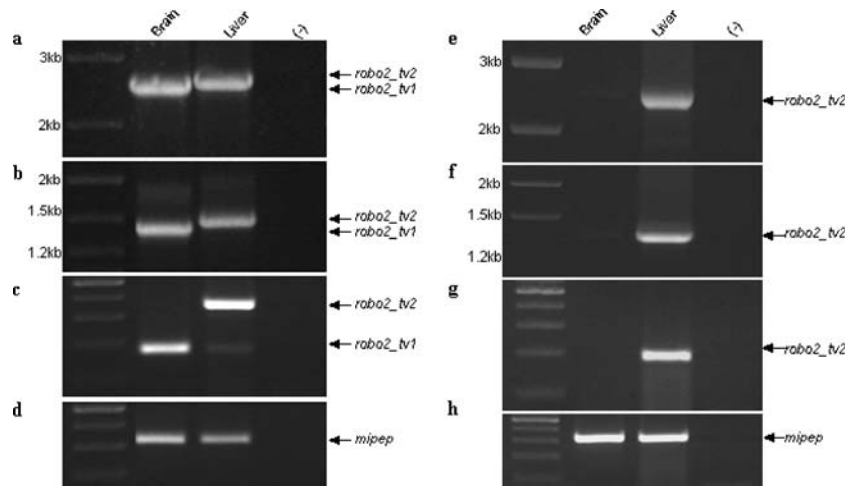


Fig. 4 The expression of *robo2_tv1* and *robo2_tv2* isoforms on zebrafish brain and liver tissues. **a** Amplification with forward primer NCBI1F (NM_131633; 1,593–1,613) and the reverse primer E22R. Expected sizes of *robo2_tv1* and *robo2_tv2* are 2,534 and 2,720 bp in brain and liver, respectively. The *first lane* corresponds to DNA Ladder Plus. **b** Amplification with NCBI2F (NM_131633; 2,692–2,712) and the reverse primer E22R. Expected sizes of *robo2_tv1* and *robo2_tv2* are 1,435 and 1,621 bp in brain and liver, respectively. The *first lane* corresponds to DNA Ladder Plus. **c** Amplification with the forward primer E21F (NM_131633; 3,955–3,974) and the reverse primer E22R (NM_131633; 4,108–4,126). Expected sizes of *robo2_tv1* and *robo2_tv2* are 172 and 358 bp in brain and liver, respectively. The *first lane* corresponds to DNA Ladder Plus; maximum size shown is 500 bp. **d, h** RT-PCR of the

same tissues performed using MIPEP F-R primers (Table 1). The *first lane* corresponds to 100 bp DNA Ladder; maximum size shown is 500 bp. **e** Amplification with forward primer NCBI1F (NM_131633; 1,593–1,613) and the reverse primer AER (Table 1). Expected sizes of *robo2_tv2* is 2,547 bp in liver. The *first lane* corresponds to DNA Ladder Plus. **f** Amplification with forward primer NCBI2F (NM_131633; 2,692–2,712) and the reverse primer AER. Expected size of *robo2_tv2* is 1,448 bp in liver. The *first lane* corresponds to DNA Ladder Plus. **g** Amplification with the forward primer E21F (NM_131633; 3,955–3,974) and the reverse primer AER. Expected size of *robo2_tv2* is 185 bp in liver. The *first lane* corresponds to DNA Ladder Plus; maximum size shown is 500 bp

Expression pattern of *robo2* isoforms in adult zebrafish and embryonic/larval development

The expression of the transcripts of potential *robo2* isoforms encompassing CAE was investigated in a series of zebrafish adult tissues including the eye, heart, spleen,

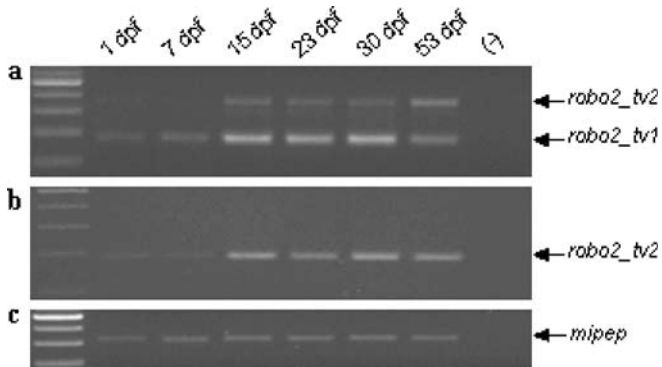


Fig. 5 The expression of *robo2* isoforms during zebrafish development. **a** RT-PCR results with the primer pair E21F–E22R (expected amplicon sizes 172 and 358, with and without CAE, respectively) performed on cDNAs from various stages: 1, 7, 15, 23, 30, and 53 dpf. The *first lane* corresponds to 100-bp DNA Ladder Plus (maximum size shown, 600 bp). **b** Amplification of RT-PCR results performed on the same set of cDNAs using the primer pair E21F–AER, AER reverse primer specifically targeting a region within CAE. A 100-bp DNA Ladder Plus (maximum size shown, 500 bp) was used. **c** RT-PCR of the same tissues was also performed with MIPEP F-R primers. The *first lane* corresponds to 100-bp DNA Ladder Plus (maximum size shown, 500 bp)

digestive tract, gills, and dorsal fin together with that of a housekeeping gene *mipep* (mitochondrial intermediate peptidase) (Fig. 3). Our findings demonstrated that CAE was either skipped or included depending on the tissue type; this produced two alternative forms herein called *robo2_tv1* and *robo2_tv2*, respectively (Fig. 3b). In many of the non-neuronal tissues, *robo2_tv2* was expressed at detectable levels while *robo2_tv1* was predominantly expressed in the eye. Furthermore, the expression of these isoforms was either very low or absent in zebrafish heart (Fig. 3b).

As *robo2_tv1* and *robo2_tv2* seem to be differentially expressed in neuronal and non-neuronal tissues of zebrafish, brain and liver were used as representative tissues for subsequent experiments. The differential expression pattern of *robo2_tv1* and *robo2_tv2* in liver and brain was confirmed in RT-PCR reactions performed with three different 5' forward primers, two of which targeted the extracellular portion of *robo2* protein and one targeted the exon 21 preceding the CAE (Table 1; Fig. 4). Our findings indicated that CAE was highly expressed in liver while its expression was absent or negligible in brain (Fig. 4e–g), whereas *robo2_tv1* was predominant in brain (Fig. 4a–c). Sequencing of both the *robo2_tv1*- and the *robo2_tv2*-specific PCR products obtained from zebrafish brain and liver tissue cDNAs confirmed the alternative pre-mRNA splicing of CAE in zebrafish (DQ481484, DQ481485). The presence of alternative splicing events concerning exons other than CAE has not been tested in this study; thus, it cannot be excluded. In general, *robo2_tv2* expression was

predominantly present in multiple non-neuronal tissues whereas zebrafish *robo2_tv1* isoform could be characterized as neurally enriched although not necessarily neurally restricted.

Considering that the relative amount of *robo2_tv1* and *robo2_tv2* expression varied among adult tissues, the possibility of differential expression during embryogenesis and at larval–juvenile stages was also assessed (Fig. 5). Our results indicated that *robo2_tv2* expression exhibited a dramatic increase starting from the mid-larval stages while it was hardly detectable during embryogenesis or early larval stages (Fig. 5a,b). *Robo2_tv1* expression, present at higher levels during embryonic and early larval stages, also displayed an induction later in development (Fig. 5a).

The two isoforms, *robo2_tv1* and *robo2_tv2*, identified in the present study differ in their use of CAE, which is located in the intracellular part of the protein. The extracellular Ig domains allow the binding of slit leucine-rich regions, whereas the intracellular portion of *robo2* relays the signal (Chen et al. 2001; Battye et al. 2001; Nguyen Ba-Charvet et al. 2001; Liu et al. 2004). It is interesting that while the extracellular domain of *robo2* is evolutionarily conserved, the intracellular domain has been shown to be divergent (Lee et al. 2001). The high-level expression of the *robo2_tv2* and the scanty expression of the *robo2_tv1* in non-neuronal tissues suggest the possible compartmentalization of the cell migratory and axon guidance roles in the developing animal. Further studies are needed to assess whether *robo2* isoforms that contain CAE could have a role beyond regulating the cytoskeleton; they perhaps act as transcriptional factors (Couch et al. 2004) regulating the extent and timing of the molecular events leading to tissue morphogenesis.

Expression pattern of *robo2* isoforms in the adult rat

Rat *robo2* mRNA (XM_213677) was reported to include the homolog of the zebrafish CAE, although some of the Ensembl-predicted isoforms differed in their usage of it (Ensnrnot00000044621, Ensnrnot00000043725, Ensnrnot00000042437, Ensnrnot00000050397). To determine whether *robo2* is also differentially and alternatively expressed in rats, RT-PCR from several rat tissues, including brain, skeletal muscle, heart, liver, spleen, kidney, lung, testes, and ovary was performed with rat-specific primers, RATF and RATRO (Fig. 6; Table 1). Our results demonstrated that, although the rat homolog of CAE (Fig. 6a) was transcribed in most of the tissues examined, exon skipping was observed to a greater degree in brain and testes (Fig. 6b). On the other hand, the CAE exclusion and inclusion events were comparable to each other in the ovary (Fig. 6b). In addition, nested PCRs on these PCR products performed with the alternative exon-specific primer pair, RATF and RATRI (Table 1), yielded the expected product (Fig. 6c). These findings support the predictions reported in Ensembl in regards to differential incorporation of the rat CAE in *robo2* transcripts (DQ481486, DQ481487).

In rats, the expression of the *robo2_tv1* isoform without CAE was found predominantly in the brain similar to that of zebrafish. Several distinguishing features between zebrafish and rats were also obvious with regards to the transcription of CAE. The expression of the CAE seemed to be more ubiquitous in rats and, thus, was found in both the neuronal and non-neuronal tissues unlike in zebrafish. Moreover, zebrafish heart contained very little of either isoforms, if any, whereas both were detectable in rats. The expression levels of *robo2_tv1* in brain and testis were relatively high compared to other tissues; this expression profile was in accord with the findings of a recent study on alternative splicing capacities of human tissues (Yeo et al. 2004). It has been reported that the brain and testis have the highest levels of exon-skipping events while the ovary is one of the tissues likely to have multiple isoforms simultaneously.

In summary, an alternative splicing event in a highly conserved predicted exon (CAE) in *robo2* genes of zebrafish and rat was identified and experimentally verified (Figs. 1 and 6). CAE of zebrafish was transcribed in most of the non-neuronal tissues while it was either absent or expressed at a very low level in the brain, eye, and heart. Furthermore, the isoform-specific expression of *robo2* appeared to be tightly controlled during development such that zebrafish larvae exhibited an expressional induction of *robo2_tv1* and *robo2_tv2* at around the mid-larval stages. These findings might implicate *robo2_tv2* in a role in tissue morphogenesis, during which cell growth/migration is

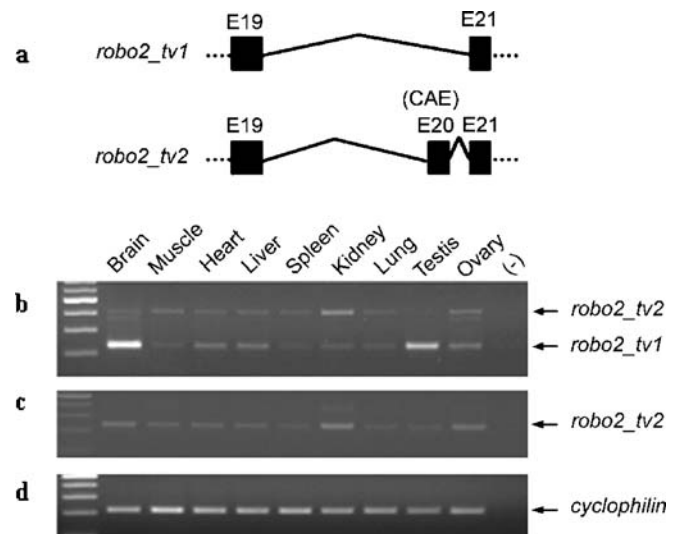


Fig. 6 Genomic representation and differential expression of *robo2* alternative isoforms in rat. **a** Genomic representation of *robo2* isoforms **b** RT-PCR result with primers RATF–RATRO, where RATF targets E19 and RATRO targets E21, on various rat tissues is shown. Expected sizes of the rat *robo2_tv1* and *robo2_tv2* are 224 bp and 407 bp, respectively. The first lane corresponds to 100 bp DNA Ladder Plus (maximum size shown, 600 bp). **c** Nested PCR result performed with primers RATF–RATRI, where RATRI targets CAE (i.e., E20) in **a**, using the reactions from RATF–RATRO PCR of **b** (expected band size, 228 bp). A 100-bp DNA Ladder Plus (maximum size shown, 500 bp) was used. **d** RT-PCR of the same samples performed with *cyclophilin* F–R primers. A 100-bp DNA Ladder Plus (maximum size shown, 500 bp) was used

temporally and spatially regulated. There is unearched evidence indicating that tissue-type-dependent and, in some cases, neuronal non-neuronal tissue-distinct expression profiles result from alternative splicing events (Fukuda 2003; Shen et al. 2002; Jin et al. 2002; Rahman et al. 2002; Ramming et al. 2000; Hu et al. 1999). Some robo family members have previously been shown to exhibit alternatively spliced isoforms leading to functional divergence within Robo paralogs (Clark et al. 2002; Challa et al. 2005; Camurri et al. 2005). The differential expression of robo2 isoforms in neuronal and non-neuronal tissues could similarly become a source of variation in the function of robo2, possibly directed by different regulatory processes. Our findings strongly point to the existence of multiple robo2 proteins; thus, it is critical to identify which form is involved in a specific developmental process or event in pathogenesis (Latil et al. 2003).

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